Radiobiology
FOR THE
Radiologist

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This book, like so many before it, grew out of a set of lecture notes. The lectures were given during the autumn months of 1969, 1970, and 1971 at the Columbia-Presbyterian Medical Center, New York City. The audience consisted primarily of radiology residents from Columbia, affiliated schools and hospitals, and various other institutions in and around the city.

To plan a course in radiobiology involves a choice between, on the one hand, dealing at length and in depth with those few areas of the subject in which one has personal expertise as an experimenter or, on the other hand, surveying the whole field of interest to the radiologist, necessarily in less depth. The former course is very much simpler for the lecturer and in many ways more satisfying; it is, however, of very little use to the aspiring radiologist who, if this course is followed, learns too much about too little and fails to get an overall picture of radiobiology. Consequently, I opted in the original lectures, and now in this book, to cover the whole field of radiobiology as it pertains to radiology. I have endeavored to avoid becoming evangelical over those areas of the subject which interest me, those to which I have devoted a great deal of my life. At the same time I have attempted to cover, with as much enthusiasm as I could muster and from as much knowledge as I could glean, those areas in which I had no particular expertise or personal experience.

This book, then, was conceived and written for the radiologist—specifically, the radiologist who, motivated ideally by an inquiring mind or more realistically by the need to pass an examination, elects to study the biological foundations of radiology. It may incidentally serve also as a text for graduate students in the life sciences or even as a review of radiobiology for active researchers whose viewpoint has been restricted to their own area of interest. If the book serves these functions, too, the author is doubly happy, but first and foremost, it is intended as a didactic text for the student of radiology.

Radiology is not a homogenous discipline. The diagnostician and therapist have divergent interests; indeed, it sometimes seems that they come together only when history and convenience dictate that they share a common course in physics or radiobiology. The bulk of this book will be of concern, and hopefully of interest, to all radiologists. The diagnostic radiologist is recommended particularly to Chapters 11, 12, and 13 concerning radiation accidents, late effects, and the irradiation of the embryo and fetus. A few chapters, particularly Chapters 8, 9, 15, and 16, are so specifically oriented towards radiotherapy that the diagnostician may omit them without loss of continuity.

A word concerning reference material is in order. The ideas contained in this book represent, in the author's estimate, the consensus of opinion as expressed in the scientific literature. For ease of reading, the text has not been broken up with a large number of direct references. Instead, a selection of general references has been included at the end of each chapter for the reader who wishes to pursue the subject further.

I wish to record the lasting debt that I owe my former colleagues at Oxford and my present colleagues at Columbia, for it is in the daily cut and thrust of debate and discussion that ideas are formulated and views tested.

Finally, I would like to thank the young men and women who have regularly attended my classes. Their inquiring minds have forced me to study hard and reflect carefully before facing them in a lecture room. As each group of students has grown in maturity and understanding, I have experienced a teacher's satisfaction and joy in the belief that their growth was due in some small measure to my efforts.

E. J. H.
New York
July 1972
The seventh edition is the most radical revision of this textbook to date and now includes color figures, a visual transformation over the sixth edition. However, we were careful to retain the same format as the sixth edition, which divided the book into two parts. Part I contains 17 chapters and represents both a general introduction to radiation biology and a complete self-contained course in the subject, suitable for residents in diagnostic radiology and nuclear medicine. It follows the format of the Syllabus in Radiation Biology prepared by the Radiological Society of North America (RSNA), and its content reflects the questions appearing in recent years in the written examination for diagnostic radiology residents given by the American Board of Radiology. Part II consists of 11 chapters of more in-depth material designed primarily for residents in radiation oncology.

We live in an exciting time, but yet a dangerous time as well. The threat of nuclear terror rears its head way too often. If such an event occurs, those trained in the radiation sciences will be called on to manage exposed individuals. For this reason, we have included a new chapter on Radiologic Terrorism (Chapter 14).

The translation of molecular imaging into the clinic is moving at a rapid pace. Therefore, we also included a chapter on fundamental concepts in molecular imaging that involves ionizing radiation such as CAT scans and PET imaging to reflect these new advances and describe the underlying biologic principles for each of these technologies (Chapter 15).

The subject of retreatment with radiotherapy is not covered in most textbooks, and, because of this void, we have dedicated a new chapter to this subject (Chapter 24). Recent clinical trials have demonstrated a clinical benefit with hyperthermia, fulfilling the promise of this form of therapy. These recent clinical studies have been included in a thorough revision of this chapter (Chapter 28). Most of the other chapters in this edition have also been revised and updated to reflect current thoughts and ideas.

With the addition of the new chapters, some of the old chapters from the sixth edition have been eliminated. For some time, we considered omitting the chapters on gene therapy and predictive assays because these areas have yet to justify their early promise. In the end, they did not make the cut for the seventh edition. A separate chapter on “Molecular Techniques in Radiobiology” has also been eliminated because we felt that we could incorporate the fundamentals of these molecular techniques into the description of the data where applicable throughout the book, and the relevance of this chapter to the diagnostic or therapeutic radiologist is questionable.

The ideas contained in this book represent, we believe, the consensus of opinion as expressed in the scientific literature. We have followed the precedent of previous editions, in that, the pages of text are unencumbered with flyspeck-like numerals referring to footnotes or original publications, which are often too detailed to be of much interest to the general reader. On the other hand, there is an extensive and comprehensive bibliography at the end of each chapter for those readers who wish to pursue the subject further.

We commend this new edition to residents in radiology, nuclear medicine, and radiation oncology, for whom it was conceived and written. If it serves also as a text for graduate students in the life sciences or even as a review of basic science for active researchers or senior radiation oncologists, the authors will be doubly happy.

Eric J. Hall
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October 2010
We would like to thank the many friends and colleagues who generously and willingly gave permission for diagrams and illustrations from their published work to be reproduced in this book.

Although the ultimate responsibility for the content of this book must be ours, we acknowledge with gratitude the help of several friends who read chapters relating to their own areas of expertise and made invaluable suggestions and additions. With each successive edition, this list grows longer and now includes Drs. Ged Adams, Philip Alderson, Sally Amundson, Joel Bedford, Roger Berry, Max Boone, Victor Bond, David Brenner, J. Martin Brown, Ed Bump, Denise Chan, Julie Choi, James Cox, Nicholas Denko, Bill Dewey, Mark Dewhirst, Frank Ellis, Peter Esser, Stan Field, Greg Freyer, Charles Geard, Eugene Gerner, Julian Gibbs, George Hahn, Simon Hall, Ester Hammond, Tom Hei, Robert Kallman, Richard Kolesnick, Adam Krieg, Dennis Leeper, Howard Lieberman, Philip Lorio, Edmund Malaise, Gillies McKenna, Mortimer Mendelsohn, George Merriam, Noelle Metting, Jim Mitchell, Anthony Nias, Ray Oliver, Stanley Order, Tej Pandita, Marianne Powell, Simon Powell, Julian Preston, Elaine Ron, Harold Rossi, Robert Rugh, Chang Song, Fiona Stewart, Robert Sutherland, Roy Tishler, Len Tolmach, Liz Travis, Lou Wagner, John Ward, Barry Winston, Rod Withers, and Basil Worgul. Of particular note are Dr. Ted Graves, who was instrumental in the development of Chapter 14 on molecular imaging, and Dr. Elizabeth Repasky, who revised Chapter 28 on hyperthermia. Without their help, this volume would be much the poorer.

The principal credit for this book must go to the successive classes of residents in radiology, radiation oncology, and nuclear medicine that we have taught over the years at Columbia and Stanford, as well as at ASTRO and RSNA refresher courses. Their perceptive minds and searching questions have kept us on our toes. Their impatience to learn what was needed of radiobiology and to get on with being doctors has continually prompted us to summarize and get to the point.

We are deeply indebted to the U.S. Department of Energy, the National Cancer Institute, and the National Aeronautical and Space Administration, which have generously supported our work and, indeed, much of the research performed by numerous investigators that is described in this book.

We owe an enormous debt of gratitude to Ms. Sharon Clarke, who not only typed and formatted the chapter revisions, but also played a major role in editing and proofreading. Our publisher, Ryan Shaw, guided our efforts at every stage and arranged for many of the figures to be updated.

Finally, we thank our wives, Bernice Hall and Jeanne Giaccia, who have been most patient and have given us every encouragement with this work.
SECTION I

For Students of Diagnostic Radiology, Nuclear Medicine, and Radiation Oncology
In 1895, the German physicist Wilhelm Conrad Röntgen discovered “a new kind of ray,” emitted by a gas discharge tube, that could blacken photographic film contained in light-tight containers. He called these rays “x-rays” in his first announcement in December 1895—the x representing the unknown. In demonstrating the properties of x-rays at a public lecture, Röntgen asked Rudolf Albert von Kölliker, a prominent Swiss professor of anatomy, to put his hand in the beam and so produced the first publicly taken radiograph (Fig. 1.1).

The first medical use of x-rays was reported in the *Lancet* of January 23, 1896. In this report, x-rays were used to locate a piece of a knife in the backbone of a drunken sailor, who was paralyzed until the fragment was removed following its location. The new technology spread rapidly through Europe and the United States, and the field of diagnostic radiology was born. There is some debate about who first used x-rays therapeutically, but by 1896, Leopold Freund, an Austrian surgeon, demonstrated before the Vienna Medical Society the disappearance of a hairy mole following treatment with x-rays. Antoine Henri Becquerel discovered radioactivity emitted by uranium compounds in 1896, and 2 years later, Pierre and Marie Curie isolated the radioactive elements polonium and radium. Within a few years, radium was used for the treatment of cancer.

The first recorded biologic effect of radiation was due to Becquerel, who inadvertently left a radium container in his vest pocket. He subsequently described the skin erythema that appeared 2 weeks later and the ulceration that developed and that required several weeks to heal. It is said that Pierre Curie repeated this experience in 1901 by deliberately producing a radium “burn” on his own forearm (Fig. 1.2). From these early beginnings, at the turn of the century, the study of radiobiology began.

**Radiobiology** is the study of the action of ionizing radiations on living things. As such, it inevitably involves a certain amount of radiation physics. The purpose of this chapter is to present, in summary form and with a minimum of mathematics, a listing of the various types of ionizing radiations and a description of the physics and chemistry of the processes by which radiation is absorbed.

### TYPES OF IONIZING RADIATIONS

The absorption of energy from radiation in biologic material may lead to *excitation* or to *ionization*. The raising of an electron in an atom or molecule to a higher energy level without actual ejection of the electron is called *excitation*. If the radiation has sufficient energy to eject one or more orbital electrons from the atom or molecule, the process is called *ionization*, and that radiation is said to be *ionizing radiation*. The important characteristic of ionizing radiation is the localized release of large amounts of energy. The energy dissipated per ionizing event is about 33 eV, which is more than enough to break a strong chemical bond; for example, the energy associated with a C=\(\text{C}\) bond is 4.9 eV. For convenience, it is usual to classify ionizing radiations as either *electromagnetic* or *particulate*.

**Electromagnetic Radiations**

Most experiments with biologic systems have involved x- or \(\gamma\)-rays, two forms of electromagnetic radiation. X- and \(\gamma\)-rays do not differ in...
electrical device that accelerates electrons to high energy and then stops them abruptly in a target usually made of tungsten or gold. Part of the kinetic energy (the energy of motion) of the electrons is converted to x-rays. On the other hand, γ-rays are emitted by radioactive isotopes; they represent excess energy that is given off as the unstable nucleus breaks up and decays in its efforts to reach a stable form. Natural background radiation from rocks in the earth also includes γ-rays. Everything that is stated about x-rays in this chapter applies equally well to γ-rays.

X-rays may be considered from two different standpoints. First, they may be thought of as waves of electrical and magnetic energy. The magnetic and electrical fields, in planes at right angles to each other, vary with time, so that the wave moves forward in much the same way as ripples move over the surface of a pond if a stone is dropped into the water. The wave moves with a velocity, \( c \), which in a vacuum has a value of \( 3 \times 10^8 \) cm/s. The distance between successive peaks of the wave, \( \lambda \), is known as the wavelength.

The number of waves passing a fixed point per second is the frequency, \( \nu \). The product of frequency times wavelength gives the velocity of the wave; that is, \( \lambda \nu = c \).

A helpful, if trivial, analogy is to liken the wavelength to the length of a person’s stride when walking; the number of strides per minute is the frequency. The product of the length of stride multiplied by the number of strides per minute gives the speed or velocity of the walker.
Like x-rays, radio waves, radar, radiant heat, and visible light are forms of electromagnetic radiation. They all have the same velocity, $c$, but they have different wavelengths and, therefore, different frequencies. To extend the previous analogy, different radiations may be likened to a group of people, some are tall, some are short, all walking together at the same speed. The tall walkers take long measured strides but make few strides per minute; to keep up, the short walkers compensate for the shortness of their strides by increasing the frequencies of their strides. A radio wave may have a distance between successive peaks (i.e., wavelength) of 300 m; for a wave of visible light, the corresponding distance is about 500 thousandths of a centimeter ($5 \times 10^{-5}$ cm). The wavelength for x-rays may be 100 millionths of a centimeter ($10^{-8}$ cm). X- and γ-rays, then, occupy the short-wavelength end of the electromagnetic spectrum (Fig. 1.3).

Second, x-rays may be thought of as streams of photons, or “packets” of energy. Each energy packet contains an amount of energy equal to $hv$, where $h$ is known as Planck’s constant and $v$ is the frequency. If a radiation has a long wavelength, it has a small frequency, and so, the energy per photon is small. Conversely, if a given radiation has a short wavelength, the frequency is large and the energy per photon is large. There is a simple numeric relationship between the photon energy (in kiloelectron volts*) and the wavelength (in angstroms†):

$$\lambda \text{Å} = 12.4/E(\text{keV})$$

For example, x-rays with wavelengths of 0.1 Å correspond to a photon energy of 124 keV.

The concept of x-rays being composed of photons is very important in radiobiology. If x-rays are absorbed in living material, energy is deposited in the tissues and cells. This energy is deposited unevenly in discrete packets. The energy in a beam of x-rays is quantized into large individual packets, each of which is big enough to break a chemical bond and initiate the chain of events that culminates in a biologic change. The critical difference between nonionizing and ionizing radiations is the size of the individual packets of energy, not the total energy involved. A simple calculation illustrates this point. It is shown in Chapter 8 that a total body dose of about 4 Gy‡ of x-rays given to a human is lethal in about half of the individuals exposed. This dose represents absorption of energy of only about 67 cal, assuming the person to be a “standard man” weighing 70 kg. The smallness of the amount of energy involved can be illustrated in many ways. Converted to heat, it would represent a temperature rise of 0.002° C, which would do no harm at all; the same amount of energy in the form of heat is absorbed in drinking one sip of warm coffee. Alternatively, the energy inherent in a lethal dose of x-rays may be compared with mechanical energy or work. It would correspond to the work done in lifting a person about 16 in. from the ground (Fig. 1.4).

Energy in the form of heat or mechanical energy is absorbed uniformly and evenly, and much greater quantities of energy in these forms are required to produce damage in living things. The potency of x-rays, then, is a function not

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*The kiloelectron volt (keV) is a unit of energy. It is the energy possessed by an electron that has been accelerated through 1,000 volts (V). It corresponds to $1.6 \times 10^{-19}$ ergs.

†The angstrom (Å) is a unit of length equal to $10^{-8}$ cm.

‡Quantity of radiation is expressed in röntgen, rad, or gray. The röntgen (R) is the unit of exposure and is related to the ability of x-rays to ionize air. The rad is the unit of absorbed dose and corresponds to energy absorption of 100 erg/g. In the case of x- and γ-rays, an exposure of 1 R results in an absorbed dose in water or soft tissue roughly equal to 1 rad. The International Commission on Radiological Units and Measurements (ICRU) recommended that the rad be replaced as a unit by the gray (Gy), which corresponds to an energy absorption of 1 J/kg. Consequently, 1 Gy = 100 rad.
Section I · For Students of Diagnostic Radiology, Nuclear Medicine, and Radiation Oncology

Electrons are small, negatively charged particles that can be accelerated to high energy to a speed close to that of light by means of an electrical device, such as a betatron or linear accelerator. They are widely used for cancer therapy.

Protons are positively charged particles and are relatively massive, having a mass almost 2,000 times greater than that of an electron. Because of their mass, they require more complex and more expensive equipment, such as a cyclotron, to accelerate them to useful energies, but they are increasingly used for cancer treatment in specialized centers because of their favorable dose distribution (see Chapter 25).

In nature, the earth is showered with protons from the sun, which represent a component of natural background radiation. We are protected on earth to a large extent by the earth's atmosphere. In addition, the earth behaves like a giant magnet so that charged particles from solar events on the sun are deflected away from the equator by the earth's magnetic field; most miss the earth altogether while others are funneled into the polar regions. This is the basis of the “aurora borealis,” or northern lights caused by intense showers of charged particles that spiral down the lines of magnetic field into the poles, ionizing the air as they do so (see Chapter 16). Protons are a major hazard to astronauts on long-duration space missions.

α-particles are nuclei of helium atoms and consist of two protons and two neutrons in close association. They have a net positive charge and, therefore, can be accelerated in large electrical devices similar to those used for protons.

α-particles are also emitted during the decay of heavy, naturally occurring radionuclides, such as uranium and radium (Fig. 1.5). α-particles are the major source of natural background radiation to the general public. Radon gas seeps out of the soil and builds up inside houses, where, together with its decay products, it is breathed in and irradiates the lining of the lung. It is estimated that 10,000 to 20,000 cases of lung cancer are caused each year by this means in the United States, mostly in smokers.

Neutrons are particles with a mass similar to that of protons, but they carry no electrical charge. Because they are electrically neutral, they cannot be accelerated in an electrical device.
absorbed in the material through which they pass, they give up their energy to produce fast-moving charged particles that in turn are able to produce damage.

The process by which x-ray photons are absorbed depends on the energy of the photons concerned and the chemical composition of the absorbing material. At high energies, characteristic of a cobalt-60 unit or a linear accelerator used for radiotherapy, the Compton process dominates. In this process, the photon interacts with what is usually referred to as a “free” electron, an electron whose binding energy is negligibly small compared with the photon energy. Part of the energy of the photon is given to the electron as kinetic energy; the photon, with whatever energy remains, continues on its way, deflected from its original path (Fig. 1.6). In place of the incident

**FIGURE 1.5** Illustration of the decay of a heavy radionuclide by the emission of an \( \alpha \)-particle. An \( \alpha \)-particle is a helium nucleus consisting of two protons and two neutrons. The emission of an \( \alpha \)-particle decreases the atomic number by 2 and the mass number by 4. Note that radium has changed to another chemical element, radon, as a consequence of the decay.

They are produced if a charged particle is accelerated to high energy and then made to impinge on a suitable target material. Neutrons are also emitted as a by-product if heavy radioactive atoms undergo fission; that is, a split to form two smaller atoms. Consequently, neutrons are present in large quantities in nuclear reactors and are emitted by some artificial heavy radionuclides. They are also an important component of space radiation and contribute significantly to the exposure of passengers and crews of high-flying jetliners.

Heavy charged particles are nuclei of elements, such as carbon, neon, argon, or even iron, that are positively charged because some or all of the planetary electrons have been stripped from them. To be useful for radiation therapy, they must be accelerated to energies of thousands of millions of volts and, therefore, can be produced in only a few specialized facilities, though the number of such centers is increasing.

Charged particles of enormous energy are encountered in space and represent a major hazard to astronauts on long missions, such as the proposed trip to Mars. During the lunar missions of the 1970s, astronauts “saw” light flashes while their eyes were closed in complete darkness, which turned out to be caused by high-energy iron ions crossing the retina.

**ABSORPTION OF X-RAYS**

Radiation may be classified as directly or indirectly ionizing. All of the charged particles previously discussed are directly ionizing; that is, provided the individual particles have sufficient kinetic energy, they can disrupt the atomic structure of the absorber through which they pass directly and produce chemical and biologic changes. Electromagnetic radiations (x- and y-rays) are indirectly ionizing. They do not produce chemical and biologic damage themselves, but when they are

**FIGURE 1.6** Absorption of an x-ray photon by the Compton process. The photon interacts with a loosely bound planetary electron of an atom of the absorbing material. Part of the photon energy is given to the electron as kinetic energy. The photon, deflected from its original direction, proceeds with longer wavelength (i.e., with reduced energy).
the mass absorption coefficient for photoelectric absorption varies rapidly with atomic number \(Z^*\) and is, in fact, about proportional to \(Z^3\).

For diagnostic radiology, photons are used in the energy range in which photoelectric absorption is as important as the Compton process. Because the mass absorption coefficient varies critically with \(Z\), the x-rays are absorbed to a greater extent by the bone because the bone contains elements with high atomic numbers, such as calcium. This differential absorption in

\*\(Z\), the atomic number, is defined as the number of positive charges on the nucleus; it is, therefore, the number of protons in the nucleus.
of electrons, spins are paired; that is, for every electron spinning clockwise, there is another one spinning counterclockwise. This state is associated with a high degree of chemical stability. In an atom or molecule with an odd number of electrons, there is one electron in the outer orbit for which there is no other electron with an opposing spin; this is an unpaired electron. This state is associated with a high degree of chemical reactivity.

For simplicity, we consider what happens if radiation interacts with a water molecule, because 80% of a cell is composed of water. As a result of the interaction with a photon of x- or γ-rays or a charged particle, such as an electron or proton, the water molecule may become ionized. This may be expressed as

\[ \text{H}_2\text{O} \rightarrow \text{H}_2\text{O}^+ + e^- \]

Materials of high Z is one reason for the familiar appearance of the radiograph. For radiotherapy, however, high-energy photons in the megavoltage range are preferred because the Compton process is overwhelmingly important. As a consequence, the absorbed dose is about the same in soft tissue, muscle, and bone, so that differential absorption in bone, which posed a problem in the early days when lower energy photons were used for therapy, is avoided.

Although the differences among the various absorption processes are of practical importance in radiology, the consequences for radiobiology are minimal. Whether the absorption process is the photoelectric or the Compton process, much of the energy of the absorbed photon is converted to the kinetic energy of a fast electron.

**DIRECT AND INDIRECT ACTION OF RADIATION**

The biologic effects of radiation result principally from damage to deoxyribonucleic acid (DNA), which is the critical target, as described in Chapter 2.

If any form of radiation—x- or γ-rays, charged or uncharged particles—is absorbed in biologic material, there is a possibility that it will interact directly with the critical targets in the cells. The atoms of the target itself may be ionized or excited, thus initiating the chain of events that leads to a biologic change. This is called direct action of radiation (Fig. 1.8); it is the dominant process if radiations with high linear energy transfer (LET), such as neutrons or α-particles, are considered.

Alternatively, the radiation may interact with other atoms or molecules in the cell (particularly water) to produce free radicals that are able to diffuse far enough to reach and damage the critical targets. This is called indirect action of radiation.* A free radical is an atom or molecule carrying an unpaired orbital electron in the outer shell. An orbital electron not only revolves around the nucleus of an atom but also spins around its own axis. The spin may be clockwise or counterclockwise. In an atom or molecule with an even number of electrons, spins are paired; that is, for every electron spinning clockwise, there is another one spinning counterclockwise. This state is associated with a high degree of chemical stability. In an atom or molecule with an odd number of electrons, there is one electron in the outer orbit for which there is no other electron with an opposing spin; this is an unpaired electron. This state is associated with a high degree of chemical reactivity.

For simplicity, we consider what happens if radiation interacts with a water molecule, because 80% of a cell is composed of water. As a result of the interaction with a photon of x- or γ-rays or a charged particle, such as an electron or proton, the water molecule may become ionized. This may be expressed as

*It is important to avoid confusion between directly and indirectly ionizing radiation, on the one hand, and the direct and indirect actions of radiation on the other.

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*Fig. 1.8: Direct and indirect actions of radiation. The structure of DNA is shown schematically. In direct action, a secondary electron resulting from absorption of an x-ray photon interacts with the DNA to produce an effect. In indirect action, the secondary electron interacts with, for example, a water molecule to produce a hydroxyl radical (OH·), which in turn produces the damage to the DNA. The DNA helix has a diameter of about 20 Å (2 nm). It is estimated that free radicals produced in a cylinder with a diameter double that of the DNA helix can affect the DNA. Indirect action is dominant for sparsely ionizing radiation, such as x-rays. S, sugar; P, phosphorus; A, adenine; T, thymine; G, guanine; C, cytosine."
H$_2$O$^+$ is an ion radical. An ion is an atom or molecule that is electrically charged because it has lost an electron. A free radical contains an unpaired electron in the outer shell, making it highly reactive. H$_2$O$^+$ is charged and has an unpaired electron; consequently, it is both an ion and a free radical. The primary ion radicals have an extremely short lifetime, on the order of $10^{-10}$ second. They decay to form free radicals, which are not charged but still have an unpaired electron. In the case of water, the ion radical reacts with another water molecule to form the highly reactive hydroxyl radical (OH·):

$$H_2O^+ + H_2O \rightarrow H_3O^+ + OH.$$  

The hydroxyl radical possesses nine electrons; therefore, one of them is unpaired. It is a highly reactive free radical and can diffuse a short distance to reach a critical target in a cell. For example, it is thought that free radicals can diffuse to DNA from within a cylinder with a diameter about twice that of the DNA double helix. It is estimated that about two thirds of the x-ray damage to DNA in mammalian cells is caused by the hydroxyl radical. The best evidence for this estimate comes from experiments using free radical scavengers, which can reduce the biologic effect of sparsely ionizing radiations, such as x-rays, by a factor of close to 3. This is discussed further in Chapter 9. Indirect action is illustrated in Figure 1.8. This component of radiation damage is most easily modified by chemical means—either protectors or sensitizers—unlike direct action.

For the indirect action of x-rays, the chain of events, from the absorption of the incident photon to the final observed biologic change, may be described as follows:

1. Incident x-ray photon
2. Fast electron ($e^-$)
3. Ion radical
4. Free radical
5. Chemical changes from the breakage of bonds
6. Biologic effects

There are vast differences in the time scale involved in these various events. The physics of the process, the initial ionization, may take only $10^{-15}$ second. The primary radicals produced by the ejection of an electron generally have a lifetime of $10^{-10}$ second. The OH· radical has a lifetime of about $10^{-9}$ second in cells, and the DNA radicals formed either by direct ionization or by reaction with OH· radicals have a lifetime of perhaps $10^{-5}$ second (in the presence of air). The period between the breakage of chemical bonds and the expression of the biologic effect may be hours, days, months, years, or generations, depending on the consequences involved. If cell killing is the result, the biologic effect may be expressed hours to days later when the damaged cell attempts to divide. If the radiation damage is oncogenic, its expression as an overt cancer may be delayed for 40 years. If the damage is a mutation in a germ cell leading to hereditary changes, it may not be expressed for many generations.

### ABSORPTION OF NEUTRONS, PROTONS, AND HEAVY IONS

In contrast to x-rays, neutrons interact not with the planetary elections, but with the nuclei of the atoms that make up the tissue (Fig 1.9) resulting in recoil protons, or in the case of higher energy neutrons “spallation products” (i.e., a high-energy neutron may hit a carbon atom which then breaks up into three α-particles), or may hit an oxygen atom to produce four α-particles.

Protons interact with both planetary electrons to ionize the atoms with which they interact and produce fast recoil electrons, and protons also interact with the nuclei of atoms in the tissue to produce heavier secondary particles. Nuclear disintegration becomes more and more likely to happen as the proton energy increases.
For heavy particles, as for x-rays, the biologic effect may be a consequence of the direct or indirect action, but there is a shift in the balance between the two modes of action. For x-rays, indirect action is dominant whereas for the neutrons or heavy ions, the direct action assumes greater importance, increasingly so as the density of ionization increases; that is, as the density of ionization increases, the probability of a direct interaction between the particle track and the target molecule increases.

It is important to note at this stage that the indirect effect involving free radicals is most easily modified by chemical means. Radioprotective compounds have been developed that work by scavenging free radicals. Such compounds, therefore, are quite effective for x- and γ-rays but of little use for neutrons, α-particles, or heavier ions.

**SUMMARY OF PERTINENT CONCLUSIONS**

- X- and γ-rays are indirectly ionizing; the first step in their absorption is the production of fast recoil electrons.
- Neutrons are also indirectly ionizing; the first step in their absorption is the production of fast recoil protons, α-particles, and heavier nuclear fragments.
- Biologic effects of x-rays may be caused by direct action (the recoil electron directly ionizes the target molecule) or indirect action (the recoil electron interacts with water to produce an OH, which diffuses to the target molecule).
- About two thirds of the biologic damage by x-rays is caused by indirect action.
- DNA radicals produced by both the direct and indirect action of radiation are modifiable with sensitizers or protectors.
- DNA lesions produced by high-LET radiations involve large numbers of DNA radicals. Chemical sensitizers and protectors are ineffective in modifying such lesions.
- The physics of the absorption process is over in \(10^{-15}\) second; the chemistry takes longer because the lifetime of the DNA radicals is about \(10^{-5}\) second; the biology takes hours, days, or months for cell killing, years for carcinogenesis, and generations for heritable effects.

**BIBLIOGRAPHY**


GENERAL OVERVIEW OF DNA STRAND BREAKS

There is strong evidence that DNA is the principal target for the biologic effects of radiation, including cell killing, carcinogenesis, and mutation. A consideration of the biologic effects of radiation, therefore, begins logically with a description of the breaks in DNA caused by charged-particle tracks and by the chemical species produced.

Deoxyribonucleic acid (DNA) is a large molecule with a well-known double helical structure. It consists of two strands held together by hydrogen bonds between the bases. The “backbone” of each strand consists of alternating sugar and phosphate groups. The sugar involved is deoxyribose. Attached to this backbone are four bases, the sequence of which specifies the genetic code. Two of the bases are single-ring groups (pyrimidines); these are thymine and cytosine. Two of the bases are double-ring groups (purines); these are adenine and guanine. The structure of a single strand of DNA is illustrated in Figure 2.1. The bases on opposite strands must be complementary; adenine pairs with thymine, and guanine pairs with cytosine. This is illustrated in the simplified model of DNA in Figure 2.2A.

Radiation induces a large number of lesions in DNA, most of which are repaired successfully by the cell and are discussed in the following sections of this chapter. A dose of radiation that induces an average of one lethal event per cell leaves 37% still viable; this is called the D₀ dose and is discussed further in Chapter 3. For mammalian cells, the x-ray D₀ usually lies between 1 and 2 Gy. The number of DNA lesions per cell detected immediately after such a dose is approximately:

- Base damage, >1,000
- Single-strand breaks (SSBs), 1,000
- Double-strand breaks (DSBs), 40

If cells are irradiated with a modest dose of x-rays, many breaks of a single strand occur. These can be observed and scored as a function of dose if the DNA is denatured and the supporting structure is stripped away. In intact DNA, however, SSBs are of little biologic consequence as far as cell killing is concerned because they are repaired readily using the opposite strand as a template (Fig. 2.2B). If the repair is incorrect (misrepair), it may result in a mutation. If both strands of the DNA are broken and the breaks are well separated (Fig. 2.2C), repair again occurs readily because the two breaks are handled separately.

By contrast, if the breaks in the two strands are opposite one another or separated by only a few base pairs (Fig. 2.2D), this may lead to a DSB (double-strand break), resulting in the cleavage of chromatin into two pieces. DSBs are believed to be the most important lesions produced in chromosomes by radiation; as described...
in the next section, the interaction of two DSBs may result in cell killing, carcinogenesis, or mutation. There are many kinds of DSBs, varying in the distance between the breaks on the two DNA strands and the kinds of end groups formed. Their yield in irradiated cells is about 0.04 times that of SSBs, and they are induced linearly with dose, indicating that they are formed by single tracks of ionizing radiation.

Both free radicals and direct ionizations may be involved in the formation of the type of strand break illustrated in Figure 2.2D. As described in Chapter 1, the energy from ionizing radiations is not deposited uniformly in the absorbing medium but is located along the tracks of the charged particles set in motion—electrons in the case of x- or γ-rays, protons and α-particles in the case of neutrons. Radiation chemists speak in terms of “spurs,” “blobs,” and “short tracks.” There is, of course, a full spectrum of energy event sizes, and it is quite arbitrary to divide them into just three categories, but it turns out to be instructive. A spur contains up to 100 eV of energy and involves, on average, three ion pairs. In the case

**FIGURE 2.1** The structure of a single strand of DNA.

**FIGURE 2.2** Diagrams of single- and double-strand DNA breaks caused by radiation. A: Two-dimensional representation of the normal DNA helix. The base pairs carrying the genetic code are complementary (i.e., adenine pairs with thymine, guanine pairs with cytosine). B: A break in one strand is of little significance because it is repaired readily using the opposite strand as a template. C: Breaks in both strands, if well separated, are repaired as independent breaks. D: If breaks occur in both strands and are directly opposite or separated by only a few base pairs, this may lead to a double-strand break in which the chromatin snaps into two pieces. (Courtesy of Dr. John Ward.)
of x- or γ-rays, 95% of the energy deposition events are spurs, which have a diameter of about 4 nm, which is about twice the diameter of the DNA double helix (Fig. 2.3). Blobs are much less frequent for x- or γ-rays; they have a diameter of about 7 nm and contain on average about 12 ion pairs with an energy range of 100–500 eV (Fig. 2.3). Because spurs and blobs have dimensions similar to the DNA double helix, multiple radical attacks occurs if they overlap the DNA helix. There is likely to be a wide variety of complex lesions, including base damage as well as DSBs. The term locally multiply damaged site was initially coined by John Ward to describe this phenomenon, but it has been replaced with the term clustered lesion. Given the size of a spur and the diffusion distance of hydroxyl free radicals, the clustered lesion could be spread out up to 20 base pairs. This is illustrated in Figure 2.3, in which a DSB is accompanied by base damage and the loss of genetic information.

In the case of densely ionizing radiations, such as neutrons or α-particles, a greater proportion of blobs are produced. The damage produced, therefore, is qualitatively different from that produced by x- or γ-rays and it is much more difficult for the cell to repair.

**MEASURING DNA STRAND BREAKS**

Over the years, various techniques have been used to measure DNA strand breaks, including sucrose gradient sedimentation, alkaline and neutral filter elution, nucleoid sedimentation, pulsed-field gel electrophoresis (PFGE), and single-cell gel electrophoresis (also known as the comet assay). Of these techniques, PFGE and single-cell gel electrophoresis are still used to measure DNA strand breaks. In addition to these past techniques, radiation-induced nuclear foci has become a popular approach to visualize DNA damage through the recruitment of DNA repair proteins to sites of DNA damage.

**PFGE** is the method most widely used to detect the induction and repair of DNA DSBs. It is based on the electrophoretic elution of DNA from agarose plugs within which irradiated cells have been embedded and lysed. PFGE allows separation of DNA fragments according to size in the megabase-pair range, with the assumption that DNA DSBs are induced randomly. The fraction of DNA released from the agarose plug is directly proportional to dose (Fig. 2.4A). The kinetics of DNA DSB rejoining exhibit a fast initial rate, which then decreases with repair time. The most widely accepted description of this kinetic behavior uses two first-order components (fast and slow) plus some fraction of residual DSBs. Studies have supported the finding that rejoining of incorrect DNA ends originates solely from slowly rejoining DSBs, and this subset of radiation-induced DSBs is what is manifested as chromosomal damage (i.e., chromosome translocations and exchanges).

**Single-cell electrophoresis (comet assay)** has the advantage of detecting differences in DNA damage and repair at the single-cell level. This is particularly advantageous for biopsy specimens from tumors in which a relatively small number of cells can be assayed to determine DNA damage and repair. Similar to PFGE (described earlier), cells are exposed to ionizing radiation, embedded in agarose, and lysed under neutral buffer conditions to quantify induction.
afforded the DNA by packaging with proteins such as histones. Certain regions of DNA, particularly actively transcribing genes, appear to be more sensitive to radiation, and there is some evidence also of sequence-specific sensitivity.

DNA damage-induced nuclear foci (radiation-induced foci assay) in response to ionizing radiation represents complexes of signaling and repair proteins that localize to sites of DNA strand breaks in the nucleus of a cell. There are several advantages of assaying for foci formation over other techniques to measure DNA strand breaks, which include the ease of the protocol and that it can be carried out on both tissue sections and individual cell preparations. Technically, cells/tissues are incubated with a specific antibody raised to the signaling/repair protein of interest, and binding of the antibody is then detected with a secondary antibody, which also carries a fluorescent tag. Fluorescence microscopy detects the location and intensity of the tag, which can then be quantified.
two proteins involved in the repair of DNA damage by homologous recombination, have been used as biomarkers in a small pilot study by Willers et al. to detect repair defects in breast cancer biopsies.

DNA REPAIR PATHWAYS

Mammalian cells have developed specialized pathways to sense, respond to, and repair base damage, SSBs, DSBs, sugar damage, and DNA–DNA crosslinks. Research from yeast to mammalian cells has demonstrated that the mechanisms used to repair ionizing radiation-induced base damage are different from the mechanisms used to repair DNA DSBs. In addition, different repair pathways are used to repair DNA damage, depending on the stage of the cell cycle.

Much of our knowledge of DNA repair is the result of studying how mutations in individual genes result in radiation hypersensitivity. Radiation-sensitive mutants identified from yeast and mammalian cells appear either to be directly involved in the repair process or to function as molecular checkpoint–controlling elements. The pathways involved in the repair of base damage, SSBs, DSBs, sugar damage, and DNA–DNA crosslinks are discussed in the next sections and represent a simplified representation of our current state of understanding.
In Chapter 18, the syndromes associated with mutations in genes involved in sensing DNA damage or repairing DNA damage are discussed in more detail.

**Base Excision Repair**

Base damage is repaired through the base excision repair (BER) pathway illustrated in Figure 2.6. Bases on opposite strands of DNA must be complementary; adenine (A) pairs with thymine (T), and guanine (G) pairs with cytosine (C). U therefore represents a putative single-base mutation that is first removed by a glycosylase/DNA lyase (Fig. 2.6A). Removal of the base is followed by the removal of the sugar residue by an apurinic endonuclease 1 (APE1), then replacement with the correct nucleotide by DNA polymerase β, and completed by DNA ligase III-XRCC1-mediated ligation. If more than one nucleotide is to be replaced (illustrated by the putative mutation UU in Fig. 2.6B), then the complex of replication factor C (RFC)/proliferating cell nuclear antigen (PCNA)/DNA polymerase δ/ε performs the repair synthesis, the overhanging flap structure is removed by the flap endonuclease 1 (FEN1), and DNA strands are sealed by ligase I (Fig. 2.6B). Although ionizing radiation–induced base damage is efficiently repaired, defects in BER may lead to an increased mutation rate, but usually do not result in cellular radiosensitivity. One exception to this is the mutation of the x-ray cross complementing factor 1 (XRCC1) gene, which confers about a 1.7-fold increase in radiation sensitivity. However, the radiation sensitivity of XRCC1-deficient cells may come from XRCC1’s potential involvement in other repair processes such as SSBs.

**Nucleotide Excision Repair**

Nucleotide excision repair (NER) removes bulky adducts in the DNA such as pyrimidine dimers. The process of NER can be subdivided into two pathways: global genome repair (GGR or GG-NER) and transcription-coupled repair (TCR or TC-NER). The process of GG-NER is genome-wide (i.e., lesions can be removed from DNA that encodes or does not encode for genes). In contrast, TC-NER only removes lesions in the DNA strands of actively

**FIGURE 2.6** Base excision repair pathways. **A:** Base excision repair of a single nucleotide. Bases on opposite strands must be complementary; adenine (A) pairs with thymine (T), and guanine (G) pairs with cytosine (C). U represents a putative mutation that is first removed through a DNA glycosylase–mediated incision step. **B:** Base excision repair of multiple nucleotides. In this case, the double UU represents a putative mutation that is first removed through apurinic endonuclease 1 (APE1). See text for details.
transcribed genes. When a DNA strand that is being actively transcribed becomes damaged, the RNA polymerase can block access to the site of damage and hence prevents DNA repair. TC-NER has evolved to prevent this blockade by RNA polymerase by effectively removing it from the site of damage to allow the repair proteins access. The mechanism of GG-NER and TC-NER differs only in the detection of the lesion; the remainder of the pathway used to repair the damage is the same for both. The essential steps in this pathway are (1) damage recognition; (2) DNA incisions that bracket the lesion, usually between 24 and 32 nucleotides in length; (3) removal of the region containing the adducts; (4) repair synthesis to fill in the gap region; and (5) DNA ligation (Fig. 2.7). Mutation in NER genes does not lead to ionizing radiation sensitivity. However, defective NER increases sensitivity to UV-induced DNA damage and anticancer agents such as alkylating agents that induce bulky adducts. Germline mutations in NER genes lead to human DNA repair deficiency disorders such as xeroderma pigmentosum in which patients are hypersensitive to ultraviolet light.

**DNA Double-Strand Break Repair**

In eukaryotic cells, DNA DSBs can be repaired by two basic processes: homologous recombination repair (HRR), which requires an undamaged DNA strand as a participant in the repair as a template, and nonhomologous end-joining (NHEJ), which mediates end-to-end joining. In lower eukaryotes such as yeast, HRR is the predominant pathway used for repairing DNA DSBs. Homologous recombination is an error-free process because
The ligation of DNA DSBs by NHEJ does not require sequence homology. However, the damaged ends of DNA DSBs cannot simply be ligated together; they must first be modified before they can be rejoined by a ligation reaction. NHEJ can be divided into five steps: (1) end recognition by Ku binding, (2) recruitment of DNA-dependent protein kinase catalytic subunit (DNA-PKcs), (3) end processing, (4) fill-in synthesis or end bridging, and (5) ligation (Fig. 2.9).

End recognition occurs when the Ku heterodimer, composed of 70-kDa and 83-kDa subunits, and the DNA-PKcs bind to the ends of the DNA DSB. Although the Ku/DNA-PKcs complex is thought to bind ends first, it is still unknown what holds the two DNA DSB ends together. Although microhomology between one to four nucleotides can aid in end alignment, there is no absolute requirement for microhomology for NHEJ. In fact, Ku does not only recruit DNA-PKcs to the DNA ends, but an additional protein, Artemis, which possesses endonuclease activity, forms a physical complex with DNA-PKcs. The Ku/DNA-PKcs complex that is bound to the DNA ends gives rise to a signal for a fill-in reaction to proceed after endonuclease processing. However, DNA polymerase μ or λ has been found to be associated with the Ku/DNA/XRCC4/DNA ligase IV complex and serves as the polymerase for the fill-in reaction. In the final step of NHEJ, ligation of nicked DNA is performed by copying information from the undamaged homologous chromatid/chromosome.
**FIGURE 2.9** Nonhomologous end-joining. DNA strand breaks are recognized by the ATM and the MRN (Mre11-Rad50-Nbs1) complex, resulting in resection of the DNA ends. Homologous recombination is inhibited by the activity of 53BP1. The initial step of the core NHEJ pathway starts with the binding of the ends at the DSB by the Ku70/Ku80 heterodimer. This complex then recruits and activates the catalytic subunit of DNA-PK (DNA-PKcs), whose role is the juxtaposition of the two DNA ends. The DNA-PK complex then recruits the ligase complex (XRCC4/XLF-LIGIV/PNK) that promotes the final ligation step.

DNA ends that have been processed is mediated by a PNK/XRCC4/DNA ligase IV/XLF complex that is probably recruited by the Ku heterodimer. Polynucleotide kinase (PNK) is a protein that has both 3'-DNA phosphatase and 5'-DNA kinase activities and serves to remove end groups that are not ligatable to allow end-joining. XRCC4-like factor (XLF) is a protein that has a similar protein structure as x-ray repair complementing defective repair in Chinese hamster cells 4 (XRCC4) and stimulates the activity of DNA ligase IV. NHEJ is error prone and plays an important physiologic role in generating antibodies through V(D)J rejoining. The error-prone nature of NHEJ is essential for generating antibody diversity and often goes undetected in mammalian cells, as errors in the noncoding DNA that composes most human genome has little consequence. NHEJ is primarily found in the G1 phase of the cell cycle, where there is no sister chromatid.

**Homologous Recombination Repair**

HRR provides the mammalian genome a high-fidelity mechanism of repairing DNA DSBs (Fig. 2.10). In particular, the increased activity of this recombination pathway in late S/G2 suggests that its primary function is to repair and restore the functionality of replication forks with DNA DSBs. Compared to NHEJ, which requires no sequence homology to rejoin broken ends, HRR requires physical contact with an undamaged chromatid or chromosome (to serve as a template) for repair to occur.
FIGURE 2.10 Homologous recombinational repair. The initial step in HR is the recognition of the lesion and processing of the double-strand DNA ends into 3’ DNA single-strand tails by the MRN (Mre11-Rad50-Nbs1) complex, which are then coated by RPA forming a nucleoprotein filament. Then, specific HR proteins are recruited to the nucleoprotein filaments, such as RAD51, RAD52, and BRCA1/2. RAD51 is a key protein in homologous recombination as it mediates the invasion of the homologous strand of the sister chromatid, leading to formation of Holliday junctions. The Holliday junctions are finally resolved into two DNA duplexes. See text for details.

During recombination, evidence exists that ATM phosphorylates the breast cancer tumor suppressor protein BRCA1, which is then recruited to the site of the DSB that has been bound by the NBS/MRE11/Rad50s protein complex (Fig. 2.10). MRE11 and perhaps other yet unidentified endonucleases resect the DNA, resulting in a 3’ single-strand DNA that serves as a binding site for Rad51. BRCA2, which is attracted to the DSB by BRCA1, facilitates the loading of Rad51 onto RPA-coated single-strand overhangs produced by endonuclease resection. Rad51 protein is a homologue of the *Escherichia coli* recombinase RecA and possesses the ability to form nucleofilaments and catalyze strand exchange with the complementary strand in the undamaged chromosome. Five additional paralogues of Rad51 also bind to the RPA-coated single-stranded region and recruit Rad52, which protects against exonucleolytic degradation. To facilitate repair, Rad54 uses its ATPase activity to unwind the double-stranded molecule. The two invading ends serve as primers for DNA synthesis, and the so-called Holliday junctions are resolved by MMS4 and MUS81 by noncrossing over, in which case, the Holliday junctions disengage and DNA strand pairing is followed by gap filling, or by crossing over of the Holliday junctions, which is followed by gap filling. The identities of the polymerase and ligase involved in these latter steps are unknown. Because inactivation of HRR genes results in radiosensitivity and genomic instability, these genes provide a critical link between HRR and chromosome stability. Dysregulated homologous recombination can also lead to cancer by loss of heterozygosity (LOH).
Crosslink Repair
Several DNA–DNA and DNA–protein crosslinks induced by ionizing radiation have not been extensively studied to arrive at a quantitative estimate. Furthermore, the genes and pathways used for DNA–DNA or DNA–protein crosslink repair are still under investigation. The current thinking is that a combination of NER and recombinational repair pathways is needed to repair DNA crosslinks (Fig. 2.11). The predominant signal from a DNA-interstrand crosslink that signals for repair is stalling of the DNA replication fork. The crosslink is removed in a multistep process, first from one strand by a second round of NER, resulting in a strand break and a DNA adduct. DNA synthesis can proceed past the lesion, resulting in a point mutation opposite the lesion. However, the SSB will become a DSB, and seems to require HRR for restitution. Finally, the adduct that remains is removed by NER. Cells with mutations in NER and HRR pathways are modestly sensitive to crosslinking agents. In contrast, individuals afflicted with the syndrome Fanconi anemia are hypersensitive to crosslinking agents. Chromatin that contains actively transcribed genes is more susceptible to DNA–protein crosslinks, and the crosslinked proteins are usually nuclear matrix proteins.

Mismatch Repair
The mismatch repair (MMR) pathway removes base–base and small insertion mismatches that occur during replication. In addition, the MMR pathway removes base–base mismatches in homologous recombination intermediates. See Figure 2.12 for schematic representation and an indication of the critical gene products. The process of MMR can be subdivided into four components: first, the mismatch must be identified by sensors that transduce the signal of a mismatched base pair; second, MMR factors are recruited; third, the newly synthesized strand harboring the mismatch is identified and the incorrect/altered nucleotides are excised; and in the fourth stage, resynthesis and ligation.

**FIGURE 2.11** DNA–DNA cross-link repair. The initial signal for DNA–DNA crosslinks is stalling of the replication fork (A). The crosslink is removed from one strand by nucleotide excision repair (B), followed by translesion synthesis, resulting in a mutation opposite the adduct (C). The resulting DNA double-strand break is repaired by homologous recombination (D) and the crosslink is removed from the DNA by another round of nucleotide excision repair (E–F). This schema for crosslink repair is still a work in progress.
of the excised DNA tract is completed. MMR was first characterized in *E. coli* by the characterization of the *Mut* genes, of which homologues of these gene products have been identified and extensively characterized in both yeast and humans. Mutations in any of the mismatch *MSH*, *MLH*, and *PSM* families of repair genes lead to microsatellite instability (small base insertions or deletions) and cancer, especially hereditary nonpolyposis colon cancer (HNPCC).

### RELATIONSHIP BETWEEN DNA DAMAGE AND CHROMOSOME ABERRATIONS

Cell killing does not correlate with SSBs, but relates better to DSBs. Agents (such as hydrogen peroxide) produce SSBs efficiently, but very few DSBs, and also kill very few cells. Cells defective in DNA DSB repair exhibit hypersensitivity to killing by ionizing radiation and increased numbers of chromosome aberrations. On the basis of evidence such as this, it is concluded that DSBs are the most relevant lesions leading to most biologic insults from radiation including cell killing. The reason for this is that DSBs can lead to chromosomal aberrations that present problems at cell division.

### CHROMOSOMES AND CELL DIVISION

The backbone of DNA is made of molecules of sugar and phosphates, which serve as a framework to hold the bases that carry the genetic code. Attached to each sugar molecule is a base: thymine, adenine, guanine, or cytosine. This whole configuration is coiled tightly in a double helix.

Figure 2.13 is a highly schematized illustration of the way an organized folding of the long DNA helix might be achieved as a closely packed series of looped domains wound in a tight helix. The degree of packing also is illustrated by the

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**Figure 2.12** Mismatch repair. The initial step in the mismatch repair pathway is the recognition of mismatched bases through either Msh2-Msh6 or Msh2-Msh3 complexes. These recognition complexes recruit MLH1-PMS2, MLH1-PMS1, and MLH1-MLH3, alongside the exonuclease EXO1 that catalyzes the excision step that follows. A gap-filling step by polymerases δ/ε, RCF, and PCNA is followed by a final ligation step.
The various events that occur during mitosis are divided into several phases. The first phase of division is called prophase. The beginning of this phase is marked by a thickening of the chromatin and an increase in its stainability as the chromosomes condense into light coils. By the end of prophase, each chromosome has a lightly staining constriction known as a centromere; extending from the centromere are the arms of the chromosome. Prophase ends when the chromosomes reach maximal condensation and the nuclear membrane disappears, as do any nucleoli.

With the disappearance of the nuclear membrane, the nuclear plasm and the cytoplasm mix. Metaphase then follows, in which two events occur simultaneously. The chromosomes move to the center of the cell (i.e., to the cell’s equator), and the spindle forms. The spindle is composed of fibers that cross the cell, linking its poles. Once the chromosomes are stabilized at the equator of the cell, their centromeres divide, and metaphase is complete.

The phase that follows, anaphase, is characterized by a movement of the chromosomes on the spindle to the poles. The chromosomes appear to be pulled toward the poles of the cell by fibers attached to the centromeres. The arms, particularly the long arms, tend to trail behind.

Anaphase is followed by the last phase of mitosis, telophase. In this phase, the chromosomes, congregated at the poles of the cell, begin to uncoil. The nuclear membrane reappears, as do the nucleoli; and as the phase progresses, the chromosome coils unwind until the nucleus regains the appearance characteristic of interphase.

THE ROLE OF TELOMERES

Telomeres cap and protect the terminal ends of chromosomes. The name telomere literally means “end part.” Mammalian telomeres consist of long arrays of TTAGGGG repeats that range in total length anywhere from 1.5 to 150 kilobases. Each time a normal somatic cell divides, telomeric DNA is lost from the lagging strand because DNA polymerase cannot synthesize

FIGURE 2.13 Illustration of the relative sizes of the DNA helix, the various stages of folding and packing of the DNA, and an entire chromosome condensed at metaphase.
new DNA in the absence of an RNA primer. Successive divisions lead to progressive shortening, and after 40 to 60 divisions, the telomeres in human cells are shortened dramatically, so that vital DNA sequences begin to be lost. At this point, the cell cannot divide further and undergoes senescence. Telomere length has been described as the “molecular clock” or generational clock because it shortens with age in somatic tissue cells during adult life. Stem cells in self-renewing tissues, and cancer cells in particular, avoid this problem of aging by activating the enzyme telomerase. Telomerase is a reverse transcriptase that includes the complementary sequence to the TTAGGG repeats and so continually rebuilds the chromosome ends to offset the degradation that occurs with each division.

In tissue culture, immortalization of cells—that is, the process whereby cells pass through a “crisis” and continue to be able to divide beyond the normal limit—is associated with telomere stabilization and telomerase activity.

Virtually all human tumor cell lines and approximately 90% of human cancer biopsy specimens exhibit telomerase activity. By contrast, normal human somatic tissues, other than stem cells, do not possess detectable levels of this enzyme. It is an attractive hypothesis that both immortalization and carcinogenesis are associated with telomerase expression.

### RADIATION-INDUCED CHROMOSOME ABERRATIONS

In the traditional study of chromosome aberrations, the effects of ionizing radiations are described in terms of their appearance when a preparation is made at the first metaphase after exposure to radiation. This is the time when the structure of the chromosomes can be discerned.

The study of radiation damage in mammalian cell chromosomes is hampered by the large number of mammalian chromosomes per cell and by their small size. Most mammalian cells currently available for experimental purposes have a diploid complement of 40 or more chromosomes. There are exceptions, such as the Chinese hamster, with 22 chromosomes, and various marsupials, such as the rat kangaroo and woolly opossum, which have chromosome complements of 12 and 14, respectively. Many plant cells, however, contain fewer and generally much larger chromosomes; consequently, until recently, information on chromosomal radiation damage accrued principally from studies with plant cells.

If cells are irradiated with x-rays, DSBs are produced in the chromosomes. The broken ends appear to be “sticky” because of unpaired bases and can rejoin with any other sticky end. It would appear, however, that a broken end cannot join with a normal, unbroken chromosome, although this is controversial. Once breaks are produced, different fragments may behave in various ways:

1. The breaks may restitute, that is, rejoin in their original configuration. In this case, of course, nothing amiss is visible at the next mitosis.
2. The breaks may fail to rejoin and give rise to an aberration, which is scored as a deletion at the next mitosis.
3. Broken ends may reassort and rejoin other broken ends to give rise to chromosomes that appear to be grossly distorted if viewed at the following mitosis.

The aberrations seen at metaphase are of two classes: chromosome aberrations and chromatid aberrations. Chromosome aberrations result if a cell is irradiated early in interphase, before the chromosome material has been duplicated. In this case, the radiation-induced break is in a single strand of chromat; during the DNA synthetic phase that follows, this strand of chromat lays down an identical strand next to itself and replicates the break that has been produced by the radiation. This leads to a chromosome aberration visible at the next mitosis because there is an identical break in the corresponding points of a pair of chromatin strands. If, on the other hand, the dose of radiation is given later in interphase, after the DNA material has doubled and the chromosomes consist of two strands of chromatin, then the aberrations produced are called chromatid aberrations. In regions removed from the centromere, chromatid arms may be fairly well separated, and it is reasonable to suppose that the radiation might break one chromatid without breaking its sister chromatid, or at least not in the same place. A break that occurs in a single chromatid arm after chromosome replication and leaves the opposite arm of the same chromosome undamaged leads to chromatid aberrations.
EXAMPLES OF RADIATION-INDUCED ABERRATIONS

Many types of chromosomal aberrations and rearrangements are possible, but an exhaustive analysis is beyond the scope of this book. Three types of aberrations that are lethal to the cell are described, followed by two common rearrangements that are consistent with cell viability but are frequently involved in carcinogenesis. The three lethal aberrations are the dicentric; the ring, which are chromosome aberrations; and the anaphase bridge, which is a chromatid aberration. All three represent gross distortions and are clearly visible. Many other aberrations are possible but are not described here.

The formation of a dicentric is illustrated in diagrammatic form in Figure 2.14A. This aberration involves an interchange between two separate chromosomes. If a break is produced in each one early in interphase and the sticky ends are close to one another, they may rejoin as shown. This bizarre interchange is replicated during the DNA synthetic phase, and the result is a grossly distorted chromosome with two centromeres (hence, dicentric). There also are two fragments that have no centromere (acentric fragment), which will therefore be lost at a subsequent mitosis. The appearance at metaphase is shown in the bottom panel of Figure 2.14A. An example of a dicentric and fragment in a metaphase human cell is shown in Figure 2.15B; Figure 2.15A shows a normal metaphase for comparison.

The formation of a ring is illustrated in diagrammatic form in Figure 2.14B. A break is induced by radiation in each arm of a single chromatid early in the cell cycle. The sticky ends may rejoin to form a ring and a fragment. Later in the cycle, during the DNA synthetic phase, the chromosome replicates. The ultimate appearance at metaphase is shown in the lower panel of Figure 2.14B. The fragments have no centromere and probably will be lost at mitosis because they will not be pulled to either pole of the cell. An example of a ring chromosome in a human cell at metaphase is illustrated in Figure 2.15C.

An anaphase bridge may be produced in various ways. As illustrated in Figure 2.14C and Figure 2.16, it results from breaks that occur late in the cell cycle (in G2) after the chromosomes have replicated. Breaks may occur in both chromatids of the same chromosome, and the sticky ends may rejoin incorrectly to form a sister union. At anaphase, when the two sets of chromosomes move to opposite poles, the section of chromatin between the two centromeres is stretched across the cell between the poles, hindering the separation into two new progeny cells, as illustrated in Figure 2.14C and Figure 2.16B. The two fragments may join as shown, but because there is no centromere, the joined fragments will probably be lost at the first mitosis. This type of aberration occurs in human cells and is essentially always lethal. It is hard to demonstrate because preparations of human chromosomes usually are made by accumulating cells at metaphase, and the bridge is only evident at anaphase. Figure 2.16 is an anaphase preparation of Tradescantia paludosa, a plant used extensively for cytogenetic studies because of the small number of large chromosomes. The anaphase bridge is seen clearly as the replicate sets of chromosomes move to opposite poles of the cell.

Gross chromosome changes of the types discussed previously inevitably lead to the reproductive death of the cell.

Two important types of chromosomal changes that are not lethal to the cell are symmetric translocations and small deletions. The formation of a symmetric translocation is illustrated in Figure 2.17A. It involves a break in two prereplication (G1) chromosomes, with the broken ends being exchanged between the two chromosomes as illustrated. An aberration of this type is difficult to see in a conventional preparation but is easy to observe with the technique of fluorescent in situ hybridization (FISH), or chromosome painting, as it commonly is called. Probes are available for every human chromosome that makes them fluorescent in a different color. Exchange of material between two different chromosomes then is readily observable (Fig. 2.18). Translocations are associated with several human malignancies caused by the activation of an oncogene; Burkitt lymphoma and certain types of leukemia are examples.

The other type of nonlethal chromosomal change is a small interstitial deletion. This is illustrated in Figure 2.17B and may result from two breaks in the same arm of the same chromosome, leading to the loss of the genetic information between the two breaks. The actual sequence of events in the formation of a deletion is easier
FIGURE 2.14  A: The steps in the formation of a dicentric by irradiation of prereplication (i.e., G₁) chromosomes. A break is produced in each of two separate chromosomes. The “sticky” ends may join incorrectly to form an interchange between the two chromosomes. Replication then occurs in the DNA synthetic period. One chromosome has two centromeres: a dicentric. The acentric fragment will also replicate and both will be lost at a subsequent mitosis because, lacking a centromere, they will not go to either pole at anaphase. B: The steps in the formation of a ring by irradiation of a prereplication (i.e., G₁) chromosome. A break occurs in each arm of the same chromosome. The sticky ends rejoin incorrectly to form a ring and an acentric fragment. Replication then occurs. C: The steps in the formation of an anaphase bridge by irradiation of a postreplication (i.e., G₂) chromosome. Breaks occur in each chromatid of the same chromosome. Incorrect rejoining of the sticky ends then occurs in a sister union. At the next anaphase, the acentric fragment will be lost, one centromere of the dicentric will go to each pole, and the chromatid will be stretched between the poles. Separation of the progeny cells is not possible; this aberration is likely to be lethal. (Courtesy of Dr. Charles Geard.)
FIGURE 2.15  Radiation-induced chromosome aberrations in human leukocytes viewed at metaphase. 
A: Normal metaphase. B: Dicentric and fragment (arrows). (Continued)
to understand from Figure 2.19, which shows an interphase chromosome. It is a simple matter to imagine how two breaks may isolate a loop of DNA—an acentric ring—which is lost at a subsequent mitosis. Deletions may be associated with carcinogenesis if the lost genetic material includes a tumor suppressor gene. This is discussed further in Chapter 10 on radiation carcinogenesis.

The interaction between breaks in different chromosomes is by no means random. There is great heterogeneity in the sites at which deletions and exchanges between different chromosomes occur; for example, chromosome 8 is particularly sensitive to exchanges. As mentioned previously, each chromosome is restricted to a domain, and most interactions occur at the edges of domains, which probably involves the nuclear matrix. Active chromosomes are therefore those with the biggest surface area to their domains.

- **CHROMOSOME ABERRATIONS IN HUMAN LYMPHOCYTES**

Chromosomal aberrations in peripheral lymphocytes have been used widely as biomarkers of radiation exposure. In blood samples obtained for cytogenetic evaluation within a few days to a few weeks after total body irradiation, the frequency of asymmetric aberrations (dicentrics and rings) in the lymphocytes reflects the dose received. Lymphocytes in the blood sample are stimulated to divide with a mitogen such as phytohemagglutinin and are arrested at metaphase, and the incidence of rings and dicentrics is scored. The dose can be estimated by comparison with in vitro cultures exposed to known doses. Figure 2.20 shows a dose-response curve for aberrations in human lymphocytes produced by γ-rays. The data are fitted by a linear-quadratic relationship, as would be expected, because rings and dicentrics result from the interaction of two chromosome breaks, as previously described. The linear component is a consequence of the two breaks resulting from a single charged particle. If the two breaks result from different charged particles, the probability of an interaction is a quadratic function of dose. This also is illustrated for the formation of a dicentric in Figure 2.20.

If a sufficient number of metaphases are scored, cytogenetic evaluations in cultured lymphocytes readily can detect a recent total body exposure of as low as 0.25 Gy in the exposed person. Such studies are useful in distinguishing between “real” and “suspected” exposures, particularly in those instances involving “black film badges” or
in potential accidents in which it is not certain whether individuals who were at risk for exposure actually received radiation doses.

Mature T lymphocytes have a finite life span of about 1,500 days and are eliminated slowly from the peripheral lymphocyte pool. Consequently, the yield of dicentrics observed in peripheral lymphocytes declines in the months and years after a radiation exposure.

During \textit{in vivo} exposures to ionizing radiation, chromosome aberrations are induced not only in mature lymphocytes but also in lymphocyte progenitors in marrow, nodes, or other organs. The stem cells that sustain asymmetric aberrations

\textbf{FIGURE 2.16} Anaphase chromosome preparation of \textit{Tradescantia paludosa}. \textbf{A}: Normal anaphase. \textbf{B}: Bridge and fragment resulting from radiation (arrow). (Courtesy of Drs. Brewen, Luippold, and Preston.)
**FIGURE 2.17** A: Formation of a symmetric translocation. Radiation produces breaks in two different prereplication chromosomes. The broken pieces are exchanged between the two chromosomes, and the “sticky” ends rejoin. This aberration is not necessarily lethal to the cell. There are examples in which an exchange aberration of this type leads to the activation of an oncogene. See Chapter 10 on radiation carcinogenesis.

B: Diagram of a deletion. Radiation produces two breaks in the same arm of the same chromosome. What actually happens is illustrated more clearly in Figure 2.18.

**FIGURE 2.18** Fluorescence *in situ* hybridization of a metaphase spread from a cell that received 4 Gy. The hybridization was performed with a cocktail of DNA probes that specifically recognize each chromosome pair. Chromosome aberrations are demarcated by the arrows. (Courtesy of Dr. Michael Cornforth.)
dicentrics underestimates the dose and only stable aberrations such as translocations give an accurate picture. Until recently, translocations were much more difficult to observe than dicentrics, but now the technique of FISH makes the scoring of such symmetric aberrations a relatively simple matter. The frequency of translocations assessed in this way correlates with total body dose in exposed individuals even after more than 50 years, as was shown in a study of the survivors of the atomic bomb attacks on Hiroshima and Nagasaki.

(such as dicentrics) die in attempting a subsequent mitosis, but those that sustain a symmetric non-lethal aberration (such as a translocation) survive and pass on the aberration to their progeny. Consequently, dicentrics are referred to as “unstable” aberrations because their number declines with time after irradiation. Symmetric translocations, by contrast, are referred to as “stable” aberrations because they persist for many years. Either type of aberration can be used to estimate dose soon after irradiation, but if many years have elapsed, scoring dicentrics underestimates the dose and only stable aberrations such as translocations give an accurate picture. Until recently, translocations were much more difficult to observe than dicentrics, but now the technique of FISH makes the scoring of such symmetric aberrations a relatively simple matter. The frequency of translocations assessed in this way correlates with total body dose in exposed individuals even after more than 50 years, as was shown in a study of the survivors of the atomic bomb attacks on Hiroshima and Nagasaki.

FIGURE 2.19 Formation of a deletion by ionizing radiation in an interphase chromosome. It is easy to imagine how two breaks may occur (by a single or two different charged particles) in such a way as to isolate a loop of DNA. The “sticky” ends rejoin, and the deletion is lost at a subsequent mitosis because it has no centromere. This loss of DNA may include the loss of a suppressor gene and lead to a malignant change. See Chapter 10 on radiation carcinogenesis.

FIGURE 2.20 The frequency of chromosomal aberrations (dicentrics and rings) is a linear-quadratic function of dose because the aberrations are the consequence of the interaction of two separate breaks. At low doses, both breaks may be caused by the same electron; the probability of an exchange aberration is proportional to dose (D). At higher doses, the two breaks are more likely to be caused by separate electrons. The probability of an exchange aberration is proportional to the square of the dose (D²).
SUMMARY OF PERTINENT CONCLUSIONS

- Ionizing radiation induces base damage, SSBs, DSBs, and DNA protein crosslinks.
- The cell has evolved an intricate series of sensors and pathways to respond to each type of radiation-induced damage.
- DNA DSBs, the most lethal form of ionizing radiation–induced damage, is repaired by nonhomologous recombination in the G2 phase of the cell cycle and homologous recombination (mainly) in the S/G2 phase of the cell cycle.
- Defective nonhomologous recombination leads to chromosome aberrations, immune deficiency, and ionizing radiation sensitivity.
- Defective homologous recombination leads to chromatid and chromosome aberrations, decreased proliferation, and ionizing radiation sensitivity.
- There is good reason to believe that DSBs rather than SSBs lead to important biological endpoints including cell death, carcinogenesis, and mutation.
- Radiation-induced breakage and incorrect rejoicing in prereplication (G1) chromosomes may lead to chromosome aberrations.
- Radiation-induced breakage and incorrect rejoicing in postreplication (late S or G2) chromosomes may lead to chromatid aberrations.
- Lethal aberrations include dicentrics, rings, and anaphase bridges. Symmetric translocations and small deletions are nonlethal.
- There is a good correlation between cells killed and cells with asymmetric exchange aberrations (i.e., dicentrics or rings).
- The incidence of most radiation-induced aberrations is a linear-quadratic function of dose.
- Scoring aberrations in lymphocytes from peripheral blood may be used to estimate total body doses in humans accidentally irradiated. The lowest single dose that can be detected readily is 0.25 Gy.
- Dicentrics are “unstable” aberrations; they are lethal to the cell and are not passed on to progeny. Consequently, the incidence of dicentrics declines slowly with time after exposure.
- Translocations are “stable” aberrations; they persist for many years because they are not lethal to the cell and are passed on to the progeny.

BIBLIOGRAPHY

**REPRODUCTIVE INTEGRITY**

A cell survival curve describes the relationship between the radiation dose and the proportion of cells that survive. What is meant by “survival”? Cell survival, or its converse, cell death, may mean different things in different contexts; therefore, a precise definition is essential. For differentiated cells that do not proliferate, such as nerve, muscle, or secretory cells, death can be defined as the loss of a specific function. For proliferating cells, such as stem cells in the hematopoietic system or the intestinal epithelium, loss of the capacity for sustained proliferation—that is, loss of reproductive integrity—is an appropriate definition. This is sometimes called reproductive death. This is certainly the end point measured with cells cultured *in vitro*.

This definition reflects a narrow view of radiobiology. A cell may still be physically present and apparently intact, may be able to make proteins or synthesize DNA, and may even be able to struggle through one or two mitoses; but if it has lost the capacity to divide indefinitely and produce a large number of progeny, it is by definition dead; it has not survived. A survivor that has retained its reproductive integrity and is able to proliferate indefinitely to produce a large clone or colony is said to be clonogenic.

This definition is generally relevant to the radiobiology of whole animals and plants and their tissues. It has particular relevance to the radiotherapy of tumors. For a tumor to be eradicated, it is only necessary that cells be “killed” in the sense that they are rendered unable to divide and cause further growth and spread of the malignancy. Cells may die by different mechanisms, as is described here subsequently. For most cells, death while attempting to divide, that is, mitotic death, is the dominant mechanism following irradiation. For some cells, programmed cell death, or apoptosis, is important. Whatever the mechanism, the outcome is the same: The cell loses its ability to proliferate indefinitely, that is, its reproductive integrity.

In general, a dose of 100 Gy is necessary to destroy cell function in nonproliferating systems. By contrast, the mean lethal dose for loss of proliferative capacity is usually less than 2 Gy.

**THE IN VITRO SURVIVAL CURVE**

The capability of a single cell to grow into a large colony that can be seen easily with the naked eye is a convenient proof that it has retained its reproductive integrity. The loss of this ability as a function of radiation dose is described by the dose-survival curve.
With modern techniques of tissue culture, it is possible to take a specimen from a tumor or from many normal regenerative tissues, chop it into small pieces, and prepare a single-cell suspension by the use of the enzyme trypsin, which dissolves and loosens the cell membrane. If these cells are seeded into a culture dish, covered with an appropriate complex growth medium, and maintained at $37^\circ$ C under aseptic conditions, they attach to the surface, grow, and divide.

In practice, most fresh explants grow well for a few weeks but subsequently peter out and die. A few pass through a crisis and continue to grow for many years. Every few days, the culture must be “farmed”: The cells are removed from the surface with trypsin, most of the cells are discarded, and the culture flask is reseeded with a small number of cells, which quickly repopulate the culture flask. These are called **established cell lines**; they have been used extensively in experimental cellular radiobiology.

Survival curves are so basic to an understanding of much of radiobiology that it is worth going through the steps involved in a typical experiment using an established cell line in culture.

Cells from an actively growing stock culture are prepared into a suspension by the use of trypsin, which causes the cells to round up and detach from the surface of the culture vessel.

The number of cells per unit volume of this suspension is counted in a hemocytometer or with an electronic counter. In this way, for example, 100 individual cells may be seeded into a dish; if this dish is incubated for 1 to 2 weeks, each single cell divides many times and forms a colony that is easily visible with the naked eye, especially if it is fixed and stained (Fig. 3.1). All cells making up each colony are the progeny of a single ancestor. For a nominal 100 cells seeded into the dish, the number of colonies counted may be expected to be in the range of 50 to 90. Ideally, it should be 100, but it seldom is for various reasons, including suboptimal growth medium, errors and uncertainties in counting the cell suspension, and the trauma of trypsinization and handling. The term **plating efficiency** indicates the percentage of cells seeded that grow into colonies. The plating efficiency is given by the formula

\[
\text{Plating Efficiency (PE)} = \frac{\text{Number of colonies counted}}{\text{Number of cells seeded}} \times 100
\]

There are 70 colonies on the control dish in Figure 3.1A; therefore, the plating efficiency is 70%. If a parallel dish is seeded with cells, exposed to a dose of 8 Gy of x-rays, and incubated for 1 to 2 weeks before being fixed and stained, then the following may be observed: (1) Some of the seeded single cells are still single and have

![Figure 3.1](image)
Chapter 3 • Cell Survival Curves

not divided, and in some instances the cells show evidence of nuclear deterioration as they die an apoptotic death; (2) some cells have managed to complete one or two divisions to form a tiny abortive colony; and (3) some cells have grown into large colonies that differ little from the unirradiated controls, although they may vary more in size. These cells are said to have survived because they have retained their reproductive integrity.

In the example shown in Figure 3.1B, 2,000 cells had been seeded into the dish exposed to 8 Gy. Because the plating efficiency is 70%, 1,400 of the 2,000 cells plated would have grown into colonies if the dish had not been irradiated. In fact, there are only 32 colonies on the dish; the fraction of cells surviving the dose of x-rays is thus

$$\frac{32}{1,400} = 0.023$$

In general, the surviving fraction is given by

$$\text{Surviving fraction} = \frac{\text{Colonies counted}}{\text{Cells seeded} \times (\text{PE}/100)}$$

This process is repeated so that estimates of survival are obtained for a range of doses. The number of cells seeded per dish is adjusted so that a countable number of colonies results: Too few reduces statistical significance; too many cannot be counted accurately because they tend to merge into one another. The technique is illustrated in Figure 3.2. This technique, and the survival curve that results, does not distinguish the mode of cell death, that is, whether the cells died mitotic or apoptotic deaths.

THE SHAPE OF THE SURVIVAL CURVE

Survival curves for mammalian cells usually are presented in the form shown in Figure 3.3, with dose plotted on a linear scale and surviving fraction on a logarithmic scale. Qualitatively, the shape of the survival curve can be described in relatively simple terms. At “low doses” for sparsely ionizing (low-linear energy transfer [LET]) radiations, such as x-rays, the survival curve starts out straight on the log-linear plot with a finite initial slope; that is, the surviving fraction is an exponential function of dose. At higher doses, the curve bends. This bending or curving region extends over a dose range of a few grays. At very high doses, the survival curve often tends to straighten again; the surviving fraction returns to being an exponential function of dose. In general, this does not occur until doses in excess of those used as daily fractions in radiotherapy have been reached.

By contrast, for densely ionizing (high-LET) radiations, such as α-particles or low-energy neutrons, the cell survival curve is a straight line from the origin; that is, survival approximates to an exponential function of dose (see Fig. 3.3).

Although it is a simple matter to qualitatively describe the shape of the cell survival curve, finding an explanation of the biologic observations in terms of biophysical events is another matter. Many biophysical models and theories have been proposed to account for the shape of the mammalian cell survival curve. Almost all can be used to deduce a curve shape that is consistent with experimental data, but it is never possible to choose among different models or theories based on goodness of fit to experimental data. The biologic data are not sufficiently precise, nor are the predictive theoretic curves sufficiently different, for this to be possible.

Two descriptions of the shape of survival curves are discussed briefly with a minimum of mathematics (see Fig. 3.3). First, the multtarget model that was widely used for many years still has some merit (Fig. 3.3B). In this model, the survival curve is described in terms of an initial slope, $D_I$, resulting from single-event killing; a final slope, $D_F$, resulting from multiple-event killing; and some quantity (either $n$ or $D_q$) to represent the size or width of the shoulder of the curve. The quantities $D_I$ and $D_F$ are the reciprocals of the initial and final slopes. In each case, it is the dose required to reduce the fraction of surviving cells to 37% of its previous value. As illustrated in Figure 3.3B, $D_I$, the initial slope, is the dose required to reduce the fraction of surviving cells to 0.37 on the initial straight portion of the survival curve. The final slope, $D_F$, is the dose required to reduce survival from 0.1 to 0.037 or from 0.01 to 0.0037, and so on. Because the surviving fraction is on a logarithmic scale and the survival curve becomes straight at higher doses, the dose required to reduce the cell population by a given factor (to 0.37) is the same at all survival levels. It is, on average, the dose required to deliver one inactivating event per cell.
When the priest was handed the badly deformed infant who was to grow up to be the hunchback, he cradled him in his arms and said, “We will call him Quasimodo—he is almost a person!” Similarly, the quasithreshold dose is almost a threshold dose. It is defined as the dose at which the straight portion of the survival curve, extrapolated backward, cuts the dose axis drawn through a survival fraction of unity. A threshold dose is the dose below which the extrapolation number, \( n \), is a measure of the width of the shoulder. If \( n \) is large (e.g., 10 or 12), the survival curve has a broad shoulder. If \( n \) is small (e.g., 1.5–2), the shoulder of the curve is narrow. Another measure of shoulder width is the quasithreshold dose, shown as \( D_q \) in Figure 3.3. This sounds like a term invented by a committee, which in a sense it is. An easy way to remember its meaning is to think of the hunchback of Notre Dame. When the priest was handed the badly deformed infant who was to grow up to be the hunchback, he cradled him in his arms and said, “We will call him Quasimodo—he is almost a person!” Similarly, the quasithreshold dose is almost a threshold dose. It is defined as the dose at which the straight portion of the survival curve, extrapolated backward, cuts the dose axis drawn through a survival fraction of unity. A threshold dose is the dose below which

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**FIGURE 3.2** The cell culture technique used to generate a cell survival curve. Cells from a stock culture are prepared into a single-cell suspension by trypsinization, and the cell concentration is counted. Known numbers of cells are inoculated into petri dishes and irradiated. They are then allowed to grow until the surviving cells produce macroscopic colonies that can be counted readily. The number of cells per dish initially inoculated varies with the dose so that the number of colonies surviving is in the range that can be counted conveniently. Surviving fraction is the ratio of colonies produced to cells plated, with a correction necessary for plating efficiency (i.e., for the fact that not all cells plated grow into colonies, even in the absence of radiation).

- **no. cells seeded:**
  - 100
  - 400
  - 1,000
  - 10,000
- **X ray dose:**
  - 0 Gy
  - 2 Gy
  - 4 Gy
  - 6 Gy
- **incubate 1–2 weeks**
- **no. colonies counted:**
  - 90
  - 72
  - 36
  - 45
- **plating efficiency:**
  - 90%
  - —
  - —
  - —
- **surviving fraction:**
  - —
  - 0.2
  - 0.04
  - 0.005

---
the square of the dose. The notion of a component of cell inactivation that varies with the square of the dose introduces the concept of dual radiation action. This idea goes back to the early work with chromosomes in which many chromosome aberrations are clearly the result of two separate breaks. (Examples discussed in Chapter 2 are dicentrics, rings, and anaphase bridges, all of which are likely to be lethal to the cell.)

By this model, the expression for the cell survival curve is

\[
S = e^{-\alpha D - \beta D^2}
\]

in which \(S\) is the fraction of cells surviving a dose \(D\), and \(\alpha\) and \(\beta\) are constants. The components of cell killing that are proportional to dose and to the square of the dose are equal if

\[
\alpha D = \beta D^2
\]

or

\[
D = \alpha/\beta
\]

There is no effect. There is no dose below which radiation produces no effect, so there can be no true threshold dose; \(D_0\), the quasithreshold dose, is the closest thing.

At first sight, this might appear to be an awkward parameter, but in practice, it has certain merits that become apparent in subsequent discussion. The three parameters, \(n\), \(D_0\), and \(D_q\), are related by the expression

\[
\log_{10} n = D_q/D_0
\]

The linear-quadratic model has taken over as the model of choice to describe survival curves. It is a direct development of the relation used to describe exchange-type chromosome aberrations that are clearly the result of an interaction between two separate breaks. This is discussed in some detail in Chapter 2.

The linear-quadratic model, illustrated in Figure 3.3A, assumes that there are two components to cell killing by radiation, one that is proportional to dose and one that is proportional to the square of the dose. The notion of a component of cell inactivation that varies with the square of the dose introduces the concept of dual radiation action. This idea goes back to the early work with chromosomes in which many chromosome aberrations are clearly the result of two separate breaks. (Examples discussed in Chapter 2 are dicentrics, rings, and anaphase bridges, all of which are likely to be lethal to the cell.)

By this model, the expression for the cell survival curve is

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\[
\alpha D = \beta D^2
\]

or

\[
D = \alpha/\beta
\]

\(D_q\), the quasithreshold dose, is the closest thing.

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By this model, the expression for the cell survival curve is

\[
S = e^{-\alpha D - \beta D^2}
\]

in which \(S\) is the fraction of cells surviving a dose \(D\), and \(\alpha\) and \(\beta\) are constants. The components of cell killing that are proportional to dose and to the square of the dose are equal if

\[
\alpha D = \beta D^2
\]

or

\[
D = \alpha/\beta
\]
This is an important point that bears repeating: The linear and quadratic contributions to cell killing are equal at a dose that is equal to the ratio of $\alpha$ to $\beta$.

A characteristic of the linear-quadratic formulation is that the resultant cell survival curve is continuously bending; there is no final straight portion. This does not coincide with what is observed experimentally if survival curves are determined down to seven or more decades of cell killing in which case the dose–response relationship closely approximates to a straight line in a log-linear plot; that is, cell killing is an exponential function of dose. In the first decade or so of cell killing and up to any doses used as daily fractions in clinical radiotherapy, however, the linear-quadratic model is an adequate representation of the data. It has the distinct advantage of having only two adjustable parameters, $\alpha$ and $\beta$.

**MECHANISMS OF CELL KILLING**

**DNA as the Target**

Abundant evidence shows that the principal sensitive sites for radiation-induced cell lethality are located in the nucleus as opposed to the cytoplasm.

Early experiments with nonmammalian systems, such as frog eggs, amoebae, and algae, were designed so that either the cell nucleus or the cytoplasm could be irradiated selectively with a microbeam. The results indicated that the nucleus was much more radiosensitive than the cytoplasm.

Evidence for chromosomal DNA as the principal target for cell killing is circumstantial but overwhelming. There is evidence that the nuclear membrane may also be involved. Indeed, the one does not exclude the other because some portions of the DNA may be intimately involved with the membrane during some portions of the cell cycle.

The evidence implicating the chromosomes, specifically the DNA, as the primary target for radiation-induced lethality may be summarized as follows:

1. Cells are killed by radioactive tritiated thymidine incorporated into the DNA. The radiation dose results from short-range $\alpha$-particles and is therefore very localized.

2. Certain structural analogues of thymidine, particularly the halogenated pyrimidines, are incorporated selectively into DNA in place of thymidine if substituted in cell culture growth medium. This substitution dramatically increases the radiosensitivity of the mammalian cells to a degree that increases as a function of the amount of the incorporation. Substituted deoxyuridines, which are not incorporated into DNA, have no such effect on cellular radiosensitivity.

3. Factors that modify cell lethality, such as variation in the type of radiation, oxygen concentration, and dose rate, also affect the production of chromosome damage in a fashion qualitatively and quantitatively similar. This is at least prima facie evidence to indicate that damage to the chromosomes is implicated in cell lethality.

4. Early work showed a relationship between virus size and radiosensitivity; later work showed a better correlation with nucleic acid volume. The radiosensitivity of a wide range of plants has been correlated with the mean interphase chromosome volume, which is defined as the ratio of nuclear volume to chromosome number. The larger the mean chromosome volume, the greater the radiosensitivity.

**The Bystander Effect**

Generations of students in radiation biology have been taught that heritable biologic effects require direct damage to DNA; however, experiments in the last decade have demonstrated the existence of a bystander effect, defined as the induction of biologic effects in cells that are not directly traversed by a charged particle, but are in proximity to cells that are. Interest in this effect was sparked by the 1992 report of Nagasawa and Little that following a low dose of $\alpha$-particles, a larger proportion of cells showed biologic damage than was estimated to have been hit by an $\alpha$-particle; specifically, 30% of the cells showed an increase in sister chromatid exchanges even though less than 1% were calculated to have undergone a nuclear traversal. The number of cells hit was arrived at by a calculation based on the fluence of $\alpha$-particles and the cross-sectional area of the cell nucleus. The conclusion was thus of a statistical nature because it was not possible to know on an individual basis which cells were hit and which were not.

This observation has been extended by the use of sophisticated single-particle microbeams,
which make it possible to deliver a known number of particles through the nucleus of specific cells, whereas biologic effects can be studied in unirradiated close neighbors. Most microbeam studies have used α-particles because it is easier to focus them accurately, but a bystander effect has also been shown for protons and soft x-rays. Using single-particle microbeams, a bystander effect has been demonstrated for chromosomal aberrations, cell killing, mutation, oncogenic transformation, and alteration of gene expression. The effect is most pronounced when the bystander cells are in gap-junction communication with the irradiated cells. For example, up to 30% of bystander cells can be killed in this situation. The bystander effect is much smaller when cell monolayers are sparsely seeded so that cells are separated by several hundred micrometers. In this situation, 5% to 10% of bystander cells are killed, the effect being due, presumably, to cytotoxic molecules released into the medium. The existence of the bystander effect indicates that the target for radiation damage is larger than the nucleus and, indeed, larger than the cell itself. Its importance is primarily at low doses, where not all cells are “hit,” and it may have important implications in risk estimation.

In addition to the experiments described previously involving sophisticated single-particle microbeams, there is a body of data involving the transfer of medium from irradiated cells that results in a biologic effect (cell killing) when added to unirradiated cells. These studies, which also evoke the term bystander effect, suggest that irradiated cells secrete a molecule into the medium that is capable of killing cells when that medium is transferred onto unirradiated cells. Most bystander experiments involving medium transfer have used low-LET x- or γ-rays.

**Apoptotic and Mitotic Death**

Apoptosis was first described by Kerr and colleagues as a particular set of changes at the microscopic level associated with cell death. The word *apoptosis* is derived from the Greek word meaning “falling off,” as in petals from flowers or leaves from trees. Apoptosis, or programmed cell death, is common in embryonic development in which some tissues become obsolete. It is the mechanism, for example, by which tadpoles lose their tails.

This form of cell death is characterized by a stereotyped sequence of morphologic events. One of the earliest steps a cell takes if it is committed to die in a tissue is to cease communicating with its neighbors. This is evident as the dying cell rounds up and detaches from its neighbors. Condensation of the chromatin at the nuclear membrane and fragmentation of the nucleus are then evident. The cell shrinks because of cytoplasmic condensation, resulting from cross-linking of proteins and loss of water. Eventually, the cell separates into several membrane-bound fragments of differing sizes termed **apoptotic bodies**, which may contain cytoplasm only or nuclear fragments. The morphologic hallmark of apoptosis is the condensation of the nuclear chromatin in either crescents around the periphery of the nucleus or a group of spheric fragments.

Double-strand breaks (DSBs) occur in the linker regions between nucleosomes, producing DNA fragments that are multiples of approximately 185 base pairs. These fragments result in the characteristic ladders seen in gels. In contrast, necrosis causes a diffuse “smear” of DNA in gels. Apoptosis occurs in normal tissues, as described previously, and also can be induced in some normal tissues and in some tumors by radiation.

As a mode of radiation-induced cell death, apoptosis is highly cell-type dependent. Hemopoietic and lymphoid cells are particularly prone to rapid radiation-induced cell death by the apoptotic pathway. In most tumor cells, mitotic cell death is at least as important as apoptosis, and in some cases, it is the only mode of cell death. Several genes appear to be involved. First, apoptosis after radiation seems commonly to be a *p53*-dependent process; Bcl-2 is a suppressor of apoptosis.

The most common form of cell death from radiation is mitotic death: Cells die attempting to divide because of damaged chromosomes. Death may occur in the first or a subsequent division following irradiation. Many authors have reported a close quantitative relationship between cell killing and the induction of specific chromosomal aberrations. The results of one of the most elegant studies by Cornforth and Bedford are shown in Figure 3.4. It should be noted that these experiments were carried out in a cell line where apoptosis is not observed. The log of the surviving fraction is plotted against the average number of putative “lethal” aberrations per cell, that is, asymmetric exchange-type aberrations...
chromosome aberrations is carried over to the cell survival curve. Autophagic Cell Death

Autophagy is literally defined as a self-digestive process that uses lysosomal degradation of long-lived proteins and organelles to restore or maintain cellular homeostasis. Autophagy is evolutionarily conserved and is considered a dynamic process that involves a unique series of steps, of which the sequestration of portions of the cytoplasm and organelles in a double-membrane vesicle, called an autophagosome, is a hallmark characteristic. These autophagosomes ultimately fuse with lysosomes, where protein and organelles are degraded and reprocessed. Autophagosomes then fuse with lysosomes, which acidify as they mature to become autolysosomes in a step called autophagic flux. Autophagy is a multistep process that is genetically regulated by a unique set of genes termed autophagy-related genes (Atgs). These Atgs were first discovered in yeast, and approximately 30 Atg orthologs have been identified in mammals that include two ubiquitin-like conjugation systems: the Atg12-Atg5 and the Atg8 (LC-3)- phosphatidylethanolamine (PE). These systems are required for the elongation of the autophagosomal membrane. However, the proteins and trafficking mechanisms involved in the autophagosomal maturation step are not completely understood.
chemotherapy and radiotherapy. Previous studies have reported that the metabolic stress observed in human tumors leads cancer cells to acquire resistance to apoptosis and to stimulate autophagy to maintain energy demand and prevent necrosis. Furthermore, chemotherapeutic agents and radiotherapy have been reported to induce autophagy and autophagic cell death. Although the mechanism underlying this form of cell death is unclear, accumulation of autophagosomes in response to chemotherapy or radiotherapy suggests that this type of cell death is associated with an inhibition of the maturation and degradation process. The signals for the induction of autophagy by radiotherapy are still under investigation, but may involve signaling from the endoplasmic reticulum, particularly, the protein kinase-like endoplasmic reticulum kinase (PERK), which is described in more detail in Chapter 26. Induction of endoplasmic reticulum stress in cells that have lost their ability to die by apoptosis when exposed to radiation results in radiosensitization. This data suggests that the combination of endoplasmic stress-inducing agents and ionizing radiation could enhance cell killing by inducing autophagic cell death. Thus, in the regulation of cancer, autophagy should be considered a new target for anticancer therapy.

**Senescence**

Cellular senescence has emerged as a programmed cellular stress response that represents a unique response to the accumulation of damage to a cell. Whether through the shortening of telomeres associated with a high number of cell divisions, activation of oncogenes, or DNA damage caused by oxidative stress, induction of senescence in primary cells leads to an irreversible cell cycle arrest that is almost invariably characterized by the activation of the p53 and retinoblastoma (Rb) proteins, and is associated with chromatin modifications that result in the silencing of genes necessary to promote transition from the G1 to S phase of the cell cycle. For these reasons, senescence has been classified as a tumor suppressor mechanism that prevents excessive cellular divisions in response to inappropriate growth signals or division of cells that have accumulated DNA damage. Because it is a genetically regulated process that involves the p53 and Rb proteins, it is understandable that loss of p53 and Rb control through gene mutation will result in...
in loss of senescence in response to DNA damage. However, although tumor cells typically lose their ability to undergo senescence in response to DNA damage caused by mutations in the p53 and Rb pathways, normal cells do not lose this ability. The induction of senescence in normal tissue is important to consider because these cells are still metabolically active, but reproductively inhibited. This is best exemplified by fibroblasts that, when irradiated in cell culture, stay attached to plates for weeks, but never divide. However, they are able to secrete growth factors and mitogens that promote the growth of tumor cells. Therefore, senescence results in a permanent cell cycle arrest, but will not eliminate the mitogenic or cytokine contribution of the arrested cell that could ultimately promote tumor regrowth.

**SURVIVAL CURVES FOR VARIOUS MAMMALIAN CELLS IN CULTURE**

Survival curves have been measured for many established cell lines grown in culture. These cell lines have been derived from the tissues of humans or other mammals such as small rodents. In some cases, the parent tissue has been neoplastic; in other cases, it has been normal. The first in vitro survival curve for mammalian cells irradiated with x-rays is shown in Figure 3.6. All mammalian cells studied to date, normal or malignant, regardless of their species of origin, exhibit x-ray survival curves similar to those in Figure 3.6; there is an initial shoulder followed by a portion that tends to become straight on a log-linear plot. The size of the initial shoulder is extremely variable. For some cell lines, the survival curve appears to bend continuously, so that the linear-quadratic relationship is a better fit and $n$ has no meaning. The $D_0$ of the x-ray survival curves for most cells cultured in vitro falls in the range of 1 to 2 Gy. The exceptions are cells from patients with cancer-prone syndromes such as ataxia-telangiectasia (AT); these cells are much more sensitive to ionizing radiations, with a $D_0$ for x-rays of about 0.5 Gy. This in vitro sensitivity correlates with a hypersensitivity to radiotherapy found in these patients.

The first in vitro survival curve was reported in 1956 and generated great excitement in the field of radiobiology. It was thought that at last, with a quantitative system available to relate absorbed dose with surviving fraction of cells, great strides would be made in understanding the effect of ionizing radiation on biologic materials. In particular, it was anticipated that significant contributions would be made toward understanding radiotherapeutic practice. This enthusiasm was not shared by everyone. Some researchers were skeptical that these in vitro techniques, which

![Figure 3.6](image-url)
involved growing cells in petri dishes in very artificial conditions, would ever benefit clinical radiotherapy. The fears of these skeptics were eloquently voiced by F.G. Spear in the MacKenzie Davidson Memorial Lecture given to the British Institute of Radiology in 1957:

An isolated cell in vitro does not necessarily behave as it would have done if left in vivo in normal association with cells of other types. Its reactions to various stimuli, including radiations, however interesting and important in themselves, may indeed be no more typical of its behavior in the parent tissue than Robinson Crusoe on his desert island was representative of social life in York in the mid-seventeenth century.

The appropriate answer to this charge was given by David Gould, then professor of radiology at the University of Colorado. He pointed out that the in vitro culture technique measured the reproductive integrity of cells and that there was no reason to suppose that Robinson Crusoe’s reproductive integrity was any different on his desert island from what it would have been had he remained in York; all that Robinson Crusoe lacked was the opportunity. The opportunity to reproduce to the limit of their capability is afforded to cells cultured in vitro if they find themselves in the petri dish, with temperature and humidity controlled and with an abundant supply of nutrients.

At the time, it required a certain amount of faith and optimism to believe that survival curves determined with the in vitro technique could be applied to the complex in vivo situation. Such faith and optimism were completely vindicated, however, by subsequent events. When techniques became available to measure cell survival in vivo, the parameters of the dose–response relationships were shown to be similar to those in vitro.

In more recent years, extensive studies have been made of the radiosensitivity of cells of human origin, both normal and malignant, grown and irradiated in culture. In general, cells from a given normal tissue show a narrow range of radiosensitivities if many hundreds of people are studied (Fig. 3.7). By contrast, cells from human tumors show a very broad range of \( D_0 \) values; some cells, such as those from squamous carcinoma cells, tend to be more radioresistant, whereas sarcomas are somewhat more radiosensitive. Each tumor type, however, has a broad spectrum of radiosensitivities that tend to overlap. Tumor cells bracket the radiosensitivity of cells from normal tissues; that is, some are more sensitive, and others are more resistant.

**SURVIVAL CURVE SHAPE AND MECHANISMS OF CELL DEATH**

Mammalian cells cultured in vitro vary considerably in their sensitivity to killing by radiation. This is illustrated in Figure 3.8A, which includes survival curves for asynchronous cultures of...
mouse tumor cells (EMT6) as well as for six cell lines derived from human tumors.

Asynchronous EMT6 cells are the most radioresistant, followed closely by glioblastoma cells of human origin; thereafter, radiosensitivity increases, with two neuroblastoma cell lines being the most sensitive. Although asynchronous cells show this wide range of sensitivities to radiation, it is a remarkable finding that mitotic cells from all of these cell lines have essentially the same radiosensitivity. The implication of this is that if the chromosomes are condensed during mitosis, all cell lines have the same radiosensitivity governed simply by DNA content; but in interphase, the radiosensitivity differs because of different conformations of the DNA. Another interesting observation comes from a comparison of the survival curves in Figure 3.8A with the DNA laddering in Figure 3.8B.
Characteristic laddering is indicative of programmed cell death or apoptosis during which the DNA breaks up into discrete lengths as previously described. Comparing Figure 3.8A and B, it is evident that there is a close and impressive correlation between radiosensitivity and the importance of apoptosis. The most radioresistant cell lines, which have broad shoulders to their survival curves, show no evidence of apoptosis; the most radiosensitive, for which survival is an exponential function of dose, show clear DNA laddering as an indication of apoptosis. The increased clarity of the laddering correlates with increasing radiosensitivity together with a smaller and smaller shoulder to the survival curve. Many of the established cell lines that have been cultured in vitro for many years, and with which many of the basic principles of radiation biology were demonstrated, show no apoptotic death and have an abrogated p53 function. Continued culture in vitro appears often to select for cells with this characteristic.

Mitotic death results (principally) from exchange-type chromosomal aberrations; the associated cell survival curve, therefore, is curved in a log-linear plot, with a broad initial shoulder. As is shown subsequently here, it is also characterized by a substantial dose-rate effect. Apoptotic death results from mechanisms that are not yet clearly understood, but the associated cell survival curve appears to be a straight line on a log-linear plot—that is, survival is an exponential function of dose. In addition, there appears to be little or no dose-rate effect, although data are sparse on this point.

Although there are some cell lines in which mitotic death dominates and others in which apoptosis is the rule, most cell lines fall somewhere in between, with contributions from both mitotic and apoptotic death following a radiation exposure, in varying proportions. It has been proposed that the dose–response relationship be described by the following relation:

\[ S = e^{-(\alpha_M + \alpha_A)D - \beta_M D^2} \]

in which \( S \) is the fraction of cells surviving a dose \( D \), \( \alpha_M \) and \( \alpha_A \) describe the contributions to cell killing from mitotic and apoptotic death that are linear functions of dose, and \( \beta_M \) describes the contribution to mitotic death that varies with the square of the dose.

### ONCOGENES AND RADIORESISTANCE

Numerous reports have appeared in the literature that transfection of activated oncogenes into cells cultured in vitro increases their radioresistance, as defined by clonogenic survival. Reports include the transfection of activated N-ras, raf, or ras + myc, a combination that is particularly effective in transforming primary explants of rodent embryo cells to a malignant state. Results, however, are equivocal and variable. The change of radiosensitivity did not correlate with cell cycle distribution or DNA DSBs or their repair; the best correlation was with the length of the G2 phase delay induced by radiation. It is by no means clear that oncogene expression is directly involved in the induction of radioresistance, and it is far less clear that oncogenes play any major role in radioresistance in human tumors.

### GENETIC CONTROL OF RADIOSENSITIVITY

The molecular biology of repair processes in lower organisms, such as yeast and bacteria, has been studied extensively. In several instances, a dramatically radioresistant mutant can result from a mutation in a single gene that functions as a repair or checkpoint gene. In mammalian cells, the situation is much more complicated, and it would appear that a large number of genes may be involved in determining radiosensitivity. Many radiosensitive mutants have been isolated from cell lines maintained in the laboratory, especially rodent cell systems. In many but not all cases, their sensitivity to cell killing by radiation has been related to their greatly reduced ability to repair DNA DSBs. Examples of these genes are Ku 80, Ku 70, and XRCC7. The first of these two genes are involved in DNA-dependent kinase activity that binds to the free ends at the site of a DSB, so that if they are defective, repair of DSBs is prejudiced. The third gene codes for a protein that is defective in mice with the “severe combined immune deficiency syndrome” that are sensitive to radiation.

Some patients who show an abnormally severe normal tissue reaction to radiation therapy exhibit the traits of specific inherited syndromes. These are listed in Table 3.1 and discussed in more detail in Chapter 18. The most striking example is AT. Fibroblasts taken from patients...
It has been well accepted that the radiosensitivity of cells changes as they undergo differentiation. If this is true, then cell death in tumors exposed to ionizing radiation in tumors should correlate with the elimination of tumor stem cells and survival of the more differentiated tumor cells that have lost their renewal capability. However, this has not what has been recently reported in the literature. In fact, cancer stem cells may be more resistant to radiation than their more differentiated counterparts. Mechanistically, cancer stem cells appear to have lower levels of reactive oxygen species because of increased levels of free radical scavengers. If this is the case, then the same increased levels of free radical scavengers that protect the cancer stem cells from the metabolic consequences of reactive oxygen species during their growth also protect them when exposed to ionizing radiation. The radiosensitivity of cancer stem cells can be increased if they are first treated with inhibitors of free radical scavengers before radiation exposure.

These rather provocative results will definitely require further investigation to determine whether all cancer stem cells possess high levels of free radical scavengers both in experimental tumor systems as well as in human biopsies. If this is the case, then a potential means of radiosensitizing these stem cells through the targeting of free radical scavengers could be tested.

### EFFECTIVE SURVIVAL CURVE FOR A MULTIFRACTION REGIMEN

Because multifraction regimens are used most often in clinical radiotherapy, it is frequently useful to think in terms of an effective survival curve.

If a radiation dose is delivered in a series of equal fractions, separated by sufficient time for repair of sublethal damage to occur between doses, the effective dose-survival curve becomes an exponential function of dose. The shoulder of the survival curve is repeated many times, so that the effective survival curve is a straight line from the origin through a point on the single-dose survival curve corresponding to the daily dose fraction. This is illustrated in Figure 3.9. The effective survival curve is an exponential function of dose whether the single-dose survival curve has a constant terminal slope (as shown) or is continuously bending, as implied by the linear-quadratic relation. The \( D_0 \) of the effective
survival curve (i.e., the reciprocal of the slope), defined to be the dose required to reduce the fraction of cells surviving to 37%, has a value close to 3 Gy for cells of human origin. This is an average value and can differ significantly for different tumor types.

For calculation purposes, it is often useful to use the $D_{10}$, the dose required to kill 90% of the population. For example:

$$D_{10} = 2.3 \times D_0$$

in which 2.3 is the natural logarithm of 10.

### CALCULATIONS OF TUMOR CELL KILL

The concept outlined previously of an effective survival curve for a multifraction radiation treatment may be used to perform simple calculations of tumor cell kill after radiotherapy. Although such calculations are greatly oversimplified, they are nevertheless instructive. Four examples are given here.

#### Problem 1

A tumor consists of $10^8$ clonogenic cells. The effective dose-response curve given in daily dose fractions of 2 Gy has no shoulder and a $D_0$ of 3 Gy. What total dose is required to give a 90% chance of tumor cure?

**Answer**

To give a 90% probability of tumor control in a tumor containing $10^8$ cells requires a cellular depopulation of $10^{-9}$. The dose resulting in one decade of cell killing ($D_{10}$) is related to the $D_0$ by the expression $D_{10} = 2.3 \times D_0$.

$$D_{10} = 2.3 \times D_0 = 2.3 \times 3 = 6.9 \text{ Gy}$$

The total dose for 9 decades of cell killing, therefore, is $9 \times 6.9 = 62.1 \text{ Gy}$.

#### Problem 2

Suppose that, in the previous example, the clonogenic cells underwent three cell doublings during treatment. About what total dose would be required to achieve the same probability of tumor control?

**Answer**

Three cell doublings would increase the cell number by

$$2 \times 2 \times 2 = 8$$

Consequently, about one extra decade of cell killing would be required, corresponding to
an additional dose of 6.9 Gy. The total dose is $62.1 + 6.9 = 69$ Gy.

**Problem 3**

During the course of radiotherapy, a tumor containing $10^9$ cells receives 40 Gy. If the $D_0$ is 2.2 Gy, how many tumor cells will be left?

**Answer**

If the $D_0$ is 2.2 Gy, the $D_{10}$ is given by

$$D_{10} = 2.3 \times D_0 = 2.3 \times 2.2 = 5 \text{ Gy}$$

Because the total dose is 40 Gy, the number of decades of cell killing is $40/5 = 8$. The number of cells remaining is $10^9 \times 10^{-8} = 10$.

**Problem 4**

If $10^7$ cells were irradiated according to single-hit kinetics so that the average number of hits per cell is one, how many cells would survive?

**Answer**

A dose that gives an average of one hit per cell is the $D_0$, that is, the dose that on the exponential region of the survival curve reduces the number of survivors to 37%. The number of surviving cells is therefore

$$10^7 \times \frac{37}{100} = 3.7 \times 10^6$$

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**THE RADIOSENSITIVITY OF MAMMALIAN CELLS COMPARED WITH MICROORGANISMS**

The final illustration in this chapter (Fig. 3.10) is a compilation from the literature of survival data for many types of cells. The steepest dose–response relationship (curve A) is an average curve for mammalian cells; it is evident that they are exquisitely radiosensitive compared with microorganisms. The bacterium *Escherichia coli* is more resistant, yeast is more resistant still, and the most resistant is *Micrococcus radiodurans*, which shows no significant cell killing even after a dose of 1,000 Gy. There are several important points to be made from this:

1. The dominant factor that accounts for this huge range of radiosensitivities is the DNA content. Mammalian cells are sensitive because they have a large DNA content, which represents a large target for radiation damage.

2. DNA content is not the whole story, however. *E. coli* and *E. coli* B/r have the same DNA content but differ in radiosensitivity because B/r has a mutant and more efficient DNA repair system. In higher organisms, mode of cell death—that is, apoptotic versus mitotic—also affects radiosensitivity.

3. Figure 3.10 explains why, if radiation is used as a method of sterilization, doses of the order of 20,000 Gy are necessary. Even if objects are socially clean, such huge doses are necessary to reduce the population of contaminating microorganisms because of their extreme radioresistance.

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**SUMMARY OF PERTINENT CONCLUSIONS**

- Cells from tumors and many normal regenerative tissues grow and form colonies *in vitro*.
- Fresh explants of normal tissues often grow well in culture for a few weeks before they...
A cell is said to have retained its reproductive integrity if it is capable of sustained proliferation, that is, if it can grow into a macroscopic colony.

A survivor that has retained its reproductive integrity is said to be clonogenic.

The percentage of untreated cells seeded that grow into macroscopic colonies is known as the plating efficiency. Thus:

\[
\text{PE} = \frac{\text{Number of colonies counted}}{\text{Number of cells seeded}} \times 100
\]

The plating efficiency may be close to 100% for some established cell lines but 1% or less for fresh explants of human cells.

The fraction of cells surviving a given dose is determined by counting the number of macroscopic colonies as a fraction of the number of cells seeded. Allowance must be made for the plating efficiency. Thus:

\[
\text{SF} = \frac{\text{Number of colonies counted}}{\text{Number of cells seeded}} \times \left(\frac{\text{PE}}{100}\right)
\]

A cell survival curve is the relationship between the fraction of cells retaining their reproductive integrity and the absorbed dose.

Conventionally, surviving fraction on a logarithmic scale is plotted on the ordinate against dose on the abscissa. The shape of the survival curve is important.

The cell survival curve for α-particles and low-energy neutrons (densely ionizing radiations) is a straight line on a log-linear plot; that is, survival approximates to an exponential function of dose.

The cell survival curve for x- or γ-rays (sparsely ionizing radiations) has an initial slope, followed by a bending region or shoulder, after which it tends to straighten again at higher doses.

Survival data are adequately fitted by many models and theories. The data are never sufficiently precise, nor are the models sufficiently different for experimental results to discriminate among models.

For the first one or two decades of survival and up to doses used in single fractions in radiotherapy, survival data are adequately represented by the linear-quadratic relationship

\[
S = e^{-\alpha D - \beta D^2}
\]

in which \(S\) is the fraction of cells surviving a dose \(D\) and \(\alpha\) and \(\beta\) are constants representing the linear and quadratic components of cell killing.

The initial slope of the cell survival curve is determined by \(\alpha\); the quadratic component of cell killing (\(\beta\)) causes the curve to bend at higher doses.

The ratio \(\alpha/\beta\) is the dose at which linear and quadratic components of cell killing are equal.

There is good evidence that the nucleus, specifically the DNA, is the principal target for radiation-induced cell lethality. Membrane damage also may be a factor.

Following exposure to radiation, cells may die attempting the next or a subsequent mitosis (mitotic death), or they may die programmed cell deaths (apoptotic death).

In cells that die a mitotic death, there is a one-to-one correlation between cell survival and the average number of putative “lethal” chromosomal aberrations per cell, that is, asymmetric exchange-type aberrations such as dicentrics and rings.

Cells that die an apoptotic death follow a stereotyped sequence of morphologic events, culminating in the breaking up of the DNA into fragments that are multiples of 185 base pairs; this leads to the characteristic DNA laddering seen in gels.

In some cell types (such as lymphoid cells), apoptotic death is dominant following irradiation. Survival is then an exponential function of dose; that is, the survival curve is straight and shoulderless on the usual log-linear plot. There is also no dose-rate effect.

In some cell types (such as Chinese hamster ovary [CHO] or V79 cells in culture), mitotic death is dominant following irradiation. Survival is then a linear-quadratic function of dose; that is, the survival curve has a shoulder on the usual log-linear plot. There is usually a large dose-rate effect.

Many cell populations die both mitotic and apoptotic deaths. There is, in general, a correlation between the importance of apoptosis and radiosensitivity. If apoptosis
is dominant, cells are radiosensitive; if apoptosis is absent, cells are radioresistant.

- In addition to mitotic and apoptotic cell death, cells exposed to ionizing radiation can die by autophagic cell death or enter senescence, which is a permanent type of growth arrest.

- Cells cultured from different tumors in humans show a broad range of radiosensitivities that bracket the sensitivity of normal cells from different people.

- There is some evidence in cells cultured in vitro that transfection of activated oncogenes in cells increases their radioresistance. It is not clear that oncogenes play a role in the radioresistance of human tumors in vivo.

- Several genes that influence the radiosensitivity of mammalian cells have been identified.

- If these genes are defective, the repair of DSBs is often prejudiced.

- Several human syndromes have been found to be associated with radiosensitivity; AT is the best example.

- There is often a link between sensitivity to killing by radiation and predisposition to cancer.

- Cancer stem cells may be more radioresistant than their differentiated tumor cell counterparts because of increased levels of reactive oxygen-specific scavengers.

- The effective survival curve for a multifraction regimen is an exponential function of dose: A straight line from the origin through a point on the single-dose survival curve corresponding to the daily dose fraction.

- The average value of the effective $D_0$ for the multifraction survival curve for human cells is about 3 Gy.

- The $D_{10}$, the dose resulting in one decade of cell killing, is related to the $D_0$ by the expression:

$$D_{10} = 2.3 \times D_0$$

- Calculations of tumor cell kill can be performed for fractionated clinical radiotherapy regimens using the concept of effective survival curve.

- Mammalian cells are exquisitely radiosensitive compared with microorganisms such as bacteria and yeast, principally because of their larger DNA content, which represents a bigger "target" for radiation damage.

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Mammalian cells propagate and proliferate by mitosis. When a cell divides, two progeny cells are produced, each of which carries a chromosome complement identical to that of the parent cell. After an interval of time has elapsed, each of the progeny may undergo a further division. The time between successive divisions is known as the mitotic cycle time or, as it is commonly called, the cell cycle time (TC).

If a population of dividing cells is observed with a conventional light microscope, the only event in the entire cell cycle that can be identified and distinguished is mitosis, or division itself. Just before the cell divides to form two progeny cells, the chromosomes (which are diffuse and scattered in the nucleus in the period between mitoses) condense into clearly distinguishable forms. In addition, in monolayer cultures of cells just before mitosis, the cells round up and become loosely attached to the surface of the culture vessel. This whole process of mitosis—in preparation for which the cell rounds up, the chromosome material condenses and the cell divides into two and then stretches out again and attaches to the surface of the culture vessel—lasts only about 1 hour. The remainder of the cell cycle, the interphase, occupies all of the intermitotic period. No events of interest can be identified with a conventional microscope during this time.

Because cell division is a cyclic phenomenon repeated in each generation of the cells, it is usual to represent it as a circle, as shown in Figure 4.1. The circumference of the circle represents the full mitotic cycle time for the cells (TC); the period of mitosis is represented by M. The remainder of the cell cycle can be further subdivided by using some marker of DNA synthesis. The original technique was autoradiography, introduced by Howard and Pelc in 1953.

The basis of the technique, illustrated in Figure 4.2, is to feed the cells thymidine, a basic building block used for making DNA, which has been labeled with radioactive tritium (H-TdR). Cells that are actively synthesizing new DNA as part of the process of replicating their chromosome complements incorporate the radioactive thymidine. Then the surplus radioactive thymidine is flushed from the system, the cells are fixed and stained so that they may be viewed easily, and the preparation of cells is coated with a very thin layer of nuclear (photographic) emulsion. β-particles from cells that have incorporated radioactive thymidine pass through the nuclear emulsion and produce a latent image. When the emulsion is subsequently developed and fixed, the area through which a β-particle has passed appears as a black spot. It is then a comparatively simple matter to view the preparation of cells and to observe that some of the cells have black spots or “grains” over them, which indicates that they were actively synthesizing DNA at the time radioactive thymidine was made available. Other cells do not have any grains over their nuclei; this is interpreted to mean that the cells were not actively making DNA when the radioactive label was made available to them. Examples of labeled cells are shown in Figure 4.3. If the cells...
are allowed to grow for some time after labeling with tritiated thymidine so that they move into mitosis before being fixed, stained, and autoradiographed, then a labeled mitotic cell may be observed (Fig. 4.3A).

The use of tritiated thymidine to identify cells in the DNA synthetic phase (S) has been replaced largely by the use of 5-bromodeoxyuridine, which differs from thymidine only by the substitution of a bromine atom for a methyl group. If this halogenated pyrimidine is fed to the cells, it is incorporated into DNA in place of thymidine, and its presence can be detected by using an appropriate stain (see Fig. 4.3B). Cells that have incorporated bromodeoxyuridine appear darkly stained a bright purple color. To identify cells that are in S phase and have incorporated bromodeoxyuridine even more readily, one can use a fluorochrome-tagged G2 G1

\( \text{FIGURE 4.1} \) The stages of the mitotic cycle for actively growing mammalian cells. M, mitosis; S, DNA synthetic phase; G1 and G2, “gaps,” or periods of apparent inactivity between the major discernible events in the cycle.

\( \text{FIGURE 4.2} \) Cell-labeling techniques. Top panels: The principle of autoradiography, which may be applied to cells in culture growing as a monolayer on a glass microscope slide or to thin sections cut from a tumor or normal tissue. Cells in the DNA synthetic phase (S) take up tritiated thymidine. After the cells are fixed and stained so that they are visible by light microscopy, they are covered with a layer of nuclear (photographic) emulsion and left for several weeks in a cool refrigerator. As \( \beta \)-particles from the tritiated thymidine pass through the emulsion, they form latent images that appear as black grains when the emulsion is subsequently developed and fixed. If cells are stained and autoradiographed immediately after incorporation of the tritiated thymidine, cells that are labeled are in S phase (top middle panel). If staining and autoradiography are delayed for 6 to 8 hours after the pulse label, some cells may move from S to M, and labeled mitotic cells are observed (top right panel). The lengths of the various phases of the cycle can be determined in this way. Bottom panels: The principle of cell cycle analysis using 5-bromodeoxyuridine as the DNA precursor instead of radioactively labeled thymidine. The bromodeoxyuridine is incorporated into cells in S. It can be recognized by the use of a Giemsa stain (which is purple) or a monoclonal antibody to bromodeoxyuridine-substituted DNA. The antibody is tagged with a fluorescing dye (e.g., fluorescein), which shows up bright green under a fluorescence microscope. If cells are stained immediately after labeling with bromodeoxyuridine, those staining darkly are in S phase (bottom middle panel). If staining is delayed for 6 to 8 hours, cells incorporating bromodeoxyuridine may move from S to M, and a darkly staining mitotic cell is seen (bottom right panel). (Courtesy of Dr. Charles Geard.)
There is an interval between mitosis and DNA synthesis in which no label is incorporated. This first “gap” in activity was named G1 by Howard and Pelc, and the nomenclature is used today. After DNA synthesis has been completed, there is a second gap before mitosis, G2.

All proliferating mammalian cells, whether in culture or growing normally in a tissue, have a cycle of mitosis (M), followed by G1, S, and G2, after which mitosis occurs again. The relative lengths of these various constituent parts of the cell cycle vary according to the particular cells studied. If cells stop progressing through the cycle (i.e., are arrested), they are said to be in G0 (see Fig. 4.4).

The use of bromodeoxyuridine has two advantages over conventional autoradiography using tritiated thymidine. First, it does not involve radioactive material. Second, it greatly shortens the time to produce a result because if cells are coated with emulsion to produce an autoradiograph, they must be stored in a refrigerator for about a month to allow β-particles from the incorporated tritium to produce a latent image in the emulsion.

By using either of these techniques, it can be shown that cells synthesize DNA only during a discrete well-defined fraction of the cycle, the S phase. There is an interval between mitosis and DNA synthesis in which no label is incorporated. This first “gap” in activity was named G1 by Howard and Pelc, and the nomenclature is used today. After DNA synthesis has been completed, there is a second gap before mitosis, G2.

All proliferating mammalian cells, whether in culture or growing normally in a tissue, have a cycle of mitosis (M), followed by G1, S, and G2, after which mitosis occurs again. The relative lengths of these various constituent parts of the cell cycle vary according to the particular cells studied. If cells stop progressing through the cycle (i.e., are arrested), they are said to be in G0 (see Fig. 4.4).

The characteristics of two cell lines commonly used for in vitro culture are summarized in Table 4.1. HeLa cells have a total cell cycle time of about 24 hours, which is more than double that of the Chinese hamster cell, which has a cell cycle time of about 11 hours. Mitosis lasts only a relatively short time, about 1 hour, and is not very different for those two cell lines or for most others. The S phase is 8 hours for HeLa cells and 6 hours for hamster cells; in all cell lines studied...
in culture or growing in vivo, the S phase never exceeds about 15 hours. The G2 period is very similar in HeLa and hamster cells; in fact, the difference in the total cell cycle time between these two cell lines is accounted for almost entirely by the difference in the length of the G1 period.

This is an important point: The difference among mammalian cell cycle times in different circumstances, varying from about 10 hours for a hamster cell grown in culture to hundreds of hours for stem cells in some self-renewal tissues, is the result of a dramatic variation in the length of the G1 period. The remaining components of the cell cycle (M, S, and G2) vary comparatively little among different cells in different circumstances.

The description of the principal phases of the cell cycle (M, G1, S, and G2) dates from Howard and Pelc in 1953, as previously discussed. During a complete cell cycle, the cell must accurately replicate the DNA once during S phase and distribute an identical set of chromosomes equally to two progeny cells during M phase. In recent years, we have learned much more about the mechanisms by which the cycle is regulated in eukaryotic cells. Regulation occurs by the periodic activation of different members of the cyclin-dependent kinase (Cdk) family. In its active form, each Cdk is complexed with a particular cyclin. Different Cdk–cyclin complexes are required to phosphorylate several protein substrates that drive key events, including the initiation of DNA replication and the onset of mitosis. Cdk–cyclin complexes are also vital in preventing the initiation of a cell cycle event at the wrong time.

Extensive regulation of Cdk–cyclin activity by several transcriptional and posttranscriptional mechanisms ensures perfect timing and coordination of cell cycle events. The Cdk catalytic subunit by itself is inactive, requiring association with a cyclin subunit and phosphorylation of a key threonine residue to become fully active. The Cdk–cyclin complex is reversibly inactivated either by phosphorylation on a tyrosine residue located in the adenosine triphosphate–binding domain, or by association with Cdk inhibitory proteins. After the completion of the cell cycle transition, the complex is inactivated irreversibly by ubiquitin-mediated degradation of the cyclin subunit.

Entry into S phase is controlled by Cdns that are sequentially regulated by cyclins D, E, and A.itable 4.1 Phases of the Cell Cycle for Two Commonly Used Cell Lines Cultured In Vitro

<table>
<thead>
<tr>
<th>PIase of the Cell Cycle for Two Commonly Used Cell Lines Cultured In Vitro</th>
<th>Hamster Cells, h</th>
<th>HeLa Cells, h</th>
</tr>
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<tbody>
<tr>
<td>Tc</td>
<td>11</td>
<td>24</td>
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<tr>
<td>Tm</td>
<td>1</td>
<td>1</td>
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<tr>
<td>Ts</td>
<td>6</td>
<td>8</td>
</tr>
<tr>
<td>Tg1</td>
<td>11</td>
<td></td>
</tr>
<tr>
<td>Tg2</td>
<td>3</td>
<td>4</td>
</tr>
</tbody>
</table>
D-type cyclins act as growth factor sensors, with their expression depending more on the extracellular cues than on the cell’s position in the cycle. Mitogenic stimulation governs both their synthesis and complex formation with Cdk4 and Cdk6, and catalytic activity of the assembled complexes persists through the cycle as long as mitogenic stimulation continues. Cyclin E expression in proliferating cells is normally periodic and maximal at the G1/S transition, and throughout this interval, it enters into active complexes with its catalytic partner, Cdk2. Figure 4.4 illustrates this view of the cell cycle and its regulation. This is, in essence, an update of Figure 4.1 and is discussed in more detail in Chapter 18.

- SYNCHRONOUSLY DIVIDING CELL CULTURES

In the discussion of survival curves in Chapter 3, the assumption was implicit that the population of irradiated cells was asynchronous; that is, it consisted of cells distributed throughout all phases of the cell cycle. Study of the variation of radiosensitivity with the position or age of the cell in the cell cycle was made possible only by the development of techniques to produce synchronously dividing cell cultures—populations of cells in which all of the cells occupy the same phase of the cell cycle at a given time.

There are essentially two techniques that have been used to produce a synchronously dividing cell population. The first is the mitotic harvest technique, first described by Terasima and Tolmach. This technique can be used only for cultures that grow in monolayers attached to the surface of the growth vessel. It exploits the fact that if such cells are close to mitosis, they round up and become loosely attached to the surface. If at this stage the growth medium over the cells is subjected to gentle motion (by shaking), the mitotic cells become detached from the surface and float in the medium. If this medium is then removed from the culture vessel and plated out into new petri dishes, the population consists almost entirely of mitotic cells. Incubation of these cell cultures at 37° C then causes the cells to move together synchronously in step through their mitotic cycles. By delivering a dose of radiation at various times after the initial harvesting of mitotic cells, one can irradiate cells at various phases of the cell cycle.

An alternative method for synchronizing cells, which is applicable to cells in a tissue as well as cells grown in culture, involves the use of a drug. Several different substances may be used. One of the most widely applicable drugs is hydroxyurea.

The dynamics of the action of hydroxyurea are illustrated in Figure 4.5. The drug is left in position for a period equal to the combined lengths of G2, M, and G1 for that particular cell line. By the end of the treatment period, all of the viable cells left in the population are situated in a narrow “window” at the end of G1, poised and ready to enter S phase. If the drug is then removed from the system, this synchronized cohort of cells proceeds through the cell cycle. For example, in hamster cells, 5 hours after the removal of the drug, the cohort of synchronized cells occupies a position late in the S phase. Some 9 hours after the removal of the drug, the cohort of cells is at or close to mitosis.

Techniques involving one or another of a wide range of drugs have been used to produce synchronously dividing cell populations in culture, in organized tissues (in a limited number of cases),

**FIGURE 4.5** Mode of action of hydroxyurea as an agent to induce synchrony. A: This drug kills cells in S phase and imposes a “block” at the end of the G1 phase. B: Cells in G2, M, and G1 accumulate at this block when the drug is added. C: If the block is removed, the synchronized cohort of cells moves on through the cycle.
1 hour after the mitotic cells are seeded into the petri dishes, when the cells are in G1, a dose of 6.6 Gy results in a surviving fraction of about 13%. The proportion of cells that survive the dose increases rapidly with time as the cells move into S phase; by the time the cells near the end of S phase, 42% of the cells survive this same dose. When the cells move out of S phase into G2 phase and subsequently to a second mitosis, the proportion of surviving cells falls again. This pattern of response is characteristic for most lines of Chinese hamster cells and has been reported by several independent investigators.

Complete survival curves at several discrete points during the cell cycle were measured by Sinclair. The data (from Sinclair) were obtained using Chinese hamster cells in culture. As can be seen from the figure, 1 hour after the mitotic cells are seeded into the petri dishes, when the cells are in G1, a dose of 6.6 Gy results in a surviving fraction of about 13%. The proportion of cells that survive the dose increases rapidly with time as the cells move into S phase; by the time the cells near the end of S phase, 42% of the cells survive this same dose. When the cells move out of S phase into G2 phase and subsequently to a second mitosis, the proportion of surviving cells falls again. This pattern of response is characteristic for most lines of Chinese hamster cells and has been reported by several independent investigators.

**THE EFFECT OF X-RAYS ON SYNCHRONOUSLY DIVIDING CELL CULTURES**

Figure 4.7 shows results of an experiment in which mammalian cells, which were harvested at mitosis, were irradiated with a single dose of 6.6 Gy at various times afterward, corresponding to different phases of the cell cycle. The data (from Sinclair) were obtained using Chinese hamster cells in culture. As can be seen from the figure,
The broken line in Figure 4.8 is the calculated survival curve that would be expected to apply for mitotic cells under conditions of hypoxia; that is, the slope is 2.5 times shallower than the solid line for mitotic cells, which applies to the aerated condition. This line is included in the figure to show that the range of sensitivity between the most sensitive cells (mitotic) and the most resistant cells (late S) is of the same order of magnitude as the oxygen effect (the oxygen effect is discussed in Chapter 6).

The experiments of Terasima and Tolmach with HeLa cells, in which a dose of 3 Gy was delivered to cultures at various intervals after mitotic harvesting of the cells, are shown in Figure 4.9. From the beginning of S phase onward, the pattern of sensitivity is very similar to that of hamster cells; the cells become progressively more resistant as they proceed toward the latter part of S, and after the cells move from S into G2, their sensitivity increases rapidly as they approach mitosis. The important difference between HeLa and hamster cells is the length of the G1 phase. The G1 of HeLa cells is appreciably long, and there appears to be a fine structure in the age-response function during this period. At the beginning of G1, there is a peak of resistance, followed by a sensitive trough toward the end of G1. This pattern cannot be distinguished in the hamster cell because G1 is too short.

**Figure 4.7** Fraction of Chinese hamster cells surviving a dose of 6.6 Gy of x-rays as a function of time. Time zero corresponds to the harvesting of mitotic cells. The surviving fraction increases to a maximum late in S phase. (Adapted from Sinclair WK, Morton RA. X-ray sensitivity during the cell generation cycle of cultured Chinese hamster cells. *Radiat Res.* 1966;29:450–474, with permission.)

**Figure 4.8** Cell survival curves for Chinese hamster cells at various stages of the cell cycle. The survival curve for cells in mitosis is steep and has no shoulder. The curve for cells late in S phase is shallower and has a large initial shoulder. G1 and early S phases are intermediate in sensitivity. The broken line is a calculated curve expected to apply to mitotic cells under hypoxia. (Adapted from Sinclair WK. Cyclic x-ray responses in mammalian cells in vitro. *Radiat Res.* 1968;33:620–643, with permission.)
Figure 4.10 compares the age-response curves for cells with short G₁, represented by V79 hamster cells, and cells with a long G₁, represented by HeLa cells. If the time scales are adjusted so that S phase has a comparable length for both cell lines, it is evident that the general pattern of cyclic variation is very similar, the only important difference being the extra structure during G₁ in the HeLa cells. In later experiments, other sub-lines of hamster cells were investigated for which G₁ had an appreciable length; an extra peak of resistance was noted for hamster cells that was similar to the one observed for HeLa cells.

The sensitivity of cells in different parts of G₂ is difficult to determine if synchrony is produced by mitotic selection because of synchrony decay during the passage of the starting population of mitotic cells through their first G₁ and S phases and because G₂ transit times are relatively short (about

![Diagram](image1)

**FIGURE 4.9** Fraction of HeLa cells surviving a dose of 3 Gy of x-rays administered at different times in the division cycle. Time zero represents mitosis. (Adapted from Terasima T, Tolmach LJ. Variations in several responses of HeLa cells to x-irradiation during the division cycle. *Biophys J.* 1963;3:11–33, with permission.)

![Diagram](image2)

**FIGURE 4.10** Age-response curves for cells with short G₁ phase, represented by hamster cells (A), and cells with long G₁ phase, represented by HeLa cells (B). The time scales have been adjusted so that S phase has a comparable length on the figure for both cell lines. (Adapted from Sinclair WK. Dependence of radiosensitivity upon cell age. In: *Proceedings of the Carmel Conference on Time and Dose Relationships in Radiation Biology as Applied to Radiotherapy*. Brookhaven National Laboratory Report 50203 (C-57). Upton, NY: 1969:97–107, with permission.)
1–2 hours). A modification of the technique, however, allows a much greater resolution for studying \( G_2 \) sensitivity. This is sometimes called “retroactive synchronization”: Cells first are irradiated, and then, as a function of time, cells arriving in mitosis are harvested by mitotic shake-off and plated for survival. In this way, it was shown that early \( G_2 \) cells are as radioresistant as late \( S \) cells and late \( G_2 \) cells are nearly as sensitive as mitotic cells; that is, a sharp transition in radiosensitivity occurs around the so-called x-ray transition point (now often called a “checkpoint”) for \( G_2 \) cell cycle delay.

The following is a summary of the main characteristics of the variation of radiosensitivity with cell age in the mitotic cycle:

1. Cells are most sensitive at or close to mitosis.
2. Resistance is usually greatest in the latter part of \( S \) phase. The increased resistance is thought to be caused by homologous recombination repair between sister chromatids that is more likely to occur after the DNA has replicated (see Chapter 2).
3. If \( G_1 \) phase has an appreciable length, a resistant period is evident early in \( G_1 \), followed by a sensitive period toward the end of \( G_1 \).
4. \( G_2 \) phase is usually sensitive, perhaps as sensitive as \( M \) phase.

Several cell lines other than HeLa and hamster have been investigated, some of which tend to agree with these results and some of which are contradictory. The summary points listed here are widely applicable, but exceptions to every one of these generalizations have been noted for one cell line or another.

### MOLECULAR CHECKPOINT GENES

Cell cycle progression is controlled by a family of genes known as **molecular checkpoint genes**. It has been known for many years that mammalian cells exposed to radiation tend to experience a block in the \( G_2 \) phase of the cell cycle. For example, the inverse dose-rate effect has been reported for cells of human origin, whereby over a limited range of dose rates around 0.30 to 0.40 Gy per hour, cells become more sensitive to radiation-induced cell killing as the dose rate is reduced, resulting in their accumulation in \( G_2 \), which is a radiosensitive phase of the cell cycle. This is described in Chapter 5. The mechanisms for this observation in human cells are not understood in detail, but the molecular genetics in yeast have been worked out, and the search is on for homologous pathways in mammalian cells.

In several strains of yeast, mutants have been isolated that are more sensitive than the wild type to both ionizing radiation and ultraviolet light by a factor between 10 and 100. The mutant gene has been cloned and sequenced and found to be a “\( G_2 \) molecular checkpoint gene.”

In the most general terms, the function of checkpoint genes is to ensure the correct order of cell cycle events, that is, to ensure that the initiation of later events depends on the completion of earlier events. The particular genes involved in radiation effects halt cells in \( G_2 \), so that an inventory of chromosome damage can be taken and repair is initiated and completed before the complex task of mitosis is attempted (Fig. 4.11). Mutant cells that lose this \( G_2 \) checkpoint gene function move directly into mitosis with damaged chromosomes and are, therefore, at a higher risk of dying—hence their greater sensitivity to radiation or, for that matter, to any DNA-damaging agent.

It has been proposed that a checkpoint control monitors spindle function during mitosis. If the spindle is disrupted by a microtubular poison, progression through mitosis is blocked. The checkpoint control is involved in this...
dependency of mitosis on spindle function. It is thought that the mechanism of action of G2 checkpoint genes involves Cdk1 (p34 protein kinase), levels of which control passage through mitosis. It is likely that mammalian cells that lack checkpoint genes would be sensitive not only to radiation-induced cell killing but also to carcinogenesis. Cells with damaged chromosomes that survive mitosis are likely to give rise to errors in chromosome segregation at mitosis, and this is one of the hallmarks of cancer.

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**THE EFFECT OF OXYGEN AT VARIOUS PHASES OF THE CELL CYCLE**

By combining the most sophisticated techniques of flow cytometry to separate cells in different phases of the cycle with the most sensitive assays for cell survival, it has been shown that the oxygen enhancement ratio (OER) varies significantly through the cycle, at least if measured for fast-growing proliferating cells cultured in vitro. The OER was measured at 2.3 to 2.4 for G2 phase cells, compared with 2.8 to 2.9 for S phase, with G1 phase cells showing an intermediate value. This is discussed in more detail in Chapter 6.

For any given phase of the cell cycle, oxygen was purely dose modifying; that is, the value of the OER was the same for all dose levels. For an asynchronous population of cells, however, the OER does vary slightly with dose or survival level. This is illustrated in Figure 6.1. The OER appears to be smaller at high levels of survival, at which the survival curve is dominated by the killing of the most sensitive moieties of the population; the OER appears to be larger at higher doses and lower levels of survival, at which the response of the most resistant (S phase) cells, which also happen to exhibit the largest OER, dominates.

This is an interesting radiobiologic observation, but the small change of OER is of little or no clinical significance in radiation therapy.

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**THE AGE-RESPONSE FUNCTION FOR A TISSUE IN VIVO**

Most studies of the variation in radiosensitivity with phase of the mitotic cycle have been done with mammalian cells cultured in vitro because of the ease with which they can be made to divide synchronously. The mitotic harvest technique is clearly only applicable to monolayer cultures, but techniques that involve a drug, such as hydroxyurea, to produce a synchronously dividing population can be applied to some organized tissues.

The epithelial lining of the mouse jejunum represents a classic self-renewal tissue. (The technique used to obtain a survival curve for the crypt cells is described in Chapter 19.) The rapidly dividing crypt cells can be synchronized by giving each mouse five intraperitoneal injections of hydroxyurea every hour. The rationale for this regimen is that all S cells are killed by the drug, and cells in other phases of the cycle are accumulated at the G1/S boundary for at least 4 hours (the overall time of the five injections).

Figure 4.12, from Withers and his colleagues, shows the response of the jejunal crypt cells to a single dose of 11 Gy of $\gamma$-rays (uppermost curve) delivered at various times after the synchronizing action of the five injections of hydroxyurea. The number of crypt cells per circumference of the sectioned jejunum varies by a factor of 100, according to the phase in the cycle at which the radiation is delivered, ranging from about 2 survivors per circumference for irradiation 2 hours after the last injection of hydroxyurea to about 200 survivors per circumference by 6 hours. The DNA synthetic activity of the synchronized jejunal mucosa was monitored by injecting groups of mice with tritiated thymidine at hourly intervals after the last injection of hydroxyurea and subsequently removing a sample of the jejunum and assaying the radioactive content. The bottom curve of Figure 4.12 shows the variation of thymidine uptake with time. The first wave of the thymidine uptake represents the period of DNA synthesis of the synchronized crypt cells. The peak coincides closely with the period of maximum resistance to $\gamma$-rays (about 5 hours after the last injection of hydroxyurea).

These data indicate clearly that the radiosensitivity of crypt cells in the mouse jejunum varies substantially with the phase of the cell cycle at which the radiation is delivered. Further, the pattern of response in this organized normal tissue, with a sensitive period between G1 and S and maximum radioresistance late in S, is very similar to that characteristic of many cell lines cultured in vitro.
In the early part of the cycle, before replication has occurred, DSBs must be repaired by nonhomologous end joining because no template exists to guide gap filling. This process is error prone. On the other hand, in S phase after replication, DSBs can be repaired by homologous recombination because a template is available (i.e., an identical sister chromatid is available). This process is less likely to result in errors. Radiosensitivity correlates with error-prone nonhomologous end joining of DSBs; radioresistance correlates with homologous recombination of DSBs, which is likely to be more faithful. For a description of homologous and nonhomologous repair, see Chapter 2.

The possible implications of the age-response function in radiotherapy

If a single dose of radiation is delivered to a population of cells that are asynchronous—that is, distributed throughout the cell cycle—the effect is different on cells occupying different phases of the cell cycle at the time of the radiation exposure. A greater proportion of cells are killed in the sensitive portions of the cell cycle, such as those at or close to mitosis; a smaller proportion of those in the DNA synthetic phase are killed. The overall effect is that a dose of
radiation, to some extent, tends to synchronize the cell population, leaving most cells in a resistant phase of the cycle. Between dose fractions, movement of cells through the cycle into more sensitive phases may be an important factor in “sensitizing” a cycling population of tumor cells to later doses in fractionated regimens. This is considered sensitization resulting from reassortment. It results in a therapeutic gain because sensitization by this mechanism occurs only in rapidly dividing cells and not in late-responding normal tissues.

SUMMARY OF PERTINENT CONCLUSIONS

- The cell cycle for mammalian cells can be divided into four phases: mitosis (M), followed by G1, followed by the DNA synthetic phase (S), then G2, and into mitosis again.
- The phases of the cycle are regulated by the periodic activation of different members of the Cdk family.
- The fastest cycling mammalian cells in culture, as well as crypt cells in the intestinal epithelium, have cycle times as short as 9 to 10 hours. Stem cells in resting mouse skin may have cycle times of more than 200 hours. Most of this difference results from the varying length of G1, the most variable phase of the cycle. The M, S, and G2 phases do not vary much.
- In general, cells are most radiosensitive in the M and G2 phases and most resistant in late S phase.
- For cells with longer cell cycle times and significantly long G1 phases, there is a second peak of resistance early in G1.
- Molecular checkpoint genes stop cells from cycling if exposed to x-rays or any other DNA-damaging agent, allowing the chromosomes to be checked for integrity before the complex task of mitosis is attempted.
- The OER varies little with phase of the cell cycle but may be slightly lower for cells in G1 than for cells in S.
- The age-response function for crypt cells in the mouse jejunal is similar to that for cells in culture. This is the only tissue in which this has been studied.

- The age-response function for neutrons is qualitatively similar to that for x-rays, but the magnitude of changes through the cycle is smaller.
- The patterns of radiosensitivity and radioresistance correlate with the mechanism of repair of DNA DSBs. Radiosensitivity correlates with nonhomologous end joining, which dominates early in the cell cycle and is error prone. Radioresistance correlates with homologous recombinational repair, which occurs after replication (in S phase) and is more faithful.
- Variations in sensitivity through the cell cycle may be important in radiation therapy because they lead to “sensitization resulting from reassortment” in a fractionated regimen.

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Operational Classifications of Radiation Damage

Radiation damage to mammalian cells can operationally be divided into three categories: (1) lethal damage, which is irreversible and irreparable and, by definition, leads irrevocably to cell death; (2) potentially lethal damage (PLD), the component of radiation damage that can be modified by postirradiation environmental conditions; and (3) sublethal damage (SLD), which under normal circumstances, can be repaired in hours unless additional SLD is added (e.g., from a second dose of radiation) with which it can interact to form lethal damage (SLD repair, therefore, is manifested by the increase in survival observed if a dose of radiation is split into two fractions separated by a time interval).

Potentially Lethal Damage Repair

Varying environmental conditions after exposure to x-rays can influence the proportion of cells that survive a given dose because of the expression or repair of PLD. This damage is potentially lethal because under ordinary circumstances, it causes cell death, but if survival is increased as a result of the manipulation of the postirradiation environment, PLD is considered to have been repaired. PLD is repaired if cells are incubated in a balanced salt solution instead of a full growth medium for several hours after irradiation. This is a drastic treatment, however, and does not mimic a physiologic condition that is ever likely to occur. Little and his colleagues chose to study PLD repair in density-inhibited stationary-phase cell cultures, which are considered a better in vitro model for tumor cells in vivo (Fig. 5.1). Cell survival was enhanced considerably if the cells were allowed to remain in the density-inhibited state for 6 or 12 hours after irradiation before being subcultured and assayed for colony-forming ability.

The relevance of PLD to radiotherapy became much more obvious when it was shown that repair, comparable in magnitude and kinetics to that found in vitro, also occurred in vivo in experimental tumors. In this case, repair took the form of significantly enhanced cell survival if several hours were allowed to elapse between irradiation of the tumor in situ and removal of the cells from the host to assess their reproductive integrity (Fig. 5.2).

To summarize the available experimental data, there is a general agreement that PLD is repaired, and the fraction of cells surviving a given dose of x-rays is enhanced if postirradiation conditions are suboptimal for growth, so that cells do not have to attempt the complex process of mitosis while their chromosomes are damaged. If mitosis is delayed by
to suppose that it does not occur in human tumors. It has been suggested that the radioresistance of certain types of human tumors is linked to their ability to repair PLD; that is, radiosensitive tumors repair PLD inefficiently, but radioresistant tumors have efficient mechanisms to recover from suboptimal growth conditions, DNA damage can be repaired.

The importance of PLD repair to clinical radiotherapy is a matter of debate. That it occurs in transplantable animal tumors has been documented beyond question, and there is no reason to suppose that it does not occur in human tumors. It has been suggested that the radioresistance of certain types of human tumors is linked to their ability to repair PLD; that is, radiosensitive tumors repair PLD inefficiently, but radioresistant tumors have efficient mechanisms to repair potentially lethal damage. (Adapted from Little JB, Hahn GM, Frindel E, et al. Repair of potentially lethal radiation damage in vitro and in vivo. *Radiology.* 1973;106:689–694, with permission.)

**FIGURE 5.1** X-ray survival curves for density-inhibited stationary-phase cells, subcultured (trypsinized and plated) either immediately or 6 or 12 hours after irradiation. Cell survival is enhanced if cells are left in the stationary phase after irradiation, allowing time for the repair of potentially lethal damage. (Adapted from Little JB, Hahn GM, Frindel E, et al. Repair of potentially lethal radiation damage in vitro and in vivo. *Radiology.* 1973;106:689–694, with permission.)

**FIGURE 5.2** Repair of potentially lethal damage in mouse fibrosarcomas. The tumors were irradiated in situ and then removed and prepared into single cell suspension. The number of survivors was determined by their ability to form colonies in vitro. The fraction of cells surviving a given dose increases if a time interval is allowed between irradiation and removal of the tumor, because during this interval, PLD is repaired. (Adapted from Little JB, Hahn GM, Frindel E, et al. Repair of potentially lethal radiation damage in vitro and in vivo. *Radiology.* 1973;106:689–694, with permission.)
repair PLD. This is an attractive hypothesis, but it has never been proven.

**Sublethal Damage Repair**

SLD repair is the operational term for the increase in cell survival that is observed if a given radiation dose is split into two fractions separated by a time interval.

Figure 5.3 shows data obtained in a split-dose experiment with cultured Chinese hamster cells. A single dose of 15.58 Gy of absorbed radiation leads to a surviving fraction of 0.005. If the dose is divided into two approximately equal fractions separated by 30 minutes, the surviving fraction is already appreciably higher than for a single dose. As the time interval is extended, the surviving fraction increases until a plateau is reached at about 2 hours, corresponding to a surviving fraction of 0.02. This represents about four times as many surviving cells as for the dose given in a single exposure. A further increase in the time interval between the dose fractions is not accompanied by any significant additional increment in survival. The increase in survival in a split-dose experiment results from the repair of sublethal radiation damage.

The data shown in Figure 5.3 were obtained with cultured mammalian cells maintained at room temperature (24°C) between the dose fractions to prevent the cells from moving through the cell cycle during this interval. This rather special experiment is described first because it illustrates repair of sublethal radiation damage uncomplicated by the movement of cells through the cell cycle.

Figure 5.4 shows the results of a parallel experiment in which cells were exposed to split doses and maintained at their normal growing temperature of 37°C. The pattern of repair seen in this case differs from that observed for cells kept at room temperature. In the first few hours, prompt repair of SLD is again evident, but at longer intervals between the two split doses, the surviving fraction of cells decreases, reaching a minimum with about a 5-hour separation.

![Figure 5.3](image-url) Survival of Chinese hamster cells exposed to two fractions of x-rays and incubated at room temperature for various time intervals between the two exposures. (Adapted from Elkind MM, Sutton-Gilbert H, Moses WB, Alescio T, Swain RB. Radiation response of mammalian cells in culture: V. Temperature dependence of the repair of x-ray damage in surviving cells [aerobic and hypoxic]. Radiat Res. 1965;25:359–376, with permission.)

![Figure 5.4](image-url) Survival of Chinese hamster cells exposed to two fractions of x-rays and incubated at 37°C for various time intervals between the two doses. The survivors of the first dose are predominantly in a resistant phase of the cycle (late S). If the interval between doses is about 6 hours, these resistant cells have moved to the G2M phase, which is sensitive. (Adapted from Elkind MM, Sutton-Gilbert H, Moses WB, et al. Radiation response of mammalian cells in culture: V. Temperature dependence of the repair of x-ray damage in surviving cells [aerobic and hypoxic]. Radiat Res. 1965;25:359–376, with permission.)
An understanding of this phenomenon is based on the age-response function described in Chapter 4. If an asynchronous population of cells is exposed to a large dose of radiation, more cells are killed in the sensitive than in the resistant phases of the cell cycle. The surviving population of cells, therefore, tends to be partly synchronized.

In Chinese hamster cells, most of the survivors from a first dose of radiation are located in the S phase of the cell cycle. If about 6 hours are allowed to elapse before a second dose of radiation is given, this cohort of cells progresses around the cell cycle and is in G2/M, a sensitive period of the cell cycle at the time of the second dose. If the increase in radiosensitivity in moving from late S to the G2/M period exceeds the effect of repair of SLD, the surviving fraction falls.

The pattern of repair shown in Figure 5.4 is therefore a combination of three processes occurring simultaneously. First, there is the prompt repair of sublethal radiation damage. Second, there is progression of cells through the cell cycle during the interval between the split doses, which has been termed reassortment. Third, there is an increase of surviving fraction resulting from cell division, or repopulation, if the interval between the split doses is from 10 to 12 hours, because this exceeds the length of the cell cycle of these rapidly growing cells.

This simple experiment, performed in vitro, illustrates three of the “four Rs” of radiobiology: repair, reassortment, and repopulation. The fourth “R,” reoxygenation, is discussed in Chapter 6. It should be emphasized that the dramatic dip in the split-dose curve at 6 hours caused by reassortment, and the increase in survival by 12 hours because of repopulation are seen only for rapidly growing cells. Hamster cells in culture have a cycle time of only 9 or 10 hours. The time sequence of these events would be longer in more slowly proliferating normal tissues in vivo.

Repair of sublethal radiation damage has been demonstrated in just about every biologic test system for which a quantitative end point is available. Figure 5.5 illustrates the pattern for repair of sublethal radiation damage in two in vivo systems in mice, P388 lymphocytic leukemia and skin cells. In neither case, there is a dramatic dip in the curve at 6 hours resulting from movement of cells through the cycle, because the cell cycle is long. In resting skin, for example, the cell cycle of stem cells may be as long as 10 days rather than 9 hours of the rapidly growing cells in Figure 5.4. The mouse tumor data show more repair in small 1-day tumors than in large hypoxic 6-day tumors; this important point illustrates that repair is an active process requiring oxygen and nutrients.

![Figure 5.5](image_url)
The various factors involved in the repair of SLD are summarized in Figure 5.6. Figure 5.6A shows that if a dose is split into two fractions separated by a time interval, more cells survive than for the same total dose given in a single fraction, because the shoulder of the curve must be repeated with each fraction. In general, there is a good correlation between the extent of repair of SLD and the size of the shoulder of the survival curve. This is not surprising because both are manifestations of the same basic phenomenon: the accumulation and repair of SLD. Some mammalian cells are characterized by a survival curve with a broad shoulder, and split-dose experiments then indicate a substantial amount of SLD repair. Other types of cells show a survival curve with a minimal shoulder, and this is reflected in more limited repair of SLD. In the terminology of the linear-quadratic ($\alpha/\beta$) description of the survival curve, it is the quadratic component ($\beta$) that causes the curve to bend and that results in the sparing effect of a split dose. A large shoulder corresponds to a small $\alpha/\beta$ ratio.

The time course of the increase in cell survival that results from the repair of SLD is charted in Figure 5.6B. As the time interval between the two dose fractions is increased, there is a rapid increase in the fraction of cells surviving owing to the prompt repair of SLD. This repair is complete by 1 or 2 hours for cells in culture but may take longer for late-responding tissues in vivo (Chapter 23). As the time interval between the two dose fractions is increased, there is a dip in the curve owing to the movement of surviving cells through the cell cycle, as explained in Figure 5.4. This occurs only in a population of fast-cycling cells. In cells that are noncycling, there can be no dip. If the time interval between the two dose

![Figure 5.6](25858_Hall_CH05.indd_71.png)

**FIGURE 5.6** Summary of the repair of sublethal damage as evidenced by a split-dose experiment. **A:** If the dose is delivered in two fractions separated by a time interval, there is an increase in cell survival because the shoulder of the curve must be expressed each time. **B:** The fraction of cells surviving a split dose increases as the time interval between the two dose fractions increases. As the time interval increases from 0 to 2 hours, the increase in survival results from the repair of sublethal damage. In cells with a long cell cycle or that are out of cycle, there is no further increase in cell survival by separating the dose by more than 2 or 3 hours. In a rapidly dividing cell population, there is a dip in cell survival caused by reassortment. However, as shown in Figure 5.7, if the time interval between the split doses exceeds the cell cycle, there is an increase in cell survival owing to proliferation or repopulation between the doses.
fractions exceeds the cell cycle, there is an increase in the number of cells surviving because of cell proliferation; that is, cells can double in number between the dose fractions.

**MECHANISM OF SUBLETHAL DAMAGE REPAIR**

In Chapter 3, evidence was summarized of the correlation between cell killing and the production of asymmetric chromosomal aberrations, such as dicentrics and rings. This, in turn, is a consequence of an interaction between two (or more) double-strand breaks in the DNA. Based on this interpretation, the repair of SLD is simply the repair of double-strand breaks. If a dose is split into two parts separated by a time interval, some of the double-strand breaks produced by the first dose are rejoined and repaired before the second dose. The breaks in two chromosomes that must interact to form a lethal lesion such as a dicentric may be formed by (1) a single track breaking both chromosomes (i.e., single-track damage), or (2) separate tracks breaking the two chromosomes (i.e., multiple-track damage).

The component of cell killing that results from single-track damage is the same whether the dose is given in a single exposure or fractionated. The same is not true of multiple-track damage. If the dose is given in a single exposure (i.e., two fractions with $t = 0$ between them), all breaks produced by separate electrons can interact to form dicentrics. But if the two dose fractions ($D/2$) are separated by, for example, 3 hours, then breaks produced by the first dose may be repaired before the second dose is given. Consequently, there are fewer interactions between broken chromosomes to form dicentrics, and more cells survive. Based on this simple interpretation, the repair of SLD reflects the repair and rejoining of double-strand breaks before they can interact to form lethal lesions. This may not be the whole story, but it is a useful picture to keep in mind.

**REPAIR AND RADIATION QUALITY**

For a given biologic test system, the shoulder on the acute survival curve and, therefore, the amount of SLD repair indicated by a split-dose experiment vary with the type of radiation used. The effect of dose fractionation with $x$-rays and neutrons is compared in Figure 5.7. For $x$-rays, dividing the total dose into two equal fractions, separated from 1 to 4 hours, results in a marked increase in cell survival because of the prompt repair of SLD. By contrast, dividing the dose into two fractions has little effect on cell survival if neutrons are used, indicating little repair of SLD.

**THE DOSE-RATE EFFECT**

For $x$- or $\gamma$-rays, dose rate is one of the principal factors that determine the biologic consequences of a given absorbed dose. As the dose rate is lowered and the exposure time extended, the biologic effect of a given dose generally is reduced.

The classic dose-rate effect, which is very important in radiotherapy, results from the repair of SLD that occurs during a long radiation exposure. To illustrate this principle, Figure 5.8 shows an idealized experiment in which each dose ($D_2, D_3, D_4, \text{and so on}$) is delivered in several equal fractions of size $D$, with a time interval between fractions that is sufficient for repair of SLD. The shoulder of the survival curve is repeated with each fraction. The broken line, $F$, shows the overall survival curve that would be
be expected, because both are expressions of the cell’s capacity to accumulate and repair sublethal radiation damage. By contrast, Chinese hamster cells have a broad shoulder to their acute x-ray survival curve and show a correspondingly large dose-rate effect. This is evident in Figure 5.10; observed if only single points were determined, corresponding to equal dose increments. This survival curve has no shoulder. Because continuous low-dose-rate (LDR) irradiation may be considered to be an infinite number of infinitely small fractions, the survival curve under these conditions also would be expected to have no shoulder and to be shallower than for single acute exposures.

**EXAMPLES OF THE DOSE-RATE EFFECT IN VITRO AND IN VIVO**

Survival curves for HeLa cells cultured in vitro over a wide range of dose rates, from 7.3 Gy/min to 0.535 cGy/min, are summarized in Figure 5.9. As the dose rate is reduced, the survival curve becomes shallower and the shoulder tends to disappear (i.e., the survival curve becomes an exponential function of dose). The dose-rate effect caused by repair of SLD is most dramatic between 0.01 and 1 Gy/min. Above and below this dose-rate range, the survival curve changes little, if at all, with dose rate.

The magnitude of the dose-rate effect from the repair of SLD varies enormously among different types of cells. HeLa cells are characterized by a survival curve for acute exposures that has a small initial shoulder, which goes hand in hand with a modest dose-rate effect. This is to be expected, because both are expressions of the cell’s capacity to accumulate and repair sublethal radiation damage. By contrast, Chinese hamster cells have a broad shoulder to their acute x-ray survival curve and show a correspondingly large dose-rate effect. This is evident in Figure 5.10;
apoptosis is an important form of cell death following radiation, whereas for hamster cells, apoptotic death is rarely seen.

FIGURE 5.11 shows survival curves for 40 different cell lines of human origin, cultured in vitro and irradiated at high dose rates (HDR) and low dose rates (LDR). At LDR, the survival there is a clear-cut difference in biologic effect, at least at high doses, between dose rates of 1.07, 0.30, and 0.16 Gy/min. The differences between HeLa and hamster cells in the size of the shoulder to the acute survival curve and the magnitude of the dose-rate effect reflect differences in the importance of apoptosis. In the case of HeLa cells, apoptosis is an important form of cell death following radiation, whereas for hamster cells, apoptotic death is rarely seen.

Figure 5.11 shows survival curves for 40 different cell lines of human origin, cultured in vitro and irradiated at high dose rates (HDR) and low dose rates (LDR). At LDR, the survival
curves “fan out” and show a greater variation in slope because, in addition to the variation of inherent radiosensitivity evident at an HDR, there is a range of repair times of SLD. Some cell lines repair SLD rapidly, some more slowly, and this is reflected in the different survival curves at LDR.

Survival curves for crypt cells in the mouse jejunum irradiated with γ-rays at various dose rates are shown in Figure 5.12. There is a dramatic dose-rate effect owing to the repair of sublethal radiation damage from an acute exposure at 2.74 Gy/min to a protracted exposure at 0.92 cGy/min. As the dose rate is lowered further, cell division begins to dominate the picture because the exposure time is longer than the cell cycle. At 0.54 cGy/min, there is little reduction in the number of surviving crypts, even for very large doses, because cellular proliferation occurs during the long exposure and makes up for cell killing by the radiation.

THE INVERSE DOSE-RATE EFFECT

There is at least one example of an inverse dose-rate effect, in which decreasing the dose rate results in increased cell killing. This is illustrated in Figure 5.13. Decreasing the dose rate for this HeLa cell line from 1.54 to 0.37 Gy/h increases the efficiency of cell killing, such that this LDR
during the radiation exposure if the dose rate is low enough and the exposure time is long compared with the length of the mitotic cycle. This may lead to a further reduction in biologic effect as the dose rate is progressively lowered, because cell birth tends to offset cell death.

BRACHYTHERAPY OR ENDOCURIETHERAPY

Implanting radioactive sources directly into a tumor was a strategy first suggested by Alexander Graham Bell in 1901. Over the years, various groups in different countries coined various names for this type of therapy, using the prefix brachy, from the Greek word for “short range,” or endo, from the Greek word for “within.” There are two distinct forms of brachytherapy, also called endocurietherapy: (1) intracavitary

THE DOSE-RATE EFFECT SUMMARIZED

Figure 5.15 summarizes the entire dose-rate effect. For acute exposures at high dose rates, the survival curve has a significant initial shoulder. As the dose rate is lowered and the treatment time protracted, more and more SLD can be repaired during the exposure. Consequently, the survival curve becomes progressively more shallow (D₀ increases) and the shoulder tends to disappear. A point is reached at which all SLD is repaired resulting in a limiting slope. In at least some cell lines, a further lowering of the dose rate allows cells to progress through the cycle and accumulate in G₂. This is a radiosensitive phase, and so the survival curve becomes steeper again. This is the inverse dose-rate effect. A further reduction in dose rate allows cells to pass through the G₂ block and divide. Proliferation then may occur during the radiation exposure if the dose rate is low enough and the exposure time is long compared with the length of the mitotic cycle. This may lead to a further reduction in biologic effect as the dose rate is progressively lowered, because cell birth tends to offset cell death.

FIGURE 5.14 The inverse dose-rate effect. A range of dose rates can be found, at least for HeLa cells, that allows cells to progress through the cycle to a block in late G₂. Under continuous low-dose-rate irradiation, an asynchronous population becomes a population of radiosensitive G₂ cells. (Adapted from Hall EJ. The biological basis of endocurietherapy: the Henschke Memorial Lecture 1984. Endocurie Hypertherm Oncol. 1985;1:141–151, with permission.)

FIGURE 5.15 The dose-rate effect resulting from repair of sublethal damage, redistribution in the cycle, and cell proliferation. The dose-response curve for acute exposures is characterized by a broad initial shoulder. As the dose rate is reduced, the survival curve becomes progressively more shallow as more and more sublethal damage is repaired, but cells are “frozen” in their positions in the cycle and do not progress. As the dose rate is lowered further and for a limited range of dose rates, the survival curve steepens again because cells can progress through the cycle to pile up at a block in G₂, a radiosensitive phase, but still cannot divide. A further lowering of dose rate below this critical dose rate allows cells to escape the G₂ block and divide; cell proliferation then may occur during the protracted exposure, and survival curves become shallower as cell birth from mitosis offsets cell killing from the irradiation. (Based on the ideas of Dr. Joel Bedford.)
Interstitial brachytherapy can be either temporary or permanent. Temporary implants in earlier times used radium, but the most widely used radionuclide at present is iridium-192. Implants at LDR are considered by many radiotherapists to be the treatment of choice for the 5% or so of human cancers that are accessible to such techniques.

The dose-rate range used in these treatments is in the region of the dose-rate spectrum in which the biologic effect varies rapidly with dose rate. The maximum dose that can be delivered without unacceptable damage to the surrounding normal tissue depends on the volume of tissue irradiated and on the dose rate, which is in turn a function of the number of radioactive sources used and their geometric distribution. To achieve a consistent biologic response, the total dose used should be varied according to the dose rate employed.

Paterson and Ellis independently published curves to relate total dose to result in normal-tissue tolerance to dose rate (Fig. 5.16); there is remarkable agreement between the two sets of data based on clinical judgment.

Quoting Paterson:

"The graph for radium implants is an attempt to set out the doses in 5 to 10 days, which are equivalent to any desired 7-day dose. In its original form, it perhaps owed more to inspiration than to science but it has gradually been corrected to match actual experience."

To an increasing extent, LDR intracavitary brachytherapy is being replaced by HDR intracavitary therapy, delivered in 3- to 12-dose fractions. Replacing continuous LDR therapy with a few large-dose fractions gives up much of the radiobiologic advantage and the sparing of late-responding normal tissues, as described in Chapter 23. It is only possible because the treatment of carcinoma of the cervix is a special case in which the dose-limiting normal tissues (e.g., bladder, rectum) receive a lower dose than the prescribed dose to the tumor (or to point A). For HDR treatments lasting a few minutes, it is possible to use retractors that result in even lower doses to the critical normal tissues than are possible with an insertion that lasts 24 hours or more. These physical advantages offset the radiobiologic disadvantages, so that the general principle that administration of a few large fractions at an HDR gives poorer results than at an LDR, does not apply to this special case.

**Interstitial Brachytherapy**

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used, so only limited conclusions can be drawn from these data. The results (Fig. 5.18), however, show a correlation between the proportion of recurrent tumors and the dose rate. For a given total dose, there were markedly fewer recur-

estimates of Paterson and of Ellis, as it should, because their judgment was stated unequivocally to be based on late effects.

In the 1990s, Mazeron and his colleagues in Paris published two papers that show clearly that a dose-rate effect is important in interstitial implants. They have, perhaps, the most experience in the world with the use of iridium-192 wire implants. Their first report describes the analysis of local tumor control and the incidence of necrosis in a large cohort of patients with T1-2 squamous cell carcinoma of the mobile tongue and the floor of the mouth who were treated with interstitial iridium-192. The data are shown in Figure 5.17. Patients were grouped according to dose rate, either more or less than 0.5 Gy/h. It is evident that there was a substantially higher incidence of necrosis in patients treated at the higher dose rates. By contrast, dose rate makes little or no difference to local control provided that the total dose is high enough, from 65 to 70 Gy, but there is a clear separation at lower doses (60 Gy), with the lower dose rate being less effective. These results are in good accord with the radiobiologic predictions.

Their second report analyzes data from a large group of patients with carcinoma of the breast who received iridium-192 implants as a boost to external beam radiotherapy. These results allow an assessment of the effect of dose rate on tumor control, but provide no information on the effect of dose rate on late effects, because there was only one case that involved necrosis. The interstitial implant was only part of the radiotherapy and a fixed standard dose was

FIGURE 5.16 Dose equivalent to 60 Gy in 7 days as proposed by Paterson (in 1963) and by Ellis (in 1968) based on clinical observation of normal-tissue tolerance or calculated from radiobiologic principles. The $\alpha/\beta$ ratios and the half-time of repair of sublethal damage were chosen for early- or late-responding tissues (Chapter 23).

FIGURE 5.17 Local tumor control and necrosis rate at 5 years as a function of dose in patients with T1–2 squamous cell carcinomas of the mobile tongue and the floor of the mouth who were treated with interstitial iridium-192 implants. The patients were grouped according to whether the implant was characterized by a high dose rate (equal to or above 0.5 Gy/h) or low dose rate (below 0.5 Gy/h). The necrosis rate was higher for the higher-dose-rate group at all dose levels. Local tumor control did not depend on dose rate, provided the total dose was sufficiently large. (Data from Mazeron JJ, Simon JM, Le Pechoux C, et al. Effect of dose rate on local control and complications in definitive irradiation of T1-2 squamous cell carcinomas of mobile tongue and floor of mouth with interstitial iridium-192. Radiother Oncol. 1991;21:39–47.)
size can be small, and (2) its lower photon energy makes radiation protection easier than with radium or cesium-137. Sources of this radionuclide are ideal for use with computer-controlled remote afterloaders introduced in the 1990s (Fig. 5.19). Catheters can be implanted into the patient while inactive and then the sources transferred from the safe by remote control after the patient has returned to his own room. The sources can be returned to the safe if the patient needs nursing care.

**Permanent Interstitial Implants**

Encapsulated sources with relatively short half-lives can be left in place permanently. There are two advantages for the patient: (1) An operation to remove the implant is not needed, and (2) the patient can go home with the implant in place. On the other hand, this does involve additional expense because the sources are not reused. The initial dose rate is high and falls off as the implanted sources decay. Iodine-125 has been used most widely to date for permanent implants. The total prescribed dose is usually about 160 Gy at the periphery of the implanted volume, with 80 Gy delivered in the first half-life of 60 days. The soft emission from iodine has a relative biologic effectiveness (explained in Chapter 7) of about 1.5; this corresponds to $80/1.5 = 53.3$ Gy of high-energy $\gamma$-rays. This is a big dose, even at an LDR and corresponds to a good level of cell kill. It is, however, spread over 60 days; consequently, the success of the implant in sterilizing the tumor depends critically on the dose rate.

The relatively short half-life of iridium-192 (70 days) means that a range of dose rates is inevitable, because the activity of the sources decays during the months that they are in use. It is important, therefore, to correct the total dose for the dose rate because of the experience of Mazeron and his colleagues described previously. Iridium-192 has two advantages: (1) The source references if the radiation was delivered at a higher dose rate rather than a lower dose rate.

![Graph showing percentage of patients with no local recurrence as a function of dose rate in treatment for breast carcinoma by a combination of external-beam irradiation plus iridium-192 interstitial implant. The implant was used to deliver a total dose of 37 Gy; the dose rate varied by a factor of 3 (30–90 cGy/hr), owing to different linear activities of the iridium-192 wire and different volumes implanted. (Data from Mazeron JJ, Simon JM, Crook J, et al. Influence of dose rate on local control of breast carcinoma treated by external beam irradiation plus iridium-192 implant. Int J Radiat Oncol Biol Phys. 1991;21:1173–1177.)](image-url)

![Diagram illustrating the use of a computer-controlled remote afterloader to minimize radiation exposure of personnel during brachytherapy. Catheters are implanted into the tumor, and radiographs are made to check the validity of the implant using “dummy” nonradioactive sources. The catheters then are connected to a shielded safe containing the radioactive (iridium-192) sources, which are transferred by remote control to the implant in the patient. The control panel is located outside a lightly shielded room. The sources can be retracted temporarily to the safe so that personnel can care for the patient, thus effectively eliminating radiation exposure to personnel.](diagram-url)
characteristics of radionuclides more commonly used for brachytherapy.

### RADIOLABELED IMMUNOGLOBULIN THERAPY FOR HUMAN CANCER

Radiolabeled immunoglobulin therapy is radiotherapy for cancer using an antibody to deliver a radioactive isotope to the tumor. Much of the pioneering work in this field was done by Stanley Order and his colleagues in the 1980s, with the primary focus on antiferritin labeled with radioactive iodine or yttrium.

Ferritin is an iron-storage protein that is synthesized and secreted by a broad range of malignancies, including hepatoma, lung cancer, neuroblastoma, acute myelogenous leukemia, cancer of the breast and pancreas, and Hodgkin disease. It is not known why ferritin is produced preferentially in tumors. This suggestion is highly speculative but consistent with the observation that ferritin is present in tumors that are suspected of having a viral cause. This connection is strongly suspected for hepatomas, which have been associated with the hepatitis B virus and probably exists for Hodgkin disease, too.

<table>
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<tr>
<th>Radionuclide</th>
<th>Average Photon Energy, keV</th>
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<td></td>
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<tr>
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<td>30 y</td>
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<tr>
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<tr>
<td>Iodine-125</td>
<td>28</td>
<td>3–35</td>
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<td>Ytterbium 169</td>
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</tr>
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</table>

*HVL, half-value layer, the thickness required to reduce the incident radiation by 50.

Data computed by Dr. Ravinder Nath, Yale University.
Although ferritin is also present in normal tissues, selective tumor targeting has been demonstrated in animal models and in clinical scanning, historically performed first for Hodgkin disease. This differential is the basis of the potential therapeutic gain, and thus, the clinical usefulness of radiolabeled immunoglobulin therapy.

In the early years of radiolabeled immunoglobulin therapy, radiolabeled polyclonal antibodies were used. These were replaced with murine monoclonal antibodies carrying iodine-131, which could be used for both diagnosis and therapy. More recently, chimeric mouse–human antibodies, which are human antibodies derived by tissue culture or produced in genetically altered mice, and synthetically derived antibodies have become available. These developments have progressively reduced the possibility of inducing an immune response, lengthened the effective half-life, and hence, increased the tumor dose.

**Radionuclides**

Early studies used iodine-131, which is easily linked to antibodies. The disadvantage of using iodine-131 is that it requires large amounts of radioactivity (about 1,000 MBq); as a consequence of this, patients must be hospitalized, self-care is needed, and pediatric patients are excluded. In addition, the dose and dose rate to the tumor are limited by the relatively weak β-emission (0.3 MeV) and by the total body dose resulting from the γ-emission, which causes systemic hematopoietic toxicity.

In more recent developments, iodine-131 has been replaced by yttrium-90, which is characterized by a pure β-emission of relatively high energy (0.9 MeV). This allows a higher tumor dose and dose rate and enables the applications to be administered on an outpatient basis. More recently, rhenium-188, rhenium-186, and phosphorus-32 have been used. New chemical linkages, including various chelates, have also been used; all seeking to bind the isotope firmly to the antibody.

**Tumor Target Visualization**

When iodine-131 was used, the γ-ray emission allowed tumor localization as well as providing the bulk of the therapeutic dose. When pure β-emitters such as yttrium-90 were first introduced, it was necessary to add a γ-emitter such as indium-111 to allow visualization. Today, it is no longer acceptable to scan with a conventional γ-camera because single photon emission computed tomography (SPECT) provides a clearer picture. The bremsstrahlung from β-emitters can be scanned by this means, so that radionuclides such as yttrium-90 can be used without the need to add a γ-emitter for visualization.

**Targeting**

The ability to target tumors with antiferritin mirrors the vascularity of the tumor nodules. In general, tumors with a high degree of vascularity are better targeted with antiferritin than less vascularized tumors. The presence of ferritin per se is not enough to ensure targeting. The need for neovasculature means that uptake tends to be greater in smaller tumors. Uptake also can be affected by radiation or hyperthermia. A dose of external radiation can act as an initiator. This first was observed empirically but now is used routinely to enhance the targeting of the radio-labeled antiferritin. This is probably a consequence of damage to tumor vasculature, which allows antiferritin to leak out of vessels and into tumor cells. The targeting ratio of a tumor to the average for normal tissue is about 2.9 for antiferritin labeled with iodine-131; the corresponding ratios are 1.2 for bone and gastrointestinal tract and 0.8 for lung.

**Clinical Results**

The most promising results have been in the treatment of unresectable primary hepatoma, for which 48% partial remission and 7% complete remission rates have been reported by the Johns Hopkins group for patients receiving iodine-131–labeled antiferritin in combination with low doses of doxorubicin (Adriamycin) and 5-fluorouracil. Some success also has been reported by other groups using similar techniques in the treatment of metastatic neuroblastoma, relapsed grade IV gliomas after radiotherapy and chemotherapy, metastatic ovarian cancer resistant to prior radiotherapy, and malignant pleural and pericardial effusions of diverse causes.

Iodine-131–labeled antiferritin led to partial remissions in patients with Hodgkin disease, but yttrium-90 antiferritin produced complete remissions, indicating the increased effectiveness of the larger doses possible with radionuclides emitting pure β-rays. Radiolabeled immunoglobulin therapy has been used with varying
degrees of success for a wide range of other malignancies, including hepatomas, ovarian cancer, gliomas, and leukemia. Although various radiolabeled antibodies have been shown to achieve remissions in lymphoma, the question of the effect of the total body exposure versus tumor targeting is still open.

**Dosimetry**

For iodine-131–labeled antiferritin treatment of unresectable primary hepatoma, about 1,000 MBq is administered on day 1, followed by about 700 MBq on day 5. Escalation of dose beyond these levels is not helpful because the deposition of labeled antiferritin becomes saturated. This translates into a peak dose rate of 45 to 50 mGy/h on days 1 and 5 and a total accumulated dose of 10 to 12 Gy by about 15 days. The corresponding dose rate to normal liver is 10 mGy/h, and the total body dose is 2 to 3 mGy/h, which results limit hematologic toxicity. It is remarkable that such a small dose at such an LDR can produce remissions in patients with tumors of 1 kg or more. This response is difficult to explain based on conventional radiobiologic data, but the clinical results are exciting.

For yttrium-90–labeled antiferritin treatment, a single application of about 700 MBq results in a peak dose rate of about 0.16 Gy/h, which decays with a tumor-effective half-life of 2 days and results in a total accumulated dose of about 20 to 35 Gy.

**SUMMARY OF PERTINENT CONCLUSIONS**

**Potentially Lethal Damage Repair**
- The component of radiation damage that can be modified by manipulation of the postirradiation conditions is known as PLD.
- PLD repair can occur if cells are prevented from dividing for 6 hours or more after irradiation; this is manifested as an increase in survival. This repair can be demonstrated *in vitro* by keeping cells in saline or plateau phase for 6 hours after irradiation and *in vivo* by delayed removal and assay of animal tumors or cells of normal tissues.
- PLD repair is significant for x-rays but does not occur after neutron irradiation.
- It has been suggested that resistant human tumors (e.g., melanoma) owe their resistance to large amounts of PLD repair. This is still controversial.

**Sublethal Damage Repair**
- SLD repair is an operational term that describes the increase in survival if a dose of radiation is split into two fractions separated in time.
- The half-time of SLD repair in mammalian cells is about 1 hour, but it may be longer in late-responding normal tissues *in vivo*.
- SLD repair occurs in tumors and normal tissues *in vivo* as well as in cells cultured *in vitro*.
- The repair of SLD reflects the repair of DNA breaks before they can interact to form lethal chromosomal aberrations.
- SLD repair is significant for x-rays, but almost nonexistent for neutrons.

**Dose-Rate Effect**
- If the radiation dose rate is reduced from about 1 Gy/min to 0.3 Gy/h, there is a reduction in the cell killing from a given dose, because SLD repair occurs during the protracted exposure.
- As the dose rate is reduced, the slope of the survival curve becomes shallower (D0 increases), and the shoulder tends to disappear.
- In some cell lines, an inverse dose-rate effect is evident (i.e., reducing the dose rate increases the proportion of cells killed) owing to the accumulation of cells in G2, which is a sensitive phase of the cycle.

**Brachytherapy**
- Implanting sources into or close to a tumor is known as brachytherapy (from the Greek word *brachy*, meaning “short”) or endocurietherapy (from the Greek word *endo*, meaning “within”).
- Intracavitary radiotherapy involves placing radioactive sources into a body cavity close to a tumor. The most common example is the treatment of carcinoma of the uterine cervix.
- Interstitial therapy involves implanting radioactive sources directly into the tumor and adjacent normal tissue.
- Temporary implants, which formerly utilized radium needles, now are performed most often with iridium-192 wires or seeds.
- If the implant is used as a sole treatment, a commonly used dose is 50 to 70 Gy in
5 to 9 days. Total dose should be adjusted for dose rate. Clinical studies show that both tumor control and late effects vary with dose rate for a given total dose. Often, the implant is used as a boost to external beam therapy, and only half the treatment is given with the implant.

- Because of their small size and low photon energy, iridium-192 seeds are suitable for use with computer-controlled remote afterloaders.
- Permanent implants can be used with radionuclides (such as iodine-125 or palladium-103) that have relatively short half-lives.
- Several novel radionuclides are being considered as sources for brachytherapy. Most emit low-energy photons, which simplifies the problems of radiation protection.

Radiolabeled Immunoglobulin Therapy

- In the early days of radiolabeled immunoglobulin therapy, radiolabeled polyclonal antibodies were used. These were replaced with murine monoclonal antibodies. More recently, chimeric mouse–human antibodies, which are human antibodies derived by tissue culture or produced in genetically altered mice, have become available. Finally, synthetically derived antibodies have been produced.
- Iodine-131 largely has been replaced by pure β-ray emitters such as yttrium-90, resulting in an increased tumor dose and decreased total body toxicity.
- SPECT can now be used to visualize the tumor, using the bremsstrahlung from the β-rays, so it is no longer necessary to add a γ-emitter when using yttrium-90.

- Radiolabeled immunoglobulin therapy has produced promising results in unresectable primary hepatoma and in patients with Hodgkin lymphoma. It has been used with varying degrees of success for a wide range of other malignancies.

BIBLIOGRAPHY


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A number of chemical and pharmacologic agents that modify the biologic effect of ionizing radiations have been discovered. None is simpler than oxygen, none produces such a dramatic effect, and, as it turns out, no other agent has such obvious practical implications.

The oxygen effect was observed as early as 1912 in Germany by Swartz, who noted that the skin reaction produced on his forearm by a radium applicator was reduced if the applicator was pressed hard onto the skin. He attributed this to the interruption in blood flow. By 1921, it had been noted by Holthusen that *Ascaris* eggs were relatively resistant to radiation in the absence of oxygen, a result wrongly attributed to the absence of cell division under these conditions. The correlation between radiosensitivity and the presence of oxygen was made by Petry in 1923 from a study of the effects of radiation on vegetable seeds. All of these results were published in the German literature but were apparently little known in the English-speaking world.

In England in the 1930s, Mottram explored the question of oxygen in detail, basing his investigations on work of Crabtree and Cramer on the survival of tumor slices irradiated in the presence or absence of oxygen. He also discussed the importance of these findings to radiotherapy. Mottram began a series of experiments that culminated in a quantitative measurement of the oxygen effect by his colleagues Gray and Read, using as a biologic test system the growth inhibition of the primary root of the broad bean *Vicia faba*.

### THE NATURE OF THE OXYGEN EFFECT

Survival curves for mammalian cells exposed to x-rays in the presence and absence of oxygen are illustrated in Figure 6.1. The ratio of doses administered under hypoxic to aerated conditions needed to achieve the same biologic effect is called the oxygen enhancement ratio (OER). For sparsely ionizing radiations, such as x- and γ-rays, the OER at high doses has a value of between 2.5 and 3.5. The OER has been determined for various chemical and biologic systems with different end points, and its value for x-rays and γ-rays always tends to fall in this range. There is some evidence that for rapidly growing cells cultured in vitro, the OER has a smaller value of about 2.5 at lower doses, on the order of the daily dose per fraction generally used in radiotherapy. This is believed to result from the variation of OER with the phase of the cell cycle: Cells in G1 phase have a lower OER than those in S, and because G1 cells are more radiosensitive, they dominate the low-dose region of the survival curve. For this reason, the OER of an asynchronous population is slightly smaller at low doses than at high doses. This result has been demonstrated for fast-growing cells cultured in vitro, for which precise survival measurements are possible, but would be difficult to
the survival curve does not have an initial shoulder. In this case, survival estimates made in the presence or absence of oxygen fall along a common line; the OER is unity—in other words, there is no oxygen effect. For radiations of intermediate ionizing density, such as neutrons, the survival curves have a much reduced shoulder. In this case, the oxygen effect is apparent, but it is much smaller than is the case for x-rays. In the example shown in Figure 6.2, the OER for neutrons is about 1.6.

In summary, the oxygen effect is large and important in the case of sparsely ionizing radiations, such as x-rays; is absent for densely ionizing radiations, such as α-particles; and has an intermediate value for fast neutrons.

THE TIME AT WHICH OXYGEN ACTS AND THE MECHANISM OF THE OXYGEN EFFECT

For the oxygen effect to be observed, oxygen must be present during the radiation exposure or, to be precise, during or within microseconds after the radiation exposure. Sophisticated experiments have been performed in which oxygen, contained in a chamber at high pressure, was allowed to “explode” onto a single layer of bacteria (and later mammalian cells) at various times before or after irradiation with a 2-μs electron pulse from a linear accelerator. It was found that oxygen need not be present during the irradiation to sensitize but could be added afterward, provided the delay was not too long. Some sensitization occurred with oxygen added as late as 5 milliseconds after irradiation.

Experiments such as these shed some light on the mechanism of the oxygen effect. There is general agreement that oxygen acts at the level of the free radicals. The chain of events from the absorption of radiation to the final expression of biologic damage has been summarized as follows: The absorption of radiation leads to the production of fast-charged particles. The charged particles, in passing through the biologic material, produce several ion pairs. These ion pairs have very short life spans (about 10^{-10} second) and produce free radicals, which are highly reactive molecules because they have an unpaired valence electron. The free radicals are important because although their life spans are only about...
cannot take place in the absence of oxygen; since then, many of the ionized target molecules are able to repair themselves and recover the ability to function normally. In a sense, then, oxygen may be said to "fix" or make permanent the radiation lesion. This is known as the oxygen fixation hypothesis. The process is illustrated in Figure 6.3.


10^{-5} second, that is appreciably longer than that of the ion pairs. To a large extent, it is these free radicals that break chemical bonds, produce chemical changes, and initiate the chain of events that result in the final expression of biologic damage; however, it has been observed that the extent of the damage depends on the presence or absence of oxygen.

If molecular oxygen is present, DNA reacts with the free radicals (R·). The DNA radical can be chemically restored to its reduced form through reaction with a sulfhydryl (SH) group. However, the formation of RO2·, an organic peroxide, represents a nonrestorable form of the target material; that is, the reaction results in a change in the chemical composition of the material exposed to the radiation. This reaction cannot take place in the absence of oxygen; since then, many of the ionized target molecules are able to repair themselves and recover the ability to function normally. In a sense, then, oxygen may be said to “fix” or make permanent the radiation lesion. This is known as the oxygen fixation hypothesis. The process is illustrated in Figure 6.3.

### THE CONCENTRATION OF OXYGEN REQUIRED

A question of obvious importance is the concentration of oxygen required to potentiate the effect of radiation. Is the amount required small or large? Many investigations have been performed using bacteria, plants, yeast, and mammalian cells, and the similarities between them are striking.
obtained under experimental conditions (10 ppm of oxygen in the gas phase). The introduction of a very small quantity of oxygen, 100 ppm, is readily noticeable in a change in the slope of the survival curve. A concentration of 2,200 ppm, which is about 0.22% oxygen, moves the survival curve about halfway toward the fully aerated condition.

Other experiments have shown that, generally, by the time a concentration of oxygen corresponding to 2% has been reached, the survival curve is virtually indistinguishable from that obtained under conditions of normal aeration. Furthermore, increasing the amount of oxygen present from that characteristic of air to 100% oxygen does not further affect the slope of the curve. This has led to the more usual “textbook representation” of the variation of radiosensitivity with oxygen concentration as shown in Figure 6.5. The term used here to represent radiosensitivity is proportional to the reciprocal of the \( D_0 \) of the survival curve. It is arbitrarily assigned a value of unity for anoxic conditions. As the oxygen concentration increases, the biologic material becomes progressively more sensitive to radiation, until, in the presence of 100% oxygen, it is about three times as sensitive as under complete anoxia. Note that the rapid change of radiosensitivity occurs as the partial pressure of oxygen is increased from zero to about 30 mm Hg (5% oxygen). A further increase in oxygen tension to an atmosphere of pure oxygen has

The simple way to visualize the effect of oxygen is by considering the change of slope of the mammalian cell survival curve. Figure 6.4 is a dramatic representation of what happens to the survival curve in the presence of various concentrations of oxygen. Curve A is characteristic of the response under conditions of equilibration with air. Curve B is a survival curve for irradiation in as low a level of hypoxia as usually can be
CHRONIC AND ACUTE HYPOXIA

It is important to recognize that hypoxia in tumors can result from two quite different mechanisms. Chronic hypoxia results from the limited diffusion distance of oxygen through tissue that is respiring. The distance to which oxygen can diffuse is largely limited by the rapid rate at which it is metabolized by respiring tumor cells. Many tumor cells may remain hypoxic for long periods. In contrast to chronic hypoxia, acute hypoxia is the result of the temporary closing of a tumor blood vessel owing to the malformed vasculature of the tumor, which lacks smooth muscle and often has an incomplete endothelial lining and basement membrane. Tumor cells are exposed to a continuum of oxygen concentrations, ranging from the highest in cells surrounding the capillaries to almost anoxic conditions in cells more distant from the capillaries. This is significant because both chronic and acute hypoxia have been shown to drive malignant progression.

Chronic Hypoxia

As already mentioned, radiotherapists began to suspect that oxygen influences the radiosensitivity of tumors in the 1930s. It was, however, a paper by Thomlinson and Gray in 1955 that triggered the tremendous interest in oxygen as a factor in radiotherapy; they described the phenomenon of chronic hypoxia that they observed in their histologic study of fresh specimens of bronchial carcinoma. Cells of the stratified squamous epithelium, normal or malignant, generally remain in contact with one another; the vascular stroma on which their nutrition depends lies in contact with the epithelium, but capillaries do not penetrate between the cells. Tumors that arise in this type of tissue often grow in solid cords that, seen in section, appear to be circular areas surrounded by stroma. The centers of large tumor areas are necrotic and are surrounded by intact tumor cells, which consequently appear as rings. Figure 6.6A, reproduced from Thomlinson and Gray, shows a transverse section of a tumor cord and is typical of areas of a tumor in which necrosis is not far advanced. Figure 6.6B shows large areas of necrosis separated from stroma by a narrow band of tumor cells about 100 μm wide.

By viewing a large number of these samples of human bronchial carcinomas, Thomlinson and Gray recognized that as the tumor cord lengthened.
went on to calculate the distance to which oxygen could diffuse in respiring tissue and came up with a distance of about 150 μm. This was close enough to the thickness of viable tumor cords on their histologic sections for them to conclude that oxygen depletion was the principal factor leading to the development of necrotic areas in tumors. Using more appropriate values grows larger, the necrotic center also enlarges, so that the thickness of the sheath of viable tumor cells remains essentially constant. This is illustrated in Figure 6.7.

The obvious conclusion was that tumor cells could proliferate and grow actively only if they were close to a supply of oxygen or nutrients from the stroma. Thomlinson and Gray then

**FIGURE 6.6** Transverse sections of tumor cords surrounded by stroma from human carcinoma of the bronchus. **A**: A typical tumor area in which necrosis is not far advanced. **B**: Large areas of necrosis separated from the stroma by a band of tumor cells about 100 μm wide. (From Thomlinson RH, Gray LH. The histological structure of some human lung cancers and the possible implications for radiotherapy. *Br J Cancer*. 1955;9:539–549, with permission.)
of oxygen diffusion coefficients and consumption values, a better estimate of the distance that the oxygen can diffuse in respiring tissue is about 70 μm. This, of course, varies from the arterial to the venous end of a capillary, as illustrated in Figure 6.8.

By histologic examination of sections, it is possible to distinguish only two classes of cells: (1) those that appear to be proliferating well and (2) those that are dead or dying. Between these two extremes, and assuming a steadily decreasing oxygen concentration, one would expect a region in which cells would be at an oxygen tension high enough for cells to be clonogenic but low enough to render the cells protected from the effect of ionizing radiation. Cells in this region would be relatively protected from a treatment with x-rays because of their low oxygen tension and could provide a focus for the subsequent regrowth of the tumor (Fig. 6.8). Based on these ideas, it was postulated that the presence of a relatively small proportion of hypoxic cells in tumors could limit the success of radiotherapy in some clinical situations.
These ideas about the role of oxygen in cell killing dominated the thinking of radiobiologists and radiotherapists in the late 1950s and early 1960s. A great deal of thought and effort was directed toward solving this problem. The solutions proposed included the use of high-pressure oxygen chambers, the development of novel radiation modalities such as neutrons, negative \(\pi\)-mesons, and heavy-charged ions, and the development of hypoxic cell sensitizers.

**Acute Hypoxia**

Regions of **acute hypoxia** develop in tumors as a result of the temporary closing or blockage of a particular blood vessel. If this blockage were permanent, the cells downstream, of course, would eventually die and be of no further consequence. There is, however, good evidence that tumor blood vessels open and close in a random fashion, so that different regions of the tumor become hypoxic intermittently. In fact, acute hypoxia results from transient fluctuations in blood flow because of the malformed vasculature. At the moment when a dose of radiation is delivered, a proportion of tumor cells may be hypoxic, but if the radiation is delayed until a later time, a different group of cells may be hypoxic. The occurrence of **acute hypoxia** was postulated in the early 1980s by Martin Brown and was later demonstrated unequivocally in rodent tumors by Chaplin and his colleagues. Figure 6.9, which illustrates how acute hypoxia is caused by fluctuating blood flow, also depicts the difference between acute and chronic hypoxia. In contrast to acutely hypoxic cells, chronically hypoxic cells are less likely to become reoxygenated and will die unless they are able to access a blood supply.

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**THE FIRST EXPERIMENTAL DEMONSTRATION OF HYPOXIC CELLS IN A TUMOR**

The dilution assay technique, described in Chapter 21, was used by Powers and Tolmach to investigate the radiation response of a solid subcutaneous lymphosarcoma in the mouse. Survival estimates were made for doses from 2 to 25 Gy. The results are shown in Figure 6.10, in which the dose on a linear scale is plotted against the fraction of surviving cells on a logarithmic scale.

The survival curve for this solid tumor clearly consists of two separate components. The first, up to a dose of about 9 Gy, has a \(D_0\) of 1.1 Gy. The second has a shallower \(D_0\) of 2.6 Gy. This biphasic survival curve has a final slope about 2.5 times shallower than the initial portion, which strongly suggests that the tumor consists of two separate groups of cells, one oxygenated and the other hypoxic. If the shallow component of the curve is extrapolated backward to cut the surviving-fraction axis, it does so at a survival level of about 1%. From this, it may be inferred that about 1% of the clonogenic cells in the tumor were deficient in oxygen.

The response of this tumor to single doses of radiation of various sizes is explained readily on
this basis. If 99% of the cells are well oxygenated and 1% are hypoxic, the response to lower doses is dominated by the killing of the well-oxygenated cells. For these doses, the hypoxic cells are depopulated to a negligibly small extent. Once a dose of about 9 Gy is exceeded, however, the oxygenated compartment of the tumor is depopulated severely, and the response of the tumor is characteristic of the response of hypoxic cells.

This biphasic survival curve was the first unequivocal demonstration that a solid tumor could contain cells sufficiently hypoxic to be protected from cell killing by x-rays but still clonogenic and capable of providing a focus for tumor regrowth.

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**PROPORTION OF HYPOXIC CELLS IN VARIOUS ANIMAL TUMORS**

Over the years, many investigators have determined the fraction of hypoxic cells in various tumors in experimental animals. The most satisfactory and most widely used method is to obtain paired survival curves (Fig. 6.11A). The steepest curve relates to a fully oxygenated population of cells; the uppermost curve, to a population made up entirely of hypoxic cells. The intermediate curves refer to mixed populations of oxygenated cells with varying proportions of hypoxic cells. At low doses, the survival curve for a mixed population closely follows that for the oxygenated population. At higher doses, the number of surviving oxygenated cells is negligible compared with the number of anoxic cells, and consequently, the curve representing the mixed population is parallel to (i.e., has the same slope as) the curve for the hypoxic population. The fraction of hypoxic cells in the tumor determines the distance between the parallel terminal slopes of the dose-response curves, as shown in Figure 6.11A. This fraction is identical to the ratio of the surviving cells from the partially hypoxic tumor to those from the entirely hypoxic tumor.

In practice, the procedure is as follows: Survival measurements are made at several dose levels under two different conditions:

1. The animal (e.g., a mouse) is asphyxiated several minutes before irradiation by breathing nitrogen. Under these conditions, all of the tumor cells are hypoxic, and the data points obtained define a line comparable to the upper curve in Figure 6.11A.
2. The animal is alive and breathing air when irradiated, so that the proportion of hypoxic cells in the tumor is at its normal level. The
Figure 6.11 A: Theoretic survival curves for cell populations containing different fractions of hypoxic cells. The fraction of hypoxic cells in each population determines the distance between its survival curve and the curve for the completely hypoxic population. From the relative radiosensitivity at any dose level at which the survival curves are approximated by parallel lines, the fraction of hypoxic cells can be determined from the ratio of survival of the completely and partially hypoxic populations, as indicated by the vertical lines A-A, B-B, and so on. This illustration is based on the model proposed by Hewitt and Wilson. (Adapted from van Putten LM, Kallman RF. Oxygenation status of a transplantable tumor during fractionated radiotherapy. J Natl Cancer Inst. 1968;40:441–451, with permission.)

B: The proportion of hypoxic cells in a mouse tumor. The biphasic curve labeled air curve represents data for cells from tumors irradiated in mice breathing air, which are therefore a mixture of aerated and hypoxic cells. The hypoxic curve is for cells irradiated in mice asphyxiated by nitrogen breathing or for cells irradiated in vitro in nitrogen, so that they are all hypoxic. The air curve is for cells irradiated in vitro in air. The proportion of hypoxic cells is the ratio of the air to hypoxic curves or the vertical separation between the curves because the surviving fraction is on a logarithmic scale. (Courtesy of Dr. Sara Rockwell; based on data of Moulder and Rockwell and of Rockwell and Kallman.)

Data points obtained define a lower line typical of a mixed population of hypoxic and oxygenated cells. The vertical separation between the two lines gives the proportion of hypoxic cells characteristic of that particular tumor.

An example of experimental data for a determination of the hypoxic fraction in a mouse tumor is shown in Figure 6.11B. Hypoxic fractions can also be calculated from a comparison of the TCD50 values (i.e., the doses at which 50%
of the tumors are locally controlled) for clamped and unclamped tumors or from a comparison of growth delays from tumors irradiated under these two conditions. Any of these methods involves several assumptions, notably that cells made hypoxic artificially have the same sensitivity as those that have respired to this condition in the tumor naturally and that the tumor is composed of two distinct populations, one aerated and the other hypoxic, with nothing falling in between. Consequently, measured values for hypoxic fractions can serve only as a guide and must not be taken too seriously.

Moulder and Rockwell published a survey of all published data on hypoxic fractions and reported that of 42 tumor types studied, 37 were found to contain hypoxic cells in at least one study. Hypoxic fractions range from 0% to 50%, with a tendency for many results to average about 15%. Comparable measurements cannot be made in human tumors, of course, to determine precisely the proportion of hypoxic cells.

**EVIDENCE FOR HYPOXIA IN HUMAN TUMORS**

Over the last decade, various techniques have been used to determine the oxygenation of human tumors, including measuring the distance between tumor cells and vessels in histologic sections, determining the oxygen saturation of hemoglobin, and monitoring changes in tumor metabolism. These techniques have been replaced by newer methods, including oxygen probes, hypoxia markers, the comet assay, and noninvasive imaging. Although each of these techniques has strengths and weaknesses, together they convincingly demonstrate that hypoxia is a common feature of human solid tumors that can influence both the malignant progression and the response of tumors to therapy. In most studies, the assessment of hypoxia in human tumors has been based largely on oxygen-probe measurements. This approach has been used to group patients based on their median pO2 values, but disregards a great deal of information that is obtained in the process, especially the heterogeneity of oxygen measurements in solid tumors. Although oxygen probes are considered the “gold standard” for measuring tumor pO2, newer noninvasive techniques will supplant them in the future.
surrounding the blood vessels are not visible because they are fully oxygenated and do not express HIF-1 or reduce the pimonidazole. As would be expected, there is a gradient in oxygen tension away from blood vessels, with therapeutically relevant hypoxic cells situated at some distance.

### REOXYGENATION

Van Putten and Kallman determined the proportion of hypoxic cells in a transplantable sarcoma in the mouse. This tumor, which was of spontaneous origin, was transplanted from one generation of animals to the next by inoculating a known number of tumor cells subcutaneously. The tumor was allowed to grow for 2 weeks, by which time it had reached a size suitable for the experiment. The tumor was irradiated in vivo and then excised and made into a suspension of cells. The proportion of hypoxic cells was determined by the method described in Figure 6.11. These experiments have far-reaching implications in radiotherapy. The fact that they are fully oxygenated and do not express HIF-1 or reduce the pimonidazole. As would be expected, there is a gradient in oxygen tension away from blood vessels, with therapeutically relevant hypoxic cells situated at some distance.

**REOXYGENATION**

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The researchers found that for this mouse sarcoma, the proportion of hypoxic cells in the untreated tumor was about 14%. The vital contribution made by van Putten and Kallman involved a determination of the proportion of hypoxic cells in this tumor after various fractionated radiation treatments. When groups of tumors were exposed to five daily doses of 1.9 Gy delivered Monday through Friday, the proportion of hypoxic cells was determined on the following Monday to be 18%. In another experiment, four daily fractions were given Monday through Thursday, and the proportion of hypoxic cells measured the following day, Friday, was found to be 14%.

These experiments have far-reaching implications in radiotherapy. The fact that the
The proportion of hypoxic cells in the tumor is about the same at the end of a fractionated radiotherapy regimen as in the untreated tumor, demonstrating that during the course of the treatment, some hypoxic cells become oxygenated. If this were not the case, then the proportion of hypoxic cells would increase during the course of the fractionated treatment because the radiation depopulates the aerated cell compartment more than the hypoxic cell compartment. This phenomenon, by which hypoxic cells become oxygenated after a dose of radiation, is termed **reoxygenation**. The oxygen status of cells in a tumor is not static; it is dynamic and constantly changing.

The process of reoxygenation is illustrated in Figure 6.13. A modest dose of x-rays to a mixed population of aerated and hypoxic cells results in significant killing of aerated cells but little killing of hypoxic cells. Consequently, the viable cell population immediately after irradiation is dominated by hypoxic cells. If sufficient time is allowed before the next radiation dose, the process of reoxygenation restores the proportion of hypoxic cells to about 15%. If this process is repeated many times, the tumor cell population is depleted, despite the intransigence to killing by x-rays of the cells deficient in oxygen. In other words, if reoxygenation is efficient between dose fractions, the presence of hypoxic cells does not have a significant effect on the outcome of a multifraction regimen.

**TIME SEQUENCE OF REOXYGENATION**

In the particular tumor system used by van Putten and Kallman, the proportion of hypoxic cells returned to its original pretreatment level by 24 hours after delivery of a fractionated dosage schedule. Kallman and Bleehen reported experiments in which the proportion of hypoxic cells in the same transplantable mouse sarcoma
was determined at various times after delivery of a single dose of 10 Gy. Their results are shown in Figure 6.14; the shape of the curve indicates that in this particular tumor, the process of reoxygenation is very rapid indeed.

Similar results subsequently have been reported by several researchers using various tumor systems. The patterns of reoxygenation after irradiation observed in several different animal tumor systems are summarized in Figure 6.15. Four of the five animal tumors show efficient and rapid reoxygenation, with the proportion of hypoxic cells returning to or even falling below the pretreatment level in a day or two. The time sequence, however, is not the same for the five types of tumors. In particular, the mammary carcinoma investigated by Howes shows a minimum proportion of hypoxic cells that is very much lower than that characteristic of the unirradiated tumor. This point is reached 3 days after the delivery of a single large dose of radiation. The only one of the five tumors that does not show any significant rapid reoxygenation is the osteosarcoma studied by van Putten, also illustrated in Figure 6.15.

■ MECHANISM OF REOXYGENATION

In experimental animals, some tumors take several days to reoxygenate; in others, the process appears to be complete within 1 hour or so. In a few tumors, both fast and slow components to reoxygenation are evident. The differences of timescale reflect the different types of hypoxia that are being reversed, chronic versus acute. In the long term, a restructuring or a revascularization of the tumor occurs as the cells killed by the radiation are broken down and removed from the population. As the tumor shrinks in size, surviving cells that previously were beyond the range of oxygen diffusion are closer to a blood supply and so reoxygenate. This slow component of reoxygenation, taking place over a period of days as the tumor shrinks, involves reoxygenation of cells that were chronically hypoxic. By contrast, the fast component of reoxygenation, which is complete within hours, is caused by the reoxygenation of acutely hypoxic cells; that is, those cells that were hypoxic at the time of irradiation because they were in regions in which a blood vessel was temporarily closed quickly reoxygenate when that vessel is reopened.

■ THE IMPORTANCE OF REOXYGENATION IN RADIOTHERAPY

The process of reoxygenation has important implications in practical radiotherapy. If human tumors do in fact reoxygenate as rapidly and efficiently as most of the animal tumors studied, then the use of a multifraction course of radiotherapy, extending over a long period, may well be all that is required to deal effectively with any hypoxic cells in human tumors.

The reoxygenation studies with mouse mammary carcinoma, included in Figure 6.15, indicate that by 2 to 3 days after a dose of radiation, the proportion of hypoxic cells is actually
lower than in untreated tumors. Consequently, it was predicted that several large doses of x-rays given at 48-hour intervals would virtually eliminate the problem of hypoxic cells in this tumor. Fowler and his colleagues indeed showed that for the eradication of this tumor, the preferred x-ray schedule was five large doses in 9 days. These results suggest that x-irradiation can be an extremely effective form of therapy, but ideally requires a sharply optimal choice of fractionation pattern. Making this choice, however, demands a detailed knowledge of the time course of reoxygenation in the particular tumor to be irradiated. Unfortunately, however, this information is available for only a few animal tumors and is impossible to obtain at present for human tumors. Indeed, in humans, it is not known with certainty whether any or all tumors reoxygenate, although the evidence from radiotherapy clinics that many tumors are eradicated with doses on the order of 60 Gy given in 30 treatments argues strongly in favor of reoxygenation because the presence of a very small proportion of hypoxic cells would make “cures” unlikely at these dose levels. It is an attractive hypothesis that some of the human tumors that do not respond to conventional radiotherapy are those that do not reoxygenate quickly and efficiently.

- HYPOXIA AND CHEMoresistance

Hypoxia can also decrease the efficacy of some chemotherapeutic agents owing to fluctuating blood flow, drug diffusion distance, and decreased proliferation. In addition, some chemotherapeutic agents that induce DNA damage, such as doxorubicin and bleomycin, are less...
efficient at killing hypoxic tumor cells in part because of decreased free-radical generation. Experimental animal studies have shown that 5-FU, methotrexate, and cisplatin are less effective at killing hypoxic cells than they are at killing normoxic tumor cells. Furthermore, hypoxic tumor regions are frequently associated with a low pH that can also diminish the activity of some chemotherapy agents. (For more about chemotherapy agents and hypoxia, see Chapter 27.)

**HYPOXIA AND TUMOR PROGRESSION**

Evidence that low-oxygen conditions play an important role in malignant progression comes from studies of the correlation between tumor oxygenation and treatment outcome in patients, as well as from laboratory studies in cells and animals. A clinical study in Germany in the 1990s showed a correlation between local control in advanced carcinoma of the cervix treated by radiotherapy and oxygen-probe measurements. Specifically, patients in whom the probe measurements indicated pO₂s greater than 10 mm Hg did better than those with pO₂s less than 10 mm Hg. This suggested that the presence of hypoxic cells limited the success of radiotherapy. Later studies, however, indicated a similar improvement in outcome for patients with better oxygenated tumors if the treatment was by surgery rather than radiotherapy. This suggests that the correct interpretation is that hypoxia is a general indicator of tumor aggression in these patients, rather than the initial view that hypoxia conferred radioresistance on some cells. A Canadian study supported the concept that hypoxia drove malignant progression of cervical carcinomas in node-negative patients, as most of the failures occurred in the poorly oxygenated tumors with distant tumor spread outside the pelvis.

Studies carried out in the United States on patients receiving radiotherapy for soft tissue sarcoma highlighted the correlation between tumor oxygenation and the frequency of distant metastases. Seventy percent of those patients with pO₂s less than 10 mm Hg developed distant metastases, versus 35% of those with pO₂s greater than 10 mm Hg. This study is particularly compelling because the primary tumor was eradicated in all patients regardless of the level of oxygenation; only the incidence of metastases varied between high and low pO₂ values. These data were subsequently confirmed in a Danish study in which 28 patients with soft tissue sarcoma exhibited an increased risk of metastatic spread if they possessed low tumor pO₂ values. This argues strongly that the level of tumor oxygenation influences the aggressiveness of the tumor. This topic is discussed further in Chapter 26.

**SUMMARY OF PERTINENT CONCLUSIONS**

- The presence or absence of molecular oxygen dramatically influences the biologic effect of x-rays.
- The OER is the ratio of doses under hypoxic to aerated conditions that produce the same biologic effect.
- The OER for x-rays is about 3 at high doses and is possibly lower (about 2) at doses less than about 2 Gy.
- The OER decreases as linear energy transfer increases. The OER approaches unity (i.e., no oxygen effect) for α-particles. For neutrons, the OER has an intermediate value of about 1.6.
- To produce its effect, molecular oxygen must be present during the radiation exposure or at least during the lifetime of the free radicals generated by the radiation.
- Oxygen “fixes” (i.e., makes permanent) the damage produced by free radicals. In the absence of oxygen, damage produced by the indirect action may be repaired.
- Only a small quantity of oxygen is required for radiosensitization; 0.5% oxygen (pO₂ of about 3 mm Hg) results in a radiosensitivity halfway between hypoxia and full oxygenation.
- There are two forms of hypoxia that are the consequence of different mechanisms: chronic hypoxia and acute hypoxia.
- Chronic hypoxia results from the limited diffusion range of oxygen through respiring tissue.
- Acute hypoxia is a result of the temporary closing of tumor blood vessels and is therefore transient.
- In either case, there may be cells present during irradiation that are at a sufficiently low oxygen tension to be intransigent to killing by x-rays but high enough to be viable.
Most transplantable tumors in animals have been shown to contain hypoxic cells that limit curability by single doses of x-rays. Hypoxic fractions vary from 0% to 50%, with a tendency to average about 15%.

There is strong evidence that human tumors contain hypoxic cells. This evidence includes histologic appearance, oxygen-probe measurements, the binding of nitroimidazoles, PET and SPECT studies, and pretreatment hemoglobin levels.

Oxygen probes with fast response times, implanted in a tumor and moving quickly under computer control, may be used to obtain the oxygen profile of a tumor.

Hypoxia in tumors can be visualized by the use of hypoxia markers such as pimonidazole or hypoxia-inducible factors.

Reoxygenation is the process by which cells that are hypoxic at the time of irradiation become oxygenated afterward.

The extent of reoxygenation and the rapidity with which it occurs vary widely for different experimental animal tumors.

If reoxygenation is rapid and complete, hypoxic cells have little influence on the outcome of a fractionated radiation schedule.

The “slow” component is caused by the reoxygenation of chronically hypoxic cells as the tumor shrinks. The “fast” component of reoxygenation is caused by the reoxygenation of acutely hypoxic cells as tumor blood vessels open and close.

Reoxygenation cannot be measured in human tumors, but presumably it occurs, at least in those tumors controlled by conventional fractionated radiotherapy.

There is clinical evidence that in addition to causing radioresistance, hypoxia may play an important role in malignant progression and in metastasis.

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If radiation is absorbed in biologic material, ionizations and excitations occur that are not distributed at random but tend to be localized along the tracks of individual charged particles in a pattern that depends on the type of radiation involved. For example, x-ray photons give rise to fast electrons, particles carrying unit electrical charge and having very small mass; neutrons, on the other hand, give rise to recoil protons, particles again carrying unit electrical charge but having mass nearly 2,000 times greater than that of the electron. $\alpha$-particles carry two electrical charges on a particle four times as heavy as a proton. The charge-to-mass ratio for $\alpha$-particles therefore differs from that for electrons by a factor of about 8,000.

As a result, the spatial distribution of the ionizing events produced by different particles varies enormously. This is illustrated in Figure 7.1. The background is an electron micrograph of a human liver cell. The white dots generated by a computer simulate ionizing events. The lowest track represents a low-energy electron, such as might be set in motion by diagnostic x-rays. The primary events are well separated in space, and for this reason, x-rays are said to be *sparsely* ionizing. The second track from the bottom represents an electron set in motion by cobalt-60 $\gamma$-rays, which is even more sparsely ionizing. For a given particle type, the density of ionization decreases as the energy goes up. The third track from the bottom represents a proton that might be set in motion by a fission spectrum neutron from a nuclear reactor; a dense column of ionization is produced, so the radiation is referred to as *densely* ionizing. The uppermost track refers to a 10-MeV proton, such as may be set in motion by the high-energy neutrons used for radiotherapy. The track is intermediate in ionization density.

**LINEAR ENERGY TRANSFER**

Linear energy transfer (LET) is the energy transferred per unit length of the track. The special unit usually used for this quantity is kiloelectron volt per micrometer (keV/μm) of unit density material. In 1962, the International Commission on Radiological Units defined this quantity as follows:

The LET ($L$) of charged particles in medium is the quotient of $dE/dl$, where $dE$ is the average energy locally imparted to the medium by a charged particle of specified energy in traversing a distance of $dl$.

That is,

$$L = \frac{dE}{dl}$$

LET is an average quantity because at the microscopic level, the energy per unit length of track varies over such a wide range. Indeed, the range is so large that some believe that the concept of LET has little meaning. This can be illustrated by the story of a Martian visitor to Earth who arrives knowing that the Earth is inhabited by living creatures with an average mass of 1 g. Not only is this information of very little use, but it also may be positively misleading, particularly if the first
living creature that the Martian encounters is an elephant. An average quantity has little meaning if individual variation is great.

The situation for LET is further complicated by the fact that it is possible to calculate an average in many different ways. The most commonly used method is to calculate the track average, which is obtained by dividing the track into equal lengths, calculating the energy deposited in each length, and finding the mean. The energy average is obtained by dividing the track into equal energy increments and averaging the lengths of the track over which these energy increments are deposited. These methods are illustrated in Figure 7.2.

In the case of either x-rays or monoenergetic charged particles, the two methods of averaging

\[ \text{LET} = \frac{\text{Average energy deposited per unit length of track (keV/µm)}}{\text{Track average}} \]

\[ \text{Energy average} \]

**FIGURE 7.1** Variation of ionization density associated with different types of radiation. The background is an electron micrograph of a human cell. The white dots are a computer simulation representing ionizations. **Top to bottom:** A 10-MeV proton, typical of the recoil protons produced by high-energy neutrons used for radiotherapy. The track is intermediate in ionization density. Also shown is a secondary 1-keV δ-ray, an electron set in motion by the proton. A 500-keV proton, produced by lower-energy neutrons (e.g., from fission spectrum) or by higher-energy neutrons after multiple collisions. The ionizations form a dense column along the track of the particle. A 1-MeV electron, produced, for example, by cobalt-60 γ-rays. This particle is very sparsely ionizing. A 5-keV electron, typical of secondary electrons produced by x-rays of diagnostic quality. This particle is also sparsely ionizing but a little denser than the higher-energy electron. (Courtesy of Dr. Albrecht Kellerer.)

**FIGURE 7.2** Linear energy transfer (LET) is the average energy deposited per unit length of track. The track average is calculated by dividing the track into equal lengths and averaging the energy deposited in each length. The energy average is calculated by dividing the track into equal energy intervals and averaging the lengths of the track that contain this amount of energy. The method of averaging makes little difference for x-rays or for monoenergetic charged particles, but the track average and energy average are different for neutrons.
a measure of the energy absorbed per unit mass of tissue. Equal doses of different types of radiation do not, however, produce equal biologic effects. For example, 1 Gy of neutrons produces a greater biologic effect than 1 Gy of x-rays. The key to the difference lies in the pattern of energy deposition at the microscopic level.

In comparing different radiations, it is customary to use x-rays as the standard. The National Bureau of Standards in 1954 defined relative biologic effectiveness (RBE) as follows:

The RBE of some test radiation \((r)\) compared with x-rays is defined by the ratio \(D_{250}/D_r\), where \(D_{250}\) and \(D_r\) are, respectively, the doses of x-rays and the test radiation required for equal biologic effect.

To measure the RBE of some test radiation, one first chooses a biologic system in which the effect of radiations may be scored quantitatively. To illustrate the process involved, we discuss a specific example. Suppose we are measuring the RBE of fast neutrons compared with 250-kV x-rays, using the lethality of plant seedlings as a test system. Groups of plants are exposed to graded doses of x-rays; parallel groups are exposed to a range of neutron doses. At the end of the period of observation, it is possible to calculate the doses of x-rays and then of neutrons that result in the death of half of the plants in a group. This quantity is known as the LD\(_{50}\), the mean lethal dose. Suppose that for x-rays, the LD\(_{50}\) turns out to be 6 Gy and that for neutrons, the corresponding quantity is 4 Gy. The RBE of

### Table 7.1: Typical Linear Energy Transfer Values

<table>
<thead>
<tr>
<th>Radiation</th>
<th>Linear Energy Transfer, keV/(\mu)m</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cobalt-60 (\gamma)-rays</td>
<td>0.2</td>
</tr>
<tr>
<td>250-kV x-rays</td>
<td>2.0</td>
</tr>
<tr>
<td>10-MeV protons</td>
<td>4.7</td>
</tr>
<tr>
<td>150-MeV proton</td>
<td>0.5</td>
</tr>
<tr>
<td>Track average</td>
<td>—</td>
</tr>
<tr>
<td>Energy average</td>
<td>—</td>
</tr>
<tr>
<td>14-MeV neutrons</td>
<td>12</td>
</tr>
<tr>
<td>2.5-MeV (\alpha)-particles</td>
<td>166</td>
</tr>
<tr>
<td>2-GeV Fe ions</td>
<td>1,000</td>
</tr>
<tr>
<td>(space radiation)</td>
<td></td>
</tr>
</tbody>
</table>

yield similar results. In the case of 14-MeV neutrons, by contrast, the track average LET is about 12 keV/\(\mu\)m and the energy average LET is about 100 keV/\(\mu\)m. The biologic properties of neutrons tend to correlate best with the energy average.

As a result of these considerations, LET is a quantity condemned by the purists as worse than useless, because it can, in some circumstances, be very misleading. It is, however, useful as a simple and naive way to indicate the quality of different types of radiation. Typical LET values for various radiations are listed in Table 7.1. Included are x- and \(\gamma\)-rays used for radiotherapy, protons, neutrons, and naturally occurring \(\alpha\)-particles, as well as high-energy heavy ions encountered by astronauts in space. Note that for a given type of charged particle, the higher the energy, the lower the LET and, therefore, the lower its biologic effectiveness. At first sight, this may be counterintuitive. For example, \(\gamma\)-rays and x-rays both give rise to fast secondary electrons; therefore, 1.1-MV cobalt-60 \(\gamma\)-rays have lower LETs than 250-kV x-rays and are less effective biologically by about 10%. By the same token, 150-MeV protons have lower LETs than 10-MeV protons and therefore are slightly less effective biologically.

### RELATIVE BIOLOGIC EFFECTIVENESS

The amount or quantity of radiation is expressed in terms of the absorbed dose, a physical quantity with the unit of gray (Gy). Absorbed dose is a measure of the energy absorbed per unit mass of tissue. Equal doses of different types of radiation do not, however, produce equal biologic effects. For example, 1 Gy of neutrons produces a greater biologic effect than 1 Gy of x-rays. The key to the difference lies in the pattern of energy deposition at the microscopic level.

In comparing different radiations, it is customary to use x-rays as the standard. The National Bureau of Standards in 1954 defined relative biologic effectiveness (RBE) as follows:

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To measure the RBE of some test radiation, one first chooses a biologic system in which the effect of radiations may be scored quantitatively. To illustrate the process involved, we discuss a specific example. Suppose we are measuring the RBE of fast neutrons compared with 250-kV x-rays, using the lethality of plant seedlings as a test system. Groups of plants are exposed to graded doses of x-rays; parallel groups are exposed to a range of neutron doses. At the end of the period of observation, it is possible to calculate the doses of x-rays and then of neutrons that result in the death of half of the plants in a group. This quantity is known as the LD\(_{50}\), the mean lethal dose. Suppose that for x-rays, the LD\(_{50}\) turns out to be 6 Gy and that for neutrons, the corresponding quantity is 4 Gy. The RBE of...
neutrons compared with x-rays is then simply the ratio 6:4 or 1.5.

The study of RBE is relatively straightforward so long as a test system with a single, unequivocal end point is used. It becomes more complicated if, instead, a test system such as the response of mammalian cells in culture is chosen. Figure 7.3A shows survival curves obtained if mammalian cells in cultures are exposed to a range of doses of, on the one hand, fast neutrons and, on the other hand, 250-kV x-rays. The RBE may now be calculated from these survival curves as the ratio of doses that produce the same biologic effect. If the end point chosen for comparison is the dose required to produce a surviving fraction of 0.01, then the dose of neutrons necessary is 6.6 Gy; the corresponding dose of x-rays is 10 Gy. The RBE, then, is the quotient of 10/6.6 or about 1.5. If the comparison is made at a surviving fraction of 0.6, however, the neutron dose required is only 1 Gy and the corresponding x-ray dose is 3 Gy. The resultant RBE is 3:1 or 3.0. Because the x-ray and neutron survival curves have different shapes, the x-ray survival curve having an initial shoulder, and the neutron curve being an exponential function of dose, the resultant RBE depends on the level of biologic damage (and therefore the dose) chosen. The RBE generally increases as the dose is decreased, reaching a limiting value that is the ratio of the initial slopes of the x-ray and neutron survival curves.

### RELATIVE BIOLOGIC EFFECTIVENESS AND FRACTIONATED DOSES

Because the RBE of more densely ionizing radiations, such as neutrons, varies with the dose per fraction, the RBE for a fractionated regimen with neutrons is greater than for a single exposure, because a fractionated schedule consists of several small doses and the RBE is large for small doses.

Figure 7.3B illustrates a hypothetical treatment with neutrons consisting of four fractions. For a surviving fraction of 0.01, the RBE
for neutrons relative to x-rays is about 2.6. The RBE for the same radiations in Figure 7.3A at the same level of survival was about 1.5 because only single exposures were involved. This is a direct consequence of the larger shoulder that is characteristic of the x-ray curve, which must be repeated for each fraction. The width of the shoulder represents a part of the dose that is “wasted”; the larger the number of fractions, the greater the extent of the wastage. By contrast, the neutron survival curve has little or no shoulder, so there is correspondingly less wastage of dose from fractionation. The net result is that neutrons become progressively more efficient than x-rays as the dose per fraction is reduced and the number of fractions is increased. The same is true, of course, for exposure to continuous low-dose-rate irradiation. The neutron RBE is larger at a low dose rate than for an acute exposure, because the effectiveness of neutrons decreases with dose rate to a much smaller extent than is the case for x- or γ-rays. Indeed, for low-energy neutrons, there is no loss of effectiveness.

### RELATIVE BIOLOGIC EFFECTIVENESS FOR DIFFERENT CELLS AND TISSUES

Even for a given total dose or dose per fraction, the RBE varies greatly according to the tissue or cell line studied. Different cells or tissues are characterized by x-ray survival curves that have large but variable shoulder regions, whereas the shoulder region for neutrons is smaller and less variable. As a consequence, the RBE is different for each cell line. In general, cells characterized by an x-ray survival curve with a large shoulder, indicating that they can accumulate and repair a large amount of sublethal radiation damage, show larger RBEs for neutrons. Conversely, cells for which the x-ray survival curve has little if any shoulder exhibit smaller neutron RBE values.

### RELATIVE BIOLOGIC EFFECTIVENESS AS A FUNCTION OF LINEAR ENERGY TRANSFER

Figure 7.4 illustrates the survival curves obtained for 250-kVp x-rays, 15-MeV neutrons, and 4-MeV α-particles. As the LET increases from about 2 keV/μm for x-rays up to 150 keV/μm for α-particles, the survival curve changes in two important respects. First, the survival curve becomes steeper. Second, the shoulder of the curve becomes progressively smaller as the LET increases. A more common way to represent these data is to plot the RBE as a function of LET (Fig. 7.5). As the LET increases, the RBE increases slowly at first and then more rapidly as the LET increases beyond 10 keV/μm. Between 10 and 100 keV/μm, the RBE increases rapidly with increasing LET and, in fact, reaches a maximum at about 100 keV/μm. Beyond this value for the LET, the RBE again falls to lower values. This is an important effect and is explained in more detail in the next section.

The LET at which the RBE reaches a peak is much the same (about 100 keV/μm) for a wide range of mammalian cells, from mouse to human, and is the same for mutation as an end point as for cell killing.

### THE OPTIMAL LINEAR ENERGY TRANSFER

It is of interest to ask why radiation with an LET of about 100 keV/μm is optimal in terms of producing a biologic effect. At this density
of ionization, the average separation between ionizing events just about coincides with the diameter of the DNA double helix (20 Å or 2 nm). Radiation with this density of ionization has the highest probability of causing a double-strand break (DSB) by the passage of a single charged particle, and DSBs are the basis of most biologic effects, as discussed in Chapter 2. This is illustrated in Figure 7.6. In the case of x-rays, which are more sparsely ionizing, the probability of a single track causing a DSB is low, and in general, more than one track is required. As a consequence, x-rays have a low biologic effectiveness. At the other extreme, much more densely ionizing radiations (with an LET of 200 keV/μm, for example) readily produce DSBs, but energy is “wasted” because the ionizing events are too close together. Because RBE is the ratio of doses producing equal biologic effect, this more densely ionizing radiation has a lower RBE than the optimal LET radiation. The more densely ionizing radiation is just as effective per track, but less effective per unit dose.

**FIGURE 7.5** Variation of relative biologic effectiveness (RBE) with linear energy transfer (LET) for survival of mammalian cells of human origin. The RBE rises to a maximum at an LET of about 100 keV/μm and subsequently falls for higher values of LET. Curves 1, 2, and 3 refer to cell survival levels of 0.8, 0.1, and 0.01, respectively, illustrating that the absolute value of the RBE is not unique but depends on the level of biologic damage and, therefore, on the dose level. (Adapted from Barendsen GW. Responses of cultured cells, tumors, and normal tissues to radiation of different linear energy transfer. *Curr Top Radiat Res Q*. 1968;4:293–356, with permission.)

**FIGURE 7.6** Diagram illustrating why radiation with a linear energy transfer (LET) of 100 keV/μm has the greatest relative biologic effectiveness (RBE) for cell killing, mutagenesis, or oncogenic transformation. For this transfer, the average separation between ionizing events coincides with the diameter of the DNA double helix (i.e., about 20 Å or 2 nm). Radiation of this quality is most likely to produce a double-strand break from one track for a given absorbed dose.
It is possible, therefore, to understand why RBE reaches a maximum value in terms of the production of DSBs, because the interaction of two DSBs to form an exchange-type aberration is the basis of most biologic effects. In short, the most biologically effective LET is that at which there is a coincidence between the diameter of the DNA helix and the average separation of ionizing events. Radiations having this optimal LET include neutrons of a few hundred kiloelectron volts, as well as low-energy protons and α-particles.

■ FACTORS THAT DETERMINE RELATIVE BIOLOGIC EFFECTIVENESS

The discussion of RBE began with a simple illustration of how this ratio may be determined for neutrons compared with x-rays using a simple biologic test system with a single, unequivocal end point, such as the LD₅₀ for plant seedlings. Under these circumstances, RBE is conceptually very simple. In the years immediately after World War II, it was commonplace to see references to the RBE for neutrons, as if it were a single, unique quantity.

Now that more information is available from different biologic systems, many of which allow the researcher to investigate the relationship between biologic response and radiation dose rather than observing one end point at a single dose; it is apparent that RBE is a very complex quantity. RBE depends on the following:

- Radiation quality (LET)
- Radiation dose
- Number of dose fractions
- Dose rate
- Biologic system or end point

Radiation quality includes the type of radiation and its energy, whether electromagnetic or particulate, and whether charged or uncharged.

RBE depends on the dose level and the number of dose fractions (or, alternatively, the dose per fraction) because in general, the shape of the dose–response relationship varies for radiations that differ substantially in their LET.

RBE can vary with the dose rate because the slope of the dose–response curve for sparsely ionizing radiations, such as x- or γ-rays, varies critically with a changing dose rate. In contrast, the biologic response to densely ionizing radiations depends little on the rate at which the radiation is delivered.

The biologic system or end point that is chosen has a marked influence on the RBE values obtained. In general, RBE values are high for tissues that accumulate and repair a great deal of sublethal damage and low for those that do not.

■ THE OXYGEN EFFECT AND LINEAR ENERGY TRANSFER

An important relationship exists between LET and the oxygen enhancement ratio (OER). Figure 7.7 shows mammalian cell survival curves for various types of radiation that have very different LETs and that exhibit very different OERs. Figure 7.7A refers to x-rays, which are sparsely ionizing, have a low LET, and consequently exhibit a large OER of about 2.5. Figure 7.7B refers to neutrons, which are intermediate in ionizing density and characteristically show an OER of 1.6. Figure 7.7D refers to 2.5-MeV α-particles, which are densely ionizing and have a high LET; in this case, survival estimates, whether in the presence or absence of oxygen, fall along a common line, and so the OER is unity. Figure 7.7C contains data for 4-MeV α-particles, which are slightly less densely ionizing; in this case, the OER is about 1.3.

Barendsen and his colleagues have used mammalian cells cultured in vitro to investigate the OER for a wide range of radiation types. Their results are summarized in Figure 7.8, in which OER is plotted as a function of LET. At low LET, corresponding to x- or γ-rays, the OER is between 2.5 and 3; as the LET increases, the OER falls slowly at first, until the LET exceeds about 60 keV/μm, after which the OER falls rapidly and reaches unity by the time the LET has reached about 200 keV/μm.

Both OER and RBE are plotted as a function of LET in Figure 7.9. (The curves are taken from the more complete plots in Figures 7.5 and 7.8.) Interestingly, the two curves are virtually mirror images of each other. The optimal RBE and the rapid fall of OER occur at about the same LET value (i.e., 100 keV/μm).

■ RADIATION WEIGHTING FACTOR (Wᵢ)

Radiations differ in their biologic effectiveness per unit of absorbed dose, as discussed previously. The complexities of RBE are too difficult
to apply in specifying dose limits in everyday radiation protection; it is necessary to have a simpler way to consider differences in biologic effectiveness of different radiations. The term radiation weighting factor \( (W_R) \) has been introduced for this purpose by International Commission on Radiological Protection (ICRP). The quantity produced by multiplying the absorbed dose by the weighting factor is called the equivalent dose. The unit of absorbed dose is the gray, and the unit of equivalent dose is the sievert (Sv). (In the old system of units, the unit of absorbed dose was the rad and the unit of equivalent dose was the rem.)

Radiation weighting factors are chosen by the ICRP based on a consideration of experimental RBE values, biased for biologic end points relevant to radiation protection, such as cancer and heritable effects, and also relevant to low doses and low dose rate. There is a considerable element of judgment involved. The radiation weighting factor is set at unity for all low-LET radiations (x-rays, γ-rays, and electrons), with a value of 20 for maximally effective neutrons and α-particles. Detailed values recommended by the ICRP are discussed in Chapter 17. Using this system, an absorbed dose of 0.1 Gy of α-particles with a radiation weighting factor of 20 would result in an equivalent dose of 2 Sv.

**SUMMARY OF PERTINENT CONCLUSIONS**

- X- and γ-rays are said to be sparsely ionizing because along the tracks of the electrons set in motion, primary ionizing events are well separated in space.
- α-particles and neutrons are densely ionizing because the tracks consist of dense columns of ionization.
- LET is the energy transferred per unit length of track. Typical values are 0.2 keV/μm for cobalt-60 γ-rays, 2 keV/μm for 250-kV x-rays, 166 keV/μm for 2.5-MeV α-particles, and 1,000 keV/μm for heavy charged particles encountered in space.
- RBE of some test radiation \( r \) is the ratio \( D_{250}/D_r \), in which \( D_{250} \) and \( D_r \) are the doses of 250-kV x-rays and the test radiation, respectively, required to produce equal biologic effect.
- RBE increases with LET to a maximum at about 100 keV/μm, thereafter decreasing with higher LET.
- For radiation with the optimal LET of 100 keV/μm, the average separation between ionizing events is similar to the
diameter of the DNA double helix (2 nm), so that DSBs can be most efficiently produced by a single track.

- The RBE of high-LET radiations compared with that of low-LET radiations increases as the dose per fraction decreases. This is a direct consequence of the fact that the dose-response curve for low-LET radiations has a broader shoulder than for high-LET radiations.
- RBE varies according to the tissue or end point studied. In general, RBE values are high for cells or tissues that accumulate and repair a great deal of sublethal damage, so that dose-response curves for x-rays have a broad initial shoulder.
- RBE depends on the following:
  - Radiation quality (LET)
  - Radiation dose
  - Number of dose fractions
  - Dose rate
  - Biologic system or end point

- The OER has a value of about 3 for low-LET radiations, falls when the LET rises more than about 30 keV/μm and reaches unity by an LET of about 200 keV/μm.
- The radiation weighting factor (WQ) depends on LET and is specified by the ICRP as a representative RBE at low dose and low dose rate for biologic effects relevant to radiation protection, such as cancer induction and heritable effects. It is used in radiologic protection to reduce radiations of different biologic effectiveness to a common scale.
- Equivalent dose is the product of absorbed dose and the radiation weighting factor. The unit of equivalent dose is the sievert (Sv) (in the old units, absorbed dose was expressed in rads and equivalent dose was expressed in rems).

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Acute Radiation Syndrome

Early Lethal Effects
The Prodromal Radiation Syndrome
The Cerebrovascular Syndrome
The Gastrointestinal Syndrome
The Hematopoietic Syndrome
The First and Most Recent Acute Radiation Syndrome Death
Mean Lethal Dose and Bone Marrow Transplants
Cutaneous Radiation Injury

Symptoms Associated with the Acute Radiation Syndrome
Treatment of Radiation Accident Victims Exposed to Doses Close to the LD50/60
Triage
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ACUTE RADIATION SYNDROME

The effect of ionizing radiation on the whole organism is discussed in this chapter. Data on the various forms of the acute radiation syndrome (ARS) have been drawn from many sources. Animal experiments provide the bulk of the data and result in a significant understanding of the mechanisms of death after exposure to total body irradiation. At the human level, data have been drawn from experiences in radiation therapy and studies of the Japanese survivors of Hiroshima and Nagasaki, the Marshallese accidentally exposed to fallout in 1954, and the victims of the limited number of accidents at nuclear installations, including Chernobyl. From these various sources, the pattern of events that follows a total body exposure to a dose of ionizing radiation has been well documented. To date, worldwide, about 400 humans have died of the ARS.

EARLY LETHAL EFFECTS

Early radiation lethality generally is considered to be death occurring within a few weeks that can be attributed to a specific high-intensity exposure to radiation. Soon after irradiation, early symptoms appear, which last for a limited period; this is referred to as the prodromal radiation syndrome. These symptoms may clear up after a few days, to be followed by a latent period before the development of the eventual life-threatening syndrome. This is illustrated in Figure 8.1. The eventual survival time and mode of death depend on the magnitude of the dose. In most mammals, three distinct modes of death can be identified, although in the circumstances of an actual accidental exposure, some overlap is frequently seen. At very high doses, in excess of about 100 Gy, death occurs 24 to 48 hours after exposure and appears to result from neurologic and cardiovascular breakdown; this mode of death is known as the cerebrovascular syndrome. At intermediate dose levels, approximately 5 to 12 Gy, death occurs in about 9 or 10 days and is associated with extensive bloody diarrhea and destruction of the gastrointestinal mucosa; this mode of death is known as the gastrointestinal syndrome. At lower dose levels, approximately 2.5 to 5 Gy, death occurs several weeks to 2 months after exposure and is caused by effects on the blood-forming organs; this mode of death has come to be known as bone marrow death or the hematopoietic syndrome.

The exact cause of death in the cerebrovascular syndrome is by no means clear. In the case of both of the other modes of death—the gastrointestinal and the hematopoietic syndromes—the principal mechanisms that lead to the death of the organism are understood. Death is caused by the depletion of the stem cells of a critical self-renewal tissue: the epithelium of the gut or the circulating blood cells, respectively. The difference in the dose level at which these two forms of death occur and the difference in the time scales involved reflect variations in the
The various symptoms making up the human prodromal syndrome vary with respect to time of onset, maximum severity, and duration, depending on the size of the dose. With doses of a few tens of gray, all exposed individuals can be expected to show all phases of the syndrome within 5 to 15 minutes of exposure. Reaction might reach a maximum by about 30 minutes and persist for a few days, gradually diminishing in intensity until the prodromal symptoms merge with the universally fatal cerebrovascular syndrome or, after a lower dose, with the fatal gastrointestinal syndrome.

At lower doses, dose-response predictions are difficult to make because of the interplay of many different factors. A severe prodromal response usually indicates a poor clinical prognosis and portends at the least a prolonged period of acute hematologic aplasia accompanied by potentially fatal infection: anemia and hemorrhage.

The signs and symptoms of the human prodromal syndrome can be divided into two main groups: gastrointestinal and neuromuscular. The gastrointestinal symptoms are anorexia, nausea, vomiting, diarrhea, intestinal cramps, salivation, fluid loss, dehydration, and weight loss. The neuromuscular symptoms include easy fatigability, apathy or listlessness, sweating, fever, headache, and hypotension. At doses that would be fatal to 50% of the population, the principal symptoms of the prodromal reaction are anorexia, nausea, vomiting, and easy fatigability. Immediate diarrhea, fever, and hypotension frequently are associated with supralethal exposure (Table 8.1). One of the Soviet firefighters at the Chernobyl reactor accident vividly described the onset of these symptoms as he accumulated a dose of several gray working in a high-dose-rate area. The prodromal phase is followed by a latent stage before the final radiation syndrome develops. In the symptom-free latent stage, the patient may seem and feel relatively well for a period of hours or even weeks. The duration of the latent stage is inversely proportional to the dose and may last a few hours for high exposures or as long as two or more weeks for lower exposures. Absence of a latent phase—that is, a progressive worsening from prodromal signs and symptoms directly into the manifest illness phase—is an indicator that the dose was probably very high.

The diagnosis of the ARS can also be based on laboratory data. During the prodromal phase, evidence of hematopoietic damage can already be observed by a drop in the lymphocyte count after an exposure as low as 0.5 Gy. The circulating lymphocytes are one of the most radiosensitive cell lines, and a fall in the absolute lymphocyte count is the best and most useful laboratory test to determine the level of radiation exposure in the early phase of observation. Among assays for biologic population kinetics of the two cell renewal systems involved and differences in the amount of the damage that can be tolerated in these different systems before death ensues.

**THE PRODROMAL RADIATION SYNDROME**

The various symptoms making up the human prodromal syndrome vary with respect to time of onset, maximum severity, and duration, depending on the size of the dose. With doses of a few tens of gray, all exposed individuals can be expected to show all phases of the syndrome within 5 to 15 minutes of exposure. Reaction might reach a maximum by about 30 minutes and persist for a few days, gradually diminishing in intensity until the prodromal symptoms merge with the universally fatal cerebrovascular syndrome or, after a lower dose, with the fatal gastrointestinal syndrome.

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**TABLE 8.1 Symptoms of the Prodromal Syndrome**

<table>
<thead>
<tr>
<th>Neuromuscular</th>
<th>Gastrointestinal</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Signs and Symptoms to be Expected at about 50% Lethal Dose</strong></td>
<td></td>
</tr>
<tr>
<td>Easy fatigability</td>
<td>Anorexia</td>
</tr>
<tr>
<td>—</td>
<td>Nausea</td>
</tr>
<tr>
<td>—</td>
<td>Vomiting</td>
</tr>
<tr>
<td><strong>Additional Signs to be Expected After Supralethal Doses</strong></td>
<td></td>
</tr>
<tr>
<td>Fever</td>
<td>Immediate diarrhea</td>
</tr>
<tr>
<td>Hypotension</td>
<td></td>
</tr>
</tbody>
</table>
disoriented, and his blood pressure could not be maintained; he died 49 hours after the accident.

In a nuclear criticality accident at Los Alamos in 1958, one worker received a total body dose of mixed neutron and \(\gamma\)-radiation estimated to be between 39 and 49 Gy. Parts of his body may have received as much as 120 Gy. This person went into a state of shock immediately and was unconscious within a few minutes. After 8 hours, no lymphocytes were found in the circulating blood, and there was virtually a complete urinary shutdown despite the administration of large amounts of fluids. The patient died 35 hours after the accident.

The exact and immediate cause of death in what is known as the cerebrovascular syndrome is not fully understood. Although death is usually attributed to events taking place within the central nervous system, much higher doses are required to produce death if the head alone is irradiated rather than the entire body; this would suggest that effects on the rest of the body are by no means negligible. It has been suggested that the immediate cause of death is damage to the microvasculature, which results in an increase in the fluid content of the brain owing to leakage from small vessels, resulting in a buildup of pressure within the bony confines of the skull.

### THE GASTROINTESTINAL SYNDROME

A total body dose of about 100 Gy of \(\gamma\)-rays or its equivalent of neutrons results in death in 24 to 48 hours. At these doses, all organ systems are also seriously damaged; both the gastrointestinal and hematopoietic systems are, of course, severely damaged and would fail if the person lived long enough, but cerebrovascular damage brings death very quickly, so that the consequences of the failure of the other systems do not have time to be expressed (i.e., death occurs before other symptoms have time to appear). The symptoms that are observed vary with the species of animal involved and also with level of radiation dose; they are summarized briefly as follows: There is the development of severe nausea and vomiting, usually within a matter of minutes. This is followed by manifestations of disorientation, loss of coordination of muscular movement, respiratory distress, diarrhea, convulsive seizures, coma, and finally death. Only a few instances of accidental human exposure have involved doses high enough to produce a cerebrovascular syndrome; two such cases are described briefly.

In 1964, a 38-year-old man working in a uranium-235 recovery plant was involved in an accidental nuclear excursion. He received a total body dose estimated to be about 88 Gy made up of 22 Gy of neutrons and 66 Gy of \(\gamma\)-rays. He recalled seeing a flash and was hurled backward and stunned; he did not lose consciousness, however, and was able to run from the scene of the accident to another building 200 yards away. Almost at once he complained of abdominal cramps and headache, vomited, and was incontinent of bloody diarrheal stools. The next day, the patient was comfortable but restless. On the second day, his condition deteriorated; he was restless, fatigued, apprehensive, and short of breath and had greatly impaired vision; his blood pressure could only be maintained with great difficulty. Six hours before his death, he became disoriented, and his blood pressure could not be maintained; he died 49 hours after the accident.

In a nuclear criticality accident at Los Alamos in 1958, one worker received a total body dose of mixed neutron and \(\gamma\)-radiation estimated to be between 39 and 49 Gy. Parts of his body may have received as much as 120 Gy. This person went into a state of shock immediately and was unconscious within a few minutes. After 8 hours, no lymphocytes were found in the circulating blood, and there was virtually a complete urinary shutdown despite the administration of large amounts of fluids. The patient died 35 hours after the accident.

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### THE GASTROINTESTINAL SYNDROME

A total body exposure of more than 10 Gy of \(\gamma\)-rays or its equivalent of neutrons commonly leads in most mammals to symptoms characteristic of the gastrointestinal syndrome, culminating in death some days later (usually between 3 and 10 days). The characteristic symptoms are nausea, vomiting, and prolonged diarrhea. People with the gastrointestinal syndrome lose their appetite and appear sluggish and lethargic. Prolonged diarrhea, extending for several days, usually is regarded as a bad sign because it indicates that the dose received was more than 10 Gy, which is inevitably fatal. The person with this syndrome shows signs of dehydration, loss of weight, emaciation, and complete exhaustion; death usually occurs in a few days. There is no instance on record of a human having survived a dose in excess of 10 Gy.

The symptoms that appear and the death that follows are attributable principally to the
Depopulation of the epithelial lining of the gastrointestinal tract by the radiation. The normal lining of the intestine is a classic example of a self-renewing tissue; Figure 8.2 shows the general characteristics of such a tissue. It is composed of a stem cell compartment, a differentiating compartment, and mature functioning cells.

The structure of the intestinal epithelium is illustrated in Figure 8.3. Dividing cells are confined to the crypts, which provide a continuous supply of new cells; these cells move up the villi, differentiate, and become the functioning cells. The cells at the top of the folds of villi are sloughed off slowly but continuously in the normal course of events, and the villi are continuously replaced by cells that originate from mitoses in the crypts. A single-cell-thick barrier separates the blood vessels in the villi from the contents of the intestine.

A dose of radiation of about 10 Gy sterilizes a large proportion of the dividing cells in the crypts; a dose of this order of magnitude does not seriously affect the differentiated and functioning cells. As the surface of the villi is sloughed off and rubbed away by normal use, there are no replacement cells produced in the crypt. Consequently, after a few days, the villi begin to shorten and shrink, and eventually, the surface lining of the intestine is completely denuded of villi.

The rate of cell loss and shrinkage depends on the dose. It occurs faster at higher doses than at lower doses. At death, the villi are very clearly flat and almost completely free from cells.

Before Chernobyl, there was probably only one example in the literature of a human suffering a gastrointestinal death as a result of radiation exposure. In 1946, a 32-year-old man was admitted to the hospital within 1 hour of a radiation accident in which he received a total body dose of neutrons and γ-rays. The dosimetry is very uncertain in this early accident, and various estimates of total body exposure range from 40 to 1700 rads (400 to 17,000 rads).

The precise time schedule of these events and the time required before the intestine is denuded of cells entirely vary with the species. In small rodents, this condition is reached between 3 and 4 days after the dose of radiation is delivered. In humans, it does not occur until about 9 to 10 days after irradiation. All of the individuals who received a dose large enough for the gastrointestinal syndrome to result in death have already received far more than enough radiation to result in hematopoietic death. Death from a denuding of the gut occurs, however, before the full effect of the radiation on the blood-forming organs has been expressed because of differences in the population kinetics of the stem cell systems involved.

Before Chernobyl, there was probably only one example in the literature of a human suffering a gastrointestinal death as a result of radiation exposure. In 1946, a 32-year-old man was admitted to the hospital within 1 hour of a radiation accident in which he received a total body dose of neutrons and γ-rays. The dosimetry is very uncertain in this early accident, and various estimates of total body exposure range from 40 to 1700 rads (400 to 17,000 rads).
The gastrointestinal epithelium is an example of a classic self-renewal tissue. Stem cells in the crypts divide rapidly and provide cells that differentiate to form the lining of the villi. A single cell layer separates the blood supply within the villus from the contents of the gastrointestinal (GI) tract. An exposure to radiation kills cells in the crypts, cutting off a supply of cells to cover the villi. As a consequence, the villi shrink and, eventually, the barrier between blood supply and the contents of the GI tract is compromised, leading to a loss of fluids and massive infections. (Courtesy of Dr. Jack Little.)

At doses of 2.5 to 5 Gy, death, if it occurs, is a result of radiation damage to the hematopoietic system. Mitotically active precursor cells are sterilized by the radiation, and the subsequent supply of mature red blood cells, white blood cells, and platelets is thereby diminished. The time of potential crisis at which circulating cells in the blood reaches a minimum value is delayed for some weeks. It is only when the mature circulating cells begin to die off and the supply of new cells from the depleted precursor population is inadequate to replace them that the full effect of the radiation becomes apparent.

The concept of the 50% lethal dose (LD$_{50}$) as an end point for scoring radiation death from
The chief symptoms of which are nausea and vomiting. A symptom-free interval, known as the latent period, follows. This is, in fact, a very inappropriate name because during this period, the most important consequences of the radiation exposure, leading to its lethal effects, are in progress. About 3 weeks after the radiation exposure, there is onset of chills, fatigue, petechial hemorrhages in the skin, and ulceration of the mouth; epilation (hair loss) also occurs at this time. These symptoms are a manifestation of the depression of blood elements: infections and fever from granulocyte depression and impairment of immune mechanisms, bleeding, and possibly anemia caused by hemorrhage resulting from platelet depression. Anemia from red blood cell depression usually does not occur. Death occurs at this stage unless the bone marrow has begun to regenerate in time. Infection is an important cause of death, but it may be controlled to a large extent by antibiotic therapy.

As a consequence of the reactor accident at Chernobyl, 203 operating personnel, firemen, and emergency workers suffering from the early radiation syndrome were hospitalized, having received doses in excess of 1 Gy. Of these, 35 had severe bone marrow failure, and 13 of them died. The remainder recovered with conservative medical care.

■ THE FIRST AND MOST RECENT ACUTE RADIATION SYNDROME DEATH

The most recent person to die of the ARS was Alexander Litvinenko, a former officer of the Russian Security Service who received political asylum in Great Britain and was assassinated by the administration of polonium-210 (Fig. 8.5). This radionuclide emits only α-particles that do not penetrate even a sheet of paper or the epidermis of human skin, so α-emitters can cause significant damage only if ingested. Litvinenko fell ill and was hospitalized on November 1, 2006, and died on November 23, just more than 3 weeks later. Scotland Yard initially investigated claims that he had been poisoned with thallium because the distinctive effects include hair loss and damage to peripheral nerves. Polonium-210 was identified only after his death. The administered activity was estimated to be about 2 GBq, which corresponds to about 10 mcg of polonium and is many times the mean lethal dose.
Studies of total body irradiation have been performed on many species; a few LD_{50} values are listed in Table 8.2, ranging from mouse to human. Such studies were popular and important in the 1950s and 1960s, supported largely by the military. In more recent years, total body irradiation has been of interest from the point of view of bone marrow transplantation. This interest may stem from the treatment of radiation accidents, such as the Chernobyl disaster, or from the rescue of patients receiving cancer therapy with total body irradiation, radiolabeled antibodies, or cytotoxic drugs.

Many attempts have been made to estimate the LD_{50/60} for humans based on the experiences at Hiroshima and Nagasaki, the total body irradiation of patients with malignant disease, and the accidents that have occurred at nuclear installations. In a careful summary of all of the available data, Lushbaugh claims that the best estimate is around 3.25 Gy for young healthy adults without medical intervention. There does exist in the literature a surprising number of instances in which young men and women have received total body irradiation up to a dose of around 4 Gy and recovered under conservative care in a modern well-equipped hospital. The LD_{50/60} for

This massive amount of polonium-210 could only be produced in a large state-controlled nuclear reactor. In retrospect, the symptoms were characteristic of the classic hematopoietic syndrome, hair loss, erythema, and death in about 3 weeks caused by loss of circulating blood elements.

The first person to die of the ARS was a 26-year-old male involved in a criticality accident at Los Alamos in March 1945. He was exposed total body to a mixture of neutrons and γ-rays, the estimated equivalent dose being 6.35 Sv. His right hand received a much higher dose of 200 Gy, and his left hand received a dose of 30 Gy. His red blood count changed little up to the time of his death. The platelet count dropped before being restored by a transfusion and then fell again. There was the characteristic early initial rise in the granulocyte count, but it fell to eventually zero by the time of his death. The most important events can be listed as follows:

Day 1: Nausea, anorexia, and vomiting
Day 2: Greatly improved, except for numbness in his hand
Day 3: Erythema on the front of the body
Day 5: Rise of temperature
Day 10: Nausea and cramps
Day 12: Acute mucositis of mouth and tongue
Day 17: Epilation of body hair
Day 24: Died with white cell count close to zero

Alexander Litvinenko, before and after radiation poisoning. Alexander Litvinenko was a former officer of the Russian Security Service who received political asylum in Great Britain. He was assassinated by the administration of a massive dose of polonium-210, presumably added to his food. He died 3 weeks later.
the threshold local dose for epilation is approximately 3 Sv and that for erythema is about 6 Sv. With increasing dose of more than 10 Sv, the injury worsens progressively, involving dry desquamation, wet desquamation, bullae (blisters) formation, ulceration, and finally necrosis. These may be debilitatingly painful like second-degree thermal burns and life threatening with concomitant infections.

**Symptoms Associated with the Acute Radiation Syndrome**

The International Atomic Energy Agency and the World Health Organization jointly sponsored a report entitled *Diagnosis and Treatment of Radiation Injuries*. Tables 8.3 and 8.4 have been adapted from that report, and the expected distribution of symptoms following whole body irradiation are summarized. Table 8.3 refers to the prodromal syndrome in the period soon after irradiation, whereas Table 8.4 refers to the later critical phase. These should not be taken too literally because the information is based on a limited number of exposed individuals over the years, but they are a useful guide. They cover the dose range from 1.0 Gy, which results in little effect, to more than 8 Gy, which is expected to result in 100% lethality. The nature of the symptoms, their severity, and the time of onset can be a useful predictor of the eventual outcome in the absence of physical dosimetry. For example, those exposed at Chernobyl was closer to 7 Gy because, although general medical care was poor, antibiotics were available.

**Cutaneous Radiation Injury**

The hematopoietic and gastrointestinal tract syndromes may be accompanied by *cutaneous radiation injury* (CRI). Radiation injury to the skin can also occur in the absence of the ARS, because nonpenetrating β-particles and low-energy photons may deposit excess dose only to the integument. Such an injury may become apparent within hours or may not be seen for weeks, depending on the dose. Findings and complaints can range from itching and tingling to epilation, erythema, edema, progressing to dry desquamation, wet desquamation, ulceration, and necrosis as the dose is increased. Chronic, possibly severe, skin infections and recurrent ulceration may complicate the process. The first person ever to die of the ARS in 1945, died a classic hematopoietic syndrome, having received a total body dose of about 6 Sv, but his hands received a much higher dose. Radiation damage to the skin and microvasculature of his hands caused great suffering before he died on Day 28 after exposure because of bone marrow failure.

**Localized radiation burns** to the skin differ from thermal and chemical burns primarily in the delay between exposure and effect, and in their tendency to undergo recurrent breakdown, even after a scar has formed. The threshold local dose for epilation is approximately 3 Sv and that for erythema is about 6 Sv. With increasing dose of more than 10 Sv, the injury worsens progressively, involving dry desquamation, wet desquamation, bullae (blisters) formation, ulceration, and finally necrosis. These may be debilitatingly painful like second-degree thermal burns and life threatening with concomitant infections.

**Table 8.2**

The 50% Lethal Doses for Various Species from Mouse to Human and the Relation between Body Weight and the Number of Cells that Need to Be Transplanted for a Bone Marrow “Rescue”

<table>
<thead>
<tr>
<th>Species</th>
<th>Average Body Weight, kg</th>
<th>50% Lethal Total Body Irradiation, Gy</th>
<th>Rescue Dose per kg $\times 10^{-3}$</th>
<th>Relative Hematopoietic Stem Cell Concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mouse</td>
<td>0.025</td>
<td>7</td>
<td>2</td>
<td>10</td>
</tr>
<tr>
<td>Rat</td>
<td>0.2</td>
<td>6.75</td>
<td>3</td>
<td>6.7</td>
</tr>
<tr>
<td>Rhesus monkey</td>
<td>2.8</td>
<td>5.25</td>
<td>7.5</td>
<td>7.3</td>
</tr>
<tr>
<td>Dog</td>
<td>12</td>
<td>3.7</td>
<td>17.5</td>
<td>1.1</td>
</tr>
<tr>
<td>Human</td>
<td>70</td>
<td>4</td>
<td>20</td>
<td>1</td>
</tr>
</tbody>
</table>

If the dose is known to have exceeded about 5 Gy, then death from the hematopoietic syndrome 3 to 4 weeks later is a real possibility. In some countries, isolation and barrier nursing—that is, isolation of patients from others so that they do not come in contact with possible infections while their blood count is low—is recommended. It has been shown in animals that the LD50 can be raised by a factor of about 2 by the use of antibiotics, and there is no reason to suppose that the same is not true in humans. Indeed, this is supported by the Chernobyl experience where the LD50 was closer to 7 Gy than 4 Gy. The important things to avoid are infection, bleeding, and physical trauma during the period in which the circulating blood elements reach a nadir, and to give the bone marrow a chance to regenerate.

The area of most discussion and disagreement is the use of bone marrow transplantation. This technique was used on four Yugoslav scientists who were exposed accidentally in the 1950s to doses initially estimated to be about

### TREATMENT OF RADIATION ACCIDENT VICTIMS EXPOSED TO DOSES CLOSE TO THE LD50/60

If the radiation exposure is known to be less than 4 to 5 Gy, most experts recommend that the patient be watched carefully but only treated in response to specific symptoms, such as antibiotics for an infection, fresh platelets for local hemorrhage, and so on. Petechial hemorrhages in skin were commonly observed in the Japanese irradiated in 1945 but are not reported so commonly among individuals exposed accidentally in nuclear power installations in the United States. Blood transfusions should not be given prophylactically because they delay the regeneration of the blood-forming organs.

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**TABLE 8.3 Latent Phase (Prodromal Syndrome) of Acute Radiation Syndrome**

<table>
<thead>
<tr>
<th>Degree of Acute Radiation Syndrome and Approximate Dose of Acute Whole Body Exposure (Gy)</th>
<th>Mild (1–2 Gy)</th>
<th>Moderate (2–4 Gy)</th>
<th>Severe (4–6 Gy)</th>
<th>Very Severe (6–8 Gy)</th>
<th>Lethal (&gt;8 Gy)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lymphocytes (G/L) (days 3–6)</td>
<td>0.8–1.5</td>
<td>0.5–0.8</td>
<td>0.3–0.5</td>
<td>0.1–0.3</td>
<td>0.0–0.1</td>
</tr>
<tr>
<td>Granulocytes (G/L)</td>
<td>&gt;2.0</td>
<td>1.5–2.0</td>
<td>1.0–1.5</td>
<td>( \leq 0.5 )</td>
<td>( \leq 0.1 )</td>
</tr>
<tr>
<td>Diarrhea</td>
<td>None</td>
<td>None</td>
<td>Rare</td>
<td>Appears on days 6–9</td>
<td>Appears on days 4–5</td>
</tr>
<tr>
<td>Epilation</td>
<td>None</td>
<td>Moderate, beginning on day 15 or later</td>
<td>Moderate or complete on days 11–21</td>
<td>Complete earlier than day 11</td>
<td>Complete earlier than day 10</td>
</tr>
<tr>
<td>Latency period (d)</td>
<td>21–35</td>
<td>18–28</td>
<td>8–18</td>
<td>7 or less</td>
<td>None</td>
</tr>
<tr>
<td>Medical response</td>
<td>Hospitalization not necessary</td>
<td>Hospitalization recommended</td>
<td>Hospitalization necessary</td>
<td>Hospitalization urgently necessary</td>
<td>Symptomatic treatment only</td>
</tr>
</tbody>
</table>

Adapted from *Diagnosis and Treatment of Radiation Injuries*, International Atomic Energy Agency, Vienna, Austria, 1998.
**TABLE 8.4** Critical Phase of Acute Radiation Syndrome

<table>
<thead>
<tr>
<th>Degree of ARS and Approximate Dose of Acute Whole Body Exposure (Gy)</th>
<th>Mild (1–2 Gy)</th>
<th>Moderate (2–4 Gy)</th>
<th>Severe (4–6 Gy)</th>
<th>Very Severe (6–8 Gy)</th>
<th>Lethal (&gt;8 Gy)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Onset of symptoms</td>
<td>&gt;30 days</td>
<td>18–28 days</td>
<td>8–18 days</td>
<td>&lt;7 days</td>
<td>&lt;3 days</td>
</tr>
<tr>
<td>Lymphocytes (G/L)</td>
<td>0.8–1.5</td>
<td>0.5–0.8</td>
<td>0.3–0.5</td>
<td>0.1–0.3</td>
<td>0–0.1</td>
</tr>
<tr>
<td>Platelets (G/L)</td>
<td>60–100 %</td>
<td>30–60 %</td>
<td>25–35 %</td>
<td>15–25 %</td>
<td>&lt;20 %</td>
</tr>
<tr>
<td>Clinical manifestations</td>
<td>Fatigue, weakness</td>
<td>Fever, infections, bleeding, weakness, epilation</td>
<td>High fever, infections, bleeding, epilation</td>
<td>High fever, diarrhea, vomiting, dizziness and disorientation, hypotension</td>
<td>High fever, diarrhea, unconsciousness</td>
</tr>
<tr>
<td>Lethality (%)</td>
<td>0 %</td>
<td>0–50 % Onset 6–8 weeks</td>
<td>20–70 % Onset 4–8 weeks</td>
<td>50–100 % Onset 1–2 weeks</td>
<td>100 % 1–2 weeks</td>
</tr>
<tr>
<td>Medical response</td>
<td>Prophylactic</td>
<td>Special prophylactic treatment from days 14–20; isolation from days 10–20</td>
<td>Special prophylactic treatment from days 7–10; isolation from the beginning</td>
<td>Special treatment from day 1; isolation from the beginning</td>
<td>Symptomatic only</td>
</tr>
</tbody>
</table>

*In very severe cases, with a dose > 50 Gy, death precedes cytopenia.

Adapted from *Diagnosis and Treatment of Radiation Injuries*, International Atomic Energy Agency, Vienna, Austria, 1998.
7 Gy. All of the grafts were rejected, but the exposed individuals survived anyway, probably because later estimates indicated that the dose received was much lower, in the region of 4 Gy. In fact, many observers claim that the scientists survived in spite of the transplants rather than because of them. Figure 8.6 shows the depression and recovery of blood elements in the Yugoslav scientists and also in victims of the famous Y-12 reactor accident at Oak Ridge, Tennessee, who received about 4 Gy.

In more recent years, bone marrow transplantation techniques have been greatly improved and, together with growth factors, have been used routinely to “rescue” patients given supralethal doses of radiation for the treatment of leukemia or in preparation for organ transplants. In such cases, of course, the dosimetry is accurate and the doses are just enough to suppress the immunologic response.

Of the Chernobyl accident victims, 13 received bone marrow transplants (some matched for immune compatibility and some are not). In addition, 6 received fetal liver transplants, but these patients all died early, some of gastrointestinal symptoms. Of the 13 who received bone marrow transplants, only 2 survived and 1 showed autologous bone marrow repopulation. There was, therefore, only one possible successful transplant that saved a life, and even that result has been questioned.

The situation was made difficult because the doses to which individuals had been exposed were not known with any precision. After doses close to the LD$_{50}$, and certainly for higher doses, peripheral lymphocytes disappear before 24 hours, and then it is not possible to estimate total body doses by counting chromosome aberrations in stimulated lymphocytes taken from peripheral blood. Because the US transplant team did not arrive in Chernobyl for some time, biologic dosimetry was never possible for those exposed to higher doses. Consequently, some victims who received bone marrow transplants were already doomed to die of the gastrointestinal syndrome, having received doses in excess of 10 Gy.

In fact, the “window” of dose within which a bone marrow transplant is useful is very small. For a dose of less than about 8 Gy, an exposed person is likely to survive with careful nursing and an antibiotic screen because the LD$_{50}$ can be approximately doubled by such conservative measures. In such cases, therefore, a transplant is not necessary. For a dose of more than about 10 Gy, death from the gastrointestinal syndrome is inevitable, and so a bone marrow transplant is of no use. This highlights the narrow “window” of dose within which a transplant can be effective (about 8–10 Gy). This is illustrated in Figure 8.7. Therefore, there is an urgent need to develop better methods of in vivo biologic dosimetry because chromosome aberrations in lymphocytes are not always useful in this dose range.

- **TRIAGE**

Following an event in which several individuals are exposed to radiation, an immediate need is to know what doses are involved. If the exposed individuals are radiation workers wearing monitors, the solution is easy. However, in general, members of the public will not be monitored. The following are several possibilities:

1. **The average time to emesis decreases with increasing dose.** The original work was performed by Ricks and Lushbaugh at Oak Ridge Associated Universities and involved 502 patients who had undergone therapeutic or accidental radiation exposure between 1964 and 1975. The data as analyzed by Goans is shown in Figure 8.8. Individual responses vary so widely that time to emesis can provide only rough guidance. For example, few individuals vomit if the acute radiation dose is less than 1 Gy, whereas most vomit if the dose is more than 2 Gy. Further, if no vomiting occurs during the first 4 hours after exposure, it is unlikely that severe clinical effects caused by radiation will follow later. On the other hand, Goans has reported that if the time to emesis is less than 2 hours after exposure, the effective whole body dose is at least 3 Gy.

2. **The decline in the lymphocyte count allows an estimate to be made of the total body radiation exposure.** Typical data for the decline of lymphocyte count with radiation dose are shown in Figure 8.9. Goans and colleagues developed an algorithm to estimate an approximate dose based on the depletion rate from serial blood counts performed at various times after exposure because, of course, the preirradiation lymphocyte count is usually unknown. Because the lymphocyte count falls more rapidly the higher the dose, the best estimate of dose and prognosis can be made at about 48 hours postexposure. Again, the estimate is only an approximation.
Comparison of platelet counts in the Y-12 patients and in 4 victims of the Vinča accident

Comparison of granulocyte counts in the Y-12 patients and in 4 victims of the Vinča accident

**FIGURE 8.7** Illustrating the narrow window of dose over which bone marrow transplants might be useful following total body irradiation. Up to about 8 Gy, most people would survive with antibiotics and careful nursing. Above about 10 Gy, most people would die because of the gastrointestinal syndrome.

**FIGURE 8.8** Dose can be estimated by the time of onset of vomiting. Early onset of vomiting indicates a high dose. However, there is a large variation between individuals. (Based on Anno GH, et al. Health Physics, 1999; 56(6):821–838, and Goans RE. Clinical care of the radiation accident patient: patient presentation, assessment, and initial diagnosis. In: Ricks RC, Berger ME, Ohara, FM Jr, eds. The Medical Basis for Radiation Accident Preparedness: The Clinical Care of Victim. Boca Raton, FL: The Parthenon Publishing; 2001.)

**FIGURE 8.9** The rate of decline and the degree of depleted lymphocytes are directly related to absorbed dose. (Based on the data from Guskova et al., 1988.)
3. If a cytogenetic laboratory is available, the best method to assess radiation exposure is to measure the incidence of chromosomal aberrations in peripheral lymphocytes stimulated to divide in vitro. Doses of more than 0.2 Gy can be accurately assessed in this way. This technique is described in more detail in Chapter 2. After high doses, of course, lymphocytes disappear quickly and so this technique has severe limitations.

**SURVIVORS OF SERIOUS RADIATION ACCIDENTS IN THE UNITED STATES**

Over the past 50 years, there have been several accidents in which small numbers of people employed in the nuclear program were exposed to total body or partial body irradiation. Most occurred in the early days of the nuclear program and involved criticality accidents. The number involved in the United States is about 70 workers in 13 separate accidents.

The long-term survivors have been studied exhaustively over the years. The medical history of these heavily irradiated people mirrors that of any aging population. The expected high incidences of shortened life span, early malignancies after a short latent period, and rapidly progressing lenticular opacities have not been observed. The numbers in any group are small, but the several malignancies, cataracts, and degenerative diseases that have been seen are no more than might be expected in a similar group of unirradiated people of the same age.

The survivors of the 1958 criticality accident at the Oak Ridge Y-12 plant are a case in point. Their blood cell counts are shown in Figure 8.6. A group of eight workers, ranging in age from 25 to 56 years, received total body doses of 0.23 to 3.65 Gy; five of them received doses of more than 2 Gy. Nevertheless, as of 1999, more than 40 years after the accident, none had died of a classic “radiogenic” cancer. There were two cases of lung cancer in very heavy smokers, a meningioma, and prostate cancer in a 70-year-old man. In fact, the only medical finding likely to be radiation related is bilateral posterior capsular cataracts in two of these patients. Three of the workers who received the biggest doses are retired and in good health.

This highlights the problem of detecting an excess cancer incidence in any small irradiated population. For example, if a group of workers receives a total body exposure of 3 Gy, the biggest dose possible without suffering early death from the hematopoietic syndrome, the excess cancer incidence would be expected to be about 24%. (The cancer risk estimates of the Committee on Biological Effects of Ionizing Radiation and the United Nations Scientific Committee on the Effects of Atomic Radiation based on the Japanese atomic bomb survivors amount to about 8% per sievert.) Thus, the biggest dose to which humans can be exposed and survive doubles the spontaneous cancer incidence. This is difficult to detect in a small group of people and is likely to be masked by other biologic factors. That is not to say that heavily irradiated individuals are not at increased risk, but an excess cancer incidence can be observed only by a careful study of a large population.

**RADIATION EMERGENCY ASSISTANCE CENTER**

In the context of radiation accidents, it should be noted that the Medical Sciences Division of the Oak Ridge Institute for Science and Education operates a Radiation Emergency Assistance Center/Training Site (REAC/TS). This is operated on behalf of the U.S. Department of Energy.

REAC/TS provides 24-hour direct or consultative assistance with medical and health physics problems associated with radiation accidents in local, national, and international incidents. The resources of REAC/TS consist of expertise in cytogenetics for dose assessment, calculation of doses from internally deposited radionuclides, and laboratory facilities that include total body counting capabilities. The regular telephone number for information is (865) 576-3131, and the 24-hour emergency number is (865) 576-1005 (ask for REAC/TS). The REAC/TS website is http://www.orau.gov/reacts.

**SUMMARY OF PERTINENT CONCLUSIONS**

- The prodromal syndrome varies in time of onset, severity, and duration.
- At doses close to the dose that would be lethal to 50% of the population (LD50), the principal symptoms of the prodromal syndrome are anorexia, nausea, vomiting, and easy fatigability.
- Immediate diarrhea, fever, or hypotension indicates a supralethal exposure.
The cerebrovascular syndrome results from a total body exposure to about 100 Gy of γ-rays and in humans results in death in 24 to 48 hours. The cause of death may be changes in permeability of small blood vessels in the brain following damage to the microvasculature.

The gastrointestinal syndrome results from a total body exposure to about 10 Gy. Death occurs in about 7 to 10 days in humans because of depopulation of the epithelial lining of the gastrointestinal tract.

The hematopoietic syndrome results from total body exposure to 2.5 to 5 Gy. The radiation sterilizes some or all of the mitotically active precursor cells. Symptoms result from lack of circulating blood elements 3 or more weeks later.

The hematopoietic or gastrointestinal syndromes may be complicated by damage to the skin from high local doses.

The LD₅₀ for humans is estimated to be between 3 and 4 Gy for young adults without medical intervention. It may be less for the very young or the very old. The LD₅₀ may be raised to 7 Gy by the use of antibiotics, as was the case at Chernobyl.

Some people who would otherwise die from the hematopoietic syndrome may be saved by antibiotics, platelet infusions, bone marrow transplants, or factors to stimulate hematopoiesis.

The dose window over which bone marrow transplants may be useful is narrow, namely, 8 to 10 Gy.

Heavily irradiated survivors of accidents in the nuclear industry have been followed for many years; their medical history mirrors that of any aging population. An expected higher incidence of shortened life span, early malignancies after a short latency, and rapidly progressing cataracts has not been observed. That is not to say that heavily irradiated individuals are not at increased risk, but an excess cancer incidence can be observed only by a careful study of a large population.

The first recorded death with the ARS was a worker at Los Alamos in 1945. The most recent was the Russian agent, Litvinenko, who was “poisoned” with polonium-210 in 2006. Worldwide, about 400 individuals have suffered from the ARS.

BIBLIOGRAPHY


THE DISCOVERY OF RADIOPROTECTORS

Some substances, although they do not directly affect the radiosensitivity of cells, nevertheless, may protect whole animals because they cause vasoconstriction or, in some way, upset normal processes of metabolism to such an extent that the oxygen concentration in critical organs is reduced. Because cells are less sensitive to x-rays under hypoxia, this confers a measure of protection. Examples of such protective substances are sodium cyanide, carbon monoxide, epinephrine, histamine, and serotonin. Such compounds are not really radioprotectors per se and are not discussed further here.

The most remarkable group of true radioprotectors is the sulfhydryl (SH) compounds. The simplest is cysteine, an SH compound containing a natural amino acid, the structure of which is

\[
\text{NH}_2 \quad \text{SH} \quad \text{CH}_2 \quad \text{CH} \quad \text{COOH}
\]

In 1948, Patt discovered that cysteine could protect mice from the effects of total body exposure to x-rays if the drug was injected or ingested in large amounts before the radiation exposure. At about the same time, Bacq and his colleagues in Europe independently discovered that cysteamine could also protect animals from total body irradiation. This compound has a structure represented by

\[
\text{SH} \quad \text{CH}_2 \quad \text{CH}_2 \quad \text{NH}_2
\]

Animals injected with cysteamine to concentrations of about 150 mg/kg require doses of x-rays 1.8 times larger than control animals to produce the same mortality rate. This factor of 1.8 is called the dose reduction factor (DRF), defined as

\[
\text{DRF} = \frac{\text{Dose of radiation in the presence of the drug}}{\text{Dose of radiation in the absence of the drug}}
\]

to produce a given level of lethality.

MECHANISM OF ACTION

Many similar SH compounds have been tested and found to be effective as radioprotectors. The most efficient SH compounds tend to have certain structural features in common: a free SH group (or potential SH group) at one end of the molecule and a strong basic function, such as amine or guanidine, at the other end, separated by a straight chain of two or three carbon atoms. SH compounds are efficient radioprotectors against sparsely ionizing radiations such as x- or γ-rays.

The mechanisms most implicated in SH-mediated cytoprotection include:

1. Free-radical scavenging that protects against oxygen-based free radical generation by ionizing radiations or chemotherapy agents such as alkylating agents
2. Hydrogen atom donation to facilitate direct chemical repair at sites of DNA damage

Chapter 1 includes a discussion of the chain of events between the absorption of a photon and the eventual biologic damage, which includes the
production of free radicals, which are highly reactive species. If these free radicals are scavenged before they can interact with biologic molecules, the effect of the radiation is reduced. This process is illustrated in Figure 9.1.

The protective effect of SH compounds tends to parallel the oxygen effect, being maximal for sparsely ionizing radiations (e.g., x- or γ-rays) and minimal for densely ionizing radiations (e.g., low-energy α-particles). It might be predicted that with effective scavenging of all free radicals, the largest possible value of DRF for sparsely ionizing radiations would equal the oxygen enhancement ratio, with a value of 2.5 to 3.0.

This simple description of the mechanism of action of SH radioprotectors is intellectually satisfying, but it is clearly not the whole story because radioprotectors of this class have more effect with densely ionizing radiations (such as neutrons) than would be expected based on this explanation alone. Other factors must be involved that are not fully understood.

## DEVELOPMENT OF MORE EFFECTIVE COMPOUNDS

The discovery in 1948 of a compound that offered protection against radiation excited the interest of the U.S. Army because the memory of Nagasaki and Hiroshima was vivid in the years immediately after World War II. However, although cysteine is a radioprotector, it is also toxic and induces nausea and vomiting at the dose levels required for radioprotection. A development program was initiated in 1959 by the U.S. Army in studies conducted at the Walter Reed Institute of Research to identify and synthesize drugs capable of conferring protection to individuals in a radiation environment, but without the debilitating toxicity of cysteine or cysteamine. More than 4,000 compounds were synthesized and tested. At an early stage, the important discovery was made that the toxicity of the compound could be greatly reduced if the SH group was covered by a phosphate group. This is illustrated for cysteamine, otherwise known as mercaptopethyamine (MEA), in Table 9.1. The 50% lethal

### Table 9.1 Effect of Adding a Phosphate-Covering Function on the Free Sulphydryl of β-Mercaptoethylamine (MEA)

<table>
<thead>
<tr>
<th>Drug</th>
<th>Formula</th>
<th>Mean 50% Lethal Dose (Range) in Mice</th>
<th>Dose Reduction Factor</th>
</tr>
</thead>
<tbody>
<tr>
<td>MEA</td>
<td>NH₂—CH—CH₂—SH</td>
<td>343 (323–364)</td>
<td>1.6 at 200 mg/kg</td>
</tr>
<tr>
<td>MEA-PO₃</td>
<td>NH₂—CH₂—CH—SH₂PO₃</td>
<td>777 (700–864)</td>
<td>2.1 at 500 mg/kg</td>
</tr>
</tbody>
</table>
Chapter 9 • Radioprotectors

The astronauts would have been exposed to a shower of high-energy protons, resulting in an estimated total body dose of several grays. The availability of a radioprotector with a $\text{DRF}$ of between 2 and 3 would have been very important in such a circumstance. As it turned out, no major solar event occurred during any manned lunar mission, thus the protectors were not used. The potential for this problem will be greatly magnified in future missions to Mars, which may take as long as 3 years.

**AMIFOSTINE (WR-2721) AS A RADIOPROTECTOR IN RADIOTHERAPY**

The only radioprotective drug approved by the U.S. Food and Drug Administration (FDA) for use in radiation therapy is amifostine (WR-2721), sold under the trade name Ethyol for use in the prevention of xerostomia in patients treated for head and neck cancer. The Radiotherapy Oncology Group (RTOG) conducted a phase III randomized clinical trial, which demonstrated the efficacy of amifostine in reducing xerostomia in patients with head and neck cancer receiving radiotherapy without prejudice to early tumor control. The drug was administered daily, 30 minutes before each dose fraction in a multifraction regimen. Three months posttreatment, the incidence of xerostomia was significantly reduced.

### TABLE 9.2

**Two Radioprotectors in Practical Use**

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Use</th>
</tr>
</thead>
<tbody>
<tr>
<td>WR-638</td>
<td>$\text{NH}_2\text{CH}_2\text{CH}_2\text{SPO}_3\text{HNa}$</td>
<td>Carried in field pack by Russian army (cystaphos)</td>
</tr>
<tr>
<td>WR-2721</td>
<td>$\text{NH}_2(\text{CH}_2)_3\text{NHCH}_2\text{CH}_2\text{SPO}_3\text{H}_2$</td>
<td>Protector in radiotherapy and carried by US astronauts on lunar trips (amifostine)</td>
</tr>
</tbody>
</table>

### Comparison of Gastrointestinal and Hematopoietic Dose Reduction Factors in Mice for these Radioprotectors (the Two Compounds Listed Previously)

<table>
<thead>
<tr>
<th>Compound</th>
<th>Drug Dose, mg/kg</th>
<th>7 Days (Gastrointestinal)</th>
<th>30 Days (Hematopoietic)</th>
</tr>
</thead>
<tbody>
<tr>
<td>WR-638</td>
<td>500</td>
<td>1.6</td>
<td>2.1</td>
</tr>
<tr>
<td>WR-2721</td>
<td>900</td>
<td>1.8</td>
<td>2.7</td>
</tr>
</tbody>
</table>

The structures of two typical compounds of more than 4,000 synthesized in the Walter Reed series are shown in Table 9.2. The first compound, WR-638, called cystaphos, was said to be carried routinely in the field pack of Soviet infantry in Europe during the Cold War for use in the event of a nuclear conflict.

The second compound, WR-2721, now known as amifostine, is perhaps the most effective of those synthesized in the Walter Reed series. It gives good protection to the blood-forming organs, as can be seen by the $\text{DRF}$ for 30-day death in mice, which approaches the theoretic maximum value of 3. It was probably the compound carried by US astronauts on their trips to the moon, to be used if a solar event occurred. On these missions, when the space vehicle left Earth’s orbit and began coasting toward the moon, the astronauts were committed to a 14-day mission because they did not have sufficient fuel to turn around without first orbiting the moon and using its gravitational field. If there had been a major solar event in that period, the astronauts would have been exposed to a shower of high-energy protons, resulting in an estimated total body dose of several grays. The availability of a radioprotector with a $\text{DRF}$ of between 2 and 3 would have been very important in such a circumstance. As it turned out, no major solar event occurred during any manned lunar mission, thus the protectors were not used. The potential for this problem will be greatly magnified in future missions to Mars, which may take as long as 3 years.
in those patients treated with amifostine. There was an improvement in the patients’ assessments of such symptoms as dry mouth and difficulty in eating or speaking and in the need for fluids and oral comfort aids. There was no difference in locoregional tumor control between patients who received the radioprotector and those who did not. Giving the amifostine only 30 minutes before each treatment was designed to exploit the slower rate at which the drug penetrates tumors relative to normal tissues.

Amifostine is a phosphorothioate that is non-reactive and does not readily permeate cells, primarily because of its terminal phosphorothioic acid group. It is therefore a “prodrug.” When dephosphorylated by the enzyme alkaline phosphatase, which is present in high concentrations in normal tissues and capillaries, it is converted to the active metabolite designated WR-1065. This metabolite readily enters normal cells by facilitated diffusion and scavenges free radicals generated by ionizing radiations or by drugs used in chemotherapy such as alkylating agents.

It might have been expected that radioprotectors would enjoy a wider use in radiation therapy, but in practice, clinical use continues to be plagued by issues relating to possible tumor protection and loss of therapeutic gain. The potential use of such protectors is based on the observation from animal studies that amifostine quickly floods normal tissues but penetrates more slowly into tumors. Consequently, if the radiation dose is given within minutes after the administration of the radioprotector, there is a differential sparing of normal tissue compared with tumor cells. Because one can never be sure that the tumor is not protected to some extent, the use of radioprotectors is not “fail safe.” For this reason, radioprotectors are not widely used in radiotherapy, indeed, in practice they are used only for the reduction of xerostomia.

### RADIOPROTECTORS AND CHEMOTHERAPY

Although SH compounds were developed initially as radioprotectors against ionizing radiation, they also protect against the cytoxic effects of several chemotherapeutic agents. The experimental clinical use of amifostine has shown that the compound offers significant protection against nephrotoxicity, ototoxicity, and neuropathy from cisplatin and hematologic toxicity from cyclophosphamide. The same experimental studies indicated no obvious antitumor activity of the radioprotector, implying a differential uptake between normal and malignant tissues.

#### AMIFOSTINE AS A PROTECTOR AGAINST CANCER

Although the emphasis for the development of amifostine was to protect against cell killing, this compound also protects against radiation-induced mutagenesis and oncogenic transformation in cells in culture and against carcinogenesis in mouse model systems. Furthermore, although a dose of about 400 mg/kg is required to demonstrate optimal cytoprotection—a dose that carries with it significant side effects—its anti-mutagenic effect persists following prolonged exposure to a dose as low as 25 mg/kg, which is nontoxic. Of even greater interest is the observation that the effect occurs even when cells are exposed to amifostine up to 3 hours following irradiation. This has led to the speculation that the antioxidant properties of amifostine may not be the only mechanism by which it protects against cancer; it has been proposed that the polyamine-like properties of the phosphorothioates may result in a stabilization of DNA-damaged sites, facilitating a slower and more error-free repair of damage.

### DIETARY SUPPLEMENTS AS COUNTERMEASURES TO RADIATION

Long-term exposure to nonlethal doses of ionizing radiation is known to result in an excess incidence of cancer and other deleterious biologic effects. To the extent that the mechanism involved may include oxidative stress, dietary supplements involving antioxidants have a potential role to play. Several possibilities have shown promise in cellular and animal systems. One such example is the soybean-derived serine protease inhibitor known as the Bowman-Birk inhibitor (BBI), which has long been proposed as a cancer chemopreventive agent. Another possibility is a cocktail of common antioxidants, including L-selenomethionine, ascorbic acid, N-acetyl cysteine, alpha-lipoic acid, vitamin E succinate, and coenzyme Q10.
Following the destruction of the World Trade Center on September 11, 2001, and the rise of a nuclear terrorism threat, there has been a revived interest in the development of novel, effective, and nontoxic radioprotectors for potential use in homeland defense as well as in medical applications. In addition, National Aeronautics and Space Administration (NASA) is interested in countermeasures to the exposure to protons and high-energy heavy ions that astronauts experience during long-term missions in space.

**Summary of Pertinent Conclusions**

- Radioprotectors are chemicals that reduce the biologic effects of radiation.
- The SH compounds, cysteine and cysteamine, were discovered early but are toxic. If the SH group is covered by a phosphate group, toxicity is reduced.
- The mechanism of action is the scavenging of free radicals and restitution of free-radical damage, although this is not the whole story.
- The DRF is the ratio of radiation doses required to produce the same biologic effect in the absence and presence of the radioprotector.
- The best available radioprotectors can attain DRF values of 2.5 to 3.0 for bone marrow death in mice irradiated with x-rays.
- DRF values close to the oxygen enhancement ratio are possible for γ-rays, but the effectiveness of radioprotectors decreases with increasing linear energy transfer.
- During the Cold War, it is said that Soviet infantry in Europe carried radioprotectors for use in a possible nuclear war. Radioprotectors were carried to the moon by US astronauts to be used in the event of a solar flare.
- More than 4,000 compounds were synthesized by the U.S. Army in studies conducted at the Walter Reed Institute of Research. Amifostine (WR-2721) appears to be the best for use in conjunction with radiotherapy.
- Amifostine, sold under the trade name Ethyl, is the only radioprotective drug approved by the FDA for use in the prevention of xerostomia in patients treated for head and neck cancer.
- An RTOG phase III trial demonstrated the efficacy of amifostine in reducing xerostomia in patients with head and neck cancer receiving radiation therapy without affecting locoregional control. The radioprotector was administered 30 minutes before radiation.
- Amifostine is a “prodrug” that is unreactive and that penetrates poorly into cells until it is dephosphorylated by the enzyme alkaline phosphatase to the active metabolite WR-1065.
- The rationale for the use of phosphorothioate radioprotectors is that they flood normal tissues rapidly after administration but penetrate tumors much more slowly. The strategy is to begin irradiation soon after administration of the drug to exploit a differential effect.
- The clinical use of radioprotectors in radiation therapy continues to be plagued by issues relating to possible tumor protection and diminution of therapeutic gain.
- Amifostine is useful as a protector for chemotherapy as well as radiotherapy. It is reported to offer protection against nephrotoxicity, ototoxicity, and neuropathy from cisplatin and hematologic toxicity from cyclophosphamide, without reduction of tumor activity.
- A dose of 400 mg/kg is required for optimal cytoprotection, which is toxic with many side effects, but its antimutagenic effect persists at a low nontoxic dose of 25 mg/kg. Furthermore, its antimutagenic effect still occurs if the drug is added 3 hours following irradiation.
- Dietary supplements, including various antioxidants, have been suggested as countermeasures to the long-term biologic effects of radiation exposure.
- Following the destruction of the World Trade Center on September 11, 2001, and the rise of a nuclear terrorism threat, there has been a revived interest in the development of novel, effective, and nontoxic radioprotectors for potential use in homeland defense. In addition, NASA is interested in countermeasures to the radiation exposure that astronauts experience on long-term space missions.


DETERMINISTIC AND
STOCHASTIC EFFECTS

If cellular damage occurs as a result of radiation and it is not adequately repaired, it may prevent the cell from surviving or reproducing or it may result in a viable cell that has been modified, that is, suffered a change or mutation that it retains as a legacy of the radiation exposure. The two outcomes have profoundly different implications for the person of whom the cell is a part.

Most organs or tissues of the body are unaffected by the loss of a few cells; but if the number of cells lost is sufficiently large, there is observable harm, reflecting the loss of tissue function. The probability of such harm is zero at small radiation doses, but above some level of dose, called the threshold dose, the probability increases rapidly with dose to 100%. Above the threshold, the severity of harm also increases with dose. Effects such as these are said to be deterministic. A deterministic effect has a threshold in dose, and the severity of the effect is dose related. Radiation-induced cataracts and late tissue fibrosis are examples of deterministic effects.

The outcome is very different if the irradiated cell is viable but modified. Carcinogenesis and heritable effects fall into this category. If somatic cells are exposed to radiation, the probability of cancer increases with dose, probably with no threshold, but the severity of the cancer is not dose related. A cancer induced by 1 Gy is no worse than one induced by 0.1 Gy, but of course, the probability of its induction is increased. This category of effect is called stochastic, a word that has been given a special meaning in radiation protection but in general, just means “random.” If the radiation damage occurs in germ cells, mutations may occur that could cause deleterious effects in future generations. Again, there is probably no threshold and the severity of heritable effects is not dose related, although the probability of it occurring is.

The belief that stochastic effects have no dose threshold is based on the molecular mechanisms involved. There is reason to believe that even a single x-ray photon could result in a base change leading to a mutation that could cause cancer or a heritable defect. For this reason, it is considered prudent and conservative to assume that no dose is too small to be effective, although this can never be proved.

The two types of effects are summarized as follows:

Deterministic effect: severity increases with dose; practical threshold; probability of occurrence increases with dose (e.g., cataract).

Stochastic effect: severity independent of dose; no threshold; probability of occurrence increases with dose (e.g., cancer).

<table>
<thead>
<tr>
<th></th>
<th>DETERMINISTIC AND STOCHASTIC EFFECTS</th>
</tr>
</thead>
<tbody>
<tr>
<td>Deterministic effect: severity increases with dose; practical threshold; probability of occurrence increases with dose (e.g., cataract).</td>
<td></td>
</tr>
<tr>
<td>Stochastic effect: severity independent of dose; no threshold; probability of occurrence increases with dose (e.g., cancer).</td>
<td></td>
</tr>
</tbody>
</table>
CANCER induction is the most important somatic effect of low-dose ionizing radiation. In sharp contrast to the case for the heritable effects of radiation (Chapter 11), risk estimates for leukemogenesis and carcinogenesis do not rely on animal data but can be based on experience in humans. There is a long history of a link between radiation exposure and an elevated incidence of cancer. Figure 10.1 is a beautiful photograph of Marie Curie and her daughter Irene, who are both thought to have died of leukemia as a result of the radiation exposure they received while conducting their experiments with radioactivity. Figure 10.2 is a photograph of the hand of a dentist in New York who held x-ray films in patients’ mouths for many years and who suffered malignant changes as a result. Quantitative data on cancer induction by radiation come from populations irradiated for medical purposes or exposed deliberately or inadvertently to nuclear weapons. Persons exposed therapeutically received comparatively high doses, and their susceptibility to the effects of radiation might have been influenced by the medical condition for which treatment was being given. Populations exposed to γ-rays and neutrons from nuclear weapons represent a wider cross section in terms of age and health and also include persons exposed to lower doses. In both cases, dose rates were high and exposure times brief.

There are a few groups of exposed persons to whom these generalizations do not apply. Examples include pitchblende and uranium miners who inhaled the radioactive gas radon and its progeny products over a prolonged period, patients injected with radium chloride or Thorotrast for medical purposes, and persons who ingested radionuclides while painting luminous dials on clocks and watches with paint containing radium. Hundreds of thousands of nuclear workers have been exposed occupationally, and useful cancer risk estimates have become available in recent years. Miners exposed to radon in the uranium mines are an excellent source of data on lung cancer.
The dosimetry in each instance is so uncertain that it is rarely possible to deduce any quantitative relationship between the dose of radiation involved and the tumor incidence.

More recent examples of the human experience with radiation-induced cancer and leukemia include the following:

1. The Japanese survivors of the atomic bomb attacks on Hiroshima and Nagasaki are the most important single group studied because of their large number, the care with which they have been followed, and the fact that people of all ages and both sexes received a wide range of doses. About 120,000 people have been followed carefully, of whom about 50,000 received doses in excess of 0.005 Sv. By 1998, there had been more than 17,000 cases of cancer, of which about 853 were considered to be caused by radiation. The weapons used on the two cities were very different. The one used on Nagasaki was of a type that would be expected to emit gamma rays with few neutrons and had been previously tested, so dosimetry is based partly on measurements. The weapon used at Hiroshima was of a type never tested before or since, so that dose estimates are based largely on computer simulations. The radiation from this weapon was a mixture of neutrons and $\gamma$-rays. The dosimetry relating to the atomic bombs has been revised several times over the years, leading to changes in the cancer risk estimates. The most recent estimates were published in the Biologic Effects of Ionizing Radiation (BEIR) VII report in 2006 and will be discussed later in this chapter.

2. In Britain, from 1935 through 1944, some 14,000 patients suffering from ankylosing spondylitis were given radiotherapy to various regions of their spine to relieve pain. A small risk of leukemia mortality has been reported in these patients. Although the spondylitic series provides one of the largest bodies of data on leukemia in humans after exposure to x- or $\gamma$-radiation, and the dosimetry is quite good, it is far from ideal because it lacks a proper control, consisting of patients with the same disease who did not receive x-ray therapy but whose treatment was otherwise the same. A possible contribution

The early human experience of radiation-induced cancer may be summarized as follows:

1. Skin cancer and leukemia were common in early x-ray workers, principally physicists and engineers who worked around accelerators before radiation safety standards were introduced.

2. Lung cancer was a frequent problem in pitchblende miners in Saxony, who dug out the ore from which radium was extracted. In the years following World War II, lung cancer also was noted in uranium miners in the central Colorado plateau. In both cases, the mines were poorly ventilated and there was a buildup of radon gas in the atmosphere of the mine; radon and its progeny were inhaled by the miners, depositing atoms of radioactive material in their lungs. The intense local $\alpha$-radiation was responsible for inducing lung tumors. Bone tumors were observed in the radium dial painters. The painters were mostly young women who worked in factories in which the luminous dials on clocks and watches were painted with a special paint preparation containing radium. The workers dipped their brushes into the radium paint and used their tongues to shape the brushes into sharp points to paint the small dials on watches. As a result, some radium was ingested, which, because it is in the same group in the periodic table as calcium, was deposited in the tips of the growing bones. The intense $\alpha$-radiation produced bone tumors. There is also history of bone tumors in people who, in the 1920s and 1930s, received injections of radium salts for the treatment of tuberculosis or ankylosing spondylitis.

3. An excess incidence of liver tumors was reported in patients in whom the contrast material Thorotrast was used. Thorotrast contains radioactive thorium, which, when deposited in the liver, produced a small incidence of liver tumors by $\alpha$-radiation.

These early examples are interesting but largely anecdotal, although they did alert scientists to the danger of excessive radiation exposure. None of these examples involved situations that now constitute a public health hazard; these problems will never happen again, and the dosimetry in each instance is so uncertain that it is rarely possible to deduce any quantitative relationship between the dose of radiation involved and the tumor incidence.
of carcinogenic drugs to the tumor incidence also has been suggested.

3. There is also documentation of an elevated incidence of leukemia in radiologists who joined learned societies before about 1922, before the introduction of radiation safety standards. This will be discussed later in the chapter.

4. Thyroid cancer has been observed in children who received radiotherapy for what was thought to be an enlarged thymus. The thyroid was included in the treatment field, and both malignant and benign thyroid tumors have been observed. Breast cancer is also elevated in these patients.

5. Until the 1950s, it was common practice to use x-rays to epilate children suffering from tinea capitis (ringworm of the scalp). An increased incidence of thyroid cancer from this practice was first reported by Modan and his colleagues in Israel, who treated more than 20,000 immigrant children from North Africa in whom ringworm of the scalp reached epidemic proportions. There was also a significantly increased risk of brain tumors (mostly meningiomas), salivary gland tumors, skin cancer, and leukemia mortality. A comparable group of children in New York for whom x-rays were used for epilation before treatment for tinea capitis show quite different results. There were only two malignant thyroid tumors in addition to some benign tumors. There is, however, an incidence of skin cancer around the face and scalp in those areas also subject to sunlight. The skin tumors arose only in white children and there were no tumors in black children in the New York series.

6. Patients with tuberculosis, who were fluoroscopyed many times during artificial pneumothorax, have shown an elevated incidence of breast cancer. This was first reported in Nova Scotia, but the report was confirmed by a similar study in New England. The doses these patients received are uncertain but must have been about 0.8 to 0.9 Gy, because some of the women developed skin changes in the chest wall on the side frequently fluoroscopyed. Patients who received radiotherapy for postpartum mastitis were also shown to have an excess incidence of breast cancer.

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**THE LATENT PERIOD**

The time interval between irradiation and the appearance of a malignancy is known as the **latent period**.

Leukemia has the shortest latent period. Excess cases began to appear in the survivors of Hiroshima and Nagasaki a few years after irradiation and reached a peak in 5 to 7 years; most cases occurred in the first 15 years. Solid tumors show a longer latency than the leukemias, on the order of anything from 10 to 60 years or more. For example, an excess incidence of solid tumors is still evident in Japanese survivors exposed to radiation from the atomic bombs in 1945. Indeed, for solid cancers, the excess risk is apparently more like a lifelong elevation of the natural age-specific cancer risk.

As the Japanese data have matured, the concept of a fixed time interval between irradiation and the appearance of the malignancy has been replaced by a combination of “age at exposure” and “time since exposure.” Regardless of the age at the time of exposure, radiation-induced solid tumors tend to be expressed later in life, at the same time as spontaneous tumors of the same type. Breast cancer in women is the most striking example. This suggests that although radiation may initiate the carcinogenic process at a young age, additional steps are required later in life, some of which may well be hormone dependent.

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**ASSESSING THE RISK**

To use the available human data to estimate risks as a function of dose, it is necessary to fit the data to a model. Several reasons for this are as follows:

1. Data obtained at relatively high doses must be extrapolated to the low doses of public health concern.

2. No large human population exposed to radiation has yet been studied for its full life span, and so estimates must be projected into the future. For example, in the year 2000, about half of the Japanese survivors irradiated in 1945 were still alive.

3. The best data pertain to the Japanese irradiated by the atomic bombs and risk estimates based on this must be transferred to other populations that have quite different characteristics, including their natural cancer incidence.
There are two types of models that are conceptually quite different: the absolute risk model and the relative risk model. The **absolute risk model** assumes that radiation induces a “crop” of cancers over and above the natural incidence unrelated to it. The **relative risk model** assumes that the effect of radiation is to increase the natural incidence at all ages subsequent to exposure by a given factor. Because the natural or spontaneous cancer incidence rises significantly in old age, the relative risk model predicts a large number of radiation-induced cancers in old age.

The model favored by recent BEIR committees, for the assessment of the cancer risks from the Japanese atomic bomb survivors is the **time-dependent relative risk model**. The excess incidence of cancer was assumed to be a function of dose, the square of the dose, age at exposure, and time since exposure. For some tumors, gender must be added as a variable—for example, in the case of breast cancer.

### COMMITTEES CONCERNED WITH RISK ESTIMATES AND RADIATION PROTECTION

There are two series of reports that analyze available data and come up with risk estimates for radiation-induced cancer. The first is the United Nations Scientific Committee on the Effects of Atomic Radiation (UNSCEAR) reports. This committee reports to the General Assembly at regular intervals; the most recent report appeared in 2000. The second is the committee of the U.S. National Academy of Sciences known as the Biologic Effects of Ionizing Radiation (BEIR). Reports appear periodically, the most recent comprehensive report (BEIR VII) appearing in 2006. To a large extent, these are “scholarly” committees, inasmuch as they are under no compulsion to draw conclusions if data are not available.

On the other hand, there are committees involved with radiation protection that cannot afford to be scholarly because they must make recommendations whether or not adequate data are available. First, there is the International Commission on Radiological Protection (ICRP). This commission was originally set up and funded by the first International Congress of Radiology. Over the years, the funding base of this commission has broadened, and it has assumed the role of an independent, self-propagating committee. At a national level in the United States, there is the National Council on Radiological Protection and Measurements (NCRP). This is an independent body chartered by Congress and is funded from industry, government grants, and professional societies. The NCRP formulates policies for radiation protection in the United States often, but not always, following the lead of the ICRP. The recommendations of the NCRP carry no weight in law but are usually adopted eventually and enforced by the regulatory agencies in the United States, although there can often be a long lag period. (See Chapter 17 on radiation protection for more regarding these committees.)

### RADIATION-INDUCED CANCER IN HUMAN POPULATIONS

Under appropriate conditions, a malignancy can be induced in essentially all tissues of the body. Some of the most common are discussed below.

#### Leukemia

The incidence of chronic lymphocytic leukemia does not appear to be affected by radiation. Acute and chronic myeloid leukemia are the types chiefly responsible for the excess incidence observed in irradiated adults. Susceptibility to acute lymphatic or stem cell leukemia seems to be highest in childhood and to decrease sharply during maturation.

Two principal population groups provide data to determine risk estimates:

1. Survivors of the atomic bomb attacks on Hiroshima and Nagasaki
2. Patients treated for ankylosing spondylitis

Leukemia was the first malignancy to be linked with radiation exposure in the A-bomb survivors and has the highest relative risk of any malignancy. Leukemia risks increased with dose up to about 3 Sv, with evidence of upward curvature; that is, a linear-quadratic function of dose fits the data significantly better than a linear function. Because of this curvature, the risk per unit of dose at 1 Sv is about three times greater than at 0.1 Sv.

For those exposed younger than age about 30, nearly all of the excess deaths occurred before 1975, but for those exposed at older ages,
the excess risk appeared to persist throughout the follow-up period. Because of these complications, simple models cannot adequately summarize leukemia risks.

**Thyroid Cancer**

The thyroid gland is an organ of high sensitivity for radiation carcinogenesis, at least in children; in adults, radiation is much less efficient in inducing thyroid cancer. The malignant tumors that have been produced, however, consistently have been of a histologically well-differentiated type, which develops slowly and often can be removed completely by surgery or treated successfully with radioactive iodine if metastasized; consequently, these tumors show a low mortality rate. It is estimated that about 5% of those with radiation-induced thyroid cancer die as a result.

The following are the principal population groups available for deriving risk estimates for thyroid cancer:

1. Survivors of the atomic bomb attacks on Hiroshima and Nagasaki.
2. Residents of the Marshall Islands exposed to external radiation and ingested iodine-131 from fallout after the 1954 testing of a thermonuclear device, in whom there was a high incidence of nodule formation and some thyroid cancer (benign as well as malignant tumors).
3. Individuals who ingested radioactive iodine as a result of the Chernobyl accident (this experience shows how very sensitive children are and that adults are relatively resistant).
4. Children treated with x-rays for an enlarged thymus.
5. Children treated for diseases of the tonsils and nasopharynx.
6. Children epilated with x-rays for the treatment of tinea capitis.
7. Children treated for cancer.

Figure 10.3 shows the relative risk for thyroid cancer after exposure to external radiation, taken from a pooled analysis of seven studies, which dramatically illustrates the importance of age at exposure. (Figure prepared by Dr. Elaine Ron, based on the data from Ron E, Lubin JH, Shore RE, et al. Thyroid cancer after exposure to external radiation: a pooled analysis of seven studies. *Radiat Res.* 1995;141:259–277.)

There are three principal exposed populations from which the risk of breast cancer incidence may be derived:

2. Female patients in a Nova Scotia sanatorium subjected to multiple fluoroscopies during artificial pneumothorax for pulmonary tuberculosis. There is doubt about the dosimetry, but the dose to breast tissue per fluoroscopy is estimated to have been 0.04 to 0.2 Gy. The number of examinations commonly exceeded 100, and in some instances, women received more than 500 fluoroscopies; three patients, in fact, developed radiation dermatitis. This group of exposed women probably constitutes the most convincing evidence of the production of cancer by fractionated x-rays used for diagnosis. This Canadian study also showed the importance of age at the time of exposure. The study was later confirmed by the follow-up of patients discharged from two tuberculosis sanatoria in Massachusetts. These patients were examined fluoroscopically at an average of 102 times over a period of years and, subsequently, were...
found to be 80% more likely to develop breast cancer than a comparable unexposed population.

3. Females treated for postpartum mastitis and other benign conditions. Patients typically received 1 to 6 Gy and showed an excess incidence of breast cancer compared with the general female population of New York State. A legitimate objection to the use of these data for risk estimates is the uncertainty of whether postpartum mastitis predisposes to breast cancer.

4. The data for excess incidence of breast cancer in these populations are shown in Figure 10.4. Several interesting points are immediately apparent. First, the data from the New York series of postpartum mastitis patients are so poor that they do not give any clue about the shape of the dose–response relationship. Second, there is a marked difference in the natural incidence of breast cancer in Japanese women in whom it is low, compared with American and Canadian women in whom it is high; nevertheless, in all cases, incidence rises with radiation dose. Third, the data for breast cancer are reasonably well fitted by a straight line.

**Lung Cancer**

Radiation is but one of a long list of carcinogens for lung cancer: Cigarette smoking, asbestos, chromium salts, mustard gas, hematite, and asphalt derivatives have also been implicated.

**FIGURE 10.4** Incidence of breast cancer as a function of dose for four human populations that allow risk estimates to be made. The data are expressed in terms of the number of cases per 100,000 women-years (WY). Note that the natural incidence of breast cancer is low in Japanese women and high in American and Canadian women. (Adapted from Boice JD Jr, Land CE, Shore RE, et al. Risk of breast cancer following low-dose exposure. *Radiology*. 1979;131:589–597, with permission.)
Radiation risk estimates come from two principal sources:

1. Individuals exposed to external sources of radiation, including the Japanese survivors and those with ankylosing spondylitis. An excess was found even when smoking was taken into account.

2. Underground miners exposed to radon in the mine atmosphere. The naturally occurring deposits of radioactive materials in the rocks of the earth decay through a long series of steps until they reach a stable isotope of lead. One of these steps involves radon, which, unlike the other elements in the decay series, is a gas. In the closed environment of a mine, workers inhale radon gas and some radon atoms decay to the next solid member of the radioactive series, which consequently is deposited on the bronchial epithelium. Subsequent steps in the radioactive decay series take place in the lungs, causing intense α-irradiation of localized surrounding tissue.

There is a clear excess of lung cancer among workers in the uranium mines of the Colorado plateau in the United States, the uranium mines in Czechoslovakia, the nonuranium mines in Sweden, and the fluorspar mines in Newfoundland. It remains difficult to separate adequately the contributory effects of radon and cigarette smoking in causing the cancers, because there are too few nonsmoking miners to form an adequate control group. In addition, the average duration of exposure usually spans from 15 to 20 years, during which standards of safety and ventilation have changed substantially. In any case, it is no easy matter to estimate the dose to the critical cells in the basal layer of the epithelium of the lung from knowledge of the radon concentration in the air that is breathed. There is also some evidence, summarized in the BEIR VI report, of an excess of lung cancer from domestic radon exposure. It is estimated that 10% of the 150,000 lung cancer deaths annually in the United States are caused by radon.

**Bone Cancer**

There is some evidence of bone cancer induced by external x-irradiation in children epilated for the treatment of tinea capitis and in patients treated for ankylosing spondylitis. The numbers are small and the risk estimates poor. The largest body of data comes from two populations, each of which ingested isotopes of radium that emit high linear energy transfer (LET) α-particles and that follow the metabolic pathways of calcium in the body to become deposited in the bone. The populations include the following:

1. Young persons, mostly women, employed as dial painters, who ingested radium as a result of licking their brushes into a sharp point for application of luminous paint to watches and clocks. In this group, there have been bone sarcomas and carcinomas of epithelial cells lining the paranasal sinuses and nasopharynx. None of these tumors occurred at doses below 5 Gy; above this level, the incidence rose sharply, particularly the sarcomas. The radium in these paints consisted of the isotopes radium-226 and radium-228, with half-lives of about 1,600 years and 6 years, respectively.

2. Patients given injections of radium-224 for the treatment of tuberculosis or ankylosing spondylitis.

There are three points that need to be emphasized. First, the dose is made up of α-particles, which have a short range and deposit their energy close to the site at which the isotope is deposited; α-particles are also more effective than x-rays by a factor of about 20. Second, osteosarcomas arise predominantly from endosteal cells, and the relevant dose for estimating the risk of sarcoma is the dose to these cells, which lie at a distance of up to 10 μm from the bone surface, rather than the mean dose throughout the bone. Radium-224 has a short half-life (3.6 days), and its radiation therefore is largely delivered while it is still present on the bone surface. This contrasts sharply with radium-226 and radium-228, which have long half-lives and, consequently, become distributed throughout the bone during their periods of radioactive decay. The dose to endosteal cells from radium-224 is about nine times larger than the dose averaged throughout bone, whereas it is about two-thirds of the mean bone value from radium-226. Consequently, it is difficult to compare data from the two groups of people who were exposed to these very different
x-ray technicians, in an era in which safety standards were virtually nonexistent. In most cases, the onset of neoplasms followed chronic radiodermatitis and a long latent period. Squamous cell and basal cell carcinomas have been most frequently observed, and occasionally, a sarcoma of the subcutaneous tissues has been seen. Since the evolution of modern safety standards, epidermoid carcinoma has ceased to be an occupational disease of radiation workers.

Radiation-induced skin cancers are diagnosed readily and treated at an early stage of development, and there is a large difference between rates of incidence and mortality. There is a small excess incidence of skin cancer in the children epilated with x-rays for the treatment of tinea capitis.

### Quantitative Risk Estimates for Radiation-Induced Cancer

Despite a diverse collection of data for cancer in humans from medical sources, both the BEIR and UNSCEAR reports elect to base their risk estimates almost entirely on the data from the survivors of the atomic bomb attacks on Hiroshima.
and Nagasaki. Figure 10.6 summarizes the study groups available from the Radiation Effects Research Foundation (RERF).

1. The Life Span Study (LSS), comprising about 120,000 people, allows estimates to be made of the radiation-induced cancer incidence and cancer mortality.
2. The in utero cohort, comprising about 3,300 people who were exposed to radiation from the bombs while in utero, allows estimates to be made of the incidence of malformations, growth retardation, microcephaly, and mental retardation.
3. The children of the exposed persons, the so-called F1 generation, allows estimates to be made of heritable effects.

Figure 10.7 charts the incidence of radiation-associated deaths following the A-bomb attacks in 1945. Leukemia was the first malignancy to be linked with radiation exposure in bomb survivors and has the highest relative risk of any malignancy. Leukemia deaths reached a peak of 5 to 7 years after irradiation, subsequently falling rapidly. For those exposed younger than the age of about 30 years, nearly all of the excess deaths occurred within 30 years, but for those exposed at older ages, the excess risk appears to persist throughout the follow-up period. An excess of solid tumors did not appear at first, but once they did, excess deaths have continued up to the present time. There are about six solid cancers for each leukemia. Since about 1990, there is evidence for the induction of noncancer effects, particularly heart disease, stroke, digestive disorders, and respiratory disease, particularly at higher doses of around 1 Sv. For these noncancer end points, it is not possible to say with any certainty whether there is a threshold, nor is it clear what cellular or tissue mechanisms might underlie such a diverse set of disorders.

Table 10.1 shows a summary of the data for cancer incidence in the atomic bomb survivors and Nagasaki. Figure 10.6 summarizes the study groups available from the Radiation Effects Research Foundation (RERF).

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**TABLE 10.1**

<table>
<thead>
<tr>
<th>Cohort</th>
<th>Size</th>
</tr>
</thead>
<tbody>
<tr>
<td>Life Span Study</td>
<td>120,000</td>
</tr>
<tr>
<td>Allows an estimate of cancer incidence and mortality</td>
<td></td>
</tr>
<tr>
<td>In-Utero Cohort</td>
<td>3,600</td>
</tr>
<tr>
<td>Allows estimates of mental retardation, microcephaly, etc.</td>
<td></td>
</tr>
<tr>
<td>Children of exposed individuals</td>
<td>77,000</td>
</tr>
<tr>
<td>Allows estimate of heritable effects</td>
<td></td>
</tr>
</tbody>
</table>
up to 1998. The raw data are shown principally to emphasize the relative poverty of the data; only a few hundred excess cancer cases caused by radiation are involved, compared with many thousands of naturally occurring malignancies—and these must be allocated to different dose groups and different sites.

Figure 10.8 shows the data for cancer incidence in the A-bomb survivors for the years 1958–1994. The relative risk is a linear function of dose up to about 2 Sv. Over the lower dose range from 0 to 0.5 Sv, there is a suggestion that the risks are slightly higher than the linear extrapolation from higher doses. There is some uncertainty in the control group (i.e., the zero-dose group) used for comparison. There are in fact two zero-dose groups; N beyond 3,000 m and survivors within 3,000 m who, for one reason or another, were not exposed (e.g., they might have been out of the city at the time). The two groups have slightly different cancer rates, which is not surprising, because one is a rural and the other, an urban population.

### DOSE AND DOSE-RATE EFFECTIVENESS FACTOR

The Japanese data relate only to high dose rates (HDR) because they are based on the atomic bomb survivors. Both the UNSCEAR and BEIR committees considered that there is a dose-rate effect for low LET radiations; that is, fewer malignancies are induced if a given dose is spread out over a period of time at low dose rate (LDR) than if it is delivered in an acute

---

**TABLE 10.1** Solid Cancer 1958 through 1998

<table>
<thead>
<tr>
<th>Dose, Gy</th>
<th>Subjects</th>
<th>Mean Dist, m</th>
<th>Cases</th>
<th>Excess</th>
</tr>
</thead>
<tbody>
<tr>
<td>No in city</td>
<td>25,427</td>
<td>—</td>
<td>3,994</td>
<td>0</td>
</tr>
<tr>
<td>&lt;0.005</td>
<td>35,545</td>
<td>3,969</td>
<td>5,603</td>
<td>3</td>
</tr>
<tr>
<td>0.005–</td>
<td>27,789</td>
<td>2,114</td>
<td>4,406</td>
<td>81</td>
</tr>
<tr>
<td>0.1–</td>
<td>5,527</td>
<td>1,608</td>
<td>968</td>
<td>75</td>
</tr>
<tr>
<td>0.2–</td>
<td>5,935</td>
<td>1,430</td>
<td>1,144</td>
<td>179</td>
</tr>
<tr>
<td>0.5–</td>
<td>3,173</td>
<td>1,260</td>
<td>688</td>
<td>206</td>
</tr>
<tr>
<td>1–</td>
<td>1,647</td>
<td>1,118</td>
<td>460</td>
<td>196</td>
</tr>
<tr>
<td>2–4</td>
<td>564</td>
<td>934</td>
<td>185</td>
<td>111</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>105,427</strong></td>
<td>—</td>
<td><strong>17,448</strong></td>
<td><strong>853</strong></td>
</tr>
</tbody>
</table>

Abbreviations: dist, distance; m, meter.

exposure. The dose and dose-rate effectiveness factor (DDREF) is defined as the factor by which radiation cancer risks observed after large acute doses should be reduced when the radiation is delivered at LDR or in a series of small dose fractions. Animal data are equivocal on the subject with experiments suggesting a DDREF in the range of 2 to 10. For purposes of radiation protection, the ICRP recommends a DDREF of 2, which reflects their policy of being conservative. BEIR VII came up with an even lower value of 1.5 based of the possible slight curvature of the dose–response relationship for solid cancers. It is also clear that the female cancer risks are significantly higher than the male cancer risks, not only because of breast cancer, but also because of lung and bladder cancers, which are affected by smoking. In Japan in 1945, smoking was common in males, but not in females.

These estimates from the BEIR committee are for all solid cancers lumped together and for exposure. The dose and dose-rate effectiveness factor (DDREF) is defined as the factor by which radiation cancer risks observed after large acute doses should be reduced when the radiation is delivered at LDR or in a series of small dose fractions. Animal data are equivocal on the subject with experiments suggesting a DDREF in the range of 2 to 10. For purposes of radiation protection, the ICRP recommends a DDREF of 2, which reflects their policy of being conservative. BEIR VII came up with an even lower value of 1.5 based of the possible slight curvature of the dose–response relationship for solid cancers.

### SUMMARY OF RISK ESTIMATES

The population averaged cancer risk estimates from the BEIR VII committee are summarized in Table 10.2. As would be expected, the radiation-induced cancer incidence at 10.8% per Sv is approximately double the cancer mortality at 5.4% per Sv. It is also clear that the female cancer risks are significantly higher than the male cancer risks, not only because of breast cancer, but also because of lung and bladder cancers, which are affected by smoking. In Japan in 1945, smoking was common in males, but not in females.

These estimates from the BEIR committee are for all solid cancers lumped together and for exposure. The dose and dose-rate effectiveness factor (DDREF) is defined as the factor by which radiation cancer risks observed after large acute doses should be reduced when the radiation is delivered at LDR or in a series of small dose fractions. Animal data are equivocal on the subject with experiments suggesting a DDREF in the range of 2 to 10. For purposes of radiation protection, the ICRP recommends a DDREF of 2, which reflects their policy of being conservative. BEIR VII came up with an even lower value of 1.5 based of the possible slight curvature of the dose–response relationship for solid cancers.
whereas adults are quite resistant. It is also dramatic for breast cancer in females; females exposed before 15 years of age are most susceptible; women 50 years of age or older show little or no excess. Figure 10.10 shows the variation of cancer incidence as a function of age for males and females as calculated by the BEIR VII committee from the A-bomb data. There are exceptions to this general rule. Susceptibility to radiation-induced leukemia is relatively constant throughout life, and susceptibility to respiratory cancers increases in middle age. The overall risk, however, drops dramatically with age; children and young adults are much more susceptible to radiation-induced cancer than the middle- and old-aged.

An important question is the lowest dose at which there is epidemiologic evidence of a radiation-induced excess cancer incidence. There is a population of about 30,000 A-bomb survivors who all age groups. The data from the A-bomb survivors also make it possible to calculate organ-specific risk estimates. These are summarized in Figure 10.9. It appears that the bladder, breast, lung, thyroid, and colon are more radiosensitive than the average, whereas the stomach and liver are less sensitive. These data are of enormous importance because they can be used, for example, to calculate cancer risks from diagnostic or therapeutic procedures where only a specific area of the body is irradiated.

As the data from Japan have matured and more detailed information has become available, it is evident that the risk of radiation-induced cancer also varies considerably with age at the time of exposure. In most cases, those exposed at an early age are much more susceptible than those exposed at later times. The difference is most dramatic for thyroid cancer; children are very radiosensitive, whereas adults are quite resistant. It is also dramatic for breast cancer in females; females exposed before 15 years of age are most susceptible; women 50 years of age or older show little or no excess. Figure 10.10 shows the variation of cancer incidence as a function of age for males and females as calculated by the BEIR VII committee from the A-bomb data. There are exceptions to this general rule. Susceptibility to radiation-induced leukemia is relatively constant throughout life, and susceptibility to respiratory cancers increases in middle age. The overall risk, however, drops dramatically with age; children and young adults are much more susceptible to radiation-induced cancer than the middle- and old-aged.

An important question is the lowest dose at which there is epidemiologic evidence of a radiation-induced excess cancer incidence. There is a population of about 30,000 A-bomb survivors who

![Figure 10.9](image_url) The study of the A-bomb survivors also makes it possible to calculate cancer risk estimates for some specific organs. In this figure, they are expressed in terms of the ERR/Sv. These figures are useful for estimating the possible risks from medical radiation where often only part of the body is exposed.

![Figure 10.10](image_url) Illustrating how cancer incidence from radiation exposure falls dramatically with age. Children are 10 times more sensitive than older adults. It is also clear that females are more radiosensitive than males. Based on data from the BEIR VII report.
lived in the outskirts of the two cities and were exposed to doses in the range of 5 to 100 mSv. This low-dose sub population has been studied for more than 50 years, and shows a small, but statistically significant increased cancer risk.

**SECOND MALIGNANCIES IN RADIOTHERAPY PATIENTS**

The risk of second malignancies after radiotherapy is a subject not without controversy. One of the reasons for the uncertainty is that patients undergoing radiotherapy are often at high risk of a second cancer because of their lifestyles, and this factor is more dominant than the radiation risk.

There are many single institution studies in the literature involving radiotherapy from various sites that conclude that there is no increase in second malignancies, although a more accurate assessment would have been that the studies had limited statistical power to detect a relatively small increased incidence of second malignancies induced by the treatment.

Whenever large studies have been performed, radiotherapy has been shown to be associated with a statistically significant, although small, enhancement in the risk of second malignancies, particularly in long-term survivors. The three requirements for a study to be credible are as follows:

1. A sufficiently large number of patients.
2. A suitable comparison group; that is, patients with the same cancer treated by some means other than radiation.
3. A sufficiently long follow-up for radiation-induced solid tumors to manifest.

Only a few studies satisfy these criteria; these will be further discussed in details.

**Second Cancers after Radiotherapy for Prostate Cancer**

Brenner and colleagues described a study using data from the National Cancer Institute's Surveillance, Epidemiology, and End Results (SEER) program. The SEER program is a set of geographically defined, population-based tumor registries, covering approximately 10% of the US population. The database contained information on 51,584 men with prostate cancer treated by radiotherapy and 70,539 men who underwent surgery. There was no evidence of a difference in the risk of leukemia for radiotherapy versus surgery patients, but the risk of a second solid tumor at any time postdiagnosis was significantly greater after radiotherapy than after surgery. The relative risk increased with time posttreatment and reached 34% after 10 years or more. The most dramatic increases in relative risk were for the bladder (77%) and the rectum (105%) for 10 years or more following diagnosis. The relative risks are shown in Figure 10.11, together with the distribution of second cancers; note that even sites remote from the treatment area (e.g., lung) show an increased incidence. The absolute risk was about 1 in 70 by 10 years posttreatment. Figure 10.12 shows the relative risk of sarcomas in the heavily irradiated tissues in or near the treatment field. It can be seen that the relative risk increases to more than 200% at 10 years or more, compared with the surgical patients.

It is interesting to note that the increase in relative risk for carcinoma of the lung, which was exposed to a relatively low dose (about 0.5 Gy), is of the same order as that for carcinomas of the bladder, rectum, and colon, all of which were subject to much higher doses (typically more than 5 Gy). This pattern may reflect the fact that carcinomas, originating in actively dividing cells or cells under hormonal control, can be efficiently induced by relatively low doses of radiation as evidenced by the atomic bomb survivors, but the cancer risk at high doses decreases because of the effects of cell killing. In contrast to this pattern for radiation-induced carcinomas, radiation-induced sarcomas appear only in heavily irradiated sites, close to the treatment volume, because large radiation doses are needed to produce sufficient tissue damage to stimulate cellular renewal in mostly dormant cells. The sarcoma data in this study appear to follow this pattern, with significant radiation-related risks being exhibited for sites in and close to the treatment volume, but no significant increases being shown for sites that are more distant.

**Radiation Therapy for Carcinoma of the Cervix**

In the largest study of its kind, Boice and colleagues studied the risk of second malignancies in a wide range of organs and tissues as a consequence of the treatment by radiation of carcinoma of the uterine cervix. This huge international study was a tour de force. The paper had 42 authors from 38 institutions representing both
equally well treated by radiation or surgery. The results can be summarized as follows:

1. Very high doses, on the order of several hundred gray, were found to increase the risk of
cancer of the bladder, rectum, vagina, and possibly bone, uterus corpus, and cecum as well as non–Hodgkin lymphoma. The risk ratios vary from a high of 4.0 for the bladder to a low of 1.3 for the bone. For all female genital cancers combined, a steep dose-response curve was observed, with a 5-fold excess at doses of more than 150 Gy.

2. Doses of several grays increased the risk of stomach cancer and leukemia.

3. Perhaps surprisingly, radiation was found not to increase the overall risk of cancers of the small intestine, colon, ovary, vulva, connective tissue, or breast or of Hodgkin disease, multiple myeloma, or chronic lymphocytic leukemia.

The overall conclusion of this study was that excess cancers certainly were associated with radiotherapy, as opposed to surgery, and that the risks were highest among long-term survivors and concentrated among women irradiated at relatively young ages.

**Second Cancers among Long-Term Survivors from Hodgkin Disease**

Second cancer represents the leading cause of death in long-term survivors of Hodgkin disease, with exceptionally high risks of breast cancer among women treated at a young age. Several studies have been reported. Bhatia and colleagues reported that 17 out of 483 girls in whom Hodgkin disease was diagnosed before the age of 16 years subsequently developed breast cancer, with radiotherapy implicated in most cases. The ratio of observed to expected cases is 75.3. Another study (by Sankila and colleagues) involved 1,641 patients treated for Hodgkin disease as children in five Nordic countries and reported a relative risk that was 17 times higher than the general population based on 16 cases of breast cancer. Travis and colleagues evaluated 3,869 women in population-based registries participating in the SEER program. All these women received radiotherapy as an initial treatment for Hodgkin disease. Breast cancer developed in 55 patients, who represents a ratio of observed to expected cases of 2.24. The risk of breast cancer, however, was 60.57% in women treated before the age of 16 years, with most tumors appearing 10 or more years later. This agrees with previous studies that have shown the female breast to be very radiosensitive to carcinogenesis at young ages. The risk of breast cancer decreased with increasing age at the time of therapy and was only slightly elevated in women who were 30 years old or older when treated. In a later study, Travis and colleagues followed 3,817 female survivors of Hodgkin disease, diagnosed at age 30 years or younger, over a long period of time. A radiation dose of 4 Gy or more delivered to the breast was associated with a 3.2-fold increase in risk. Risk increased 8-fold with a dose of more than 40 Gy.

Hormonal stimulation appears to be important for the development of radiation-induced breast cancer, as evidenced by the reduced risk in patients who received alkylating agents, as well as radiation, which caused ovarian damage.

These studies clearly show that if an adequate cohort can be studied, there is a clear excess of second cancers induced by radiotherapy.

The data confirm previous studies that show that in the young, the breast is especially sensitive to the carcinogenic effects of radiation. In addition, excess cancers develop with a latency of 10 years or more and persist for decades after exposure.

**DOSE–RESPONSE RELATIONSHIP FOR RADIATION CARCINOGENESIS AT HIGH DOSES**

In the 1960s, Gray proposed that the dose–response relationship for radiation-induced malignancies would be bell-shaped, as illustrated in Figure 10.13; that is, the incidence would rise at low doses but fall at high doses. He explained this shape by the concurrent presence of two phenomena: (1) a “dose-related” increase in the proportion of normal cells that are transformed to a malignant state and (2) a dose-related decrease in the probability that transformed cells may survive the radiation exposure. Gray argued that whatever sequence of changes has taken place in the course of cell transformation, the changes must have been such as to leave the cell capable of indefinite proliferation; that is, with full reproductive integrity. The balance between transformation and cell killing leads to the overall shape, with cell killing becoming dominant at increasingly high doses. With Figure 10.13, Gray was specifically attempting to explain the shape of the dose–response relationship for the induction of leukemia in mice exposed to total body irradiation, which is why the dose goes up only to 5 Gy, but it has been tacitly assumed ever since that this bell-shaped curve applies to radiation-induced carcinogenesis in general. However, several recent studies challenge the validity of this assumption by...
examining whether the linear dose response for radiation-induced cancer, evident in the A-bomb survivors at doses up to 2 Sv, extends to the higher dose ranges used for radiotherapy. Two studies involved the incidence of breast cancer in women treated for Hodgkin disease with a mantle field, which results in a large dose gradient across the breast (3 to 42 Gy). There was an increasing risk of breast cancer over this entire dose range. Some of the data from the Hodgkins patients, together with data from the A-bomb survivors, are shown in Figure 10.14, taken from a paper by Brenner and Sachs. It clearly shows that for the Excess Relative Risk (ERR) for high-fractionated doses is larger than at the low doses received by the A-bomb survivors. It certainly does not fall as would be predicted by the Gray model.

Another study from St. Jude Children’s Research Hospital evaluated 1,612 patients with acute lymphoblastic leukemia, whose primary treatment was chemotherapy, but who also received prophylactic cranial irradiation because many chemotherapy agents do not effectively cross the blood–brain barrier (BBB). An excess of high-grade gliomas and meningiomas were evident during the first decade of follow-up, whereas an increased risk of low-grade brain tumors was observed at later follow-up intervals. The risk of brain tumors increased significantly with increasing radiation dose, as shown in Figure 10.15, but there is no sign of the cancer incidence falling at high doses. There is some indication of a plateau, but no fall as would be predicted as cell killing takes over. As a consequence of these studies, prophylactic cranial radiotherapy (PCR) in children with leukemia has been largely replaced by intrathecal methotrexate.
These examples are further evidences that the incidence of radiation-induced solid cancers does not fall at the high-fractionated doses typically used therapeutically and accords with the clinical observation that second cancers often occur in or near the treatment field in high-dose areas, as well as in more remote locations.

**CANCER RISKS IN NUCLEAR INDUSTRY WORKERS**

The International Agency for Research on Cancer (IARC), a branch of the World Health Organization (WHO), conducted an impressive epidemiologic study of cancer mortality among 400,000 nuclear workers from 15 countries. The importance of this study stems from the fact that nuclear workers receive protracted exposures to multiple low doses of radiation over many years, in contrast to the acute instantaneous dose received by the Japanese A-bomb survivors. The surprising result was a statistically significant excess of solid cancers for a mean dose of only 19.4 mSv. The data are shown in Figure 10.16. Furthermore, the excess relative risk per sievert (ERR/Sv) was 0.97, more than three times larger than the corresponding quantity for the A-bomb survivors (see Table 10.3).

However, these results need to be viewed with caution for two reasons.

1. While data from 15 nations were pooled, the overall solid cancer risk is driven by the Canadian data, which is evident from Figure 10.16. Indeed, if the data from Canada are excluded, the excess of solid cancer deaths no longer has significance.

2. Lung cancer is prominent in the excess solid cancers, suggesting a confounding effect of smoking, a possibility recognized by the authors of the study.

**FIGURE 10.16** Excess relative risk illustrating the data on cancer mortality from the 15-country study of nuclear workers by the IARC. With all countries combined, the ERR is statistically significant. However, the result is driven by the Canadian data, which makes a disproportionate contribution and casts some doubt on the validity of the study. In addition, there are a disproportionate number of cancers of the lung, which raise the possibility that the confounding effect of smoking has not been adequately accounted for. (Adapted from Cardis E, Vrijheid M, Blettner M, et al. Risk of cancer after low doses of ionising radiation: retrospective cohort study in 15 countries BMJ. 2005;331:77, with permission.)
The data from the A-bomb survivors, one involving a protracted exposure and the other one an acute instantaneous exposure, supports the low DDREF values suggested by both the ICRP and BEIR VII committees (i.e., a given dose appears to result in a similar cancer mortality whether delivered in an instantaneous acute exposure, or spread out over a protracted period of years).

### MORTALITY PATTERNS IN RADIOLOGISTS

An extensive and interesting report was published by Sir Richard Doll and colleagues in 2003 that assessed 100 years of observations in terms of mortality from cancer and other causes in British radiologists entering the field from 1897 to 1997. Table 10.4 shows the trend in the Standard Mortality Ratio (SMR) over the years. There was a clear excess of cancer in the early radiologists in the years before the introduction of radiation safety standards. This is not surprising, in that, estimated annual doses to early radiologists were typically in the range of 1 Gy per year. What is surprising is that the SMR for the post-war period, 1955 to 1979 is much smaller than unity, due largely to a statistically significant lower rate of noncancer deaths. This attracted much interest, leading to speculation by some that low doses of radiation may be beneficial and may actually lead to a longer life!

A weakness of the British study is that the data for the control group, labeled “all-male medical practitioners” were in fact estimated indirectly from census data, with adjustments made...

<table>
<thead>
<tr>
<th>Years</th>
<th>Standard Mortality Ratio</th>
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<tbody>
<tr>
<td>1897–1920</td>
<td>1.75</td>
</tr>
<tr>
<td>1921–1935</td>
<td>1.24</td>
</tr>
<tr>
<td>1936–1954</td>
<td>1.12</td>
</tr>
<tr>
<td>1955–1979</td>
<td>0.71</td>
</tr>
<tr>
<td>All post-920</td>
<td>1.04</td>
</tr>
</tbody>
</table>

that radiation was the causative agent. They concluded that:

- Low-dose irradiation of the fetus in utero, particularly in the last trimester, causes an increased risk of childhood malignancies.
- An obstetric x-ray examination, even though the dose is only about 10 mGy, increases the risk of childhood cancer by 40%.
- The excess absolute risk is about 6% per gray. The relative risk of 40% is very high because, of course, cancer is relatively rare in children. The absolute risk works out to be about 6% per gray, which is not very different from the cancer risk calculated for the atomic bomb survivors following adult exposure.

### NONNEOPLASTIC DISEASE AND RADIATION

A link between exposure to high doses of ionizing radiation and damage to the heart and coronary arteries has been well established for many years, based on the treatment of breast cancer during the era of cobalt-60 units, characterized by a large penumbra to the treatment field, which inevitably resulted in an appreciable dose to the heart. An association between lower doses of radiation and late occurring cardiovascular and other diseases emerged in the 1990s from a study of the A-bomb survivors (see Fig. 10.7). Statistically, significant associations were seen for the categories of heart disease, stroke, and diseases of the digestive, respiratory, and hematopoietic systems. However, the data are inadequate to distinguish between a linear dose...
The shortest latency is for leukemia, with a peak of 5 to 7 years. For solid tumors, the latency may extend for 60 years or more.

Regardless of the age at exposure, radiation-induced malignancies tend to appear at the same age as spontaneous malignancies of the same type. Indeed, for solid cancers, the excess risk is apparently more like a lifelong elevation of the natural age-specific cancer risk.

To determine risk estimates for radiation-induced cancer from observed data (the Japanese atomic bomb survivors), a model must be assumed because of the following:
1. Data must be extrapolated from relatively high doses to the low doses of public health concern.
2. Data must be projected out to a full life span, because large exposed populations, such as the A-bomb survivors, have not yet lived out their life span.
3. Risks must be “transferred” from the Japanese population to, for example, a Western population with different natural cancer rates.

There are two principal risk models: The absolute risk model assumes that radiation produces a discrete “crop” of cancers, over and above the spontaneous level and unrelated to the spontaneous level. The relative risk model assumes that radiation increases the spontaneous incidence by a factor. Because the natural cancer incidence increases with age, this model predicts excess cancers appearing late in life after irradiation.

The assessment of radiation-induced cancer risks by both the BEIR and UNSCEAR committees is based on a time-related relative risk model. Excess cancer deaths were assumed to depend on dose, square of the dose, age at exposure, time since exposure, and, for some cancers, gender.

For solid tumors, the A-bomb data show that both the excess cancer incidence and mortality are a linear function of dose up to about 2 Sv.

Leukemia data were best fitted by a linear-quadratic function of dose (i.e., an upward curvature), so that the risk per unit of dose at 1 Sv is about three times that at 0.1 Sv.
The Japanese atomic bomb data refer to acute exposure at an HDR. A DDREF is needed to convert risk estimates to the low dose and LDR encountered in radiation protection. From animal studies, this is anywhere from 2 to 10. The ICRP conservatively assumes a value of 2, whereas the BEIR VII committee assumes a value of 1.5.

The BEIR VII committee suggests a risk estimate of excess cancer incidence of 10.8% per sievert and excess cancer mortality of 5.4% per sievert, including a DDREF of 1.5. These figures represent a population average, with risks for females slightly higher than for males.

There is a marked reduction with age of the risk of both cancer incidence and cancer mortality. Children are so much more radiosensitive than adults.

The ICRP estimates that, on average, 13 to 15 years of life are lost for each radiation-induced cancer and that death occurs at age 68 to 70 years.

There is a clear excess of second cancers induced by radiation therapy, both in heavily irradiated tissue and in more remote organs. This is evident if a sufficiently large number of patients and an adequate control group are available for study and if there is a sufficiently long follow-up for solid tumors to manifest.

Large studies show a clear excess of second cancers after radiotherapy for prostate cancer, carcinoma of the cervix, and Hodgkin lymphoma. An excess has also been shown following radiation therapy for breast cancer, carcinoma of the testes, and various childhood malignancies.

The IARC studied 400,000 nuclear workers from 15 countries and found a statistically significant excess of solid cancers at a mean dose of 19.4 mSv. However, this result must be viewed with caution because the result is dominated by the Canadian data, and in addition, the large incidence of lung cancer suggests that smoking may be a confounding factor. The UK NRRW studied about 175,000 nuclear workers in the UK over a very long period of time. The study showed the usual “healthy worker effect,” in that, rates of mortality from all causes were significantly lower than those expected from national mortality data, but the cancer risk increased with cumulative dose and the slope of this trend (ERR/Sv) is very similar to the corresponding figure for the A-bomb survivors.

Early radiologists who practiced prior to the 1920s showed an excess of malignancies. No excess is evident in radiologists in recent years. The report that British radiologists live longer is not confirmed in later studies.

Irradiation in utero by diagnostic x-rays appears to increase the spontaneous incidence of leukemia and childhood cancers by a factor of about 1.4. This is a high relative risk because malignancies in children are rare, but the absolute risk is about 6% per gray—not very different from the risk estimate calculated for the A-bomb survivors following adult exposure.

From a study of both radiotherapy patients and the A-bomb survivors, it is evident that doses of more than about 0.5 Sv can also result in an excess of nonneoplastic diseases, including cardiovascular diseases.

**BIBLIOGRAPHY**


Gray LH. Radiation biology and cancer. In: Cellular Radiation Biology: A Symposium Considering Radiation Effects in the Cell and Possible Implications for Cancer Therapy; A Collection of Papers. Published for the University of Texas MD Anderson Hospital and Tumor Institute, Baltimore, MD, Lipphart Williams & Wilkins; 1965:25.


In the male mammal, spermatozoa arise from the germinal epithelium in the seminiferous tubules of the testes, and their production is continuous from puberty to death. The spermatogonial (stem) cells consist of several different populations that vary in their sensitivity to radiation. The postspermatogonial cells pass through several stages of development: primary spermatocytes, secondary spermatocytes, spermatids, and finally spermatozoa. The division of a spermatogonium to the development of mature sperm involves a period of 6 weeks in the mouse and 10 weeks in the human. The effect of radiation on fertility is not apparent immediately because the postspermatogonial cells are relatively resistant compared with the sensitive stem cells. After exposure to a moderate dose of radiation, the person remains fertile as long as mature sperm cells are available, but decreased fertility or even temporary sterility follows if these are used up. The period of sterility lasts until the spermatogonia are able to repopulate by division.

Radiation doses as low as 0.15 Gy result in oligospermia (diminished sperm count) after a latent period of about 6 weeks. Doses greater than 0.5 Gy result in azoospermia (absence of living spermatozoa) and therefore temporary sterility. The duration of azoospermia is dose dependent; recovery can begin within 1 year after doses of less than 1 Gy but requires 2 to 3.5 years after a dose of 2 Gy. The original single-dose data came from the irradiation of prisoners, which showed that a dose in excess of 6 Gy is needed to result in permanent sterility. In contrast to most organ systems where fractionation of dose results in sparing, fractionated courses cause more gonadal damage than a single dose. Studies of patients receiving radiation therapy indicate that permanent sterility can result from 2.5 to 3 Gy in a fractionated regime over 2 to 4 weeks. The induction of sterility by radiation in human males does not produce significant changes in hormone balance, libido, or physical capability.

Gonadal kinetics in women are opposite to those in men, as the germ cells are nonproliferative. All cells in the oogonial stages progress to the oocyte stage in the embryo. By 3 days after birth, in the mouse or human, all of the oocytes are in a resting phase and there is no cell division. Consequently, in the adult, there are no stem (oogonial) cells, but there are three types of follicles: immature, nearly mature, and mature. At birth, a woman has about 1 million oocytes, which are reduced to about 300,000 at puberty.

In women, radiation is highly effective at inducing permanent ovarian failure, but there is a marked age dependence in sensitivity. The dose required to induce permanent sterility varies from 12 Gy prepubertal to 2 Gy premenopausal. Pronounced hormonal changes, comparable to those associated with the natural menopause, accompany radiation-induced sterilization in females.

Overall, radiation sterility is very different between men and women, and these differences are compared and contrasted in Table 11.1.
sexes, plus a pair of sex chromosomes—the X and Y. Males have 22 pairs of autosomes plus an X and a Y. Females have 22 pairs of autosomes plus a pair of Xs. One chromosome of each pair is derived from each parent.

The human genome is composed of the DNA of chromosomes and, to a minute extent, the DNA of mitochondria. The 46 chromosomes contain about $\frac{6}{1100}$ of $10^9$ base pairs of DNA, with each chromosomal arm including a single supercoiled molecule of DNA associated with chromosomal proteins. The total number of protein-encoding genes is in the range of 25,000 to 50,000 per haploid set of chromosomes. The study of individual genes hardly has begun, but it is apparent that some genes are smaller than this and at least one, whose mutation can cause Duchenne-type muscular dystrophy, has been reported to contain more than $10^3$ kb of DNA. Because most protein products of genes are less than 300 kDa, the translated portions of genes are seldom larger than 10 kb, so a major part of the genome appears to be untranslated. Some of this DNA consists of introns that reside between translated exons.

### REVIEW OF BASIC GENETICS

The study of the inheritance of observable characteristics includes molecular as well as morphologic and some behavioral traits. The chromosomes carry, in code form, all of the information that specifies a particular human, with all of his or her individual characteristics. The chromosomes are long threadlike structures, the essential ingredient of which is DNA, which is itself a long complex molecule with a sugar-phosphate backbone. Attached to each sugar molecule is an organic base; these come in four varieties: thymine, adenine, guanine, and cytosine. This whole configuration is tightly coiled in a double helix, rather like a miniature spiral staircase, with chains of sugar molecules linked by phosphates forming the rails on either side, bridged at regular intervals by pairs of bases, which form the steps. The order or sequence of the bases contains the genetic information in code form.

A gene is a finite segment of DNA specified by an exact sequence of bases. Genes occur along chromosomes in linear order like beads on a string, and the position of a gene is referred to as its locus.

The human chromosome complement consists of 22 pairs of autosomes present in both sexes, plus a pair of sex chromosomes—the X and Y. Males have 22 pairs of autosomes plus an X and a Y. Females have 22 pairs of autosomes plus a pair of Xs. One chromosome of each pair is derived from each parent.

The human genome is composed of the DNA of chromosomes and, to a minute extent, the DNA of mitochondria. The 46 chromosomes contain about $6 \times 10^9$ base pairs of DNA, with each chromosomal arm including a single supercoiled molecule of DNA associated with chromosomal proteins. The total number of protein-encoding genes is in the range of 25,000 to 50,000 per haploid set of chromosomes. The study of individual genes hardly has begun, but it is apparent that some genes are smaller than this and at least one, whose mutation can cause Duchenne-type muscular dystrophy, has been reported to contain more than $10^3$ kb of DNA. Because most protein products of genes are less than 300 kDa, the translated portions of genes are seldom larger than 10 kb, so a major part of the genome appears to be untranslated. Some of this DNA consists of introns that reside between translated exons.

| TABLE 11.1 Radiation Sterility—Comparing Male and Female |
|----------------------------------|----------------------------------|
| **Male** | **Female** |
| Self-renewal system: | Gonadal kinetics opposite of males: |
| Spermatogonia $\rightarrow$ spermatocytes $\rightarrow$ spermatids $\rightarrow$ spermatozoa | by 3 days after birth, all cells progressed to oocyte stage; no further cell division |
| Latent period between irradiation and sterility | Neither latent period nor temporary sterility in females |
| Oligospermia and reduced fertility: 0.15 Gy | — |
| Azoospermia and temporary sterility: 0.5 Gy | — |
| Recovery is dose dependent (1 yr after 2 Gy) | — |
| Permanent sterility: | Radiation can induce permanent ovarian failure; marked age dependence |
| 6 Gy, single dose | Permanent sterility: 12 Gy, prepuberty |
| 2.5–3 Gy, fractionated, 2–4 wks | 2 Gy, premenopausal |
| Induction of sterility does not affect hormone balance, libido, or physical capability | Radiation sterility produces hormonal changes like those seen in natural menopause |
In addition, much of the DNA outside the exons is involved in gene regulation and RNA polymerase attachment.

Not only does this genome recombine in each generation, but it also may undergo mutation, a term used here to denote any change in chromosomes, their genes, and their DNA. Thus, alterations in chromosome number and structure would be included along with changes not visible microscopically. These latter changes include an array of changes in DNA, such as deletion, rearrangement, breakage in the sugar-phosphate backbone, and base alterations. Gene function can be disturbed not only by loss or by modification of translated exons, but also through alteration of nonexonic sites that regulate transcription and translation. Mutation occurs in both germ cells and somatic cells, although it is much less apparent in the latter unless it occurs under conditions of clonal proliferation, as happens with cancer. On the other hand, many mutations in the germ line are lethal during embryonic development.

In humans, every normal cell has 46 chromosomes, 23 derived from the mother and 23 from the father. The two members of a pair of chromosomes normally have the same genes for given characteristics lined up in the same sequence. In this case, the two chromosomes are said to be homologous. The pair of chromosomes that determine sex are XX in the female and XY in the male; in the case of the male, therefore, the two chromosomes of this pair are heterologous; they do not contain parallel genes. If the two members of a pair of genes are alike, the person is said to be homozygous for that pair of genes; if they are different, the person is said to be heterozygous.

The fact that pairs of chromosomes contain corresponding sets of genes introduces the idea of dominant and recessive genes. A dominant gene, by definition, expresses itself if its corresponding gene is recessive, the recessive gene in this case being either ineffective or suppressed. A completely recessive gene is expressed only if both corresponding genes of a pair of chromosomes are recessive (i.e., the person must be homozygous for the recessive gene) or if the recessive gene is on the X chromosome in a male.

Eye color is the simplest example. The gene for blue eyes is recessive; that for brown eyes is dominant. A child will have blue eyes only if he or she receives the gene for blue eyes from both parents. If both or only one of the genes that determine eye color is for brown eyes, then the child will have brown eyes because this gene is dominant. It should be pointed out that not all genes are completely dominant; some permit expression of the recessive counterpart to a varying extent, depending on the particular characteristics involved.

The Y sex chromosome in humans has genes that determine maleness but appears to have few other genes. The X chromosome, on the other hand, has many genes. If a mother carries a recessive mutant gene on the single X chromosome that she donates to her son, there is no matching gene from the father, and consequently, the recessive gene is expressed. If the offspring is a daughter, there may well be a dominant gene on the X chromosome supplied by the father, which would suppress the expression of the recessive mutant. The daughter, however, could transmit the mutant gene to her son in whom it would be expressed. Characteristics that result from recessive genes on the X chromosome, so that they are expressed almost exclusively in male children, are said to be sex-linked. The most common examples are color blindness and hemophilia.

An elementary discussion of genetics, such as presented here, may give the impression that each characteristic of a person is determined by a single pair of genes. On the contrary, this is the exception rather than the rule because most characteristics are the result of an interplay in the expression of many genes.

**Mutations**

Exposure of a population to radiation can cause adverse health effects in the descendants as a consequence of mutations induced in the germ cells. Heritable diseases, also known as genetic diseases,* may result when mutations occurring in the germ cells of parents are transmitted to progeny; in contrast, most cancers result from mutations in somatic cells. Because the human

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*Such diseases were formerly called genetic diseases, but the term heritable is more descriptive, reflecting the transfer from one generation to the next, to distinguish them from cancer, which is also, in essence, genetic.
It is a commonly held view that radiation produces bizarre mutations or monsters that may be recognized readily. This view is not true. Radiation increases the incidence of the same mutations that occur spontaneously in a given population. The study of radiation-induced heritable effects is difficult because the mutations produced by the radiation must be identified on a statistical basis in the presence of a high natural incidence of the same mutations.

The genome includes between 25,000 to 50,000 genes, the potential number of mutations, and thus heritable diseases, is staggering.

It is a commonly held view that radiation produces bizarre mutants and monsters, as illustrated in Figure 11.1. This view is absolutely false. Radiation does not result in heritable effects that are new or unique but rather increases the frequencies of the same mutations that already occur spontaneously or naturally in that species.

Heritable diseases are classified into three principal categories: mendelian, chromosomal, and multifactorial (Table 11.2). The baseline frequencies of these different classes of heritable diseases in the human population have been estimated by the United Nations Scientific Committee on the

<table>
<thead>
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<th>Table 11.2 Heritable Effects of Radiation</th>
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<tr>
<td>Heritable Effect</td>
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<tr>
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</tr>
<tr>
<td>Gene mutations*</td>
</tr>
<tr>
<td>Single dominant 736 (753)</td>
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<td>Recessive 521 (596)</td>
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<tr>
<td>Sex-linked 80 (60)</td>
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<tr>
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<td>Too many or too few</td>
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<td>Multifactorial</td>
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<tr>
<td>Congenital abnormalities present at birth</td>
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<td>Chronic diseases of adult onset</td>
</tr>
</tbody>
</table>

*The numbers following types of gene mutations refer to several human diseases known to be caused by such a mutation. The numbers in parentheses refer to additional possible diseases.
Effects of Atomic Radiation (UNSCEAR) 2001 committee and are summarized in Table 11.3.

**Mendelian**
Mendelian diseases are those caused by mutations in single genes located on either the autosomes or the sex chromosomes. The mutation may be a change in the structure of DNA, which may involve either the base composition, the sequence, or both. An alteration so small that it involves the substitution, gain, or loss of a single base can be the cause of significant heritable changes. A striking example is sickle cell anemia, which results from the substitution of only one base. The important point with respect to mendelian diseases is that the relationship between mutation and disease is simple and predictable.

These diseases are subdivided into autosomal dominant, autosomal recessive, and X-linked conditions, depending on which chromosome the mutant genes are located on and the pattern of transmission. In the case of dominant diseases, one mutant gene received from either one of the parents is sufficient to cause disease, although the copy of the gene from the other parent is normal.

A dominant gene mutation is expressed in the first generation after its occurrence. More than 700 such conditions have been identified with certainty, and an additional 700 or more are less well established. Some examples are polydactyly, achondroplasia, and Huntington chorea. By contrast, recessive mutations, unless sex-linked, require that the gene be present in duplicate to produce the trait, which means that the mutant gene must be inherited from each parent; consequently, many generations may pass before it is expressed. If one copy of the gene is mutant and the other is normal, the person is not affected. More than 500 recessive diseases are known and another 600 are suspected. Some examples are sickle cell anemia, cystic fibrosis, and Tay-Sachs disease.

**X-linked** recessive diseases are caused by mutations in genes located on the X chromosome. The Y chromosome contains far fewer genes than the X. Because males have only one X chromosome, all males having a mutation in the X chromosome show the effect of mutation; like dominant mutations, they are expressed soon after the mutation occurs. Because females have two X chromosomes, they need two mutant genes to show the effect of an X-linked recessive mutation. The best known examples of sex-linked disorders are hemophilia, color blindness, and a severe form of muscular dystrophy, but altogether, there are more than 80 well-established and another 60 probable conditions of this sort.

<table>
<thead>
<tr>
<th>Disease Class</th>
<th>Frequency per Million&lt;sup&gt;a&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mendelian diseases</td>
<td>24,000</td>
</tr>
<tr>
<td>Autosomal dominant diseases</td>
<td>15,000</td>
</tr>
<tr>
<td>X-linked diseases</td>
<td>1,500</td>
</tr>
<tr>
<td>Autosomal recessive diseases</td>
<td>7,500</td>
</tr>
<tr>
<td>Chromosomal diseases</td>
<td>4,000</td>
</tr>
<tr>
<td>Multifactorial diseases</td>
<td>710,000</td>
</tr>
<tr>
<td>Chronic diseases</td>
<td>650,000</td>
</tr>
<tr>
<td>Congenital abnormalities</td>
<td>60,000</td>
</tr>
<tr>
<td>Total</td>
<td>738,000</td>
</tr>
</tbody>
</table>

<sup>a</sup>For mendelian and chromosomal diseases and for congenital abnormalities, the frequencies are per million live births; for chronic multifactorial diseases, the frequency is per million of the population.

In the case of mendelian diseases, about 67% are caused predominantly by point mutations (base pair changes in the DNA), 22% by both point mutations and DNA deletions within genes (i.e., they are intragenic), and 13% by intragenic deletions and large multilocus deletions. In some genes, the mutational sites of mutations are distributed throughout the gene; in a large proportion, however, these are nonrandomly distributed—that is, restricted to specific sites along the gene (specificities). Likewise, the break points of deletions are also nonrandomly distributed, showing specificities.

Some dominant and some recessive mutant genes cause traits that are regarded by society as normal or acceptable, such as different eye colors or blood groups. The majority, however, cause diseases ranging from mild to severe in their impact on the person.

The three types of mendelian diseases are summarized as follows:

Autosomal dominant: The disease is caused by a mutation in a single gene on one chromosome.

Autosomal recessive: The disease is caused by a defective copy of the same gene from each parent.

Sex-linked: Males have one X chromosome, so one mutation can cause the disease; females have two Xs, so two mutant genes are needed to cause the disease.

Cytogenetic Changes

Chromosomal diseases are caused by gross abnormalities either in the structure of the chromosomes or in the number of chromosomes (too many or too few). With a few important exceptions, gross chromosome changes that can be seen under the microscope are not compatible with viability in the fertilized egg. Down syndrome is the best known example: It results from an extra chromosome 21. It is estimated that at least 40% of the spontaneous abortions that occur from the 5th through the 28th week of gestation and about 6% of stillbirths are associated with chromosomal anomalies. This kind of chromosome error is not believed to be influenced strongly by radiation, particularly at low doses. Chromosome breakage is less frequent than aberrations among spontaneous instances of severe human anomalies, but radiation is much more effective at breaking chromosomes than in causing errors in chromosome distribution. Chromosomes that are broken may rejoin in various ways (Chapter 2). A translocation, for example, involves the reciprocal exchange of parts between two or more chromosomes and is not necessarily harmful as long as both rearranged chromosomes are present and contain the full gene complement. Children of a person with a translocation often receive only one of the rearranged chromosomes, and their cells are therefore genetically unbalanced. The nature and extent of the abnormality may vary enormously, and the harm to the person ranges from rather mild to very severe. Chromosome imbalance, if it does not cause the death of the embryo, typically leads to physical abnormalities, usually accompanied by mental deficiency.

Robertsonian translocations are the most common type found in normal humans. These are fusions of two chromosomes, each having a spindle attachment at the end of the chromosome, to produce a single chromosome with the spindle attachment in the center. The children of a person with this type of translocation are usually normal because they inherit either the translocated pair or a pair of normal chromosomes.

Radiation does not appear to be a major cause of robertsonian translocations but rather tends to induce those of the reciprocal-exchange type.

Multifactorial

The term multifactorial is a general designation assigned to a disease known to have a genetic component but whose transmission patterns cannot be described as simple mendelian. The common congenital abnormalities that are present at birth (e.g., neural tube defects, cleft lip with or without cleft palate) and many chronic diseases of adult onset, such as diabetes, essential hypertension, and coronary heart disease, are examples of multifactorial diseases. These diseases result from several causes, both genetic and environmental, the nature of which can vary among persons, families, and populations. For these diseases, there is no simple relationship between mutation and disease, but the fact that genetic factors are involved is evident from observations of familial clustering; that is, these diseases run in families, but the recurrence risk to first-degree relatives is in the range of 5% to 10%, depending on the multifactorial disease, but never close to the 25% to 50% characteristic of mendelian diseases. The potential role of some of the environmental factors has...
been delineated for only a few of these diseases; for example, excess caloric intake rich in saturated fat is an environmental risk factor for coronary heart disease, as are environmental allergens for asthma. Mendelian and chromosomal diseases are rare and account for only a very small proportion of heritable diseases in the population; the major load is from multifactorial diseases.

Characteristics of multifactorial diseases include the following:

- Known to have a genetic component
- Transmission pattern not simple mendelian
- Congenital abnormalities: cleft lip with or without cleft palate, neural tube defects
- Adult onset: diabetes, essential hypertension, coronary heart disease
- Interaction with environmental factors

### RADIATION-INDUCED HERITABLE EFFECTS IN FRUIT FLIES

As early as 1927, Müller reported that exposure to x-rays could cause readily observable mutations in the fruit fly, *Drosophila melanogaster*. These included a change of eye color from red to white, the ebony mutant with its jet-black color, the “vestigial wing” mutant, and the easiest of all to observe, the recessive lethal mutation. The fact that mutants produced by radiation cannot be identified as different compared with those that occur spontaneously makes their study particularly difficult. Sample sizes must be sufficiently large to detect the small excess by radiation, which made *Drosophila* an attractive and economical biologic test system because huge numbers can be accommodated in a small space (compared, for example, with mice). Indeed, it was heritable data from the fruit fly that led to the recommendation of a 5-R maximum permissible annual dose for radiation workers in the 1950s. The units have changed from roentgen (R) to rem to millisievert (mSv), and the justification has also changed, but the quantity remains to this day. In the 1950s, heritable changes were considered the principal hazard of exposure to ionizing radiation. There were three reasons for this:

1. A low **doubling dose** (5–150 R) for mutations was estimated from fruit fly experiments. (The doubling dose is the dose required to double the spontaneous mutation rate.)

2. Based again in the fruit fly experiments, it was thought that heritable effects were cumulative; that is, a little radiation now, some next week, and some next year all added up and contributed to the genetic load carried by the human race.

3. In the 1950s, little was known of the carcinogenic potential of low doses of radiation. An excess incidence of leukemia was evident in the Japanese survivors of the A-bomb attacks, but the much larger number of solid cancers did not appear until many years later.

Over the past 50 years, the heritable risks of radiation have been progressively reduced, replaced by a concern over carcinogenesis as the data from the A-bomb survivors have matured.

### RADIATION-INDUCED HERITABLE EFFECTS IN MICE

The most common way to estimate the heritable risks of radiation is to compare radiation-induced mutations with those that occur spontaneously and to express the results in terms of the doubling dose—the amount of radiation required to produce as many mutations as would occur spontaneously in a generation. The idea is based on the assumption that “if nature can do it, radiation can do it, too.” This is the **relative mutation risk**.

In the years following World War II, the husband-and-wife team of Russell and Russell, working at Oak Ridge National Laboratory, mounted an experiment of heroic proportions to determine specific locus mutation rates in the mouse under various irradiation conditions. This experiment often is referred to as the “megamouse project” because of the enormous number of animals involved. Before the study ended, 7 million mice had been used.

An inbred mouse strain was chosen in which seven specific locus mutations occur, six involving a change of coat color and one expressed as a stunted ear. Figure 11.2 shows three coat-color variations: a piebald, a light honey, and a darker brown. These mutations occur spontaneously, and their incidence is increased by radiation.

These extensive studies included the irradiation of both male and female mice with a range of doses, dose rates, and fractionation...
patterns. Five major conclusions are pertinent to the radiologist:

1. The radiosensitivity of different mutations varies by a significant factor of about 35, so that it is only possible to speak in terms of average mutation rates. We now know that this is caused simply by a size difference among the various genes involved.

2. In the mouse, there is a substantial dose-rate effect, so that spreading the radiation dose over a period results in fewer mutations for a given dose than in an acute exposure. This is in complete contrast to the data on *Drosophila*, where fractionated doses are cumulative. The big dose-rate effect discovered in the mouse was attributed to a repair process. The data are shown in Figure 11.3.

3. Essentially, all of the radiation-induced heritable data come from experiments with male mice. In the mouse, the oocytes are exquisitely radiosensitive and are readily killed by even low doses of radiation. For this reason, the mouse was an unfortunate choice as an experimental animal.

4. The heritable consequences of a given dose can be reduced greatly if a time interval is allowed between irradiation and conception, presumably caused by repair.

   This information is already used in genetic counseling given to people who have received radiation. In the mouse, an interval between irradiation and insemination of 2 months in the male and rather longer in the female is sufficient to produce a maximum reduction in the effect of radiation. Although data are not available for humans, by analogy, a period of 6 months usually is recommended. Consequently, if people are exposed to significant doses of radiation, either accidentally or because of their occupation, it is recommended that 6 months be allowed to elapse between the exposure to the radiation and a planned conception, to minimize the genetic consequence. This would be good

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**FIGURE 11.2** In the megamouse project, seven specific locus mutations were used to study radiation-induced hereditary effects. This photo shows three of the mutations, which involve changes of coat color. (Courtesy of Dr. William L. Russell, Oak Ridge National Laboratory.)
advice to a person accidentally exposed to, say 0.1 Gy, to young patients with Hodgkin disease receiving radiotherapy, or even to patients subjected to diagnostic x-ray procedures involving the lumbar spine or the lower gastrointestinal tract in which a large exposure is used and the gonads must be included within the radiation field.

5. The estimate of the doubling dose favored by the Committee on the Biological Effects of Ionizing Radiation (BEIR V) and the UNSCEAR 1988 is 1 Gy, based on low-dose-rate exposure. This is a calculated rather than a measured quantity, based on the measured mutation rate per locus in the mouse adjusted for the estimated comparable number of loci in the human. It also allows for the fact that the human is usually exposed at low dose rate, whereas the mouse data reflect acute exposures.

### Radiation-Induced Heritable Effects in Humans

To estimate the risk of heritable effects in the human population caused by exposure to radiation, two basic pieces of data are needed: first, the baseline spontaneous mutation rate, which is known for humans, and second, the doubling dose, which can only come from mouse experiments (1 Gy). Two correction factors must then be applied, which were derived by a task force of the International Commission on Radiological Protection (ICRP) in 2000. The first must allow for the fact that not all mutations lead to a disease. This so-called mutation component (MC) varies for different classes of heritable diseases; it is simple for autosomal recessive diseases and very complex for multifactorial diseases. For the first generation following radiation exposure, the MC is about 0.3 for autosomal dominant and X-linked diseases, close to zero for autosomal recessive diseases, and about 0.01 to 0.02 for chronic multifactorial diseases. The second correction factor must allow for the fact that the seven specific locus mutations used to assess the doubling dose in the megamouse project are not representative of the spectrum of inducible heritable diseases in humans. The success of experimental radiation mutagenesis studies in the mouse is mainly caused by the fortunate choice of genes that are nonessential for survival of the animal or the cell and also happen to be located in nonessential regions of the genome. Most human disease-causing genes are not of this type. Among the human autosomal and X-linked genes studied, only 15% to 30% may be

![Figure 11.3](image-url) Mutations in mice as a function of dose, delivered at high and low dose rates. (Courtesy of Dr. William L. Russell, Oak Ridge National Laboratory.)
responsive to induced mutations that are potentially recoverable in live births. For chronic multifactorial diseases, which involve several genes, the fraction recoverable in live births would be even lower.

The UNSCEAR 2001 estimates of hereditary risks for the first generation and first two generations of an irradiated population are listed in Table 11.4. The risk of autosomal dominant and X-linked diseases for the first generation after irradiation is about 750 to 1,500 cases per million progeny per gray of chronic low linear energy transfer (LET) radiation (compared with the baseline of 16,500 cases per million). The risk of autosomal recessive diseases is essentially zero (compared with the baseline of 7,500 per million). The risk of chronic diseases is about 250 to 1,200 cases per million (compared with the baseline of 650,000 per million). The risk of multisystem developmental or congenital abnormalities may be about 2,000 cases per million. Note that the total risk per gray is only about 0.41% to 0.64% of the baseline risk of 738,000 per million live births— that is, a relatively small proportion.

### INTERNATIONAL COMMISSION ON RADIOLOGICAL PROTECTION

**ESTIMATES OF HEREDITARY RISKS**

The strategy of ICRP to estimate hereditary risks is based on the data for the first two generations calculated by UNSCEAR 2001, shown in Table 11.4. These data refer to a “reproductive” population and apply when the radiation doses received by all people in the population are genetically significant. However, when the total population of all ages is considered, the genetically significant dose will be markedly lower than the total dose received over a lifetime. Genetic damage sustained by germ cells of persons who are beyond the reproductive period or who are not procreating for any reason poses no heritable risks. Assuming an average life expectancy of 75 years, with mean reproductive age stopping at 30 years, the risk coefficients for

<table>
<thead>
<tr>
<th>TABLE 11.4</th>
<th>Current Estimates of Genetic Risks from Continuing Exposure to Low-LET, Low-Dose, or Chronic Irradiation (from UNSCEAR 2001) (Assumed Doubling Dose: 1 Gy)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Disease Class</strong></td>
<td><strong>Based Frequency per 10^6 Live Births</strong></td>
</tr>
<tr>
<td>Mendelian</td>
<td></td>
</tr>
<tr>
<td>Autosomal dominant and X-linked</td>
<td>16,500</td>
</tr>
<tr>
<td>Autosomal recessive</td>
<td>7,500</td>
</tr>
<tr>
<td>Chromosomal</td>
<td>4,000</td>
</tr>
<tr>
<td>Multifactorial</td>
<td></td>
</tr>
<tr>
<td>Chronic</td>
<td>650,000</td>
</tr>
<tr>
<td>Congenital abnormalities</td>
<td>60,000</td>
</tr>
<tr>
<td>Total</td>
<td>738,000</td>
</tr>
<tr>
<td>Total risk per Gy expressed as percent of baseline</td>
<td>—</td>
</tr>
</tbody>
</table>

*Assumed to be subsumed in part under the risk of autosomal dominant and X-linked diseases and in part under congenital abnormalities.*

a total population of all ages will be 30/75, that is, 40% of that for a reproductive population. This rounds out at 0.2% per sievert. For a working population, the relevant age range is only from 18 to 30 years because of the fact that no one is allowed to be a radiation worker before the age of 18 and the reproductive cycle is assumed to end at the age of 30; consequently, the risk coefficients will be 12/75, or 16% of that for a reproductive population, which rounds out at 0.1% per sievert. These figures are summarized in Table 11.5.

**MUTATIONS IN THE CHILDREN OF THE A-BOMB SURVIVORS**

The survivors of the atomic bomb attacks on Hiroshima and Nagasaki constitute the largest irradiated human population studied carefully for heritable effects. Over the years, a cohort of 31,150 children born to parents receiving significant amounts of radiation (within 2 km of the bomb’s hypocenter) and a somewhat larger control cohort (41,066) have been studied with respect to various indicators: first, in the early years, for congenital defects, gender of child, physical development, and survival; then, in the middle years, for cytogenetic abnormality; and, more recently, for the occurrence of malignant disease and for electrophoretic or functional defects in a battery of some 30 serum proteins or erythrocyte enzymes. None of these indicators was related significantly to parental radiation exposure, but the net regression was slightly positive.

For these various measures of heritable effects, the differences between the children of proximally and distally exposed survivors is in the direction expected if a heritable effect did result from the radiation, but in fact, none of the findings are statistically significant. Nevertheless, the differences between these groups of children were used to calculate doubling doses.

Only three of these indicators lend themselves to an estimate of doubling dose, and the results are shown in Table 11.6, taken from a study in the early 1980s. The simple average of the three estimates is 1.56 Sv. In a more recent review paper, Neel estimated the doubling dose for the human to be about 2 Sv; this is subject to considerable uncertainties, with a lower limit of 1 Sv and an upper limit that is indeterminate. This, of course, refers to an acute radiation dose because it is based on the Japanese survivors.

The sparse human data support the current impression that the doubling dose derived from the mouse specific-locus experiments is too low. The lack of a statistically significant excess of heritable effects in the children of the A-bomb survivors is consistent with the numeric risk estimates developed by ICRP. The number of children involved and the range of doses to which their parents were exposed are too small for a statistically significant heritable effect to be expected.

**CHANGING CONCERNS FOR RISKS**

In 1956, the ICRP reduced the dose limit for radiation workers to, effectively, what it is today—a maximum of 50 mSv per year. This was based
entirely on heritable effects in the fruit fly, Drosophila. In the half century or so that has elapsed since that time, the level of concern involving genetic effects, or heritable effects as we call them, has declined steadily—first, because of the availability of mouse data, and more recently, with a reassessment of the importance of multifactorial diseases and doubt about the relevance of the specific locus mutations in mice. As a consequence, the percentage of radiation detriment attributed to the genetic component in the view of ICRP has declined from 100% in 1955, to 25% in 1977, to 18% in 1991, and to only 4% in 2007. In the meantime, the level of concern involving radiation carcinogenesis has increased as more and more solid tumors have appeared in the Japanese A-bomb survivors. This trend is illustrated in Figure 11.4. In the 1950s, genetic effects were considered to be the most important because solid tumors had not then appeared in large numbers in the A-bomb survivors. Over the years, concern has switched entirely so that at the present time, radiation carcinogenesis is considered to be by far the most important consequence of low doses of radiation. Meanwhile, radiation protection standards have changed little.

### EPIGENETICS

In this chapter so far, it has been tacitly assumed that a transgenerational heritable effect must, of necessity, involve a change in the DNA sequence. The emerging study of epigenetics implies that this is not necessarily the case.

The Greek prefix *epi*- in epigenetics implies features that are “on top of” or “in addition to” genetics; the term is now generally used to refer to changes in gene expression that takes place without a change in the DNA sequence. Epigenetic changes can result from various molecular modifications, but the most extensively studied and best understood are as follows:

1. DNA methylation, which takes place at the carbon-5 position of cytosine in CpG dinucleotides, and,
2. Changes to chromatin packaging of DNA by posttranslational histone modifications.

Both of these mechanisms can have a profound effect on gene expression, including the silencing of the gene. Other epigenetic mechanisms that are less well understood include regulation of gene expression by noncoding RNAs, including micro RNAs, and mechanisms that control the higher level organization of chromatin within the nucleus.

There is increasing evidence from animal studies that prenatal and early postnatal environmental factors can result in altered epigenetic programming and subsequent changes in the risk of developing disease later in life. Environmental factors studied to date include nutritional supplements, xenobiotic chemicals and, interestingly enough, exposure to ionizing radiation. The radiation studies showed that exposure of adult male or female mice led to transgenerational genome instability in the offspring, resulting from a significant loss of DNA methylation in somatic tissue. In addition, there is some evidence from animal studies that epigenetic alterations may be inherited transgenerationally, thereby affecting the health of future generations.
Imprinted Genes

Most autosomal genes are expressed from the alleles of both parents; however, about 1% of autosomal genes in humans are “imprinted” (i.e., expression is from only one parental allele with the other allele silenced). This leads to a nonmendelian germ line inherited form of gene regulation that involves heritable DNA methylation and histone modification. In addition, expression of the single functioning allele is parent-of-origin-dependent. Consequently, expression of an imprinted gene in the present generation depends on the environment that it experienced in the previous generation. Several human syndromes, and even some cancers, result from genetic and epigenetic modifications at imprinted loci.

An interesting example in the human involves children exposed prenatally to the Dutch famine at the end of World War II that now have an increased incidence of metabolic disorders, cardiovascular disease, and mental disorders in adulthood. They also show significant alterations in DNA methylation at both imprinted and nonimprinted loci, decades after birth.

The study of radiation on epigenetics is in its infancy, but it is a factor that may influence the perception of radiation-induced heritable effects in the future. Mutations in DNA are no longer the whole story.

SUMMARY OF PERTINENT CONCLUSIONS

- In the male, doses as low as 0.15 Gy result in oligospermia (diminished sperm count) after a latent period of about 6 weeks. Doses greater than 0.5 Gy result in azoospermia (absence of living spermatozoa) and therefore temporary sterility. Recovery time depends on dose.
- Permanent sterility in the male requires a single dose in excess of 6 Gy.
- In the male, fractionated doses cause more gonadal damage than a single dose. Permanent sterility can result from a dose of 2.5 to 3 Gy in a fractionated regime over 2 to 4 weeks.
- In the female, radiation is highly effective in inducing permanent ovarian failure, with a marked age dependence on the dose required.
- The dose required for permanent sterility in the female varies from 12 Gy prepubertal to 2 Gy premenopausal.
- The induction of sterility in males does not produce significant changes in hormone balance, libido, or physical capability, but in the female, it leads to pronounced hormonal changes comparable to natural menopause.
- Exposure of a population can cause adverse health effects in the descendants because of mutations induced in germ cells. These used to be called “genetic” effects but are now more often called “heritable” effects.
- Heritable diseases are classified into three principal categories: mendelian, chromosomal, and multifactorial.
- Radiation does not produce new, unique mutations but increases the incidence of the same mutations that occur spontaneously.
- Information on the heritable effects of radiation comes almost entirely from animal experiments.
- The earlier mutation experiments were carried out with the fruit fly, D. melanogaster.
- Relative mutation rates have been measured in the megamouse project by observing seven specific locus mutations. This leads to an estimate of the “doubling dose.”
- The doubling dose is the dose required to double the spontaneous mutation incidence; put another way, it is the dose required to produce an incidence of mutations equal to the spontaneous rate. Based on the mouse data, the doubling dose for low dose-rate exposure in the human is estimated to be 1 Gy.
- Not more than 1% to 6% of spontaneous mutations in humans may be ascribed to background radiation.
- To estimate the risk of radiation-induced heritable diseases in the human, two quantities are required: (1) the baseline mutation rate for humans, which is estimated to be 738,000 per million, and (2) the doubling dose from the mouse data, which is about 1 Gy.
- Two correction factors are needed: (1) to allow that not all mutations lead to a disease—this is MC, which varies for different classes of heritable diseases; and (2) to allow for the fact that the seven specific locus mutations used in the mouse
experiments are not representative of inducible heritable diseases in the human because they are all nonessential for the survival of the animal or cell.

- The ICRP estimates that the heritable risk of radiation is about 0.2% per sievert for the general population and about 0.1% per sievert for a working population.

- In terms of detriment, expressed in years of life lost or impaired, congenital anomalies (i.e., resulting from effects on the developing embryo and fetus) are much more important than heritable disorders.

- Children of the atomic bomb survivors have been studied for several indicators, including congenital defects, gender ratio, physical development, survival, cytogenetic abnormalities, malignant disease, and electrophoretic variants of blood proteins. A recent paper estimated the doubling dose to be about 2 Sv, with a lower limit of 1 Sv and an upper limit that is indeterminate because the increase in mutations is not statistically significant.

- Since the 1950s, concern for the heritable effects of radiation has declined continuously and has been replaced by carcinogenesis as the principal effect of low doses of radiation.

- Epigenetics refers to changes in gene expression that take place without a change in the DNA sequence. The most studied mechanisms include DNA methylation and changes in chromatin packaging by posttranslational histone modification.

- There is evidence from animal studies that prenatal and early postnatal environmental factors, including exposure to radiation, can alter epigenetic programming with subsequent changes in the risk of developing disease in later life.

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HISTORICAL PERSPECTIVE

In the early years of the 20th century, case reports began to appear in the medical literature that described mental retardation in children with small head size, as well as other gross malformations, born to mothers who had received pelvic radiotherapy before realizing that they were pregnant. As early as 1929, Goldstein and Murphy reviewed 38 such cases. Interestingly enough, they concluded that large doses were needed to produce such effects and did not consider diagnostic pelvic irradiation of the mother to be a hazard.

We now have a great deal of information concerning the effects of radiation on the developing embryo and fetus from both animal experiments and the human experience.

OVERVIEW OF RADIATION EFFECTS ON THE EMBRYO AND FETUS

Among the somatic effects of radiation other than cancer, developmental effects on the unborn child are of greatest concern. The classic effects are listed as follows:

1. **Lethal effects** are induced by radiation before or immediately after implantation of the embryo into the uterine wall or are induced after increasingly higher doses during all stages of intrauterine development, to be expressed either before birth (prenatal death) or at about the time of birth (neonatal death).

2. **Malformations** are characteristic of the period of major organogenesis in which the main body structures are formed, and especially of the most active phase of cell multiplication in the relevant structures.

3. **Growth disturbances and growth retardation, without malformations** are induced at all stages of development but particularly in the latter part of pregnancy.

The principal factors of importance are the **dose** and the **stage of gestation at which it is delivered**. **Dose rate** is also of significance because many pathologic effects on the embryo are reduced significantly by reducing the dose rate.

It should be recognized that congenital anomalies arise in all animal species, even in the absence of any radiation beyond that received from natural sources. The incidence depends largely on the time at which the anomalies are scored. In humans, the average incidence of malformed infants at birth is about 6%. Some malformations disappear after birth, but more become evident later that are not scored at birth. The global incidence roughly doubles to 12% if grown children rather than infants are examined. Any assessment of the effectiveness of radiation in inducing damage in utero must be viewed against this natural level of inborn defects and its variable expression.
DATA FROM MICE AND RATS

Most experimental data on the effect of radiation on the developing embryo or fetus have been obtained with the mouse or rat, animals that reproduce in quantity with relatively short gestation periods. Russell and Russell divided the total developmental period in utero into three stages: (1) preimplantation, which extends from fertilization to the time at which the embryo attaches to the wall of the uterus; (2) organogenesis, the period during which the major organs are developed; and (3) the fetal period, during which growth of the structures already formed takes place. There is a very large variability in the relative duration of these periods among animal species, as well as in the total duration of intrauterine life. In addition, at any given stage of development, the state of differentiation or maturation of any one structure, with respect to all the others, varies considerably in different species.

In the mouse, preimplantation corresponds to days 0 through 5; organogenesis, days 5 through 13; and the fetal period, day 13 through full term, which is about 20 days. The effect of about 2 Gy of x-rays delivered at various times after conception is illustrated in Figure 12.1. The lower scale contains Rugh’s estimates of the equivalent stages for the human embryo, based solely on comparable stages of organ development. It is a nonlinear match because preimplantation, organogenesis, and the fetal period in the mouse are about equal in length, whereas the fetal period in the human is proportionately much longer.

Figure 12.2 is taken from the work of Brent and Ghorson, who performed an extensive series of experiments with rats. It shows the various periods during gestation in which the principal effects of radiation are most evident. The horizontal scale refers to the times of the major events during gestation for the rat and gives an estimate of the comparable stages for the human. The following discussion of the principal effects of radiation delivered during preimplantation, organogenesis, and the fetal stages represents a consensus view, combining conclusions from various experiments with either rats or mice.

Preimplantation

Preimplantation is the most sensitive stage to the lethal effects of radiation. The high incidence of prenatal death may be expressed in a decrease in litter size. Growth retardation is not observed after irradiation at this stage; if the embryo survives, it grows normally in utero and afterward. Few, if any, abnormalities are produced by irradiation at this stage. The International Commission on Radiological Protection (ICRP) Publication 103 confirms embryonic susceptibility to the lethal effects of irradiation in the preimplantation period of embryonic development, but suggests that at doses less than 100 milligrays (mGy), such lethal effects will be very infrequent in the human.

FIGURE 12.1 Incidence of abnormalities and of prenatal and neonatal death in mice given a dose of about 2 Gy of x-rays at various times after fertilization. The lower scale consists of Rugh’s estimates of the equivalent stages for the human embryo. (Data from Russell LB, Russell WL. An analysis of the changing radiation response of the developing mouse embryo. J Cell Physiol. 1954;43[suppl 1]:130–149.)
For Students of Diagnostic Radiology, Nuclear Medicine, and Radiation Oncology

During organogenesis, most of the embryonic cells are in their blastula or differentiating stage and are particularly sensitive. This corresponds to the period in the human at which the tranquilizer thalidomide produced such disastrous effects (i.e., at about 35 days after conception), and it is also the time of maximum risk of deleterious effects from the rubella virus.

Examples of gross anomalies resulting from irradiation during the period of organogenesis are shown in Figures 12.4 and 12.5. It is characteristic of mice and rats that various structural malformations are seen. The production of a specific defect is associated with a definite time during this period of organogenesis, usually the time of the first morphologic evidence of differentiation in the organ or portion of the organ involved. On the basis of animal experiments, there appears to be a dose threshold of about 100 mGy for the induction of malformations during organogenesis, so that for practical purposes, the risks of
malformations in the human, for doses well below 100 mGy, would not be expected (ICRP 103).

Embryos exposed during early organogenesis also exhibit the greatest intrauterine growth retardation. This is expressed as a weight reduction at term and is a phenomenon resulting from cell depletion. Animals show a remarkable ability to recover from the growth retardation produced by irradiation during organogenesis, and although they may be smaller than usual at birth, they may achieve a normal weight as adults. There is an association between growth retardation and teratogenesis: Irradiated embryos that show major congenital anomalies also suffer an overall reduction of growth. In animals, a dose of about 1 Gy of x-rays produces growth retardation if delivered at any stage of gestation (except during preimplantation); 0.25 Gy does not produce an observable effect, even at the most sensitive stage.

If death occurs because of irradiation in organogenesis, it is likely to be neonatal death—occurring at or about the time of birth. The transition from prenatal death from irradiation mainly during preimplantation to neonatal death resulting from irradiation mainly in organogenesis is very clear from Figure 12.1. The neonatal deaths peak at 70% for mice receiving about 2 Gy on about the 10th day.

**FIGURE 12.3** During preimplantation, the embryo consists of a limited number of cells. A: Newly fertilized mouse egg. B: By the 3rd day, the mouse embryo consists of only 16 cells. About 5 days after conception in the mouse, which corresponds to about 9 or 10 days in the human, the embryo becomes embedded in the wall of the uterus, and at about this time, cells begin to differentiate to form specific tissues and organs. (Courtesy of Dr. Pedersen, University of California at San Francisco.)
The Fetal Period
The remainder of pregnancy, the fetal period, extends from about day 13 onward in the mouse; this corresponds to about 6 weeks onward in the human. Various effects have been documented in the experimental animal after irradiation during the fetal stages, including effects on the hematopoietic system, liver, and kidney, all occurring, however, after relatively high radiation doses. The effects on the developing gonads have been documented particularly well, both morphologically and functionally. There appears at present to be little correspondence between the cellular and functional damage as a function of dose, but doses of a few tenths of a gray as a minimum are necessary to produce fertility changes in various

FIGURE 12.4  Litter from a female mouse irradiated with x-rays during organogenesis and sacrificed at 19 days. Several anomalies are demonstrated in this litter. There are four resorbed embryos (bottom) and five fetuses that would have been born alive (top). From left to right, the first shows exencephaly; the second, exencephaly and evisceration; the third is apparently normal; and the remaining two are anencephalics with stunting. (Photograph by Dr. Roberts Rugh.)

FIGURE 12.5  Two rats from the same litter exposed to a dose of about 1 Gy of x-rays 9.5 days after conception. The rat on the left has a normal right eye and microphthalmus of the left. The rat on the right shows anophthalmia of both eyes. (From Rugh R, Caveness WF, Duhamel L, Schwarz GS. Structural and functional [electroencephalographic] changes in the post-natal mammalian brain resulting from x-irradiation of the embryo. Mil Med. 1963;128:392–408, with permission.)
animal species. The effects of radiation on humans were discussed in more detail in Chapter 11.

Much higher doses of radiation are required to cause lethality during this period than at earlier stages of development, although the irradiated early fetus exhibits the largest degree of permanent growth retardation, in contrast to the embryo in early organogenesis, which exhibits the most temporary growth retardation, which is evident at term but from which the animal is able to recover later.

**EXPERIENCE IN HUMANS**

Information on the irradiation of human concepti comes from two major sources: studies of atomic bomb survivors in Japan and medical exposures (particularly therapeutic irradiations), especially during the early part of the century, when hazards were not yet fully appreciated. These will be discussed in turn.

**Survivors of the A-Bomb Attacks on Hiroshima and Nagasaki Irradiated In Utero**

The growth to maturity of children exposed in utero at Hiroshima and Nagasaki has been studied carefully. There are difficulties associated with the dosimetry, but the conclusions have far-reaching implications.

Data on the children exposed in utero in Hiroshima and Nagasaki show too few individuals who were younger than 4 weeks of gestational age at the time the bomb was dropped. This deficiency presumably results from increased fetal loss or infant mortality rate. This stage of development is so early that damage to a single cell or group of cells is likely to impair the function of all the progeny cells and leads to death of the embryo. In accord with this reasoning is the observation that no birth defects were found because of irradiation before 15 days of gestational age. This is in accord with the experimental data for rats and mice in which exposure during preimplantation had an all-or-nothing effect: death of the embryo or normal development.

Exposure to radiation resulted in growth retardation (Table 12.1). Children exposed as embryos closer than 1,500 m to the hypocenter of the atomic explosion were shorter, weighed less, and had head diameters significantly smaller than children who were more than 3,000 m from the hypocenter and received negligibly small doses. It is of interest to note that there was no catch-up growth because the smallness in head size was maintained into adulthood.

**TABLE 12.1 Growth Retardation at Hiroshima from In Utero Irradiation: Comparison of Those Exposed within 1,500 m of the Hypocenter with Those More than 3,000 m from the Hypocenter**

<p>| | | |</p>
<table>
<thead>
<tr>
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<tbody>
<tr>
<td>Height</td>
<td>2.25 cm shorter</td>
<td></td>
</tr>
<tr>
<td>Weight</td>
<td>3 kg lighter</td>
<td></td>
</tr>
<tr>
<td>Head diameter</td>
<td>1.1 cm smaller</td>
<td></td>
</tr>
</tbody>
</table>

80% of 1,613 children exposed in utero followed to age 17 years.

Average kerma, 0.25 Gy.


1. **Microcephaly.** A significant effect of radiation on the frequency of small heads is observed only in the periods 0 to 7 and 8 to 15 weeks postovulation. No significant excess was seen among persons exposed at 16 weeks or more. The proportion of exposed persons with microcephaly increases with dose, and there is little evidence for a threshold in dose (Fig. 12.7). As pointed out earlier, radiation-related small head size is related to a generalized growth retardation, including reduced height and weight.
and absorbed dose for this most sensitive period. The relationship appears to be linear, and the data are consistent with a probability of occurrence of mental retardation of 40% at a dose of 1 Gy. The data are clearly consistent with a threshold in dose because the incidence of severe mental retardation at the two lower doses in Figure 12.8 is not statistically significant. A dose threshold is also consistent with the presumed deterministic nature of mental retardation that would require the killing of a minimum number of cells to be manifest. ICRP Publication 90 (2003) offers the view that the induction of severe mental retardation during this most sensitive period has a threshold of at least 0.3 Gy, with the absence of risk at lower doses.

MICROCEPHALY AND MENTAL RETARDATION DIFFER SOMEWHAT IN THE GESTATIONAL AGES AT WHICH THEY OCCUR. BOTH OCCUR IN THE 8- TO 15-WEEK PERIOD, BUT IN THE MOHAMAD PERIOD (0-7 WEEKS), MICROCEPHALY APPEARS WITHOUT MENTAL RETARDATION, WHEREAS IN

2. **Mental retardation.** A child was deemed to be severely retarded if he or she was “unable to perform simple calculations, to make simple conversation, to care for himself or herself, or if he or she was completely unmanageable or has been institutionalized.” Most of these children were never enrolled in public schools, but among the few who were, the highest IQ was 68. In all, 30 children were judged to be severely mentally retarded; in 5 of these children, causes other than radiation were considered likely, including Down’s syndrome, neonatal jaundice, encephalitis, or birth trauma. Nevertheless, the remaining number represents an incidence far higher than normal. Severe mental retardation was not observed to be induced by radiation before 8 weeks after conception or after 25 weeks. The most sensitive period is 8 to 15 weeks after conception; for exposure during weeks 16 to 25, the risk is four times smaller. Figure 12.8 shows the relation between the incidence of mental retardation and gestational age.
the later period (16–25 weeks), mental retardation is observed in the absence of microcephaly. The highest risk of mental retardation occurs at a gestational age at which the relevant tissue, that is, the brain cortex, is being formed. It is thought to be associated with impaired proliferation, differentiation, and, most of all, migration of cells from their place of birth to their site of function. Cells killed before 8 weeks of gestation can cause small head size without mental retardation because the neurons that lead to the formation of the cerebrum are at a stage not yet sensitive to impairment by radiation. Glial cells that provide structural support for the brain are, however, susceptible to depletion. Magnetic resonance images of persons irradiated in utero at 8 to 15 weeks of gestation show evidence of massive impairment of cells to migrate from proliferative zones. A typical distribution of gray matter is often seen in patients with spontaneous mental retardation, but it is usually unilateral; that caused by radiation exposure is bilateral.

Although severe mental retardation requiring the children to be institutionalized has been known for many years in those exposed in utero at Hiroshima, later studies have shown mental impairment of less severity, indicated by IQ test scores. For irradiation during the sensitive period of 8 to 15 weeks after conception, the observed shift in intelligence test scores corresponds to about 25 IQ points per Gy. ICRP regards these data as more difficult to interpret than the severe mental retardation data, and the possibility of a nonthreshold response cannot be ruled out. However, even in the absence of a true dose threshold, any effects on IQ following in utero doses less than about 0.1 Gy would be of no practical significance (ICRP Publication No. 103, 2003).

Exposure to Medical Radiation

A relationship between microcephaly and x-irradiation during intrauterine life has been recognized since Goldstein and Murphy first focused attention on the subject in 1929. The numbers are small and the doses are not known with any certainty, although most were in the therapeutic range. Microcephaly was reported as well as mental retardation and various defects, including spina bifida, bilateral clubfoot, ossification defects of the cranial bones, deformities of the upper extremities, hydrocephaly, alopecia of the scalp, divergent squint, and blindness at birth.

Dekaban surveyed the literature for instances of pelvic x-irradiation in pregnant women. Based on the available data, the following generalizations were proposed:

1. Large doses of radiation (2.5 Gy) delivered to the human embryo before 2 to 3 weeks of gestation are not likely to produce severe abnormalities in most children born, although a considerable number of the embryos may be resorbed or aborted.
2. Irradiation between 4 and 11 weeks of gestation would lead to severe abnormalities of many organs in most children.
3. Irradiation between 11 and 16 weeks of gestation may produce a few eye, skeletal, and genital organ abnormalities; stunted growth, microcephaly, and mental retardation are frequently present.

| Figure 12.8 | The frequency of mental retardation as a function of dose among those exposed in utero to atomic bomb radiation. The data are pooled from Hiroshima and Nagasaki for those exposed at 8 to 15 weeks gestational age. The vertical bars represent the 90% confidence intervals. There was no risk at 0 to 8 weeks after conception, and for exposure at later periods during gestation (16+ weeks), the excess is barely significant even at the higher doses. (Adapted from Otake M, Schull WJ. In utero exposure to A-bomb radiation and mental retardation: a reassessment. Br J Radiol. 1984;57:409–414, with permission.) |
4. Irradiation of the fetus between 16 and 25 weeks of gestation may lead to a mild degree of microcephaly, mental retardation, and stunting of growth.

5. Irradiation after 30 weeks of gestation is not likely to produce gross structural abnormalities leading to a serious handicap in early life but could cause functional disabilities.

**COMPARISON OF HUMAN AND ANIMAL DATA**

Figure 12.9 is an attempt to summarize the data for the effects of radiation on the developing embryo and fetus, comparing the information from animals and human A-bomb survivors.

Exposure to radiation during preimplantation leads to a high incidence of embryonic death, but embryos that survive develop normally. This has been shown clearly in experiments with both rats and mice and is consistent with the data from Japan. In animals, irradiation during organogenesis leads to neonatal death, temporary growth retardation, and, above all, a wide range of malformations affecting many different limbs and organs. By contrast, the principal effect in the Japanese survivors of the atomic bomb attacks is microcephaly with or without mental retardation, and this begins in the 8th week, that is, after the period classically described as organogenesis. The wide array of congenital malformations found in rats and mice irradiated during organogenesis (and in humans in medical exposures) was not reported in the Japanese survivors. Much has been made of this difference. On the one hand, it has been suggested that the gross structural deformities in Japan simply were not recorded in the chaos that followed the dropping of the atomic bombs. On the other hand, it is argued that humans differ from rats and mice in that the period of susceptibility to a wide array of congenital malformations (i.e., during organogenesis) is short compared with the period during which mental retardation can be induced. Because the number of children involved is quite small, it might be expected that effects on the central nervous system, which is developing over a larger period, would dominate. The situation is different in laboratory animals: Their susceptibility to radiation-induced small head size is of similar duration to that for the induction of other deformities. The data from patients exposed to therapeutic doses of

![Figure 12.9](image-url)
medical radiation show a range of congenital malformations that more closely mirrors the animal results, although the numbers are small and the doses are high. To be on the safe side, it must be assumed that the entire period of gestation from about 10 days to 25 weeks is sensitive to the induction of malformations by radiation.

Table 12.2 summarizes the lowest doses at which effects on the embryo and fetus have been observed. This table summarizes the conclusions of the third report of the Committee on the Biological Effects of Ionizing Radiation (BEIR III). Readily measurable damage can be observed at doses less than 0.1 Gy delivered at sensitive stages of gestation. The principal abnormalities produced by radiation are almost certainly the consequence of damage to many cells (i.e., deterministic effects) and, therefore, would be consistent with a threshold in dose. There is also some indication of this in the data for mental retardation.

**TABLE 12.3** Childhood Cancer and Irradiation In Utero

| Number of children with leukemia or cancer before age 10 years | 7,649 |
| Number of x-rayed in utero | 1,141 |
| Number of matched controls | 7,649 |
| Number of controls irradiated in utero | 774 |
| Number of films | 1–5 |
| Fetal dose per film | 4.6–2 mGy |
| Relative cancer risk estimate assuming radiation to be the causative agent | 1.52 |

over the period of the study. A subsequent study in New England by MacMahon also reported an association between prenatal x-rays and childhood leukemia.

This subject has been the source of great controversy for many years. No one seriously doubts the association between in utero irradiation and childhood cancer; the debate is whether the radiation is causative or whether it involves the selection of a particular group of children prone to cancer.

In a careful paper in 1997, Doll and Wakeford summarized all of the studies of in utero exposure and came to the following conclusions:

- **Low-dose irradiation of the fetus in utero** causes an increased risk of childhood malignancies. Most of the data refer to exposure in the third trimester.
- An obstetric x-ray examination results in a 40% increase in the risk of childhood cancer over the spontaneous level.
- Radiation doses of around 10 mGy increase the risk.
- The excess absolute risk is about 6% per Gy.
- These risk estimates are highly uncertain except to say that they are not zero.

In a later (2003) paper, Wakeford and Little compared the risk of childhood cancer per unit dose of radiation received in utero for the obstetric x-ray examinations and the Japanese A-bomb survivors. They concluded that once all sources of uncertainty are taken into account, the risks are not inconsistent, which supports a causal explanation for the cancers seen in the Oxford survey.

**OCCUPATIONAL EXPOSURE OF WOMEN**

The National Council on Radiological Protection and Measurements in its NCRP Report 116 recommends a monthly limit of 0.5 mSv to the embryo or fetus once pregnancy is declared. This recommendation is designed to limit the risk of mental retardation, congenital malformations, and carcinogenesis. NCRP no longer recommends specific controls for occupationally exposed women until a pregnancy is declared. Once a pregnancy is declared, the radiation worker should be interviewed by the radiation safety officer or the chair of the radiation safety committee to discuss the advisability of changing or curtailing duties to limit exposure.

**THE PREGNANT OR POTENTIALLY PREGNANT PATIENT**

Most practicing radiologists at some time in their careers are faced with a patient who has discovered in retrospect that she was pregnant at a time when extensive x-ray procedures were performed involving the pelvis or lower abdomen.

The only completely satisfactory solution to this problem is to ensure that the situation never occurs in the first place. Patients should always be asked if they are, or may be, pregnant, and in the case of procedures involving larger doses of radiation to the pelvis, a pregnancy test may be in order.

Despite the best-laid plans and the most careful precautions, there still are occasional instances in which, because of clinical urgency or unusual accident, an early developing embryo is exposed to a substantial dose of radiation amounting to several tens of millisieverts or more. The first step in evaluating whether or not damage may have been done to the embryo is to estimate the dose involved. It is sometimes useful to solicit the help of an experienced medical physicist to make measurements in a phantom after carefully reconstructing the setup that was used. No dose level can be regarded as completely safe. Congenital abnormalities occur in 5% to 10% of the human population anyway, so it is impossible in retrospect to attribute a given anomaly to a small dose of radiation received by an embryo or fetus. All that can be said is that radiation increases the probability of an anomaly and that this increase is a function of dose.

Doses less than about 100 mSv during pre-implantation and organogenesis, and perhaps 200 mSv during the fetal period pose a low risk of deleterious effects, except the very small possibility of carcinogenesis, which is difficult to quantify. Much higher doses than this to the developing embryo or fetus may be justification for considering a therapeutic abortion. Not everyone would agree with this view, but if the dose involved is sufficiently large, it may be prudent to consider the relative merits of terminating the pregnancy in consultation with the referring physician as well as with the patient and her family.
The effects depend on the stage of gestation, the dose, and the dose rate. Gestation is divided into preimplantation, organogenesis, and the fetal period. In humans, these periods correspond to about 0 through 9 days, 10 days through 6 weeks, and 6 weeks through term, respectively.

The principal effects of radiation on the developing embryo and fetus, aside from cancer, are embryonic, fetal, or neonatal death; congenital malformations; growth retardation; and functional impairment, such as mental retardation.

Irradiation during preimplantation leads to potential death of the embryo. At doses less than 100 mGy, such lethal effects will be infrequent in humans. Growth retardation or malformations are not seen in animals from irradiation at this time. The human data are consistent with this conclusion.

In animals, lethality from irradiation varies with stage of development. The embryonic lethality from irradiation varies with stage of development. The embryonic lethality from irradiation varies with stage of development. The embryonic lethality from irradiation varies with stage of development. The embryonic lethality from irradiation varies with stage of development. The embryonic lethality from irradiation varies with stage of development. The embryonic lethality from irradiation varies with stage of development.

There are several factors to consider in conjunction with the dose. These include the hazard of the pregnancy to the expectant mother, the probability of future pregnancies, the extent to which the prospective parents want the unborn infant, their mental outlook on the possibility of a deformed child, and the ethnic and religious background of the family. The exact dose level at which it is justifiable to terminate the pregnancy may be flexible within broad limits around the guideline figure, depending on a combination of these other circumstances.

There are special problems involved in the use of nuclear medicine procedures in pregnant or potentially pregnant females. This is particularly true in the case of radionuclides that are able to cross the placenta. This topic is discussed in Chapter 16.

Table 12.4 is a historical summary of events in our gradual understanding of radiation effects on the developing embryo and fetus.

### SUMMARY OF PERTINENT CONCLUSIONS

- Doses that have little effect on adults can produce catastrophic effects on the developing embryo and fetus.
- The effects depend on the stage of gestation, the dose, and the dose rate.
- Irradiation during preimplantation leads to potential death of the embryo. At doses less than 100 mGy, such lethal effects will be infrequent in humans. Growth retardation or malformations are not seen in animals from irradiation at this time. The human data are consistent with this conclusion.
- In animals, lethality from irradiation varies with stage of development. The embryonic lethality from irradiation varies with stage of development. The embryonic lethality from irradiation varies with stage of development.

### TABLE 12.4 Major Events in Understanding Effects of Radiation on the Developing Embryo and Fetus

<table>
<thead>
<tr>
<th>Investigators</th>
<th>Year</th>
<th>Observations</th>
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</thead>
<tbody>
<tr>
<td>Goldstein and Murphy</td>
<td>1929</td>
<td>Various abnormalities, including mental retardation and small head diameter in children born to mothers who received pelvic radiation therapy during pregnancy</td>
</tr>
<tr>
<td>Job et al.</td>
<td>1935</td>
<td>Recognition that different periods of gestation differ in radiosensitivity</td>
</tr>
<tr>
<td>Russell</td>
<td>1950</td>
<td>Nature of developmental abnormality determined by gestational age at exposure</td>
</tr>
<tr>
<td>Russell and Russell</td>
<td>1952</td>
<td>Clinical implications of irradiation in pregnancy</td>
</tr>
<tr>
<td>Plummer</td>
<td>1952</td>
<td>Mental retardation and microcephaly observed in children of atomic bomb survivors</td>
</tr>
<tr>
<td>Stewart and Kneale</td>
<td>1952</td>
<td>Leukemia and childhood cancer in children irradiated in utero with diagnostic x-rays</td>
</tr>
<tr>
<td>Otake and Schull</td>
<td>1984</td>
<td>Mental retardation caused by irradiation at 8–15 weeks of pregnancy in Japanese survivors</td>
</tr>
</tbody>
</table>
50% lethal dose is lowest during early pre-implantation; at this stage, embryos killed by radiation suffer a prenatal death and are resorbed. In organogenesis, prenatal death is replaced by neonatal death—death at or about the time of birth. During the fetal stage, the 50% lethal dose approaches that of the adult.

- In animals, the peak incidence of teratogenesis, or gross malformations, occurs if the fetus is irradiated in organogenesis. For practical purposes, the risk of malformations for doses well below 100 mGy would not be expected.
- In contrast to what is observed in experimental animals, radiation-induced malformations of body structures other than the central nervous system are uncommon in the Japanese survivors irradiated in utero, although they have been reported in patients exposed to therapeutic doses of medical radiation.
- In the Japanese survivors, irradiation in utero resulted in small head size (microcephaly) and mental retardation.
- Mental retardation from irradiation occurred primarily at 8 to 15 weeks of gestational age, with a smaller excess at 16 to 25 weeks. It is thought to be caused by radiation effects on cell migration within the brain.
- The incidence of severe mental retardation as a function of dose is apparently linear at 8 to 15 weeks, with a risk coefficient of 0.4 per Gy. The incidence is about four times lower at 16 to 25 weeks. The data are also consistent with a dose threshold of 0.3 Gy.
- Small head circumference was more common than mental retardation.
- Data on atomic bomb survivors indicate that microcephaly can result from exposure at 0 to 7 and 8 to 15 weeks postovulation, but not at later times. There is little evidence for a threshold in dose.
- Various effects have been documented in experimental animals after irradiation during fetal stages, including effects on the hematopoietic system, liver, and kidney, all occurring, however, after quite high radiation doses.
- There is an association between exposure to diagnostic x-rays in utero and the subsequent development of childhood malignancies, although there are particular uncertainties concerning the risk involved.
- The original study of diagnostic x-ray exposure in utero and subsequent malignancies, principally leukemia, was done by Stewart and Kneale at Oxford University, but the same association was observed in the United States by MacMahon. If x-rays are the causative agent, these studies imply that radiation at low doses in utero increases the spontaneous cancer incidence in the first 10 to 15 years of life by 50%—that is, by a factor of 1.5.
- It has been argued for years whether radiation is the causative agent or whether there are other factors involved, such as the selection of a particular group of children prone to cancer.
- Doll and Wakeford in 1997 summarized all of the evidence for and against and concluded that an obstetric x-ray examination, particularly in the third trimester, increased the risk of childhood cancer by 40%. The risk is increased by a dose of only 10 mGy. The excess absolute risk is about 6% per Gy, which is not very different from the risk estimates from the atomic bomb survivors for adult exposure.
- Until a pregnancy is declared, no special limits apply to women other than those applicable to any radiation worker. Once a pregnancy is declared, the maximum permissible dose to the fetus is 0.5 mSv per month.
- Once a pregnancy is declared, the duties of a radiation worker should be reviewed to ensure that this limit is not exceeded.
- A sufficiently large dose to the embryo or fetus during the sensitive period of gestation (10 days to 25 weeks) may be justification for considering a therapeutic abortion to avoid the possibility of an anomalous child. Not everyone would agree with this, and the decision to terminate a pregnancy should be flexible and must depend on many factors in addition to dose.

**BIBLIOGRAPHY**


Committee for the Compilation of Materials on Damage


CHAPTER 13

Radiation Cataractogenesis

Cataracts of the Ocular Lens

Lens Opacification in Experimental Animals

Cataracts in Humans

The Degree of Opacity

The Latent Period

Dose–Response Relationship for Cataracts in Humans

Summary of Pertinent Conclusions

Bibliography

CATARACTS OF THE OCULAR LENS

The word cataract is used to describe any detectable change in the normally transparent lens of the eye. The effect may vary from tiny flecks in the lens to complete opacification, resulting in total blindness. Cataracts are usually associated with old age or, less commonly, with some abnormal metabolic disorder, chronic ocular infection, or trauma. It is also well known that sufficient exposure to ionizing radiations (such as x- or γ-rays, charged particles, or neutrons) may cause a cataract.

The ocular lens is enclosed in a capsule (Fig. 13.1); the lens itself consists largely of fiber cells and is covered with an epithelium anteriorly. The lens has no blood supply. Dividing cells are limited to the preequatorial region of the epithelium. The progeny of these mitotic cells differentiate into lens fibers and accrete at the equator.

Cell division in the lens continues throughout life, but it is a most curious cellular system in that there is no mechanism for cell removal. If dividing cells are injured by radiation, the resulting abnormal fibers are not removed from the lens but migrate toward the posterior pole; because they are not translucent, they constitute the beginning of a cataract.

LENSES OPACTION IN EXPERIMENTAL ANIMALS

Some species of animals, especially the mouse, are very sensitive to radiation as far as lens opacification is concerned. A large proportion of a mouse population naturally develop opacifications as they become older. A dose of a few tens of milligray (mGy) of x-rays or a fraction of 1 mGy of fast neutrons produces readily discernible changes in the lens. As the dose is increased, the latent period (i.e., the time that elapses before an opacity of given severity is evident) becomes shorter. Put another way, radiation advances in time, a process that occurs normally late in life.

Neutrons and other densely ionizing radiations are very effective at inducing cataracts, as evidenced by several physicists and engineers who developed cataracts as a result of working around high-energy accelerators in the early days before safety procedures were introduced. The relative biologic effectiveness (RBE) of fast neutrons is a strong function of dose, with a value of about 10 pertaining to high-dose levels on the order of several grays relative to x-rays, but rising to 50 or more for small doses of less than 10 mGy. Worgul and his associates have reported similar RBEs for lens damage in rat eyes exposed to accelerated heavy ions. The increase in RBE at low doses is caused largely by the sharply declining effectiveness of x-rays with decreasing dose, rather than an increase in effect per unit dose of neutrons or charged particles.

CATARACTS IN HUMANS

Radiologists have known for many years that the lens of the eye may be damaged by radiation. A study of patients treated with x- or γ-rays in which a proportion of the dose reached the eye has provided some insight into radiation cataractogenesis in humans. Figure 13.2 shows a typical cataract in a patient who had undergone radiotherapy. An early radiation cataract viewed through an ophthalmoscope may appear
person with a radiation history strongly suggests radiation as the causative agent. Similarly, an absence of this sequence of events would exclude radiation as a cause. In other words, although it is never possible to state unequivocally that a given cataract is radiation induced, it is possible to say with some certainty that some cataracts—for example, nuclear cataracts—do not have a radiation etiology. Depending on the dose, the cataract frequently remains stationary at this stage, confined to the posterior subcapsular region. If it continues to progress, it becomes nonspecific and cannot be distinguished from other types of cataracts.

### The Degree of Opacity

Figure 13.3 shows the system of cataract classification devised in the 1950s by Merriam and Focht. The accumulation of some opaque fibers at the posterior pole is labeled a 1+ cataract; as the severity of this opacity increases and some impaired fibers show up in the anterior part of the lens, the score edges up progressively to 4+.

The severity of the cataract can be assessed quantitatively and objectively by using the Scheimpflug imaging system. This device provides a distortion-free digitized image for densitometric analysis of the cataract. Figure 13.4 shows a cross section of the lens of one of the “liquidators” who worked on the roof of the reactor at Chernobyl and accumulated a significant opacity as a dot, usually situated at the posterior pole. As it enlarges, small granules and vacuoles appear around it. With further enlargement, to the point at which the opacity is several millimeters in diameter, it may develop with a relatively clear center, so that it is shaped like a doughnut. At the same time, granular opacities and vacuoles may appear in the anterior subcapsular region, usually in the pupillary area. This sequence of events is not unique to radiation, but its appearance in a

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**Figure 13.1** Diagram of a sagittal section of a human lens, illustrating the various cellular relationships. Cells are produced by mitosis in the germination zone (GZ) of the epithelium. They begin to differentiate into lens fibers at the meridional rows (MR) and accumulate at the equator. Cells in the central zone (CZ) do not normally divide. (Adapted from Merriam GR, Worgul BV. Experimental radiation cataract: its clinical relevance. *Bull NY Acad Med.* 1983;59:372–392, with permission.)

**Figure 13.2** Cataract in the posterior subcapsular region 4 years after a dose of 24 Gy of x-rays to a patient on radiotherapy. (From Merriam GR, Worgul BV. Experimental radiation cataract: its clinical relevance. *Bull NY Acad Med.* 1983;59:372–392, with permission.)

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of between 6.51 to 11.5 Gy, the average latent period was reduced to about 4 years. This and other evidence indicate that the latent period becomes shorter as dose is increased.

### DOSE–RESPONSE RELATIONSHIP FOR CATARACTS IN HUMANS

Both the National Council on Radiation Protection and Measurements (NCRP) and the International Commission on Radiological Protection (ICRP) have categorized a radiation-induced cataract as a “deterministic effect.” That is, a cataract is characterized by the following:

- A threshold in dose below which the effect does not occur.
- The severity of the effect increases with dose above the threshold.
- The effect is thought to be caused by damage to many cells.

The threshold suggested by ICRP is 2 Gy, delivered in a single exposure, or a larger dose (5–8 Gy) for a prolonged or fractionated exposure. These recommendations were based largely on early work by Merriam and Focht and others, which involved the study of a limited number of people (mostly radiotherapy patients) exposed to relatively large radiation doses and followed for relatively short periods. For example, Merriam, Szechter, and Focht reviewed the case histories of 233 patients on radiotherapy who received radiation to the lens of the eye and for whom dose estimates were available. Of these patients, 128 developed cataracts, 105 did not. Britten and his colleagues reported 14 cases of dose. The degree of opacity, measured with the Scheimpflug system, is shown in the lower panel.

#### THE LATENT PERIOD

Time to onset of clinically relevant disabling opacities ultimately requiring cataract extraction surgery is variable, but in general dose related; the latency gets shorter with increasing dose. This latency has been reported in the literature variously to be from 6 months to more than 50 years. In radiotherapy patients who had received 2.5 to 6.5 Gy to the eye, the average latent period was about 8 years. At higher doses

![Figure 13.3](image)  
**FIGURE 13.3** The system of cataract classification devised by Merriam and Focht, illustrating the arbitrary numeric scores assigned to progressive severities of cataracts. (Courtesy of Dr. Basil Worgul.)

![Figure 13.4](image)  
**FIGURE 13.4** Top: Photograph of the lens of a “liquidator” who worked on top of the reactor at Chernobyl and accumulated a substantial radiation dose. Bottom: Degree of opacity through the lens measured with the Scheimpflug imaging system. This equipment gives a quantitative and objective assessment of the severity of the cataract. The area under the curve represents a densitometric reading of the lens. The region of greatest opacification is under the posterior capsule. (Courtesy of Dr. Basil Worgul.)
These and other recent data call into question the classification of an ocular cataract as a “deterministic effect,” because vision-impairing cataracts appear following quite low doses of radiation in persons who are exposed at younger ages and followed for a sufficiently long period. This view is supported by the unexpected appearance of cataracts in astronauts, clean-up workers from Chernobyl, and radiation technologists, all of whom received doses well below the purported threshold of 2 Gy. Historically, radiation-induced cataracts appeared to be deterministic if the persons exposed were mature adults who survived for a relatively short period (e.g., radiotherapy patients in the 1950s). In this scenario, it required a dose of several grays to produce a vision-impairing cataract quickly. It is likely that the standard-setting bodies, such as ICRP and NCRP, will revise their recommendations regarding cataracts in the near future.

**SUMMARY OF PERTINENT CONCLUSIONS**

- A cataract is an opacification of the normally transparent lens of the eye.
- Dividing cells are limited to the preequatorial region of the epithelium. The progeny of these mitotic cells differentiate into lens fibers and accrete at the equator. It is the failure of these cells to differentiate correctly that leads to a cataract, whether spontaneous or radiation induced.
- A unique feature of the lens is that there is no mechanism for the removal of dead or damaged cells.
The latent period between irradiation and the appearance of a lens opacity is dose related. The latency is about 8 years after exposure to a dose in the range of 2.5 to 6.5 Gy.

It is never possible to state unequivocally that a given cataract is radiation induced; however, the appearance of a cataract at the posterior pole of the lens in a person with a radiation history strongly suggests radiation as the causative agent. On the other hand, it is possible to say with some certainty that some cataracts—for example, nuclear cataracts—do not have a radiation etiology.

Neutrons or heavy ions are very effective at producing cataracts. The RBE may be as high as 50 or more for small doses.

There is evidence of early cataracts in astronauts, particularly those exposed to heavy ions.

Both ICRP and NCRP classify a radiation-induced cataract as a deterministic effect, with a threshold of 2 Gy for a single acute exposure and a larger dose (5–8 Gy) for a protracted exposure. This is based on data from adults exposed late in life and followed for a relatively short period.

There is evidence from the A-bomb survivors that much smaller doses of radiation increase the incidence of cataracts that need to be removed surgically if younger persons are exposed and followed until they reach the age at which some cataracts appear anyway in unirradiated persons.
In the years following the 2001 attack on the World Trade Center in New York City, and the resulting tension between the Western world and militant elements of Islam, there has been much talk of the possibility of an attack involving radiation and/or radioactive materials.

Several different scenarios are possible, varying widely in the probability that they will occur and will result in very different consequences. The various possibilities and the consequences are discussed in more detail in the subsequent paragraphs.

- **POSSIBLE SCENARIOS FOR RADIOLOGIC TERRORISM**

1. **The detonation of a nuclear weapon, in or close to a major city.** This is considered “highly unlikely,” but not impossible because several portable “suitcase bombs” are said to be missing, following the breakup of the former Soviet Union. These weapons might perhaps be comparable to the bombs used on Hiroshima and Nagasaki that ended World War II. If such an event were to take place, the consequences would be catastrophic.
   - Individual doses are unlikely to be high enough to cause the ARS.
   - There would be a long-term risk of leukemia and solid cancers in those exposed to external radiation or in those ingesting and/or inhaling radioactive material.
   - There would certainly be chaos and enormous economic loss because cleanup would be a long and slow process.

2. **An attack on a nuclear power station.** It is claimed (although, of course, never tested) that the nuclear reactor itself is so massively built that it would not be destroyed if a jetliner, full of fuel, were deliberately flown into it. What may be more vulnerable are the used fuel elements, often stored in “swimming pool” facilities close to the reactor because, to date, no long-term storage facilities are in widespread use. An attack of this nature is usually considered unlikely, but if it occurred, the huge amount of radioactive material in the used fuel elements (with both long and short half-lives) would spread over the surrounding countryside. The consequences might be the following:
   - Thousands of people would be killed by blast and heat.
   - Hundreds to thousands would be killed or made ill by the acute radiation syndrome (ARS) because of exposure to γ-rays and neutrons.
   - A long-term risk of leukemia and solid cancers in those who survive the acute effects would result principally because of exposure to γ-rays and neutrons and also to fallout.

3. **The detonation of a “dirty bomb,” or to give its official description, a radiologic dispersal device (RDD).** This consists of a relatively small amount of radioactive material attached to a conventional explosive material, such as a plastic explosive or even a quantity of fertilizer (Fig. 14.1). The principle
In the year 2003, British intelligence found a diagram illustrating the principle of an RDD in Herat, Afghanistan, and concluded that Al Qaeda had succeeded in constructing a small dirty bomb, although the device was never found (Fig. 14.2). The consequences of a “dirty bomb” might be as follows:

- It is unlikely that any person will receive a dose of radiation sufficient to cause the ARS.
- The most likely scenarios would be a small number of persons contaminated with radioactive materials, either on their clothes and skin, or inhaled or ingested. The possibility of a minimal long-term risk of leukemia or solid cancers could not be ruled out.
- The certain consequence would be chaos, psychological terror, and widespread fear.

4. **Hidden radiation exposure device (RED).** An alternative to the “dirty bomb” would be to place a γ-emitting source, such as cesium-137, in a locker or garbage can in a busy public place, such as Grand Central Station, and then make a public disclosure of its presence some days or weeks later. In the intervening period, thousands of people passing close by may have been exposed to a dose of radiation. Like the “dirty bomb,” the probability of a RED is considered to be “highly likely.” The consequences of a RED are as follows:

- Fear and psychological chaos
- Unlikely that any person would receive sufficient radiation to result in the ARS
- Very small long-term possibility of radiation-induced malignancies

is that the conventional explosive material scatters the radioactive material over a wide area. Ideally, the radioactive material should be vaporized, but in practice, this is difficult to achieve except perhaps for cesium-137.

The probability of a “dirty bomb” is considered to be “highly likely”; indeed, some wonder why such an event has not yet occurred. For example, in the year 2003, British intelligence found a diagram illustrating the principle of an RDD in Herat, Afghanistan, and concluded that Al Qaeda had succeeded in constructing a small dirty bomb, although the device was never found (Fig. 14.2). The consequences of a “dirty bomb” might be as follows:

- It is unlikely that any person will receive a dose of radiation sufficient to cause the ARS.
- The most likely scenarios would be a small number of persons contaminated with radioactive materials, either on their clothes and skin, or inhaled or ingested. The possibility of a minimal long-term risk of leukemia or solid cancers could not be ruled out.
- The certain consequence would be chaos, psychological terror, and widespread fear.

**FIGURE 14.1** A radioactive dispersal device (RDD), commonly known as a “dirty bomb,” consists simply of a small amount of radioactive material attached to a conventional explosive material, such as Semtex or even fertilizer. When the explosive material is detonated, the radioactive material is scattered over a wide area.

**FIGURE 14.2** Diagram of a “dirty bomb” discovered in 2003 by British intelligence in Afghanistan. It was assumed that Al Qaeda had succeeded in constructing a small “dirty bomb,” but the device was never found.
Chapter 14 • Radiologic Terrorism

AVAILABILITY OF RADIOACTIVE MATERIAL

As was previously discussed in scenario 1, the detonation of a nuclear weapon would involve a leftover “suitcase bomb” from the former Soviet Union, or the production or theft of fissile material. All possible, but very unlikely. The attack on a nuclear power station, as discussed in scenario 2, would necessitate the hijacking of a commercial airliner. This is again possible, but unlikely because of improved security. Scenarios 3 and 4 require the availability of only relatively small quantities of radioactive material. In both cases, the fear and psychological chaos are not directly related to the quantity of radioactivity or the magnitude of the radiation dose involved. If a Geiger counter ticks, that would be sufficient to cause chaos. Small amounts of radioactive material are readily available.

There are several examples. First, in 2004 in London, an Islamic terrorist cell had collected several thousand household smoke detectors, readily available from hardware stores (Fig. 14.3). Each contains a tiny amount (few thousand becquerels) of americium-241, which is an α-emitter, only dangerous if inhaled or ingested. Imagine the chaos if these were dispersed in a crowded public place.

Second, moisture density gauges, used in the laying of tarmacadam on road surfaces, contain small amounts of americium-241 and cesium-137 (Fig. 14.4). More than 23,000 are in regular use in the United States and about 50 per year are lost or reported missing.

Third, cesium-137 sources are widely used in hospitals and medical centers (Fig. 14.5). Cesium is a very suitable material for a “dirty bomb” because it has a relatively long half-life of about 30 years and, more importantly, it vaporizes readily and can be widely spread by an explosive device. Large cesium sources such as blood or total body animal irradiators are now the subject of intense security and would be difficult to acquire illegally. On the other hand, the smaller sources used for the brachytherapy of cancer do get lost occasionally. In 1998, 19 small tubes of cesium-137, with total activity of about 22 GBq, were reported missing from a medical center in Greensboro, North Carolina.

FIGURE 14.3 An Islamic terrorist cell in London in 2004 had collected several thousand household smoke detectors, which can be readily purchased from hardware stores. Each contains a tiny amount of americium-241, which is an α-emitter. This could be combined with a conventional explosive to produce a “dirty bomb.” Imagine the chaos that would result if this radioactive material were dispersed in a busy public place.
FIGURE 14.5 Cesium-137 would be a very suitable material for a "dirty bomb" because it has a relatively long half-life (about 30 years) and can be widely disseminated because it vaporizes readily. For this reason, security of large cesium-137 irradiators in medical centers has been greatly improved in recent years. Nevertheless, small sources used in the brachytherapy of cancer do get lost from time to time. For example, 19 small cesium-137 sources with a total activity of about 22 GBq were reported missing from a medical center in North Carolina in 1998.

HEALTH EFFECTS OF RADIATION

The health effects of radiation can be divided into two types: deterministic and stochastic, which have quite different characteristics.

Deterministic effects only occur at relatively high doses and result from the killing of many cells in a tissue or organ. Above a tissue-specific dose threshold, the severity of the damage increases with increasing dose as more and more cells are killed. As long as the dose threshold is not exceeded, the effect will not be seen.

Some deterministic effects do not manifest until years later (e.g., cataract, fibrosis) and are therefore known as late effects. By contrast, those that manifest in days to weeks are called early effects, or acute effects. A sufficiently large radiation dose over a short time can result in the ARS, which may result in death. This results from the killing of many cells in the bone marrow or lining of the intestines. Examples of this occurred at Hiroshima and Nagasaki, and also following the reactor accident at Chernobyl. A dose of several grays is needed to result in these effects, which is possible following the detonation of a nuclear weapon or the meltdown of a nuclear reactor, but would be unlikely to occur following a dirty bomb. Because a whole chapter is devoted to the ARS (Chapter 8), no more will be said about it here.

Stochastic effects, of which carcinogenesis is the most important, are all-or-nothing effects...
EXTERNAL CONTAMINATION

External contamination refers to radioactive material on the surface of the body. Most external contamination (up to 90%), following an event like a dirty bomb, can be disposed of by removing the clothing, which should be placed in a plastic bag and labeled. A Geiger counter can then be used to survey the patient. The top priority to decontaminate is open wounds because they offer a fast direct route for internalizing radioactive materials and transportation to critical organs. Next is the nose and mouth, followed by intact skin. Contaminated areas should be carefully washed with soap and water. The survey–scrub–rinse sequence of a wound or intact skin should be repeated until the readings on the survey meter drop to twice background or until further efforts do not result in a decrease in radiation levels. Too vigorous an effort should be avoided because damaging the skin could create a pathway for internal contamination.

INTERNAL CONTAMINATION

The possibility of significant internal contamination is a much more difficult problem. Contaminants gain entry into the body by several routes, such as through inhalation into the lungs, ingestion into the gastrointestinal (GI) tract, or percutaneous or transdermal absorption through intact skin and particularly through open wounds. Some radionuclides tend to remain in the body, often concentrating in a particular tissue or organ. Iodine in the thyroid or transuranics such as plutonium or americium in the bone are some examples. Others may be eliminated in urine, feces, or perspiration. Radionuclides within the body that are $\gamma$-emitters can be detected and measured with a whole body counter. Quantities of $\beta$-emitters can only be estimated by measuring amounts in excreted materials. Not much can be done to counter the effects of internal contamination, except in a few specific cases. One such case is radioactive iodine. If taken within 4 to 6 hours of contamination, stable iodine in the form of potassium iodide saturates binding sites in the thyroid and inhibits the incorporation of radioactive iodine. The Chernobyl experience has shown clearly that radioactive iodine can cause hypothyroidism and can also result in cancer. This is particularly important for children as well as for the developing embryo or fetus. Radioactive
iodine is unlikely to be released from an RDD, but certain to be present with a nuclear weapon or a serious accident at a nuclear reactor because it is a fission product.

The only other countermeasure approved by the U.S. Food and Drug Administration (FDA) is the use of Prussian blue (ferric III hexacyanoferrate II) to block the uptake of cesium-137 from the intestine. This could be important because, as previously mentioned, cesium-137 would be well suited for an RDD.

### MEDICAL MANAGEMENT ISSUES IN THE EVENT OF RADIOLOGIC TERRORISM

Many major medical centers have put together a “radiation casualty team” consisting of health providers, physicists, social workers, and administrators to cope with the possibility of a radiation accident or an act of radiologic terrorism. In most circumstances that can be imagined, there is likely to be a large number of people who are frightened and worried, a much smaller number that may actually be contaminated with radioactive material, and an even smaller number who need treatment for a significant radiation exposure.

The highest priority is to give critical care to those who have suffered life-threatening injuries (Table 14.1). This should not be delayed because of the possibility of radioactive contamination. Radiation exposure and contamination are secondary considerations. Once life-threatening injuries are taken care of, the next step is to use a sensitive Geiger counter to identify those persons who are contaminated with radioactive materials, which in most cases, will be in the form of dust particles on the body and/or clothing. Contaminated clothing should be removed and sealed in plastic bags; this is likely to account for 80% to 90% of the contamination. The next priority is to decontaminate any open wounds because this can be a conduit for radioactive material to be internalized. This can be achieved by gentle irrigation. Lastly comes the decontamination of intact skin. Overly aggressive methods should be avoided and decontamination efforts should be stopped when radiation levels are less than about twice background. Urine samples should be collected to identify persons who have internalized radioactive material by inhalation or ingestion, but there are very limited treatments available as discussed previously.

### FURTHER INFORMATION

More detailed information on this topic is available from several sources, such as the following:

The Department of Health and Human Services (DHHS) established the Radiation Emergency Medical Management website (http://www.remm.nlm.gov) in 2007 to give guidance to healthcare workers on radiation health effects. The Radiation Studies Branch of the Center for Disease Control has a website (http://www.bt.cdc.gov/radiation/clinicians.asp) suitable for both professionals and the public, giving information on protective measures during a nuclear attack or a radiologic event. They also offer a handbook, which is available at http://www.bt.cdc.gov/radiation/pocket.asp.

The Department of Energy’s Radiation Emergency Assistance Center Training Site (REAC/TS) in Oak Ridge, Tennessee provides online guidance for medical management of radiation events.

The American College of Radiology (ACR) and the American Society of Therapeutic Radiology and Oncology (ASTRO) have prepared a primer entitled “Disaster Preparedness of Radiology Professional: Response to Radiological Terrorism,” which is available at http://www.astro.org/GovernmentRelations/RadiationDisasterManagement.
The National Council on Radiation Protection and Measurements (NCRP) prepared NCRP Report No. 138, entitled “Management of Terrorist Events Involving Radioactive Material,” which aims to provide guidance to those responsible for responding to terrorist events.

**SUMMARY OF PERTINENT CONCLUSIONS**

The following are several possible scenarios for radiologic terrorism:

- **Detonation of a nuclear weapon**
  - **Risk**
  - Exposure to γ-rays and neutrons
  - Fallout of fission products
  - **Outcome**
  - Large number of acute deaths
  - Long-term carcinogenesis
  - **Likelihood**
  - Remote

- **Attack on a nuclear power plant**
  - **Risk**
  - Attack on reactor itself
  - Attack on stored used fuel elements
  - Release of fission products
  - **Outcome**
  - Unlikely to involve acute deaths
  - Long-term carcinogenesis
  - **Likelihood**
  - Extremely unlikely

- **Dirty bomb (RDD)**
  - **Risk**
  - Release of radioactive material
  - Small number of contaminated people
  - Large number of slightly contaminated people
  - Psychological chaos, many frightened people
  - **Outcome**
  - Unlikely to involve acute deaths
  - Small risk of long-term carcinogenesis
  - **Likelihood**
  - Likely

- **Hidden RED**
  - **Risk**
  - Large number of people exposed to small doses of radiation
  - Psychological chaos, many frightened people

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**Outcome**
- Unlikely to result in acute deaths
- Small risk of long-term carcinogenesis

**Likelihood**
- Likely

**Availability of radioactive material**
- Small amounts readily available from smoke detectors, humidity gauges, and lost or stolen medical sources

**Health effects of radiation**
- Deterministic effects occur at high doses, such as cataract, fibrosis, or the ARS.
- Stochastic effects, including carcinogenesis and heritable effects, are important even at lower doses and occur much later.

**External exposure to radiation** refers to irradiation from an outside source that never comes in contact with the body. **External contamination** refers to radioactive material on the skin or clothing. **Internal contamination** refers to radioactive materials that are inhaled, ingested, or internalized through open wounds.

**Medical management issues in the event of radiologic terrorism**
- Standard medical triage; attend first to critical injuries
- Decontaminate
- Remove clothing, survey with Geiger counter
- Decontaminate
- Open wounds
- Mouth and nose
- Intact skin
- Cease decontamination efforts when:
  - Further efforts do not reduce count
  - Count less than twice background
- Collect urine sample to detect internal contamination
- Potassium iodide (KI) tablets to stop uptake of radioactive iodine
- Prussian blue to prevent absorption of cesium-137

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**BIBLIOGRAPHY**


The challenge of noninvasively mapping biologic tissue has occupied radiologists and engineers for over a century. Shortly after Roentgen’s discovery of the x-ray in 1895, it was applied toward medical imaging by directing x-rays through his wife’s hand and onto a piece of film, which revealed the bony structure of her hand. In the time since that landmark discovery, medical imaging has evolved to encompass multiple modalities and mechanisms of signal generation and reconstruction. X-ray radiography matured over the first half of the 20th century, while the first medical ultrasonic measurement was made in 1954, followed by the invention of the Anger camera in 1964 and the advancement of nuclear medicine, the introduction of the clinical x-ray computed tomography (CT) scanner in 1972, the acquisition of the first magnetic resonance image (MRI) in 1973, and the introduction of the first clinical positron emission tomography (PET) device in 1975. These methods have developed to offer high imaging performance in terms of scanning time, spatial resolution, and image quality, and now are commonplace in medical settings.

Imaging techniques were first exploited by the radiation oncologist for the identification and targeting of tumors to be irradiated. With the use of radiographic imaging to localize a neoplastic mass within a patient, radiotherapy can be more accurately delivered to the intended target while sparing surrounding tissues. Furthermore, as radiation therapy physics progressed toward the calculation of delivered doses using patient-specific information, three-dimensional x-ray CT was used as a model of patient structure. This modality is especially well suited to radiotherapy planning as the image intensities reflect x-ray absorption and can therefore be used to accurately predict dose deposition. For these reasons, x-ray radiography and CT are now essential tools in the radiation oncology clinic. As MRI methods have advanced, this imaging modality and its superior soft tissue contrast relative to x-ray CT have also become incorporated into many radiation oncology departments. However, a shortcoming of these modalities is that their source of contrast is gross tissue anatomy or structure. X-ray CT differentiates tissues based on their density and x-ray absorption, whereas MRI detects differences in proton density and magnetic relaxation. These properties are not directly related to the physiologic or functional aspects of the tissue being imaged. This limitation of conventional imaging prompted the development of molecular imaging, generally defined as the noninvasive detection, localization, and quantification of specific molecular entities or physiologic processes in a living organism.

Over the last 20 years, the field of molecular imaging has undergone tremendous growth, with significant advances in the interrelated areas of imaging technology, imaging target identification, molecular probe development, and image analysis.
As such, these imaging methods are now being adopted into the practice of radiation oncology to better stage and plan tumors for treatment and to monitor their response to therapy. Despite encouraging preliminary results of studies employing molecular imaging in the diagnosis, staging, and treatment of cancer, many challenges remain to be overcome to fully exploit the potential of this technology in this setting. The imaging methods with the most utility for oncology must be identified, and quantitative and/or volumetric metrics that can be extracted from these images must be established and incorporated into the staging, treatment planning, and posttreatment monitoring of radiation oncology patients. Many molecular imaging modalities have been engineered, including nuclear medicine techniques, optical imaging, and functional MRI methods including fMRI and magnetic resonance spectroscopy; however, this chapter will focus on two imaging techniques with widespread utility in radiology and radiation oncology: x-ray CT and PET. Although CT is commonly used as an anatomic imaging modality, developments with x-ray contrast agents and dynamic acquisition protocols have allowed the collection of functional information with this imaging method. Alternately, PET imaging of radioactive imaging agents has reached a point where this technique is now routinely being used to complement conventional imaging. The physical aspects of each of these modalities will be summarized, after which their role in clinical imaging will be described, including discussion of their use in clinical tumor staging, treatment planning, and follow-up.

**X-RAY COMPUTED TOMOGRAPHY**

Imaging of biologic tissues with electromagnetic radiation is among the oldest medical imaging methods and has a long history in its application to oncology. Although two-dimensional x-ray radiographs were first applied to the visualization of tissues of interest, the continued advancement of x-ray imaging technology has resulted in the widespread adoption of both two-dimensional x-ray CT and three-dimensional x-ray CT in the modern oncology clinic.

**Basic Physics**

*X-ray Radiography and Computed Tomography*

Consider a beam of x-rays passing through the chest of a patient, striking a film on the far side. Ignoring scatter in this treatment, one can consider the x-ray beam to be made up of several smaller, parallel beams, each passing through specific tissue volumes on their way to the film. Because each tissue volume has its own x-ray absorption properties depending on the atomic composition of the tissue, the image formed on the film will vary in brightness depending on how the patient attenuated the x-ray beam at different spatial locations. This concept of measuring differential x-ray absorption is the basis of all x-ray imaging methods. If the measurement geometry and the intensity of the source are known, the average mass attenuation coefficient (\(\mu/\rho\)) along the path traversed by each x-ray path may be calculated. In biologic subjects, bony tissues are typically strongly absorbing and have high \(\mu/\rho\) for x-rays at the energy of typical imaging systems, whereas low-density tissues, such as lungs, are poorly absorbing and have low \(\mu/\rho\).

Two-dimensional x-ray radiography is commonly used for various medical applications, including the diagnosis of bone fractures, pneumonia and lung diseases, and congestive heart failure. This technique has been extended using fast digital detectors to allow dynamic imaging of a subject, known as x-ray fluoroscopy. This method facilitates image-based monitoring of dynamic physiologic processes such as in angiography, the assessment of interventional procedures such as device implantation, and precise patient positioning such as that needed for radiotherapy setup. However, two-dimensional x-ray imaging collapses all information along the third dimensions, providing a projection through a subject instead of resolving objects in three dimensions. Several situations may not be amenable to this technique, for example, in the detection of a lung lesion that is masked by highly absorbing ribs in the same x-ray projection. To overcome this limitation, three-dimensional x-ray imaging methods employing tomography have been developed. The mathematics of x-ray tomography were elucidated as early as the 1930s. Tomography involves the acquisition of multiple two-dimensional projection views of a subject, with each view acquired through a different projection angle as demonstrated in Figure 15.1A. These views are then combined using a tomographic reconstruction algorithm to mathematically resolve the three-dimensional structure of the subject. X-ray CT has matured through
FIGURE 15.1 X-ray computed tomography image acquisition and reconstruction. A: A CT imaging system consists of an x-ray source (top) and detector (bottom) that can rotate around a subject to collect multiple x-ray projection views. B: Backprojection reconstruction of CT data involves the smearing of a measured one-dimensional x-ray projection along the line of projection, shown here for two projection views. When expanded to many views, the backprojected data approximates the structure of the object for which data were collected. (Courtesy of Dr. E.E. Graves.)
metal such as copper or aluminum in the beam path. This has the effect of preferentially removing low-energy photons from the beam and increasing the average energy of the beam. Beam profiles for CT scanning are generally either fans (wide angular dispersion, narrow profile) designed to function with a correspondingly narrow one-dimensional detector array, or cones (rectangular profile) for irradiation of a rectangular planar two-dimensional detector array.

The simplest x-ray detector is radiographic film, which darkens upon absorption of x-rays. However, the need for a reusable, digital detector capable of rapid and repeat imaging has encouraged the development of electronic x-ray detectors. Although detectors such as scintillators that convert x-rays into visible photons and subsequently to electrical signals are common, many modern x-ray imaging systems now employ solid-state semiconductor detectors because they are capable of direct digital detection of x-rays without the need for conversion of the incoming photons into an intermediate form, such as visible photons. Research is ongoing to engineer new detector materials capable of measuring very small numbers of photons to reduce the dose required for x-ray imaging. In modern CT scanners, multiple detector rows are now commonly used to collect multiple CT slices in a single acquisition, resulting in scanners generally referred to as multidetector CT (MDCT). These advances have resulted in scan times for x-ray CT as short as 0.5 seconds per frame. Helical scanning protocols, in which the subject is translated longitudinally through the CT ring as the gantry rotates and projection data are recorded, has further accelerated the acquisition of volumetric CT data. In addition to advances in fan beam architectures for CT, cone beam scanners have emerged following the development of high-performance flat panel imaging arrays and effective cone beam volumetric reconstruction methods, and are now being integrated in many linear accelerator designs to facilitate the collection of CT data during radiotherapy.

**Image Reconstruction and Analysis**

The most common reconstruction algorithm applied to x-ray CT data is the filtered backprojection technique, a form of the inverse radon transform. This method uniformly distributes each measured projection back along the line of several data collection geometries and reconstruction methods since its introduction in 1972; however, the fundamental tomographic concept remains the same, as does the source of contrast (x-ray attenuation) in the images.

**Imaging Hardware**

X-ray imaging and CT employ an x-ray source that produces a beam that is directed through a subject, and an x-ray detector that measures several x-rays that traverse the subject. The geometry of the CT scanner has evolved through several forms as this imaging modality has been developed. Hounsfield’s original scanner, known as “first generation” CT, included only one or two spatial detector channels, which therefore measured x-ray penetration through the subject at only a few discrete locations. This detector setup, paired with a tightly collimated x-ray beam that only irradiated these detectors, was translated across the patient to measure x-ray fluence along a single projection angle at multiple locations. After recording this projection, this apparatus was rotated and the process repeated to record a new projection. CT scanners have evolved greatly since this original instrument, with wide-angle x-ray sources and detectors minimizing the time required to collect a full set of tomographic projections. In most modern CT scanners, the source and detector are mounted on a rotating gantry as depicted in Figure 15.1A. The most common x-ray source used in imaging systems is the x-ray tube. This device is a type of vacuum tube, consisting of a cathode that generates electrons and an anode toward which the electrons are accelerated through a voltage applied across the cathode/anode pair. The anode absorbs the energy of the accelerated electrons and emits a portion of it in the form of x-rays. The x-rays produced are known as Bremsstrahlung or braking radiation, because the incident electrons are slowed and the energy lost is emitted in the form of an x-ray. The emitted x-rays range in energy from near zero to a maximum corresponding to the voltage applied to the tube. Generally, the mean photon energy produced by an x-ray tube is approximately one-third of the voltage applied to the tube. X-rays produced by the anode are then emitted through a window in the vacuum tube and collimated to a desired beam profile. The beam is commonly also “filtered” by the insertion of thin sheets of
from which it was acquired, as demonstrated in Figure 15.1B. This results in an image of the original object when sufficient projections are used. Because this technique introduces a blurring into the reconstructed image, the projections are sharpened through application of a ramp filter in the frequency domain prior to backprojection. Because of the filter that is used, filtered backprojection is sensitive to high-frequency noise, which commonly necessitates the use of smoothing to reduce noise at the cost of spatial resolution. To overcome the shortcomings of this method, several iterative techniques have been proposed that improve reconstruction performance at the cost of increased computational complexity. Iterative routines function by modeling the data acquisition process and applying this model to an initial guess at the reconstructed image distribution, producing a set of simulated measurements. These simulations are then compared to the actual measurements and the reconstructed image is correspondingly adjusted. This process is then repeated until there is acceptable agreement between the measured and simulated data. Although computationally expensive, these methods allow sophisticated modeling of the data acquisition process, permitting consideration of scatter, noise, and other factors.

Pixel intensities in reconstructed CT images are typically converted into the Hounsfield unit (HU) scale, a standardized intensity range constructed to permit comparison of x-ray opacity between images and scanners. In this scale, all measured x-ray absorption values \( \mu \) are related to that of water \( \mu_{\text{water}} \) by the relation

\[
\text{HU} = 1,000 \times \left( \frac{\mu - \mu_{\text{water}}}{\mu_{\text{water}}} \right)
\]

This is a linear function; therefore, HUs are proportional to the measured x-ray absorption \( \mu \). Examining the given expression, it can be seen that the HU of water is 0, whereas the HU of air, with \( \mu \approx 0 \), is −1,000. Water and water-based soft tissues demonstrate HU of 0 to 100, whereas fatty tissues occupy the HU range −100 to 0. Strongly absorbing bones exhibit HU of 500 or greater. Modern CT scanners are capable of detecting differences in HU on the order of 5 to 10 HU.

**X-ray Contrast Agents**
Agents that alter x-ray absorption after delivery to a subject have been developed nearly as long as x-ray imaging itself to improve on the minimal x-ray contrast between soft tissues. Because of the linear relationship between concentration and x-ray attenuation, and between x-ray attenuation and HU, the effectiveness of an x-ray contrast material is dictated by the x-ray absorption of its constituent materials and the concentration at which it can be delivered. Strongly absorbing elements including barium (\( Z = 56 \)), gadolinium (\( Z = 64 \)), gold (\( Z = 79 \)), bismuth (\( Z = 83 \)), and tungsten (\( Z = 74 \)) have been studied as potential x-ray contrast materials; however, iodine (\( Z = 53 \)) has emerged as the predominant component of clinical x-ray contrast agents because of its biocompatibility and its ease of incorporation into labeling chemistries. Sodium iodide was administered as an oral contrast agent for x-ray imaging in 1918, and subsequently various iodine-containing compounds were engineered to include more iodine per mole, improve pharmacokinetics, and reduce toxicity. Although ionic compounds prevailed through the first half of the 20th century, the synthesis of nonionic multi-iodinated agents such as iopamidol and iohexol in the 1970s greatly reduced the toxicity of iodinated contrast agents, and these have become the clinical standard. Gases including krypton (\( Z = 36 \)) and xenon (\( Z = 54 \)) have also been used as inhaled x-ray contrast agents. However, the limiting factor with all x-ray contrast agents is the large concentrations required to produce a measurable change in imaging signal. For iodine, the concentration required to produce a CT change of 100 HU is approximately 10 mM.

**Nanomaterials**
The development of nanotechnology over the last 10 years has encouraged engineering of new x-ray contrast materials by overcoming the limitations of previous contrast agents. Nanoparticles facilitate the use of otherwise toxic or biologically incompatible materials in contrast agent formulations, while also permitting the packing of a large number of contrast-producing atoms in a single composite structure. In addition, the large surface area and functionalized status of many nanoparticle constructs allows the conjugation of other components to the nanoparticle to alter molecular targeting, biodistribution, and/or pharmacokinetics.

Iodine-based or iodine-containing nanoparticles have been a topic of intense interest, with
multiple synthetic strategies having emerged including highly iodinated fullerenes, polymeric macromolecules containing triiodinated moieties, crystalline iodinated nanoparticles dispersed with surfactant, and polyethylene glycol (PEG)-based shells enclosing iodinated materials at high concentrations.

More recently, significant work has been performed to evaluate the use of gold nanoparticles for CT contrast. Hainfield and colleagues showed that small (~100 μm) blood vessels can be visualized within minutes of intravenous delivery of gold nanoparticles. As with iodine nanoparticles, PEG has also been incorporated into gold nanoparticles to alter their pharmacokinetics. Although concentration limits have hampered the use of contrast CT for molecular imaging, the use of gold nanoparticles in this context has been examined in vitro through the coupling of UM-A9 antibodies specific to head and neck squamous cell carcinoma to gold nanorods. Although measurements of cells with and without the target antigen demonstrated a fivefold difference in HU, it remains unclear whether this level of contrast will be sufficient in vivo. Targeting of gold nanoparticles to folic acid receptor has also been investigated as a second molecular-specific CT imaging agent.

Nanoparticles synthesized from bismuth sulphide have also recently been evaluated for in vivo CT vascular imaging. This agent has been seen to be tolerable to mice and rabbits at a dose containing 0.2 M bismuth, giving CT numbers in excess of 500 HU in vivo. Toxicity for bismuth nanoparticles was seen to be significantly less than equivalent concentrations of bismuth salts. Other heavy metals including gadolinium, dysprosium, erbium, europium, and lutetium have been incorporated into fullerene structures to produce water-soluble x-ray contrast materials. These molecules represent additional pathways toward novel x-ray contrast materials that may yet evolve into valuable tools for CT imaging.

**Radiotherapy Dose Enhancement**

A potentially useful secondary property of x-ray contrast materials is their ability to enhance the cell kill produced by high-energy radiation. This effect is mediated by the electrons produced on photoelectric absorption of high-energy photons by high-Z materials and is produced both by endogenous materials such as bone as well as exogenous contrast agents. Although pair production for megavoltage photons commonly used in radiotherapy may also enhance radiation dose effect, even at this energy range, photoelectrons are the primary cause of this enhancement. This enhancement has been studied by several groups through simulations and animal experiments; however, it has a limited history of clinical application with only preliminary studies of kilovoltage radiotherapy of human brain tumors after intravenous injection of an iodinated contrast agent having been conducted. Recent developments with nanoparticle technology may yet exploit this phenomenon, however.

**Clinical Applications of X-ray Computed Tomography**

**Radiotherapy Planning**

One of the principle tasks of radiotherapy planning is the delineation of patient anatomy, including the desired radiation target as well as organs at risk that should be spared from radiation dose. These identified volumes are then used to compute an optimal radiation treatment strategy, employing the degrees of freedom available to the specific radiotherapy treatment device. X-ray CT has become the primary imaging modality used for this process, both because of its high spatial and temporal resolution and lack of image distortion artifacts, as well as the fact that CT image intensities denote x-ray absorption and can be used to predict dose deposition by a treatment beam within the patient. However, the low CT contrast between soft tissues complicates the task of the radiation oncologist and has raised interest in use of other imaging modalities such as PET to improve radiotherapy planning. This is discussed further in the subsequent paragraphs.

**Vascular and Perfusion Imaging**

Because of the physiologic and clinical significance of the vasculature within tumors, efforts to image this aspect of the cancer microenvironment have been widespread. With CT, one mechanism to achieve this goal is the intravenous delivery of a contrast agent followed by repeat imaging as the agent passes through the vasculature. Through analysis of the collected images, various parameters including blood flow, blood volume, and vascular permeability may be derived. This method is generally called dynamic contrast-enhanced (DCE) imaging. In its
simplest form, this method can be applied following an intravenous bolus contrast injection to acquire images timed to when the contrast material is passing through a specific portion of the vasculature. For example, commonly contrast-enhanced CT includes the acquisition of an “arterial phase” image as the agent is passing through the arterial system, followed by a later “venous phase” image. These images may then be considered either independently or together to assess vascular structure and function, as shown in Figure 15.2. More sophisticated DCE-CT approaches however involve the acquisition of images with greater temporal resolution over a longer period before and after contrast injection. The signal curve obtained for a given pixel over time may be analyzed in several ways to estimate specific vascular parameters. Empirical assessment of the CT signal versus time curve has been evaluated, examining measurements such as the maximum uptake of contrast agent, the time to reach maximum uptake, or the maximum slope of the initial signal rise, and many of which have been used to identify and characterize malignant lesions. Alternately, several quantitative models that describe the pharmacokinetics of contrast agents have been developed. These models can be fit to experimentally measured signal curves to estimate the parameters of the model, which may include vascular volume, perfusion, vascular permeability, and vascular surface area depending on the specific model and acquisition sequence. This imaging protocol and analysis is demonstrated in Figure 15.3.

**Tumor Diagnosis, Staging, and Response Assessment**

X-ray CT has become a routine clinical procedure in the management of oncology patients. Lung tumors are particularly suited toward diagnosis with x-ray imaging methods because of the large contrast between low-density, air-filled

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**FIGURE 15.2** Contrast-enhanced x-ray computed tomography. Arterial (top row) and venous (bottom row) phase images were acquired from a patient with pancreatic cancer at set delays after intravenous delivery of an iodinated x-ray contrast agent. Axial (left), coronal (middle), and sagittal (right) views are shown for each dataset. Increases in CT signal are apparent in the aorta in the arterial phase axial view (solid arrows), whereas the vascularized portion of the pancreatic lesion can be differentiated between the arterial and venous phases (dashed arrows). A pancreatic stent is prominent in the coronal images, as is the delivery and clearance of the contrast material in the blood pool of the heart (solid arrows). (Courtesy of Dr. A.C. Koong.)
lungs and solid lung tumor masses. Fast CT scanning technology as well as methods for managing breathing motion, including breath-hold CT and four-dimensional (time-resolved) CT, have further improved the utility of this imaging modality for interrogating thoracic lesions and especially for delineating spatial targets for radiotherapy. The high spatial resolution of x-ray CT has encouraged the use of CT-based measures of tumor volume as a prognostic marker. However, the usefulness of this measure is limited for small nodules, which has encouraged the use of contrast-enhanced CT to evaluate malignancy.

Multiphase contrast CT is commonly used to diagnose and characterize various tumor types, including lesions of the liver, pancreas, and kidney. Commonly, two or more images are acquired at standardized delays after contrast delivery, focusing on times at which the contrast agent highlights specific components of the vascular system, including the arterial system, the portovenous system, the pancreatic vasculature, and the corticomedullary vasculature, depending on the site being studied. For example, contrast agent washout in the venous phase has been reported to be a specific hallmark of malignant lesions within the liver. Reports of the accuracy of this staging method have commonly noted an improvement in performance when imaging lesions greater than 1 to 2 cm in diameter.
X-ray mammography is the dominant screening method in use for early breast cancer detection. Although this method is inherently two-dimensional, volumetric x-ray imaging method for breast is currently in development, with the goal of improving lesion detection while not significantly increasing the radiation dose of the procedure relative to mammography. Devices for multiple view stereoscopic digital mammography and digital breast tomosynthesis as well as dedicated breast CT devices have been reported but are yet to enter into widespread clinical use. Contrast agents have been used in conjunction with breast x-ray imaging to improve diagnostic ability. In addition, dynamic x-ray imaging has been applied to the characterization of breast lesions to identify axillary metastasis based on vascular properties.

**POSITRON EMISSION TOMOGRAPHY**

Although x-ray imaging has evolved to facilitate high spatial and temporal resolution imaging of biologic subjects in three dimensions, it is fundamentally limited by the minimal contrast in terms of x-ray attenuation between tissues. Although the development of x-ray contrast materials has alleviated this limitation for certain applications, the low contrast of x-ray CT continues to impose limitations on the use of this imaging modality for various tasks, including tumor diagnosis and target definition for radiotherapy planning. Emerging molecular imaging modalities have the potential to overcome this obstacle by generating image contrast through other functional and molecular mechanisms. PET is such a molecular imaging technique that matured from its roots in nuclear medicine through the development of a dedicated clinical scanning device in the 1970s. PET scanners are now common components of radiology and nuclear medicine departments and are beginning to be deployed in other medical settings as well. The field of radiation oncology has begun to embrace PET as a means of staging tumors, planning treatments, and assessing response using functional and metabolic information in addition to the anatomic information provided by CT.

**Basic Physics**

**Positron Decay**

PET is based on the decay process of positron emission undergone by several unstable radioactive isotopes including carbon-11, nitrogen-13, fluorine-18, copper-64, and iodine-124. In this process, a proton in the nucleus of the isotope is converted into a positron that is ejected from the atom, as well as a neutron and a neutrino. The ejected positron then travels some distance, generally in the range of 1 to 10 mm depending on its energy, before annihilating with its opposite particle—the electron. In this annihilation, the positron and electron energy are converted into a pair of 511 keV photons, emitted in opposite directions.

**Tomography and Imaging Hardware**

The tomographic principle applied to imaging positron emission events involves detection of the annihilation events produced, summarized in Figure 15.4A. PET scanners rely on the concept of coincidence detection to localize the source of positron emission. Because each positron emission event results in two photons traveling at 180 degrees to each other, the PET detector is designed to report on instances where two photons are recorded at separate locations on the detector within a narrow space of time. These pairs of events are termed “coincidence events” and are assumed to represent two photons that were generated through a single positron emission event and subsequent annihilation. By measuring several recorded coincidence events for each pair of detectors, this data may then be back-projected in a manner similar to that described previously for CT to reconstruct the distribution of positron-emitting radioisotopes. This tomographic concept was first proposed in the late 1950s and early 1960s, but it was not until the mid-1970s before a working prototype scanner was built and validated. Modern clinical PET scanners demonstrate spatial resolutions of 4 to 6 mm and can record 200 to 2,000 counts per second for a $\mu$Ci point source. This is caused by advances in scanner geometry, electronics, and detector materials.

The performance of a PET scanner is dictated by several factors. In reality, the angle between the emitted annihilation photons is slightly noncollinear, with a divergence of 0.5 to 1 degree. This deviation from linearity causes a reduction in PET spatial resolution, with the problem worsening with increasing scanner bore size. In addition, uncertainty as to where a given photon was absorbed in a thick detector also decreases the spatial resolution of PET scanning.
FIGURE 15.4  Positron emission tomography image acquisition and examination protocol. A: A probe containing a radioactive isotope that decays by positron emission, in this case, 2-deoxyglucose labeled with fluorine-18 ($^{18}$F), emits a positron. The positron annihilates locally with an electron, producing two anti-parallel 511 keV photons. These photons are measured as near-simultaneous events on a detector ring. B: A PET examination involves the injection of a probe to a subject that then distributes according to its biologic activity, accumulating in sites of interest as well as background organs. For [$^{18}$F]fluoro-2-deoxyglucose (FDG), intense uptake caused by high glucose utilization is seen in tumors and the brain, whereas strong PET signals are seen in the kidneys and bladder due to excretion of the probe. (Courtesy of Dr. E.E. Graves.)
Currently, several groups are pursuing “depth of interaction” detector technology to estimate where within a crystal an event occurred to mitigate this effect. Scattering of annihilation photons between their generation and detection by a scanner also limits PET performance, as does the occurrence of random coincidences when two photons from two separate annihilation events strike the detector array simultaneously. Strategies for correcting for these phenomena are discussed in the subsequent paragraphs.

Although conceived of during the initial development of PET, recent advances in detector technology has facilitated the implementation of “time of flight” (TOF) PET. In this strategy, the difference in arrival time of two annihilation photons is recorded in addition to their spatial orientation. If the time difference is known with high precision, it is possible to explicitly calculate where along the line connecting the detectors the annihilation photon pair were generated. Current detector technology can localize the positron event down to a 10-cm range, which does not allow complete image reconstruction from TOF data alone but does significantly enhance the quality of images when incorporated into conventional reconstruction schemes. Furthermore, incorporation of respiratory monitoring equipment with PET scanners has facilitated the acquisition of motion-resolved, 4D PET datasets using methods analogous to 4D CT.

Integration with Computed Tomography

Although PET offers a radically different contrast mechanism than conventional anatomic imaging modalities such as CT and MRI, this can hinder image interpretation because of the lack of anatomic information. PET data can be aligned with such anatomic images using various image registration techniques; however, this procedure is dependent on the accuracy of the registration algorithms and can be computationally expensive. A more attractive solution is the acquisition of both anatomic and functional imaging data in a single examination without moving the patient between imaging systems, allowing direct overlay of the two datasets. This strategy has been realized through the combination of CT and PET imaging hardware in a single system known as PET/CT (Fig. 15.5). This engineering challenge is solved by creating a scanner encompassing a CT gantry and a PET gantry, each with their respective imaging components, mounted coaxially with a patient bed sliding through both scanners. With this hybrid system, a whole body x-ray CT scan may be acquired and used to select a region for subsequent PET scanning, which can then be conducted by moving the patient bed such that the appropriate anatomic site now lies within the PET subsystem.

As described earlier, a key motivation for PET/CT is the acquisition of intrinsically aligned PET and CT images in a single examination. The merits of this practice have been reported for various oncologic imaging applications, demonstrating the improved diagnostic ability for multimodal PET/CT image datasets beyond PET or CT alone. However, a second advantage of PET/CT is the ability to use the high-resolution CT dataset to correct the PET image for attenuation of the annihilation photons as they travel out from the subject. This is accomplished by estimating x-ray attenuation along the line connecting each detector pair and correcting the measured coincidence count rate for that pair by the expected attenuation factor. This was previously done in stand-alone PET scanners using low-resolution transmission scans acquired using a radioactive source; however, the improved accuracy of CT has greatly enhanced this procedure in PET/CT systems. In radiation oncology settings, the emergence of PET/CT has facilitated the use of PET/CT in treatment planning as the functional PET data may be considered alongside the CT data that is used for treatment planning as described earlier.

Image Reconstruction and Analysis

A PET acquisition results in a dataset consisting of several coincidence events recorded for each pair of detectors. For 4D PET, individual coincidence events are associated with their time of measurement so that they can be retrospectively associated with an external measurement of breathing motion so as to reconstruct PET images associated with each breathing phase, just as is done with x-ray data in 4D CT (Fig. 15.6). A “delayed window” method is commonly used to reduce the influence of random coincidences by estimating the frequency of random coincidences for each detector pair and by subtracting this rate from the measured count rate. The influence of scattered photons on the measured count rate for each detector pair is more difficult
FIGURE 15.5 Combined PET/CT. A: A clinical PET/CT scanner. The individual gantry rings for the PET and CT subsystems can be seen within the bore as the thick and thin rings, respectively. B: Sagittal slices from CT, PET, and fused PET/CT images collected with a combined PET/CT system. The acquisition of CT and PET data during the same examination allows direct overlay of the two images, typically shown with the CT in grayscale and the PET in pseudocolor as seen at right. (Courtesy of Dr. E.E. Graves.)
PET systems, including maximum-likelihood expectation maximization (MLEM), ordered subset expectation maximization (OSEM), and row-action maximization-likelihood (RAMLA). After reconstruction, the pixel intensities of the images are calibrated to a meaningful physical scale, similar to the calibration of CT images to the HU scale. For PET, the physical property of interest is the amount of radiotracer in each pixel, which is calculated by multiplying the reconstructed image in units of counts by an experimentally measured calibration factor to produce pixel intensities in units of activity per unit volume (mCi/mL). After representation of the image in physically meaningful units, the image intensities are commonly further manipulated into the standardized uptake value (SUV) scale. This is typically done by dividing the amount of radiotracer in each pixel by the metabolic rate of the tissue in which it is located.

FIGURE 15.6 Four-dimensional PET and CT imaging. The top row shows three slices from a 4D CT dataset acquired at expiration (left), mid-inspiration (middle), and full inspiration (right). The tumor volume delineated on the expiration scan is shown in red. In the bottom row, the corresponding 4D PET scan is shown at the same phases of the breathing cycle. (Courtesy of Dr. L. Xing.)
measured image intensities by the injected dose divided by the patient weight. Although the accuracy of this value has been called into question because of its sensitivity to various factors, the SUV remains a standard method of quantitating the degree of radiotracer uptake observed in a PET scan.

Positron Emission Tomography Radiotracers

The PET acquisition and image reconstruction process is not specific to any radionuclide or radiopharmaceutical, it can be applied to any subject containing a positron-emitting compound. Therefore, in conjunction with PET system design, there has been an equal effort to design compounds labeled with positron-emitting radioisotopes that can be administered to a subject and that will distribute according to a specific physiologic or molecular process, so that the PET image can be used to quantitate said process. As picomolar concentrations of positron-emitting radioisotopes can be detected with PET, PET imaging probes are called “radiotracers” because they are given in trace doses in order to sample rather than perturb the physiology of the subject. A typical in vivo PET examination involves the systemic injection of a radiotracer at a dose of ~7.4 MBq for mice and 200 to 600 MBq for humans, and imaging at one or more time points after injection to measure the distribution of the probe, as shown in Figure 15.4B. Dynamic imaging may be performed as described for CT to map the pharmacokinetics of the probe and the estimated specific physiologic parameters, or alternately imaging may be done at a single time point when the probe has distributed according to its biologic activity and PET signal intensities are well associated with the biologic parameter of interest. In general, this association improves with increasing circulation time as the probe associates with its target while nontarget probe accumulation is cleared; however, because of the finite half-life of PET radiotracers, a tradeoff must be made between imaging at early times when background signals are high versus at late times when more signal is lost because of radioactive decay of the probe. Various currently developed PET radiotracers are summarized in Table 15.1.

A fluorinated analog of glucose, 2-\[^{18}\text{F}\]fluoro-2-deoxyglucose (FDG), is the predominant PET radiotracer in use today. FDG differs from glucose in that the hydroxy group typically found in the 2 position of the six-member ring has been replaced with fluorine-18. The nonfluorinated version of this molecule, 2-deoxyglucose (2DG), was originally intended for use as an anticancer agent, designed to take advantage of the well-known Warburg effect that postulates that cancer cells are critically dependent on glycolysis for energy production. Both 2DG and FDG are taken into cells via glucose transporters and undergo the initial step of glycolysis, phosphorylation of the CH2OH group at the 6 position of the ring by the enzyme hexokinase. However, because of the missing hydroxy group, the molecule cannot undergo isomerization in the next step of glycolysis, and is therefore trapped in the phosphorylated form, unable to exit the cell membrane because of the charge of the phosphate group. Both 2DG and FDG therefore accumulate within glycolytically active cells. 2DG failed as a cancer therapeutic because of the large doses required for efficacy and their associated toxicity; however, use of tracer doses of FDG as a PET-imaging agent has been enormously successful. A clinical dose of FDG for PET imaging is typically 15 mCi of radioactivity, with a mass dose of approximately 75 nmol. FDG has established uses in various oncologic imaging applications as discussed later in this chapter and shown in Figure 15.7. However, this imaging agent has several notable drawbacks, including accumulation in nonmalignant inflammatory processes as well as limited sensitivity for certain cancers such as prostate.

The search for improved PET radiotracers beyond FDG has taken several forms. In general, next generation PET radiotracers aim for improved physiologic specificity and/or improved clinical sensitivity. For example, there has been great effort to produce the so-called super FDG, that is, a radiotracer that has the broad spectrum of application of FDG but with improved sensitivity and specificity. Alternately, many groups have produced PET radiotracers that report specifically on a particular physiologic or molecular parameter, for example, PET probes that specifically bind to epidermal growth factor receptor (EGFR). These two goals are not necessarily mutually exclusive, but have been difficult to realize simultaneously with a single agent. One agent that has generated significant interest in the PET community is \[^{18}\text{F}\]fluorothymidine (FLT), a thymidine analog that is phosphorylated by thymidine kinases that are upregulated...
hypoxic cells. Molecules of this type include the 2-nitroimidazole derivatives pimonidazole and 2-(2-nitro-1H-imidazol-1-yl)-N-(2,2,3,3,3-[18F]pentafluoropropyl)-acetamide ([18F]EF5), both of which have been used as immunohistochemical probes for hypoxia. Labeling of these molecules with positron-emitting radioisotopes has allowed their use as PET hypoxia probes. Hypoxia radio-tracers of this type that have been studied clinically include [18F]fluoromisonidazole (FMISO), [18F]fluoroazomycin-arabinoside (FAZA), 2-(2-nitro-1H-imidazol-1-yl)-N-(2,2,3,3,3-[18F]pentafluoropropyl)-acetamide ([18F]EF5), both of which have been used as immunohistochemical probes for hypoxia. Labeling of these molecules with positron-emitting radioisotopes has allowed their use as PET hypoxia probes. Hypoxia radiotracers of this type that have been studied clinically include [18F]fluoromisonidazole (FMISO), [18F]fluoroazomycin-arabinoside (FAZA), 2-(2-nitro-1H-imidazol-1-yl)-N-(2,2,3,3,3-[18F]pentafluoropropyl)-acetamide ([18F]EF5), both of which have been used as immunohistochemical probes for hypoxia. Labeling of these molecules with positron-emitting radioisotopes has allowed their use as PET hypoxia probes. Hypoxia radiotracers of this type that have been studied clinically include [18F]fluoromisonidazole (FMISO), [18F]fluoroazomycin-arabinoside (FAZA), 2-(2-nitro-1H-imidazol-1-yl)-N-(2,2,3,3,3-[18F]pentafluoropropyl)-acetamide ([18F]EF5), both of which have been used as immunohistochemical probes for hypoxia. Labeling of these molecules with positron-emitting radioisotopes has allowed their use as PET hypoxia probes. Hypoxia radiotracers of this type that have been studied clinically include [18F]fluoromisonidazole (FMISO), [18F]fluoroazomycin-arabinoside (FAZA), 2-(2-nitro-1H-imidazol-1-yl)-N-(2,2,3,3,3-[18F]pentafluoropropyl)-acetamide ([18F]EF5), both of which have been used as immunohistochemical probes for hypoxia. Labeling of these molecules with positron-emitting radioisotopes has allowed their use as PET hypoxia probes. Hypoxia radiotracers of this type that have been studied clinically include [18F]fluoromisonidazole (FMISO), [18F]fluoroazomycin-arabinoside (FAZA), 2-(2-nitro-1H-imidazol-1-yl)-N-(2,2,3,3,3-[18F]pentafluoropropyl)-acetamide ([18F]EF5), both of which have been used as immunohistochemical probes for hypoxia. Labeling of these molecules with positron-emitting radioisotopes has allowed their use as PET hypoxia probes. Hypoxia radiotracers of this type that have been studied clinically include [18F]fluoromisonidazole (FMISO), [18F]fluoroazomycin-arabinoside (FAZA), 2-(2-nitro-1H-imidazol-1-yl)-N-(2,2,3,3,3-[18F]pentafluoropropyl)-acetamide ([18F]EF5), both of which have been used as immunohistochemical probes for hypoxia. Labeling of these molecules with positron-emitting radioisotopes has allowed their use as PET hypoxia probes. Hypoxia radiotracers of this type that have been studied clinically include [18F]fluoromisonidazole (FMISO), [18F]fluoroazomycin-arabinoside (FAZA), 2-(2-nitro-1H-imidazol-1-yl)-N-(2,2,3,3,3-[18F]pentafluoropropyl)-acetamide ([18F]EF5), both of which have been used as immunohistochemical probes for hypoxia. Labeling of these molecules with positron-emitting radioisotopes has allowed their use as PET hypoxia probes. Hypoxia radiotracers of this type that have been studied clinically include [18F]fluoromisonidazole (FMISO), [18F]fluoroazomycin-arabinoside (FAZA), 2-(2-nitro-1H-imidazol-1-yl)-N-(2,2,3,3,3-[18F]pentafluoropropyl)-acetamide ([18F]EF5), both of which have been used as immunohistochemical probes for hypoxia. Labeling of these molecules with positron-emitting radioisotopes has allowed their use as PET hypoxia probes. Hypoxia radiotracer
FIGURE 15.7 Metabolic and hypoxic PET imaging. The columns show PET scans acquired using [(^{18}F)fluoro-2-deoxyglucose (FDG) (left) and [(^{18}F)fluorooazomycin-arabinoside (FAZA) (right) from a patient with a primary head and neck cancer and locally involved lymph nodes. The increased signal-to-noise ratio and reduced background of FDG PET relative to FAZA PET are apparent. (Courtesy of Dr. R. Hicks.)
acquired using this agent have shown an association with patient outcome in initial clinical cancer imaging studies; however, the specificity of this agent for hypoxia has been called into question.

Although many PET radiotracers have been developed that employ some form of intracellular trapping to generate imaging signal, there are equally many PET agents that bind to specific molecules, commonly cell surface proteins. This approach has been used to produce PET agents for imaging angiogenesis by radiolabeling arginine-glycine-aspartic acid (RGD) peptides. This peptide sequence has been shown to bind to \( \alpha_\text{v}\beta_3 \) integrins expressed on endothelial cells of angiogenic vessels. Furthermore, various antibodies specific to oncologic targets have been radiolabeled and used as PET radiotracers, allowing the development of imaging approaches to function in conjunction with antibody-based therapies. The long circulation half-lives of antibodies generally necessitate that they be labeled with a long-lived positron-emitting isotope such as copper-64 or iodine-124, although antibody fragments have been engineered to tailor the pharmacokinetics of these agents toward those more suitable for imaging probes.

Finally, several PET agents have been developed for use as perfusion probes, including \([^{13}\text{N}]\text{NH}_3\) and \([^{15}\text{O}]\text{H}_2\text{O}\). Although the radiotracers discussed earlier are usually imaged at late time points after injection so as to minimize detection of nonspecific probe accumulation, perfusion PET probes are generally imaged at one or more time points immediately after injection, when the probe is still confined to the vasculature. This imaging strategy therefore suits the short half-lives of radioisotopes such as nitrogen-13 (9.96 minutes) and oxygen-15 (2.03 minutes). As in dynamic CT, dynamic PET images may be acquired and fit to compartmental models to extract estimates of vascular volume and/or perfusion from the measured time-activity data.

**Clinical Applications of Positron Emission Tomography**

**Tumor Staging**

The value of \(^{18}\text{F}\) FDG PET and PET/CT has been studied extensively for various cancers in several clinical scenarios. Currently, \(^{18}\text{F}\) FDG PET and PET/CT scanning is used routinely for the evaluation of several tumors that are typically avid for FDG. These malignancies are lung cancer, melanoma, breast cancer, esophageal and colorectal cancers, head and neck cancer, thyroid cancer, and lymphoma. Additionally, PET is well suited for detecting distant metastases because it is routinely performed as a whole body exam (from the base of the skull to the mid-thighs).

FDG PET has a sensitivity of greater than 96% for detecting and characterizing malignant lung nodules greater than 1 cm, with an overall specificity of 77%. In addition to this setting, FDG PET has been incorporated into the standard staging workup for cancers of the esophagus and head and neck, as well as for lymphoma. However, this modality has limited use for brain tumors because of the high uptake of FDG in normal brain, and for prostate cancer because of the low uptake of FDG by these tumors. FDG PET has little role for the initial screening and detection of melanoma because of limits of detection, but it is commonly used to stage metastasis.

**Radiotherapy Planning**

Although x-ray CT remains a crucial element of patient simulation for the planning of radiation therapies, the improved contrast between tumors and normal tissues offered by PET has encouraged its incorporation into radiotherapy planning. PET may improve radiation treatment outcomes both by allowing more sensitive detection of small tumor deposits in the volume receiving the highest radiation dose, or the gross tumor volume (GTV) in International Commission on Radiation Units and Measurements (ICRU) terminology, as well as by identifying occult or microscopic tumor spread, allowing inclusion of these regions in the GTV or the clinical target volume (CTV) where appropriate. The combination of PET and CT imaging technology in hybrid PET/CT scanners has further accelerated the consideration of this molecular imaging method in radiotherapy planning. Although the concept of defining biologically conformal radiation therapy (BCRT) using multiple molecular imaging measurements to map various tumor-associated functional and physiologic parameters has been forwarded, at present, researchers are addressing more immediate challenges such as how to best define a radiation target based on a single FDG PET image.

Definition of targets for radiation therapy involves clinical judgment and is prone to subjectivity. Despite tremendous efforts to standardize
FIGURE 15.8 Use of FDG PET/CT to assess treatment response. Fused PET/CT scans (CT: grayscale, PET: black-red-white pseudocolor) acquired from a patient with pancreatic cancer before (top) and 4 months after (bottom) treatment with a single radiation dose of 25 Gy and gemcitabine chemotherapy. The maximum standardized uptake value (SUV) within the primary lesion measured by FDG PET decreased from 16.4 prior to treatment to <2 at the follow-up time point. (From Koong AC, Christofferson E, Le Q-T, et al. Phase II study to assess the efficacy of conventionally fractionated radiotherapy followed by a stereotactic radiosurgery boost in patients with locally advanced pancreatic cancer. Int J Radiat Oncol Biol Phys. 2005;63:320–323, with permission.)
CONCLUSIONS

Imaging is now a required component of any oncology clinic. Although anatomic x-ray imaging is the dominant method in use in diagnosing and staging cancer, the emergence of molecular imaging as a clinically feasible means of mapping tissue function and physiology has clear potential to improve the clinical management of cancer patients. Molecular imaging, in its loosest definition, includes both functional imaging methods such as perfusion CT as well as truly molecular techniques such as PET. At present, the incorporation of molecular imaging into clinical practice is very much a work in progress. Many of the possible roles for molecular imaging in radiation therapy have been evaluated only in pilot studies, and larger clinical trials using standardized image acquisition and analysis methods as well as prospective treatment strategies are still forthcoming. Several studies examining how radiation treatment plans could be constructed incorporating functional imaging data have been performed but have yet to be delivered to actual patients. The term biologically conformal radiation therapy was coined by Ling et al. in 2000 as they developed a formalism for using imaging data to construct radiation treatments that are optimally tailored to a specific patient’s disease. Currently, though, we are caught between a “bottom-up” approach to this problem, in which imaging methods are developed to provide estimates of specific molecular and cellular parameters to apply a BCRT treatment model, and a “top-down” approach, in which existing imaging methods such as FDG PET are used to develop treatment strategies in the absence of a rigorous radiobiologic model at the molecular level. As is commonly the case, the most efficient path forward appears to be to combine these two paths for development. In the next 10 years, we can expect modest improvements in the performance of imaging methods such as CT and PET, in terms of quantitative accuracy, sensitivity, and spatial resolution. Many PET radiotracers are currently in evaluation in preclinical settings that may allow quantitation of specific parameters of radiobiologic interest, moving us closer to the bottom-up BCRT model as described earlier. However, several clinical trials evaluating molecular imaging-based diagnostic, staging, and treatment strategies are also in progress, ensuring we are most fully exploiting the clinical power of existing imaging modalities.
SUMMARY OF PERTINENT CONCLUSIONS

- Imaging of tumors using two-dimensional x-ray fluoroscopy and three-dimensional x-ray computed tomography is a standard procedure in cancer diagnosis, staging, and treatment monitoring.
- X-ray imaging involves the measurement of the x-ray absorption properties of an object, typically over either two (fluoroscopy) or three (tomography) dimensions.
- Modern clinical x-ray CT scanners can achieve imaging times as short as 0.5 seconds per frame and spatial resolutions as small as 250 μm.
- X-ray contrast agents have been formulated incorporating x-ray-attenuating materials such as barium, iodine, and gold.
- X-ray contrast agents are limited by the concentration of agent required to produce a measurable change in the x-ray attenuation of a tissue.
- Nanoparticle technology has revitalized the development of x-ray contrast agents by allowing the delivery of large concentrations of x-ray-attenuating materials in biologically tolerable formulations.
- Radiation doses delivered by beams of x-rays are enhanced when the absorbing material contains a strongly attenuating x-ray contrast material. This fact has been exploited for radiation therapy.
- X-ray CT is used to plan radiation therapies because the x-ray absorption measured in these images can be used to predict dose deposition by radiation treatment beams.
- Dynamic imaging after delivery of a contrast agent can be used to image vasculature and quantify perfusion and vessel permeability.
- X-ray CT is a clinically standard imaging method that is used to diagnose, stage, and monitor tumors from various sites.
- PET involves the localization and quantitation of exogenously delivered imaging agents that undergo nuclear decay through the ejection of a positron. Events are collected through the measurement of two antiparallel annihilation photons, which can then be traced along the line connecting them to find the source location.
- PET has been integrated with x-ray CT to produce hybrid PET/CT scanners, allowing the acquisition of functional PET information in the same examination as radiation treatment planning CT images.
- PET may be used to image any positron-emitting radiopharmaceutical delivered to a patient.
- PET radiotracers specific to glucose metabolism, cellular proliferation, hypoxia, angiogenesis, and various cellular proteins have been developed and used for imaging cancer.
- PET using the standard radiotracer FDG has been shown to enhance tumor staging and treatment response assessment for various tumor types.
- Methods for defining radiation target volumes based on PET images are currently in development.

BIBLIOGRAPHY


The purpose of this chapter is to review the doses involved and to estimate the associated risks in radiology, cardiology, and nuclear medicine. The bulk of radiation exposure is received by patients as part of their diagnosis or treatment, so there is a tangible medical benefit to balance against the risk; but medical radiation exposure is also conducted for medicolegal reasons and on volunteers (patients or healthy persons) for research purposes, and here, the risk–benefit equation is quite different. However, to put things in perspective, we first summarize the radiation doses from background sources that everyone receives naturally. This is usually regarded as an important benchmark, because life on earth has evolved with this continuous background radiation.

**DOSES FROM NATURAL BACKGROUND RADIATION**

Natural sources of radiation include cosmic rays from outer space and from the sun, terrestrial radiation from natural radioactive materials in the ground, and radiation from radionuclides naturally present in the body, ingested from food, or inhaled. The sources of natural background radiation are illustrated in Figure 16.1.

**Enhanced natural sources** are sources that are natural in origin but to which exposure is increased because of human activity (inadvertent or otherwise). Examples include air travel at high altitude, which increases cosmic ray levels, and movement of radionuclides on the ground in phosphate mining, which can increase the terrestrial component to persons living in houses built on waste landfills. Indoor radon exposure might be considered in some instances an enhanced natural source, inasmuch as it is not natural to live in an insulated house. In a sense, also, all operations associated with the nuclear fuel cycle, starting with mining, involve natural radionuclides, but these are more generally classified as a consequence of human activity.

**Cosmic Radiation**

Cosmic rays are made up of radiations originating from outside the solar system and from charged particles (largely protons) emanating from the surface of the sun. The intensity of cosmic rays arriving at the earth’s surface varies with both latitude and altitude above sea level.
States is illustrated in Figure 16.3 As would be expected, the highest doses are in the Rocky Mountains at high elevations.

Long flights at high altitudes involve some increased dose, too. For example, the extra dose from cosmic rays received by a passenger on a commercial flight flying from the United States to Europe is about 0.05 mSv. Flight crews on northerly routes accumulate larger doses than most radiology staff in hospitals; in fact, airline crews are already classified as radiation workers in Europe, but that is not yet the case in the United States.

**Natural Radioactivity in the Earth’s Crust**

Naturally occurring radioactive materials are widely distributed throughout the earth’s crust, and humans are exposed to the γ-rays from them. In the United States, there is a big variation between the Colorado plateau area, where the rocks and soil contain relatively more radioactive
that does little harm itself, because it is breathed in and breathed out again. However, in the confined space of an underground mine or the basement of a house, it decays with a 3-day half-life to form solid progeny that stick to dust or moisture particles and, if inhaled, become lodged on the surface of the bronchus or lung. Radon progeny emit $\alpha$-particles that, it is believed, are responsible for lung cancer. Radon levels in houses vary enormously, but the average concentration in the United States is about 37 Bq per cubic meter in aboveground living areas and much more in basements. It is a sobering thought that in an average home, in every cubic meter of air, 37 atoms of radon decay each second, producing radioactive progeny. Only the bronchi and lungs are irrigated by this source, but $\alpha$-particles are highly effective and have a radiation weighting factor of 20 (radiation weighting factor is explained in Chapters 7 and 17). This translates into an annual average effective dose of about 2 mSv. There is no question that radon is by far the largest component of natural background radia-

FIGURE 16.4 Color plot of the annual cosmic radiation doses (in microsievert) in North America. The variation with altitude is very clear, with the highest doses in the Rocky Mountains. (From Grasty RL, Lamarre JR. The annual effective dose from natural sources of ionizing radiation in Canada. Radiat Prot Dosim. 2004;108:215–226, with permission.)

Internal Exposure
Small traces of radioactive materials are normally present in the human body, ingested from the tiny quantities present in food or inhaled as airborne particles. Radioactive thorium, radium, and lead can be detected in most persons, but the amounts are small and variable, and the figure usually quoted for the average dose rate resulting from these deposits is less than 10 $\mu$Sv per year. Only radioactive potassium-40 makes an appreciable contribution to human exposure from ingestion. The dose rate is about 0.2 mSv per year, which cannot be ignored as a source of mutations in humans.

The biggest source of natural background radiation is radon gas, which seeps into the basements of houses from rocks underground. Radon, a decay product in the uranium series, is a noble gas that does little harm itself, because it is breathed in and breathed out again. However, in the confined space of an underground mine or the basement of a house, it decays with a 3-day half-life to form solid progeny that stick to dust or moisture particles and, if inhaled, become lodged on the surface of the bronchus or lung. Radon progeny emit $\alpha$-particles that, it is believed, are responsible for lung cancer. Radon levels in houses vary enormously, but the average concentration in the United States is about 37 Bq per cubic meter in aboveground living areas and much more in basements. It is a sobering thought that in an average home, in every cubic meter of air, 37 atoms of radon decay each second, producing radioactive progeny. Only the bronchi and lungs are irradiated by this source, but $\alpha$-particles are highly effective and have a radiation weighting factor of 20 (radiation weighting factor is explained in Chapters 7 and 17). This translates into an annual average effective dose of about 2 mSv. There is no question that radon is by far the largest component of natural background radia-
from background radiation. Undoubtedly, the highest natural background radiation is in Kerala, India, where more than 100,000 people receive an average annual dose of about 13 mSv, reaching a high in certain locations on the coast of 70 mSv. Many studies have been made of these human populations who have lived for many generations in areas of high natural background radiation. So far, no excess incidence of cancer or heritable anomalies has been observed that can reasonably be attributed to the radiation. Such studies, of course, are beset with difficulties.

**Areas of High Natural Background**

There are several inhabited areas of the world where background radiation is considerably higher than average because of radioactivity in rocks, soil, or in building materials from which houses are made. These areas are in Brazil, France, India, Niue Island (in the South Pacific), and Egypt.

In Brazil, some 30,000 people who live in coastal areas are exposed to dose rates of 5 mSv per year. About one-sixth of the population of France live in areas, largely in the Burgundy wine-growing district, in which the rocks are principally granite, and they receive 1.8 to 3.5 mSv per year from background radiation. Undoubtedly, the highest natural background radiation is in Kerala, India, where more than 100,000 people receive an average annual dose of about 13 mSv, reaching a high in certain locations on the coast of 70 mSv.

Many studies have been made of these human populations who have lived for many generations in areas of high natural background radiation. So far, no excess incidence of cancer or heritable anomalies has been observed that can reasonably be attributed to the radiation. Such studies, of course, are beset with difficulties.

**COMPARISON OF RADIATION DOSES FROM NATURAL SOURCES AND HUMAN ACTIVITIES**

In addition to natural background radiation, the human population is exposed to various sources of radiation resulting from human activities, as illustrated in Figure 16.5. Figure 16.6 shows the contribution of various sources of exposure to the total collective effective dose (1,870,000...
The annual effective dose resulting from human activities is now almost exactly equal to the total of all natural sources. Second, radon represents the largest source of natural background radiation, whereas medical radiation dominates the contribution from human activities.

It is important to realize that the populations exposed to these two major sources are not the same. The entire US population in age, gender, and health status is exposed to background radiation. By contrast, persons exposed to medical radiation are “patients,” and, as such, are skewed to older ages and to those with health problems. The principal exceptions are x-rays used for screening problems (e.g., mammography) and for trauma in children.

**FIGURE 16.5** The various sources of radiation resulting from human activity to which the human population is exposed. In developed countries, the effective dose is dominated by medical radiation.

**FIGURE 16.6** Percentage contribution of the various sources of exposure to the collective effective dose (1,870,000 person-Sv) and the average total effective dose per person in the US population (6.2 mSv) for 2006. Medical radiation and natural background radiation make almost equal contributions. (Data from National Council on Radiation Protection and Measurements. *I onizing Radiation Exposure of the Population of the United States. Report 160.* Bethesda, MD: NCRP; 2009.)
Apart from this important exception, the potential deleterious consequences of diagnostic radiology involve *stochastic effects*, that is, carcinogenesis and heritable effects. The characteristic of stochastic effects is that there is no threshold in dose; that is, there is no dose below which the effect does not occur, and the probability of carcinogenesis or heritable effects increases with dose. A stochastic effect may result from irradiation of one or a few cells, and the severity of the response is not dose related. As a consequence, *absorbed dose* to a limited portion of a person’s body does not provide by itself the overall perspective on risk associated with a given procedure.

*Effective dose* is a more relevant quantity; it takes into account the tissues and organs irradiated, as well as the dose involved. This is important because some tissues and organs are more susceptible than others to radiation. (Effective dose is discussed in detail in Chapter 17.) The technical definition of effective dose is the sum of the equivalent doses to each tissue and organ exposed multiplied by the appropriate tissue weighting factors (WT). What

### TABLE 16.1 Overview of the Practice of Radiology

<table>
<thead>
<tr>
<th>Country</th>
<th>Population $\times 10^6$</th>
<th>Mammography Units (per $10^6$ Population)</th>
<th>CT Scanners—Total (per $10^6$ Population)</th>
<th>Medical X-rays—Number of Annual Radiation Exams and Treatments $\times 10^6$ (per $10^6$ Population)</th>
<th>Physicians Conducting Radiology (per $10^6$ Population)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Canada</td>
<td>27.9</td>
<td>20.2</td>
<td>223 (8.0)</td>
<td>24.9 (0.89)</td>
<td>74</td>
</tr>
<tr>
<td>France</td>
<td>57.7</td>
<td>42.2</td>
<td>561 (9.7)</td>
<td>92.0 (1.59)</td>
<td>119</td>
</tr>
<tr>
<td>Germany</td>
<td>81.5</td>
<td>43.6</td>
<td>1,400 (17.2)</td>
<td>102.2 (1.25)</td>
<td>405</td>
</tr>
<tr>
<td>Japan</td>
<td>125.0</td>
<td>11.7</td>
<td>7,959 (63.7)</td>
<td>184.7 (1.48)</td>
<td>94</td>
</tr>
<tr>
<td>Sweden</td>
<td>8.8</td>
<td>19.3</td>
<td>115 (13.1)</td>
<td>5.0 (0.57)</td>
<td>125</td>
</tr>
<tr>
<td>United Kingdom</td>
<td>58.2</td>
<td>4.4</td>
<td>350 (6.0)</td>
<td>28.9 (0.50)</td>
<td>41</td>
</tr>
<tr>
<td>United States</td>
<td>260.0</td>
<td>38.6</td>
<td>6,800 (26.2)</td>
<td>250.0 (0.96)</td>
<td>92</td>
</tr>
</tbody>
</table>

Based on UNSCEAR 2000, which in turn is based on UNSCEAR surveys 1991–1996.
### TABLE 16.2 Entrance Skin Exposure and Absorbed Doses to Various Organs from Radiographic Studies in Adults

<table>
<thead>
<tr>
<th>Examination and View</th>
<th>Free-in-Air Exposure at Skin Entrance, mR</th>
<th>Active Bone Marrow</th>
<th>Thyroid</th>
<th>Breast</th>
<th>Lungs</th>
<th>Ovaries</th>
<th>Testes</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Chest</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Posteroanterior</td>
<td>20</td>
<td>0.02 (2)</td>
<td>0.01 (1)</td>
<td>0.01 (1)</td>
<td>0.07 (7)</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td>Lateral</td>
<td>65</td>
<td>0.02 (2)</td>
<td>0.07 (7)</td>
<td>0.15 (15)</td>
<td>0.12 (12)</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td>Series</td>
<td>—</td>
<td>0.04 (4)</td>
<td>0.07 (7)</td>
<td>0.16 (16)</td>
<td>0.19 (19)</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td><strong>Skull</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Anteroposterior</td>
<td>330</td>
<td>0.08 (8)</td>
<td>0.06 (6)</td>
<td>—</td>
<td>N</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td>Lateral</td>
<td>190</td>
<td>0.05 (5)</td>
<td>0.21 (21)</td>
<td>—</td>
<td>N</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td>Series</td>
<td>—</td>
<td>0.24 (24)</td>
<td>0.34 (34)</td>
<td>—</td>
<td>0.01 (1)</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td><strong>Cervical spine</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Anteroposterior</td>
<td>150</td>
<td>0.02 (2)</td>
<td>1.00 (100)</td>
<td>—</td>
<td>0.02 (2)</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td>Lateral</td>
<td>100</td>
<td>0.02 (2)</td>
<td>0.06 (6)</td>
<td>—</td>
<td>0.02 (2)</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td>Series</td>
<td>—</td>
<td>0.09 (9)</td>
<td>2.60 (260)</td>
<td>—</td>
<td>0.11 (11)</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td><strong>Thoracic spine</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Anteroposterior</td>
<td>280</td>
<td>0.05 (5)</td>
<td>0.25 (25)</td>
<td>0.95 (95)</td>
<td>0.35 (35)</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td>Lateral</td>
<td>630</td>
<td>0.12 (12)</td>
<td>0.05 (5)</td>
<td>0.05 (5)</td>
<td>0.75 (75)</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td>Series</td>
<td>—</td>
<td>0.17 (17)</td>
<td>0.30 (30)</td>
<td>1.00 (100)</td>
<td>1.10 (110)</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td><strong>Lumbar spine</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Anteroposterior</td>
<td>640</td>
<td>0.18 (18)</td>
<td>N</td>
<td>—</td>
<td>0.40 (40)</td>
<td>1.10 (110)</td>
<td>0.02 (2)</td>
</tr>
<tr>
<td>Lateral</td>
<td>2,300</td>
<td>0.44 (44)</td>
<td>N</td>
<td>—</td>
<td>0.30 (30)</td>
<td>0.90 (90)</td>
<td>0.02 (2)</td>
</tr>
<tr>
<td>Series</td>
<td>—</td>
<td>1.10 (110)</td>
<td>N</td>
<td>—</td>
<td>1.70 (170)</td>
<td>3.70 (370)</td>
<td>0.06 (6)</td>
</tr>
<tr>
<td><strong>Urography</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>KUB (anteroposterior)</td>
<td>600</td>
<td>0.20 (20)</td>
<td>N</td>
<td>—</td>
<td>0.07 (7)</td>
<td>1.30 (130)</td>
<td>0.10 (10)</td>
</tr>
<tr>
<td>Series</td>
<td>—</td>
<td>0.90 (90)</td>
<td>N</td>
<td>—</td>
<td>0.27 (27)</td>
<td>5.50 (550)</td>
<td>0.40 (40)</td>
</tr>
<tr>
<td>Series + 4 tomograms</td>
<td>—</td>
<td>1.70 (170)</td>
<td>N</td>
<td>—</td>
<td>0.54 (54)</td>
<td>6.50 (650)</td>
<td>0.50 (50)</td>
</tr>
<tr>
<td>Mammography</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Upper gastrointestinal series</td>
<td>3.00 (300)</td>
<td>0.03 (3)</td>
<td>0.50 (50)</td>
<td>1.00 (100)</td>
<td>12.00 (1,200)</td>
<td>0.80 (80)</td>
<td></td>
</tr>
<tr>
<td>Barium enema series</td>
<td>—</td>
<td>5.20 (520)</td>
<td>N</td>
<td>—</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*Values given are exposures and doses received by some patients at some facilities. Values can be much higher or lower depending on patient size, the technology employed, and the examination protocols established by the radiologist. Key: —, no estimate is made; N, negligible dose (<0.01 mGy).

*Two-view screening with film-screen grid.

this amounts to in simpler terms is that effective dose is the whole body dose of x-rays that would have to be delivered to produce the same stochastic risk as the partial body dose that actually was delivered. This quantity provides an easy assessment of overall risk and makes comparison of risks much simpler; for example, risk from a diagnostic examination is more readily compared with that from background radiation if effective dose is quoted. The unit of absorbed dose is the gray, whereas the unit of effective dose is the sievert. Most current reports in the literature use effective dose in discussing the potential health consequences of diagnostic radiology.

Last but not the least, the overall population impact of diagnostic radiology can be assessed in terms of the collective effective dose, the product of effective dose and the number of persons exposed. In this case, the unit is the person-sievert. This quantity is a surrogate for “harm” resulting from a given event involving radiation exposure. For example, the collective effective dose from the Chernobyl accident multiplied by the risk coefficient (5% per sievert for fatal cancer) gives an estimate of the number of cancer cases resulting from the accident, and is therefore a measure of the harm done. Later in this chapter, the consequences of the collective effective dose from diagnostic radiology will be discussed.

These three quantities—dose, effective dose, and collective effective dose—are discussed in turn. (In general, whenever the term dose is used alone, it refers to the absorbed dose.)

**Dose**

Table 16.2 is a summary of entrance skin exposures, as well as absorbed doses to various organs, characteristic of a representative sample of standard diagnostic procedures. The data do not contain any big surprises. As would be expected, radiographs of the lumbar spine, barium enema series, and upper gastrointestinal (GI) series involve substantial doses of radiation because of the need to penetrate these thick and dense regions of the body. These are the procedures, too, that inevitably lead to large gonadal doses.

The Nationwide Evaluation of X-ray Trends (NEXT) is a series of reports that give doses for several common examinations:

- The 1992 report summarized mammography x-ray data. It was shown that over the years, the mean glandular dose per examination has fallen and is now about 2 mGy while the image quality has improved. This is illustrated in Figure 16.7.
- The 1994 report summarized data for adult chest x-rays. The average entrance air kerma was 0.14 mGy at an average clinical kVp (kilovolt peak) of 101, with an average exposure time of 31 ms.
- The 1995 report referred to abdomen and lumbosacral spine x-ray data, which are shown in Table 16.3.
- The 1998 report focused on pediatric chest x-ray data. In contrast to the adult chest x-ray data of 1994, the average entrance air kerma was 0.05 mGy at an average kVp of 71, with an average exposure time of 12 ms.
- Dental x-ray data were the subject of the 1999 survey. The average entrance air kerma was 1.6 mGy with an average clinical kVp of 71.

**FIGURE 16.7** Graph showing how the mean glandular dose for mammography fell dramatically between 1970 and 1990 and is now about 1.4 mGy. Meanwhile, the image quality improved. Image quality is measured in terms of phantom score; a phantom contains fiber specks and spheres of different sizes, and the score indicates how many can be seen. (Adapted with permission from Nationwide Evaluation of X-ray Trends. Twenty-five years of NEXT. February 2003. Available at: http://www.crcpd.org/pubs/nextrifolds/25yrsofnext%20trifold-rev2-4-03.pdf.)
Some of the largest doses in diagnostic radiology are associated with fluoroscopy. In this case, the dose rate is greatest at the skin, where the x-ray beam first enters the patient. Dose rates from fluoroscopy from the NEXT 1996 upper GI fluoroscopy survey is shown in Table 16.4. Although dose rates in the literature are now reported in the new Système International d’Unités (SI) unit of milligray per minute, existing regulations still specify limits in terms of an exposure rate (roentgen per minute). The entrance exposure limit for standard operation of a fluoroscope is 10 R per minute. Some fluoroscopes are equipped with a high output or “boost” mode, and the limit for operation in this mode on state-of-the-art equipment is 20 R per minute. There is no limit on entrance exposure rate during any type of recorded fluoroscopy, such as cinefluorography or digital acquisitions.

A typical fluoroscopic entrance exposure rate for a medium-built person is approximately 3 R per minute (corresponding to an absorbed dose rate of about 30 mGy per minute). Much higher dose rates may be encountered during recorded interventional and cardiac catheterization studies, such as those that involve a series of multiple still-frame image acquisitions.

The number of CT scanners in clinical use has risen steadily over the years, and varies enormously between various countries with very different health care systems. This is evident from Table 16.5 taken from the UNSCEAR review in 2000, which is already way out of date. Figure 16.8 shows the dramatic increase in the number of

### Table 16.3: Abdomen and Lumbosacral Spine

<table>
<thead>
<tr>
<th></th>
<th>1995 Abdomen</th>
<th>1995 Lumbosacral Spine</th>
</tr>
</thead>
<tbody>
<tr>
<td>Entrance air kerma (mGy)</td>
<td>2.8</td>
<td>3.2</td>
</tr>
<tr>
<td>Clinical kVp</td>
<td>76</td>
<td>78</td>
</tr>
<tr>
<td>Exposure time (ms)</td>
<td>145</td>
<td>247</td>
</tr>
<tr>
<td>Percent using grids</td>
<td>97</td>
<td>96</td>
</tr>
<tr>
<td>Phantom film optical density</td>
<td>1.74</td>
<td>1.32</td>
</tr>
</tbody>
</table>

From the NEXT 1995 Abdomen and LS Spine X-ray Data Survey.

### Table 16.4: Dose Rates from Fluoroscopy

<table>
<thead>
<tr>
<th></th>
<th>1996 Gastrointestinal</th>
<th>1996 Cardiac Cath Labs</th>
<th>1996 C-Arm Units</th>
</tr>
</thead>
<tbody>
<tr>
<td>Entrance air kerma (mGy/min) d</td>
<td>45</td>
<td>38</td>
<td>22</td>
</tr>
<tr>
<td>Clinical kVp</td>
<td>99</td>
<td>82</td>
<td>78</td>
</tr>
<tr>
<td>Fluoroscope tube current (mA)</td>
<td>2.3</td>
<td>5.1</td>
<td>3.0</td>
</tr>
<tr>
<td>Air kerma rate with contrast b (mGy/min) e</td>
<td>67</td>
<td>71</td>
<td>41</td>
</tr>
<tr>
<td>Maximum air kerma rate e</td>
<td>70</td>
<td>74</td>
<td>44</td>
</tr>
</tbody>
</table>

* Determined at 1 cm of the table top and does not include contributions from over-table units.
* Copper is used to simulate the presence of barium contrast medium.

From the NEXT 1996 Upper GI Fluoroscopy Survey.
CT scans performed in the United States and in the United Kingdom in the past 25 years. By 2006, the number of CT scans performed in the United States reached 67 million, with about 10% of them in children. The rate of increase in the United Kingdom is just as rapid, but, taking into account the relative population sizes of the two countries, the frequency of use of CT (in terms of the number of scans/person/year) is about five times lower in the United Kingdom than in the United States. For CT scanning, organs in the beam can receive doses in the range of 10 to 100 mGy, but are usually in the range of 15 to 30 mGy for each single CT sequence. However, surveys emphasize the fact that doses can vary by a factor of 6 between different manufacturers, and by a factor of 5 for the same make scanner between different departments.

**Effective Dose**

Most recent surveys of diagnostic radiology quote effective dose, because this is related to the risk of stochastic effects, such as the induction of cancer or heritable effects. Table 16.6 shows representative values (together with range of values) of effective doses from representative radiologic procedures reported in the literature and summarized by Mettler and colleagues in 2008.

![Graph showing frequency of CT scans per year](image)

**FIGURE 16.8** Estimated number of CT scans performed annually in the United States and in the United Kingdom from 1980 until 2006. Note that although the rate of increase is similar in the United Kingdom to the United States, the number of scans per person is five times lower in the United Kingdom.
It is not difficult to understand why CT scans involve relatively larger effective doses, because larger volumes of tissue are exposed to higher doses than with plain x-rays. Table 16.7 shows representative values and ranges of values for effective doses from various CT procedures, again taken from the literature and summarized by Mettler and colleagues in 2008.

Pediatric CT scans are a special case. If the same parameters (kV and mA) are used for babies and small children as for adults, much larger doses and effective doses are received by the pediatric cases. This was common practice until 2001, when it was pointed out that children are at least 10 times as sensitive as adults to radiation-induced cancer. As a consequence, a major effort has been made by pediatric radiologists to tailor the appropriate parameters to the size of the person being scanned. This has led to a substantial dose reduction in children receiving CT scans. All major radiologic societies now support the “Image Gently” campaign to reduce doses in diagnostic radiology and to eliminate unnecessary procedures.

**Collective Effective Dose**

Next to be considered is the effect of diagnostic radiology on the population as a whole, rather than...
and used to quantify the harmful effects of radiation exposure to different parts of the body, taking into account the severity of the disease in terms of lethality, loss of quality of life, and years of life lost. Detriment includes a small component for heritable effects, a large component for fatal cancers, and an allowance for nonfatal cancers that, although they do not cause death, nevertheless affect quality of life. ICRP has suggested the detriment-adjusted risk coefficients for stochastic effects after exposure of the whole population to radiation at low dose rate to be 5.5% per sievert for cancer (fatal and nonlethal combined), and 0.2% per sievert for heritable effects, making a total of 5.7% per sievert. A collective effective dose of 899,000 person-Sv translates into 49,445 cases of cancer (fatal plus nonlethal), together with 1,798 cases of severe heritable effects resulting from 1 year’s practice of radiology. This is almost certainly a gross overestimate because most radiologic procedures are performed late in life or during a terminal illness.

Second, occupational exposure. The total collective effective dose in the United States in 2006 from occupational exposure to radiation was estimated to be 1,399 person-Sv. The two largest (and almost equal) contributors were medical and aviation workers, amounting to 77% of the total. However, the number of workers occupationally exposed in a medical setting

<table>
<thead>
<tr>
<th>Examination</th>
<th>Average Effective Dose (mSv)</th>
<th>Values Reported in Literature (mSv)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Head</td>
<td>2</td>
<td>0.9–4.0</td>
</tr>
<tr>
<td>Neck</td>
<td>3</td>
<td>—</td>
</tr>
<tr>
<td>Chest</td>
<td>7</td>
<td>4.0–18.0</td>
</tr>
<tr>
<td>Chest for pulmonary embolism</td>
<td>15</td>
<td>13–40</td>
</tr>
<tr>
<td>Abdomen</td>
<td>8</td>
<td>3.5–25</td>
</tr>
<tr>
<td>Pelvis</td>
<td>6</td>
<td>3.3–10</td>
</tr>
<tr>
<td>Three-phase liver study</td>
<td>15</td>
<td>—</td>
</tr>
<tr>
<td>Spine</td>
<td>6</td>
<td>1.5–10</td>
</tr>
<tr>
<td>Coronary angiography</td>
<td>16</td>
<td>5.0–32</td>
</tr>
<tr>
<td>Calcium scoring</td>
<td>3</td>
<td>1.0–12</td>
</tr>
<tr>
<td>Virtual colonoscopy</td>
<td>10</td>
<td>4.0–13.2</td>
</tr>
</tbody>
</table>

is much greater than in the aviation industry, so that the average effective dose to the person in aviation is four times larger than in medical.

**INTERVENTIONAL RADIOLOGY AND CARDIOLOGY**

Interventional fluoroscopy refers to any procedure in which the use or application of a medical device is fluoroscopically guided in the body and includes procedures that may be for diagnostic or therapeutic purposes. The past two decades have witnessed a major increase in such procedures in the United States. The collective effective dose has more than doubled in the past 10 years, making interventional radiology one of the fastest growing areas of medical radiation. These procedures include cardiac radiofrequency ablation, coronary artery angioplasty and stent placement, neuroembolization, and transjugular intrahepatic portosystemic shunt (TIPS) placement. Such procedures tend to be lengthy and involve fluoroscopy of a single area of the anatomy for a prolonged period—frequently for longer than 30 minutes and occasionally for over an hour. In addition, the need for multiple sequential treatment sessions can occur.

Procedures have evolved to include increasingly complex curative interventions that are associated with higher radiation exposures to both patients and health care workers. Angiography consists of inserting a catheter into a patient, guiding it along with the aid of fluoroscopy, and injecting contrast material into the vascular system. Stenosis, or blockage of one or more of these vessels, can lead to a myocardial infarction, but it can be visualized and treated with surgery or coronary angioplasty. Coronary angiography is a procedure that uses cineangiography in angulated projections, which can expose the operator to higher doses than when the x-ray equipment is used in the standard posteroanterior position.

Percutaneous transluminal coronary angioplasty is a therapeutic procedure to open blocked arteries by inflating a small balloon inside the artery, compressing and fracturing the obstruction, or using rotating blades to cut and remove the obstruction. It often requires the deployment of stents as well to maintain vessel patency. During conventional coronary angioplasty, prolonged fluoroscopy in severely angulated positions increases the dose to the operator and the patient. For coronary angioplasty, the overall potential radiation exposure to the operators is medium to high. The usual arterial accesses are the femoral artery in the groin and the brachial artery in the shoulder. Large doses are also involved in the placement of

---

**FIGURE 16.9** Percentage contribution of the various CT categories to the total collective effective dose from CT. Note that abdominal and pelvic scans account for the largest contribution. (Data from National Council on Radiation Protection and Measurements. *Ionizing Radiation Exposure of the Population of the United States. Report 160*. Bethesda, MD: NCRP; 2009.)
In addition, a large number of nonvascular interventional procedures using radiation are performed, such as the drainage of a blocked kidney or ablation of liver cancer.

**Patient Doses and Effective Doses**

Radiation doses received by patients from interventional radiology and cardiology are much higher than from general diagnostic radiology; so much so that there is a risk of deterministic effects, such as early or late skin damage. During these procedures, typical fluoroscopic absorbed dose rates to the skin can range from 20 to more than 50 mGy per minute. There are reports in the literature of several dozen cases of skin damage following fluoroscopically guided interventional procedures. Also frequently reported are cases showing an acute phase involving erythema and deep ulceration, followed by a late phase involving telangiectasia, and/or hyperpigmentation. Less frequent and following more than 2 hours of fluoroscopy, erythema, desquamation, and later a moist ulcer with tissue necrosis have been reported, requiring a skin graft. Table 16.8 is a

**TABLE 16.8 Potential Effects of Fluoroscopic Exposures on the Reaction of the Skin**

<table>
<thead>
<tr>
<th>Effect</th>
<th>Approximate Threshold Dose, Gy</th>
<th>Time of Onset</th>
</tr>
</thead>
<tbody>
<tr>
<td>Early transient erythema</td>
<td>2</td>
<td>2–24 h</td>
</tr>
<tr>
<td>Main erythema reaction</td>
<td>6</td>
<td>~1.5 wk</td>
</tr>
<tr>
<td>Temporary epilation</td>
<td>3</td>
<td>~3 wk</td>
</tr>
<tr>
<td>Permanent epilation</td>
<td>7</td>
<td>~3 wk</td>
</tr>
<tr>
<td>Dry desquamation</td>
<td>14</td>
<td>~4 wk</td>
</tr>
<tr>
<td>Moist desquamation</td>
<td>18</td>
<td>~4 wk</td>
</tr>
<tr>
<td>Secondary ulceration</td>
<td>24</td>
<td>&gt;6 wk</td>
</tr>
<tr>
<td>Late erythema</td>
<td>15</td>
<td>8–10 wk</td>
</tr>
<tr>
<td>Ischemic dermal necrosis</td>
<td>18</td>
<td>&gt;10 wk</td>
</tr>
<tr>
<td>Dermal atrophy (first phase)</td>
<td>10</td>
<td>&gt;12 wk</td>
</tr>
<tr>
<td>Dermal atrophy (second phase)</td>
<td>10</td>
<td>&gt;52 wk</td>
</tr>
<tr>
<td>Telangiectasia</td>
<td>10</td>
<td>&gt;52 wk</td>
</tr>
<tr>
<td>Delayed necrosis</td>
<td>12?</td>
<td>&gt;52 wk (related to trauma)</td>
</tr>
<tr>
<td>Skin cancer</td>
<td>Not known</td>
<td>&gt;15 y</td>
</tr>
</tbody>
</table>

Adapted from Wagner LK, Archer BR. *Minimizing Risks from Fluoroscopic X-rays*. 2nd ed. Houston, TX: Partners in Radiation Management; 1998, with permission, and modified by Hopewell (personal communication).
responding collective effective dose of 128,394 person-Sv, slightly more than half of this coming from cardiac procedures. Table 16.10 shows typical effective doses from various interventional radiology procedures, collected by Mettler and colleagues in 2008. Long interventional procedures can result in effective doses of 5 to 70 mSv, with TIPS being perhaps the highest. Because the patients are, in general, older and suffering from life-threatening medical conditions, the possibility of radiation-induced cancer 10, 20, or 30 years down the road is largely academic. The immediate threat of deterministic effects, however, is very real and can affect quality of life in a serious way.

**Dose to Personnel**

Physicians involved in cardiology, angiography, and fluoroscopically guided interventional work routinely receive radiation doses higher than any other staff in a medical facility and comparable to doses received in the nuclear industry (Fig. 16.10 and Table 16.11). Frequently, doses received by interventional radiologists are close to the annual dose limits, and there is also evidence that ocular cataracts are not uncommon. This is principally because of prolonged fluoroscopy.

### Table 16.9 Risks from Fluoroscopically Guided Interventional Procedures

<table>
<thead>
<tr>
<th>Benefit/risk ratio is high . . . . . . . BUT . . .</th>
</tr>
</thead>
<tbody>
<tr>
<td>■ 1.5% of patients develop an erythema.</td>
</tr>
<tr>
<td>■ 0.25% of patients develop some late skin damage.</td>
</tr>
<tr>
<td>■ 1 in 10,000 to 1 in 50,000 show severe late damage.</td>
</tr>
</tbody>
</table>

Courtesy of Dr. Lou Wagner.

### Table 16.10 Adult Effective Doses for Various Interventional Radiology Procedures

<table>
<thead>
<tr>
<th>Examination</th>
<th>Average Effective Dose (mSv)*</th>
<th>Values Reported in Literature (mSv)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Head and/or neck angiography</td>
<td>5</td>
<td>0.8–19.6</td>
</tr>
<tr>
<td>Coronary angiography (diagnostic)</td>
<td>7</td>
<td>2.0–15.8</td>
</tr>
<tr>
<td>Coronary percutaneous transluminal angioplasty, stent placement, or radiofrequency ablation</td>
<td>15</td>
<td>6.9–57</td>
</tr>
<tr>
<td>Thoracic angiography of pulmonary artery or aorta</td>
<td>5</td>
<td>4.1–9.0</td>
</tr>
<tr>
<td>Abdominal angiography or aortography</td>
<td>12</td>
<td>4.0–48.0</td>
</tr>
<tr>
<td>Transjugular intrahepatic portosystemic shunt placement</td>
<td>70</td>
<td>20–180</td>
</tr>
<tr>
<td>Pelvic vein embolization</td>
<td>60</td>
<td>44–78</td>
</tr>
</tbody>
</table>

*Values can vary markedly on the basis of the skill of the operator and the difficulty of the procedure.

NUCLEAR MEDICINE

Nuclear medicine is the medical specialty in which unsealed radionuclides, chemically manipulated to form radiopharmaceuticals, are used for diagnosis and therapy. Radiopharmaceuticals localize in various target tissues and organs, and although nuclear medicine images have less spatial and anatomic resolution than do radiographic or magnetic resonance images, they are better able to display physiology and metabolism.

Historical Perspective

The first person to suggest using radioactive isotopes to label compounds in biology and medicine was the Hungarian chemist Georg von Hevesy, whose work, beginning before World War II, earned him a Nobel Prize in 1943 (Fig. 16.11). The concept of using radioactive tracers in medicine could not be exploited until the means to produce artificial isotopes were readily available. The cyclotron was invented and developed by Ernest Lawrence in the 1930s, also leading to a Nobel Prize, and devices of this type have been used to produce short-lived isotopes and positron emitters (Fig. 16.12). Nuclear reactors were developed during World War II and are used to produce most medically used radioactive isotopes, all of which are electron or γ-ray emitters.

For the reasons mentioned earlier, nuclear medicine was a late starter compared with radiation therapy and x-ray diagnosis. Radiopharmaceuticals of adequate quality and consistency were not available until 1946, but since then, nuclear medicine has grown into a specialty in its own right and is now one of the most rapidly growing areas of radiation medicine largely because of nuclear cardiology. A broad array of pharmaceuticals, coupled with the development of sophisticated hardware, has made possible a widening diversity of applications. Positron emission tomography (PET) scanning has opened up a whole new area of rapid growth that is discussed later in this chapter.

Most nuclear medicine procedures are diagnostic examinations. Therapeutic nuclear medicine procedures represent only 1% to 2% of all radionuclide use, involving principally the treatment of hyperthyroidism, thyroid cancer, and bone metastases. Because of the high doses involved in therapy, the quantity “effective dose” is not appropriate to use for these procedures,
### Table 16.11 Estimated Dose to Staff during Typical Cardiac Studies

<table>
<thead>
<tr>
<th>Category of Staff</th>
<th>One Catheterization, mSv</th>
<th>One Angioplasty, mSv</th>
<th>One Pacemaker Implant (No Cine), mSv</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Weighted Surface Dose</td>
<td>Weighted Surface</td>
<td>Weighted Surface</td>
</tr>
<tr>
<td></td>
<td>with No Apron</td>
<td>Dose with Apron</td>
<td>Dose, with Apron</td>
</tr>
<tr>
<td></td>
<td>Hands</td>
<td>Eyes</td>
<td>Hands</td>
</tr>
<tr>
<td>Cardiologist</td>
<td>1.6</td>
<td>0.09</td>
<td>2.1</td>
</tr>
<tr>
<td>Nurse or anesthetist</td>
<td>0.3</td>
<td>0.02</td>
<td>0.4</td>
</tr>
<tr>
<td>Technologist</td>
<td>0.08</td>
<td>&lt;0.01</td>
<td>0.09</td>
</tr>
<tr>
<td>Nurse or anesthetist</td>
<td>0.04</td>
<td>&lt;0.01</td>
<td>—</td>
</tr>
</tbody>
</table>

FIGURE 16.11  The great Hungarian chemist Georg von Hevesy (1885–1966), whose work, beginning before World War II, earned him a Nobel Prize in 1943. He was the first to conceive of using radioactive isotopes to label compounds for biology and medicine. (Courtesy of the University of California Lawrence Berkeley Laboratory.)

and the therapeutic use of radionuclides will be discussed separately later in this chapter.

Effective Dose and Collective Effective Dose

The NCRP estimated that, in 2006, there were 19.7 million nuclear medicine procedures performed in the United States, with more than three-quarters of these procedures being performed in patients older than 45 years. The resulting collective effective dose was 22,000 person-Sv, up about 600% from the 1980s. Figure 16.13 shows the distribution of collective effective dose from nuclear medicine procedures, with nuclear cardiology accounting for 85% of the total. According to the 2008 UNSCEAR report, although the United States represents 5% of the world population, it receives 50% of the world's nuclear medicine procedures—a staggering figure. These estimates from NCRP involve patients, but do not include the many thousands of research procedures that use radionuclides because such data is hard to come by.

Table 16.12 shows representative effective doses from nuclear medicine procedures, collected from the literature by Mettler and colleagues in 2008. Some figures are as high as 40 mSv for a single procedure.

FIGURE 16.12  The concept of using radioactive isotopes as tracers in medicine was not fully explored until the invention of the cyclotron in 1931. Its inventor, Ernest O. Lawrence, is seen here (right) with his second cyclotron in 1934. Many short-lived isotopes that are positron emitters are made with a device of this sort. (Courtesy of the University of California Lawrence Berkeley Laboratory.)
The dramatic increase in the use of nuclear medicine (and the concomitant rise in the collective effective dose) is driven largely by cardiology. Numerous reviews have appeared in recent years attempting to document this trend. One such review (Bedetti et al., 2008) drew attention to the fact that many cardiac patients undergo as many as 36 or more examinations, with a median effective dose of more than 60 mSv. Of course, most of these patients are elderly and have a life-threatening illness, so that the risk of radiation-induced cancer 20 years down the road is largely academic. Fazel et al. (2009) performed a huge study of “nonelderly adults” and concluded that approximately 4 million Americans every year are subject to procedures that result in cumulative effective doses that exceed 20 mSv.

Principles in Nuclear Medicine

A wide range of radionuclides are used in diagnostic nuclear medicine that meet the necessary requirements for effective and efficient imaging. All are produced artificially, using four principal routes of manufacture: (1) cyclotron bombardment (producing, for example, gallium-67, indium-111, thallium-201, cobalt-57, iodine-123, carbon-11, oxygen-15, nitrogen-13, and fluorine-18), (2) reactor irradiation (e.g., chromium-51, selenium-75, iron-59, cobalt-58, iodine-125, and iodine-131), (3) fission products (e.g., iodine-131, xenon-133, and strontium-90), and (4) generators that provide secondary decay products from longer lived parent radionuclides. The most common example of the latter is the column generator incorporating molybdenum-99 for the provision of technetium-99m, which, because of its highly suitable physical characteristics for a wide range of applications, forms the basis for more than 80% of the radiopharmaceuticals used in nuclear medicine. Most technetium-99m generators use fission-produced molybdenum-99, although techniques of neutron irradiation could provide a viable alternative source of this important parent radionuclide. Other generators include those incorporating tin-113 (for the provision of indium-113m), rubidium-81 (for krypton-81m), and germanium-68 (for gallium-68).

The use of radiopharmaceuticals for diagnosis or therapy is based on the accumulation or concentration of the isotope in the organ of interest, referred to as the target organ. A radiopharmaceutical may have an affinity for a certain organ that is not necessarily the organ of interest, in which case this organ is termed a critical organ. Often the dose to a critical organ limits the amount of radioisotope that may be administered. The risk to which the patient is subjected is clearly a function of the doses received in all organs and is expressed in terms of the effective dose. The risk must be balanced against the expected advantages and benefits rendered by the procedure.
### Table 16.12 Effective Doses for Adults from Various Nuclear Medicine Examinations

<table>
<thead>
<tr>
<th>Examination</th>
<th>Effective Dose (mSv)</th>
<th>Administered Activity (MBq)</th>
<th>Effective Dose (mSv/MBq)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Brain (^{99m})Tc-HMPAO-exametazime</td>
<td>6.9</td>
<td>740</td>
<td>0.0093</td>
</tr>
<tr>
<td>Brain (^{99m})Tc-ECD-Neurolite</td>
<td>5.7</td>
<td>740</td>
<td>0.0077</td>
</tr>
<tr>
<td>Brain (^{99m})F-FDG</td>
<td>14.1</td>
<td>740</td>
<td>0.019</td>
</tr>
<tr>
<td>Thyroid scan (sodium iodine-123)</td>
<td>1.9</td>
<td>25</td>
<td>0.075 (15% uptake)</td>
</tr>
<tr>
<td>Thyroid scan (^{131m})Tc-pertechnetate</td>
<td>4.8</td>
<td>370</td>
<td>0.013</td>
</tr>
<tr>
<td>Parathyroid scan (^{99m})Tc-sestamibi</td>
<td>6.7</td>
<td>740</td>
<td>0.009</td>
</tr>
<tr>
<td>Cardiac stress-rest test</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(thallium-201 chloride)</td>
<td>40.7</td>
<td>185</td>
<td>0.22</td>
</tr>
<tr>
<td>Cardiac rest-stress test</td>
<td>9.4</td>
<td>1100</td>
<td>0.0085 (0.0079 stress, 0.0090 rest)</td>
</tr>
<tr>
<td>((^{99m})Tc-sestamibi 1-day protocol)</td>
<td>12.8</td>
<td>1500</td>
<td>0.0085 (0.0079 stress, 0.0090 rest)</td>
</tr>
<tr>
<td>Cardiac rest-stress test</td>
<td>11.4</td>
<td>1500</td>
<td>0.0076</td>
</tr>
<tr>
<td>((^{99m})Tc-sestamibi 2-day protocol)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cardiac ventriculography</td>
<td>7.8</td>
<td>1110</td>
<td>0.007</td>
</tr>
<tr>
<td>((^{99m})Tc-labeled red blood cells)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cardiac (^{99m})Tc-FDG</td>
<td>14.1</td>
<td>740</td>
<td>0.019</td>
</tr>
<tr>
<td>Lung perfusion (^{99m})Tc-MAA</td>
<td>2.0</td>
<td>185</td>
<td>0.011</td>
</tr>
<tr>
<td>Lung ventilation (xenon-133)</td>
<td>0.5</td>
<td>740</td>
<td>0.00074</td>
</tr>
<tr>
<td>Lung ventilation (^{99m})Tc-DTPA</td>
<td>0.2</td>
<td>1300</td>
<td>0.0049 (40 actually inhaled)</td>
</tr>
<tr>
<td>Liver-spleen (^{99m})Tc-sulfur colloid</td>
<td>2.1</td>
<td>222</td>
<td>0.0094</td>
</tr>
<tr>
<td>Biliary tract (^{99m})Tc-disofenin</td>
<td>3.1</td>
<td>185</td>
<td>0.017</td>
</tr>
<tr>
<td>Gastrointestinal bleeding</td>
<td>7.8</td>
<td>1110</td>
<td>0.007</td>
</tr>
<tr>
<td>((^{99m})Tc-labeled red blood cells)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Gastrointestinal emptying</td>
<td>0.4</td>
<td>14.8</td>
<td>0.024</td>
</tr>
<tr>
<td>((^{99m})Tc-labeled solids)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Renal (^{99m})Tc-DTPA</td>
<td>1.8</td>
<td>370</td>
<td>0.0049</td>
</tr>
<tr>
<td>Renal (^{99m})Tc-MAG3</td>
<td>2.6</td>
<td>370</td>
<td>0.007</td>
</tr>
<tr>
<td>Renal (^{99m})Tc-DMSA</td>
<td>3.3</td>
<td>370</td>
<td>0.0088</td>
</tr>
<tr>
<td>Renal (^{99m})Tc-glucoheptonate</td>
<td>2.0</td>
<td>370</td>
<td>0.0054</td>
</tr>
<tr>
<td>Bone (^{99m})Tc-MDP</td>
<td>6.3</td>
<td>1110</td>
<td>0.0057</td>
</tr>
<tr>
<td>Gallium-67 citrate</td>
<td>15</td>
<td>150</td>
<td>0.100</td>
</tr>
<tr>
<td>Pentreotide (^{111})In</td>
<td>12</td>
<td>222</td>
<td>0.054</td>
</tr>
<tr>
<td>White blood cells (^{99m})Tc</td>
<td>8.1</td>
<td>740</td>
<td>0.011</td>
</tr>
</tbody>
</table>

*(Continued)*
In addition to conventional planar imaging, techniques have also been developed to allow emission tomography that, like x-ray CT, can demonstrate internal structures or functional information from cross-sectional slices of the patient. Two basic modalities have evolved. First, there is single photon emission computed tomography (SPECT). This uses conventional γ-emitting radiopharmaceuticals and is often performed in combination with planar imaging. SPECT imaging requires a scanning system incorporating a circular array of detectors or, more often, a rotating γ-camera system with up to four detector heads. The second modality is the more specialized technique of PET. This is based on the simultaneous detection of the pairs of photons (511 keV) arising from positron annihilation and mostly uses the short-lived biologically active radionuclides oxygen-15, carbon-11, fluorine-18, and nitrogen-13. Dedicated PET scanners use a circular array of detectors, and PET scanning is now usually performed in conjunction with a CT scan, so that the PET images, displaying information about metabolic and physiologic function, can be fused with the CT images that depict anatomy.

In general, there are three doses of interest after the administration of a given amount of a radiopharmaceutical:

1. **Effective dose**, because this determines the risk of stochastic effects (cancer and heritable effects).
2. **Dose to the target organ**.
3. **Dose to the critical organ**, because this may be many times larger than the total body dose, and it is known that certain tissues are particularly susceptible to radiation-induced cancer.

### Positron Emission Tomography

The important and unique feature of PET studies is that they document physiologic abnormalities, or changes in metabolism, rather than simply alterations in anatomy.

The principle of PET imaging is that the scanner locates the tracer by detecting the colinear pairs of 0.511-MeV photons emitted if a positron annihilates after uniting with an electron. A positron is a particle with the same mass and magnitude of charge as an electron, except that the charge is positive. A positron cannot exist at rest if it has lost all its kinetic energy; it is electrostatically attracted to an electron with which it annihilates to produce two antiparallel 0.511-MeV photons. Radionuclides that emit positrons have excesses of protons in their nuclei and are produced by bombarding stable elements in a cyclotron. Positron emitters do not occur in nature.

Examples of radionuclides used for PET imaging include oxygen-15, carbon-11, and fluorine-18; these radionuclides have short half-lives of 2, 20, and 110 minutes, respectively, so that the PET facility is frequently close to the cyclotron that produces the radionuclides. The most commonly administered positron-emitting radionuclide is fluorine-18, which is used for the production of \(^{18}F\)-2-deoxy-2-fluoro-D-glucose, usually referred to as FDG. This material, used routinely in clinical care, highlights areas of metabolism and therefore

**TABLE 16.12 Effective Doses for Adults from Various Nuclear Medicine Examinations (Continued)**

<table>
<thead>
<tr>
<th>Examination</th>
<th>Effective Dose (mSv)</th>
<th>Administered Activity (MBq)</th>
<th>Effective Dose (mSv/MBq)</th>
</tr>
</thead>
<tbody>
<tr>
<td>White blood cells ((^{111})In)</td>
<td>6.7</td>
<td>18.5</td>
<td>0.360</td>
</tr>
<tr>
<td>Tumor ((^{11}F)-FDG)</td>
<td>14.1</td>
<td>740</td>
<td>0.019</td>
</tr>
</tbody>
</table>

\(^{1}DMSA\), dimercaptosuccinic acid; DTPA, diethylenetriaminepentaacetic acid; ECD, ethyl cysteinate dimer; \(^{18}F\), fluorine-18; FDG, fluorodeoxyglucose; HMPAO, hexamethylpropyleneamine oxime; \(^{111}\)In, indium-111; MAA, macroaggregated albumin; MAG3, mercaptoacetyltriglycine; MDP, methylene diphosphonate; \(^{99m}\)Te, technetium-99m.

\(^{2}\)Recommended ranges vary, although most laboratories tend to use the upper end of suggested ranges.

can be used to detect the metastatic spread of cancer. \(^{18}\text{F}\)-FDG has also found a use in the early diagnosis of Alzheimer disease through the direct visualization of amyloid plaques in the brain. Better imaging agents are available for this purpose, but they are based on carbon-11 which, because of its 20-minute half-life, cannot be used as widely as \(^{18}\text{F}\)-FDG with its longer (2 hour) half-life. Other metabolic radiopharmaceuticals are rapidly finding a place in clinical practice—for example, \(^{18}\text{F}\)-fluorothymidine (FLT) to map areas of rapid cell proliferation within tumors, or compounds such as \(^{18}\text{F}\)-fluoromisonidazole (FMISO), \(^{18}\text{F}\)- fluoroazomycin-arabinoside (FAZA), or 64-Cu-diacetyl-bis(N4-methylthiosemicarbazone) (64-Cu-ATSM) to highlight areas in tumors that are hypoxic. Many of the other positron-emitting radionuclides are used for the production of experimental compounds used in research studies. For example, oxygen-15 is used to label water for blood flow studies. In addition, there are more than a dozen radiopharmaceuticals labeled with carbon-11 that are used in brain research.

The doses delivered to patients or research subjects in PET studies are relatively low because of the short half-lives of the radionuclides, even though the administered amounts of radioactivity are high to allow rapid and detailed imaging.

Table 16.13 shows organ doses, as well as effective doses for studies with \(^{18}\text{F}\)-FDG and for \(^{15}\text{O}\)-H\(_2\)O. For procedures using \(^{18}\text{F}\)-FDG, the agent most commonly used in routine clinical care, the bladder wall receives the highest dose; the heart, brain, and kidney also receive relatively high absorbed doses. A typical administered activity may be 370 MBq, which results in an effective dose of about 11 mSv. This is equal to about 4 months of natural background radiation in the United States, and an associated risk of fatal radiation-induced cancer might be about 4 cases per 10,000. Of course, the PET procedure is nowadays accompanied by a CT scan, which further increases the effective dose.

PET technologists often have higher radiation exposures than other workers in nuclear medicine, and pharmacists at cyclotron facilities have even higher exposures, especially to their hands. This is a function of two factors: (1) the relatively high energy, and therefore penetrating nature, of the photons emitted by the radionuclides used (0.511 MeV); and (2) because of the short half-lives of the commonly used positron-emitting radionuclides, the large initial activities must be prepared so that a sufficient amount is left by the time the patient is imaged (this is especially true for oxygen-15, which has a half-life of only 2 minutes).

### Table 16.13: Organ Doses and Effective Doses for Positron Emission Tomography Compounds

<table>
<thead>
<tr>
<th></th>
<th>(^{18}\text{F})-FDG</th>
<th>(^{15}\text{O})-H(_2)O</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>mGy/MBq × 10(^{-2})</td>
<td>rad/mCi × 10(^{-2})</td>
</tr>
<tr>
<td>Brain</td>
<td>1.9</td>
<td>7.0</td>
</tr>
<tr>
<td>Heart wall</td>
<td>6.0</td>
<td>22.0</td>
</tr>
<tr>
<td>Kidneys</td>
<td>2.0</td>
<td>7.4</td>
</tr>
<tr>
<td>Ovaries</td>
<td>1.7</td>
<td>6.3</td>
</tr>
<tr>
<td>Red marrow</td>
<td>1.3</td>
<td>4.8</td>
</tr>
<tr>
<td>Spleen</td>
<td>3.7</td>
<td>14.0</td>
</tr>
<tr>
<td>Testes</td>
<td>1.3</td>
<td>4.8</td>
</tr>
<tr>
<td>Thyroid</td>
<td>1.0</td>
<td>3.9</td>
</tr>
<tr>
<td>Bladder wall</td>
<td>19.0</td>
<td>70</td>
</tr>
<tr>
<td>Effective dose</td>
<td>3.0</td>
<td>11.0</td>
</tr>
</tbody>
</table>

Data from the Oak Ridge Institute for Science and Education (ORISE).
The Therapeutic Use of Radionuclides

The most common form of nuclear medicine therapy is the use of radioactive iodine-131 for the treatment of hyperthyroidism. A therapeutic treatment with iodine-131 involves an absorbed dose to the thyroid gland that varies with the person and is very nonuniform within the tissue itself but is on the order of many tens of grays. In addition, there is a total body dose of typically 70 to 150 mGy, which results from the isotope circulating in the blood. Because radiation is known to be a potent carcinogen, the risk of leukemia or of thyroid cancer following iodine-131 therapy has been appreciated from the outset, but quite large studies have failed to show an excess. Pregnancy is, of course, a contraindication to the treatment of hyperthyroidism with iodine-131. Treatment of fertile women should be preceded by the taking of a careful history and a pregnancy test. Treatment should be delayed, if possible, to eliminate the potential effects during pregnancy.

The second most common form of therapy with unsealed radionuclides is the treatment of thyroid cancer. Following complete surgical removal of the cancer and the thyroid gland, radioactive iodine may be given to destroy any residual iodine-accumulating cancer cells that had spread to lymph nodes, lungs, or bone. Such treatments involve large doses of iodine-131 that may result in a total body dose of 0.5 to 1.0 Gy, which is sufficient to cause severe depression of the bone marrow.

The therapy of bone metastases. Several cancers, including prostate and breast, have a predilection for diffuse spread throughout the skeleton. There are several radiopharmaceuticals (such as strontium-89 chloride) that will localize in the metastatic lesions to provide palliation (but not cure).

Polycythemia vera is a relatively rare disease that is characterized by overproduction of red and white blood cells by the bone marrow. Phosphate-32 is given intravenously, which localizes in the bone so that the β-rays emitted result in a mild bone marrow suppression and reduction in the production of many blood elements.

Radioimmunotherapy uses radiolabeled antibodies directed against specific antigens. These agents can be used for the treatment of chemotherapy-resistant lymphomas. The antibodies are most commonly labeled with iodine-131 or yttrium-90 and injected intravenously in relatively large activities.

MEDICAL IRRADIATION OF CHILDREN AND PREGNANT WOMEN

Irradiation of Children

The hazards associated with medical radiation in children are basically the same as in adults—namely, cancer and heritable effects—except that the risks associated with a given absorbed dose of radiation are higher because of an increased sensitivity in younger persons. There is good evidence for this. The Japanese survivors of the atomic bomb attacks represent the most carefully studied human population exposed to radiation. There is a marked change in sensitivity to radiation-induced malignancies with increasing age, with young children being more radiosensitive than older adults by a factor of 10 to 15 (Chapter 10). Concern for possible heritable effects induced by radiation is likewise greater in children, because they have their entire reproductive lives ahead of them.

In pediatric nuclear medicine, as in pediatric radiology, the general principle is that radiation exposures should be kept at the lowest practical level. In each case, the expected benefit should exceed the risks clearly. Physicians and patients alike are much more cautious about nuclear medicine procedures in children than about diagnostic x-rays, even if the dose levels are similar.

Table 16.14 compares the effective dose in children of various ages with adults for various diagnostic nuclear medicine procedures. In general, effective doses are larger in young children for the same procedure, even if the administered activity is adjusted for body weight.

The implication of the review of carcinogenesis and heritable effects on humans (Chapters 10 and 11) is that any amount of radiation, no matter how small, has a deleterious effect. This conclusion is based on the assumption of a linear, nontreshold, dose-effect model that has been adopted by most standard-setting bodies as the most conservative basis for risk estimates. This philosophy requires that the physicians have some reasonable indications that the potential gain for the patient from the use of a procedure in nuclear medicine exceeds the risks.
<table>
<thead>
<tr>
<th>Radiopharmaceutical</th>
<th>Activity for Adult Patient, MBq</th>
<th>Effective Dose per Procedure by Patient Age$^a$ (mSv)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Adult 70 kg [1.0]</td>
<td>15-Year-Old 55 kg [0.9]</td>
</tr>
<tr>
<td>$^{99m}$Tc MAG3 (normal renal function)</td>
<td>100</td>
<td>0.7</td>
</tr>
<tr>
<td>$^{99m}$Tc MAG3 (abnormal renal function)</td>
<td>100</td>
<td>0.6</td>
</tr>
<tr>
<td>$^{99m}$Tc DTPA (normal renal function)</td>
<td>300</td>
<td>1.6</td>
</tr>
<tr>
<td>$^{99m}$Tc DTPA (abnormal renal function)</td>
<td>300</td>
<td>1.4</td>
</tr>
<tr>
<td>$^{99m}$Tc DMSA (normal renal function)</td>
<td>80</td>
<td>0.7</td>
</tr>
<tr>
<td>$^{99m}$Tc pertechnetate (no thyroid block)</td>
<td>80</td>
<td>1.0</td>
</tr>
<tr>
<td>$^{99m}$Tc IDA (normal biliary function)</td>
<td>150</td>
<td>2.3</td>
</tr>
<tr>
<td>$^{99m}$Tc HMPAO</td>
<td>500</td>
<td>4.7</td>
</tr>
<tr>
<td>$^{99m}$Tc leukocytes</td>
<td>200</td>
<td>2.2</td>
</tr>
<tr>
<td>$^{99m}$Tc erythrocytes</td>
<td>800</td>
<td>5.3</td>
</tr>
<tr>
<td>$^{99m}$Tc phosphates</td>
<td>600</td>
<td>3.6</td>
</tr>
<tr>
<td>$^{99m}$Tc MIBI (resting)</td>
<td>400</td>
<td>3.3</td>
</tr>
<tr>
<td>$^{201}$Tl chloride</td>
<td>80</td>
<td>20</td>
</tr>
<tr>
<td>$^{123}$I iodide (55% thyroid uptake)</td>
<td>20</td>
<td>7.2</td>
</tr>
<tr>
<td>$^{123}$I iodide (total thyroid block)</td>
<td>20</td>
<td>0.2</td>
</tr>
<tr>
<td>$^{125}$I MIBG (no impurity)</td>
<td>400</td>
<td>5.6</td>
</tr>
<tr>
<td>$^{67}$Ga citrate</td>
<td>150</td>
<td>15</td>
</tr>
</tbody>
</table>

*Figures in brackets are scaling factors for activity based on body weights shown. Doses are calculated using age-specific coefficients.

MAG3, mercaptoacetyltriglycine; DTPA, diethylenetriamine pentaacetic acid; DMSA, dimercaptosuccinic acid; IDA, iminodiacetic acid; HMPAO, hexamethylpropyleneamine oxime; MIBI, methoxy-isobutyl-isonitrile; MIBG, metaiodobenzylguanidine.

Based on UNSCEAR 2000.
Irradiation of Pregnant Women

The risks involved in exposure to radiation of the embryo or fetus are discussed in detail in Chapter 12. They may be summarized as follows:

1. For the first 10 days following conception (i.e., during preimplantation), the most significant effect of radiation may be to kill the embryo, leading to resorption.
2. Between 10 days and 8 weeks postconception (i.e., during organogenesis), the risks include congenital malformations and small head size, as well as carcinogenesis.
3. Between 8 and 15 weeks, and to a lesser extent 15 to 25 weeks, the risks include mental retardation, as well as small head size and carcinogenesis.
4. Beyond 25 weeks, the only risk of externally delivered diagnostic radiation is carcinogenesis, which is much reduced, compared with the risk during the first trimester.

Radiation-induced carcinogenesis is considered a stochastic effect; that is, there is no threshold and the risk increases with dose. One obstetric examination involving a few films may increase the relative risk of leukemia and childhood cancer by about 40%; however, because malignancies are relatively rare in children, the absolute risk is small. The other serious effects, such as mental retardation and congenital malformations, are considered deterministic; that is, there is a dose threshold of about 0.3 Gy. It is against the background of these possible risks that the irradiation of the pregnant or potentially pregnant woman must be considered.

Some radiologists instruct their technologists to ask all female patients about possible pregnancy before these women have abdominal or pelvic radiographic examinations. This would appear to be prudent and expedient. Indeed, if time permits, a pregnancy test might be desirable. If a woman requires an emergency radiologic examination, however, there should be no hesitation to do the study. The health of the woman is of primary importance, and if serious injury or illness is suspected, this takes priority in determining the need for a study. The risk to a possible conceptus must be weighed against the risks of not performing the examination.

The NCRP in 1977 recommended for pregnant women that if, in the best judgment of the attending physician, a diagnostic examination or nuclear medicine procedure at that time is deemed advisable to the medical well-being of the patient, it should be carried out without delay, with special efforts being made, however, to minimize the dose received by the lower abdomen (uterus).

In the case of nonemergency examinations, it sometimes may be prudent to consider delaying the proposed procedure. The physician contemplating the delay of a study on a woman early in pregnancy should consider the consequences in view of the possibility that the diagnostic examination might become necessary later in the pregnancy, when the risks are much greater. For example, during the first 10 days postconception, the risks are possible carcinogenesis and resorption of the embryo.

If the study is delayed, however, but becomes essential during the 8th through the 15th week, the risks include small head size and carcinogenesis; in addition, this is the peak of sensitivity to mental retardation. Delay compounds the problem. Conversely, if the patient is already in this peak period of radiosensitivity when a procedure is contemplated, then a delay until after the 25th week would be an advantage, because radiation risks during this period may be at their smallest.

The effects of radionuclides on the developing embryo or fetus have not been studied as extensively as the consequences of externally administered x-rays. The biologic effects may depend on many factors, including the chemical form of the isotope, the type and energy of the radiation emitted, whether the compounds containing the radioactivity cross the placenta, and whether they tend to be concentrated in specific target organs.

The metabolism of the radiopharmaceutical may cause high concentrations of the radionuclide in organs of a conceptus if the material crosses the placenta. This may result in dysfunctioning fetal organs. The classic example of this effect involves the uptake of iodine-131 in the thyroid of the developing embryo and fetus. Up to about 12 weeks, the fetal thyroid does not take up iodine. After this time, iodine concentrates in the fetal thyroid in amounts considerably greater than those in the maternal thyroid (Table 16.15). Several cases from the 1950s through the 1980s have documented the induction of hypothyroidism and cretinism from doses of iodine-131 to the fetal thyroid.

Although pregnant women receive diagnostic x-rays occasionally, it is rare for them to be given radioactive isotopes. In general, physicians and patients alike are much more wary about nuclear medicine procedures than about diagnostic x-rays, even if dose levels may be similar. Never
scans, (2) nuclear medicine (including nuclear cardiology), and (3) interventional radiology and cardiology. This is illustrated in Figure 16.14.

The countless millions of conventional radiographs, including chest x-rays and mammography, account for no more than 10% of the total collective effective dose.

**SUMMARY**

The three principal contributors to the collective effective dose from medical radiation are (1) CT scans, (2) nuclear medicine (including nuclear cardiology), and (3) interventional radiology and cardiology. This is illustrated in Figure 16.14.

The countless millions of conventional radiographs, including chest x-rays and mammography, account for no more than 10% of the total collective effective dose.

**TABLE 16.15** Thyroidal Radioiodine Dose to the Fetus

<table>
<thead>
<tr>
<th>Gestation Period</th>
<th>Fetal/Maternal Ratio (Thyroid Gland)</th>
<th>Dose to Fetal Thyroid, rad/µCi&lt;sup&gt;a&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>10–12 weeks</td>
<td>—</td>
<td>0.001 (precursors)</td>
</tr>
<tr>
<td>12–13 weeks</td>
<td>1.2</td>
<td>0.7</td>
</tr>
<tr>
<td>Second trimester</td>
<td>1.8</td>
<td>6</td>
</tr>
<tr>
<td>Third trimester</td>
<td>7.5</td>
<td>—</td>
</tr>
<tr>
<td>Birth imminent</td>
<td>—</td>
<td>8</td>
</tr>
</tbody>
</table>

<sup>a</sup>Rad/µCi of iodine-131 ingested by mother.

Courtesy of Dr. J. Keriakes, unpublished data.

**FIGURE 16.14** Contributions to the total collective effective dose from the medical uses of radiation (899,000 person-Sv). The biggest contributions come from CT, nuclear medicine, and interventional procedures, in that order. The millions of conventional procedures (mammograms, chest x-rays, etc.) account for only 10% of the total. (Data from National Council on Radiation Protection and Measurements. *Ionizing Radiation Exposure of the Population of the United States. Report 160*. Bethesda, MD: NCRP; 2009).
SUMMARY OF PERTINENT CONCLUSIONS

- Everyone is exposed to radiation from unperturbed natural sources, enhanced natural sources, and sources resulting from human activity, including medical x-rays.
- Natural background radiation comes from cosmic rays, terrestrial radiation from the earth’s crust, and inhaled or ingested radioactivity.
- Cosmic ray levels vary with altitude and latitude.
- Terrestrial radiation levels vary widely with locality.
- Radon and its progeny result in irradiation of lung tissue with α-particles; this is the largest source of natural radiation.
- Medical radiation is by far the largest source of radiation resulting from human activities.
- By the year 2006, the collective effective dose to the US population from medical radiation had increased by about 700% from the 1980s, and now approximately equals that from natural background radiation.
- The radiation doses involved in diagnostic radiology, except for interventional procedures, do not result in deterministic effects; the risks are stochastic effects (i.e., carcinogenesis and heritable effects).

Effective Dose and Cancer

- The cancer risk to a person is expressed in terms of the effective dose, the equivalent dose to the various organs and tissues exposed, multiplied by the appropriate tissue weighting factors (WT).
- The collective effective dose is the product of the effective dose and the number of persons exposed. It gives an indication of the harm or “detriment” to an exposed population.
- The collective effective dose to the US population in 2006 from medical radiation was estimated by NCRP to be 899,000 person-Sv, a 700% increase from the previous comprehensive survey conducted in the 1980s.
- The “detriment” from medical radiation may be calculated from the collective effective dose and the risk estimate of 5.5% per sievert for cancer (fatal and nonfatal) and 0.2% per sievert for serious heritable effects.
- Detriment includes an allowance for loss of quality of life as well as for death.
- The use of medical radiation for 1 year in the United States may benefit more than half of the population, but may result in about 50,000 cases of cancer (fatal plus nonfatal) and about 1,798 cases of serious heritable effects.

Interventional Procedures

- The past two decades have witnessed a major increase in high-dose fluoroscopically guided interventional procedures in medicine. Interventional radiology and cardiology now represent one of the three fastest growing areas of radiation medicine (together with CT and nuclear medicine).
- Radiation doses to patients from interventional radiology and cardiology are, in general, much higher than from general diagnostic radiology.
- NCRP estimated that, in 2006, there were 16.7 million interventional procedures performed in the United States, with a corresponding collective effective dose of 128,394 person-Sv.
- Long interventional procedures can result in effective doses to the individual of 5 to 70 mSv, with TIPS being one of the largest.
- Because patients undergoing interventional procedures are, in general, older and/or
In general, there are three doses of interest following a nuclear medicine procedure.
1. Dose to the target organ
2. Dose to the critical organ, which may be much higher
3. Effective dose, which determines the risk of stochastic effects (cancer and heritable effects)

PET displays physiologic and metabolic data that can be fused with the anatomic data from a CT scan.

The most commonly administered PET agent is $^{18}$F-FDG that highlights areas of metabolism and can detect cancer metastases.

A PET image with $^{18}$F-FDG results in an effective dose of about 11 mSv. However, this is usually accompanied by a CT scan, which increases the effective dose by an amount that depends on the site being imaged.

Other PET agents used in oncology can detect areas of rapid cellular proliferation in a tumor, or areas of hypoxia. Still other agents can be used to measure blood flow in cardiology.

The most common form of nuclear medicine therapy is the use of iodine-131 for the treatment of hyperthyroidism. The dose to the thyroid may be many tens of grays. In addition, there is a total body dose of 70 to 150 mGy, which is caused by the circulation of the radioactive iodine in the bloodstream. The second most common nuclear medicine therapy procedure is the treatment of thyroid cancer. Following surgical removal of the cancer and the thyroid gland, radioiodine may be used to destroy any residual iodine-accumulating cancer cells that had spread to lymph nodes, lung, or bone. The third most common therapy with radionuclides is the treatment of bony metastases. Radioimmunotherapy uses antibodies labeled with iodine-131 or yttrium-90, and injected against specific antigens to treat various malignancies.

### Medical Radiation of Children and Pregnant Women

Children are more sensitive than adults to radiation-induced malignancies by a factor of 10 to 15. Physicians and patients alike are more cautious about nuclear medicine with life-threatening illnesses, the possibility of radiation-induced malignancy 20 years down the road is largely academic. By contrast, doses are occasionally so high that deterministic effects can be an immediate problem.

There are reports in the literature of serious skin damage resulting from fluoroscopically guided interventional procedures, including erythema, deep ulceration, and occasionally necrosis requiring a skin graft. It is estimated that 1.5% of patients develop an erythema, 0.25% develop late skin damage, and between 1 in 10,000 and 1 in 50,000 develop severe late effects.

Doses to personnel involved in interventional procedures are among the highest recorded routinely in medical centers, and there is evidence that radiation-induced cataracts are not uncommon.

### Nuclear Medicine

Nuclear medicine is the medical specialty in which unsealed radionuclides, chemically manipulated to form radiopharmaceuticals, are used for diagnosis and therapy.

Most nuclear medicine procedures are diagnostic examinations. Therapeutic procedures account for only about 1% to 2%.

NCRP estimated that, in 2006, there were 19.7 million nuclear medicine procedures performed in the United States, three-quarters of them in persons older than 45 years. This resulted in a collective effective dose of 22,000 person-Sv, an increase of about 600% from the 1980s, with cardiology accounting for much of this trend.

UNSCEAR estimated that, with 5% of the world population, the United States was responsible for 50% of the nuclear medicine procedures performed in the world.

Cardiac patients frequently undergo as many as 36 examinations, with a median effective dose of greater than 60 mSv.

In a 2009 survey, Fazel et al. concluded that every year, about 4 million “nonelderly” adults are subject to nuclear cardiology procedures resulting in an effective dose in excess of 20 mSv.
procedures in children than about diagnostic x-rays, even when the radiation is comparable.
- It is rare for radionuclides to be administered to a pregnant woman; great care should be exercised if in an emergency situation, this is deemed to be essential. Particular care is needed if the radionuclide involved is iodine, because after the 12th week of gestation, the fetal thyroid avidly takes up iodine and can be seriously damaged.

**Summary**
- The three principal contributors to the collective effective dose from medical radiation are (1) CT scans, (2) nuclear medicine (including nuclear cardiology), and (3) interventional radiology and cardiology. The millions of conventional radiographs, including chest x-rays and mammography, account for no more than 10% of the total collective effective dose.

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CHAPTER 17

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THE ORIGINS OF RADIATION PROTECTION

At the Second International Congress of Radiology (ICR) in Stockholm in 1928, member countries were invited to send representatives to prepare x-ray protection recommendations. The British recommendations were adopted because they were most complete; guidelines on radiation protection had been set up in that country as early as 1915.

The 1928 Congress set up the International X-Ray and Radium Protection Committee, which, after World War II, was remodeled into two commissions that survive to this day:

The International Commission on Radiological Protection (ICRP)
The International Commission on Radiation Units and Measurements (ICRU)

The US representative to the 1928 Congress was Dr. Lauristor Taylor who brought back to the United States the agreed radiation protection criteria and set up a national committee, the Advisory Committee on X-Ray and Radium Protection, under the auspices of the Bureau of Standards, which was perceived to be a “neutral territory” by the various radiologic societies of the day; this committee operated until World War II. In 1946, it was renamed the National Council on Radiation Protection and Measurements (NCRP), eventually receiving a charter from Congress as an independent body to provide advice and recommendations on matters pertaining to radiation protection in the United States. NCRP reports still form the basis of radiation protection policy in the United States today, although legal responsibility for the implementation of radiation safety is variously in the hands of the Nuclear Regulatory Commission (NRC), the Department of Energy (DOE), and state or city bureaus of radiation control.

ORGANIZATIONS

The organization of radiation protection and the interrelation of various committees, whose reports are quoted, deserve a brief explanation.

First, there are the committees that summarize and analyze data and suggest risk estimates for radiation-induced cancer and heritable effects. At the international level, there is the United

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Nations Scientific Committee on the Effects of Atomic Radiation, usually known as UNSCEAR. This committee has wide international representation, being composed of scientists from 21 member states. Comprehensive reports appeared at intervals over the years since 1958, with the latest report in 2000. The United States committee is appointed by the National Academy of Sciences and is now known as the Biological Effects of Ionizing Radiations (BEIR) Committee. The first report appeared in 1956, when it was known as the Biological Effects of Atomic Radiations (BEAR) Committee. Subsequent comprehensive reports appeared in 1972 (BEIR II), 1980 (BEIR III), 1990 (BEIR V), and 2006 (BEIR VII). BEIR VI, entitled The Health Effects of Exposure to Indoor Radon, appeared in 1999.

These committees are “scholarly” committees in the sense that if information is not available on a particular topic, they do not feel compelled to make a recommendation. Because they do not serve an immediate pragmatic aim, they are not obliged to make a “best guess” estimate if data are uncertain.

Second, there are the committees that formulate the concepts for use in radiation protection and recommend maximum permissible levels. These committees serve more pragmatic aims and, therefore, must make best estimates even if good data are unavailable. At the international level, there is the ICRP, which, together with ICRU, was established in 1928 after a decision by the Second ICR, as mentioned earlier. In 1950, the ICRP was restructured and given its present name. The ICRP often takes the lead in formulating concepts in radiation protection and in recommending dose limits. As an international body, it has no jurisdiction over anyone and can do no more than recommend; it has established considerable credibility, however, and its views carry great weight. Its most recent comprehensive report is ICRP Publication No. 103, published in 2007. The United Kingdom, most of Europe, and Canada follow ICRP recommendations. In the United States, there is the NCRP (also mentioned earlier) chartered by Congress to be an “impartial” watchdog and consisting of 100 experts from the radiation sciences—who are, therefore, not impartial at all. The NCRP often, but not always, follows the lead of ICRP. Their most recent comprehensive report on dose limits (NCRP Report No. 116,
Chapter 17 • Radiation Protection

In the United States, the Environmental Protection Agency (EPA) has the responsibility for providing guidance to federal agencies; it is the EPA that sets, for example, the action level for radon. Each state can formulate its own regulations for x-rays and radiations produced by sources other than reactors. In agreement states, the NRC formulates rules for by-product materials from reactors. Figure 17.1 shows the agreement status as of 2009. In other states, this responsibility falls on the U.S. Department of Labor Occupational Safety and Health Administration (OSHA). The DOE is responsible for radiation safety regulations at all of its facilities operated by contractors. Up to the present, the various regulating bodies in the United States have accepted, endorsed, and used the reports issued by the NCRP, but they are not obliged to do so, and they are often slow to adopt the latest reports.

QUANTITIES AND UNITS

Dose
The quantity used to measure the “amount” of ionizing radiation is the absorbed dose, usually termed simply as dose. This is defined as the energy absorbed per unit mass, and its unit is joules per kilogram, which is given a special name, the gray (Gy), named after the British physicist who contributed to the development of ionization chamber theory. The unit used in the past was the radiation absorbed dose (rad), defined as an energy absorption of 100 erg/g. Consequently, 1 Gy equals 100 rad.

Radiation Weighting Factor
The probability of a stochastic effect, such as the induction of cancer or of heritable events, depends not only on the dose, but also on the type and energy of the radiation; that is, some radiations are biologically more effective for a given dose than others. This is taken into account by weighting the absorbed dose by a factor related to the quality of the radiation. A radiation weighting factor (WR) is a dimensionless multiplier used to place biologic effects (risks) from exposure to different types of radiation on a common scale. The WRs are chosen by the ICRP as representative of relative biologic effectiveness (RBE) applicable to low doses and low dose rates (LDR), and for biologic end points relevant to stochastic late effects. They can be traced ultimately to experimentally determined RBE values, but a large judgmental factor is involved in their choice. The weighting factors recommended by the ICRP for photons, electrons, protons and α-particles, fission fragments, heavy ions are shown in Table 17.1. For neutrons, a continuous curve as a function of neutron energy is recommended (Fig. 17.2) with the most biologically effective neutrons having a WR of 20.

Equivalent Dose
In radiologic protection, the equivalent dose is the product of the absorbed dose averaged over the tissue or organ and the WR selected for the type and energy of radiation involved. Thus:

$$\text{Equivalent dose} = \text{absorbed dose} \times \text{WR}$$

If absorbed dose is measured in Gy, the equivalent dose is measured in sievert (Sv), named after the Swedish physicist who designed early ionization chambers. Although 1 Gy of neutrons does not produce the same biologic effect as 1 Gy of x-rays, 1 Sv of either neutrons or x-rays

<table>
<thead>
<tr>
<th>Radiation Type</th>
<th>Radiation Weighting Factors</th>
</tr>
</thead>
<tbody>
<tr>
<td>Photons</td>
<td>1</td>
</tr>
<tr>
<td>Electrons and muons</td>
<td>1</td>
</tr>
<tr>
<td>Protons and charged pions</td>
<td>2</td>
</tr>
<tr>
<td>α-particles, fission fragments, heavy ions</td>
<td>20</td>
</tr>
<tr>
<td>Neutrons</td>
<td>A continuous curve as a function of neutron energy</td>
</tr>
</tbody>
</table>

*All values relate to the radiation incident on the body or, for internal sources, emitted from the source.

From ICRP 2007.
tissue weighting factor \((W_T)\), which represents the relative contribution of each tissue or organ to the total detriment resulting from uniform irradiation of the whole body. Table 17.2 lists the \(W_T\) values recommended by the ICRP in 2007.

The sum of all of the weighted equivalent doses in all the tissues or organs irradiated is called the **effective dose**, which is expressed by the formula

\[
\text{Effective dose} = \sum \text{absorbed dose} \times W_R \times W_T
\]

for all tissues or organs exposed. Effective dose is in principle, as well as in practice, a nonmeasurable quantity.

### Effective Dose

If the body is uniformly irradiated, the probability of the occurrence of stochastic effects (cancer and hereditary effects) is assumed to be proportional to the equivalent dose, and the risk can be represented by a single value. In fact, truly uniform total body exposures are rare, particularly if irradiation is from radionuclides deposited in tissues and organs. Sometimes, equivalent doses to various tissues differ substantially and it is well established that different tissues vary in their sensitivities to radiation-induced stochastic effects. For example, it is difficult to produce heritable effects by irradiation of the head or hands. On the other hand, the thyroid and breast appear to be particularly susceptible to radiation-induced cancer. To deal with this situation, the ICRP introduced the concept of the **tissue weighting factor** \((W_T)\), which represents the relative contribution of each tissue or organ to the total detriment resulting from uniform irradiation of the whole body. Table 17.2 lists the \(W_T\) values recommended by the ICRP in 2007.

The sum of all of the weighted equivalent doses in all the tissues or organs irradiated is called the **effective dose**, which is expressed by the formula

\[
\text{Effective dose} = \sum \text{absorbed dose} \times W_R \times W_T
\]

for all tissues or organs exposed. Effective dose is in principle, as well as in practice, a nonmeasurable quantity.

### Committed Equivalent Dose

In the case of external irradiation, the absorbed dose is delivered at the time of exposure; but for irradiation from internally deposited radionuclides, the total absorbed dose is distributed over time as well as to different tissues in the body. The dose rate falls off depending on the physical and biologic half-lives of the radionuclide.

To take into account the varying time distributions of dose delivery, the ICRP defined the **committed equivalent dose** as the integral over 50 years of the equivalent dose after intake of a radionuclide. This time was chosen to correspond to the working life of a person. For radionuclides with effective half-lives of up to about 3 months, the committed equivalent dose is essentially equal to the annual equivalent dose in the year of intake; but for radionuclides with longer effective half-lives, it is greater because it reflects the dose that will accrue over future years.
dose over the entire population out to a period of 50 years is called the **collective committed effective dose**.

These collective quantities can be thought of as representing the total consequences of exposure of a population or group and they can be thought of as surrogates for “harm.” For example, the annual collective effective dose to the US population from medical radiation is about 899,000 person-Sv. Such collective quantities are much beloved by the bureaucrats because they make it possible to compare different activities or accidents, inasmuch as each can be described by a single number. The danger is that the next step is to convert the collective dose into the number of cancers or heritable effects produced, which, of course, assumes proportionality between dose and biologic effect, which is seldom true. The quantities certainly are used widely to give a rough guide to the probability of cancer and heritable effects in a population, and in particular, they can be used to compare the approximate impact of different types of radiation accidents in terms of several health effects that might arise in that population.

**Summary of Quantities and Units**

Table 17.3 is a summary of the quantities and units that have been described here, showing how they build logically on one another. If on reading this section the reader gains the impression that the bureaucrats have taken over, it is because they have—at least in the field of...
The fundamental aim of radiation protection has been summed up by the ICRP as follows:

The primary aim of radiologic protection is to provide an appropriate standard of protection for man without unduly limiting the beneficial actions giving rise to radiation exposure. This aim cannot be achieved based on scientific concepts alone. All those concerned with radiologic protection have to make value judgments about the relative importance of different kinds of risk and about the balancing of risks and benefits. In this, they are no different from those working in other fields concerned with the control of hazards.

As stated by the NCRP, the objectives of radiation protection are the following:

1. To prevent clinically significant radiation-induced deterministic effects by adhering to dose limits that are below the apparent or practical threshold, and
2. To limit the risk of stochastic effects (cancer and heritable effects) to a reasonable level.

### AIMS AND OBJECTIVES OF RADIATION PROTECTION

Table 17.3: Quantities and Units Used in Radiation Protection

<table>
<thead>
<tr>
<th>Quantity</th>
<th>Definition</th>
<th>Unit</th>
</tr>
</thead>
<tbody>
<tr>
<td>Absorbed dose</td>
<td>Energy per unit mass</td>
<td>Gray</td>
</tr>
<tr>
<td>For individuals</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Equivalent dose</td>
<td>Average dose × radiation weighting factor</td>
<td>Sievert</td>
</tr>
<tr>
<td>(radiation weighted dose)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Effective dose</td>
<td>Sum of equivalent doses to organs and tissues exposed, each multiplied by</td>
<td>Sievert</td>
</tr>
<tr>
<td></td>
<td>the appropriate tissue weighting factor</td>
<td></td>
</tr>
<tr>
<td>Committed equivalent dose</td>
<td>Equivalent dose integrated over 50 years (relevant to incorporated</td>
<td>Sievert</td>
</tr>
<tr>
<td></td>
<td>radionuclides)</td>
<td></td>
</tr>
<tr>
<td>Committed effective dose</td>
<td>Effective dose integrated over 50 years (relevant to incorporated radionuclides)</td>
<td>Sievert</td>
</tr>
<tr>
<td>For populations</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Collective effective dose</td>
<td>Product of the average effective dose and the number of individuals exposed</td>
<td>Person-sievert</td>
</tr>
<tr>
<td>Collective committed</td>
<td>Integration of the collective dose over 50 years (relevant to incorporated</td>
<td>Person-sievert</td>
</tr>
<tr>
<td>effective dose</td>
<td>radionuclides)</td>
<td></td>
</tr>
</tbody>
</table>

Radiation protection. An elaborate set of definitions has been produced based on the assumption of linearity between dose and risk. The whole business needs to be taken with a generous grain of salt because it is like a house of cards, based on somewhat shaky premises.

The concept of collective effective dose does allow a rough and quick estimate to be made of the potential health hazards to a population from, for example, an accidental release of radioactivity from a nuclear reactor. It must be emphasized again that these concepts can be used only under conditions in which it is reasonable to assume linearity between risk and dose—that is, that risks are directly proportional to the summation of doses from different sources. Exposures that are within the administratively allowed dose limits may cause an increased incidence of stochastic effects, such as cancer and heritable effects, but are much below the thresholds for early deterministic effects. In the case of larger accidental releases in which doses to some people might be high enough to exceed these thresholds to the point of causing early death, collective effective dose is an inappropriate quantity.
in relation to societal needs, values, and benefits gained.

The difference in shape of the dose–response relationships for deterministic and stochastic effects is illustrated in Figure 17.3. The objectives of radiation protection can be achieved by reducing all exposure to as low as reasonably achievable (ALARA) and by applying dose limits for controlling occupational and general public exposures. For radiation protection purposes, it is assumed that the risk of stochastic effects is strictly proportional to dose without threshold throughout the range of dose and dose rates of importance in radiation protection. Furthermore, the probability of response (risk) is assumed to accumulate linearly with dose. This is not true at higher doses characteristic of radiation accidents in which more complex (nonlinear) dose–risk relationships may apply.

Given these assumptions, any selected dose limit has an associated level of risk. Consequently, it is necessary to justify any use of radiation in terms of a benefit to a person or to society.

Justification of exposure is one of the basic principles of radiation protection. The concept was described in 1977 by the ICRP: A practice involving exposure to radiation should produce sufficient benefit to the exposed individual or to society to offset the radiation detriment it causes. This concept is sometimes difficult to put into practice in various situations in which individuals are exposed as follows:

1. In the case of patients, the diagnostic or therapeutic benefit should outweigh the risk of detriment.
2. In the case of occupational exposure, the radiation risk must be added to and compared with other risks in the workplace.
3. The most difficult situation is exposure for the sake of research, where volunteer subjects may fall into one of three categories: Patients who may benefit, patients who may receive no benefit, and healthy volunteers. In cases in which the individual receives no benefit, the benefit to society must outweigh the risks.

### BASIS FOR EXPOSURE LIMITS

Exposure limits have changed over the years in step with evolving information about the biologic effects of radiation and with changes in the

![Deterministic and Stochastic Effects](image-url)
social philosophy within which recommended exposure limits are developed.

In the 1930s, the concept of a **tolerance dose** was used, a dose to which workers could be exposed continuously without any evident deleterious acute effects such as erythema of the skin.

By the early 1950s, the emphasis had shifted to late effects. The **maximum permissible dose** (MPD) was designed to ensure that the probability of the occurrence of injuries was so low that the risk would be readily acceptable to the average person. At about that time, based on the results of genetic studies in *Drosophila* and mice, the occupational limit was reduced substantially and a limit for exposure of the public introduced. Subsequently, the heritable effects were found to be smaller, and cancer risks larger than were thought at the time.

By the 1980s, the NCRP was comparing the probability of radiation-induced cancer death in radiation workers with annual accidental mortality rates in “safe” industries. Exposure standards, therefore, are necessarily based partly on observed effects, but with a great deal of judgment involved.

Earlier chapters described the deleterious effects of radiation in terms of heritable effects, carcinogenesis, and effects on the developing embryo and fetus. The risk estimates derived are summarized in Table 17.4. By far, the largest risk estimate is 40% per Sv for severe mental retardation for the most sensitive period of gestation and above a threshold of at least 0.3 Gy. Next comes carcinogenesis as 5% per Sv, corresponding to exposure of the general population to low doses and dose rates. Last are heritable effects, lowered in 2004 by the ICRP to 0.2% per Sv for the general population.

### LIMITS FOR OCCUPATIONAL EXPOSURE

The NCRP recommends the limits described in the following sections (and summarized in Table 17.5). These limits do not include natural background radiation or radiation for medical purposes.

**Stochastic Effects**

1. No occupational exposure should be permitted until the age of 18 years.
2. The effective dose in any year should not exceed 50 mSv (5 rem).
3. The individual worker’s lifetime effective dose should not exceed age in years $/10000$ mSv.

These limits apply to the sum of the effective dose from external radiation and the committed effective dose from internal exposures.

**Deterministic Effects**

1. 150 mSv per year for the lens of the eye.
2. 500 mSv per year for localized areas of the skin and the hands and feet.

These additional limits for deterministic effects are required because the weighting factors for, for example, the hands and the feet, are so small that huge doses could be given before cancer induction became a problem. Other deterministic effects are limiting at lower doses.

### AS LOW AS REASONABLY ACHIEVABLE

The dose limits referred to previously are all upper limits and subject to the concept of ALARA. The recommendation that standard setting committees would like to make for personnel protection is zero exposure. This is not feasible, however, if society is to realize the enormous benefits derived from the uses of radiations and radioactive materials.
exposure of personnel by a given amount? As a rule of thumb in the nuclear power industry in the United States, ALARA has a cash value of about $1,000 per 10 mSv. If the exposure of one person to 10 mSv can be avoided by the expenditure of this amount of money, it is considered reasonable. If the cost is more, it is considered unreasonable, and the exposure is allowed. However, the $1,000 per 10 mSv figure applies specifically to low-dose levels. At higher dose levels at which the accumulation of an additional exposure may threaten a worker’s job by exceeding the lifetime dose limit, then the cash value of avoiding a 10 mSv exposure

Radiation is potentially harmful, and exposure to it should be monitored continually and controlled. No unnecessary exposure should be allowed. Equipment and facilities should be designed so that exposure of the personnel and the public is kept to a minimum and not up to a standard. No exposure at all should be permitted without considering the benefits that may be derived from that exposure and the relative risks of alternative approaches.

Of course, the ultimate problem is determining what is “reasonable.” There is also the question: How much expense is justified to reduce the exposure of personnel by a given amount? As a rule of thumb in the nuclear power industry in the United States, ALARA has a cash value of about $1,000 per 10 mSv. If the exposure of one person to 10 mSv can be avoided by the expenditure of this amount of money, it is considered reasonable. If the cost is more, it is considered unreasonable, and the exposure is allowed. However, the $1,000 per 10 mSv figure applies specifically to low-dose levels. At higher dose levels at which the accumulation of an additional exposure may threaten a worker’s job by exceeding the lifetime dose limit, then the cash value of avoiding a 10 mSv exposure

<table>
<thead>
<tr>
<th>NCRP</th>
<th>ICRP (If Different)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Occupational Exposure:</strong></td>
<td></td>
</tr>
<tr>
<td>Stochastic effects: effective dose limits</td>
<td></td>
</tr>
<tr>
<td>Cumulative</td>
<td>10 mSv × age</td>
</tr>
<tr>
<td>Annual</td>
<td>50 mSv/y</td>
</tr>
<tr>
<td>Deterministic effects: dose equivalent limits for tissues and organs (annual):</td>
<td></td>
</tr>
<tr>
<td>Lens of eye</td>
<td>150 mSv/y</td>
</tr>
<tr>
<td>Skin, hands, and feet</td>
<td>500 mSv/y</td>
</tr>
<tr>
<td><strong>Embryo/Fetus Exposure:</strong></td>
<td></td>
</tr>
<tr>
<td>Effective dose limit after pregnancy declared</td>
<td>0.5 mSv/month</td>
</tr>
<tr>
<td><strong>Public Exposure (annual):</strong></td>
<td></td>
</tr>
<tr>
<td>Effective dose limit, continuous or frequent exposure</td>
<td>1 mSv/y</td>
</tr>
<tr>
<td>Effective dose limit, infrequent exposure</td>
<td>5 mSv/y</td>
</tr>
<tr>
<td>Dose equivalent limits; lens of the eye</td>
<td>15 mSv/y</td>
</tr>
<tr>
<td>Skin and extremities</td>
<td>50 mSv/y</td>
</tr>
<tr>
<td><strong>Education and Training Exposure (annual):</strong></td>
<td></td>
</tr>
<tr>
<td>Effective dose limit</td>
<td>1 mSv/y</td>
</tr>
<tr>
<td>Dose equivalent limit for lens of eye</td>
<td>15 mSv/y</td>
</tr>
<tr>
<td>Skin and extremities</td>
<td>50 mSv/y</td>
</tr>
<tr>
<td><strong>Negligible Individual Dose (annual):</strong></td>
<td></td>
</tr>
<tr>
<td></td>
<td>0.01 mSv/y</td>
</tr>
</tbody>
</table>

may be closer to $10,000. This sort of choice seldom has to be made in a hospital setting except, for example, in the purchase of remote afterloading equipment for brachytherapy.

- **PROTECTION OF THE EMBRYO/FETUS**

The NCRP recommends a monthly limit of 0.5 mSv to the embryo or fetus once a woman declares her pregnancy. In contrast to this, the ICRP recommends a limit of 2 mSv to the surface of the woman’s abdomen (lower trunk) for the remainder of her pregnancy. These recommendations are essentially similar and are designed to limit the risk of mental retardation, other congenital malformations, and carcinogenesis. The NCRP and ICRP no longer recommend specific controls for occupationally exposed women until the pregnancy is declared. There is a provision that a declared pregnancy can later be “undeclared” if the female worker so desires.

Internally deposited radionuclides pose special problems for protection of the embryo or fetus. Some remain in the body for long periods, and the doses delivered to fetal organs are not well known for all radionuclides. Consequently, particular care should be taken to limit the intake of radionuclides by pregnant women so that the equivalent dose to the embryo or fetus would not exceed the recommended limit.

- **EMERGENCY OCCUPATIONAL EXPOSURE**

Under normal conditions, only actions involving the saving of life justify acute exposures in excess of the annual effective dose limit. The use of volunteers for exposures during emergency actions is desirable. If possible, older workers with low lifetime accumulated effective doses should be chosen from among the volunteers. Exposure during emergency actions that do not involve the saving of life should be controlled, to the extent possible, at the occupational exposure limits. If this cannot be accomplished, the NCRP and ICRP recommendation of 0.5 Sv should be applied.

If, for lifesaving or equivalent purposes, the exposure may approach or exceed 0.5 Sv to a large portion of the body, the worker not only needs to understand the potential for acute effects, but also should have an appreciation of the substantial increase in his or her lifetime risk of cancer. If the possibility of internal exposures also exists, this should also certainly be taken into account.

- **EXPOSURE OF PERSONS YOUNGER THAN 18 YEARS OF AGE**

For educational and training purposes, it may be necessary and desirable to accept occasional exposure of persons younger than the age of 18 years, in which case an annual effective dose limit of 1 mSv should be maintained (NCRP).

- **EXPOSURE OF MEMBERS OF THE PUBLIC (NONOCCUPATIONAL LIMITS)**

The limitation of radiation exposure for members of the public from human-made sources is inevitably arbitrary because it cannot be based on direct experience. Various risks of accident and death regularly faced by members of the public vary greatly; the numbers range from $10^{-11}$ to $10^{-6}$ per year. Depending on their nature, these risks seem to be accepted without much thought. At the same time, everyone is exposed to natural background radiation of about 1 mSv annually, excluding radon, which may result in a mortality risk of $10^{-4}$ to $10^{-5}$ annually.

Based on these considerations, the NCRP recommended limits for human-made sources other than medical are as follows: For continuous or frequent exposure, the annual effective dose should not exceed 1 mSv. It is clear, however, that larger exposures to more limited groups of people are not especially hazardous, provided they do not occur often to the same groups. Consequently, a maximum permissible annual effective dose equivalent of 5 mSv is recommended as a limit for infrequent exposure. Medical exposures are excluded from these limitations because it is assumed that they confer personal benefit to the exposed person.

Because some organs and tissues are not necessarily protected against deterministic effects in the calculation of effective dose, the hands and feet, localized areas of the skin, and the lens of the eye are also subject to an annual dose limit of 50 mSv.

The fact that the terms frequent and infrequent in the public dose limits are not defined has caused some confusion. Nevertheless, the intention of the NCRP is laudable, namely, that exceptions to the 1 mSv per year for members of the public may be justified on the basis of
significant benefit either to those exposed or to society as a whole. Here are three examples:

1. For workers who come into contact with a co-worker who is a radionuclide therapy patient, the annual effective dose limit of 1 mSv may be exceeded under carefully controlled conditions for a small number of such workers who may receive up to 5 mSv annually.

2. For adult family members exposed to a patient who has received radionuclide therapy, the annual effective dose limit is 50 mSv. Thus, adult family members under this circumstance are considered separate from other members of the public and should receive appropriate training and individual monitoring.

3. Another example is the inadvertent irradiation of a stowaway in a cargo container irradiated with a pulsed fast neutron analysis system to assess the contents of the container. The NCRP has recommended that such systems be designed and operated in a manner such that the exposure of a stowaway would result in an effective dose of less than 1 mSv for that occurrence. However, an effective dose of up to 5 mSv would be permissible for such an occurrence if necessary to achieve national security objectives.

A more contentious issue is the exposure of members of the public to scattered radiation in a radiology department. For example, exposure of an individual member of the public to scattered radiation in the waiting room of a radiology facility is infrequent for a given individual. On the other hand, a secretary or receptionist may be exposed frequently or continuously, so the desk area must be protected to a lower level, which can be an expensive proposition. It might be tempting to reclassify the office personnel as “radiation workers,” but to do so would offend all the basic principles of radiation protection.

**EXPOSURE TO INDOOR RADON**

Radon levels vary enormously with different localities, depending on the composition of the soil and the presence of cracks or fissures in the ground, which allow radon to escape to the surface. Many homes in the United States and Europe consequently contain an appreciable quantity of radon gas, which enters the living quarters through the basement. Insulating and sealing houses increased greatly because of the escalating cost of heating oil in the 1970s, and this has exacerbated the radon problem because a well-sealed house allows fewer exchanges of air with the outside and consequently results in a greater concentration of radon. Radon is a noble gas and is itself relatively nonhazardous because if breathed in, it is breathed out again without being absorbed. In a confined space such as a basement, however, the decay of radon leads to the accumulation of progeny that are solids, which stick to particles of dust or moisture and tend to be deposited on the bronchial epithelium. These progeny emit α-particles and cause intense local irradiation.

An extreme example is the famous case of Stanley Watras who wanted to work in a nuclear power station but was turned away because the radiation monitors were set off as he entered the plant by the accumulation of radon progeny products deposited in his body and on his clothes that came from his home.

Indoor radon currently is perceived to be an important problem involving radiation exposure of the public. In the United States and most European countries, the mean radon concentration in homes is in the range of 20 to 60 Bq/m³, with higher mean values of about 100 Bq/m³ in Finland, Norway, and Sweden. Converting radon concentrations into dose to the bronchial epithelium involves many uncertainties, because such conversion depends on the model used and the assumptions made. One widely used conversion factor equates an air concentration of 20 Bq/m³ with an effective dose to the bronchial epithelium of 1 mSv per year.

The EPA has set the “action level” at about 148 Bq/m³, suggesting that remedial action should be taken to reduce radon levels if they are higher than this. The action level is about four times the average radon concentration in homes, but it is estimated that about 1 in 12 homes in the United States—about 6 million in all—have radon concentrations above this action level. In the past, other countries, including Germany, Great Britain, and Canada had much higher action levels, but these are all now under review.

The BEIR VI Committee of the National Academy of Sciences published a report on the health effects of radon in 1999. The committee’s preferred central estimates, depending on which of two models is used, are that about 1 in 10 or 1 in 7 of all lung cancer deaths—amounting to 15,400 or 21,800 per year in the United States—can be attributed to radon. There are considerable uncertainties involved, and the number could be as low
Radiation detriment is a concept used to quantify the harmful effects of radiation exposure to different parts of the body, taking into account the severity of the disease in terms of lethality, loss of quality of life, and years of life lost. Detriment includes a small component for heritable effects, a large component for lethal cancers, and an allowance for nonlethal cancers, which, although they do not cause death, nevertheless affect quality of life. ICRP has suggested the detriment-adjusted risk coefficients for stochastic effects after exposure of the whole population to radiation at LDR to be 5.5% per Sv for cancer (lethal and nonlethal combined) and 0.2% per Sv for heritable effects, making a total of 5.7% per Sv. Recent surveys indicate that the average annual dose to monitored radiation workers with measurable exposures is about 2 mSv. This results in a detriment of about 1 in 10,000, which is comparable to the death rate in what are considered to be “safe” industries such as trade and government service.

**DE MINIMIS DOSE AND NEGLIGIBLE INDIVIDUAL DOSE**

Collective dose to a population has little meaning without the concept of **de minimis dose**. The idea is to define some very low threshold below which it would make no sense to make any additional effort to reduce exposure levels further. For example, suppose there is a release of radioactivity from a reactor that dissipates into the atmosphere, blows around the world, and eventually exposes many hundreds of millions of people to very low doses. The doses may be so low that the biologic effects are negligible, but because the number of persons involved is so large, the product of the dose and the number of persons would dominate the collective dose. The term **de minimis** comes from the legal saying De minimis non curat lex, which roughly translates to “The law does not concern itself with trifles.”

Dr. Merril Eisenbud in an NCRP publication quotes this limerick of dubious origin.

There was a young lawyer named Rex, who was very deficient in sex
When charged with exposure
He said with composure
De minimis non curat lex

The concept of **de minimis** dose has been espoused by the NCRP in the form of **negligible individual dose**, defined here to be the dose below which further efforts to reduce radiation exposure to the person are unwarranted. The NCRP considers an annual effective dose of 0.01 mSv to be a negligible individual dose. This dose is associated with a risk of death between $10^{-6}$ and $10^{-7}$, which is considered trivial compared with the risk of fatality associated with ordinary and normal societal activities and, therefore, can be dismissed from consideration of additional radioprotective measures.

**RADIATION DETRIMENT**

Radiation detriment is a concept used to quantify the harmful effects of radiation exposure to different parts of the body, taking into account the severity of the disease in terms of lethality, loss of quality of life, and years of life lost. Detriment includes a small component for heritable effects, a large component for lethal cancers, and an allowance for nonlethal cancers, which, although they do not cause death, nevertheless affect quality of life. ICRP has suggested the detriment-adjusted risk coefficients for stochastic effects after exposure of the whole population to radiation at LDR to be 5.5% per Sv for cancer (lethal and nonlethal combined) and 0.2% per Sv for heritable effects, making a total of 5.7% per Sv.

Recent surveys indicate that the average annual dose to monitored radiation workers with measurable exposures is about 2 mSv. This results in a detriment of about 1 in 10,000, which is comparable to the death rate in what are considered to be “safe” industries such as trade and government service.

**NATIONAL COUNCIL ON RADIATION PROTECTION AND MEASUREMENTS AND THE INTERNATIONAL COMMISSION ON RADIOLOGICAL PROTECTION COMPARED**

At present, there is a difference in the recommendations of the national and international bodies regarding the maximum permissible effective dose for occupational exposure (stochastic effects). The differences are highlighted in Table 17.5. Both bodies recommend a maximum of 50 mSv in any 1 year, but the NCRP adds a lifetime cumulative limit of the person’s age times 0.1 mSv, and the ICRP adds a limit of 20 mSv per year averaged over defined periods of 5 years.

The practical consequence of this difference is that a radiation worker starting at, for example, age 18 years can accumulate a larger dose under the NCRP recommendations in the early years up to an age in the mid-30s, but later in life could accumulate a larger dose under the ICRP recommendations. Under NCRP recommendations, a new radiation worker could receive 50 mSv in each of several consecutive years until the limit of age times 10 mSv kicks in. Under ICRP rules, the average cannot exceed 20 mSv per year over a 5-year period, so one or two 50-mSv years would have to be followed by several years at very low exposure levels. If individuals were exposed
throughout their working lives to the maximum permissible dose, the excess risk of stochastic effects (cancer and heritable effects) would be about the same under NCRP or ICRP recommendations. Under the NCRP, a person occupationally exposed from 18 to 65 years of age could receive a total dose of 650 mSv. Under the ICRP, the same person could receive 940 mSv, but less would be received in the early years and more at later ages.

The NCRP scheme is less restrictive for a few workers in the nuclear power industry who tend to receive large effective doses in their early years working on nuclear reactors. Later in life, these individuals tend to occupy supervisory or administrative positions and receive little, if any dose. To cope with those who do not, the NCRP has added the extra recommendation that this limit, age \( \times 10 \) mSv, can be relaxed in individual cases after counseling, if implementation of the recommendation would mean loss of a job.

It should be emphasized that few persons exposed occupationally in a medical setting receive doses anywhere near the recommended limits. Some interventional radiologists may well receive more than 50 mSv per year to a monitor worn outside the lead rubber apron or to a monitor worn at neck level or on the forearm. But the recommended maximum permissible levels refer to effective dose, which takes into account the parts of the body exposed.

### THE HISTORY OF THE CURRENT DOSE LIMITS

In 1956, the ICRP reduced the dose limit for radiation workers from 0.3 R per week to 0.1 R per week. This corresponds to 5 R per year, which is still the maximum permissible dose allowed in 1 year to radiation workers today, except that the unit has changed and it is now called 50 mSv. This dose limit suggested by ICRP was based entirely on genetic effects in the fruit fly, *Drosophila*.

In the half century or so that has elapsed since then, concern for genetic effects, or heritable effects as we now call them, has declined steadily; first because of the availability of mouse data and more recently, because of doubts about the relevance of specific locus mutations in mice. In the meantime, concern for radiation-induced carcinogenesis has increased as more and more solid cancers appeared in the A-bomb survivors. In the 1950s, genetic effects were considered to be the most important consequence of low doses of ionizing radiation.

To cope with these changing perceptions, and what was considered to be an alarming increase in cancer among the A-bomb survivors, ICRP in 1991 introduced a second limit. As well as the 50 mSv limit in any 1 year, they required that the average over 5 years should not exceed 20 mSv per year. NCRP in 1993 coped with the perceived increase in cancer risk by adding a cumulative limit of age \( \times 10 \) mSv to the existing annual limit of 50 mSv. Although these differ in detail, with NCRP allowing workers more doses in earlier years and less later on, the respective limits recommended by the two organizations are quite similar, the NCRP being a little more restrictive with respect to overall lifetime risks. Both organizations aimed to make the risks to radiation workers comparable to other “safe” industries, and both sets of recommendations would result in a radiation-induced cancer mortality risk of about 3%.

However, although the NCRP Report No. 116 was published in 1993 and included the cumulative limit of age \( \times 10 \) mSv, the Council only makes “recommendations” because the legal responsibility for the implementation of radiation safety is in the hands of the NRC, the DOE, and state or city bureaus of radiation control. In fact, the United States NRC has never adopted the cumulative limit and to this day, the annual limit is a total effective dose equivalent of 50 mSv. Consequently, if a radiation worker starts at age 18 years and works at the dose limit until retiring at age 65 years, he or she would face a radiation-induced cancer incidence of 19% and a cancer mortality of 10.8% (Table 17.6). This is in marked

### TABLE 17.6 Cancer Risks for a Radiation Worker Receiving the Maximum Permissible Dose from Age 18 to 65 years

<table>
<thead>
<tr>
<th>Rule</th>
<th>Total Dose</th>
<th>Cancer Incidence</th>
<th>Cancer Mortality</th>
</tr>
</thead>
<tbody>
<tr>
<td>NRC 50 mSv/y</td>
<td>2.35 Sv</td>
<td>19.0</td>
<td>10.8</td>
</tr>
<tr>
<td>NCRP 10 mSv ( \times ) age</td>
<td>0.65 Sv</td>
<td>6.1</td>
<td>3.3</td>
</tr>
</tbody>
</table>
FIGURE 17.4  This chart compiled by Dr. Noelle Metting, Office of Science of the U.S. Department of Energy, puts into perspective the different dose ranges relevant to radiation therapy, diagnostic radiology, and background radiation.
Effective dose is the sum of the weighted equivalent doses for all irradiated tissues and organs multiplied by the appropriate $W_T$.

Committed equivalent dose is the integral over 50 years of the equivalent dose after the intake of a radionuclide.

Committed effective dose is the integral over 50 years of the effective dose in the case of an incorporated radionuclide.

Collective effective dose is a quantity for a population and is the sum of effective doses to all members of that population. The unit is person-sievert.

Collective committed effective doses applies to a population ingesting or inhaling radionuclides and is the integral over 50 years of the effective dose over the entire population.

All radiation exposures are governed by the ALARA principle.

No occupational exposure should be permitted before 18 years of age.

The effective dose in any 1 year should not exceed 50 mSv (NCRP).

The individual worker’s cumulative lifetime effective dose should not exceed age in years $\times 10$ mSv (NCRP). However, to date, the NRC has not adopted this cumulative limit.

To limit deterministic effects, the dose limit to the lens of the eye is 150 mSv per year, and the dose limit to localized areas of the skin, hands, and feet is 500 mSv per year.

Once a pregnancy is declared, the NCRP recommends a monthly limit of 0.5 mSv to the embryo or fetus.

Specific controls for occupationally exposed women are no longer recommended until a pregnancy is declared.

Internally deposited radionuclides pose a special problem for protection of the embryo or fetus; particular care should be taken to limit intake.

Emergency occupational exposures normally justify doses in excess of the recommended limits only if life-saving actions are involved. Volunteers from among older workers with low lifetime accumulated effective doses should be chosen in emergencies in which the exposure may be up to 0.5 Sv. If the exposure may exceed 0.5 Sv, the worker should be counseled about the short-term and long-term possible consequences.

Contrast to the corresponding figures that would be applicable if the ICRP or NCRP limitations were followed, when the radiation-induced cancer incidence would be 6%, and mortality would be 3%. These estimates are based on data published in the BEIR VII report. It is not widely appreciated that radiation workers in the United States are allowed significantly higher cancer risks than other industrialized countries that, by and large, adopt and enforce the recommendations of ICRP.

■ **DOSE RANGES**

Doses to which individuals are exposed vary enormously by several orders of magnitude. Figure 17.4 attempts to put this into perspective by comparing the ranges of doses used in medicine with doses received occupationally and from natural sources.

**SUMMARY OF PERTINENT CONCLUSIONS**

- The objectives of radiation protection are to prevent clinically significant deterministic effects by keeping doses below the practical threshold and to limit the risk of stochastic effects (cancer and heritable effects) to a reasonable level in relation to societal needs, values, and benefits gained.

- Justification is one of the basic principles of radiation protection; a practice involving exposure to radiation should produce sufficient benefit to the exposed individual or to society to offset the radiation detriment it causes.

- $W_R$ are approximate values of the RBE, applicable to low doses and relevant to carcinogenesis and heritable effects. $W_R$ is chosen by the ICRP based on experimental RBE values with a large judgmental factor.

- Equivalent dose is the product of absorbed dose and $W_R$. The unit is sievert for an absorbed dose in Gy. ICRP has recommended a new name for this quantity—radiation weighted dose—and is considering a new name for the unit.

- $W_T$ reflects the susceptibility of different organs or tissues to carcinogenesis or heritable effects.
For educational or training purposes, it may sometimes be desirable to accept radiation exposures of persons younger than 18 years of age, in which case the annual effective dose limit of 1 mSv should be maintained.

The annual effective dose limit for members of the public is 1 mSv, except for infrequent exposures in which the limit may be 5 mSv. Medical x-rays are excluded from these limitations because they are assumed to confer personal benefit.

For deterministic effects, the dose limit for members of the general public is 50 mSv to the hands and feet, to localized areas of the skin, or to the lens of the eye.

Indoor radon is perceived to be the most important problem involving radiation exposure of the general public. Remedial action in homes is recommended by the EPA if the radon concentration exceeds 148 Bq/m³.

Negligible individual dose is the dose below which further expenditure to improve radiation protection is unwarranted. The negligible individual dose is an annual effective dose of 0.01 mSv, which carries a risk of between $10^{-6}$ and $10^{-7}$ of carcinogenesis or heritable effects.

ICRP introduced the concept of “detriment” to quantify the harmful effects of radiation exposure in different parts of the body, taking account of the severity of the disease in terms of lethality, loss of quality of life, and years of life lost.

A uniform whole body equivalent dose of 1 Sv to an adult radiation worker is assumed to result in a total detriment of about 5.7% per Sv. This is made up of a risk of fatal and nonfatal cancer together with a small contribution from severe heritable effects.

The average annual equivalent dose to monitored radiation workers is about 2 mSv. This involves a total detriment of about one in 10,000, which is comparable to the annual risk of a fatal accident in a “safe” industry such as trade or government service.

The NCRP and ICRP differ in two important recommendations:

1. The effective dose limit for occupational exposure (stochastic effects). The NCRP recommends a lifetime cumulative limit of age $\times$ 10 mSv, with a limit in any year of 50 mSv. The ICRP recommends a limit of 20 mSv per year averaged over defined periods of 5 years, with a limit in any year of 50 mSv.

2. The dose limit to the developing embryo or fetus once a pregnancy is declared. The NCRP recommends a monthly limit of 0.5 mSv to the embryo or fetus. The ICRP recommends a limit of 2 mSv to the surface of the woman’s abdomen for the remainder of pregnancy.

- **GLOSSARY OF TERMS**

**absorbed dose:** The energy imparted to matter by ionizing radiation per unit mass of irradiated material at the place of interest. The unit is gray (Gy), defined as an energy absorption of 1 J/kg.

**ALARA** (as low as reasonably achievable): Economic and social factors being taken into account. This is identical to the principle of optimization of protection used by the ICRP.

**annual limit on intake:** The activity of a radioisotope taken into the body during a year that would provide a committed equivalent dose to a person, represented by a reference “man,” equal to the occupational dose limit set by recommending and regulating bodies. The annual limit normally is expressed in becquerel (Bq).

**becquerel (Bq):** The special name for the unit of activity. $3.7 \times 10^{-10}$ Bq = 1 Ci.

**collective committed effective dose:** Applies to a population ingesting or inhaling radionuclides that deposit their dose over a prolonged period of time and is the integral of the effective dose over the entire population out to a period of 50 years.

**collective effective dose:** Applies to a group of persons and is the sum of the products of the effective dose and the number of persons receiving that effective dose.

**collective equivalent dose:** Applies to a group of persons and is the sum of the products of the equivalent dose and the number of persons receiving that equivalent dose.

**committed effective dose:** The sum of the committed organ or tissue equivalent doses resulting from an intake multiplied by the appropriate tissue weighting factors.
committed equivalent dose: The equivalent dose averaged throughout a specified tissue in the 50 years after intake of a radionuclide into the body.

deterministic effects: Effects for which the severity of the effect in affected persons varies with the dose and for which a threshold usually exists. These were formerly known as nonstochastic effects. An example is a cataract.

effective dose: The sum over specified tissues of the products of the equivalent dose in a tissue and the appropriate WT for that tissue.

equivalent dose: A quantity used for radiation protection purposes that takes into account the different probability of effects that occur with the same absorbed dose delivered by radiations of different quality. It is defined as the product of the averaged absorbed dose in a specified organ or tissue and the WT. The unit of equivalent dose is the sievert (Sv). The ICRP is now recommending that this be called the radiation weighted dose.

genetically significant dose (GSD): The dose to the gonads weighted for the age and sex distribution in those members of the population expected to have offspring. The genetically significant dose is measured in sievert.

gray (Gy): The special name for the SI unit of absorbed dose, kerma, and specific energy imparted. 1 Gy = 1 J/kg.

negligible individual dose: A level of effective dose that can be dismissed as insignificant and below which further efforts to improve radiation protection are not justified. The recommended negligible individual dose is 0.01 mSv per year.

nonstochastic effects: Previous term for deterministic effects.

organ or tissue weighting factor (WT): See tissue weighting factor.

rad: The old unit for absorbed dose, kerma, and specific energy imparted. One rad is 0.01 J absorbed per kilogram of any material (also defined as 100 erg/g). The term is being replaced by the gray: 1 rad = 0.01 Gy.

radiation weighted dose: New name recommended by ICRP for equivalent dose.

radiation weighting factor (WR): A factor used for radiation protection purposes that accounts for differences in biologic effectiveness between different radiations. The WR is independent of the WT.

relative biologic effectiveness (RBE): A ratio of the absorbed dose of a reference radiation to the absorbed dose of a test radiation to produce the same level of biologic effect, other conditions being equal. It is the quantity that is measured experimentally.

rem: The old unit of equivalent dose or effective dose. It is the product of the absorbed dose in rad and modifying factors and is being replaced by the sievert.

sievert (Sv): The unit of equivalent dose or effective dose in the SI system. It is the product of absorbed dose in gray and modifying factors. 1 Sv = 100 rem.

stochastic effects: Effects for which the probability of their occurring, rather than their severity, is a function of radiation dose without threshold. More generally, stochastic means random in nature.

tissue weighting factor (WT): A factor that indicates the ratio of the risk of stochastic effects attributable to irradiation of a given organ or tissue to the total risk if the whole body is uniformly irradiated. Organs that have a large WT are those that are susceptible to radiation-induced carcinogenesis (such as the breast or thyroid) or to hereditary effects (the gonads).

working level: The amount of potential α-particle energy in a cubic meter of air that results in the emission of $2.08 \times 10^{-3}$ J of energy.

working level month: A cumulative exposure, equivalent to exposure to one working level for a working month (170 hours), that is, $2 \times 10^{-5}$ J·m$^{-3} \times 170 = 3.5 \times 10^{-3}$ J·h·m$^{-3}$.

\[ \text{BIBLIOGRAPHY} \]


For Students of Radiation Oncology
Tissue homeostasis depends on the regulated cell division and self-elimination (programmed cell death) of each of its constituent members except its stem cells. In fact, a tumor arises because of uncontrolled cell division and failure for self-elimination. One can consider cancer as a Darwinian-like process whereby the fittest cells reproduce to become the dominant population of a tumor. Alterations in three groups of genes are responsible for the deregulated control mechanisms that are the hallmarks of cancer cells:

- **Proto-oncogenes** are components of signaling networks that act as positive growth regulators in response to mitogens, cytokines, and cell-to-cell contact. A gain-of-function mutation in only one copy of a proto-oncogene results in a dominantly acting oncogene that often fails to respond to extracellular signals.

- **Tumor suppressor genes** are also components of the same signaling networks as proto-oncogenes, except that they act as negative growth regulators. They modulate proliferation and survival by antagonizing the biochemical functions of proto-oncogenes or responding to unchecked growth signals. In contrast to oncogenes, inactivation of both copies of tumor suppressor genes is required for loss of function in most cases.

- **DNA stability genes** form a class of genes involved in both monitoring and maintaining the integrity of DNA. Loss of these genes results in defective sensing of DNA lesions as well as improper repair of the damaged template.

The malignant progression from normal tissue to tumor to metastasis occurs in various “steps” over a period of time. These steps, which are the result of mutations, deletions,
Initially, it was thought that cancer was the result of deregulated growth signals by oncogenes, a concept supported by increased proliferation in many types of cancer. In the last decade, the finding that many cancers possess diminished apoptotic (programmed cell death) programs or loss of cell cycle control has led to the concept that mutations in proto-oncogenes and tumor suppressor genes that inhibit apoptosis provide a selective growth advantage to a premalignant cell that allows it to clonally expand. Mutations in DNA stability genes increase the rate of acquiring genetic mutations that will result in a malignant tumor. Thus, although tumor cells are considered clonal in origin, most tumors contain heterogeneous populations of cells that differ in their ability to repopulate the tumor or form metastases. In fact, only a small percentage of tumor cells possess the ability to form a tumor, leading to the concept that tumors possess “stem cells” and that elimination of these stem cells is essential for controlling tumor growth.

**MECHANISMS OF CARCINOGENESIS**

A single genetic alteration that leads to the activation of an oncogene or loss of a tumor suppressor gene does not, by itself, lead to the formation of a solid tumor. Instead, carcinogenesis appears to be a multistep process with multiple genetic alterations occurring over an extended period of time; at least, that is how it appears. Sometimes, these genetic alterations are carried in the germ line, like, for example, in the cancer-predisposing syndrome retinoblastoma; however, heritable mutations are rare. Most alterations that lead to cancer are acquired in the form of somatic mutations: chromosomal translocations, deletions, inversions, amplifications, or simple point mutations.

**ONCOGENES**

The first demonstration that a tumor was initiated by a cellular component found in tumor cells but not in normal cells was shown by Rous in the early 1900s. His landmark studies demonstrated that cell-free extracts derived from chicken sarcomas could cause a new sarcoma, if injected into healthy chickens. In the 1970s,
From these early studies, several important conclusions could be derived. These are the following:

1. Cancer can be caused by a genetically transmissible agent—in the case of chicken sarcoma, by a retrovirus containing a unique piece of genetic information that was later designated as the src gene;
2. Only a certain region of a retrovirus is needed for transformation; and
3. The region of the viral genome necessary for transformation is not involved in the normal replicative life cycle.

Huebner and Todaro later proposed that cancer-causing viral genes such as src are normally inactive but can be activated when they recombine with a retroviral genome. Once they do so, they pass from being a benign proto-oncogene (i.e., c-src) to a malignant form (v-src) capable of causing cancer when introduced into the appropriate host cell. Although we now know that viruses represent only one of several mechanisms that cause the deregulated expression of a proto-oncogene, these studies helped to define oncogenes as mutant forms of normal cellular genes that are altered in expression and/or function by various agents, including radiation, chemicals, and viruses. Consequently, very different agents produce tumors that are indistinguishable from one another. This is illustrated in Figure 18.2.

![Figure 18.2](image_url)
MECHANISMS OF ONCOGENE ACTIVATION

Although many mechanisms are involved in oncogene activation, transcriptional deregulation by overexpression or abnormal expression of the messenger RNA (mRNA) of a proto-oncogene is a common theme. At least four mechanisms exist for oncogene activation in human neoplasms (Fig. 18.3).

Retroviral Integration through Recombination

Retroviral integration of proto-oncogene sequences in retroviral genomes occurs through two possible recombination mechanisms. First, mRNAs from a proto-oncogene recombine with viral genomic RNAs. During the recombination process, the proto-oncogene mRNA becomes deregulated as it comes under the control of the viral promoter, termed long terminal repeat (LTR). However, the probability of RNA recombination events between proto-oncogene mRNA and viral mRNA generating an oncogenic retrovirus is quite low and undermines the importance of this mechanism.

A second more probable mechanism is as follows: First, a retroviral genome integrates in proximity to a proto-oncogene, where the proto-oncogene is under the transcriptional control of the retrovirus LTR promoter. Then the viral and proto-oncogene sequences become closely associated through a DNA recombination event that permits the production of mRNAs that contain both viral and proto-oncogene sequences. In this scenario, the proto-oncogene becomes transcriptionally deregulated because it is under the control of the viral promoter LTR. In addition, it can acquire mutations in its coding sequence. Although the proto-oncogene can become mutated during the recombination process, the key point is that its

![Diagram of oncogene activation mechanisms](image_url)
Several steps are needed for the molecular cloning of an oncogene from transformed rodent cells using the rodent fibroblast transformation assay. These are the following:

1. The human DNA containing the transforming oncogene is transfected into mouse cells.
2. The DNA from the transformed mouse cells is serially transfected to reduce the amount of human DNA that is not associated with the transforming oncogene.
3. After several rounds of transfection, the DNA is isolated from a soft agar colony and digested with restriction enzymes to make a genomic DNA library.
4. The library is then screened with a human-specific repetitive probe that does not cross-react with the mouse's DNA, thereby identifying human sequences in a mouse background.

**DNA Mutation of Regulatory Sites**

The union of the technique for gene transfer with mouse transformation assays facilitated the isolation of human oncogenes that were activated by DNA mutation. Transfection of human DNA into immortalized but untransformed mouse cells was first used to isolate the H-ras oncogene from bladder carcinoma cells. The key to this approach is that only transformed cells possess the ability to grow in soft agar (Fig. 18.4). The implicit assumption is that a specific gene (or more) is responsible for causing the bladder carcinoma, and that it will act in a dominant fashion to induce a tumor. Indeed, multiple groups were successful in isolating the H-ras oncogene by this approach.

**FIGURE 18.4** Schematic diagram of a typical DNA transfection protocol in which oncogenes can be isolated from cells transformed in vitro by either radiation or chemical carcinogens. DNA sequences are then characterized using Southern blot hybridization.
Chromosome Translocation

It had long been known that tumors possessed abnormal karyotypes. However, the chromosome content of many solid tumors is unstable, making it difficult to determine which cytogenetic alterations are causative for tumorigenesis and which are the consequence of the neoplastic process. The first real breakthrough in identifying tumor-specific chromosome alterations occurred in the late 1950s when Dr. Peter Nowell found a consistent shortened version of chromosome 22 in individuals afflicted with chronic myelogenous leukemia (CML). Because many patients with CML possess an abnormal chromosome 22 in their leukemic cells, this was a strong indication that a specific chromosome alteration is involved in the pathogenesis of this malignancy. With the advent of more sophisticated cytogenetic and molecular techniques, it was discovered that this shortened version of chromosome 22 is caused by a symmetric translocation with chromosome 9. It was hypothesized, therefore, that the translocation between chromosomes 9 and 22 gives rise to CML. Further molecular analysis revealed that the \textit{bcr} gene on chromosome 9 translocates in front of the \textit{abl} gene on chromosome 22, producing a fusion transcript with abnormal expression (Fig. 18.5).

Gene Amplification

Oncogene amplification occurs through breakage-fusion-bridge cycles in anaphase during mitosis. In contrast to the other mechanisms discussed that involve transcriptional deregulation as a key mechanism of oncogene activation, gene amplification represents an alternative means of increasing proto-oncogene expression by increasing the number of DNA copies of the proto-oncogene. Gene amplification can result in an increased number of copies of extrachromosomal molecules called \textit{double minutes} or can result in intrachromosomal amplified regions called \textit{homogeneously staining regions} (HSR), both of which are detectable by fluorescence \textit{in situ} hybridization or Giemsa banding of chromosomes. The N-\textit{myc} oncogene is a classic example of an oncogene amplified in leukemia, neuroblastoma, and breast cancer.
Chapter 18 • Cancer Biology

Their activation even though the other copy of the gene is unchanged. This concept led to speculation that another class of genes, termed “antioncogenes,” suppresses the effect of oncogenes on transformation and tumor formation. The existence of tumor suppressor genes was supported by cell fusion studies between tumor cells and normal cells and by the family history studies of people afflicted with inherited cancer-prone disorders such as retinoblastoma or Li–Fraumeni syndrome. Although one mutated version of an oncogene is sufficient to drive malignant progression, one functional copy of a tumor suppressor gene is sufficient to suppress transformation, suggesting that both copies of a tumor suppressor gene must be inactivated to inhibit tumor growth (Fig. 18.6). Insight into the mechanism of tumor suppressor gene inactivation came from Knudson’s epidemiologic studies of families in which retinoblastoma appeared to be inherited in an autosomal dominant manner. Patients with familial retinoblastoma develop bilateral or multifocal disease at an earlier age than

map near translocation breakpoints, as markers to identify potential translocation partners. Although numerous translocation breakpoints have been identified in hematopoietic neoplasms, few consistent translocations have been found in solid tissue tumors. The reason for this is still unclear, but may be attributed to the fact that hematopoietic cancers require fewer alterations for neoplasia than solid tumors. Table 18.1 provides examples of the chromosomal changes that result in oncogene activation and the associated human malignancies. Interestingly, there are no known examples of oncogenes activated by retroviruses in human malignancies.

### MUTATION AND INACTIVATION OF TUMOR SUPPRESSOR GENES

#### The Retinoblastoma Paradigm

Oncogenes result from a mutation, deletion, or alteration in the expression of one copy of a gene. Thus, oncogenes are dominant genes because a mutation in only one copy will cause their activation even though the other copy of the gene is unchanged. This concept led to speculation that another class of genes, termed “antioncogenes,” suppresses the effect of oncogenes on transformation and tumor formation. The existence of tumor suppressor genes was supported by cell fusion studies between tumor cells and normal cells and by the family history studies of people afflicted with inherited cancer-prone disorders such as retinoblastoma or Li–Fraumeni syndrome. Although one mutated version of an oncogene is sufficient to drive malignant progression, one functional copy of a tumor suppressor gene is sufficient to suppress transformation, suggesting that both copies of a tumor suppressor gene must be inactivated to inhibit tumor growth (Fig. 18.6). Insight into the mechanism of tumor suppressor gene inactivation came from Knudson’s epidemiologic studies of families in which retinoblastoma appeared to be inherited in an autosomal dominant manner. Patients with familial retinoblastoma develop bilateral or multifocal disease at an earlier age than

<table>
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<tr>
<th>Oncogene</th>
<th>Chromosomal Change</th>
<th>Cancer</th>
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<tr>
<td>int-1</td>
<td>Proviral insertion</td>
<td>Murine breast carcinoma</td>
</tr>
<tr>
<td>int-2</td>
<td>Proviral insertion</td>
<td>Murine breast carcinoma</td>
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<tr>
<td>pim-1</td>
<td>Proviral insertion</td>
<td>Murine T-cell lymphoma</td>
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<tr>
<td>N-ras</td>
<td>Point mutation (1)</td>
<td>Melanoma</td>
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<td>K-ras</td>
<td>Point mutation (12)</td>
<td>Pancreas carcinoma</td>
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<td>H-ras</td>
<td>Point mutation (11)</td>
<td>Colon carcinoma</td>
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<td>neu</td>
<td>Point mutation (17)</td>
<td>Neuroblastoma</td>
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<td>N-myc</td>
<td>Gene amplification (8)</td>
<td>Neuroblastoma</td>
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<td>L-myc</td>
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<td>neu</td>
<td>Gene amplification (17)</td>
<td>Breast carcinoma</td>
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<td>EGFR</td>
<td>Gene amplification (7)</td>
<td>Squamous cell carcinoma</td>
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<tr>
<td>bcr-abl</td>
<td>Translocation (9–22)</td>
<td>Chronic myelogenous leukemia</td>
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<tr>
<td>c-myc</td>
<td>Translocation (8–22)</td>
<td>Burkitt lymphoma</td>
</tr>
<tr>
<td>bcl-2</td>
<td>Translocation (14–18)</td>
<td>Diffuse large B-cell lymphoma</td>
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*Human oncogenes activated by retroviruses have not yet been found in human malignancies, only murine cancers.*
tumor suppressor genes are recessive genes that require the inactivation of both functional gene copies before malignancies develop, whereas loss of one functional copy results only in increased cancer susceptibility. In addition, it is often the case that the same tumor suppressor gene involved in hereditary cancer syndromes, such as retinoblastoma, is also inactivated in other forms of cancer. The Rb gene itself has now been implicated in several other human cancers, which indicates that it may play a generalized role in tumor growth suppression in various tissues. For example, patients who are cured of familial retinoblastoma are at increased risk of osteosarcoma, small cell lung cancer, and breast cancer; although the loss of the Rb gene alone is sufficient for retinoblastoma, further changes are required for the development of these other tumors.
The Li–Fraumeni Paradigm

The Li–Fraumeni syndrome (LFS) is a rare autosomal, dominantly inherited disease that predisposes individuals to develop osteosarcomas, soft-tissue sarcomas, rhabdomyosarcomas, leukemias, brain tumors, and carcinomas of the lung and breast (Fig. 18.7). Initial attempts to identify the genetic mutations that underlie LFS were unsuccessful because of the rarity of the syndrome and the high mortality of the patients. The major insight into the underlying cause of LFS came when it was found that mice that overexpress a mutant version of the p53 tumor suppressor gene in the presence of the wild-type p53 gene develop a spectrum of tumors similar to that seen in patients with LFS. Sequencing of p53 in affected family members revealed germ line missense mutations in the p53 tumor suppressor gene located on chromosome 17p13 that resulted in its inactivation, and tumors derived from affected individuals had lost the remaining wild-type allele of p53. Similar to retinoblastoma, loss or inactivation of both wild-type copies of p53 is needed for tumor formation. However, functional loss of one germ line inherited copy of mutant p53 accelerates the onset of spontaneous tumor formation. Therefore, patients with LFS follow a similar paradigm as patients with retinoblastoma in developing spontaneous tumors, but unlike retinoblastoma in which germ line mutations mainly give rise to retinal tumors, loss of p53 results in a wide spectrum of tumors. Table 18.2 provides examples of other cancer predisposition genes and their associated syndromes.

Familial Breast Cancer, BRCA1 and BRCA2

The breast cancer susceptibility genes (BRCA1 and BRCA2) are found mutated in 5% and 10% of all breast cancer cases and are also associated with familial ovarian and prostate cancers. Familial breast cancer can be distinguished from sporadic breast cancer by its earlier age of onset, an increased frequency of bilateral tumors, and a higher incidence of cancer, in general, within an affected family. Histopathologically and anatomically, familial and sporadic cases of breast cancer are indistinguishable. Although BRCA1 and BRCA2 mutations account for the most inherited forms of breast cancer, they are rarely found in sporadic cases of breast cancer. BRCA1 and BRCA2 have been implicated in DNA damage, repair, cell cycle progression, transcription, ubiquitination, apoptosis, and in the determination of stem-cell fate. The most well-studied roles of BRCA1 and BRCA2 are in homologous recombination (see Chapter 2). Phenotypically, cells deficient in BRCA1 or BRCA2 exhibit elevated levels of genomic instability, which is consistent with their roles in homologous recombination. This conclusion is also supported.

**FIGURE 18.7** Pedigree analysis of familial cancer history of LFS. Symbols: B, breast cancer; G, glioblastoma; L, leukemia; Lu, lung cancer; P, pancreatic cancer; S, sarcoma; W, Wilms tumor. (Adapted from Li FP, Fraumeni JF. Prospective study of a family cancer syndrome. *J Amer Med Assoc.* 1982;247:2692–2694, with permission.)
SOMATIC HOMOZYGOSITY

How are recessive tumor suppressor genes lost? Cytogenetic studies are used to identify chromosomal changes in peripheral blood lymphocytes or fibroblasts from patients with cancer, especially those with a family history of cancer, to identify chromosomal rearrangements or deletions. At the subchromosomal level, a genomewide linkage analysis is used to determine that a certain chromosome region is tightly linked with cancer predisposition. Both copies of a suppressor gene in the sporadic form of retinoblastoma and other solid tumors may result from two independent allelic mutations, but in practice, it occurs more often by the process of somatic homozygosity. The steps appear to be as follows: One chromosome of a pair is lost, a deletion occurs in the remaining chromosome, and the chromosome with the deletion replicates. Instead of having each of the two alleles contributed by different parents, the cell has both alleles from the same parent, with loss of a vital piece containing the tumor suppressor gene (Fig. 18.8). This process has been documented for chromosome 13 in the case of retinoblastoma, chromosome 11 in Wilms tumor, chromosome 3 for small cell lung cancer, and chromosome 5 for colon cancer. Most interesting of all, perhaps, is the case of
THE MULTISTEP NATURE OF CANCER

Perhaps the most pervasive dogma in cancer research is that carcinogenesis is a multistage process. The implication is that there are several distinct events that may be separated in time. This idea is almost 70 years old and is exemplified by the skin cancer experiments in mice that introduced the concepts of initiation, promotion, and progression as stages in tumor development.

Genetic analysis of cells from solid tumors also suggests alterations, mutations, or deletions in multiple signaling genes, either oncogenes or suppressor genes; 6 to 12 mutations have been suggested for the formation of a carcinoma. In the case of colorectal cancer, a model has been proposed that correlates a series of chromosomal and molecular events with the changes in the histopathology of normal epithelium during the multistage formation of colorectal cancer and metastatic carcinoma. This concept is illustrated in Figure 18.9.

A more general model of the series of events in carcinogenesis is shown in Figure 18.10. The first event, from whatever cause (including ionizing radiation), causes a mutation in a gene in one of the families responsible for the stability of the genome. This leads to a mutator phenotype, so that with many cells dividing, multiple mutations are likely in cancer-associated genes, both oncogenes and tumor suppressor genes. This in turn leads to progression of the cancer and ultimately its invasive and metastatic properties.

Therefore, it is not surprising that restoration of one copy of a tumor suppressor gene is sometimes not sufficient to restore tumor suppressor activity because solid tumors accumulate mutations that can make them refractory to the restoration of a single tumor suppressor gene. Although tumor suppressor genes are functionally quite different from oncogenes, they modulate similar cellular targets as oncogenes.
p53
p16

Cancer
Progression
Invasion
Metastasis

Mutations in cancer-associated genes

 ras

DNA repair
Mismatch repair
DNA replication
Chromosomal segregation

An early mutation in gene responsible for the stability of the genome

Mutator phenotype leading to multiple mutations

Oncogenes and tumor-suppressor genes

FIGURE 18.10 Illustrating the multistep nature of carcinogenesis and the concept of the mutator phenotype. The first step in carcinogenesis by radiation or any other agent may be a mutation in one of the gene families responsible for the stability of the genome. This may be a DNA repair gene, an MMR gene, or a gene in a family as yet unidentified. This leads to the mutator phenotype, with multiple mutations possible in both oncogenes and tumor suppressor genes. This then leads to a series of steps that result in an invasive metastatic cancer. Not all the same mutations need to be present in every case.

FUNCTION OF ONCOGENES AND TUMOR SUPPRESSOR GENES

The myriad of genetic and epigenetic changes that drive tumor evolution is a systems biology problem in which cells can be thought of as circuits, where an alteration of the circuit can lead to increased output, decreased output, complete loss of output, or no change. Therefore, cancer biologists attempt to determine how a specific gene, when mutated, alters normal tissue function. To understand how oncogenes and tumor suppressor genes lead to neoplasia, we need to understand how each of these circuits impacts normal cellular physiology. What cellular functions are disrupted by oncogene activation and tumor suppressor gene inactivation, and how do these disrupted functions affect the differentiation, growth, and death of cells? What follows is a description of the general categories of cell functions that are perturbed by deregulation of these two classes of genes during malignant progression.

Deregulated Proliferation

The loss of proliferative control of cancer cells is evident to all who study cancer. In fact, the earliest concept suggested by tumor biologists was that cancer was a disease of uncontrolled proliferation. Untransformed cells respond to extracellular growth signals known as mitogens through a transmembrane receptor that signals to intracellular circuits that increase growth. Thus, the growth factor, the receptor, and the intracellular circuits can all lead to self-sufficiency when deregulated. Typically, one cell secretes a mitogenic signal to stimulate the proliferation of another cell type. For example, an epithelial cell can secrete a signal to stimulate fibroblasts to proliferate. In contrast to untransformed cells, transformed cells have become autonomous in regulating their growth by responding to the mitogenic signals they themselves produce. In this manner, they use an autocrine circuit to escape the need for other cell types. For example, mesenchymal cells are responsive to transformation by v-sis, because they possess receptors for platelet-derived growth factor (PDGF), and breast epithelial cells are responsive to int-2, because they possess receptors for fibroblast growth factor (FGF).

If overexpression of growth factors can lead to uncontrolled proliferation, then continuous activation or overexpression of growth factor receptors will do the same. Several well-known oncogenes, such as v-erb-2 (HER-2/neu) and v-fms, encode growth factor receptors. These receptors are mutated at their amino terminal...
residues so that they no longer require their respective growth factor (ligand) to signal induction of their kinase activity. Growth factor receptors can structurally be divided into extracellular ligand-binding domains (LBDs), transmembrane-spanning domains (TMS), and intracellular kinase domains. Although mutations have been found in all three domains, mutations in the ligand-binding domains are a common alteration that results in constitutive kinase activity that transduces the signal for the cell to proliferate. In addition to structural alterations in the receptor, some tumors overexpress growth factor receptors that make them hyperresponsive to physiologic levels of growth factor stimulation. In contrast to mitogenic-responsive growth factor receptors, a second class of receptors that transmit signals from the extracellular matrix can also regulate proliferation. Integrin receptors are the prototypical example of this class of regulators that transmit signals from different components of the extracellular matrix (ECM) to signal proliferation or quiescence.

There are numerous intracellular circuits that transduce the signal from the cell surface to the nucleus of the cell. The src, ras, and abl proteins are all members of this group. Most members of this group are tyrosine kinases or serine/threonine kinases. Src and abl are tyrosine kinases located on the cytoplasmic side of the cell membrane. H-ras, K-ras, and N-ras are a family of GTP-binding proteins also located on the cytoplasmic side of the cell membrane and are the most frequently mutated oncogene family in human cancers. The Ras/Raf/ERK, Ras/Ral-GDS/Ral, and Ras/PI(3)K/Akt/TOR pathways play critical roles in transducing signals from growth factor receptors at the cell surface to the nucleus in untransformed cells. Oncogenic forms of ras bypass the normal growth regulatory signals of a cell by being locked in an “on” state, obviating the need for external signals from growth factors to activate them (Fig. 18.11).

**Failure to Respond to Growth-Restrictive Signals**

If signal transduction cascades initiate at the cell membrane, then they end in the nucleus. Normally, various cells in the body are at rest in a nonproliferative state (G0). The oncogenic activation of these nuclear oncogenes stimulates the cell into the synthetic phase (S phase), where it duplicates its genetic material before cell division. Nuclear control proteins that regulate entry into S phase include transcription factors such as c-myc, c-rel, c-jun, and c-fos, and cell cycle regulatory proteins such as E2F and cyclin D1 (PRAD1). These nuclear proto-oncogenes can work as transcription factors by binding to DNA in a sequence-specific manner and forming complexes with themselves or other proteins that will increase mRNA transcription of genes such as cyclin D that promotes cell division.

To understand how tumor cells evade antiproliferative signals, one must appreciate how the cell cycle is regulated—in particular, the G1 phase. It is during a cell’s transit through the G1 phase that it makes the decision to continue to the S phase and duplicate its genetic material or enter into a reversible state of quiescence (reversible growth arrest) or enter a permanent state of senescence (irreversible growth arrest). The Rb family of proteins is the most critical determinants of the fate of a G1 phase cell. When Rb protein is in a hypophosphorylated state, Rb protein blocks progression into S phase by sequestering E2F transcription factors that regulate the expression of genes that are essential for the transition from G1 to S phase. As previously discussed, the ECM can signal cell proliferation through integrin receptors. In addition, it can also produce antiproliferative signals through the secreted protein TGF-β. At the molecular level, TGF-β inhibits Rb protein phosphorylation and prevents the release and activation of the E2F family of transcription factors. It does this through the induction of inhibitors such as p21 and p15INK4B that inhibit the activity of the kinases that are essential to phosphorylate Rb protein. Thus, loss of Rb or failure to induce p21 and p15INK4B will aid cells in escaping antiproliferative signals.

In addition to extracellular antiproliferation signals, endogenous proteins that down-regulate signal-amplifying kinases are also important in keeping signal transduction cascades in check. For example, the NF1 protein is a GTPase-activating protein that facilitates the hydrolysis of guanosine triphosphate (GTP) by ras. When ras protein is complexed with guanosine diphosphate (GDP), it is inactive; when it is bound to GTP, it is active. Individuals afflicted with neurofibromatosis lack NF1 protein and have lower levels of GTPase activity and more ras protein complexed
with GTP, resulting in a more active ras protein. Therefore, lack of NF1 protein activity will result in enhanced activity of ras protein similar to what is found in ras-transformed cells as already described. The NF1 protein and the ras protein are excellent examples of how a tumor suppressor gene and an oncogene work in unison to control a signal transduction pathway.

Failure to Commit Suicide (Apoptosis)

Two major pathways that mediate cell death emanate either from the cell membrane or from the mitochondrion (Fig. 18.12). The signals transmitted by each pathway results in the activation of intracellular cysteine proteases, termed caspases, that cleave a diverse and ever-increasing number of substrates, including themselves, at aspartic acid residues. Caspases can broadly be divided into initiator caspases and effector caspases. The binding of ligands such as Fas to specific death receptors on the cell surface induces receptor activity and the recruitment of the initiator procaspase 8. The recruitment of procaspase 8 proteins near to each other results in active caspase 8 and effector caspases such as caspases 3, 6, and 7, which are responsible for the ultrastructural changes in the cell. In response to mitochondrial-dependent cell death,
activation of initiator caspases (e.g., caspase 9) is achieved by proteolytic cleavage of their inactive pro-forms through their recruitment and interaction with specific adapter proteins. The adapter proteins are in turn regulated by the mitochondria through the release of cytochrome \( c \). The release of cytochrome \( c \) from the mitochondria is controlled by the Bcl-2 protein family, which is composed of proapoptotic regulators such as Bax, Bak, Bid, and Bim and antiapoptotic family members Bcl-2, Bcl-xl, and Mcl-1. The Bcl-2 oncogene is the prototypical example of a membrane-associated oncogene whose overexpression protects the cell from death-inducing stimuli by various agents.

How does Bcl-2 protect cells from undergoing apoptosis? Both proapoptotic and antiapoptotic Bcl-2 family members can form dimers with themselves (homodimers) or with other family members (heterodimers). This ability to form dimers with a pro-death-promoting family member has been proposed as one mechanism of how Bcl-2 prevents apoptotic cell death that is signaled by the release of cytochrome \( c \) from the mitochondria. In the cytoplasm, cytochrome \( c \) forms a complex with the adapter protein Apaf-1, and together they recruit the inactive form of the initiator caspase, procaspase 9. In this complex, caspase 9 becomes activated and in turn activates effector caspases such as caspases 3, 6, and 7 through proteolytic cleavage. Cell lines deficient in caspases 3 and 9 exhibit substantially reduced levels of apoptosis during development and in response to exogenous stress-inducing stimuli (Fig. 18.13).

The \( p53 \) tumor suppressor gene is an important modulator of oncogene-induced apoptosis. Levels of \( p53 \) are kept low in unstressed cells through the binding of a specific E3-like ubiquitin ligase, murine double minute 2 (Mdm2). Binding of Mdm2 to the N-terminus of \( p53 \) results in the complex being shuttled to the cytoplasm, where it is quickly degraded by the proteosome. However, in response to various stresses, including ionizing radiation, serum starvation, and hypoxia, \( p53 \) protein levels increase both through protein stabilization and increased protein synthesis. Stabilization of \( p53 \) in response to
stress is thought to occur through several mechanisms, including prevention of Mdm2 binding and phosphorylation of p53. Once stabilized, p53 is a powerful proapoptotic molecule capable of transcriptionally activating gene expression by sequence-specific DNA binding to regulatory sequences. Transcriptional targets of p53 that induce apoptosis include bax, puma, noxa, and perp and provide a link between the tumor suppressor activity of p53 and apoptosis. This list of p53-regulated apoptotic genes is always growing and very dependent on the cell type and stress. Many of the mutations in the p53 gene found in human tumors are found within the DNA-binding domain (DBD), highlighting the importance of this region to the role of p53 as a tumor suppressor and its ability to induce apoptosis. However, recent reports in the literature indicate a new role for p53 in the mitochondria, where it may function directly to release cytochrome c to initiate the caspase cascade and apoptosis, bypassing the need for its transcriptional activity.

Seemingly paradoxical to its role in proliferation and oncogenic transformation is the fact that overexpression of the myc oncogene primes cells for apoptotic cell death under growth-restrictive conditions generated by nutrient deprivation or low oxygen conditions (Fig. 18.14). It is this paradox that has set forth the hypothesis that myc deregulation results in a cellular state where increased proliferation or apoptotic death are both equally possible, depending on the cellular microenvironment and the activity of certain crucial genetic determinants such as the p53 tumor suppressor gene. Evidence has accumulated that oncogenes such as myc and the adenovirus E1A gene increase p53 protein stabilization and sensitize cells to killing by growth-restrictive conditions. Loss of p53 through mutation or functional inactivation severely attenuates the sensitivity of these same oncogene-expressing cells to stress-induced apoptosis. Analysis of cells deficient in p19ARF (a cell cycle inhibitor protein) indicates that myc signals to p53 through p19ARF. Loss of p19ARF attenuates the sensitivity of myc-expressing cells to apoptosis even in the presence of wild-type p53, suggesting that myc needs to signal p19ARF to activate p53-dependent apoptosis. Furthermore, genetic analysis of tumor cells indicates that they possess either p53 mutations or p19ARF mutations, but rarely both. Implicit in this observation is that myc deregulation favors proliferation and that a growth-restrictive state induced by DNA
into senescence. The observations that led to this discovery were based on the ability of the \( \text{ras} \) oncogene alone to transform and immortalize primary rodent fibroblasts only if they lacked the \( p53 \) tumor suppressor gene or the cell cycle inhibitor \( p16 \) that regulates the \( Rb \) pathway. The loss of \( p53 \) and \( p16 \) is also important for human cell immortalization. In addition, senescent cells have elevated levels of \( p53 \) or \( p16 \), suggesting that senescence is ultimately an irreversible cell cycle arrest. In primary cells, such as fibroblasts, the activation of the constitutive growth signal by the \( \text{ras} \) oncogene will induce the activity of \( p16 \) and \( p53 \) that will counter this signal and result in the induction of cellular senescence. Therefore, for a cell to develop independence of extracellular mitogenic growth signals, it must develop mutations in pathways that send a continuous proliferation signal as well as in pathways that attempt to restrict this signal. Cell immortalization can be viewed as a competing process that requires both the activation of dominant activating oncogenes to induce proliferation and the loss of recessive tumor suppressor genes that induce a cell cycle arrest in response to this constitutive activating signal.

Although cellular senescence can be delayed by mutations in the \( p53 \) and \( Rb \) pathways, cells will ultimately encounter another junction in their road to transformation, namely, crisis (Fig. 18.15). This term crisis is highly appropriate, as this roadblock to cell immortalization results in chromosomal rearrangements and cell death. Less than 1 in 10 million cells that enter crisis survives and gains the ability to replicate indefinitely. One clue to what drives cells to the crisis stage comes from the end-to-end fusions of chromosomes. From this, it is apparent that crisis results from the progressive shortening of the protective caps (telomeres) on the ends of chromosomes.

Mammalian telomeres consist of long arrays of the repeat sequence TTAGGG that range in length anywhere from 1.5 to 150 kb. Each time a normal somatic cell divides, the terminal end of the telomere is lost; successive divisions lead to progressive shortening, and after 40 to 60 divisions, vital DNA sequences are lost. At this point, the cell cannot divide further and undergoes senescence.
Telomere length has been described as the “molecular clock,” because it shortens with age in somatic tissue cells during adult life. Stem cells in self-renewing tissues, and cancer cells in particular, avoid this process of aging by activating the enzyme telomerase. Telomerase is a reverse transcriptase that polymerizes TTAGGG repeats to offset the degradation of chromosome ends that occurs with successive cell divisions; in this way, the cell becomes immortal. Telomere shortening inhibits tumor expansion when the $p53$ tumor suppressor gene is intact. Although it has not been proven, telomeres seem to engage the $p53$ pathway by inducing a damage response signaled through the ATM pathway (Chapter 2). Studies have shown that telomere shortening leads to senescence and tumor suppression in cells with an intact $p53$ pathway. Surprisingly, telomere shortening accelerates tumorigenesis in cells that were deficient in $p53$ activity. The ability of shortened telomeres to accelerate tumorigenesis in $p53$-deficient cells results from increased chromosomal instability and rearrangements and gene amplification.

**Angiogenesis**

Angiogenesis—the recruitment of new blood vessels to regions of chronically low blood supply—is essential for the progression of solid tumors to malignancy. Increasing evidence supports the hypothesis that tumor angiogenesis is controlled by an “angiogenic switch,” a physiologic mechanism involving a dynamic balance of angiogenic factors that include both inhibitors and inducers. Numerous angiogenic factors have been identified, including specific endothelial cell growth factors (e.g., vascular endothelial growth factor, or VEGF), cytokines and inflammatory agents (e.g., tumor necrosis factor $\alpha$, or TNF-$\alpha$, and interleukin-8, or IL-8), fragments of circulatory system proteins (e.g., angiotatin and endostatin), and extracellular matrix components (e.g., thrombospondins, or TSPs). Presumably, this diversity of angiogenic factors reflects a strict requirement for controlling angiogenesis under normal physiologic conditions and in response to oncogenic events by modulating the expression of both angiogenic inducers and inhibitors.

Although the list of proangiogenic growth factors is expanding, VEGF was the first growth factor isolated that could stimulate endothelial proliferation and migration. In tumors, VEGF can be regulated by oncogenic stimuli such as $ras$ and $raf$, hypoxia, and deregulated growth factor receptor signaling. Both tumor cells and host stromal cells produce VEGF. However, the target for VEGF lies on the cell surface of endothelial cells. These cells possess several specific transmembrane receptor tyrosine kinases.
that bind VEGF, which in turn initiate endothelial migration and proliferation. Regarding neoangiogenesis, VEGF receptor II is the most important for stimulating new blood vessel formation. Studies have shown that blocking the binding of VEGF to its receptor inhibits tumor angiogenesis and tumor growth. These findings have led to the development of new antibody approaches for antiangiogenesis therapy for clinical use.

Tumors such as renal cell carcinomas that possess mutations in the von Hippel-Lindau gene (VHL) exhibit high aerobic expression of proangiogenic genes such as VEGF, whereas reintroduction of wild-type VHL substantially reduces VEGF levels to those found in untransformed cells. The mechanism underlying this observation is that VHL inhibits hypoxia-inducible factor (HIF) levels under aerobic conditions by targeting HIF for ubiquitin-mediated degradation. Cells that have lost VHL are impaired in their ability to degrade HIF and have constitutive elevated levels of HIF and HIF target genes, of which VEGF is one under aerobic conditions (see Chapter 26 for further explanation).

Thrombospondin 1 (TSP-1) is a secreted adhesive glycoprotein that has been shown to have antiangiogenic activity by preventing angiogenic factors such as VEGF from binding to their target receptors. Compelling evidence of an antiangiogenic function for TSP-1 is provided by studies showing that activation of the angiogenic switch in Li–Fraumeni fibroblasts lacking functional p53 is associated with diminished TSP-1 expression. Furthermore, introduction of functional p53 into carcinoma cells generates an antiangiogenic activity involving induction of TSP-1 expression. Reports that endogenous TSP-1 expression can be induced by p53, repressed by oncogenic signals such as c-jun overexpression, and silenced by DNA methylation, indicate that down-regulation of TSP-1 contributes to oncogenesis. As both p53 and c-jun are inducible by hypoxia, the regulation of TSP-1 expression by these proteins suggests that it is also a hypoxia-responsive angiogenic inhibitor.

In summary, TSP-1 and VEGF are genetically controlled by both tumor suppressor gene and proto-oncogene activity, providing molecular mechanisms that could contribute to the switch to the angiogenic phenotype when these controls are deregulated during oncogenesis.

**Invasion and Metastasis**

Many years ago, Paget realized that cancer spreads in defined patterns and is influenced by both lymph and blood flow patterns as well as the tissue being invaded. He proposed that a metastatic cell is analogous to a vegetable seed, in that, without the right soil conditions, it would never grow. Malignant tumor cells become locally invasive and escape their tissue confines by invading the substratum beneath them, before they can colonize to distant tissues (Fig. 18.16).

Local invasion necessitates the breakdown of epithelial integrity that is influenced by cell–cell and cell–matrix interactions through the loss of adhesion molecules such as E-cadherin (E-CAD). Decreased expression or impaired function of E-CAD leads to deregulated intercellular adhesion and increases the invasive growth and spread of the primary tumor, whereas overexpression reduces the invasive and metastatic growth of transformed cells. In gastric carcinomas and lobular breast carcinomas, E-CAD has been found to be mutated and functionally inactivated. Thus, loss of E-CAD can permit local invasion of tumor cells and may be a common step in invasion. In addition to E-CAD, cell adhesion molecules such as neural cell adhesion molecule (NCAM) also appear to play an important role in invasion. Their expression is decreased in invasive cells and when overexpressed can decrease invasion and metastasis. One final group of proteins that contribute to the invasive capability of tumor cells are integrins, which relay signals from the ECM to epithelial cells. In this case, the cell surface repertoire of integrins changes when tumor cells become invasive.

The genetic circuits that regulate metastasis remain mainly undiscovered and elusive. Some broad concepts about metastasis have been proposed, such as the need for proteases to degrade the ECM. However, which proteases and how they are regulated differently in tumor cells compared with untransformed cells are unknown. An important question is whether or not tumor cells must acquire additional mutations to invade and metastasize. In returning to Paget’s concept of the seed and the soil, only a small percentage of tumor cells are able to metastasize.
The identity of gatekeepers varies with each tissue, such that inactivation of a given gene predisposes to specific forms of cancer; inherited mutations in adenomatous polyposis coli (APC) predispose to colon cancer, for example; mutations in VHL predispose to kidney cancers; and so on. Because these gatekeeper genes are rate limiting for tumor initiation, they tend to be mutated in many cancers. They can arise both through somatic or germ line mutations.

By contrast, inactivation of caretaker genes does not directly promote the growth of tumors, but leads, instead, to genomic instability that only indirectly promotes growth by causing an increase in mutation rate. This increase in genetic instability can greatly accelerate the development of cancers, especially those that require numerous mutations for their full development. Colon cancer is a good example. The targets of the accelerated mutation rate that occurs in cells with defective caretakers are the gatekeeper tumor suppressor genes, oncogenes, or both. Table 18.3 provides examples of cancer predisposition syndromes caused by mutations.

Some die rapidly by apoptosis, some remain in the circulation, and a small number invade and colonize other tissues. One noteworthy point is that loss of apoptosis in response to detachment from neighboring cells and the ECM (anoikis) is essential for metastatic spread. Metastasis represents a major challenge in the treatment of cancer, especially as the ability to control local tumor growth is increasing through the combination of surgery and radiotherapy. Metastasis research represents the most critical frontier in cancer research.

**THE CONCEPT OF GATEKEEPERS AND CARETAKERS**

It appears that most tumor suppressor genes can be broadly divided into two classes that have been called “gatekeepers” and “caretakers.” Gatekeepers are genes that directly regulate the growth of tumors by inhibiting cell division or promoting cell death. The function of these genes, therefore, is rate limiting for tumor growth; both alleles (maternal and paternal) must be lost or inactivated for a tumor to develop. Predisposed individuals inherit one damaged copy of such a gene and so require only one additional mutation for tumor initiation. The identity of gatekeepers varies with each tissue, such that inactivation of a given gene predisposes to specific forms of cancer; inherited mutations in adenomatous polyposis coli (APC) predispose to colon cancer, for example; mutations in VHL predispose to kidney cancers; and so on. Because these gatekeeper genes are rate limiting for tumor initiation, they tend to be mutated in many cancers. They can arise both through somatic or germ line mutations.

By contrast, inactivation of caretaker genes does not directly promote the growth of tumors, but leads, instead, to genomic instability that only indirectly promotes growth by causing an increase in mutation rate. This increase in genetic instability can greatly accelerate the development of cancers, especially those that require numerous mutations for their full development. Colon cancer is a good example. The targets of the accelerated mutation rate that occurs in cells with defective caretakers are the gatekeeper tumor suppressor genes, oncogenes, or both. Table 18.3 provides examples of cancer predisposition syndromes caused by mutations.
in DNA repair and stability genes. The evidence is highly suggestive that mutations in these predisposition genes increase the rate of acquiring cancer after exposure to DNA-damaging agents such as ionizing radiation.

Evidence for this comes from a review by Swift of 161 families affected by AT. In this prospective study, new cases of cancers were observed in blood relatives of persons with AT (of whom about half may be heterozygotic), in those who are definite heterozygotes (obligates), and in spouses who were assumed to be normal but who lived in the same environment. This extensive study also divided blood relatives of AT homozygotes into those with and those without a “radiation history.” A radiation history was interpreted loosely as fluoroscopy of the chest, back, or abdomen, therapeutic irradiation, or occupational exposure. Table 18.4 shows the results of the survey: 53% of blood relatives with cancer had a radiation history, compared with 19% of those without cancer.

From these data, the study purported to show that AT heterozygotes are very sensitive to radiation-induced cancer; a control study of this kind does not provide proof of this, but the possibility certainly exists. It is a challenging and sobering thought to diagnostic radiologists that a proportion of the women

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**TABLE 18.3** DNA Repair and Stability Genes and Their Associated Syndromes

<table>
<thead>
<tr>
<th>Suppressor</th>
<th>Syndrome</th>
<th>Tumor</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>ATM</strong></td>
<td>Ataxia-telangiectasia</td>
<td>Leukemia, lymphoma</td>
</tr>
<tr>
<td><strong>XP</strong></td>
<td>Xeroderma pigmentosum</td>
<td>Skin</td>
</tr>
<tr>
<td><strong>BRCA1</strong></td>
<td>Hereditary breast cancer 1</td>
<td>Breast</td>
</tr>
<tr>
<td><strong>BRCA2</strong></td>
<td>Hereditary breast cancer 2</td>
<td>Breast, ovary</td>
</tr>
<tr>
<td><strong>FANC</strong></td>
<td>Fanconi anemia</td>
<td>Leukemia</td>
</tr>
<tr>
<td><strong>NBS</strong></td>
<td>Nijmegen breakage syndrome</td>
<td>Lymphoma</td>
</tr>
<tr>
<td><strong>hMSH2</strong></td>
<td>Hereditary nonpolyposis colorectal cancer</td>
<td>Colon</td>
</tr>
<tr>
<td><strong>hMLH1</strong></td>
<td>Hereditary nonpolyposis colorectal cancer</td>
<td>Colon</td>
</tr>
<tr>
<td><strong>hMSH6</strong></td>
<td>Hereditary nonpolyposis colorectal cancer</td>
<td>Colon</td>
</tr>
<tr>
<td><strong>hPMS1</strong></td>
<td>Hereditary nonpolyposis colorectal cancer</td>
<td>Colon</td>
</tr>
<tr>
<td><strong>hPMS2</strong></td>
<td>Hereditary nonpolyposis colorectal cancer</td>
<td>Colon</td>
</tr>
</tbody>
</table>

---

**TABLE 18.4** Breast Cancer and Radiation in 161 Families with Ataxia-Telangiectasia

<table>
<thead>
<tr>
<th>Blood Relatives with Cancer</th>
<th>Blood Relatives without Cancer</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number</td>
<td>19</td>
</tr>
<tr>
<td>With Radiation Historya</td>
<td>10/19 = 53%</td>
</tr>
</tbody>
</table>

routinely screened by mammography may be exquisitely sensitive to radiation-induced carcinogenesis because of repair deficiencies associated with being heterozygotic for AT.

**MISMATCH REPAIR**

Interest in MMR genes heightened with the discovery that they were responsible for the mutator phenotype associated with a predisposition for hereditary nonpolyposis colon cancer (HNPCC) and possibly other familial cancers. The initial clue to this novel molecular mechanism was the discovery of deletions of long monotonic (dA-dT) runs in a subset of human colon cancers. Soon after, insertions or deletions at mononucleotide, dinucleotide, and trinucleotide repeat sequences were discovered in subsets of colon tumors as well as in various colon cancers from individuals with HNPCC. This phenotype also has been detected in several other types of human malignancies, especially those associated with type 2 Lynch syndrome. These various investigations culminated in the identification and cloning of the human *hMSH2* gene, which maps to a locus linked to HNPCC on chromosome 2p21–22 and whose homologues in *Saccharomyces cerevisiae* and *Escherichia coli* are involved in the process of DNA MMR.

The primary function of MMR genes in *E. coli* appears to be to scan the genome as it replicates and to spot errors of mismatch as the DNA is replicated, that is, as the new strand is laid down using the stable methylated strand as a template. A growing number of human genes have been associated with HNPCC by means of linkage analysis and studies of mutational mapping. Table 18.5 lists human MMR genes associated with HNPCC. The MMR process in yeast and bacteria involves a large number of proteins, and so it is likely that additional causes of HNPCC remain to be uncovered.

Cells with defective or nonfunctioning MMR genes can be identified by two quite different techniques:

1. By using a selectable reporter system that inserts an exogenous long repeat sequence into the cells in question and measures the mutation rate in it.
2. By measuring the mutation rate in one or more of the many endogenous repeat sequences that already exist in every human cell—the so-called microsatellite instability assay. Microsatellite instability appears to be a factor of some importance in various human tumors.

Both techniques have strengths and weaknesses and are far from perfect.

**HERITABLE SYNDROMES THAT AFFECT RADIOSensitivity, GENOMIC INSTABILITY, AND CANCER**

The DNA damage response in mammalian cells is comprised of multiprotein complexes that sense, signal, and respond to DNA strand breaks. Disruption of the function of these multiprotein complexes by mutation of a single gene leads to cancer-prone syndromes that are characterized by hypersensitivity to DNA damage and genomic instability. Syndromes that exhibit hypersensitivity to ionizing radiation are discussed here.

**Ataxia Telangiectasia**

AT is a rare autosomal recessive disease in which afflicted individuals present with progressive cerebellar ataxia caused by increased loss of...
the ATM kinase is activated by DNA damage and chromatin alterations throughout the cell cycle, it uses different downstream effectors to mediate a checkpoint response in each of the different phases of the cell cycle. For example, the activation of a G1 checkpoint is in large part mediated by ATM signaling to p53, which results in the transcriptional induction of the p21 cell cycle inhibitor. Perhaps the most distinctive hallmark of AT cells is their failure to arrest in S phase in response to DNA damage. This phenomenon has been termed radioreistant DNA synthesis, and at least two pathways regulated by ATM control the passage of cells through S phase.

Much is known about the way in which ATM and its family members control cell cycle checkpoints, but the mechanism by which loss of ATM leads to increased radiosensitivity is still under investigation. It may involve both

Purkinje cells in the cerebellum as well as oculo-cutaneous telangiectasia. Patients with AT are immune deficient and have a high incidence of cancer, especially of the reticular endothelial system. Patients with AT exhibit a hypersensitive skin reaction to ionizing radiation and DNA breaking agents but not to ultraviolet light. Both lymphocytes and fibroblasts derived from these individuals are also hypersensitive to killing by ionizing radiation.

The genetic defect responsible for the AT phenotype is the result of mutations in the ATM gene. The ATM gene encodes for a kinase that phosphorylates a serine or threonine that is followed by a glutamine motif (S/T-Q) in target proteins (Fig. 18.17). ATM is involved in the rapid response of cells to DNA DSBs (it signals to the DNA repair machinery) as well as the activation of cell cycle checkpoints. However, it is important to recognize that although

![Diagram](image_url)

**FIGURE 18.17** Activation of the ATM pathway by an ionizing radiation-induced DNA DSB. The model proposed by Kastan and Bakkenist indicates that ATM exists as a dimer and undergoes rapid autophosphorylation in response to DNA strand breaks and chromatin conformation changes. This results in dissociation of the dimer into a monomer and the phosphorylation of numerous substrates involved in DNA repair (H2AX, SMC1, BRCA1, and p53BP1) and cell cycle control (p53 and Chk2). Recent studies have shown that the MRN complex can function upstream of ATM by sensing and localizing to DNA breaks and can also be a target of ATM.
Nijmegen Breakage Syndrome

NBS is a direct phosphorylation target of ATM and forms a complex with Rad50 and MRE11. This complex possesses nuclease activity that is necessary for DNA DSB repair. The relationship between NBS and ATM is complex, as ATM phosphorylates NBS in response to DNA damage, but NBS may also be required for activation of some ATM checkpoint responses. Thus, the mechanism of AT, ATLD1, and NBS sensitivity to ionizing radiation is probably the same.

Fanconi Anemia

Eight different proteins give rise to the autosomal recessive disorder Fanconi anemia (FA), which is characterized by spontaneous chromosomal instability, sensitivity to interstrand DNA crosslinks and, in some cases, sensitivity to ionizing radiation. The Fanconi anemia, complementation group D2 (FANCD2) gene is thought to be a key player in the FA pathway because it is the only FANC gene conserved through evolution, and most of the other FANC proteins form a complex that results in the monoubiquitination of FANCD2 that results in its localization to foci that contain the breast cancer tumor suppressor genes BRCA1 and BRCA2 (also known as FANCD1). The identification of BRCA2 as an FA gene, together with its localization in foci with FANCD2, suggests an important role for FANC genes in homologous recombination. In addition, FANCD2 has been implicated in the DNA damage checkpoint response and is phosphorylated by the ATM kinase on serine 222. The roles of FANCD2 in the radiation response and crosslinks response are separable because inhibition of phosphorylation on serine 222 inhibits the FA-mediated radiation response, and inhibition of ubiquitination of lysine 561 results in hypersensitivity to crosslinking agents.

Clinical reports have suggested that tumors derived from patients with FA are hypersensitive to radiotherapy. However, fibroblasts derived from individuals that possess mutated forms of each of the eight different FA proteins were not always found to exhibit radiosensitivity and never to the same level as AT cells. Therefore, the clinical response of patients with FA to radiotherapy does not always correlate with intrinsic radiosensitivity of fibroblasts derived from the same patients, suggesting that additional genetic alterations in tumor cells from patients with FA

Seckel Syndrome

In mammalian cells, ATM belongs to the phosphatidylinositol 3-kinase-related kinase (PIKK) family and shares homology with two other family members—ATR and DNA-dependent protein kinase catalytic subunit (DNA-PKcs)—that are also activated by DNA strand breaks. At the molecular level, ATR and DNA-PKcs like ATM regulate DNA repair and cell cycle regulatory proteins. Interestingly, although ATM is not an essential gene, ATR is essential. Some individuals afflicted with Seckel syndrome—a rare autosomal recessive disorder characterized by microcephaly and abnormal development—can possess an alteration in the ATR gene that reduces its absolute quantity, but does not eliminate its activity. Patients with Seckel syndrome do not appear to be radiosensitive, because they possess some ATR activity. The third member of this family, DNA-PKcs, is an essential component of nonhomologous recombination and is the defect in murine severe combined immunodeficiency (SCID) syndrome. However, human patients with SCID do not possess DNA-PKcs mutations, but instead possess mutations in Artemis, a target of DNA-PKcs. Artemis-deficient human SCID cells are radiosensitive, and fibroblasts from afflicted individuals exhibit increased chromosomal instability.

Ataxia-Telangiectasia-Like Disorder

ATLD is an autosomal recessive disease caused by mutations in the gene MRE11 that forms a complex with Rad50 and NBS in irradiated cells. Cells derived from patients with ATLD are radiosensitive but repair their DNA DSBs similar to wild-type levels, leaving the mechanism for their radiosensitivity in question. ATLD1 cells are also defective in their checkpoint response following DNA damage.

direct mechanisms, such as direct phosphorylation of the cohesin family member structural maintenance of chromosomes (SMC) or Nijmegen breakage syndrome (NBS) protein, and indirect mechanisms, such as dysregulated homologous recombination. Lymphoid cells in ATM homozygotes often exhibit increased chromosomal instability, and the lymphomas that develop demonstrate the importance of DNA repair and chromosome maintenance in tumor suppression.
RADIATION-INDUCED SIGNAL TRANSDUCTION

Ionizing radiation can regulate the expression of early response genes such as c-fos, c-jun, and c-myc, genes that control lipid signaling such as acid sphingomyelinase, and the PIKK family that includes ATM and DNA-PK.

Early Response Genes

The activation of early response genes by ionizing radiation suggests that radiation in some way mimics the mitogenic activation of quiescent cells. Evidence for the importance of such effects is that the overexpression of proto-oncogenes such as c-ras and c-raf, whose products are intermediates in the pathways of mitogenic signal transduction, is associated with radiation resistance. The induction of these pathways results in gene activation through elements such as AP-1, serum response element, and cAMP response element-binding protein (CREB).

The role of these early response genes is not clear at the present time, but the stimulation of signal transduction pathways and activation of transcription factors may enhance secondarily the response of the cell to radiation in terms of repair and cell cycle arrest. In addition, the activation of these early response genes, in turn, could provide a mechanism for secondary stimulation of various late response genes such as TNF-α, PDGF, FGF, and IL-1. Activation of these genes may allow the cell to adapt to acute changes in the microenvironment and may be responsible for some of the chronic responses of the cell to ionizing radiation, including apoptosis.

The Ceramide Pathway

Exposure of cells to DNA damage induces the production of ceramide through two major pathways. In one pathway that is activated by ionizing radiation and DNA-damaging agents, ceramide is generated from sphingomyelin hydrolysis by the enzyme acid sphingomyelinase (Fig. 18.18). This results in the catalysis of up to one-half of total cellular sphingomyelin, most of which would be presumably associated with the plasma membrane. Ceramide can also be generated by ceramide synthase, which is inhibited by the ATM kinase. Therefore, in response to ionizing radiation, individuals who have lost ATM have elevated levels of ceramide.
Synthase and increased apoptosis of certain cell types (e.g., crypt cells). It is important to note that ionizing radiation can activate both nuclear and membrane–cytoplasmic signal transduction pathways that can lead to either cell cycle arrest or, alternatively, apoptosis.

In summary, although there is little doubt that the lethal effects of ionizing radiation result from extensive DNA damage leading to chromosomal aberrations, radiation can also stimulate signal transduction pathways that can lead to cell cycle arrest or cell death by apoptosis, depending on the dose of radiation and genetic background of the cell.

**THERAPEUTIC EXPLOITATION OF RADIATION-ACTIVATED SIGNAL TRANSDUCTION PATHWAYS**

The molecular exploitation of oncogenic protein products should soon be possible. For example, it is still controversial whether the Erb-2 receptor is predictive of a poor prognosis for breast cancer. However, as already stated, many growth factor receptors possess truncated extracellular ligand-binding domains. Therefore, it is possible to specifically target monoclonal antibodies coupled to radionuclides or chemotherapeutic agents to these truncated receptors. It also seems
possible to reverse or decrease the oncogenic effects of certain oncogenes by altering essential interactions with subcellular compartments. For example, the ras oncogene protein product requires the addition of an isoprenoid lipid (a process termed prenylation) so that it can attach to the cell membrane where it is active. Prenylation of ras is performed by farnesyltransferase or geranylgeranyltransferase. Past studies have shown that the first generation of farnesyltransferase inhibitors was able to alter oncogenic ras activity and decrease growth of transformed cells. Because ras is found to be mutated in various human tumors, and ras requires this modification for activity, the farnesyltransferase would therefore be an excellent candidate enzyme to inhibit.

A second example of an oncogenic target is the NF-κB transcription factor (a member of the c-rel family of proto-oncogenes), which is found to be constitutively active in various tumor cells by multiple mechanisms (Fig. 18.19). If NF-κB activity is inhibited in these tumor cells, they fail to proliferate in soft agar, become more sensitive to apoptotic cell death induced by ionizing radiation or chemotherapy, and exhibit decreased migration and apoptosis. What makes this transcription factor a good target for therapy is that it is normally held in the cytoplasm by an inhibitory molecule called IκB (inhibitor of κB). If IκB is overexpressed or prevented from degradation, then NF-κB will be held in check in the cytoplasm, where it is inactive and kept out of the nucleus. At present, several drugs exist that could therapeutically inhibit IκB degradation. These are but several examples of molecular strategies that target key components of oncogene regulation for the treatment of cancer.

SUMMARY OF PERTINENT CONCLUSIONS

- Cancer is thought to be a clonal disorder.
- The control of cell proliferation is the consequence of signals that may be positive or negative.
- Gain-of-function mutations can activate oncogenes, which are positive growth regulators; tumor suppressor genes are a negative growth regulator.
- Oncogenes are genes found in either a mutant or an abnormally expressed form in many human cancers.
- Oncogenes can be activated by retroviral integration, point mutation, a chromosomal rearrangement such as a translocation, or gene amplification.
- Some human leukemias and lymphomas appear to be caused by specific chromosomal translocations that lead to oncogene activation in several different ways.
- Knudson postulated that all types of retinoblastoma involve two separate mutations. In sporadic retinoblastoma, both mutations occur somatically in the same retinal cell; therefore, this condition is rare. In the heritable form, one of the two mutations is inherited from a parent and is present in all retinal cells, so that the second mutation would occur somatically in any of these cells; hence, the incidence is close to 100%.
- There are many tumor suppressor genes whose location and function are known; the two most intensively studied are p53 and Rb.
- Because oncogenes are gain-of-function mutations, only one copy needs to be activated; that is, they act in a dominant fashion. Tumor suppressor genes involve loss-of-function mutations, so that both copies must be lost; that is, they act in a recessive fashion.
- Somatic homozygosity is the process by which one chromosome of a pair is lost, a deletion occurs in the remaining chromosome, and the chromosome with the deletion replicates.
- Carcinogenesis appears to be a multistep process with multiple genetic alterations occurring. An attractive model of carcinogenesis includes the idea that an early step causes a mutation in a gene in one of the families responsible for the stability of the genome. This leads to a mutator phenotype, so that multiple further changes are likely in the progression of the cancer.
- Telomeres cap the ends of chromosomes; they are long arrays of TTAGGG repeats. Each time a normal somatic cell divides, the terminal end of the telomere is lost; after 40 to 60 divisions, the cell undergoes senescence. Stem cells and cancer cells activate telomerase, which maintains telomere length, so the cell becomes immortal.
- AT is an autosomal recessive disorder that is caused by a defect in the ATM kinase. AT cells fail to activate checkpoints in re-
response to DNA damage, exhibit increased genomic instability at the chromosome level, and have an increased risk of lymphomas. AT cells and individuals are hypersensitive to ionizing radiation.

- The **ATR** gene expression (AT and Rad3 related) is decreased in patients with Seckel syndrome. Although it belongs to the same PIKK family as ATM, it is an essential gene at the cellular level. It has an important role in responding to DNA breaks in S phase.

- RecQ genes encode helicases that play a critical role in protecting replication forks. Decreased expression of RecQ genes results in aberrant DNA replication and genomic instability.

- There are at least three human syndromes involved in RecQ deficiency: BLM, WS, and RTS. A common feature of all these syndromes is increased chromosomal instability. These disorders do not result in hypersensitivity to ionizing radiation. However, BLM cells are sensitive to alkylating agents and mitomycin C.

- NBS is a rare disorder that results in increased cancer incidence. Cells defective in NBS lack an S phase checkpoint and are radiosensitive.

- Patients with ATLD are clinically similar to patients with AT except that their defect lies in the **MRE11** gene. Cells from these patients are also sensitive to ionizing radiation.

- Patients with FA are characterized by their hypersensitivity to crossinglinking agents. Although fibroblasts derived from these patients are not sensitive to ionizing radiation, tumors arising in these patients are hypersensitive. The reasons for this are currently unknown.

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Oncogenes and Tumor Suppressor Genes


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**An Introduction to Radiation Oncology**


Genes and Ionizing Radiation


DOSE–RESPONSE RELATIONSHIPS

Radiation biology applied to clinical radiotherapy is concerned with the relationship between a given absorbed dose of radiation and the consequent biologic response; of particular interest are factors that modify this relationship. With increasing radiation dose, radiation effects may increase in severity (i.e., grade), in frequency (i.e., incidence), or both. In most cases, it is the relationship between dose and incidence that is important. Such dose–response curves have a sigmoid (S) shape, with the incidence tending to zero as dose tends to zero and the incidence tending to 100% at very large doses. This applies to both tumor control and normal tissue complications.

A simple example is shown in Figure 19.1. Tumor control probability (TCP) is plotted as a function of total dose, and the incidence of normal tissue complications is also plotted as a function of dose. What is illustrated is a favorable situation where the tumor is more radiosensitive than the normal tissue. In the case of tumor control, the shape can be explained solely from the random nature of cell killing (or clonogen survival) after irradiation and the need to kill every single cell to achieve a cure.

For most normal tissue end points, the biologic interpretation of the S shape of the relationship is not obvious. Some researchers have evoked a hypothetical tissue rescue unit (TRU), arguing that tissue breakdown occurs when the number of TRUs falls below a critical level; however, this explanation is questionable.

Therapeutic Ratio (Therapeutic Index)

The ratio of the tumor response for a fixed level of normal tissue damage has been called either the therapeutic ratio or the therapeutic index. In the hypothetical example in Figure 19.1, there is a favorable therapeutic ratio, because a 30% probability of tumor control is possible for a 5% incidence of complications.

The time factor is the one parameter that has been most often manipulated to increase this ratio; hyperfractionation, for example, produces a greater sparing of late-responding normal tissue than tumor control. Another strategy often quoted, although seldom achieved in practice, is to add a drug or radiosensitizer that potentiates the tumor control without potentiating the radiation damage to normal tissue. In practice, it does not need to be as clear-cut as this; it would suffice for the drug to increase tumor control to a greater extent than it increases normal tissue damage. This would result in a therapeutic gain. This is illustrated in Figure 19.2. The addition of the drug moves the tumor control curve to the left farther than the normal tissue damage curve; that is, the drug has greater cytotoxic effect on the tumor than on the normal tissue. Consequently, with the combined modalities, an improved tumor control probability is possible for the same probability of normal tissue injury.
mechanism and importance are still under investigation. One additional point to consider is that ionizing radiation also induces a form of senescence or permanent growth arrest in which cells are still metabolically active, but reproductively inhibited. This is best exemplified by fibroblasts that when irradiated in cell culture, stay attached to plates for weeks but never divide. However, they are able to secrete growth factors and mitogens that promote the growth of tumor cells. Senescence has been largely studied in cell culture in the laboratory, and only recently has evidence been accumulating that it occurs in tissues. Senescence is also discussed in more detail in Chapter 18.

Most cell lines cultured in vitro die a mitotic death after irradiation; that is, they die attempting to divide. This does not necessarily occur at the first postirradiation mitosis; the cell may struggle through one, two, or more mitoses before the damaged chromosomes cause it to die, attempting the complex task of cell division. Time-lapse films of irradiated cells cultured in vitro clearly show this process of mitotic death, which is the dominant cause of death if reproductive integrity is assessed in vitro as described in Chapter 3. It is not, however, the only form of cell death. Programmed cell death, or apoptosis, occurs in normal tissues and neoplasms, in mammals and amphibians, in the embryo, and the adult. It is implicated, for example, in tissue involution such as the regression of the tadpole tail during metamorphosis. It is the programmed cell death that is common during embryonic development. It also can occur after irradiation. Apoptosis, like mitosis, comes from the Greek word meaning “falling off,” as of petals from flowers or leaves from trees.
Apoptosis is characterized by a stereotyped sequence of morphologic events, which take place in two discrete phases. In the first phase, cells condense and bud to produce many membrane-enclosed bodies. In the second phase, these bodies are phagocytized and digested by nearby tissue cells. The characteristic “laddering” of DNA that occurs during apoptotic death is illustrated in Chapter 3. Apoptosis characteristically affects scattered individual cells. If apoptosis affects cells in tissues, the resulting apoptotic bodies are squeezed along the intercellular spaces and are either shed from the epithelial surface or rapidly phagocytized by nearby cells. The cells surrounding those being deleted merely close ranks, and there is no tissue disorganization that occurs after necrosis.

Autophagic cell death has been classified with apoptotic cell death but, in fact, is distinct and is controlled by a unique set of genes. Apoptosis and autophagy may be linked because cells defective in some forms of apoptosis exhibit increased levels of autophagy. The term implies that cells cannibalize themselves in the hope of generating adenosine triphosphate (ATP) and macromolecule precursors to survive. Growth factor withdrawal, inhibition of protein synthesis, hypoxia, and ionizing radiation are all potent inducers of autophagy. Although autophagy can be found in cells dying by stress, it is unclear whether it represents a drastic means for the cell to survive by digesting part of itself, or whether it actually promotes cell death. Evidences for both possibilities exist in different cell types and will require further studies to clarify its role in irradiated cells.

Necrotic cell death occurs when cells are devoid of precursors to generate energy, such as glucose, are exposed to decreased pH, or exhibit changes in osmolarity. Ionizing radiation is also able to induce necrotic cell death. It differs from other forms of cell death because it does appear to have a definitive genetic program that controls it, although there have been reports that prove otherwise.

The long-standing concept that ionizing radiation kills cells through direct exposure requires amendment. The demonstration that cells not exposed to ionizing radiation can be killed by being close to irradiated cells has coined the term “bystander induced cell death.” Therefore, the possibility exists that ionizing radiation can influence cell killing outside the field of radiation. This has been experimentally shown for tumors and is known as the abscopal effect. Such effects for normal tissues probably also exist in lymphopenia found after radiotherapy to non-lymphoid organs, and most probably represent an immune response.

**ASSAYS FOR DOSE–RESPONSE RELATIONSHIPS**

Most experimental techniques are available to obtain dose–response relationships for the cells of normal tissues. First, there are a limited number of clonogenic assays—techniques in which the end point observed depends directly on the reproductive integrity of individual cells. These systems are directly analogous to cell survival *in vitro*. The techniques developed by Withers and his colleagues are based on their observation of a clone of cells regenerating *in situ* in irradiated tissue. Skin colonies, regenerating crypts in the jejunum, testes stem cells, and kidney tubules are described briefly later in this chapter. It is also possible to obtain dose–response curves for the cells of the epithelial lining of the colon or stomach, but the method used is essentially the same as for the jejunum. Kember described a system for scoring regenerating clones in cartilage at about the same time as Withers’ skin colony system, but it has not been used widely and is not discussed here.

The assay system for the stem cells in the bone marrow or cells of the thyroid and mammary gland depends on the observation of the growth of clones of cells taken from a donor animal and transplanted into a different tissue in a recipient animal. In Till and McCulloch’s bone marrow assay, colonies of bone marrow cells are counted in the spleens of recipient animals. Dose–response curves for mammary and thyroid cells have been obtained by Gould and Clifton by observing colonies growing from cells transplanted into the fat pads of recipient animals.

Second, dose–response relationships that are repeatable and quantitative, but that depend on functional end points, can be obtained. These include skin reactions in rodents or pigs (e.g., erythema and desquamation), pneumonitis or fibrosis in mouse lungs reflected in an increased breathing rate, myelopathy of the hind limbs from damage to the spinal cord, and deformities...
to the feet of mice. The end points observed tend to reflect the minimum number of functional cells remaining in a tissue or organ, rather than the fraction of cells retaining their reproductive integrity.

Finally, one can infer a dose-response curve, for a tissue, which cannot be observed directly by assuming the form of the dose-response curve (linear quadratic) and performing a series of multifraction experiments. This procedure, first suggested by Douglas and Fowler, has been used widely to infer values for $\alpha$ and $\beta$ in the dose–response relationships for normal tissues in which the parameters cannot be measured directly.

This chapter includes assays for both early- and late-responding tissues. The skin, intestinal epithelium, and bone marrow cells, for example, are rapidly dividing self-renewal tissues and respond early to the effects of radiation. The spinal cord, lung, and kidney, by contrast, are late-responding tissues. This reflects the current philosophy that the radiation response of all tissues results from the depletion of the critical parenchymal cells and that the difference in time at which early- and late-responding tissues express radiation damage is a function simply of different cell turnover rates. Many older papers in the literature ascribe the response of late-responding tissues to vascular damage rather than to depletion of parenchymal cells, but this thesis is becoming increasingly difficult to accept.

The various types of normal tissue assay systems are described briefly. The reader who is content with the summary already given may wish to skip the remainder of this chapter.

### CLONOGENIC END POINTS

#### Clones Regrowing In Situ

**Skin Colonies**

Withers developed an ingenious technique, shown in Figure 19.3, to determine the survival curve for mouse skin cells. The hair was plucked from an area on the back of the mouse, and a superficial x-ray machine was used to irradiate an annulus of skin to a massive dose of 30 Gy. This produced a “moat” of dead cells, in the center of which was an isolated island of intact skin that had been protected during the first exposure to low voltage x-rays by a small metal sphere. The intact skin is then given a test dose (D) and observed for nodules of regrowing skin. Figure 19.4 shows nodules regrowing in mouse skin. To obtain a survival curve, it was necessary to repeat this operation with several different areas of skin. A range of ball bearings was used to shield a small area of skin in the middle of the “moat.” The resulting survival data are shown in Figure 19.5 in which the dose (D) to the control area is plotted against the number of surviving cells per square centimeter of skin.
**FIGURE 19.4** Photograph of nodules of mouse skin regrowing from a single surviving cell in the treated area. (Courtesy of Dr. H.R. Withers.)

**FIGURE 19.5** Single-dose and two-dose survival curves for epithelial cells of mouse skin exposed to 29 kVp x-rays. The 37% dose slope \((D_0)\) is 1.35 Gy. The ordinate is not the surviving fraction, as in the survival curves for cells cultured *in vitro*, but is the number of surviving cells per square centimeter of skin. In the two-dose survival curve, the interval between dose fractions was always 24 hours. The curves are parallel, their horizontal separation being equal to about 3.5 Gy; this corresponds to \(D_q\). From a knowledge of \(D_q\) and the slope of the survival curve, \(D_0\), the extrapolation number, \(n\), may be calculated. (Adapted from Withers HR. Recovery and repopulation in vivo by mouse skin epithelial cells during fractionated irradiation. *Radiat Res*. 1967;32:227–239; and Withers HR. The dose–survival relationship for irradiation of epithelial cells of mouse skin. *Br J Radiol*. 1967;40:187–194, with permission.)
There are practical limits to the range in which the dose–response relationship can be determined. At one extreme, it is not possible to irradiate too large an area on the back of the mouse to produce the moat of sterilized skin. At the other extreme, the smallest area that can be used is determined by the fact that even 30 kV radiation scatters laterally to some extent. As can be seen in Figure 19.5, the technique results in a single-dose survival curve that extends from about 8 to 25 Gy. Over this range, with dose plotted on a linear scale and number of surviving cells per square centimeter plotted on a logarithmic scale, the survival curve is straight and has a $D_0$ of 1.35 Gy. This $D_0$ value is very similar to that obtained with mammalian cells cultured in vitro.

The extrapolation number cannot be obtained directly with this technique; the ordinate is the number of surviving cells per square centimeter of skin, and this cannot be converted to the surviving fraction because it is not known with any accuracy how many skin stem cells there are per unit area. It is, however, possible to make an indirect estimate of the extrapolation number by obtaining the survival curve for doses given in two fractions separated by 24 hours. The survival curve obtained in this way is also shown in Figure 19.5. It is parallel to that obtained for single doses, but is displaced from it toward higher doses. As explained in Chapter 3, this lateral displacement in a direction parallel to the dose axis is a measure of $D_q$, the quasithreshold dose. The $D_q$ for mouse skin is about 3.5 Gy, which is very similar to the value for human skin estimated from split-dose experiments.

Crypt Cells of the Mouse Jejunum

A technique perfected by Withers and Elkind makes it possible to obtain the survival characteristics of the crypt cells of the mouse jejunum. The lining of the jejunum is a classic example of a self-renewal system. The cells in the crypts divide rapidly and provide a continuous supply of cells that move up the villi, differentiate, and become the functioning cells. The cells at the top of the folds of the villi are slowly but continuously sloughed off in the normal course of events and are replaced continuously by cells that originate from mitoses in the crypts.

Figure 19.6, an electron micrograph, dramatically shows the three-dimensional structure of the lining of the intestinal epithelium. Mice are given a total body dose of 11 to 16 Gy, which sterilizes a significant proportion of the dividing cells in the crypts, but has essentially no effect on the differentiated cells in the villi. Consequently, crypt degeneration appears early after irradiation, and the villi remain long and their epithelial covering of differentiated cells shows little change. With the further passage of time, the tips of the villi continue to be sloughed away by normal use, but no replacement cells are available from the depopulated crypts, and so the villi begin to shorten and shrink. At sufficiently high doses, the surface lining of the jejunum is completely denuded of villi.

To obtain a survival curve for the jejunal crypt cells, groups of animals are exposed to graded total body doses of radiation. After 3.5 days, each animal is sacrificed and sections are made of the jejunum (Fig. 19.7A). At this time,
FIGURE 19.7  A: Section of mouse jejunum taken 3.5 days after a total body dose in excess of 10 Gy. Note the shortened villi and the regenerating crypts. B: Regenerating crypts shown at a higher magnification. (From Withers HR. Regeneration of intestinal mucosa after irradiation. Cancer. 1971;28:78–81, with permission.)
Surviving cells per circumference

<table>
<thead>
<tr>
<th>Dose (Gy)</th>
<th>100</th>
<th>10</th>
<th>1</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>12</td>
<td>3</td>
<td>5</td>
</tr>
<tr>
<td></td>
<td>10</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Jejunum $^{60}$Co

**FIGURE 19.8** Survival curves for crypt cells in the mouse jejunum exposed to single or multiple doses of $\gamma$-rays (1–20 fractions). The score of radiation damage is the number of surviving cells per circumference (i.e., the number of regenerating crypts per circumference of the sectioned jejunum) counted from sections such as those shown in Figure 19.7. This quantity is plotted on a logarithmic scale against radiation dose on a linear scale. The $D_0$ for the single-dose survival curve is about 1.3 Gy. The shoulder of the survival curve is very large. The separation between the single-dose survival and two-dose survival curves indicates that the $D_q$ is 4 to 4.5 Gy. (Adapted from Withers HR, Mason K, Reid BO, et al. Response of mouse intestine to neutrons and gamma rays in relation to dose fractionation and division cycle. Cancer. 1974;34:39–47, with permission.)

crypts are just beginning to regenerate and it is relatively simple to identify them. Figure 19.7B shows several regenerating crypts at a higher magnification. These pictures also show the shortened villi and the greatly reduced density of cells lining the surface. The score of radiation damage is the number of regenerating crypts per circumference of the sectioned jejunum. This quantity is plotted as a function of dose and yields a survival curve as shown in Figure 19.8. The single-dose survival curve has a $D_0$ (for $\gamma$-rays) of about 1.3 Gy. Also shown in Figure 19.8 are survival curves for radiation delivered in multiple fractions, from 2 to 20. The separation between the single- and two-dose survival curves gives a measure of $D_q$, which has the very large value of between 4 and 4.5 Gy.

This technique has two limitations. First, the quantity plotted on the ordinate is the number of surviving crypts per circumference, not the surviving fraction. Second, experiments can be done only at doses of about 10 Gy or more, at which there is a sufficient level of biologic damage for individual regenerating crypts to be identified. The doses can be delivered, however, in several smaller fractions, as long as the total results in sufficient biologic damage to be scored. The shape of the entire survival curve, then, can be reconstructed from the multifraction data if it is assumed that in a fractionated regimen each dose produces the same amount of cell killing and if an estimate is made of the number of clonogens at risk per crypt. This has been done by Withers and his colleagues; the resultant survival curve is shown in Figure 19.9.

**Testes Stem Cells**

A technique to measure the radiation response of testicular cells capable of sustaining spermatogenesis (i.e., the stem cells) was devised by Withers and his colleagues. About 6 weeks after irradiation, mouse testes are sectioned and examined histologically. Sections of normal and irradiated testes are shown in Figure 19.10. The proportion of tubules containing spermatogenic epithelium is counted and plotted as a function of dose in Figure 19.11. As in many *in vivo* assays, relatively high single doses are used.
**FIGURE 19.9** Effective single-dose survival curve reconstructed from multi-fraction experiments for clonogenic cells of the jejunal crypts of mice. The numbers on the curve refer to the number of fractions used to reconstruct that part of the curve. The initial and final slopes are about 3.57 and 1.43 Gy, respectively. The quasi-threshold dose is 4.3 Gy. The data are equally well fitted by the linear-quadratic formulation. (Adapted from Thames HD, Withers HR, Mason K, et al. Dose survival characteristics of mouse jejunal crypt cells. *Int J Radiat Oncol Biol Phys.* 1981;7:1591–1597, with permission.)

**FIGURE 19.10** A: Histology of normal testis. B: Histology of testis 35 days after a dose of 9 Gy of γ-radiation. Some tubules are completely devoid of spermatogenic epithelium and some are not. (Sertoli’s cells persist in the tubules sterilized of spermatogenic cells.) Foci of spermatogenesis can be derived from single surviving stem cells. (Magnification × 200.) (From Withers HR, Hunter N, Barkley HT Jr, et al. Radiation survival and regeneration characteristics of spermatogenic stem cells of mouse testis. *Radiat Res.* 1974;57:88–103, with permission.)
of 8 to 16 Gy are necessary so that the level of damage is sufficient to be scored. In this dose range, \( D_0 \) is about 1.68 Gy. If the split-dose technique is used, the \( D_q \) is about 2.7 Gy. It is possible to estimate the effect of small doses and reconstruct a complete survival curve by giving large doses in multiple small fractions and assuming that the response to each fraction is the same. The result of this reconstruction is shown in Figure 19.12.

**FIGURE 19.11** Single- and split-dose survival curves for spermatogenic stem cells of the mouse testis. The \( D_0 \) is about 1.68 Gy. The \( D_m \), assessed from the horizontal separation of the single- and split-dose curves, is about 2.7 Gy. (Adapted from Withers HR, Hunter N, Barkley HT Jr, et al. Radiation survival and regeneration characteristics of spermatogenic stem cells of mouse testis. *Radiat Res.* 1974;57:88–103, with permission.)

**FIGURE 19.12** Survival curve for testis stem cells reconstructed from multifraction experiments, assuming that each fraction produces the same biologic effect. The numbers on the curve refer to the number of fractions used to reconstruct that portion of the curve. The \( D_0 \) is about 1.6 Gy and the \( D_q \) is about 3.92 Gy. (Adapted from Thames HD, Withers HR. Test of equal effect per fraction and estimation of initial clonogen number in microcolony assays of survival after fractionated irradiation. *Br J Radiol.* 1980;53:1071–1077, with permission.)
Kidney Tubules

A technique using kidney tubules, again developed by Withers and his colleagues, is the first clonal assay for a late-responding tissue. One kidney per mouse is irradiated with a small field and removed for histologic examination, 60 weeks later. Figure 19.13 shows sections of normal and irradiated kidneys. For ease of scoring, only those tubules touching the renal capsule are scored, and a tubule is considered fully regenerated only if it

**FIGURE 19.13** Photomicrographs of mouse kidney. **A:** Normal, showing proximal tubules in contact with the capsule. (Hematoxylin–eosin stain, magnification × 400.) **B:** Sixty weeks after irradiation with 13 Gy. Note normal proximal tubules and glomeruli amid ghosts of de-epithelialized tubules. One epithelialized tubule is in contact with the capsule. (Hematoxylin–eosin stain, magnification × 200.) (From Withers HR, Mason K, Thames HD. Late radiation response of kidney assayed by tubule cell survival. *Br J Radiol.* 1986;59:587–595, with permission.)
is lined with well-differentiated cuboidal or columnar cells with a large amount of eosinophilic cytoplasm. By 60 weeks, tubules either have no surviving epithelial cells or are lined completely with epithelium that has regenerated from a small number of surviving cells, usually one. The number of tubules regenerating in several arbitrary sections counted is plotted as a function of radiation dose. The result is shown in Figure 19.14; $D_0$ is about 1.53 Gy.

The radiosensitivity of the cells of this late-responding tissue is not very different from that of early-responding tissues such as the skin or intestinal epithelium. The rate of response, however, is quite different. The time required for depletion of the epithelium after a single dose of 14 Gy is about 3 days in the jejunum, 12 to 24 days in the skin, and 30 days in the seminiferous tubules of the testes, but 300 days in the kidney tubules. These results argue strongly that radiation injury in the kidney results from depletion of parenchymal cells and that the slow expression of injury merely reflects the slow turnover of this cell population. Vascular injury is unlikely to be the mechanism underlying the destruction of renal tubules.

**FIGURE 19.14** Dose-survival curve for tubule-regenerating cells. The $D_0$ is 1.53 Gy. (Adapted from Withers HR, Mason K, Thames HD Jr. Late radiation response of kidney assayed by tubule cell survival. *Br J Radiol*. 1986;59:587–595.)

**FIGURE 19.15** Till and McCulloch’s technique. From the donor mouse, a cell suspension is made of nucleated isologous bone marrow. A known number of cells are injected into recipient mice previously irradiated with a 9-Gy total body dose. The spleen is removed from each recipient mouse 9 or 10 days later, and the number of nodules are counted. (Adapted from Till JE, McCulloch EA. In: Cameron IL, Padilla GM, Zimmerman AM, eds. *Developmental Aspects of the Cell Cycle*. New York, NY: Academic Press; 1971:297–313, with permission.)
Cells Transplanted to Another Site

**Bone Marrow Stem Cells**

Till and McCulloch developed a system to determine a survival curve for colony-forming bone marrow cells (Fig. 19.15). Recipient animals first are irradiated supralethally with a dose of 9 to 10 Gy, which sterilizes their spleens. Nucleated isologous bone marrow cells taken from another animal are then injected intravenously into the recipient animals. Some of these cells lodge in the spleen, where they form nodules or colonies 10 to 11 days later, because the spleen cells of the recipient animals have been sterilized previously by the large dose of radiation. At this time, the spleens are removed and the colonies counted. Figure 19.16 is a photograph of a spleen showing the colonies to be counted.

About $10^4$ cells must be injected into a recipient animal to produce one spleen colony because most of the cells in the nucleated isologous bone marrow are fully differentiated cells and would never be capable of forming a colony. To obtain a surviving fraction, bone marrow cells, a donor animal is irradiated to some test dose, and the suspension of cells from the bone marrow is inoculated into groups of recipient animals that previously had been irradiated supralethally. By counting the colonies in the spleens of the recipient animals, and with a knowledge of the number of cells required to produce a colony in an unirradiated animal (plating efficiency), the surviving fraction may be calculated as follows:

\[
\text{Surviving fraction for a dose } D = \frac{\text{colonies counted}}{\text{cells inoculated} \times \text{plating efficiency}}
\]

This procedure is repeated for a range of doses, and a survival curve is obtained (Fig. 19.17). These bone marrow stem cells are very sensitive with a $D_0$ of about 0.95 Gy and little or no shoulder to the survival curve.

![Figure 19.16](image1.png)

**FIGURE 19.16** Photograph of a mouse’s spleen. The mouse was irradiated supralethally to sterilize all the cells of the spleen. The nodules of regrowth originate from intravenously injected bone marrow cells from another animal. (Courtesy of Dr. A. Carsten, Brookhaven National Laboratory.)

![Figure 19.17](image2.png)

**FIGURE 19.17** γ-ray survival curve for the colony-forming ability of mouse bone marrow cells. The cells are irradiated in vivo in the donor animal and grow into colonies in the spleens of supralethally irradiated recipient animals. (Adapted from McCulloch EA, Till JE. The sensitivity of cells from normal mouse bone marrow to gamma radiation in vitro and in vivo. *Radiat Res* 1962;16:822–832, with permission.)
Mammary and Thyroid Cells

Clifton and Gould and their colleagues developed very useful clonogen transplant assays for epithelial cells of the mammary and thyroid glands. They have been used largely for cell survival studies, which are described later, but the initial motivation for their development was to study carcinogenesis in a quantitative system. Most in vitro transformation assays involve fibroblasts, and the bulk of human cancers arise in epithelial cells—hence, the importance and interest in these two systems.

The techniques for these two systems are much the same. To generate a survival curve for mammary or thyroid gland cells in the rat, cells may be irradiated in vivo before the gland is removed from donor animals and treated with enzymes to obtain a monodispersed cell suspension. Known numbers of cells are injected into the inguinal or interscapular white fat pads of recipient animals.

Under appropriate host conditions and grafted cell numbers, the injection of mammary cells gives rise to mammary structures that are morphologically and functionally normal. One such mammary structure may develop from a single cell. By 3.5 weeks after the injection of mammary cells, positive growth is indicated by alveolar units. An example of a milk-filled alveolar unit is shown as an inset in Figure 19.18. If thyroid cells are injected, thyroid follicular units develop (Fig. 19.19).

With either type of cell, a larger number must be injected to produce a growing unit, if the cells

---


are irradiated first to a given dose. In practice, some fancy statistics are involved, a discussion of which is beyond the scope of this chapter; in essence, the ratio of the number of irradiated to unirradiated cells required to produce one growing unit (thyroid follicular unit or alveolar unit) is a measure of the cell-surviving fraction corresponding to the dose. This procedure must be repeated for a range of graded doses to generate a survival curve. The resultant survival curve for mammary cells is shown in Figure 19.18. The characteristics of the curve are unremarkable: \( D_0 \) is about 1.27 Gy, and the extrapolation number is about 5, quite typical of rodent cells cultured \textit{in vitro}. The corresponding survival curve for thyroid cells is shown in Figure 19.19. \( D_0 \) is a little larger than for mammary glands assayed in a similar way, implying that the cells are a little more resistant. Figures 19.18 and 19.19 also show data for cells left \textit{in situ} for 24 hours after irradiation, before being removed and assayed. If this is done, the shoulder of the survival curve is larger because of the repair of potentially lethal damage. This is discussed in more detail in Chapter 5.

An interesting use of these clonogen transplant assays is that the physiologic states of either donor or recipient animals can be manipulated hormonally. For the mammary cell assay, cells may be taken from inactive, slowly dividing glands of virgin rats, from rapidly dividing glands of rats in midpregnancy, or from milk-producing glands of lactating rats. For the thyroid cell assay, the physiologic states of both donor and recipient can be manipulated by control of the diet or by partial thyroidectomy.

**SUMMARY OF DOSE–RESPONSE CURVES FOR CLONOGENIC ASSAYS IN NORMAL TISSUES**

The survival curves for all of the clonogenic assays in normal tissues are plotted together in Figure 19.20. There is a substantial range of radiosensitivities, with shoulder width being the principal variable. \textit{In vitro} curves for cells from patients with ataxia telangiectasia (AT) also are shown because these are probably the most radiosensitive mammalian cells.

![Figure 19.20](image-url)
DOSE–RESPONSE RELATIONSHIPS FOR FUNCTIONAL END POINTS

Pig Skin

Pig skin has been used widely in radiobiologic studies because it has many features in common with human skin such as color, hair follicles, sweat glands, and a layer of subcutaneous fat. In view of these structural similarities, it is not surprising that the response of pig skin to radiation closely resembles that of human skin, both qualitatively and quantitatively.

Fowler and his colleagues pioneered the use of pig skin as a radiobiologic test system. Several small rectangular fields on the pig’s flank were irradiated with graded doses of x-rays, and the reactions were scored daily using the arbitrary scale shown in Table 19.1. After a single dose of radiation, the reaction becomes apparent after about 15 days and develops as shown in Figure 19.21.

**Figure 19.21** Development of skin reactions in the pig after graded doses of x-rays, delivered as a single exposure (A) or as multiple fractions spaced over time (B). (Adapted from Fowler JR, Morgan RL, Silvester JA, et al. Experiments with fractionated x-ray treatment of the skin of pigs: 1. Fractionation up to 28 days. *Br J Radiol*. 1963;36:188–196, with permission.)

### Table 19.1 Radiation Reactions in Pig Skin

<table>
<thead>
<tr>
<th>Arbitrary Score</th>
<th>Reaction</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>No visible reaction</td>
</tr>
<tr>
<td>1</td>
<td>Faint erythema</td>
</tr>
<tr>
<td>2</td>
<td>Erythema</td>
</tr>
<tr>
<td>3</td>
<td>Marked erythema</td>
</tr>
<tr>
<td>4</td>
<td>Moist desquamation of less than half the irradiated area</td>
</tr>
<tr>
<td>5</td>
<td>Moist desquamation of more than half the irradiated area</td>
</tr>
</tbody>
</table>

Two phases of the reaction can be distinguished. First, an early wave of erythema occurred (at 10–40 days), which was variable from one animal to another. This represents the uncomfortable “acute” reaction sometimes seen in patients on radiotherapy at about the end of a course of treatment. Second, a more gradual increase to a second broad wave of moderately severe reactions took place (at 50–100 days), representing a more permanent kind of damage. This second wave shows the tolerance of skin to a more serious type of long-term damage and is a more repeatable and consistent index of radiation damage. It was subsequently found to correlate well with longer term damage (up to 2 years) and with subcutaneous damage.

The “score” of radiation damage is taken to be the average skin reaction occurring between certain time limits that encompass the medium-term reactions. After a single dose, this might be a 35-day period between 50 and 85 days after irradiation. For a protracted fractionated regimen, this period of reaction may come later, between days 65 and 100. The average skin reaction in the chosen time period then is plotted as a function of dose; examples of dose–response curves obtained this way are shown in Figure 19.22 for single and fractionated doses.

Late effects also have been studied in pig skin by measuring the contraction that results from fibrosis a year or more after irradiation. A square is tattooed on the skin of the animal in the irradiated field, and the dimensions of this square are recorded as a function of dose as the contraction occurs. This is a primitive but effective measure of late effects.

Many of the important early studies on the fractionation effects of x-rays and the comparison of x-rays with fast neutrons were performed with this biologic system. One overwhelming advantage is that data obtained this way can be extrapolated to the human with a high degree of confidence. The disadvantage is that the animals are large and awkward to work with, and their maintenance involves a considerable expense.

Rodent Skin

Because of the inconvenience and expense of using pigs, the skin of the mouse leg and foot is commonly used instead. One hind leg of each animal is irradiated; the other serves as a control. The skin response is observed each day after irradiation and is scored according to the arbitrary scale shown in Table 19.2. Various doses are used. The progressive development of the reaction after ten doses of 6 Gy each is illustrated in Figure 19.23; each point represents the mean of several animals. Reactions appear by about the 10th day, peak by 20 to 25 days, and then subside. The second wave of the reaction, noted for pig skin, is not seen in mice but is observed in rats. A dose-response curve is obtained by averaging the skin reaction over a period of time and plotting this average as a function of dose.

Early and Late Response of the Lung Based on Breathing Rate

Travis and her colleagues developed a noninvasive assay of breathing frequency to assess both early and late damage in mouse lungs. Breathing frequency increases progressively with dose after...
These data also show the dramatic sparing that results from fractionation; this is discussed further in another section of this chapter.

The various syndromes of radiation-induced injury in rodent brain and spinal cord are very similar to those described in humans. Lesions observed within approximately the first 6 months after irradiation are limited primarily to the white matter and range between early diffuse or focal demyelination and extensive necrosis. Different pathogenic pathways toward the development of white matter necrosis have been proposed, with the glial and vascular tissue components as the major targets. The most common type of late delayed injury peaks at 1 to 2 years postirradiation and almost certainly has a vascular basis. Another type of late injury that has been described more recently in various species, including humans, is slowly progressive glial atrophy. This lesion is not associated with

### TABLE 19.2 Radiation Reactions in Mouse Leg Skin

<table>
<thead>
<tr>
<th>Arbitrary Score&lt;sup&gt;a&lt;/sup&gt;</th>
<th>Observations</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.5</td>
<td>50/50; doubtful if any difference from normal or not</td>
</tr>
<tr>
<td>1−</td>
<td>Because 1 covers a wide range of reddening, even before reaching the severity or additional factors requiring 1+, it is necessary to have 1− for “definite reddening (i.e., definitely not normal), but only a very slight degree”</td>
</tr>
<tr>
<td>1</td>
<td>Definite abnormality; definite reddening, top or bottom of leg; “clean” appearance means not greater than 1</td>
</tr>
<tr>
<td>1+</td>
<td>Severe reddening or reddening with definite white marks in creases under foot; query breakdown; query puffiness</td>
</tr>
<tr>
<td>1.5</td>
<td>Some breakdown of skin (usually seen on bottom of foot first); scaly or crusty appearance; definite puffiness, plus (query) breakdown; very marked white marks in creases plus puffiness or severe redness</td>
</tr>
<tr>
<td>1.5+</td>
<td>Query possibly moist desquamation in small areas</td>
</tr>
<tr>
<td>2</td>
<td>Breakdown of large areas of skin or toes stuck together; possibly moist in places but not all moist</td>
</tr>
<tr>
<td>2.5</td>
<td>Breakdown of large areas of skin with definite moist exudates</td>
</tr>
<tr>
<td>3</td>
<td>Breakdown of most of the skin with moist exudates</td>
</tr>
<tr>
<td>3.5</td>
<td>Complete necrosis of limb (rarely seen so far)</td>
</tr>
</tbody>
</table>

<sup>a</sup> and − are equivalent to 0.25.


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a threshold of about 11 Gy (Fig. 19.24). The increased breathing frequency in rodent lungs at 16 and 36 weeks is associated with the early response (i.e., pneumonitis); by 52 weeks, the elevated breathing frequency is associated with the late response (i.e., fibrosis). This is a simple but highly quantitative and reproducible system.

### Spinal Cord Myelopathy

A dose–response relationship can be determined for late damage caused by local irradiation of the spinal cords of rats. Several investigators have worked with this system, notably van der Kogel. After latent periods of 4 to 12 months, symptoms of myelopathy develop, the first signs of which are palpable muscle atrophy, followed some time later by impaired use of the hind legs. Figure 19.25 shows the steep dose–response relationship for hind limb paralysis following the irradiation of a section of the spinal cord in rats.
FIGURE 19.23  Daily skin reaction scores for mice receiving 60 Gy in 10 equal fractions to the right hind leg. Each point represents the mean score of six animals; the vertical lines represent the standard errors of the mean. (Adapted from Brown JM, Goffinet DR, Cleaver JE, et al. Preferential radiosensitization of mouse sarcoma relative to normal skin by chronic intraarterial infusion of halogenated pyrimidine analogs. JNCI. 1971;47:75–89, with permission.)

FIGURE 19.24  Breathing frequency in mice as a function of dose measured (left to right) 16, 36, and 52 weeks after irradiation with x-rays. Breathing frequency is expressed as a percentage increase above the age-related control value. (Adapted from Travis EL, Down JD, Holmes SJ, et al. Radiation pneumonitis and fibrosis in mouse lung assayed by respiratory frequency and histology. Radiat Res. 1980;84:133–142, with permission.)
necrosis but occurs diffusely and at lower doses. With improvements in diagnostic procedures such as magnetic resonance imaging, glial atrophy may become a more frequently recognized adverse effect of brain tumor therapy.

**Latency**

Over a dose range of about 25 to 60 Gy, delivered in single doses, the general tendency is a decreasing latency with increases in dose of approximately 2 days per Gy. There is a considerable variation with animal strain, as well as with the region of the cord irradiated. In terms of mechanisms, demyelination or slowly progressive atrophy is probably a consequence of interference with the slow continuous turnover of oligodendrocytes by killing of glial progenitor cells. Vascular injury may accelerate, precipitate, or even initiate the white matter changes leading to necrosis. This is an area of some controversy.

**Fractionation and Protraction**

The effect of dose fractionation and protraction on tolerance to radiation has been investigated extensively in the rat spinal cord and to a lesser extent in the mouse, monkey, and guinea pig. Because these systems turn over slowly, there is little influence of overall treatment time up to any conventional clinical regimen of 6 to 8 weeks. On the other hand, dose per fraction is very important (Fig. 19.25), with the dose to produce paralysis increasing dramatically with number of fractions. The effect of a large number of very small fractions has also been investigated. Figure 19.26 shows the relation between total dose and dose per fraction to produce paralysis in 50% of rats from irradiation of a short length of cervical spine. The smooth curve is

![Graph showing dose-response curves for the induction of hind leg paralysis in rats following irradiation of a section of the spinal cord (L2–L5). Note how the dose necessary to produce paralysis increases rapidly with increasing numbers of fractions. (Adapted from van der Kogel AJ. Late Effects of Radiation on the Spinal Cord. Rijswijk, the Netherlands, the Radiobiological Institute of the Organization for Health Research TNO; 1979:1–160, with permission.)](image1)

![Graph showing data points for the relation between total dose, as a function of dose per fraction, to produce paralysis in 50% of rats after irradiation of the spinal cord. The curve is an isoeffect relationship based on the linear-quadratic equation with an α/β value of 1.5 Gy. The experimental data suggest that the linear-quadratic model overestimates tolerance for dose-per-fraction values less than 2 Gy. This may be a result of incomplete repair because the interfraction interval was only 4 hours. (Adapted from van der Kogel AJ. Central nervous system radiation injury in small animal models. In: Gutin PH, Leibel SA, Sheline GE, eds. Radiation Injury to the Nervous System. New York, NY: Raven Press; 1991:91–112, with permission.)](image2)
matter necrosis shows a marked dependence on the length of cord irradiated. Late vascular injury shows less dependence on cord length. Beyond a few centimeters, the tolerance is virtually independent of the length of cord irradiated. This would be predicted from the linear arrangement of the functional subunits. A chain is broken whether one, two, three, or more links are removed.

Retreatment after Long Time Intervals

The spinal cord does recover to some extent after long time periods following irradiation. The extent of the recovery depends, of course, on the first treatment—that is, what fraction of tolerance was involved. Experiments with rats indicate that after an initial treatment to 50% tolerance, the retreatment tolerance approaches 90% of the tolerance of the untreated control group by about a year after the initial irradiation. If the initial treatment represented a larger fraction of tolerance, the retreatment that can be tolerated is reduced.

### INFERRING THE RATIO $\alpha/\beta$ FROM MULTIFRACTION EXPERIMENTS IN NONCLONOGENIC SYSTEMS

The parameters of the dose-response curve for any normal tissue system for which a functional end point can be observed may be inferred by performing a multifraction experiment. Take, for example, an experiment in which mouse foot skin reaction is scored. Doses that result in the same skin reaction (e.g., moist desquamation of more than 50% of the area irradiated) if delivered as a single exposure in a multifraction regimen (e.g., 5, 10, or 20 fractions) must be determined experimentally. Several assumptions must be made as follows:

1. The dose–response relationship is represented adequately by the LQ formulation:
   $$ S = e^{-\alpha D - \beta D^2} $$
   in which $S$ is the fraction of cells surviving a dose, $D$, and $\alpha$ and $\beta$ are constants.
2. Each dose in a fractionated regimen produces the same biologic effect.
3. Full repair of sublethal damage takes place between dose fractions, but no cell proliferation occurs.
SUMMARY OF PERTINENT CONCLUSIONS

- The relationship between dose and incidence is sigmoid for both tumor control and normal tissue damage.
- The ratio of tumor response to normal tissue damage is called the therapeutic ratio or therapeutic index.
- The therapeutic index can be manipulated by dose fractionation or by the use of drugs that preferentially increase tumor response.
- After irradiation, most cells die a mitotic death; that is, they die in attempting the next or a later mitosis. In some tissues, cells die by apoptosis, which is a programmed cell death.
- Systems involving clonogenic end points (i.e., cell survival) for cells of normal tissues include some in which cells regrow in situ and some in which cells are transplanted to another site.
- In situ regrowth techniques include skin colonies, crypts in the jejunum, testes stem cells, and kidney tubules. Single dose experiments can yield the slope ($D_0$) of the dose-response curve over a range of high doses. Multifraction experiments allow the whole dose-response curve to be reconstructed.
- Systems in which cell survival is assessed by transplantation into another site include bone marrow stem cells, thyroid cells, and mammary cells.
- A dose-response curve for bone marrow stem cells can be obtained by allowing cells...
from the donor animal to lodge and grow in the spleens of recipient animals. These are very sensitive cells with a $D_0$ close to 1 Gy, with little or no shoulder.

Dose–response curves for mammary and thyroid cells can be obtained by transplanting them into fat pads of recipient animals.

The radiosensitivity of cells from normal tissues varies widely. The width of the shoulder of the curve is the principal variable. Jejunal crypt cells have a very large shoulder; bone marrow stem cells have little, if any, shoulder. Most other cell types studied in clonogenic assays fall in between.

Dose–response curves for functional end points, distinct from cell survival, can be obtained as follows:
1. Pig skin and rodent skin—by measuring skin reactions.
2. Early and late response of the lung—by measuring breathing rate.
3. Spinal cord—by observing myelopathy:
   a. Paralysis develops after a latency of months to years.
   b. Early lesions are limited to white matter; late delayed injury may have a vascular basis.
   c. Spinal cord damage is very sensitive to fractionation—$\alpha/\beta$ of about 1.5 Gy.
   d. Sublethal damage repair probably has “fast” and “slow” components.
   e. If multiple fractions per day are used, the interfraction interval should be at least 6 to 8 hours.
   f. Functional subunits are arranged serially like links in a chain.
   g. For short lengths of cord, tolerance dose varies markedly with cord length irradiated; for cord lengths greater than a few centimeters, tolerance dose is virtually independent of cord length.

The shape of the dose–response relationship for functional end points, obtained from multifraction experiments, is more pertinent to radiotherapy than clonogenic assays.

The ratio $\alpha/\beta$ (the dose at which the linear and quadratic components of radiation damage are equal) may be inferred from multifraction experiments in systems scoring nonclonogenic end points.

### BIBLIOGRAPHY


Most of the effects of radiation therapy on normal tissues can be attributed to cell killing, but there are some that cannot. Examples include the following:

- Nausea or vomiting that may occur a few hours after irradiation of the abdomen.
- Fatigue felt by patients receiving irradiation to a large volume, especially within the abdomen.
- Somnolence that may develop several hours after cranial irradiation.
- Acute edema or erythema that results from radiation-induced acute inflammation and associated vascular leakage.

It is thought that these effects are mediated by radiation-induced inflammatory cytokines. These effects aside, most effects of radiation on normal tissues result from the depletion of a cell population by cell killing.

The cells of normal tissues are not independent but form a complete integrated structure. There is a delicate balance between cell birth and cell death to maintain tissue organization and the number of cells. The response to damage is governed by (1) the inherent cellular radiosensitivity, (2) the kinetics of the tissue, and (3) the way cells are organized in that tissue. If the fate of individual cells is studied, as described in Chapter 3, there is a continuous monotonic relationship between the magnitude of the dose and the fraction of cells that are “killed” in the sense that they lose their reproductive integrity—the ability to divide indefinitely. By contrast, no effects are seen in tissues after small doses, although effects of increasing severity become apparent if the dose rises above a threshold level. The reason is, of course, that killing a small number of cells in a tissue matters very little; visible damage is evident only if a large enough proportion of the cells are killed and removed from the tissue. The threshold dose below which no effect is seen and the delay between irradiation and the time at which the damage becomes observable vary greatly among different tissues.

Cell death after irradiation occurs mostly as cells attempt to divide. In tissues with a rapid turnover rate, damage becomes evident quickly—in a matter of hours in the intestinal epithelium and bone marrow, and in a matter of days in the skin and mucosa. In tissues in which cells divide rarely, radiation damage to cells may remain latent for a long period and be expressed very slowly. Radiation damage to cells that are already on the path to...
Radiation effects are commonly divided into two categories, early and late, which show quite different patterns of response to fractionation; their dose-response relations are characterized by different $\alpha/\beta$ ratios, as described in more detail in Chapter 23. Late effects are much more sensitive to changes in fractionation than early effects. Early, or acute, effects result from the death of a large number of cells and occur within a few days or weeks of irradiation in tissues with a rapid rate of turnover. Examples include effects in the epidermal layer of the skin, gastrointestinal epithelium, and hematopoietic system, in which the response is determined by a hierarchical cell lineage composed of stem cells and their differentiating offspring. The time of onset of early reactions correlates with the relatively short life span of the mature functional cells; the identity of the target cells is usually obvious.

Late effects appear after a delay of months or years and occur predominantly in slowly proliferating tissues, such as tissues of the lung, kidney, heart, liver, and central nervous system. The difference between the two types of lesions lies in their progression: Acute damage is repaired rapidly because of the rapid proliferation of stem cells and may be completely reversible. By contrast, late damage may improve but is never completely repaired. A late effect may result from a combination of vascular damage and loss of parenchymal cells. Clearly, vascular damage is not the dominant factor in every case, because if it were, the dose-effect relationship would be the same for all tissues, and that is not the case. It may be true for some tissues, however, including the spinal cord. If intensive fractionation protocols deplete the stem cell population below levels needed for tissue restoration, an early reaction in a rapidly proliferating tissue may persist as a chronic injury. This has been termed a consequential late effect, that is, a late effect consequent to, or evolving out of, a persistent severe early effect. The earlier damage is most often attributable to an overlying acutely responding epithelial surface—for example, fibrosis or necrosis of skin consequent to desquamation and acute ulceration.

### Functional Subunits in Normal Tissues

The fraction of cells surviving determines the success or failure of a treatment regimen as far as the tumor is concerned, because a single surviving cell may be the focus for the regrowth of the tumor. For normal tissues, however, this is not the whole story. The tolerance of normal tissues for radiation depends on the ability of the clonogenic cells to maintain a sufficient number of mature cells suitably structured to maintain organ function. The relationship between the survival of clonogenic cells and organ function or failure depends also on the structural organization of the tissue. Many tissues may be thought of as consisting of functional subunits (FSUs).

In some tissues, the FSUs are discrete, anatomically delineated structures whose relationship to tissue function is clear. Obvious examples are the nephron in the kidney, the lobule in the liver, and perhaps the acinus in the lung. In other tissues, the FSUs have no clear anatomic demarcation. Examples include the skin, the mucosa, and the spinal cord. The response to radiation of these two types of tissue—with structurally defined or structurally undefined FSUs—is quite different.

The survival of structurally defined FSUs depends on the survival of one or more clonogenic cells within them, and tissue survival in turn depends on the number and radiosensitivity of these clonogens. Although such tissues are composed of a large number of FSUs, each is a small self-contained entity independent of its neighbors. Surviving clonogens cannot migrate from one to the other. Because each FSU is both small and autonomous, low doses deplete the clonogens.
in it. Each kidney, for example, is composed of a large number of relatively small FSUs, each of which is a self-contained structural entity independent of its neighbors. Consequently, survival of a nephron after irradiation depends on the survival of at least one clonogen within it and therefore on the initial number of renal tubule cells per nephron and their radiosensitivity. Because this FSU is relatively small, it is completely depleted of clonogens by low doses, which accounts for the low tolerance to radiation of the kidney. Other organs that resemble the kidney in having structurally defined FSUs not repopulated from adjacent FSUs may be those with a branching treelike system of ducts and vasculature that ultimately terminate in “end structures” or lobules of parenchymal cells. These can be visualized as independent structurally defined FSUs. Examples of organs with this tissue architecture include the lung, liver, and exocrine organs. At least some of these also have low tolerance to radiation.

By contrast, the clonogenic cells that can repopulate the structurally undefined FSUs after depletion by radiation are not confined to one particular FSU. Rather, clonogenic cells can migrate from one FSU to another and allow repopulation of a depleted FSU. For example, reepithelialization of a denuded area of skin can occur either from surviving clonogens within the denuded area or by migration from adjacent areas.

A concept proposed to link the survival of clonogenic cells and functional survival is the tissue rescue unit, defined as the minimum number of FSUs required to maintain tissue function. This model assumes that the number of tissue rescue units in a tissue is proportional to the number of clonogenic cells, that FSUs contain a constant number of clonogens, and that FSUs can be repopulated from a single surviving clonogen.

Some tissues defy classification by this system. The crypts of the jejunum, for example, are structurally well-defined subunits, but surviving crypt cells can and do migrate from one crypt to another to repopulate depleted neighbors.

**THE VOLUME EFFECT IN RADIOTHERAPY: TISSUE ARCHITECTURE**

It is generally observed in clinical radiotherapy that the total dose that can be tolerated depends on the volume of tissue irradiated. Tolerance dose has been defined as the dose that produces an acceptable probability of a treatment complication. This definition includes objective criteria, such as the radiobiology involved, and subjective factors that may be socioeconomic, medicolegal, or psychological.

The spatial arrangement of the FSUs in the tissue is critical. In the case of tissues in which the FSUs are arranged in a series, like the links of a chain, the integrity of each is critical to organ function, and elimination of any one FSU results in a measurable probability of a complication. The spinal cord is the clearest example in which specific functions are controlled by specific segments arranged linearly, or serially. Because impulses must pass along the cord, death of critical cells in any one segment results in complete failure of the organ. Radiation damage to such tissues is expected to show a binary response, with a threshold dose below which there is normal function and above which there is loss of function (e.g., radiation-induced myelopathy). This is illustrated in Figure 20.1. As the field size increases to include a greater number of FSUs—1, 4, or 16 in this example—the curve relating probability of a complication to dose rises much more steeply with dose and moves to lower doses. This explains the important volume effect found in, for example, the spinal cord in which FSUs are arranged in series and loss of any one may result in myelopathy.

Clinical tolerance also depends strongly on the volume irradiated in the kidney and lung; both of these organs are very sensitive to irradiation of their entire volume, but small volumes can be treated to much higher doses. This is because there is considerable functional reserve capacity, with only about 30% of the organ required to maintain adequate function under normal physiologic conditions. The large reserve capacity and increased tolerance to partial-volume irradiation are caused by the parallel organization of functional nephrons and alveolar subunits. Inactivation of a small number of FSUs does not lead to loss of organ function. Functional damage will not occur until a critical number of FSUs are inactivated by irradiation. This implies that there should also be a threshold volume of irradiation below which no functional damage will develop, even after high-dose irradiation. Above this threshold, damage is usually exhibited as a graded response—increasing severity
ulceration in a smaller area. In other words, although the severity of a skin reaction is relatively independent of the area irradiated because healing occurs by regeneration of surviving clonogens scattered throughout the treated area, the tolerability is not. Therefore, there is a volume effect in clinical practice, but it is not based on an increased probability of injury as it is in tissues in which FSUs are arranged serially.

### RADIATION PATHOLOGY OF TISSUES

As previously stated, the response of a tissue or organ to radiation depends primarily on three factors: (1) the inherent sensitivity of the individual cells, (2) the kinetics of the tissue as a whole of which the cells are a part, and (3) the way the cells are organized in that tissue. These factors combine to account for the substantial variation in response to radiation characteristic of different tissues.

In the case of tissues composed of highly differentiated cells performing specialized functions, cell survival curves (Chapter 3) are largely irrelevant, because these cells have no mitotic future. Little information is available at the cellular level concerning the effects of radiation on differentiated cells. All that can be said is that, in general, the amount of radiation needed to destroy the functioning ability of a differentiated cell is far greater than that necessary to stop the mitotic activity of a dividing cell.

A closed static population, composed entirely of mature differentiated cells, is therefore very resistant to radiation. In the case of self-renewing tissues, the Achilles heel is the dividing cell: Loss of reproductive ability in an appreciable fraction of these cells occurs after a moderate dose of a few grays. Whether the tissue or organ as a whole appears to be affected to a small or large extent—and is consequently labeled as sensitive or resistant—depends on the extent to which the tissue involved can continue to function adequately with a reduced number of cells.

Another factor that is evident from even this most elementary consideration of population kinetics is that the time interval between the delivery of the radiation insult and its expression in tissue damage is very variable for different populations. This time interval is determined by the normal life span of the mature functional cells and the time it takes for a cell “born” in the stem cell compartment to mature to a functional state. For
example, mature erythrocytes in circulating blood have a relatively long life span and are separated from the primitive stem cell compartment by several transit compartments, so that time is required for a cell to pass through the various stages of differentiation and maturation. Consequently, a considerable time interval elapses between the depopulation of the stem cell compartment and the final expression of this injury in terms of a reduced peripheral blood cell count. By contrast, in the case of the intestinal epithelium, the mature functional cells on the surface of the villi have a short life span, and the time interval between the “birth” of a new cell in the stem compartment of the crypt and its appearance as a mature functional cell is very short, on the order of a few days. As would be expected, therefore, radiation damage is expressed correspondingly quickly in this tissue. Two systems are typically used to classify tissue radiosensitivity in terms of population kinetics and tissue architecture: Casarett’s classification and Michalowski’s classification.

**CASARETT’S CLASSIFICATION OF TISSUE RADIOSENSITIVITY**

The limitation of our knowledge of cellular population kinetics is remedied to some extent by a wealth of information on the relative sensitivities of various tissues based on histopathologic observations. It must be emphasized that these data are based on entirely different end points than those with which previous chapters have been concerned. To score a cell as “dead” by observing a fixed and stained section of tissue through a microscope is quite different from the experimental test of cell death in terms of loss of reproductive capacity, which has been used previously. Nevertheless, the study of radiation pathology provides data that are highly relevant to clinical radiotherapy.

Casarett has suggested a classification of mammalian cell radiosensitivity based on histologic observation of early cell death. He divided parenchymal cells into four major categories, numbered I through IV (Table 20.1). The supporting structures, such as the connective tissue and the endothelial cells of small blood vessels, are regarded as intermediate in sensitivity between groups II and III of the parenchymal cells.

One of the most sensitive cells to radiation, in fact, defies all the “laws” and systems of classification; it is the small lymphocyte. This cell, it is believed, never divides at all, or at least only in exceptional circumstances. Small lymphocytes disappear from circulating blood after very small doses of radiation, and it is believed that they suffer an interphase death (by the process of apoptosis). Most of various tissues based on histopathologic observations. It must be emphasized that these data are based on entirely different end points than those with which previous chapters have been concerned. To score a cell as “dead” by observing a fixed and stained section of tissue through a microscope is quite different from the experimental test of cell death in terms of loss of reproductive capacity, which has been used previously. Nevertheless, the study of radiation pathology provides data that are highly relevant to clinical radiotherapy.

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Sensitive cells die a mitotic death after irradiation; most cells that never divide require very large doses to kill them. The small lymphocyte breaks both of these rules, inasmuch as it does not usually divide, dies of interphase death, and yet is one of the most sensitive mammalian cells.

Group I of Casarett’s classification, the most sensitive group, consists of vegetative intermitotic cells and includes the stem cells of the classic self-renewing systems, such as the basal layers of the epidermis and the intestine, the erythroblasts (precursors of red blood cells), intestinal crypt cells, and the primitive cells of the spermatogenic series. The stem cells divide regularly and provide a steady and abundant supply of progeny, some of which differentiate and mature into functioning cells. A reservoir of primitive dividing stem cells is maintained and, in some cases, can be triggered to divide more rapidly in response to a need. The primitive dividing stem cells are vulnerable to radiation; a moderate dose causes a proportion of them to “die” in attempting the next or a subsequent mitosis. The time of crisis for the organism as a whole occurs if the supply of functioning cells is inadequate: a shortage of circulating red and white blood cells in the case of the blood, and a shortage of mature covering dermal cells in the case of the skin. The time interval between irradiation and the crisis is about equal to the life span of the mature functioning cells. As the functioning cells die off at the end of their natural life span, there are none to take their place if a dose of radiation previously has depopulated the stem cell compartment. Depending on the size of the dose, the organ or tissue may not survive the critical time at which the number of functioning cells reaches a minimum value.

Group II consists of cells that divide regularly but that also mature and differentiate between divisions. Cells in this category are relatively short-lived as individuals and are produced by division of vegetative intermitotic cells. These cells usually complete a limited number of divisions and differentiate to some extent between successive mitoses. This group includes cells of the hematopoietic series in the intermediate stages of differentiation and likewise the more differentiated spermatogonia and spermatocytes.

Group III, the reverting postmitotic cells, is relatively resistant and as individuals have relatively long lives. Ordinarily, the cells in this category do not undergo mitosis, but they are capable of dividing with the appropriate stimulus, which is usually damage or loss of many of their own kind. The liver cells are a good example of this category. In the adult, there is normally little or no cell division, but if a large part of the liver is removed by surgery, the remaining cells are triggered to divide and make good the loss. Other examples in this category include the cells of the kidney and pancreas and of various glands, such as the adrenal, thyroid, and pituitary.

Group IV consists of the fixed postmitotic cells. The cells in this group are generally considered the most resistant to radiation. They are highly differentiated and appear to have lost the ability to divide. Some have a long life span, such as the neurons. Others have a short life span, such as the granulocytes, which have to be continually replaced by the division of more primitive cells. The superficial epithelial cells of the gut also fall into this category. In the normal course of events, they are sloughed off the tops of the villi and replaced by cells dividing in the crypts.

**MICHALOWSKI’S H- AND F-TYPE POPULATIONS**

Michalowski classified tissues as following either a “hierarchical” model or a “flexible” model. Within tissues, three distinct categories of cells can be identified. First are the stem cells, which are capable of unlimited proliferation and escape senescence because of telomere shortening by the enzyme telomerase. Examples include the crypt cells in the intestinal mucosa. The cells produced by stem cell proliferation both maintain the stem cell pool and provide candidates for differentiation. Second, at the other extreme, are functional cells, which are fully differentiated; they are usually incapable of further division and die after a finite life span, though the life span varies enormously among different cell types. Examples include circulatory granulocytes and the cells that make up the villi of the intestinal mucosa. Between these two extremes are maturing partially differentiated cells; these are descendants of the stem cells, still multiplying as they complete the process of differentiation. In the bone marrow, for example, the erythroblasts and granuloblasts represent intermediate compartments. Many tissues represent this hierarchical model (H-type populations), including the hematopoietic bone marrow, intestinal epithelium, and epidermis.
Other tissues, such as liver, thyroid, and dermis, are composed of cells that rarely divide under normal conditions but can be triggered to divide by damage to the tissue or organ. These flexible tissues (F-type populations) have no compartments and no strict hierarchy. After damage to the tissue, all cells, including those that are functional, enter the cell cycle. The time interval before damage becomes evident is a function of dose. If the dose is small, the expression of damage is delayed because cells divide infrequently. Consequently, the damage may be hidden for a long time.

Many tissues are hybrids of these two extreme models, with most cells able to make a few divisions and a minority of the population behaving as stem cells.

### GROWTH FACTORS

Radiation induces interleukin-1 as well as interleukin-6. Interleukin-1 acts as a radioprotectant of hematopoietic cells by increasing both the shoulder and the $D_0$ of the survival curve. Basic fibroblast growth factor induces endothelial cell growth, inhibits radiation-induced apoptosis, and therefore protects against microvascular damage. This growth factor is produced in response to stress (heat, hypoxia, chemicals, radiation) and tends to reduce late effects. Microvascular protection is more effective in branching midsize capillaries (in which higher concentrations of basic fibroblast growth factor are seen) than in nonbranching capillaries. To the extent that radiation-induced late effects are mediated by damage to blood vessels, radiation tolerance is high in organs with large blood vessels (corresponding to high levels of growth factors) and lower near nonbranching capillaries.

Platelet-derived growth factor $\beta$ increases damage to vascular tissue. Transforming growth factor $\beta$ (TGF-$\beta$) induces a strong inflammatory response—for example, in pneumonitis. It stimulates the growth of connective tissue and tends to inhibit epithelial cell growth. Consequently, fibrosis and vascular changes associated with late radiation effects are linked with this factor. TGF-$\beta$ may down-regulate interleukin-1 and tumor necrosis factor (TNF) and increase damage to hematopoietic tissue.

TNF is a cytotoxic agent that mediates the inflammatory response produced by monocytes and tumor cells by binding to cell-surface receptors that initiate signal transduction pathways. TNF induces proliferation of fibroblasts, inflammatory cells, and endothelial cells and so is associated with complications. In clinical trials, the administration of TNF causes fatigue, anorexia, weight loss, and transient leukopenia. TNF protects hematopoietic cells and sensitizes tumor cells to radiation. Serum concentrations of TNF correlate with severity of pneumonitis, hepatic dysfunction, renal insufficiency, and demyelination. TNF may contribute to the pathophysiology of radiation-induced central nervous system symptoms. The expression of TNF following radiation is believed to be regulated at the transcriptional level and involves the protein kinase C-dependent pathway.

### SPECIFIC TISSUES AND ORGANS

Table 20.2 is a compilation of tissue and organ sensitivities. Important examples will be discussed in turn.

**Skin**

The skin is composed of the outer layer, the epidermis, which is the site of early radiation reactions, and the deeper layer, the dermis, which is the site of late radiation reactions (Fig. 20.2).

The epidermis (30–300 $\mu$m thick) is derived from a basal layer of actively proliferating cells, which is covered by several layers of nondividing differentiating cells to the surface, at which the most superficial keratinized cells are desquamated. It takes about 14 days from the time a newly formed cell leaves the basal layer to the time it is desquamated from the surface. The target cells for radiation damage are the dividing stem cells in the basal layer.

The dermis is a dense connective tissue (1–3 mm thick) within which scattered fibroblasts produce most of the dermal proteins. The vasculature of the dermis plays a major role in the radiation response. The target cells are thus the fibroblasts and the vascular endothelial cells.

A few hours after doses greater than 5 Gy, there is an early erythema similar to sunburn, which is caused by vasodilation, edema, and loss of plasma constituents from capillaries. Reactions resulting from stem cell death take longer to develop. When orthovoltage ($250$-kV) x-rays were the modality commonly used, skin was frequently dose limiting, because the full dose is deposited...
# Table 20.2: A Compilation of Tissue and Organ Sensitivities

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<tr>
<th>Injury</th>
<th>TD&lt;sub&gt;5/5&lt;/sub&gt;, Gy</th>
<th>TD&lt;sub&gt;50/5&lt;/sub&gt;, Gy</th>
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<td>Bone marrow</td>
<td>Aplasia, pancytopenia</td>
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<td>Intestine</td>
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<tr>
<td>Stomach</td>
<td>Perforation, ulcer, hemorrhage</td>
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<td>Brain</td>
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<td>75</td>
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<tr>
<td>Spinal cord</td>
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<td>Pericarditis and pancarditis</td>
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<th>TD&lt;sub&gt;50/5&lt;/sub&gt;, Gy</th>
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</tr>
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<td>Esophagus</td>
<td>Esophagitis, ulceration</td>
<td>55</td>
<td>50</td>
</tr>
<tr>
<td></td>
<td></td>
<td>60</td>
<td>70</td>
</tr>
<tr>
<td>Rectum</td>
<td>Ulcer, stenosis, fistula</td>
<td>60</td>
<td>80</td>
</tr>
<tr>
<td>Salivary glands</td>
<td>Xerostomia</td>
<td>32</td>
<td>46</td>
</tr>
<tr>
<td>Bladder</td>
<td>Contracture</td>
<td>65</td>
<td>80</td>
</tr>
<tr>
<td></td>
<td></td>
<td>80</td>
<td>85</td>
</tr>
<tr>
<td>Ureters</td>
<td>Stricture</td>
<td>70</td>
<td>100</td>
</tr>
<tr>
<td>Testes</td>
<td>Sterilization</td>
<td>1</td>
<td>2</td>
</tr>
<tr>
<td>Ovaries</td>
<td>Sterilization</td>
<td>2–3</td>
<td>6–12</td>
</tr>
</tbody>
</table>

(Continued)
### TABLE 20.2
A Compilation of Tissue and Organ Sensitivities (Continued)

<table>
<thead>
<tr>
<th>Injury</th>
<th>TD$_{5/5}$, Gy</th>
<th>TD$_{50/5}$, Gy</th>
<th>Field Size</th>
</tr>
</thead>
<tbody>
<tr>
<td>Growing cartilage, child bone</td>
<td>Growth arrest, dwarfing</td>
<td>10</td>
<td>30</td>
</tr>
<tr>
<td>Mature cartilage, adult bone</td>
<td>Necrosis, fracture, sclerosis</td>
<td>60</td>
<td>100</td>
</tr>
<tr>
<td></td>
<td></td>
<td>60</td>
<td>100</td>
</tr>
<tr>
<td>Eye</td>
<td>Retina</td>
<td>Blindness</td>
<td>45</td>
</tr>
<tr>
<td></td>
<td>Cornea</td>
<td></td>
<td>50</td>
</tr>
<tr>
<td></td>
<td>Lens</td>
<td>Cataract</td>
<td>10</td>
</tr>
<tr>
<td>Endocrine</td>
<td>Thyroid</td>
<td>Hypothyroidism</td>
<td>45</td>
</tr>
<tr>
<td></td>
<td>Adrenal</td>
<td>Hypoadrenalism</td>
<td>60</td>
</tr>
<tr>
<td></td>
<td>Pituitary</td>
<td>Hypopituitarism</td>
<td>45</td>
</tr>
<tr>
<td>Peripheral nerves</td>
<td>Neuritis</td>
<td>60</td>
<td>100</td>
</tr>
<tr>
<td>Ear</td>
<td>Middle</td>
<td>Serous otitis</td>
<td>30</td>
</tr>
<tr>
<td></td>
<td>Vestibular</td>
<td>Meniere syndrome</td>
<td>60</td>
</tr>
<tr>
<td>Class III organs</td>
<td>Muscle</td>
<td>Child</td>
<td>Atrophy</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Adult</td>
<td>Fibrosis</td>
</tr>
<tr>
<td></td>
<td>Lymph nodes and lymphatics</td>
<td>Atrophy, sclerosis</td>
<td>50</td>
</tr>
<tr>
<td></td>
<td>Large arteries and veins</td>
<td>Sclerosis</td>
<td>80</td>
</tr>
<tr>
<td></td>
<td>Uterus</td>
<td>Necrosis, perforation</td>
<td>100</td>
</tr>
<tr>
<td></td>
<td>Vagina</td>
<td>Ulcer, fistula</td>
<td>90</td>
</tr>
<tr>
<td>Breast</td>
<td>Child</td>
<td>No development</td>
<td>10</td>
</tr>
<tr>
<td></td>
<td>Adult</td>
<td>Atrophy, necrosis</td>
<td>50</td>
</tr>
</tbody>
</table>

if they are spread out over 6 to 8 weeks, because a substantial amount of stem cell proliferation can occur during this time. For skin, as for oral mucosa, the total dose tolerated depends more on overall time than on fraction size. Because for high-energy x-rays $D_{\text{max}}$ occurs at a depth below the surface, late damage may occur in the dermis in the absence of early reactions in the epidermis. The clinical appearance of radiation fibrosis results from atrophy, leading to contraction of the irradiated area. Telangiectasias developing more than a year after irradiation reflects late-developing vascular injury.

**Skin Appendages: A Special Case**

Within a few days after irradiation, the death of germinal cells results in hair dysplasia (i.e., short, thin hair). The proportion of dysplastic hair is dose dependent. Epilation occurs during the 3rd week, and regrowth may occur after 1 to 3 months.
Sebaceous glands are as sensitive as hair, but sweat glands are less radiosensitive. Regenerated skin may be dry and hairless. An objective measure of skin damage may be obtained by a determination of electrical conductivity, which is influenced by sweat production.

**Hematopoietic System**

Hematopoietic tissues are located primarily in the bone marrow, with 60% located in the pelvis and vertebrae and the remainder in the ribs, skull, sternum, scapula, and proximal sections of the femur and humerus. A tiny fraction of stem cells are found in the circulation. In the normal healthy adult, the liver and spleen have no hematopoietic activity, but they can become active in some circumstances—for example, after partial body irradiation. The pluripotent stem cells go through a period of multiplication and maturation, followed by differentiation without division, before they become mature circulating blood elements of the various types.

The stem cells are particularly radiosensitive. The survival curve has little or no shoulder and a $D_0$ of slightly less than 1 Gy (Chapter 19). There is little sparing from either fractionating the dose or lowering the dose rate. The transit time from stem cell to fully functioning cell, however, differs for the various circulatory blood elements, and these differences account for the complex changes in blood count seen after irradiation.

**Blood Cell Counts after Total Body Irradiation**

A dose as low as 0.3 Gy leads to a reduction in the number of lymphocytes, because they are among the most sensitive cells in the body. After larger doses, the number of all blood cells is altered; lymphopenia is followed by granulopenia, then thrombopenia, and finally anemia.

Following a total body dose of 4 to 6 Gy, there is a temporary increase in the number of granulocytes because of the mobilization of the reserve pool, followed by a rapid fall by the end of the 1st week. The number then remains almost constant before falling again to a minimum value at 18 to 20 days after irradiation. After 1 week of aplasia, regeneration is rapid and takes place more or less simultaneously in platelets, reticulocytes, and granulocytes. After higher doses, the cell minimum is reached earlier and the period of aplasia lasts longer, increasing the possibility of hemorrhage and/or infection, which could prove fatal. At lower doses, around 1 Gy, the depression in granulocyte count is less marked and regeneration less rapid. The general pattern of the blood counts after a modest dose of radiation is illustrated in Figure 20.3.

The survival of stem cells determines the subsequent performance of the bone marrow after total body irradiation, because in the first few hours there is a sudden decrease in the number of pluripotent stem cells and progenitor cells. If the number of stem cells falls below a critical level, production of functional cells essentially stops until partial regeneration of the stem cell compartment occurs and differentiation is allowed to resume. Administration of hematopoietic growth factors can shorten the period of aplasia markedly and accelerate regeneration of all blood cells.

**FIGURE 20.3** The pattern of depletion and recovery of the principal circulating elements of the blood following an intermediate dose of total body radiation. The curves are purely illustrative. The time at which the nadir occurs is a combination of the radiosensitivity of the stem cells and the lifetime of the mature functional cells.
Partial Body Irradiation

In the irradiated volume, the effects of partial body irradiation are analogous to those following total body irradiation. In the unirradiated marrow, the stem cells start dividing within a few hours, and a compensatory hyperplasia attempts to maintain the total production of blood elements. There may also be an extension of hemopoiesis into the long bones, spleen, and liver, which are not normally hemopoietic in the adult human.

With fractionated radiation therapy, the pool of stem cells in the unirradiated volume falls progressively as differentiation is accelerated in an attempt to maintain the circulating blood count. Doses greater than about 30 Gy may cause permanent aplasia in the irradiated volume; hyperplasia and extension of the active bone marrow in the unirradiated volume may persist indefinitely.

Irradiation always reduces the number of stem cells in the bone marrow, and the return to normal may take a long time. This explains why patients remain sensitive to a new insult for months or even years following irradiation.

Radiation and Chemotherapy Agents

Some cytotoxic drugs act essentially on only those cells in cycle, and they have little effect on hematopoietic stem cells, because 90% of them are out of cycle unless the marrow is regenerating following a previous insult. This explains why these drugs show extra toxicity if administered shortly after radiotherapy. The marrow of patients irradiated to a large volume is always more sensitive to cytotoxic drugs, partly because the pool of stem cells is reduced and partly because a greater proportion of stem cells are dividing actively.

The addition of chemotherapy may also have different effects on acute and late reactions. Although many chemotherapeutic agents are dose limited by their toxicity to rapidly proliferating tissues, such as gut mucosa and bone marrow, others (bleomycin, doxorubicin, and cis-platinum) have specific toxicities to slow turnover tissues, such as lung, heart, or kidney. The additive toxicities may therefore also differ for acute and late reactions of tissues included in the irradiated volume. (For more on chemotherapy, see Chapter 27.)

Lymphoid Tissue and the Immune System

The immune system is composed of macrophages and lymphocytes. Macrophages are derived from the same progenitors as granulocytes. These give rise to monocytes, which are transformed into macrophages. This cell line is less radiosensitive than lymphocytes, which are derived, however, from the same pluripotent stem cells.

The B line gives rise to B lymphocytes and plasmocytes, which are responsible for humoral immunologic responses and have life spans of 7 weeks and 2 to 3 days, respectively. Cells of the T line pass through the thymus, where they mature to become T lymphocytes. These cells have a life span of about 3 months and are responsible for cellular immunity and for secreting lymphokines. There are also other types of lymphocytes, including killer cells, responsible for antibody-dependent cytotoxic reactions, and natural killer cells, the function of which is not fully understood.

Total body irradiation leads to a rapid fall in the number of circulating B and T lymphocytes, with the number returning to normal in a few weeks, depending on the dose. The lymphoid tissues (e.g., nodes, spleen) are very radiosensitive and are depleted of cells by quite small radiation doses. Lymphocytes are very radiosensitive, largely because of apoptosis. B cells are more radiosensitive than T cells, and overall, their radiosensitivity, as measured by a clonogenic assay, is similar to that of hematopoietic stem cells.

The effect of irradiation on immune function is complex, depending on the volume irradiated and the number of surviving cells, as well as their capacity to migrate and become lodged in the microenvironment. Total body irradiation is used to inhibit the immune system in preparing patients for an organ transplant, such as a kidney or bone marrow transplant. A total body dose of 3.5 to 4.5 Gy inhibits the immune response against a new antigen, although it is much less effective against an antigen to which the individual is already sensitized. The graft-versus-host reaction after bone marrow transplantation is relatively radioresistant. Partial body irradiation, characteristic of ordinary radiation therapy, has only a limited effect on the immune response, and whether it influences metastatic dissemination is controversial. Total lymphoid irradiation to a dose of 30 to 40 Gy is used for the treatment of lymphomas and leads to a long-lasting T-cell lymphopenia. It can be used to treat autoimmune diseases and also to prepare patients for organ transplants.
Digestive Tract

Oral Mucosa

The cellular organization of the mucosa is similar to that of the skin in that cells multiply in the basal layer and then migrate toward the surface as they differentiate. The life span of the differentiated cells, however, is much shorter than in the epidermis, so there is a more rapid reaction to radiation. The intensity of early mucous membrane reactions is a major factor limiting daily and weekly dose accumulation in the treatment of cancer of the head and neck (Table 20.2). For example, a schedule of 70 Gy delivered in 2-Gy fractions over 7 weeks leads to spotty-confluent mucositis in most patients, which approaches maximum tolerance if the schedule is accelerated to 5.5 weeks by, for example, the use of a concomitant boost technique.

The oral cavity contains various tissue types as well as the mucous membrane. The tongue consists of muscle bundles as well as mucosa with taste buds. The muscles undergo mild progressive fibrosis and fiber atrophy after irradiation. The tonsils are lymphatic tissues and function as sites of antigen process and recognition. Following radiation exposure to a maximum tolerance dose, desquamation of the oral cavity occurs by about day 12, with recovery in 2 to 3 weeks. Desquamation occurs first in the soft palate, followed by the hypopharynx, vallecula, floor of the mouth, cheeks, epiglottis, base of tongue, vocal chords, and dorsum of tongue.

The sequence of events that occurs during radiation therapy for head and neck cancer is so important for the comfort and welfare of the patient that it is worth spelling out in detail. The order of events reflects the different kinetics of the cell populations involved:

1st week: Asymptomatic to slight focal hyperemia and edema caused by dilatation of capillaries in sensitive patients. Sensitivity may be associated with alcohol or tobacco use, chemotherapy, infection (oral candidiasis, herpes simplex virus), or immunosuppression (HIV).

2nd week: Increasing pain and loss of desire to eat. Sense of taste is altered; bitter and acid flavors are most changed, with less change with salty and sweet tastes. Erythema and edema increase, and early desquamative mucositis occurs. Basal cell division has been affected; this layer is being denuded, and vasculoconnective tissue damage becomes apparent. Mucositis is patchy.

3rd week: Mucositis and swelling with depletion of gland secretions leading to difficulty in swallowing. Mucositis plaques are confluent.

4th week: Progression of signs. Confluent mucositis sloughs, resulting in denuded lamina propria. Mucosa becomes covered by fibrin and polymorphonuclear leukocytes.

5th week: Maximum radiation damage apparent by this time. Extreme sensitivity to touch, temperature, and grainy food. Recovery of epithelial layer may begin during therapy. Post-therapy, the basal cells migrate into the area and proliferate. In 2 to 4 weeks, complete resolution is observed. The serous acinar cells of the parotid and submaxillary salivary glands undergo interphase death, and hence salivary dysfunction appears after irradiation, with no threshold dose and little sparing by fractionation. Xerostomia is the main clinical effect that can interfere with nutrition, deteriorate oral hygiene, and predispose a patient to dental problems. TD₅/₅ (the tolerance dose for 5% complication in 5 years) is 32 Gy, and TD₅₀/₅ (the tolerance dose for 50% complication in 5 years) is 46 Gy. Impairment of taste acuity occurs during the 3rd week of a multifraction radiotherapy regime.

Esophagus

The mucosa consists of rapidly dividing cells. After radiation, the esophagus displays an acute mucosal response of esophagitis and increased thickness of the squamous layer. Symptoms appear that include substernal burning with pain on swallowing at about 10 to 12 days after the start of therapy, with a return to normal within a week of the end of therapy. Late effects are related to the muscle layer; they include necrosis and a thickening of the epithelium. This leads to symptoms of difficulty on swallowing and possibly ulcerations after high doses. The tolerance dose is 57.5 Gy in 10 fractions (acute effects limit).

Stomach

Irradiation of the stomach often causes nausea and vomiting immediately afterward. The precursor cells of the gastric glands give rise to mucin-secreting surface columnar cells with short life spans (about 3 or 4 days) and to acid-secreting parietal and pepsinogen-secreting chief cells that have long life spans (hundreds of days). The precursor cells are radiosensitive,
Lungs

The lung is an intermediate- to late-responding tissue. Two waves of damage can be identified: acute pneumonitis at 2 to 6 months after treatment, and fibrosis, which may develop slowly over a period of several months to years. The only symptom of early acute pneumonopathy may be an opacity on a chest x-ray, though it may be accompanied by functional signs, including cough, dyspnea, and respiratory difficulties. Progressive pulmonary fibrosis develops in most patients, including those who previously were asymptomatic, beginning about a year after irradiation. Difficulties in respiratory function increase in severity with time and are generally irreversible. Their severity depends on three factors: volume irradiated, dose, and fraction size. The lung is particularly sensitive to fractionation, with an $\alpha/\beta$ estimated to be about 3 Gy. The most likely target cells are the pulmonary endothelial cells and the type II pneumocytes (cells of the alveolar wall). Type II cells are associated with the production of surfactant during the first few days after irradiation.

Kidneys

Together with the lung, the kidney is among the more radiosensitive late-responding organs. The FSU in the lung is the pulmonary lobule, consisting of the terminal bronchioli and respiratory parenchyma that it serves. The FSUs are arranged in parallel, with a large number of bronchi and alveoli working together; consequently, volume as well as dose are important. Because of this organization of the FSUs, the lung is only dose limiting if large volumes are irradiated and if the remaining lung is not capable of providing adequate function.

The lung is among the most sensitive of late-responding organs. The FSU in the lung is the pulmonary lobule, consisting of the terminal bronchioli and respiratory parenchyma that it serves. The FSUs are arranged in parallel, with each containing only about 1,000 stem and their death leads to early depletion of the surface columnar cell epithelium. Delayed gastric emptying and epithelial denudement are the two main early radiation effects. Peptic ulceration is seen in patients receiving more than 40 Gy to the upper abdomen.

Dyspepsia may be evident in 6 months to 4 years and gastritis in 1 to 12 months. Acute ulceration may occur shortly after the completion of treatment but rarely leads to perforation. At about 5 months, late ulceration and submucosal fibrosis leading to antral fibrosis may occur. Tolerance doses range from 40 to 50 Gy.

Small and Large Intestines

As with the skin and oral mucosa, both early and late complications are observed in the gastrointestinal tract. Acute mucositis frequently occurs, with symptoms such as diarrhea or gastritis, depending on the treatment field. If the dose is limited to 50 to 54 Gy in 2-Gy fractions, acute reactions are seldom dose limiting, and if they do occur, interruption of treatment for a few days usually alleviates the problem. Much more serious are the long-term late sequelae, which may develop either from persistent severe early reactions (consequential late effects) or independently of acute damage in the submucosal, muscular, or serosal layers.

In the small intestine, stem cells are located toward the bottom of the crypts of Lieberkühn. Atrophy of the villus occurs about 2 to 4 days postirradiation. Epithelial denudation is responsible for the acute gut reactions. A regenerative response appears rapidly, and within 2 to 4 days, microcolonies and macrocolonies are detectable. The surviving crypts have at least the same radiosensitivity to reirradiation as the unirradiated crypts, and very little dose is “remembered.”

Late bowel reactions involve all tissue layers and are caused by atrophy of the mucosa caused by vascular injury, with subsequent breakdown resulting from mechanical irritation and bacterial infection, which leads to an acute inflammatory response. In addition, overgrowth of the fibromuscular tissue with stenosis and serosal breakdown and adhesion formation may occur, which may be predisposed to by previous surgery and is related to inflammatory mediators. Fibrosis and ischemia are typical late effects. Tolerance dose is about 50 Gy for the small intestine and slightly higher for the large intestine. Rectal tolerance is about 70 Gy.

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Neurons are nonproliferating end cells in adults; glial cells have a slow rate of turnover, with a small precursor (stem cell) compartment of only about 1%. Endothelial cells also have a slow turnover but can proliferate rapidly after injury. The most important injuries to the brain by radiation are all late syndromes, developing months to years after exposure. Some reactions occur within the first 6 months, including transient demyelination (somnolence syndrome) or the much more serious leukoencephalopathy. Typical radiation necrosis may become evident as early as 6 months, but may be delayed as long as 2 to 3 years. Histopathologic changes that occur within the first year are most likely to involve white matter, whereas for times beyond 6 to 12 months, the gray matter usually shows changes accompanied by vascular lesions such as telangiectasia and focal hemorrhages. A mixture of histologic characteristics is likely to be associated with radionecrosis, which manifests from 1 to 2 years postirradiation, accompanied by cognitive defects.

Spinal Cord
Radiation-induced changes in the spinal cord are similar to those seen in the brain as far as latency, tolerance dose, and histology are concerned. Lhermitte's sign is a demyelinating injury that develops early, by several months after treatment, persists for a few months to a year, but is usually reversible. It may occur at doses as low as 35 Gy, well below the tolerance dose for permanent radiation myelopathy, and its appearance does not predict later more serious problems.

Late damage includes two principal syndromes. The first, occurring from about 6 to 18 months, involves demyelination and necrosis of the white matter; the second is mostly a vasculopathy and has a latency of 1 to 4 years. For the spinal cord, the TD5/5 is about 50 Gy for a 10-cm length irradiated and 55 Gy for a 5-cm length. By 70 Gy, the incidence of myelopathy would be about 50%.

Liver
In terms of radiosensitivity, the liver ranks immediately below the kidney and lung. It shares with these organs the fact that its FSUs are arranged in parallel, so that much larger doses are tolerated if only part of the organ is exposed.

Liver tolerance is dose limiting only if the whole organ is irradiated, as in, for example, total body irradiation prior to bone marrow transplantation. The life span of a hepatocyte is about 1 year, so that, under normal conditions, the cell renewal rate in the liver is very slow. Even large doses apparently are tolerated for a few months, but then hepatic function deteriorates progressively. Fatal hepatitis may result from a fractionated protocol of only 35 Gy if the whole organ is irradiated.

Central and Peripheral Nervous Systems
The nervous system is less sensitive to radiation than other late-responding organs and tissues such as the kidney or lung. Although tolerance doses are frequently quoted at the 5% complication level (i.e., TD5), wide margins of safety in dose usually are included, because damage to these tissues results in severe consequences, including paralysis.

Brain
Three main categories of cells are involved: neurons, vascular endothelial cells, and glial cells. Neurons are nonproliferating end cells in adults; glial cells have a slow rate of turnover, with a small precursor (stem cell) compartment of only about 1%. Endothelial cells also have a slow turnover but can proliferate rapidly after injury. The most important injuries to the brain by radiation are all late syndromes, developing months to years after exposure. Some reactions occur within the first 6 months, including transient demyelination (somnolence syndrome) or the much more serious leukoencephalopathy. Typical radiation necrosis may become evident as early as 6 months, but may be delayed as long as 2 to 3 years. Histopathologic changes that occur within the first year are most likely to involve white matter, whereas for times beyond 6 to 12 months, the gray matter usually shows changes accompanied by vascular lesions such as telangiectasia and focal hemorrhages. A mixture of histologic characteristics is likely to be associated with radionecrosis, which manifests from 1 to 2 years postirradiation, accompanied by cognitive defects.
There is evidence of two components of repair, one with a halftime less than 1 hour and one with a halftime close to 4 hours. The spinal cord is the clearest example of a tissue in which FSUs are arranged in series. The probability of a myelopathy depends critically on the length irradiated for very small lengths, but once the length of the field exceeds a few centimeters, the treatment volume has little effect.

Caution must be exercised in combining radiation with chemotherapy agents, because neurotoxic agents such as methotrexate, cisplatinum, vinblastine, and AraC reduce the tolerance to radiation delivered simultaneously or sequentially.

As far as retreatment is concerned, animal data suggest that by about 2 years, most of the damage from a prior exposure has been repaired; the extent of the repair depends very much on the level of the initial injury.

Peripheral Nerves
Radiation injury of peripheral nerves is probably more common than effects on the spinal cord. It is often said that peripheral nerves are more radioresistant than the cord or brain, but there are few quantitative data to support this. A dose of 60 Gy in a conventional regimen of 2-Gy fractions may lead to a 5% probability of injury, with the probability rising steeply thereafter with increasing dose.

Testes
The seminiferous tubules are composed of two types of cells: sertoli cells, which secrete a hormone that controls the secretion of follicle-stimulating hormone by the hypophysis; and the germinal cells, the hierarchy of which is strictly defined. The stem cells, the type A spermatogonia, have a long cell cycle and divide infrequently. The process of differentiation proceeds through several types of spermatogonia to the spermatocytes, which are the cells in which meiosis occurs. Each spermatocyte gives rise to four spermatids, which finally result in spermatozoa. In humans, the transit time from stem cell to spermatozoa is about 74 days. There is considerable cell loss along the way, so that the amplification factor is much less than might be calculated from the number of divisions that occur.

Leydig cells, which secrete testosterone, also are found in the testis, and their function is regulated by pituitary gonadotropins, prolactin, and luteinizing hormone. This is important in the use of neoadjuvant hormone therapy during the treatment of prostatic cancer.

In humans, a dose as low as 0.1 Gy leads to a temporary reduction in the number of spermatozoa, and 0.15 Gy leads to temporary sterility. Azoospermia lasting for several years occurs after 2 Gy, and permanent azoospermia occurs after about 6 to 8 Gy in 2-Gy fractions. On the other hand, even much larger doses have little effect on the Leydig cells in the adult, so that although irradiation of the testes may lead to sterility, it has little or no effect on the libido.

The stem cells appear to be more radiosensitive than the differentiating spermatogonia, which explains why the duration of azoospermia increases as the dose is increased. Fractionated or continuous low-dose-rate irradiation is more effective than a single acute exposure, because a large proportion of the stem cells are in a radioresistant phase of the cell cycle. If irradiation is protracted, it affects stem cells as they move through the cell cycle into more radiosensitive phases. This accounts for the long-lasting azoospermia seen after relatively low daily doses of scattered radiation reaching the testes during irradiation of the pelvis and also the occurrence of testicular dysfunction seen after years of occupational exposure to ionizing radiation.

Several cytotoxic drugs have substantial effects on spermatogenesis. For example, the alkylating agents included in MOPP (mechlorethamine [Mustargen], vincristine [Oncovin], procarbazine, and prednisone), the combination of chemotherapy agents used at one time for the treatment of Hodgkin lymphoma, led to sterility in almost all patients. Of course, the drug treatment was prolonged and simulated low-dose-rate irradiation, killing stem cells as they came into cycle.

Ovaries
The effects of radiation on the ovaries are quite different from those on the testes, because after the fetal stage, the oocytes no longer divide. They are all present at birth, and their number diminishes steadily with age, reaching zero by the time of menopause. Oocytes are extremely radiosensitive to cell killing; like lymphocytes, they die an interphase death, with a D_0 of only 0.12 Gy. There is little effect of fractionation. Mature follicles and those in the process of maturation are damaged equally by radiation, so that sterilization is immediate (i.e., there is no latent
period, as in the male). Because hormonal secretion is associated with follicular maturation, sterilization by radiation leads to a loss of libido and all of the changes associated with menopause.

**Female Genitalia**

The skin of the vulva reacts like skin elsewhere, but because of moisture and friction, a tolerance dose of 50 to 70 Gy in conventional fractions is considered on the high side.

Acute effects of irradiation of the vagina include erythema, moist desquamation, and confluent mucositis, leading to the loss of vaginal epithelium that may persist for 3 to 6 months. Gross abnormalities in the vagina may include pale color, a thin atrophic mucosa, inflammation, and tissue necrosis with ulceration leading to a fistula. Tolerance doses, however, are high: 90 Gy before ulceration and 100 Gy for the development of a fistula. From intracavitary treatments, doses to the cervix and uterus may reach as high as 200 Gy. Effects seen include atrophy of the endometrial glands and stroma as well as ulceration.

**Blood Vessels and the Vascular System**

The effects of radiation on blood vessels is particularly important, because late damage to many different tissues and organs is mediated to some extent by effects on the vasculature. Blood vessels have a complex structure. A monolayer of endothelial cells lines the interior surface, resting on a connective tissue, the thickness of which depends on the type of vessel. Under normal circumstances, the rate or proliferation of endothelial cells is low, so that following exposure to radiation, cell loss occurs over a period as cells enter mitosis. Regions of constriction appear because of the abnormal proliferation of surviving cells. Denudation of the surface of blood vessels leads to the formation of thromboses and capillary necroses. In the smooth muscle cells that make up the wall of blood vessels, the proportion of cells cycling is very low, so that it takes several years for the number of cells to diminish significantly following irradiation. The loss of muscular fibers plays an important role in the development of late damage that may become evident several years later. Muscle cells are replaced by collagen fibers, vessel walls lose their elasticity, and blood flow is diminished.

Arterial damage may occur after doses of 50 to 70 Gy delivered in conventional fractionation patterns, but capillaries are damaged by doses above about 40 Gy. In general, veins are less sensitive to radiation than arteries.

**Heart**

In its tolerance to radiation, the heart is intermediate between the kidney or lung and the central nervous system. The most common radiation-induced heart injury is acute pericarditis, which seldom occurs during the first year posttherapy. It varies in severity from transient pericarditis, which runs a benign course, to dense sclerosis with cardiac constriction. Anterior chest pain with shortness of breath and low-grade fever may be observed. The threshold dose may be as low as 20 Gy if more than 50% of the heart is irradiated, but higher for partial exposure. A dose of 45 to 50 Gy in conventional fractions produces an 11% incidence. The α/β ratio for the heart is low (about 1 Gy), so that fractionation results in a substantial sparing effect.

Radiation-induced cardiomyopathy results from dense and diffuse fibrosis; it is a slowly evolving lesion that develops over a period of many years and leads to impaired function. Reduced cardiac function is seen in some patients with Hodgkin disease who receive a dose of about 30 Gy to most of the heart. Protection of part of the heart greatly reduces the incidence of symptoms.

The chemotherapy agent Adriamycin (doxorubicin) increases the severity of radiation-induced complications. In addition, Adriamycin may reveal latent radiation damage many years after radiation therapy.

**Bone and Cartilage**

In children, growing cartilage is particularly radiosensitive. Doses as low as 10 Gy can slow growth because of the death of chondroblasts. Above about 20 Gy, the deficit in growth is irreversible. The effects of radiation on bone growth are more serious for higher doses and for younger ages. Sequelae are particularly serious in children younger than 2 years of age, and radiation can affect stature adversely up to the time of puberty.

In the adult, osteonecrosis of the lower maxilla may be a serious complication following radiation therapy for cancer of the buccal cavity. The TD_{5/5} is 50 to 60 Gy; the TD_{50/5} is about 70 Gy for large irradiated volumes. Fractures of the humeral and femoral head are observed if the dose, in conventional fractions, is high. The TD_{5/5} is 52 Gy and the TD_{50/5} is 65 Gy. Fractures of the ribs and clavicle are sometimes seen in patients.
receiving radiotherapy for breast cancer but are generally not serious complications.

**QUANTITATIVE ANALYSIS OF NORMAL TISSUE EFFECTS IN THE CLINIC**

Over the past several decades, with the development of more sophisticated three-dimensional treatment planning systems, numerous studies in the literature have reported associations between dosimetric parameters and normal tissue outcomes. The Quantitative Analysis of Normal Tissue Effects in the Clinic (QUANTEC) article, which appeared in the *International Journal of Radiation Oncology, Biology, and Physics* in 2010 summarized the available data in a clinically useful format. Table 20.3 is the summary table from that publication. The authors concede that the data are far from ideal and that there are many limitations. First and foremost is the fact that the information is extracted largely from publications. Beyond that, there is the problem of evolving fractionation schemes, combined modality therapy, host factors, and so on. For all of these reasons, care must be taken when applying the QUANTEC data in the clinic. It is intended to be an update of the data published by Emami and colleagues in 1991, which is widely used despite the fact that it has often been criticized.

**LENT AND SOMA**

The two large organizations that initiate and coordinate multicenter clinical trials in Europe and North America, namely, the European Organization for Research and Treatment of Cancer (EORTC) and the Radiation Therapy Oncology Group (RTOG), formed working groups to update their system for assessing late injury to normal tissues. This led to the *Late Effects of Normal Tissue (LENT)* conference in 1992. This conference led to the introduction of the SOMA classification for late toxicity. SOMA is an acronym for subjective, objective, management criteria with analytic laboratory and imaging procedures. These scales, specific for each organ, form a scaffold for understanding the expression of late injury because they are the substance of LENT expression. The SOMA scales have been formulated for all anatomic sites listed in Table 20.4. An example for the central nervous system is given in Table 20.5.

**The SOMA Scoring System**

The SOMA scales have been designed to allow the acquisition of data by several different methods, which it is hoped are not inevitably dependent one upon the other:

- **Subjective**—in which the injury, if any, will be recorded from the subject's point of view, that is, as perceived by the patient. This information can be elicited during interviews or derived by asking the patient to complete a carefully designed questionnaire or diary.

- **Objective**—in which the morbidity is assessed as objectively as possible by the clinician during a clinical examination. In this case, the clinician may be able to detect signs of tissue dysfunction that are still below the threshold that will give the patient symptoms but are an indication of how close to tissue tolerance the treatment is or that may be early indicators of more serious problems that are developing and will be expressed later.

- **Management**—which indicates the active steps that may be taken in an attempt to ameliorate the symptoms.

- **Analytic**—involving tools by which tissue function can be assessed even more objectively or with more biologic insight than by simple clinical examination. It is recognized that the tools available for such analysis may differ widely from one center to another and may evolve as the clinical trials progress. The invasiveness and cost of any tool used to quantify the late effects must be reasonable and proportional to the severity of the symptoms and the possible therapeutic consequences. The scales list the techniques that could yield valuable data, but it is not envisaged that all such tests would be feasible or even desirable in all studies.

**APPLICATION OF STEM CELLS TO REGENERATE RADIATION-SENSITIVE ORGANS—SALIVARY GLAND REGENERATION**

No matter how much effort is put into sparing normal tissue toxicity, invariably certain tissues
<table>
<thead>
<tr>
<th>Organ</th>
<th>Volume Segmented</th>
<th>Irradiation Type (Partial Organ Unless Otherwise Stated)</th>
<th>End Point</th>
<th>Dose (Gy), or Dose/Volume Parameters</th>
<th>Rate (%)</th>
<th>Notes on Dose/Volume Parameters</th>
</tr>
</thead>
<tbody>
<tr>
<td>Brain</td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Symptomatic necrosis</td>
<td>$D_{\text{max}} &lt; 60$</td>
<td>$&lt; 3$</td>
<td>Data at 72 and 90 Gy extrapolated from BED models</td>
</tr>
<tr>
<td></td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Symptomatic necrosis</td>
<td>$D_{\text{max}} = 72$</td>
<td>$5$</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Symptomatic necrosis</td>
<td>$D_{\text{max}} = 90$</td>
<td>$10$</td>
<td></td>
</tr>
<tr>
<td>Whole organ</td>
<td>SRS (single fraction)</td>
<td>Symptomatic necrosis</td>
<td>$V_{12} &lt; 5–10 \text{ cc}$</td>
<td>$&lt; 20$</td>
<td></td>
<td>Rapid rise when $V_{12} &gt; 5–10 \text{ cc}$</td>
</tr>
<tr>
<td>Brain stem</td>
<td>Whole organ</td>
<td>Whole organ</td>
<td>Permanent cranial neuropathy or necrosis</td>
<td>$D_{\text{max}} &lt; 54$</td>
<td>$&lt; 5$</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Permanent cranial neuropathy or necrosis</td>
<td>$D_{1-10\text{cc}} \leq 59$</td>
<td>$&lt; 5$</td>
<td></td>
</tr>
<tr>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Permanent cranial neuropathy or necrosis</td>
<td>$D_{\text{max}} &lt; 64$</td>
<td>$&lt; 5$</td>
<td></td>
<td>Point dose $&lt; 1 \text{ cc}$</td>
</tr>
<tr>
<td>Whole organ</td>
<td>SRS (single fraction)</td>
<td>Permanent cranial neuropathy or necrosis</td>
<td>$D_{\text{max}} &lt; 12.5$</td>
<td>$&lt; 5$</td>
<td></td>
<td>For patients with acoustic tumors</td>
</tr>
<tr>
<td>Optic nerve/chiasm</td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Optic neuropathy</td>
<td>$D_{\text{max}} &lt; 55$</td>
<td>$&lt; 3$</td>
<td>Given the small size, 3D-CRT is often whole organ</td>
</tr>
<tr>
<td></td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Optic neuropathy</td>
<td>$D_{\text{max}} = 55–60$</td>
<td>$3–7$</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Optic neuropathy</td>
<td>$D_{\text{max}} &gt; 60$</td>
<td>$&gt; 7–20$</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Optic neuropathy</td>
<td>$D_{\text{max}} \leq 12$</td>
<td>$&lt; 10$</td>
<td></td>
</tr>
<tr>
<td>Spinal cord</td>
<td>Partial organ</td>
<td>3D-CRT</td>
<td>Myelopathy</td>
<td>$D_{\text{max}} = 50$</td>
<td>$0.2$</td>
<td>Including full cord cross section</td>
</tr>
<tr>
<td></td>
<td>Partial organ</td>
<td>3D-CRT</td>
<td>Myelopathy</td>
<td>$D_{\text{max}} = 60$</td>
<td>$6$</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Partial organ</td>
<td>3D-CRT</td>
<td>Myelopathy</td>
<td>$D_{\text{max}} = 69$</td>
<td>$50$</td>
<td></td>
</tr>
<tr>
<td>Partial organ</td>
<td>SRS (single fraction)</td>
<td>Myelopathy</td>
<td>$D_{\text{max}} = 13$</td>
<td>$1$</td>
<td></td>
<td>Partial cord cross section irradiated</td>
</tr>
<tr>
<td>Partial organ</td>
<td>SRS (hypofraction)</td>
<td>Myelopathy</td>
<td>$D_{\text{max}} = 20$</td>
<td>$1$</td>
<td></td>
<td>3 fractions, partial cord cross section irradiated</td>
</tr>
</tbody>
</table>

(Continued)
<table>
<thead>
<tr>
<th>Organ</th>
<th>Volume Segmented</th>
<th>Irradiation Type (Partial Organ Unless Otherwise Stated)</th>
<th>End Point</th>
<th>Dose (Gy), or Dose/Volume Parameters</th>
<th>Rate (%)</th>
<th>Notes on Dose/Volume Parameters</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cochlea</td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Sensory neural hearing loss</td>
<td>Mean dose ≤45</td>
<td>&lt;30%</td>
<td>Mean dose to cochlear, hearing at 4 kHz</td>
</tr>
<tr>
<td></td>
<td>Whole organ</td>
<td>SRS (single fraction)</td>
<td>Sensory neural hearing loss</td>
<td>Prescription dose ≤14</td>
<td>&lt;25%</td>
<td>Serviceable hearing</td>
</tr>
<tr>
<td>Parotid</td>
<td>Bilateral whole parotid glands</td>
<td>3D-CRT</td>
<td>Long-term parotid salivary function reduced to ≤25% of pre-RT level</td>
<td>Mean dose &lt;25</td>
<td>&lt;20%</td>
<td>For combined parotid glands</td>
</tr>
<tr>
<td>Unilateral whole parotid gland</td>
<td>3D-CRT</td>
<td>Long-term parotid salivary function reduced to ≤25% of pre-RT level</td>
<td>Mean dose ≤20</td>
<td>&lt;20%</td>
<td></td>
<td>For single parotid gland</td>
</tr>
<tr>
<td></td>
<td>Bilateral whole parotid glands</td>
<td>3D-CRT</td>
<td>Long-term parotid salivary function reduced to ≤25% of pre-RT level</td>
<td>Mean dose ≤39</td>
<td>&lt;50%</td>
<td>For combined parotid glands (per figure 3 in paper)</td>
</tr>
<tr>
<td>Pharynx</td>
<td>Pharyngeal constrictors</td>
<td>Whole organ</td>
<td>Sympathetic dysphagia and aspiration</td>
<td>Mean dose ≤50</td>
<td>&lt;20%</td>
<td>Based on Section B4 of paper</td>
</tr>
<tr>
<td>Larynx</td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Vocal dysfunction</td>
<td>$D_{max}$ ≤66</td>
<td>&lt;20%</td>
<td>With chemotherapy, based on single study (see Section A4.2 in paper)</td>
</tr>
<tr>
<td></td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Aspiration</td>
<td>Mean dose ≤50</td>
<td>&lt;30%</td>
<td>With chemotherapy, based on single study (see Fig. 1 of paper)</td>
</tr>
<tr>
<td></td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Edema</td>
<td>Mean dose V50 ≤44</td>
<td>&lt;20%</td>
<td>Without chemotherapy, based on single study in patients without larynx cancer</td>
</tr>
<tr>
<td></td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Edema</td>
<td>Mean dose V50 ≤27%</td>
<td>&lt;20%</td>
<td></td>
</tr>
<tr>
<td>Tissue</td>
<td>Whole organ</td>
<td>Treatment</td>
<td>Symptom</td>
<td>Grade</td>
<td>Acute</td>
<td>Mean Dose</td>
</tr>
<tr>
<td>-----------------</td>
<td>-------------</td>
<td>-----------</td>
<td>---------</td>
<td>-------</td>
<td>-------------------</td>
<td>-----------</td>
</tr>
<tr>
<td>Lung</td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Symptomatic pneumonitis</td>
<td>V20</td>
<td>≤30%</td>
<td>≤20%</td>
</tr>
<tr>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Symptomatic pneumonitis</td>
<td>Mean dose</td>
<td>7</td>
<td>13</td>
<td>20</td>
</tr>
<tr>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Symptomatic pneumonitis</td>
<td>Mean dose</td>
<td>7</td>
<td>13</td>
<td>20</td>
</tr>
<tr>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Symptomatic pneumonitis</td>
<td>Mean dose</td>
<td>7</td>
<td>13</td>
<td>20</td>
</tr>
<tr>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Symptomatic pneumonitis</td>
<td>Mean dose</td>
<td>7</td>
<td>13</td>
<td>20</td>
</tr>
<tr>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Symptomatic pneumonitis</td>
<td>Mean dose</td>
<td>7</td>
<td>13</td>
<td>20</td>
</tr>
<tr>
<td>Esophagus</td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Grade ≥3 acute esophagitis</td>
<td>V20</td>
<td>≤34%</td>
<td>5–20</td>
</tr>
<tr>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Grade ≥2 acute esophagitis</td>
<td>Mean dose</td>
<td>V35</td>
<td>&lt;50%</td>
<td>&lt;30</td>
</tr>
<tr>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Grade ≥2 acute esophagitis</td>
<td>Mean dose</td>
<td>V50</td>
<td>&lt;40%</td>
<td>&lt;30</td>
</tr>
<tr>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Grade ≥2 acute esophagitis</td>
<td>Mean dose</td>
<td>V70</td>
<td>&lt;20%</td>
<td>&lt;30</td>
</tr>
<tr>
<td>Heart</td>
<td>Pericardium</td>
<td>3D-CRT</td>
<td>Pericarditis</td>
<td>V20</td>
<td>≤15%</td>
<td>&lt;15</td>
</tr>
<tr>
<td>Pericardium</td>
<td>3D-CRT</td>
<td>Pericarditis</td>
<td>Mean dose</td>
<td>V30</td>
<td>&lt;26%</td>
<td>&lt;15</td>
</tr>
<tr>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Long-term cardiac mortality</td>
<td>V25</td>
<td>&lt;10%</td>
<td>&lt;15</td>
<td>Overly safe risk estimates based on model predictions</td>
</tr>
<tr>
<td>Liver</td>
<td>Whole liver-GTV</td>
<td>3D-CRT or whole organ</td>
<td>Classical RILD</td>
<td>Mean dose</td>
<td>30–32</td>
<td>&lt;5</td>
</tr>
<tr>
<td>Whole liver-GTV</td>
<td>3D-CRT</td>
<td>Classical RILD</td>
<td>Mean dose</td>
<td>30–32</td>
<td>&lt;5</td>
<td></td>
</tr>
<tr>
<td>Whole liver-GTV</td>
<td>3D-CRT</td>
<td>Classical RILD</td>
<td>Mean dose</td>
<td>30–32</td>
<td>&lt;5</td>
<td></td>
</tr>
<tr>
<td>Whole liver-GTV</td>
<td>3D-CRT</td>
<td>Classical RILD</td>
<td>Mean dose</td>
<td>30–32</td>
<td>&lt;5</td>
<td></td>
</tr>
<tr>
<td>Whole liver-GTV</td>
<td>SBRT (hypofraction)</td>
<td>Classical RILD</td>
<td>Mean dose</td>
<td>30–32</td>
<td>&lt;5</td>
<td>3 fractions, for primary liver cancer</td>
</tr>
<tr>
<td>Whole liver-GTV</td>
<td>SBRT (hypofraction)</td>
<td>Classical RILD</td>
<td>Mean dose</td>
<td>30–32</td>
<td>&lt;5</td>
<td>6 fractions, for primary liver cancer</td>
</tr>
</tbody>
</table>

(Continued)
### TABLE 20.3 QUANTEC Summary Table: Approximate Dose/Volume/Outcome Data for Several Organs Following Conventional Fractionation (Unless Otherwise Noted)" (Continued)

<table>
<thead>
<tr>
<th>Organ</th>
<th>Volume Segmented</th>
<th>Irradiation Type (Partial Organ Unless Otherwise Stated)</th>
<th>End Point</th>
<th>Dose (Gy), or Dose/Volume Parameters</th>
<th>Rate (%)</th>
<th>Notes on Dose/Volume Parameters</th>
</tr>
</thead>
<tbody>
<tr>
<td>Whole liver–GTV</td>
<td></td>
<td>SBRT (hypofraction)</td>
<td>Classical RILD</td>
<td>Mean dose: 15, 20</td>
<td>&lt;5, 5</td>
<td>3 fractions, for liver metastases</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Kidney</td>
<td>Bilateral whole kidney</td>
<td>SBRT (hypofraction)</td>
<td>Classical RILD</td>
<td>Mean dose: 15, 18</td>
<td>&lt;5</td>
<td>Critical volume based in 3–5 fractions</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Bilateral whole organ or 3D-CRT</td>
<td>Clinically relevant renal dysfunction</td>
<td>Mean dose: 28</td>
<td>&lt;50</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>For combined kidney</td>
</tr>
<tr>
<td>Stomach</td>
<td>Whole organ</td>
<td>Whole organ</td>
<td>Ulceration</td>
<td>D100&lt;sup&gt;c&lt;/sup&gt;: 45</td>
<td>&lt;7</td>
<td></td>
</tr>
<tr>
<td>Small bowel</td>
<td>Individual small bowel loops</td>
<td>3D-CRT</td>
<td>Grade ≥3 acute toxicity&lt;sup&gt;d&lt;/sup&gt;</td>
<td>V15: 120 cc</td>
<td>&lt;10</td>
<td>Volume based on segmentation of the individual loops of bowel, not the entire potential space within the peritoneal cavity</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>3D-CRT</td>
<td>Grade ≥3 acute toxicity&lt;sup&gt;d&lt;/sup&gt;</td>
<td>V45: 195 cc</td>
<td>&lt;10</td>
<td>Volume based on the entire potential space within the peritoneal cavity</td>
</tr>
<tr>
<td>Organ</td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Grade ≥2 late rectal toxicity</td>
<td>Grade ≥3 late rectal toxicity</td>
<td>V50</td>
<td>≤50%</td>
</tr>
<tr>
<td>----------------</td>
<td>-------------</td>
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<td>-----------------------------</td>
<td>-----------------------------</td>
<td>-----</td>
<td>------</td>
</tr>
<tr>
<td>Rectum</td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Grade ≥2 late rectal toxicity</td>
<td>Grade ≥3 late rectal toxicity</td>
<td>V60</td>
<td>≤35%</td>
</tr>
<tr>
<td></td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Grade ≥2 late rectal toxicity</td>
<td>Grade ≥3 late rectal toxicity</td>
<td>V65</td>
<td>≤25%</td>
</tr>
<tr>
<td></td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Grade ≥2 late rectal toxicity</td>
<td>Grade ≥3 late rectal toxicity</td>
<td>V70</td>
<td>≤20%</td>
</tr>
<tr>
<td></td>
<td>Whole organ</td>
<td>3D-CRT</td>
<td>Grade ≥2 late rectal toxicity</td>
<td>Grade ≥3 late rectal toxicity</td>
<td>V75</td>
<td>≤15%</td>
</tr>
</tbody>
</table>

| Bladder        | Whole organ | 3D-CRT | Grade ≥3 late RTOG D_max | ≤65 | ≤6 |
| Penile bulb    | Whole organ | 3D-CRT | Severe erectile dysfunction | Mean dose to 95% of gland | ≤50 | ≤35 |

| Whole organ | 3D-CRT | Severe erectile dysfunction | D90 | ≤50 | ≤35 |
| Whole organ | 3D-CRT | Severe erectile dysfunction | D60–70 | ≤70 | ≤55 |

Prostate cancer treatment

Variations in bladder size/shape/location during RT hamper ability to generate accurate data

Prostate cancer treatment

Based on current RTOG 0415 recommendation

All data are estimated from the literature summarized in the QUANTEC reviews (unless otherwise noted). Clinically, these data should be applied with caution. Clinicians are strongly advised to use the individual QUANTEC articles to check the applicability of these limits to the clinical situation at hand. They largely do not reflect modern IMRT.

All at standard fractionation (i.e., 1.8–2.0 Gy per daily fraction) unless otherwise noted.

Non-TBI

With combined chemotherapy

Dx, minimum dose received by the “hottest” X % (or X cc) of the organ

Severe xerostomia is related to additional factors including the doses to the submandibular glands.

Estimated by Dr. Eisbruch

Classic RT-induced liver disease (RILD) involves anicteric hepatomegaly and ascites, typically occurring between 2 weeks to 3 months after therapy. Classic RILD also involves elevated alkaline phosphatase (more than twice the upper limit of normal or baseline value).

For optic nerve, the cases of neuropathy in the 55–60 Gy range received ~59 Gy (see optic nerve paper for details). Excludes patients with pituitary tumors where the tolerance may be reduced.
cannot be spared and are subject to radiation-induced damages. In the treatment of head and neck cancer, a major complication of normal tissues is loss of salivary gland function, resulting in xerostomia. Salivary glands can be considered a radiation-sensitive tissue that differs from other radiation-sensitive tissues in not being rapidly proliferating and is well differentiated. Furthermore, the contralateral gland does not compensate for loss of function of the irradiated gland. The clinical response of the salivary gland differs in different individuals with doses as low as 10 Gy, leading to a marked reduction in salivary excretion. The variation in radiosensitivity of salivary glands is not well understood, but seems to involve apoptotic cell death.

Over the years, different approaches have been used to prevent or restore salivary gland function, including radioprotectors, in situ manipulation of ductal cells, and implantation of an artificial salivary gland. All of these past approaches have met with failure or have failed to be clinically adopted. These past failures and the advent of stem cell biology has brought forth the
### TABLE 20.5 Central Nervous System SOMA

<table>
<thead>
<tr>
<th>Subjective</th>
<th>Objective</th>
<th>Management</th>
<th>Analytic</th>
</tr>
</thead>
<tbody>
<tr>
<td>Headache</td>
<td>Neurologic deficit</td>
<td>Anticonvulsives</td>
<td>MRI</td>
</tr>
<tr>
<td>Somnolence</td>
<td>Cognitive function</td>
<td>Steroids</td>
<td>CT</td>
</tr>
<tr>
<td>Intellectual deficit</td>
<td>Mood and personality changes</td>
<td>Sedation</td>
<td>MRS</td>
</tr>
<tr>
<td>Functional competence</td>
<td>Seizures</td>
<td></td>
<td>PET</td>
</tr>
<tr>
<td>Memory</td>
<td></td>
<td></td>
<td>Cerebrospinal fluid</td>
</tr>
</tbody>
</table>


### TABLE 20.6 LENT and SOMA Scoring System and Grading Categories

<table>
<thead>
<tr>
<th>Grade 1</th>
<th>Grade 2</th>
<th>Grade 3</th>
<th>Grade 4</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Subjective:</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ascending order of severity of symptoms perceived by patient (e.g., pain)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Occasional and minimal</td>
<td>Intermittent and tolerable</td>
<td>Persistent and intense</td>
<td>Refractory and excruciating</td>
</tr>
<tr>
<td><strong>Objective:</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Signs that can be assessed by clinician (e.g., neurologic deficit)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Barely detectable</td>
<td>Easily detectable</td>
<td>Focal motor signs, vision disturbances, etc.</td>
<td>Hemiplegia, hemisensory deficit, etc.</td>
</tr>
<tr>
<td><strong>Management:</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Active steps taken to ameliorate symptoms (e.g., pain)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Occasional nonnarcotic</td>
<td>Regular nonnarcotic</td>
<td>Regular narcotic</td>
<td>Surgical intervention</td>
</tr>
<tr>
<td><strong>Analytic:</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Findings that are quantifiable (e.g., CT and MRI, special laboratory tests)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

possibility of using autologous stem cell transfer as a means of regenerating salivary glands after the completion of radiotherapy. Experimentally, salivary stem cells have been isolated, cultured in vitro, and reinjected into mice after irradiation. These reinjected stem cells were shown to repopulate the salivary glands and increased the salivary production of these mice. Therefore, the proof of concept experiment in rodents has demonstrated the feasibility of such an approach. Although this is an exciting result, there is still work that needs to be addressed to apply this to head and neck cancer patients. First, the in vitro culture of salivary stem cells needs to take place at least for several months, and the spontaneous differentiation of these cells in cell culture must be prevented. Second and probably more problematic is overcoming radiation-induced fibrosis in the area of the exposed gland. Both of these problems are not insurmountable as better culture methods are being developed to prevent stem cell differentiation in vitro, and small molecule inhibitors of fibrosis are making their way into clinical trials. Thus, the first application of stem cell biology to regenerate irradiated salivary glands and restore their function will most likely occur within the next decade.

SUMMARY OF PERTINENT CONCLUSIONS

- Most effects of radiation on normal tissues are caused by cell killing, but some, such as nausea, vomiting, or fatigue experienced by patients following irradiation of large volumes including the abdomen, may be mediated by radiation-induced inflammatory cytokines.
- Apparent radioresponsiveness of a tissue depends on inherent sensitivity of cells, kinetics of the tissue or cell population, and the way cells are organized in that tissue.
- Sensitivity of actively dividing cells is expressed by their survival curve for reproductive integrity.
- The radiation dose needed to destroy the functioning ability of a differentiated cell is far greater than that necessary to stop the mitotic activity of a dividing cell.
- The shape of the dose–response relationship for functional end points, obtained from multifraction experiments, is more pertinent to radiotherapy than clonogenic assays.
- The time interval between irradiation and its expression in tissue damage depends on the life span of mature functional cells and the time it takes for a cell born in the stem compartment to mature.
- Hyperthermia damage is expressed early compared with radiation damage (Chapter 28).
- Both early and late effects may develop in one organ system because of injury to different target cell populations or tissue elements.
- The ratio $\alpha/\beta$ (the dose at which the linear and quadratic components of radiation damage are equal) may be inferred from multifraction experiments in systems scoring nonclonogenic end points.
- Tolerance doses for late effects are more sensitive to changes in dose per fraction (low $\alpha/\beta$ value) compared with tolerance doses for early effects.
- Spatial arrangement of FSUs is critical to the tolerance of some normal tissues.
- In some tissues (e.g., spinal cord), the FSUs are arranged serially (like links in a chain), and the integrity of each is critical to organ function.
- Tissues with a serial organization (e.g., spinal cord) have little or no functional reserve, and the risk of developing a complication is less dependent on volume irradiated than for tissues with a parallel organization. The risk of complication is strongly influenced by high-dose regions and hot spots.
- A tissue with intrinsically high tolerance may fail as a result of the inactivation of a small segment (as in the spinal cord); a tissue with an intrinsically low tolerance (kidney and lung) may lose a substantial number of its functional units without impact on clinical tolerance.
- Casarett’s classification of tissue radiosensitivity is based on histopathologic observations.
- In terms of radiosensitivity based on histologic observation of cell death, parenchymal cells fall into four categories, from most sensitive to most resistant:
  1. Stem cells of classic self-renewal tissues, which divide regularly
  2. Differentiating intermitotic cells, which divide regularly but in which there is
some differentiation between divisions and which are variably differentiated

3. Reverting postmitotic cells, which do not divide regularly but can divide under the appropriate stimulus

4. Fixed postmitotic cells, which are highly differentiated and appear to have lost the ability to divide

- Connective tissue and blood vessels are intermediate in radiosensitivity between groups 2 and 3.
- Michalowski’s classification divides tissues into H- and F-type populations, which respond differently to radiation.
- Many tissues are a hybrid of H-type and F-type.
- The response of a tissue is influenced greatly by a host of growth factors, including interleukin-1 and 6, basic fibroblast growth factor, platelet-derived growth factor β, TGF-β, and TNF.
- Early radiation response in the skin is caused by damage to the epidermis; the late response reflects damage to the dermis.
- The hematopoietic system is very sensitive to radiation, especially the stem cells. The complex changes seen in peripheral blood count after irradiation reflect differences in transit time from stem cell to functioning cell for the various circulatory blood elements.
- The effect of irradiation on the immune function is complex, depending on the volume irradiated and the number of surviving cells. A total body dose of 3.5 to 4.5 Gy inhibits the immune response against a new antigen.
- The cellular organization of the lining of the gastrointestinal tract is similar to that of the skin, but the life span of the differentiated cells is shorter. Both early and late sequelae can occur.

**Stomach:** Irradiation of the stomach often leads to nausea and vomiting. Tolerance doses range from 40 to 50 Gy.

**Small and large intestines:** Both early and late complications can occur. Tolerance dose is about 50 Gy for the small intestine, slightly higher for the large intestine, and 70 Gy for the rectum.

- The lung is an intermediate- to late-responding tissue. Two waves of damage can be identified: an acute pneumonitis and a later fibrosis. The lung is among the most sensitive late-responding organs. Pulmonary damage also may occur following chemotherapy.
- Together with the lung, the kidney is among the more radiosensitive late-responding critical organs. FSUs are in parallel, with only about 1,000 stem cells in each. A dose of about 30 Gy in 2-Gy fractions to both kidneys results in nephropathy.
- In terms of radiosensitivity, the liver ranks immediately below the kidney and lung. FSUs are in parallel, so that much larger doses are tolerated if only part of the organ is exposed. Fatal hepatitis may result from 35 Gy (conventional fractionation) to the whole organ.
- Cell renewal is low in the bladder epithelium, so proliferation following irradiation is delayed. Frequency of urination increases in parallel with loss of surface cells. Absence of surface cells explains irritation by urine.
- The nervous system is less sensitive to radiation than other late-responding organs, such as the kidney or lung.

**Brain:** Histopathologic changes that occur in the first year are most likely to involve white matter; at later times, gray matter usually shows changes accompanied by vascular lesions. Radionecrosis may occur accompanied by cognitive defects.

**Spinal cord:** Early demyelinating injuries may develop after doses as low as 35 Gy but are usually reversible. For late damage, the TD5/5 is about 50 Gy for a 10-cm length of cord. By 70 Gy in conventional fractions, the incidence of myelopathy would be 50%. FSUs are in series, but once the field exceeds a few centimeters, the treatment volume has little effect. Tolerance dose shows little dependence on overall...
time but depends critically on dose per fraction (\(\alpha/\beta\) is low). If two doses per day are used, the interfractionation interval must be more than 6 hours, because there is a slow component of repair.

- In the testes, a dose of 0.1 to 0.15 Gy leads to temporary sterility. A dose of 6 to 8 Gy in 2-Gy fractions leads to permanent sterility. Such doses have little effect on libido. The stem cells are more radiosensitive than the differentiated cells, so continuous or fractionated radiation is more effective than a single acute dose.

- Sterilization by radiation to the ovaries is immediate (no latent period, as in the male) and leads to all the changes associated with menopause.

- Among the female genitalia, tolerance doses for the vagina are high: 90 Gy before ulceration and 100 Gy for the development of a fistula. For intracavitary treatment, doses to the cervix and uterus may reach 200 Gy.

- Late damage to many different tissues and organs is mediated to some extent by effects on the vasculature. Arterial damage may occur after fractionated doses of 50 to 70 Gy, but capillaries are damaged by doses above about 40 Gy.

- In its tolerance to radiation, the heart is intermediate between the kidney or lung and the central nervous system. The most common radiation-induced heart injury is acute pericarditis, which seldom occurs in the first year posttherapy. A dose of 40 to 50 Gy in conventional fractions induces about an 11% incidence. The \(\alpha/\beta\) ratio is low (1 Gy), so that fractionation results in a substantial sparing. Protection of part of the heart reduces symptoms.

- Growing cartilage is particularly radiosensitive in children: 10 Gy can slow growth, and deficits in growth are irreversible above about 20 Gy. In the adult, osteoporosis of the lower mandible may be a serious complication following radiotherapy for cancer of the buccal cavity. Fractures of the humeral or femoral head may occur; the TD50/5 is about 65 Gy.

- Over the past several decades, with the development of more sophisticated three-dimensional treatment planning systems, numerous studies in the literature have reported associations between dosimetric parameters and normal tissue outcomes. QUANTEC summarized the available data in a clinically useful format. It is intended to be an update of the data published by Emami and colleagues in 1991, which is widely used despite the fact that it has often been criticized.

- The RTOG and EORTC introduced the SOMA classification for LENT: SOMA is an acronym for subjective, objective, management criteria with analytic laboratory and imaging procedures.

**BIBLIOGRAPHY**


A wide range of experimental tumors of various histologic types have been developed for radiobiologic studies. To produce a large number of virtually identical tumors, propagation by transplantation from one generation of animals to the next is used, which makes it mandatory that the animals be isologous. In practice, pure inbred strains of rats or mice are used and are maintained by brother–sister mating, which also serves the function of reducing the variability among animals to a minimum.

The tumor from a donor animal is removed aseptically and, if possible, prepared into a single-cell suspension; this is accomplished by separating the cells with an enzyme such as trypsin and then forcing them through a fine wire mesh. To effect a transplant, $10^4$ to $10^6$ cells are inoculated subcutaneously into each of a large group of recipient animals of the same strain. The site of transplantation varies widely; the flank or back is commonly used, but sometimes a special tumor requires a particular site, such as the brain. Some tumors cannot be handled in this way and must be propagated by transplanting a small piece of tumor rather than a known number of single cells; this is obviously less quantitative. Within days or weeks, depending on the type of tumor and the strain of animals, palpable tumors appear in the recipient animals that are uniform in size, histologic type, and so on. Hundreds to thousands of animals can be used, which makes it possible to design highly quantitative studies of tumor response to different radiations, fractionation regimens, sensitizers, and combinations of radiation and chemotherapeutic agents.

There are five commonly used techniques to assay the response of solid tumors to a treatment regimen:

1. Tumor growth measurements
2. Tumor cure (TCD50) assay
3. Tumor cell survival determined in vivo by the dilution assay technique
4. Tumor cell survival assayed by the lung colony assays
5. Tumor cell survival using in vivo treatment followed by in vitro assay

Each of these methods is discussed briefly in this chapter. These assays take into account both intrinsic cell sensitivity to ionizing radiation as well as the influence of the microenvironment. Before we discuss the techniques used to assess tumor response to radiation, we should briefly revisit how the type of cell death and the tumor microenvironment can affect tumor response to therapy, in particular the role of apoptotic cell death.

**APOPTOSIS IN TUMORS**

It is generally thought that irradiated cells die in attempting the next or a subsequent mitosis. However, this is not the only form of cell death. Programmed cell death, or *apoptosis*, also occurs in both normal tissues and tumors, spontaneously,
because of irradiation, and can be triggered by changes in the tumor microenvironment.

In Chapter 22, it is pointed out that tumors grow much more slowly than would be predicted from the cell cycle time of the individual cells and the fraction of cells actively dividing. One of the reasons for this “cell loss,” as it is called, is random cell death resulting from apoptosis.

Studies with transplanted mouse tumors, as well as human tumors growing as xenografts in nude mice, have shown that the importance of apoptosis as a mechanism of cell death after x-irradiation varies substantially. Apoptosis was most important in lymphomas, essentially absent in sarcomas, and intermediate and very variable in carcinomas. In a mouse lymphoma, for example, 50% to 60% of the cells may show signs of dying an apoptotic death by 3 hours after irradiation, whereas in a sarcoma, there may be so few apoptotic cells that the process is of little significance. If a tumor responds rapidly to a relatively low dose of radiation, it generally means that apoptosis is involved, because the process peaks at 3 to 5 hours after irradiation.

Susceptibility to the induction of apoptosis also may be an important factor determining radiosensitivity, because programmed cell death appears to be a prominent early effect in radiosensitive mouse tumors and essentially absent in radioresistant tumors. In particular, the transformation of mouse cells of different histologic origins with oncogenes in cell culture makes them particularly sensitive to DNA damage-induced apoptosis. The concern with using cell lines that are highly susceptible to apoptosis is that cell culture will act as a selective pressure over time to select for variants of these cells that have lost their apoptotic sensitivity, thus changing their in vivo sensitivity to radiation and cytotoxic drugs used in cancer therapy.

Changes in the microenvironment can also influence the sensitivity of tumor cells to therapy (Fig. 21.1). Studies have shown that changes in tumor oxygenation, pH, and growth factors can induce apoptotic cell death in a subset of tumor cells that represent a therapeutically resistant population. Therefore, cells that are sensitive to apoptosis will be more susceptible to killing by

**FIGURE 21.1** Regions of hypoxia form within solid tumors as a result of the inefficient and disorganized vasculature. Hypoxic regions represent a gradient of oxygen concentrations, the highest being nearest the vessels and the lowest being the farthest away. In the most hypoxic regions, p53 is stabilized and induces apoptosis. A selection pressure therefore exists to loose p53 and allows the clonal expansion of cells. These cells are also resistant to chemotherapy and radiotherapy, both of which require an efficient blood supply. (From Hammond EM, Giaccia AJ. Hypoxia-inducible factor-1 and p53: friends, acquaintances, or strangers? *Clin Cancer Res.* 2006;12:5007–5009, with permission.)

Clonal expansion of cells with mutant p53
the environmentally restrictive conditions in experimental tumors and will be better able to be controlled by radiotherapy and chemotherapy. Cells with diminished apoptotic potential, which represent most tumor cell lines maintained in culture, will have lost this rapid response to radiotherapy but can still die by a mitotic cell death. Cells with diminished apoptotic sensitivity caused by genetic mutations in critical apoptotic pathways, such as the \( p53 \) tumor suppressor gene or the Bcl-2 oncogene (see Chapter 18), are probably more reflective of human solid tumors that are treated clinically.

**TUMOR GROWTH MEASUREMENTS**

Tumor growth measurement is possibly the simplest end point to use and involves the daily measurement of each tumor to arrive at a mean diameter. For tumor growth experiments, a large number of transplanted tumors are prepared as previously described. When they have grown to a specified size (e.g., a diameter of 8 to 10 mm in rats or 2 to 4 mm in mice), they are treated according to the plan of the particular experiment. Figure 21.2 illustrates the variation of tumor size with time for unirradiated controls and tumors given a single dose of x-rays. The untreated tumors grow rapidly at a relatively uniform rate; the radiation treatment causes a temporary shrinkage of the tumor, followed by regrowth.

Two different methods have been used to score the tumor response. Barendsen and his colleagues have used growth delay, illustrated in Figure 21.2, as the time taken after irradiation for the tumor to regrow to the size it was at the time of irradiation. Clearly, this index of response is only suitable for tumors that shrink significantly after irradiation. For tumors that do not shrink so obviously, a more convenient index of growth delay is the time taken for the irradiated tumor to grow to some specified size after exposure, compared with controls. Either index of growth delay increases as a function of radiation dose. Figure 21.3A shows growth curves for a rat rhabdomyosarcoma irradiated with various doses of x-rays or fast neutrons. In Figure 21.3B, growth delay is expressed as a function of radiation dose.

**TUMOR CURE (TCD\(_{50}\)) ASSAY**

Tumor control provides data of most obvious relevance to radiotherapy. In experiments of this kind, a large number of animals with tumors of uniform size are divided into separate groups, and the tumors are irradiated locally with graded doses. The tumors subsequently are observed regularly for recurrence or local control. The proportion of tumors that are locally controlled can be plotted as a function of dose, and data of this kind are amenable to a sophisticated statistical analysis to determine TCD\(_{50}\),
Figure 21.3

A: Volume changes of rhabdomyosarcomas in rats after irradiation. Curve 1 represents the growth of the unirradiated control tumors. Curves 2, 4, 6, and 7 refer to tumors irradiated with 10 to 40 Gy of 300-kV x-rays. Curves 3 and 5 refer to tumors irradiated with 4 and 8 Gy of 15-MeV d\(^{10}\) + T fast neutrons. B: Growth delay of rhabdomyosarcomas in rats as a function of dose of x-rays (curve 2) or fast neutrons (curve 1). A and C indicate the doses of neutrons and x-rays, respectively, required to "cure" 90% of the tumors, calculated based on the cell survival curves. B indicates the observed TCD\(_{90}\) for x-rays. Note the good agreement between calculated and observed values of the TCD\(_{90}\) for x-rays. (Adapted from Barendsen GW, Broerse JJ. Experimental radiotherapy of a rat rhabdomyosarcoma with 15 MeV neutrons and 300 kV x-rays. I. Effects of single exposures. Eur J Cancer. 1969;5:373–391, with permission.)
rising to 51.1 Gy for 2 fractions and to 84 Gy if the radiation is delivered in 10 equal fractions. This indicates that a marked and extensive repair of sublethal damage has taken place during a multifraction regimen.

DILUTION ASSAY TECHNIQUE

The dilution assay technique was devised by Hewitt and Wilson, who used it to produce the first \textit{in vivo} survival curve in 1959. They used a lymphocytic leukemia of spontaneous origin in mice. A single-cell suspension can be prepared from the infiltrated liver of an animal with advanced disease and the tumor transplanted by injecting known numbers of cells into the peritoneal cavities of recipient mice, which subsequently develop leukemias. The leukemia can be transmitted, on average, by the injection of only two cells; this quantity—the number of cells required to transmit the tumor to 50\% of the animals—is known as the dose at which 50\% of the tumors are locally controlled. This quantity is highly repeatable from one experiment to another in an inbred strain of animals.

Suit and his colleagues, over a period of more than 30 years, have made an extensive study of the response to radiation of a mammary carcinoma in C3H mice. Data from a typical experiment are presented in Figure 21.4. Tumors were propagated by transplanting $4 \times 10^9$ cells into the outer portion of the mouse ear, and irradiations were performed when the tumors had grown to a volume of about 4 mm$^3$. A brass circular clamp was fitted across the base of the ear and maintained for at least a minute before the initiation of the irradiation, so that the tumors were uniformly hypoxic. Single-dose, two-dose, and ten-dose experiments were performed, with a 24-hour interval between dose fractions. Tumor control results are shown in Figure 21.4. The TCD$_{50}$ for a single treatment is 45.75 Gy, rising to 51.1 Gy for 2 fractions and to 84 Gy if the radiation is delivered in 10 equal fractions. This indicates that a marked and extensive repair of sublethal damage has taken place during a multifraction regimen.
as TD_{50}. The dilution assay technique became the basis for obtaining an \textit{in vivo} cell survival curve.

The procedure used, illustrated in Figure 21.5, is as follows. An animal containing the tumor may be irradiated to a given dose of radiation, for example, 10 Gy. A single-cell suspension is then prepared from the infiltrated liver, the cells are counted and diluted, and various numbers of these cells are injected intraperitoneally into groups of recipient animals. It is then a matter of observation and calculation to determine how many irradiated cells are required to produce a tumor in half of the animals inoculated with that given number of cells. Suppose, for instance, that it takes 20 irradiated cells, on average, to transmit the tumor; because it is known that only 2 clonogenic cells are needed to transmit the tumor, it is a simple matter to decide that in the irradiated population of cells,
survival curve. To obtain a survival curve characteristic of aerated conditions, it is necessary either to remove the cells from the donor animal and irradiate them in a petri dish in which they are in contact with air or to inject hydrogen peroxide into the peritoneal cavity of the mouse before irradiation so that oxygen is available to the tumor cells during the irradiation. If this is done, $D_0$ is about 1.3 to 1.6 Gy.

**LUNG COLONY ASSAY**

Hill and Bush have devised a technique to assay the clonogenicity of the cells of a solid tumor irradiated *in situ* by injecting them into the recipient animals and counting the number of lung colonies produced. The general principles of the method are illustrated in Figure 21.7. The tumor used in these studies was the KHT sarcoma, which is a transplantable tumor that arose originally in a C3H mouse and which has been propagated serially through many generations. Tumors are irradiated *in situ*, after which they are removed and made into a preparation of single cells by a combined trypsinization and mechanical procedure. A known number of cells then are mixed with a large number of heavily irradiated tumor cells and injected intravenously.
Cells can be readily transferred from *in vivo* to *in vitro* and back. In one generation, they may grow as a solid tumor in an animal, and in the next, as a monolayer in a petri dish. The three most commonly used systems are a rhabdomyosarcoma in the rat (Hermens and Barendsen), a fibrosarcoma in the mouse (McNally), and the EMT6 mammary tumor in the mouse (Rockwell and Kallman).

The steps involved in this method are illustrated in Figure 21.8. This method combines many of the advantages of the *in vitro* and *in vivo* techniques. The tumors are treated *in vivo* in a natural environment, so that the cellular response is modified by the various factors that are important in determining gross tumor response. After treatment, each tumor is removed and prepared into a single-cell suspension, and the cell suspension is injected into recipient mice. About 3 weeks later, these mice are sacrificed, and the colonies formed in the lungs are readily countable. The number of lung colonies is a measure of the number of surviving clonogenic cells in the injected suspension.

The lung colony technique is not confined to the KHT sarcoma but has been used with other tumor cells. For example, the demonstration of the absence of repair of potentially lethal damage after neutron irradiation involved the use of the Lewis lung carcinoma, and the fraction of surviving cells was assayed by counting lung colonies.

**IN VIVO/IN VITRO ASSAY**

A limited number of cell lines have been adapted so that they grow either as a transplantable tumor in an animal or as clones in a petri dish. These cell lines can be readily transferred from *in vivo* to *in vitro* and back. In one generation, they may grow as a solid tumor in an animal, and in the next, as a monolayer in a petri dish. The three most commonly used systems are a rhabdomyosarcoma in the rat (Hermens and Barendsen), a fibrosarcoma in the mouse (McNally), and the EMT6 mammary tumor in the mouse (Rockwell and Kallman).

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**FIGURE 21.7** The lung colony assay system. The tumor is irradiated *in situ*, after which it is excised and made into a single-cell suspension. A known number of cells are then injected intravenously into the recipient animals. About 3 weeks later, the recipient animals are sacrificed and the colonies that have formed in the lungs are counted. The number of lung colonies is a measure of the number of surviving clonogenic cells in the injected suspension. (Adapted from Hill RP, Bush RS. The effect of continuous or fractionated irradiation on a murine sarcoma. *Br J Radiol*. 1973;46:167–174, with permission.)

**FIGURE 21.8** The principle of the *in vivo/in vitro* assay system using the rhabdomyosarcoma in the rat. The solid tumor in the animal can be removed and the tumor cells assayed for colony formation in petri dishes. This cell line can be transferred back and forth between the animal and the petri dish. (Adapted from a drawing courtesy of Drs. G.W. Barendsen and J.J. Broerse.)
concentration is counted in a hemocytometer or electronic cell counter. Known numbers of cells can then be transferred to petri dishes containing fresh growth medium, and the proportion of clonogenic cells can be determined by counting colonies 10 days later. The speed, accuracy, and relative economy of the in vitro system replace the expense and inconvenience of the recipient animals in the dilution assay technique.

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### XENOGRAFTS OF HUMAN TUMORS

A **xenograft** is a transplant from one species to another. In the cancer field, this usually refers to a human tumor transplanted into a laboratory animal. If the recipient animal has a normal immune system, a xenograft should not grow, but there are two main ways in which growth has been achieved. First, animal strains have been developed that are congenitally immune deficient. Best known are nude mice, which, in addition to being hairless, also lack a thymus. Many human tumors grow under the skin of nude mice. More recently, there have been nude rats and severe combined immune-deficient (SCID) mice, which suffer from the severe combined immunodeficiency syndrome and are deficient in both B cell and T cell immunity. Second, it is possible to severely immunosuppressed mice by the use of radiation or drugs or a combination of both, to the point at which they accept human tumor grafts. It is important to recognize that neither type of host completely fails to reject the human tumor cells: Rejection processes are still present, and these complicate the interpretation of in situ tumor therapeutic studies.

Despite the limitations, various human tumor cells have been grown as xenografts in immune-deficient animals. Steel has estimated that more than 300 individual human tumors have been investigated in this way. Breast and ovarian tumors generally have been difficult to graft, with grafting of melanomas and tumors of the colon and bronchus being relatively successful.

Xenografts retain human karyotypes through serial passages and maintain some of the response characteristics of the individual source human tumors; to this extent, they have great advantages over mouse tumors. There are, however, certain drawbacks. First, there is a tendency for the tumor to be rejected, so that observing tumor control as an end point can be misleading. Growth delay and cell survival studies, on the other hand, are probably less affected. Second, human tumor cells do undergo kinetic changes and cell selection if transplanted into mice. For example, xenografts commonly have doubling times about one-fifth of the values observed in humans, so that increased responsiveness should be expected to proliferation-dependent chemotherapeutic agents. Third, although the histologic characteristics of the human source tumors are usually well maintained by xenografts, the stromal tissue is of mouse origin. Consequently, xenografts of human tumor cells are not much more valid than murine tumors for any studies in which the vascular supply plays an important role. For example, the fraction of hypoxic cells in xenografts is much the same as in mouse tumors.

Steel and colleagues reviewed the field in 1983 and concluded that xenografts generally maintain the chemotherapeutic response characteristics of the class of tumors from which they are derived. There is good evidence, too, for individuality of response among xenografts. For example, in studying melanomas, one was responsive clinically, but another was not, and the cell survival curve after therapy with melphalan was twice as steep in the xenograft of the cells from the responsive tumor.

Figure 21.9 summarizes the correlation between growth delay in the xenograft and clinical remission of the donor patient. In the figure, the growth delay in xenografts for maximum-tolerated treatment with the single chemotherapeutic agents that are in common clinical use against the disease is plotted against the clinic complete response rate for that category of tumor. The correlation between these parameters is good. Testicular tumors are the most responsive in xenografts or in the clinic; small-cell lung cancer and breast tumors occupy an intermediate position; and the other three tumor types are unresponsive, either clinically or experimentally. This consistency of agreement between patient and xenograft responses to chemotherapeutic agents is encouraging for various human tumor types tested. Similarly, studies of radiation response indicate that measurements of growth delay in xenografts rank tumors in the same order as clinical responsiveness: Response is greater in testicular teratoma than in pancreatic carcinoma, which is greater than in bladder carcinoma, for example.
However, autochthonous mice also suffer similar limitations regarding the timing of tumor development and variation in the number of primary tumors. An additional limitation of using transgenic mice is that no one transgenic mouse is a good model for human cancer. Even those mouse models that relate well to their human tumor counterparts fail to fully recapitulate the systemic spread in the mouse that is found in humans, a finding that is less of concern in experiments examining the effect of dose on tumor control. A more significant problem for radiation experiments lies in targeting the dose to spare normal tissue effects. This is not so inconsequential a problem and requires more sophisticated technology than the current animal irradiators that are used.

### SPHEROIDS: AN IN VITRO MODEL TUMOR SYSTEM

Mammalian cells in culture may be grown either as a monolayer attached to a glass or plastic surface or in suspension, in which case they are prevented from settling out and attaching to the surface of the culture vessel by continual gentle stirring. Most cells in suspension, or in “spinner culture,” as it is often called, remain as single cells; at each mitosis, the progeny cells separate, and although the cell concentration increases with time, it continues to consist of individual separate cells.

Some cells, however, notably several rodent tumor cell lines such as Chinese hamster V79 lung cells, mouse EMT6 mammary cells, radiation-induced fibrosarcoma (RIF) cells, and rat 9L brain tumor cells, do not behave in this way...
but instead grow as spheroids. At each successive division, the progeny cells stick together, and the result is a large spheric clump of cells that grows bigger with time. A photograph of a large spheroid consisting of about $8 \times 10^4$ cells is shown in Figure 21.10. Five days after the seeding of single cells into suspension culture, the spheroids have a diameter of about 200 μm; by 15 days, the diameter may exceed 800 μm. Oxygen and nutrients must diffuse into the spheroids from the surrounding tissue culture medium. In the center of a spheroid, there is a deficiency of oxygen and nutrients and a buildup of waste products because of diffusion limitations. Eventually, central necrosis appears and the mean cell cycle lengthens. Mature spheroids contain a heterogeneous population of cells resulting from many of the same factors, as in a tumor in vivo.

The spheroid system is simpler, more reproducible, less expensive, and easier to manipulate than animal tumors, and yet the cells can be studied in an environment that includes the complexities of cell-to-cell contact and nutritional stress from diffusion limitations that are characteristic of a growing tumor. Spheroids are irradiated intact and then separated into single cells by the use of trypsin and gentle agitation before being plated out into petri dishes to be assayed for the reproductive integrity of individual cells.

Mature spheroids consist of three populations of cells with varying radiosensitivity. Starting from the outside and working toward the center, they are asynchronous, aerobic cycling cells, aerated noncycling G1-like cells, and noncycling G1-like hypoxic cells. Very large spheroids may contain about 20% hypoxic cells, similar to many animal tumors. By gently trypsinizing the spheroids for varying periods, the spheroid can be peeled like an onion and these three cell populations are separated out. Using more sophisticated methods, such as centrifugal elutriation and flow cytometry, it is possible to separate many more cell subpopulations based on location in the spheroid, cell cycle, or other parameters. Figure 21.11 is a cross section through a large spheroid, showing clearly the development of a central necrotic area, which occurs when the spheroid’s size is such that

**Figure 21.10** Photograph of an 800-μm spheroid containing about $8 \times 10^4$ cells. (Courtesy of Dr. R.M. Sutherland.)
single-cell in vitro culture and tumors in experimental animals. The spheroid system has been applied to several problems in radiobiology and in the study of pharmacologic agents, such as radiosensitizers or chemotherapeutic agents. A major problem in the application of these drugs to human tumors is the presence of resistant cells that are resting or non-cycling, often located away from blood vessels. Drugs are required to diffuse in effective concentration to these cells through layers of growing, actively dividing cells, which may inactivate the drug through their metabolism. The spheroid system mimics many of these tumor characteristics and provides a rapid, useful, and economic method for screening sensitizers and chemotherapeutic agents because it is intermediate in complexity between single-cell in vitro culture and tumors in experimental animals.

**SPHEROIDS OF HUMAN TUMOR CELLS**

Many types of human tumor cells can be cultured as spheroids, with a wide spectrum of morphologic appearances and growth rates. In general, cells from disaggregated surgical specimens form spheroids if cultured in liquid suspension above a nonadhesive surface, which can be a thin layer of agar or agarose gel or the bottom of a culture dish not prepared for cell culture.

Only if the spheroid is formed and grown to a certain size can it be transferred to a spinner culture vessel and grown in the same way as spheroids.
of established rodent cell lines. Human tumors successfully grown as spheroids include thyroid cancer, renal cancer, squamous carcinoma, colon carcinoma, neuroblastoma, human lung cancer, glioma, lymphoid tumors, melanoma, and osteosarcoma. There appears to be no general pattern. One glioma line might form and grow as spheroids; another might not. The same applies to other tumor types. Thus, it seems that the capacity to form and grow as spheroids is not a general property of tumor cells. Many nontumor cells also form spheroids, but only the spheroids of lymphoid origin continue to grow to any size.

Morphologic studies of spheroids of human tumor cells show that they maintain many characteristics of the original tumor specimens taken from the patient and of the cells if grown as a xenograft in nude mice. Radiobiologic studies show that in addition to maintaining histologic characteristics of individual tumors, spheroids of human cells preserve characteristic radiosensitivity because dose-response curves for spheroids are virtually identical to those for cells growing as xenografts in nude mice.

- COMPARISON OF THE VARIOUS MODEL TUMOR SYSTEMS

In all transplantable systems described, the tumor is treated in situ, with all of the realism and complexities of the in vivo milieu, such as cell-to-cell contact and the presence of hypoxic cells, factors that cannot be fully simulated in a petri dish. The tumor cure (TCD$_{50}$) and growth delay systems share the additional advantage that they are left in situ undisturbed after treatment. In the other techniques, the tumor must be removed, minced, and prepared into a single-cell suspension by the use of an enzyme, such as trypsin, before survival is assessed. Although this step does not appear to affect the assessment of the effects of radiation, it can result in artifacts in the case of other agents, such as chemotherapeutic drugs or hyperthermia, in which the cell membrane may be involved in the cellular response. The procedure of breaking up the tumor and partially dissolving the cell membrane with a digestive enzyme may influence results. For this reason, in the testing and evaluation of a new drug, one tumor system involving the determination of growth delay or TCD$_{50}$ is always included. For the same reason, these same systems are very expensive because they require a large number of animals for the amount of information produced. The determination of TCD$_{50}$ is perhaps ideal for producing data relevant to clinical radiotherapy. It is certainly the most expensive; to produce a single TCD$_{50}$ value for one of the lines in Figure 21.4, six to eight groups of up to 10 animals must be kept and observed for weeks. The same information can be obtained in 10 days with one or two mice and six petri dishes using the in vivo/in vitro technique.

The dilution assay technique allows clonogenic cell survival to be assessed over a large range of doses and for tumors that cannot be grown in culture. It, too, is relatively expensive, because a whole group of recipient animals must be used and kept for weeks to obtain the same information obtained from one petri dish. Unquestionably, the most rapid and efficient technique is the in vivo/in vitro technique, which combines the realism of irradiation in vivo with the speed and efficiency of in vitro plating to assess clonogenic survival. The concomitant disadvantage is that any tumor that can be switched from petri dish to animal in alternate passages is so undifferentiated and anaplastic that it bears little resemblance to a spontaneous tumor in the human.

To some extent, the same criticism can be levied at all transplantable tumor systems. They are not only highly quantitative, but they are also very artificial. Having been cultured in vitro for many generations, they tend to be highly undifferentiated, and they grow as encapsulated tumors in a muscle or beneath the skin rather than in the tissue of origin. In addition, some have produced misleading results because they are highly antigenic, which, in general, human tumors are not.

In short, transplantable tumors in laboratory animals are model systems; they must be used with care, and the results must not be overinterpreted. Used with caution, these systems have provided invaluable quantitative data and helped to establish important radiobiologic principles. They also, however, have “led us up the garden path” on several occasions in the past (Fig. 21.12). For all of the reasons listed previously, they differ in important ways from spontaneous human tumors, and for the testing of drugs at the National Cancer Institute, they have been largely replaced by a battery of cells of human origin cultured in vitro.

Xenografts of human tumors so far have been used on a much more limited scale. Because they are grown in the absence of an immune response,
or cycling versus noncycling, can be separated out. Human cell spheroids have only been used on a limited scale, but it is clear that the cells retain many of the characteristics of the tumor from which they were taken. Spheroids are much less expensive than xenografts in immunosuppressed animals and perform much the same function.

SUMMARY OF PERTINENT CONCLUSIONS

- A wide range of tumors of different histologic types can be grown in laboratory animals and propagated by transplantation.
- Transplanted tumor systems can be highly quantitative, but, in general, the more quantitative the system, the more artificial it is, because the tumors are highly undifferentiated and encapsulated.
- The five assays in common use are tumor growth delay measurements, tumor cure (TCD₅₀) assay, tumor cell survival determined by the dilution assay technique, the production of lung colonies, and in vivo treatment followed by in vitro assay.
- In all five assays, the cells can be irradiated in situ with all the realism and complexity of in vivo conditions.
- If tumor cure (TCD₅₀) or growth delay is scored, the tumor is left undisturbed after...
treatment. This avoids artifacts involved in disaggregating the tumor, especially in the study of some chemicals or hyperthermia, in which cell membrane effects are important.

- The dilution assay technique, the lung colony assay, and the in vivo/in vitro assay all measure the cell-surviving fraction; that is, they are clonogenic assays. They require fewer animals and are therefore more efficient than the scoring of tumor cure or growth delay. All three assays require, however, that a single-cell suspension be prepared from the tumor, and this may result in artifacts.

- Transplantable tumors in small laboratory animals have been used to establish many radiobiologic principles, but they are highly artificial and must be used with care. They have “led us up the garden path” on several occasions.

- Many human tumor cells can be grown as xenografts in immune-deficient animals.
- Although the histologic characteristics of the human source tumor are maintained, the stroma is of mouse origin.
- Xenografts of human tumor cells are not much better than mouse tumors for studies in which the vascular supply is important.
- Human tumor cells undergo kinetic changes and selection if transplanted into immune-deficient mice.
- Xenografts generally maintain the chemotherapeutic response characteristics of the class of tumors from which they are derived. There is evidence, too, of individuality of response.

- Spheroids of established rodent cells can be grown in suspension culture (i.e., “spinner culture”). Oxygen and nutrients must diffuse into the spheroid from the surrounding culture medium. Oxygen deficiency and a buildup of waste products result, just as in a tumor.

- Mature spheroids contain a heterogeneous population of cells, much like a tumor, but are more quantitative and more economical to work with.

- Starting from the outside and working toward the center, spheroids consist of asynchronous aerated cells, noncycling G1-like aerated cells, noncycling G1-like hypoxic cells, and necrotic cells.

- Spheroids are intermediate in complexity between monolayer cell cultures in vitro and transplantable tumors in experimental animals.

- Many types of human tumor cells grow as spheroids and maintain many characteristics of the original tumor from the patient or of the same cells grown as xenografts.

- Programmed cell death, or apoptosis, occurs after irradiation in many animal tumors, as well as in human xenografts in nude mice.

- Apoptosis is most important in lymphomas, essentially absent in sarcomas, and intermediate and variable in carcinomas.

- Cells may show signs of dying an apoptotic death by 3 hours after irradiation.

**BIBLIOGRAPHY**


THE CELL CYCLE

The ability of cells to produce exact, accurate copies of themselves is essential to the continuance of life; it is accomplished through highly organized processes, well conserved through evolution. Lack of fidelity in cellular reproduction as manifested by DNA and chromosome alterations is a hallmark of cancer.

The only event in the cell cycle that can be identified with a simple light microscope is the condensation of the chromosomes during mitosis (M); this was observed in the late 19th century. Using autoradiography, Howard and Pelc in the early 1950s divided the cell cycle by showing that DNA was synthesized only during a discrete time interval, which they called the S phase. Between mitosis and the S phase was the “first gap in activity” (G1), and between S phase and the next mitosis was the “second gap in activity” (G2). If the cells stop progressing through the cycle—that is, if they are arrested—they are said to be in G0. The cell cycle was discussed in detail in Chapter 4.

Howard and Pelc also showed that it was in these gaps that radiation affects cell cycle progression, because in their early studies, it was obvious that cells arrest cell cycle progression after low-dose radiation damage not in S or M but in either G1 or G2. It was subsequently recognized that these arrests also were related to the process of malignancy, because primary cells would arrest in both G1 and G2, but tumor cells often would show only the G2 arrest point. Breakthroughs in understanding these events and the nature of the cell cycle itself came with the discovery of the cyclins, the cyclin-dependent kinases, and the cyclin–kinase inhibitors and with the elaboration by Weinert and Hartwell of the concept of cell cycle checkpoints. The current concept of the cell cycle and its regulation is illustrated in Figure 22.1.

CYCLINS AND KINASES

Regulation of the complex processes that occur as a cell passes through the cycle is a result of a series of changes in the activity of intracellular enzymes known as cyclin-dependent kinases (Cdks). The active forms of these enzymes exist in protein complexes with a cell cycle phase-specific protein known as a cyclin. Transitions from one phase to the next in the cycle occur only if the enzymatic activity of a given kinase activates the proteins required for progression.

In mammals, cyclins A through H have been described. Each cyclin protein is synthesized at a discrete phase of the cycle: cyclin D and E in G1, cyclin A in S and G2, and cyclin B in G2 and M. Cyclin levels oscillate with phase of the cycle.

Seven Cdks have been described. Cdk levels are constant throughout the cell cycle, but their activity is regulated by cyclin-dependent activating kinases, the protein level of cyclin regulatory subunits, and association with Cdk inhibitors.

Molecular events in G1 prepare the cell for DNA synthesis. There is a stage in G1, known as the G1 restriction point, after which cells are committed to enter the S phase and no longer respond to growth conditions. Prior to this point,
cells may take several routes: They may progress, differentiate, senesce, or die, depending on external signals. Key players in the G1 restriction point include the protein of the retinoblastoma (Rb) gene, D-type cyclins, and Cdk4 and Cdk6, as well as Cdk inhibitors (Fig. 22.1).

If extracellular signals stimulate a cell to enter the cycle from quiescence, D-type cyclins are stimulated and continue through G1 and form a complex with Cdk4 or Cdk6. The activated cyclin–Cdk4 or cyclin–Cdk6 complex then phosphorylates the Rb protein, which releases it from E2F and its growth-suppressive function. E2F that is released from the Rb protein binds to the promoter of the cyclin E gene, resulting in increased cyclin E messenger ribonucleic acid (mRNA) and protein. There is more cyclin E available to bind Cdk2 and phosphorylate Rb, resulting in a positive feedback loop that is now refractory to mitogenic signals. Although numerous studies have documented the importance of all three D-type cyclins, two E-type cyclins, and Cdk2, Cdk4, and Cdk6, gene knockout studies in mice have indicated that all of these cyclins and their dependent kinases are not essential for normal cell cycle progression. However, the ability of cells to be transformed by oncogenes is dependent on G1 cyclins and their dependent kinases. The current thinking is that untransformed cells require a lower level of G1 cyclins to proliferate and differentiate, whereas transformation requires a quantitatively different level of G1 cyclins. Thus, if increased G1 cyclin activity is needed for transformation, then loss or diminished G1 cyclin activity could act to suppress tumor formation.

Once a cell has committed to entering S, it must begin the incredibly difficult task of accurately copying more than three billion bases of the genome; this feat is completed in a matter of a few hours. DNA polymerases are the enzymes involved in this copying process, which must be completed with high fidelity, aided by repair and misrepair genes that remove and replace mismatched DNA bases. Cyclin A is maximally expressed in S phase and enhances transition of the cell through this phase of the cycle.

After the cell has copied its entire genome, the next important task is to segregate the two copies of the DNA equally into the progeny cells. There is a gap (G2), however, between the end of all detectable DNA synthesis and the beginning of cell division at which the process of condensing and segregating the chromosomes begins. Events during this period are controlled by Cdk activity analogous to that occurring at the G1/S transition, but this time it is a complex of cyclins B and A with Cdk1.

Although the cell is progressing through this complicated process of DNA replication...
and division, it must respond constantly to extracellular signals concerning nutrient status, cell-to-cell contact, and so forth that arrive at the nucleus through one or another signal transduction pathway.

- **CHECKPOINT PATHWAYS**

Events in the cell cycle must take place in a specific order, and it is the function of several checkpoint genes to ensure that the initiation of late events is delayed until earlier events are complete.

There are three principal places in the cell cycle at which checkpoints function:

1. **G1/S checkpoint**
2. **S phase checkpoint**
3. **G2/M checkpoint**

If DNA is damaged, normal cells stop progressing through the cycle and are arrested at one of these checkpoints, depending on their position in the cell cycle at the time at which the damage occurs (Fig. 22.2).

Cells with damaged DNA in G1 avoid replicating that damage by arresting at the G1/S interface, or if they have already passed the restriction point governed by phosphorylation of Rb, they will transiently arrest in the S phase. Avoiding the replication of damaged DNA and allowing time for repair prevent cell death and the accumulation of heritable mutations. The tumor suppressor gene p53 is critical in the pathway that leads to G1 arrest (Fig. 22.2). DNA damage initiates a chain of events: First, ataxia-telangiectasia mutated (ATM) autophosphorylates and releases an active monomer that can directly phosphorylate p53 and murine double minute 2 (Mdm2), the ubiquitin ligase that targets p53 for degradation. In addition, the checkpoint kinases (Chk)—also targets of ATM—can also phosphorylate p53 and Mdm2. Phosphorylation of both p53 and Mdm2 results in increased levels of p53 protein. Activated p53 enhances p21WAF1/CIP1 gene expression, which results in a sustained inhibition of G1 cyclin/Cdkks. G1 cyclin inhibition prevents phosphorylation of Rb and progression from G1 into S. Mutations in p53 (which are present in so many human tumors) clearly compromise this checkpoint function. A second more rapid but transient checkpoint is also induced by DNA damage through Chk1 phosphorylation of the Cdc25A phosphatase and the inhibition of cyclin E–Cdk2 and cyclin A–Cdk2 complexes. This later checkpoint works independently of p53 (Fig. 22.2).

Control of the S phase checkpoint is in part mediated by the Cdc25A phosphatase inhibiting Cdk2 activity and the loading of Cdc45 onto chromatin. Failure to load Cdc45 onto chromatin prevents the recruitment of DNA polymerase α and replicon initiation (Fig. 22.2). A second mechanism for S phase arrest is signaled by phosphorylation of Nijmegen breakage syndrome (NBS) by ATM. The importance of the S phase checkpoint is in protecting replication forks from trying to replicate through DNA strand breaks.

The arrest of cells in G2 following DNA damage is observed readily in mammalian cells and was studied by radiation biologists for decades before checkpoints were understood at the molecular level. The arrest occurs after the levels of cyclin A increase in quantity but before cyclin B increases. The function of this checkpoint in normal cells is to prevent cells with damaged chromosomes from attempting the complex process of mitosis; they are arrested in G2 to allow DNA repair to be completed. It follows, therefore, that cells lacking the G2 checkpoint are radiosensitive, because they cannot repair all of their damaged chromosomes before entering mitosis. At the molecular level, multiple kinase signaling pathways have been implicated in regulating this checkpoint (Fig. 22.2). For example, ATM and Chk target the Cdc25C phosphatase and prevent cyclin B/A–Cdk1 activation. In addition, other regulatory proteins have been implicated in G2 arrest, such as polo-like kinases, BrcA1, and p53bp1. It appears that the G2 checkpoint is the most regulated of all checkpoints and probably the most important in preventing the inappropriate entry of damaged cells into mitosis. Consequently, targeting the inhibition of key components of the G2 checkpoint could increase radiosensitization.

The hallmark of cancer is a lack of the ability to respond to signals that normally would cause the cell to stop progressing through the cycle and dividing. Checkpoint proteins provide an important mechanism by which a cell can temporarily halt its transit through the cell cycle and attempt to restore chromosome integrity.
FIGURE 22.2 Diagram depicting the critical role of ATM and its downstream effectors in regulating the cell cycle. In response to DNA damage, ATM becomes activated and phosphorylates different key targets that will inhibit cell cycle progression in different phases of the cell cycle. In G1 phase, ATM phosphorylates p53 and Mdm2, resulting in increased p53 transcriptional activation of the inhibitor p21, which inhibits the cyclin D/E-Cdk4/6 complex. In late G1/early S phase, ATM phosphorylates Chk2 that in turn phosphorylates Cdc25A, a phosphatase that inhibits cyclin E/A-Cdk2 activity. In S phase, ATM phosphorylates NBS and SMC proteins, and transiently inhibits DNA synthesis. In G2 phase, ATM phosphorylates Chk1 that in turn phosphorylates Cdc25C, a phosphatase that inhibits cyclin B-Cdk1 activity. This diagram is included to reinforce the importance of ATM in inhibiting cell cycle progression in response to DNA damage. These interactions are dynamic and simplified and are, in reality, a great deal more complex with many additional accessory proteins.
Given the assumption that all the cells are dividing with the same cell cycle, then

$$LI = \lambda T_S/T_C$$

where $T_S$ is the duration of the DNA synthetic period and $T_C$ is the total cell cycle time.

In practice, these two quantities—the mitotic index and the labeling index—can be determined from a single specimen by counting the proportion of cells in mitosis and the proportion of cells that are labeled. This is a very important consideration in human studies in which it is usually not practical to obtain a large number of serial specimens of tumor or normal tissue material. Although these measurements yield ratios of the duration of mitosis and DNA synthesis as fractions of the total cell cycle, they do not give the absolute duration of any part of the cycle.

**THE PERCENT-LABELED MITOSES TECHNIQUE**

A complete analysis of the cell cycle to obtain the length of each phase is only possible by labeling a cohort of cells in one phase of the cycle and observing the progress of this labeled cohort through a "window" in some other readily observable phase of the cycle. In practice, the easiest phase to label is S and the easiest to observe is M.

As stated previously, the labeling can be achieved by using either tritiated thymidine, identifiable by autoradiography, or bromodeoxyuridine, identifiable by a specific stain or antibody. The basis of the technique, therefore, is to feed the population of cells a label that is taken up in S and then to observe the appearance of that label in mitotic cells as they move around the cycle from S to M. To avoid confusion, the technique involving tritiated thymidine is described in detail, partly because it is the original and classic technique and partly because pictures of autoradiographs show up well in black and white. The technique works equally well if bromodeoxyuridine is used. Bromodeoxyuridine-containing DNA can be stained and shows up well in color under a microscope, but does not reproduce well in black and white.

The percent-labeled mitoses technique is laborious and time-consuming and requires a large number of serial samples. It is readily applicable in vitro, for which it is not difficult to obtain a large...
must be removed, fixed, and stained and an autoradiograph is then prepared. This is continued for a total time longer than the cell cycle of the population under study. For each sample, the percentage of mitotic cells that carry a radioactive label must then be counted; this is the percentage of labeled mitoses. A photomicrograph of a cell preparation is shown in Figure 22.4. This is a particularly laborious process because only 1% or 2% of the cells are in mitosis in any case, and only a fraction of these will be labeled.

The basis for this type of experiment, if applied to an idealized population of cells that all have identical cell cycles, is illustrated in Figure 22.5, a plot of the percentage of labeled mitoses as a function of time. The cells that are in S while the radioactive thymidine is available take up the label. After the label is removed, cells progress through their cell cycles. At regular intervals, usually of 1 hour, a specimen of the cell population must be removed, fixed, and stained and an autoradiograph is then prepared. This is continued for a total time longer than the cell cycle of the population under study. For each sample, the percentage of mitotic cells that carry a radioactive label must then be counted; this is the percentage of labeled mitoses. A photomicrograph of a cell preparation is shown in Figure 22.4. This is a particularly laborious process because only 1% or 2% of the cells are in mitosis in any case, and only a fraction of these will be labeled.

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All of the parameters of the cell cycle may be calculated from Figure 22.5. The time interval before the appearance of the first labeled mitosis, the length $ab$, is in fact the length of G2 or TG2. The time it takes for the percent-labeled mitoses curve to rise from 0% to 100% ($bc$) corresponds to the time necessary for the leading edge of the labeled cohort of cells to proceed through mitosis and is therefore equal to the length of mitosis, $T_M$. The duration of DNA synthesis ($T_S$) is the time taken for the cohort of labeled cells to pass the beginning of mitosis ($bd$). Likewise, it is the time required for the labeled cohort to pass the end of mitosis ($ce$). In practice, $T_S$ usually is taken to be the width of the curve at the 50% level, as marked in Figure 22.5.

The total cycle ($T_C$) is the distance between corresponding points on the first and second wave ($bf$, $cg$, $db$, or $ej$) or the distance between the centers of the two peaks as marked on the figure. The remaining quantity, $T_{G1}$, usually is

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FIGURE 22.5 Percent-labeled mitoses curve for an idealized cell population in which all of the cells have identical mitotic cycle times. The cell population is flash-labeled with tritiated thymidine, which labels all cells in S. The proportion of labeled mitotic cells is counted as a function of time after labeling. The circles at the top of the figure indicate the position of the labeled cohort of cells as it progresses through the cycle. The length of the various phases (e.g., $T_{G1}$, $T_M$) of the cycle ($T_C$) may be determined as indicated.
calculated by subtracting the sum of all the other phases of the cycle from the total cell cycle, or

$$T_{G_i} = T_C - (T_S + T_{G_i} + T_M)$$

Experimental data are never as clear-cut as the idealized picture in Figure 22.5. Points such as $b$ and $e$, at which the curve begins to rise and reaches zero, are poorly defined. A more typical experimental result is illustrated in Figure 22.6. The only points that can be defined with any certainty are the peaks of the curves and the 50% levels, and these may be used to give a rough estimate of the lengths of the various phases of the cycle. The $S$ period ($T_S$) is given approximately by the width of the first peak, from the 50% level on the ascending portion of the wave to the corresponding point on the descending curve. The total cell cycle, $T_C$, is readily obtained as the time between successive peaks. In a separate experiment, the mitotic index may be counted, which is equal to $\lambda T_M/T_{G2}$ because $T_C$ is known, $T_M$ may be calculated. The time from flash labeling to the point at which the curve passes the 50% level in Figure 22.6 is $T_{G_0} + 0.5 T_M$; because $T_M$ already is known, $T_{G_0}$ may be calculated. The remaining quantity, $T_{G_i}$, is deduced by subtraction, because the total cycle time and all other phases are known.

A careful examination of an actual set of experimental data makes it plain that the first wave in the percent-labeled mitoses curve is not symmetric; the downswing is shallower than the upswing. This observation, coupled with the fact that the second peak is much smaller than the first, indicates that the population is made up of cells with a wide range of cycle times. In many instances, particularly if the population of cells involved is an in vivo specimen of a tumor or normal tissue, the spread of cell cycle times is so great that a second peak is barely discernible.

In practice, therefore, the constituent parts of the cycle are not simply read off the percent-labeled mitoses curve. Instead, a complex computer program is used to try various distributions of cell cycle times and to calculate the curve that best fits the experimental data. In this way, an estimate is obtained of the mean cell cycle time and of the range of cell cycle times in the population.

### EXPERIMENTAL MEASUREMENTS OF CELL CYCLE TIMES IN VIVO AND IN VITRO

A vast number of cell cycle measurements have been made. Only a few representative results are reviewed here to highlight the most important points.

Figure 22.7 shows the percent-labeled mitoses data for two transplantable rat tumors with very different growth rates. The tumor represented in the upper panel (Fig. 22.7A) has a gross doubling time of about 20 hours, which can be judged easily from the separation of the first and second waves of labeled mitotic cells.

For the tumor illustrated in the lower panel (Fig. 22.7B), there is no discernible second peak in the percent-labeled mitoses curve because of the large range of cell cycle times among the cells of the population. To obtain an estimate of the average cell cycle in this case, it is necessary to pool information from the percent-labeled mitoses curve and from the knowledge of the labeling index. The width of the first wave of the percent-labeled mitoses curve indicates that the length of the DNA synthetic phase ($T_S$) is about 10 hours.

**FIGURE 22.6** Typical percent-labeled mitoses curve obtained in practice for the cells of a tissue or tumor. It differs from the idealized curve in Figure 22.5 in that the only points that can be identified with precision are the peaks of the curve and the 50% levels. The first peak is not perfectly symmetric, and the second peak is lower than the first because the cells of a population have a range of cell cycle times.
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There is a wide range of cell cycle times, from less than 10 to more than 40 hours, with a modal value of about 19 hours. This range of cycle times explains the damped labeled mitoses curve and the fact that the first peak is not symmetric.

Table 22.1 is a summary of the cell cycle parameters for cell lines in culture and some of the tissues and tumors for which percent-labeled mitoses curves have been shown in this chapter. The top row of Table 22.1 shows the data for Chinese hamster cells in culture. These cells are characterized by a short cell cycle of only 10 hours and a minimal G1 period. The second row of the table gives the comparable figures for HeLa cells. From a comparison of these two in vitro cell lines, a very important point emerges. The cell cycles of the two cell lines differ by a factor of more than 2, nearly all of which results from a difference in the length of G1. The other phases of the cycle are very similar in the two cell lines.

To obtain the average cell cycle time ($T_C$), it is essential in this situation to know the labeling index. For this tumor, the labeling index is about 3.6%. The average cell cycle time ($T_C$) can then be calculated from the following equation:

$$LI = \lambda T_S/T_C$$

Therefore,

$$T_C = (0.693 \times 10)/(3.6/100) = 192.5 \text{ hours}$$

The absence of a second peak is a clue to the fact that there is a wide range of cell cycle times for the cells of this population, so 192.5 hours is very much an average value.

A computer analysis makes it possible to estimate the distribution of cell cycle times in a population. For example, Figure 22.8 shows the percent-labeled mitoses curve for a transplantable mouse tumor (Fig. 22.7B), together with an analysis of cell cycle times based on a mathematical model (Fig. 22.8A). There is a wide range of cell cycle times, from less than 10 to more than 40 hours, with a modal value of about 19 hours. This range of cycle times explains the damped labeled mitoses curve and the fact that the first peak is not symmetric.

Table 22.1 is a summary of the cell cycle parameters for cell lines in culture and some of the tissues and tumors for which percent-labeled mitoses curves have been shown in this chapter. The top row of Table 22.1 shows the data for Chinese hamster cells in culture. These cells are characterized by a short cell cycle of only 10 hours and a minimal G1 period. The second row of the table gives the comparable figures for HeLa cells. From a comparison of these two in vitro cell lines, a very important point emerges. The cell cycles of the two cell lines differ by a factor of more than 2, nearly all of which results from a difference in the length of G1. The other phases of the cycle are very similar in the two cell lines.
Also included in Table 22.1 are data for the cells of the normal cheek pouch epithelium in the hamster and a chemically induced carcinoma in the pouch. These are representative of several studies in which cells from a solid tumor have been compared with their normal tissue counterparts. In general, it is usually found that the malignant cells have the shorter cycle time.

In reviewing the data summarized in Table 22.1, it is at once evident that although the length of the cell cycle varies enormously between populations, particularly in vivo, the lengths of G₂, mitosis, and

<table>
<thead>
<tr>
<th>Authors</th>
<th>Cell or Tissue</th>
<th>$T_C$</th>
<th>$T_S$</th>
<th>$T_M$</th>
<th>$T_{G_2}$</th>
<th>$T_{G_1}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bedford</td>
<td>Hamster cells (in vitro)</td>
<td>10</td>
<td>6</td>
<td>1</td>
<td>1</td>
<td>2</td>
</tr>
<tr>
<td></td>
<td>HeLa cells (in vitro)</td>
<td>23</td>
<td>8</td>
<td>1</td>
<td>3</td>
<td>11</td>
</tr>
<tr>
<td>Steel</td>
<td>Mammary tumors in the rat</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>BICR/M1</td>
<td>19</td>
<td>8</td>
<td>$\sim 1$</td>
<td>2</td>
<td>8</td>
</tr>
<tr>
<td></td>
<td>BICR/A2</td>
<td>63</td>
<td>10</td>
<td>$\sim 1$</td>
<td>2</td>
<td>50</td>
</tr>
<tr>
<td>Quastler and Sherman</td>
<td>Mouse intestinal crypt</td>
<td>18.75</td>
<td>7.5</td>
<td>0.5</td>
<td>0.5–1.0</td>
<td>9.5</td>
</tr>
<tr>
<td>Brown and Berry</td>
<td>Hamster cheek pouch epithelium</td>
<td>120–15</td>
<td>28.6</td>
<td>1.0</td>
<td>1.9</td>
<td>108–140</td>
</tr>
<tr>
<td></td>
<td>Chemically induced carcinoma in pouch</td>
<td>10.7</td>
<td>5.9</td>
<td>0.4</td>
<td>1.6</td>
<td>2.8</td>
</tr>
</tbody>
</table>
S are remarkably constant. The vast bulk of the cell cycle variation is accounted for by differences in the length of the G₁ phase.

- **PULSED FLOW CYTOMETRY**

During the past several decades, classic autoradiography largely has been replaced by pulsed flow cytometry (Fig. 22.9). The conventional techniques of autoradiography give precise, meaningful answers, but they are laborious and slow. Techniques based on flow cytometry provide data that are available within a few days. Detailed cell kinetic data can be obtained by such techniques, including an analysis of the distribution of cells in the various phases of the cycle (Fig. 22.10).

- **THE GROWTH FRACTION**

Central to an appreciation of the pattern of growth of solid tumors is the realization that at any given moment, not all of the tumor cells that are viable and capable of continued growth are actually proceeding through the cell cycle. The population consists of proliferating (P) cells and quiescent (Q) cells. The growth fraction (GF), a term introduced by Mendelsohn, is defined as the ratio of the number of proliferating cells to the total number of cells (P + Q), or

\[ GF = \frac{P}{P + Q} \]

There are various ways to estimate growth fraction. One method consists of injecting tritiated thymidine into an animal with a tumor, and then several cell generations later, preparing an autoradiograph from sections of the tumor. The growth fraction is given by the expression

\[ GF = \frac{\text{fraction of cells labeled}}{\text{fraction of mitoses labeled}} \]

This method assumes that there are two distinct subpopulations, one growing with a uniform cell cycle, the other not growing at all.

Continuous labeling is an alternative way to provide an approximate measure of the proportion of proliferating cells. Tritiated thymidine is infused continuously for a time equal to the cell cycle minus the length of the S phase. The fraction of labeled cells then approximates to the growth fraction.

Table 22.2 is a summary of growth fractions measured for various solid tumors in experimental animals, which frequently fall between 30% and 50%, even though the tumors vary widely in degree of differentiation, arise in different species, and are of varied histologic types. As a tumor outgrows its blood supply, areas of necrosis often develop that are accompanied by hypoxic cells, the proportion of which for many solid tumors is about 15%. This accounts for part, but not all, of the quiescent cell population.

- **CELL LOSS**

The overall growth of a tumor is the result of a balance achieved between cell production from division and various types of cell loss. In most
cases, tumors grow much more slowly than would be predicted from a knowledge of the cycle time of the individual cells and the growth fraction. The difference is a result of cell loss. The extent of the cell loss from a tumor is estimated by comparing the rate of production of new cells with the observed growth rate of the tumor. The discrepancy provides a measure of the rate of cell loss. If $T_{pot}$ is the potential tumor doubling time ($T_{pot} = \lambda T_S/LI$; where $T_S$ is the length of the DNA synthetic period, $LI$ is the labeling index, and $\lambda$ is a correction factor

<table>
<thead>
<tr>
<th>Tumor</th>
<th>Author</th>
<th>Growth Fraction, %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Primary mammary carcinoma in the mouse (G1,H)</td>
<td>Mendelsohn</td>
<td>35–77</td>
</tr>
<tr>
<td>Transplantable sarcoma in the rat (RIB2)</td>
<td>Denekamp</td>
<td>55</td>
</tr>
<tr>
<td>Transplantable sarcoma in the rat (SSO)</td>
<td>Denekamp</td>
<td>47</td>
</tr>
<tr>
<td>Transplantable sarcoma in the rat (SSB1)</td>
<td>Denekamp</td>
<td>39</td>
</tr>
<tr>
<td>Mammary carcinoma in the mouse (C1,H)</td>
<td>Denekamp</td>
<td>30</td>
</tr>
<tr>
<td>Chemistry induced carcinoma in the hamster cheek pouch</td>
<td>Brown</td>
<td>29</td>
</tr>
</tbody>
</table>
between 0.67 and 1), and $T_d$ is the actual tumor doubling time, obtained from simple direct measurements on the diameter of the tumor mass, the cell loss factor ($\phi$) has been defined by Steel to be

$$\phi = 1 - \frac{T_{pot}}{T_d}$$

The cell loss factor represents the ratio of the rate of cell loss to the rate of new cell production. It expresses the loss of growth potential by the tumor. A cell loss factor of 100%, for instance, indicates a steady state of neither growth nor regression.

Cells in tumors can be lost in several ways:

1. Death from inadequate nutrition: As the tumor outgrows its vascular system, rapid cell proliferation near capillaries pushes other cells into regions remote from blood supply, in which there is an inadequate concentration of oxygen and other nutrients. These cells die, giving rise to a progressively enlarging necrotic zone.
2. Apoptosis, or programmed cell death: This form of cell death is manifested by the occurrence of isolated degenerate nuclei remote from regions of overt necrosis.
3. Death from immunologic attack.
4. Metastasis, including all processes by which tumor cells are lost to other parts of the body, such as spread through the bloodstream and lymphatic system.
5. Exfoliation, which would not apply to most model tumors in experimental animals but which could be an important mechanism of cell loss from, for example, carcinoma of the gastrointestinal tract in which the epithelium is renewed at a considerable rate.

There are limited data on the relative importance of these different processes in different tumor types, but it is clear that death from inadequate nutrition—by entry into necrotic areas—is often a major factor. It reflects the latent inability of the vascular system to keep up with the rate of cell production. There is still a great deal to be learned about the occurrence of cell loss from tumors, the mechanisms by which it occurs, and the factors by which it can be controlled. It is clear, however, that any understanding of the growth rate of tumors at the cellular level must include a consideration of this often dominant factor.

### DETERMINATIONS OF CELL LOSS IN EXPERIMENTAL ANIMAL TUMORS

The cell-loss factor has been estimated in a considerable number of tumors in experimental animals. Some of the results are listed in Table 22.3. Values for the cell-loss factor vary from 0% to more than 90%. In reviewing the literature on this subject, Denekamp pointed out that sarcomas tended to have low cell-loss factors, but carcinomas tended to have high cell-loss factors. All the sarcomas investigated had cell-loss factors less than 55%; most carcinomas had cell-loss factors in excess of 70%. Therefore, cell loss appears to be a dominant factor in the growth of carcinomas and of considerably less importance for sarcomas. This pattern correlates with the importance of apoptosis as a mode of cell death. Apoptosis is quite common in carcinomas and rare in sarcomas. If this is found to be a general phenomenon, it might be attributed to the origin of carcinomas from continuously renewing epithelial tissues in which the cell-loss factor is 100%.

This difference between sarcomas and carcinomas may also account for their differing responses to radiation. In carcinomas in which the production of new cells is temporarily stopped or reduced by a dose of radiation, cells continue to be removed from the tumor because of the high

<table>
<thead>
<tr>
<th>Tumor</th>
<th>Author</th>
<th>$\phi$, %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mouse sarcoma</td>
<td>Frindel</td>
<td>0</td>
</tr>
<tr>
<td>3-day-old tumor</td>
<td></td>
<td></td>
</tr>
<tr>
<td>7-day-old tumor</td>
<td></td>
<td>10</td>
</tr>
<tr>
<td>20-day-old tumor</td>
<td></td>
<td>55</td>
</tr>
<tr>
<td>Rat carcinoma</td>
<td>Steel</td>
<td>9</td>
</tr>
<tr>
<td>Rat sarcoma</td>
<td>Steel</td>
<td>0</td>
</tr>
<tr>
<td>Mouse carcinoma</td>
<td>Mendelsohn</td>
<td>69</td>
</tr>
<tr>
<td>Hamster carcinoma</td>
<td>Brown</td>
<td>75</td>
</tr>
<tr>
<td>Rat sarcoma</td>
<td>Hermens</td>
<td>26</td>
</tr>
<tr>
<td>Hamster carcinoma</td>
<td>Reiskin</td>
<td>81–93</td>
</tr>
<tr>
<td>Mouse carcinoma</td>
<td>Tannock</td>
<td>70–92</td>
</tr>
</tbody>
</table>
cell-loss factor, and the tumor shrinks. In sarcomas, however, even if a large proportion of the cells are sterilized by a dose of radiation, they do not disappear from the tumor mass as quickly.

It would be simple, then, to explain why two tumors, one a carcinoma and one a sarcoma, containing the same number of cells and exposed to the same radiation dose, would appear to behave quite differently. The carcinoma might shrink dramatically soon after the radiation dose, whereas the sarcoma would not appear as affected by the radiation. In the long term, the “cure” rates of both tumors may well be identical, but in the short term, the carcinoma would be said to have “responded” to the radiation, whereas the sarcoma might be said to be unresponsive or resistant to radiation.

**GROWTH KINETICS OF HUMAN TUMORS**

The first quantitative study of the growth rate of human tumors was done by Collins and his colleagues in 1956. They observed the growth of pulmonary metastases from serial chest radiographs. It is now possible to gather information from the literature on the doubling time of more than 1,000 human tumors. Most of the data were obtained either by measurements from radiographs or by direct measurements of skin tumors or metastases in soft tissue. The doubling time of human tumors varies widely from patient to patient and is, on the average, very long; Tubiana and Malaise have estimated that the median value is about 2 months.

Tumors of the same histologic type arising in different patients differ widely in growth rate. By contrast, metastases arising in the same patient tend to have similar rates of growth. The latter observation is the basis for using patients with multiple skin or pulmonary metastases to test and compare new treatment modalities, such as high-linear energy transfer radiations or hyperthermia. There is certainly a correlation between histologic type and growth rate. Tubiana and Malaise have collected values for the doubling time in 389 patients with pulmonary metastases, classified into five histologic categories. They can be arranged in order of doubling time as follows: embryonic tumors, 27 days; malignant lymphomas, 29 days; mesenchymal sarcomas, 41 days; squamous cell carcinomas, 58 days; and adenocarcinomas, 82 days. In addition, the degree of differentiation seems to be related to the doubling time, with poorly differentiated cancers generally progressing more rapidly.

In addition to growth rate measurements, studies of cell population kinetics also have been performed on a limited number of human tumors. Studies of this kind raise practical and ethical problems. The ethical problems stem from the fact that *in vivo* experiments require an injection of tritiated thymidine or bromodeoxyuridine, which limits such studies to patients who have short life expectancies and in whom the injection of the label does not raise any problems of possible genetic consequences. The practical problems arise because the percent-labeled mitoses technique, which is the most satisfactory way to obtain the duration of the various phases of the cell cycle, requires a large number of sequential samples to be taken for several days after the injection of the label.

In mice or rats, the multiple samples are obtained from a large number of identical animals bearing transplanted tumors of the same size and type by sacrificing one or more animals each time. In humans, each spontaneous tumor is unique, so the multiple samples must be obtained by repeated biopsies at frequent intervals from the same tumor. This heroic procedure is uncomfortable and inconvenient for the patient and is only practical with large superficial skin tumors, which may not be truly representative of human tumors in general. Nevertheless, a surprisingly large number of human tumors have been studied in this way. The percent-labeled mitoses curves obtained are similar to those for laboratory animals, although the second wave of labeled mitoses is rarely distinct and is usually altogether absent.

Tubiana and Malaise surveyed the field and reported 40 cases in which the cell cycle of solid tumors in humans had been evaluated with the percent-labeled mitoses technique. The cell cycles observed were between 15 and 125 hours in 90% of the cases, with a modal value of 48 hours (Table 22.4). The duration of S ($T_S$) was less variable than the total cell cycle, with 90% of the values falling between 9.5 and 24 hours and a modal value of about 16 hours. As a first approximation, it can be assumed that $T_S$ has a duration of about 16 hours and that the mean duration of the cell cycle is about three times the duration of $T_S$. 


cells should be labeled. This method of continuous labeling can be performed only with a small number of patients who are in no way representative. An alternative procedure is to estimate the growth fraction by assuming that the proportion of cells in cycle is about equal to three times the labeling index, an assumption based on the notion that the cell cycle is three times the length of the S phase. The growth fraction calculated in this way correlates well with the tumor doubling time: It is 0.9 in malignant lymphomas and embryonic tumors and less than 0.06 in adenocarcinomas. The relation between the growth rate and growth fraction appears to be much closer in human tumors than in animal tumors.

Of the various parameters that characterize tumor kinetics, the cell-loss factor is, in general, the most difficult to evaluate. The cell-loss factor for human tumors generally has been calculated by comparing the observed tumor volume-doubling time with the potential doubling time, which is the time required for the population of cells to double, assuming that all the cells produced are retained in the tumor. Tubiana and Malaise calculated a mean value of the cell-loss factor for five histologic groups of human tumors, assuming the duration of S to be 16 hours. Their results suggest that, in general, the mean cell-loss factor exceeds 50%. Furthermore, it appeared to be higher if the tumor was growing quickly and if its growth fraction was high. In humans, the smallest cell-loss factors seem to be associated with those histologic types of tumors that have the slowest rate of growth. The cell-loss factor, therefore, tends to reduce the spread of growth rates that results from the differences in growth fraction of the various types of tumors.

Steel has estimated independently the extent of cell loss in human tumors by comparing the potential doubling time with observed tumor growth rates. The relevant data on the volume doubling time for six groups of human tumors are given in Table 22.5. They consist mostly of measurements of primary and secondary tumors of the lung. There are differences between individual series, which indeed may reflect significant differences in the growth rates of the various types of tumors, but if the results are all pooled, they yield an average median doubling time of 66 days, with 80% of the values falling in the range between 18 and 200 days.

### Table 22.4

<table>
<thead>
<tr>
<th>Authors</th>
<th>Tc, h</th>
</tr>
</thead>
<tbody>
<tr>
<td>Frindel et al. (1968)</td>
<td>97, 51.5, 27.5, 48, 49.8</td>
</tr>
<tr>
<td>Bennington (1969)</td>
<td>15.5, 14.9</td>
</tr>
<tr>
<td>Young and de Vita (1970)</td>
<td>42, 82, 74</td>
</tr>
<tr>
<td>Shirakawa et al. (1970)</td>
<td>120, 144</td>
</tr>
<tr>
<td>Weinstein and Frost (1970)</td>
<td>217</td>
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<tr>
<td>Terz et al. (1971)</td>
<td>44.5, 31, 14, 25.5, 26</td>
</tr>
<tr>
<td>Peckham and Steel (1973)</td>
<td>59</td>
</tr>
<tr>
<td>Estevez et al. (1972)</td>
<td>37, 30, 48, 30, 38, 96, 48</td>
</tr>
<tr>
<td>Terz and Curutchet (1974)</td>
<td>18, 19, 19.2, 120</td>
</tr>
<tr>
<td>Malaise et al. (unpublished data)</td>
<td>24, 33, 48, 42</td>
</tr>
<tr>
<td>Muggia et al. (1972)</td>
<td>64</td>
</tr>
<tr>
<td>Bresciani et al. (1974)</td>
<td>82, 50, 67, 53, 58</td>
</tr>
</tbody>
</table>

* Measured by the mean grain count halving time.

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Taking the median values for the labeling index, doubling time, and S phase as suggested by Steel, the median cell-loss factor in all human tumors studied is 77%. It thus would appear that for human tumors, cell loss is generally the most important factor determining the pattern of tumor growth.

The high rate of cell loss in human tumors largely accounts for the great disparity between the cell cycle time of the individual dividing cells and the overall doubling time of the tumor. Although the tumor doubling time is characteristically 40 to 100 days, the cell cycle time is relatively short, 1 to 5 days. This has important implications, which often are overlooked, in the use of cycle-specific chemotherapeutic agents or radiosensitizing drugs for which it is the cell cycle time that is relevant.

Because Bergonié and Tribondeau established a relation between the rate of cell proliferation and the response to irradiation in normal tissues, it might be supposed that this would be the same for tumors. It is of interest to note that the histologic groups of human tumors that have the most rapid mean growth rates and the highest growth fractions and cell turnover rates are indeed those that are the most radiosensitive. There is also a correlation between, on the one hand, the growth rate and the labeling index or the cell loss and, on the other hand, the reaction to chemotherapy. This is not surprising, because most drugs act essentially on cells in S phase. It is remarkable, however, that the only human tumors in which it is possible to achieve cures by chemotherapy are the histologic types with high labeling indexes. Furthermore, a high level of cell loss appears to favor the response to chemotherapy, and in humans, this occurs especially in tumors with high labeling indexes.

**SUMMARY OF PERTINENT CONCLUSIONS**

- The division of the cell cycle into its constituent phases, M, G1, S, and G2 was accomplished in the 1950s.
- The arrest of cells at various positions in the cycle by the action of “checkpoint genes” is an important response to DNA damage. The two principal checkpoints are the G1/S and the G2/M boundary, but there is also a checkpoint in S. G2/M is the most important checkpoint following radiation damage; cells pause at G2/M to repair radiation-induced damage before attempting the complex process of mitosis.
- Progression through the cell cycle is governed by protein kinases, activated by cyclins. Each cyclin protein is synthesized at a discrete phase of the cycle: cyclin D and E in G1, cyclin A in S and G2, and cyclin B in G2 and M. Transitions in the cycle occur only if a given kinase activates the proteins required for progression.
- Most of the difference in cell cycle between fast- and slow-growing cells is a result of differences in G1, which varies from less than 1 hour to more than a week.

<table>
<thead>
<tr>
<th>Table 22.5 Volume Doubling Times of Human Tumors</th>
</tr>
</thead>
<tbody>
<tr>
<td>Authors</td>
</tr>
<tr>
<td>------------------</td>
</tr>
<tr>
<td>Breuer</td>
</tr>
<tr>
<td>Collins et al.</td>
</tr>
<tr>
<td>Collins</td>
</tr>
<tr>
<td>Garland</td>
</tr>
<tr>
<td>Schwartz</td>
</tr>
<tr>
<td>Spratt</td>
</tr>
</tbody>
</table>

The mitotic index (MI) is the fraction of cells in mitosis

\[ MI = \frac{\lambda T_M}{T_C} \]

The labeling index (LI) is the fraction of cells that take up tritiated thymidine (i.e., the fraction of cells in S)

\[ LI = \frac{\lambda T_S}{T_C} \]

The percent-labeled mitoses technique allows an estimate to be made of the lengths of the constituent phases of the cell cycle. The basis of the technique is to label cells with tritiated thymidine or bromodeoxyuridine in S phase and time their arrival in mitosis.

Flow cytometry allows a rapid analysis of the distribution of cells in the cycle. Cells are stained with a DNA-specific dye and sorted based on DNA content.

The bromodeoxyuridine–DNA assay in flow cytometry allows cells to be stained simultaneously with two dyes that fluoresce at different wavelengths: One binds in proportion to DNA content to indicate the phase of the cell cycle, and the other binds in proportion to bromodeoxyuridine incorporation to show if cells are synthesizing DNA.

The growth fraction is the fraction of cells in active cell cycle (i.e., the fraction of proliferative cells).

In animal tumors, the growth fraction frequently ranges from 30% to 50%.

The cell-loss factor (ϕ) is the fraction of cells produced by cell division lost from the tumor.

In animal tumors, ϕ varies from 0% to more than 90%, tending to be small in small tumors and to increase with tumor size.

The cell-loss factor ϕ tends to be large for carcinomas and small for sarcomas.

The observed volume doubling time of a tumor is the gross time for it to double overall in size as measured, for example, in serial radiographs.

Tumors grow much more slowly than would be predicted from the cycle time of individual cells. One reason is the growth fraction, but the principal reason is the cell-loss factor.

The overall pattern of tumor growth may be summarized as follows. A minority of cells (the growth fraction) are proliferating rapidly; most are quiescent. Most new cells produced by mitosis are lost from the tumor.

In general, the cell cycle time of malignant cells is appreciably shorter than that of their normal tissue counterparts.

In general, irradiation causes an elongation of the cell cycle time in tumor cells and a shortening of the cell cycle in normal tissues.

In 90% of human tumors, the cell cycle time has a modal value of 48 hours (a range of 15 to 125 hours).

In human tumors, \( T_S \) has a modal value of about 16 hours (a range of 9.5 to 24 hours).

As a first approximation, the mean duration of the cell cycle in human tumors is about three times the duration of the S phase.

Growth fraction is more variable in human tumors than in rodent tumors and correlates better with gross volume doubling time.

Cell-loss factor for human tumors has been estimated by Tubiana and Malaise to have an average value for a range of tumors in excess of 50%. Steel's estimate for a median value for all human tumors studied is 77%.

**BIBLIOGRAPHY**


THE INTRODUCTION OF FRACTIONATION

The multifraction regimens commonly used in conventional radiation therapy are a consequence largely of radiobiologic experiments performed in France in the 1920s and in the 1930s. It was found that a ram could not be sterilized by exposing its testes to a single dose of radiation without extensive skin damage to the scrotum, whereas if the radiation was spread out over a period of weeks in a series of daily fractions, sterilization was possible without producing unacceptable skin damage (Fig. 23.1). It was postulated that the testes were a model of a growing tumor, whereas the skin of the scrotum represented a dose-limiting normal tissue. The reasoning may be flawed, but the conclusion proved to be valid: Fractionation of the radiation dose produces, in most cases, better tumor control for a given level of normal tissue toxicity than a single large dose.

THE FOUR Rs OF RADIOBIOLOGY

Now, more than 80 years later, we can account for the efficacy of fractionation based on more relevant radiobiologic experiments. We can appeal to the “four Rs” of radiobiology to be as follows:

- Repair of sublethal damage
- Reassortment of cells within the cell cycle
- Repopulation
- Reoxygenation

The basis of fractionation in radiotherapy can be understood in simple terms. Dividing a dose into several fractions spares normal tissues because of repair of sublethal damage between dose fractions and repopulation of cells if the overall time is sufficiently long. At the same time, dividing a dose into several fractions increases damage to the tumor because of reoxygenation and reassortment of cells into radiosensitive phases of the cycle between dose fractions.
three or five fractions per week, overall time in this plot contains, by implication, the number of fractions as well. It commonly was found in these plots that the slope of the isoeffect curve for skin was about 0.33; that is, the total dose for an isoeffect was proportional to $T^{0.33}$.

The most important contribution in this area, made by Ellis and his colleagues with the introduction of the nominal standard dose (NSD) system, was the recognition of the importance of separating overall time from the number of fractions. According to this hypothesis, total dose for the tolerance of connective tissue is related to the number of fractions ($N$) and the overall time ($T$) by the relation

$$\text{Total dose} = (\text{NSD})T^{0.11}N^{0.24}$$

The NSD system has been discussed extensively. It does enable predictions to be made of equivalent dose regimens, provided that the range of time and number of fractions are not too great and do not exceed the range over which the data are available. For example, in changing a treatment protocol from five to four fractions per week, the formula can be used to calculate the size of dose fractions needed to result in the same normal tissue tolerance with the two different protocols. Of course, because the system is based ultimately on skin reaction data, it does not, in any way, predict late effects.

An obvious weakness of the NSD system is that time is allowed for in terms of a single power function, in which the nominal single dose is proportional to $T^{0.11}$. In fact, biologic experiments with small animals have shown that this relationship is far from accurate. Proliferation does not affect the total dose required to produce a given biologic reaction at all until some time after the start of irradiation but, then, the dependence in time is much greater than al-

The advantages of prolongation of treatment are to spare early reactions and to allow adequate reoxygenation in tumors. Excessive prolongation, however, allows surviving tumor cells to proliferate during treatment.

### THE STRANDQUIST PLOT AND THE ELLIS NOMINAL STANDARD DOSE SYSTEM

Early attempts to understand and account for fractionation gave rise to the well-known Strandquist plot, in which total dose was plotted as a function of the overall treatment time (Fig. 23.2). Because all treatments were given as...
would not be seen until a longer period into a fractionation regimen because of the slower response of the human skin and the longer cell cycle of the individual cells. Figure 23.3 is not meant to be quantitative, but to indicate that the shape of the curve relating extra dose to proliferation is sigmoidal. This illustrates immediately that the method of allowing for overall time in the NSD system is incorrect or, at best, is a very crude approximation.

A further consideration is that all normal tissues are not the same. In particular, there is a clear distinction between tissues that are early responding, such as the skin, mucosa, and intestinal epithelium, and those that are late responding, such as the spinal cord. Figure 23.4 shows the extra dose required to produce a given level of damage for a fractionated protracted regimen in the case of representative tissues from the allowed for by the Ellis formula. For these and other reasons, the NSD system is seldom used nowadays.

PROLIFERATION AS A FACTOR IN NORMAL TISSUES

Experimental evidence indicates that the total dose required to produce a given biologic effect is not a power function of time, as postulated by the Ellis NSD system, but turns out to be more complex. The extra dose required to counter proliferation and result in a given level of skin damage in mice does not increase at all until about 12 days into a fractionated regimen but, then, it increases very rapidly as a function of time. The shape of the curve is roughly sigmoidal (Fig. 23.3). If similar data were available for humans, the effects of proliferation would not be seen until a longer period into a fractionation regimen because of the slower response of the human skin and the longer cell cycle of the individual cells. Figure 23.3 is not meant to be quantitative, but to indicate that the shape of the curve relating extra dose to proliferation is sigmoidal. This illustrates immediately that the method of allowing for overall time in the NSD system is incorrect or, at best, is a very crude approximation.

A further consideration is that all normal tissues are not the same. In particular, there is a clear distinction between tissues that are early responding, such as the skin, mucosa, and intestinal epithelium, and those that are late responding, such as the spinal cord. Figure 23.4 shows the extra dose required to produce a given level of damage for a fractionated protracted regimen in the case of representative tissues from the
early- and late-responding groups. This diagram compares mouse skin, representative of early-responding tissues, and rat spinal cord, representative of late-responding tissues. It is recognized that these may not be ideal examples, but suitable data for more relevant systems are simply not available; comparable quantitative data certainly are not available for humans. The point made by this figure is that the time after the start of a fractionated regimen at which extra dose is required to compensate for cellular proliferation is quite different for late- versus early-responding tissues. The other point made, of course, is that these are data from rodents and that in the case of humans, the time scales (although they are not known with any precision) are likely to be very much longer. In particular, the time at which extra dose is required to compensate for proliferation in late-responding tissues in humans is far beyond the overall time of any normal radiotherapy regimen.

Figure 23.5 is an attempt to convert the experimental laboratory data contained in Figure 23.4 into a general principle that can be applied to clinical practice. Early-responding tissues are triggered to proliferate within a few weeks of the start of a fractionated regimen so that the “extra dose to counter proliferation” increases with time, certainly during conventional radiotherapeutic protocols. By contrast, conventional radiotherapy extending to 6 or 8 weeks is never long enough to allow the triggering of proliferation in late-responding tissues. These considerations lead to the following important axiom:

Prolonging overall time within the normal radiotherapy range has little sparing effect on late reactions, but a large sparing effect on early reactions.

This has far-reaching consequences in radiotherapy. Early reactions, such as reactions of the skin or of the mucosa, can be dealt with easily by the simple expedient of prolonging the overall time. Although such a strategy overcomes the problem of the early reactions, it has no effect whatsoever on the late reactions.

- THE SHAPE OF THE DOSE–RESPONSE RELATIONSHIP FOR EARLY- AND LATE-RESPONDING TISSUES

Clinical and laboratory data suggest that there is a consistent difference between early- and late-responding tissues in their responses to changing fractionation patterns. If fewer and larger dose fractions are given, late reactions are more severe, even though early reactions are matched by an appropriate adjustment in total dose. This dissociation can be interpreted as differences in repair capacity or shoulder shape of the underlying dose-response curves. The dose–response relationship for late-responding tissues is more curved than that for early-responding tissues. In terms of the linear–quadratic relationship between effect and dose, this translates into a larger $a/\beta$ ratio for early effects than for late effects. The difference in the shapes of the dose–response relationships is illustrated in Figure 23.6. The $a/\beta$ ratio is the dose at which cell killing by the linear ($a$) and the quadratic ($\beta$) components are equal. (See Fig. 23.12)

For early effects, $a/\beta$ is large; as a consequence, $a$ dominates at low doses, so that the dose-response curve has a marked initial slope and does not bend until higher doses. The linear and quadratic components of cell killing are not equal until about 10 Gy. For late effects, $a/\beta$ is small, so that the $\beta$ term has an influence at low doses. The dose-response curve bends at lower
Second, clinical trials of hyperfractionation, in which two doses are delivered per day for 6 or 7 weeks, appear to result in greatly reduced late effects if the total dose is titrated to produce equal or possibly slightly more severe acute effects. Tumor control is the same or slightly improved. This important clinical observation has been made in several centers and, again, is compatible with the same difference in shape of dose-response curves between early- and late-responding tissues. Late-responding tissues are more sensitive to changes in fractionation patterns than are early-responding tissues.

Third, in experiments with small laboratory animals, the isoeffect curves (i.e., dose vs. number of fractions to produce an equal biologic effect) are steeper for a range of late effects than for various acute effects. The data are shown in Figure 23.7, in which early effects are represented by, for example, skin desquamation or jejunal crypt colonies, and late effects are represented by, for example, lung or spinal cord injury.

Table 23.1 is a summary of the values of \( \alpha/\beta \) for several early- and late-responding tissues. The important result is that for early-responding tissues, \( \alpha/\beta \) (i.e., the dose at which single- and multiple-event cell killing is about equal) occurs at the dose of about 10 Gy. By contrast, \( \alpha/\beta \) for late-responding tissues is about 2 Gy. The values of \( \alpha/\beta \) shown in Table 23.1 come from experiments in which the reciprocal of the total dose is plotted against the quadratic relationship in biologic systems in which it is possible to observe equal effects from various fractionation regimens, even though single cell dose-survival curves cannot be generated. (See Fig. 19.28)

The parameters derived from curves reconstructed from multifraction experiments are specifically relevant to the end point measured in each experiment, whether it is a proportion of clonogenic cells or a stated reduction in organ function. The dose-response curve constructed from multifraction experimental data by making simple assumptions is a functional dose-response curve, deduced from data in which repair after each fractional dose is basically the quantity being measured. It is just such functional dose-response curves that are required to elucidate the relationship between tolerance dose in radiotherapy and size of dose per fraction, with overall time considered separately.
A possibile explanation for the difference in shape of dose–response relationships for early- and late-responding tissues

The radiosensitivity of a population of cells varies with the distribution of cells through the cycle. In general, cells are most resistant in late S phase; slowly growing cells with a long cycle, however, may have a second resistant phase in the early G1 phase, which may be termed G0 if the cells are out of cycle. Thus, two quite different cell populations may be radioresistant.

1. A population proliferating so fast that S phase occupies a major portion of the cycle.
2. A population proliferating so slowly that many cells are in early G1 or not proliferating at all, so that many resting cells are in G0.
It is thought that many late-responding normal tissues are resistant, owing to the presence of many resting cells. This type of resistance applies particularly to small doses per fraction and disappears at larger doses per fraction.

If resistance results from the presence of many cells in S phase in a rapidly proliferating population, redistribution occurs through all the phases of the cell cycle, which can be considered as a “self-sensitizing” activity. The fast proliferation itself is a form of resistance because the new cells produced by division offset those killed by the dose fractions. This applies to acutely responding tissues and also to tumors. Proliferation occurring during a protracted, fractionated regimen helps to spare normal tissues but, of course, is a potential danger as far as the tumor is concerned. This is discussed later in this chapter.

**FRACTION SIZE AND OVERALL TREATMENT TIME: INFLUENCE ON EARLY- AND LATE-RESPONDING TISSUES**

The difference in shape of the dose–response relationship for early- and late-responding tissues leads to an important axiom:

Fraction size is the dominant factor in determining late effects; overall treatment time has little influence. By contrast, fraction size and overall treatment time both determine the response of acutely responding tissues.

It is remarkable that neither clinical radiation oncologists nor experimental radiobiologists came to recognize this simple fact before it was described by Withers in the 1980s.

**ACCELERATED REPOPULATION**

Treatment with any cytotoxic agent, including radiation, can trigger surviving cells (clonogens) in a tumor to divide faster than before. This is known as accelerated repopulation.

Figure 23.8 illustrates this phenomenon in a transplanted rat tumor. Figure 23.8A shows the overall growth curve for this tumor, together with the shrinkage and regrowth that occurs after a single dose of 20 Gy of x-rays. Figure 23.8B shows the proliferation of individual surviving cells (i.e., clonogenic cells), which, after treatment, are dividing with a cycle time of 12 hours. The important point to note is that during the time that the tumor is overtly shrinking and regressing, the surviving clonogens are dividing and increasing in number more rapidly than before treatment.

There is evidence for a similar phenomenon in human tumors. Withers and his colleagues surveyed the literature on radiotherapy for head and neck cancer and estimated the dose to achieve local control in 50% of cases as a function of the overall duration of fractionated treatment. The results are summarized in Figure 23.9. The analysis suggests that clonogen repopulation in this rapidly growing human cancer accelerates at about 28 days after the initiation of radiotherapy in a fractionated regimen. A dose increment of about 0.6 Gy per day is required to compensate for this repopulation. Such a dose increment is consistent with a 4-day clonogen doubling rate, compared with a median of about 60 days for unperturbed growth.

The conclusion to be drawn from this is that radiotherapy, at least for head and neck cancer and probably for other fast growing tumors, should be completed as soon after it has begun, as is practicable. It may be better to delay initiation of treatment than to introduce delays during treatment. If overall treatment time is too long, the effectiveness of later dose fractions is compromised because the surviving clonogens in the tumor have been triggered into rapid repopulation.

The experimental data referred to here all relate to radiotherapy. It might be anticipated, however, that similar considerations would apply to chemotherapy or to a combination of radiotherapy and chemotherapy. There is evidence in some human malignancies that radiotherapy produces poorer results if preceded by a course of chemotherapy. It may be that accelerated repopulation, triggered by the chemotherapy, is the explanation.

**MULTIPLE FRACTIONS PER DAY**

We are now in a position to sum up the pros and cons of fractionation and prolongation of treatment in a much more sophisticated way than would have been possible at the beginning of this chapter. The advantages of prolongation of treatment are to spare early reactions and to allow adequate reoxygenation in tumors. Excessive prolongation, however, has two disadvantages, they are as follows:

1. It can decrease deceptively the acute reactions without sparing late injury, and
2. It allows the surviving tumor cells to proliferate during treatment.
FIGURE 23.8 Accelerated repopulation. Growth curves of a rat rhabdomyosarcoma showing the shrinkage, growth delay, and subsequent recurrence following treatment with a single dose of 20 Gy of x-rays. A: Curve 1: Growth curve of unirradiated control tumors. Curve 2: Growth curve of tumors irradiated at time t = 0, showing tumor shrinkage and recurrence. B: Variation of the fraction of clonogenic cells as a function of time after irradiation, obtained by removing cells from the tumor and assaying for colony formation in vitro. The surviving clonogenic cells are dividing rapidly with a cell cycle of about 12 hours. (Adapted from Hermens AF, Barendsen GW. Changes of cell proliferation characteristics in a rat rhabdomyosarcoma before and after x-irradiation. Eur J Cancer. 1969;5:173–189, with permission.)

There are two separate strategies that use multiple treatments per day: hyperfractionation and accelerated treatment.

The aims and objectives of these strategies are quite different.

**Hyperfractionation**

The basic aim of hyperfractionation is to further separate early and late effects. “Pure” hyperfractionation might be defined as keeping the same total dose as in a conventional regimen in the same overall time but delivering it in twice as many fractions by the expedient of treating twice per day. This would not be satisfactory because the total dose would need to be increased if the dose per fraction is decreased. In practice, then, “impure” hyperfractionation involves an increase in the total dose and sometimes a longer overall time as well as many more fractions delivered twice per day. The intent is to further reduce late effects but achieve the same or better tumor control and the same or slightly increased early effects.

A large controlled clinical trial of hyperfractionation was conducted in the 1990s by the European Cooperative Group (EORTC 22791)
Unequivocal advantage for hyperfractionation in oropharyngeal cancer

Two fractions per day may not be the limit of hyperfractionation: A further sparing of late effects by splitting the dose into more and more fractions of smaller and smaller size would occur as long as the dose fraction is still on the curved portion of the dose-response curve. In order not to prolong overall treatment time too much, these small doses per fraction would necessitate three or even four fractions per day.

Accelerated Treatment

The alternative strategy to hyperfractionation is accelerated treatment. “Pure” accelerated treatment might be defined as the same total dose delivered in half the overall time by the expedient of giving two or more fractions each day. In practice, it is never possible to achieve this because the acute effects become limiting. It is necessary either to interpose a rest period in the treatment of head and neck cancer. A hyperfractionated schedule of 80.5 Gy delivered in 70 fractions (1.15 Gy twice per day) over a period of 7 weeks was compared with a conventional regimen of 70 Gy delivered in 35 fractions of 2 Gy over 7 weeks. Local tumor control at 5 years was increased from 40% with the conventional regimen to 59% with hyperfractionation, and this was reflected in improved survival. There was no increase reported in late effects or complications. It was concluded that hyperfractionation confers an unequivocal advantage in the treatment of oropharyngeal cancer.

The hyperfractionation results of the cooperative trial are summarized as follows:

- Comparing 80.5 Gy in 70 fractions (1.15 Gy twice per day), 7 weeks, with 70 Gy in 35 fractions, 7 weeks
- Local tumor control, at 5 years, increased from 40% to 59%, reflected also in improved survival
- No increase in side effects
- Unequivocal advantage for hyperfractionation in oropharyngeal cancer

Two fractions per day may not be the limit of hyperfractionation: A further sparing of late effects by splitting the dose into more and more fractions of smaller and smaller size would occur as long as the dose fraction is still on the curved portion of the dose-response curve. In order not to prolong overall treatment time too much, these small doses per fraction would necessitate three or even four fractions per day.
the middle of the treatment or to reduce the dose slightly with acute effects as the limiting factor. The intent of this accelerated treatment strategy is to reduce repopulation in rapidly proliferating tumors. There should be little or no change in the late effects because the number of fractions and the dose per fraction are unaltered.

A large prospective randomized clinical trial of accelerated treatment for head and neck cancer, except oropharynx, was carried out in the 1990s by the European Cooperative Group (EORTC 22851). The accelerated treatment consisted of 72 Gy in 45 fractions (three fractions of 1.6 Gy per day) over a total time of 5 weeks, with a rest period of 2 weeks in the middle. The conventional control arm consisted of 35 fractions of 2 Gy, with a total dose of 70 Gy in 7 weeks. The results of this trial showed a 15% increase in locoregional control, which did not translate, however, into a survival advantage. As expected, acute effects were increased significantly, but the observed increase in late effects was decidedly not expected; some involved complications that proved lethal.

This EORTC trial and several other trials testing accelerated treatment show that attempting to keep the total dose as high as 66 to 72 Gy, but shortening the overall time by as much as 2 to 3 weeks from a conventional time of 6 or 7 weeks leads to serious late complications. There are probably two reasons for this: First, the late effects observed are “consequential” late damage, that is, late damage developing out of the very severe acute effects. Second, there is incomplete repair between dose fractions if several fractions per day are given. This is especially likely for protocols involving three fractions per day, in which any unrepaired damage in the first interval accumulates in the second interval in each day and also because intervals between fractions of only 4 hours were used in the early years of the EORTC trial.

This cooperative trial’s results for accelerated treatment are summarized as follows:

- Unexpected increase in late effects, including lethal complications
- Evidence that pure accelerated treatment must be used with extreme caution

### Continuous Hyperfractionated Accelerated Radiation Therapy

A unique and most interesting study of accelerated treatment was carried out in the 1990s at the Mount Vernon Hospital (United Kingdom), in association with the Gray Laboratory. This trial is known as **continuous hyperfractionated accelerated radiation therapy (CHART)**. The protocol consisted of 36 fractions over 12 consecutive days, with three fractions delivered daily with an interfraction interval of 6 hours. The dose per fraction was 1.4 to 1.5 Gy to a total dose of 50 to 54 Gy. By conventional standards, the total dose was very low, but it was delivered in a very short time. The strategy was based on a low dose per fraction to minimize late effects and a very short overall time to minimize tumor proliferation. The results of the CHART protocol showed good local tumor control with severe acute reactions. It was claimed that patients favored the protocol because treatment was concluded quickly. The incidence of late effects in general did not increase and by some measures actually decreased. The notable exception was damage to the spinal cord. Several myelopathies were recorded at total doses of 50 Gy, the probable cause being that an interfraction interval of 6 hours is not sufficient for the full repair of sublethal damage in this tissue.

Characteristics of CHART included the following:

- **Low dose/fractionation**: 36 fractions
- **Short overall time**: 12 consecutive days
- **No gap in treatment**: three fractions per day at 6-hour intervals
- **Three fractions per day**: 1.4 to 1.5 Gy per fraction, 50 to 54 Gy total

CHART is the only one of the “new” fractionation schedules that results in a lower incidence of late complications. CHART’s severe but tolerable acute reactions did not translate into late sequelae, probably because the total dose (50 to 54 Gy) was so low. Effectiveness in tumor control was not lost even at this low dose because the shortening of overall time was extreme,
minimizing tumor cell proliferation. Compliance was also high with CHART because acute reactions did not peak and become uncomfortable until after the end of treatment.

The results of CHART can be summarized as follows:
- Good local tumor control owing to short overall time
- Acute reactions that are brisk, but peak after treatment is completed
- Most late effects acceptable because of small dose per fraction
- Exception: spinal cord, with several myelopathies occurring at 50 Gy because the time between fractions (6 hours) was too short

### Accelerated Hyperfractionated Radiation Therapy while Breathing Carbogen and with the Addition of Nicotinamide

The last experimental protocol that deserves mention is accelerated hyperfractionated radiation therapy while breathing carbogen and with the addition of nicotinamide (ARCON). The strategy was to accelerate treatment to avoid tumor proliferation, hyperfractionate (small doses per fraction) to minimize late effects, and add carbogen breathing to overcome chronic hypoxia and nicotinamide to overcome acute hypoxia. Clinical trials to test this complex but imaginative protocol are under way in Europe. Early results of a trial of ARCON in the Netherlands, involving advanced laryngeal cancer, showed spectacular results compared with historical controls. Results of a prospective randomized trial have yet to be published.

Characteristics of ARCON are summarized as follows:
- Accelerated treatment to overcome proliferation
- Hyperfractionated to spare normal tissues
- Carbogen breathing to overcome chronic hypoxia
- Nicotinamide to overcome acute hypoxia

### The Time Interval between Multiple Daily Fractions

One thing the two strategies of hyperfractionation and accelerated treatment have in common is that both involve multiple fractions per day and, in this context, it is important to ensure that the fractions are separated by a sufficient time interval for the effects of the doses to be independent—that is, for the repair of sublethal damage from the first dose to be complete before the next dose is delivered.

The most pertinent and remarkable evidence comes from twice-a-day trials by the Radiation Therapy Oncology Group (RTOG) that indicate that for a given total dose delivered in a given number of fractions, the incidence of late effects is worse for interfraction intervals less than 4 hours compared with interfraction intervals longer than 6 hours. These data imply that the repair of sublethal damage in late-responding tissues is slow, and so current wisdom dictates an interfraction interval of 6 hours or more if multiple fractions per day are used. Indeed, the CHART pilot study clearly indicated that even 6 hours is not sufficient for the spinal cord, a late-responding tissue in which it appears that sublethal damage repair has a very slow component. This is radiobiology learned from the clinic.

### LESSONS LEARNED FROM FRACTIONATION STUDIES

Several lessons have been learned from the clinical trials that have been performed to test the usefulness of altered fractionation patterns, they are the following:

**First,** hyperfractionation appears to confer an unequivocal benefit in the treatment of head and neck cancer, in terms of both local control and survival, without a significant increase in late sequelae. By contrast, caution is needed in the application of accelerated treatment because the EORTC trials showed an unexpected increase in serious complications, both early and late. Particular caution is necessary if the spinal cord is in the treatment field for twice-a-day treatments because repair of sublethal damage has a slow component in this tissue.

**Second,** late effects depend primarily on total dose and dose per fraction; overall time within the usual therapeutic range has little influence.

**Third,** overall treatment time affects both acute effects and tumor control. Gaps in treatment should be avoided because they lead to an increase in overall time with a concomitant decrease in tumor control.

The importance of overall treatment time is illustrated dramatically by the retrospective
These most interesting data must be viewed with some caution because the three treatment arms were not in a single randomized study but came from different trials over a period of years, the initial purpose of which was to investigate the usefulness of hypoxic cell radiosensitizers. They indicate strongly, nevertheless, that in the case of relatively rapidly growing tumors, such as head and neck cancer, overall treatment time can be a dominant factor in determining outcome (Table 23.2).

One of the major lessons to be learned from fractionation studies is that local control is lost if overall treatment time is prolonged. Since it first was proposed independently in the 1980s by Withers and by Fowler, it is now well documented.

Analysis by Overgaard of three consecutive trials of the Danish cooperative group. All three trials involved a total dose of 66 to 68 Gy. The first trial was of a split course regimen that extended over a total of 9.5 weeks. The 3-year local control was 32%. The second trial involved five fractions per week over a treatment time of 6.5 weeks, with a 3-year local control of 52%. The third trial included six fractions per week, reducing the overall treatment time to 5.5 weeks and improving the 3-year local control to 62%. There was no change in late effects, but as would be expected, the acute reactions became brisker as the overall time was shortened. The protocols and results of these three trials are illustrated in Figure 23.10.

Table 23.2 Importance of Overall Treatment Time

<table>
<thead>
<tr>
<th>Total Dose, Gy</th>
<th>Dose, Gy</th>
<th>Comment</th>
<th>Overall Time, Weeks</th>
<th>3-Year Local Control</th>
</tr>
</thead>
<tbody>
<tr>
<td>66–68</td>
<td>2</td>
<td>Split course</td>
<td>9.5</td>
<td>32%</td>
</tr>
<tr>
<td>66–68</td>
<td>2</td>
<td>5fr/wk</td>
<td>6.5</td>
<td>52%</td>
</tr>
<tr>
<td>66–68</td>
<td>2</td>
<td>6fr/wk</td>
<td>5.5</td>
<td>62%</td>
</tr>
</tbody>
</table>

Note: DAHANCA trials show improved locoregional control with shorter overall time—no increase in late effects.
for head and neck cancer that local control is reduced by about 1.4% (range of 0.4% to 2.5%) for each day that the overall treatment time is prolonged. This does not differ much from the other way of expressing the same problem—namely, that (after the first 4 weeks of a fractionated schedule) the first 0.61 Gy of each day’s dose fraction is required to overcome proliferation from the previous day. An equally solid estimate can be made from data for carcinoma of the cervix, in which a mean of 0.5% local control (range of 0.3% to 1.1%) is lost for each day that the overall time is prolonged.

Such rapid proliferation does not occur for carcinoma of the breast or prostate, so overall treatment time is not so critical. Although the potential tumor doubling time ($T_{pot}$) has not proved useful as a predictive assay for individual patients, mean $T_{pot}$ values for groups of patients are in accord with the importance, or otherwise, of overall treatment time. For example, the $T_{pot}$ for prostate cancer is about 40 days and for non-inflammatory breast cancer is about 14 days; in both cases, overall treatment time has not been found to be critical. This can be contrasted with head and neck cancer in which the mean $T_{pot}$ can be as short as 4 days and, as we have seen, overall treatment time is an important factor governing tumor control.

### Hypofractionation: Renewed Interest

During the 1970s and the 1980s, the trend in radiotherapy, particularly in the United States, has been to increase the number of fractions more and more to reduce the severity of late effects by exploiting the difference in shape of the dose–response relationship between early- and late-responding tissues, as described in detail previously. However, a dramatic recent trend in fractionation is the exact opposite, namely, a renewed interest in dose fractions much larger than 2 Gy for curative radiotherapy. Four lines of research all point in this direction.

First, evidence has accumulated to show that in the special case of prostate cancer, the $\alpha/\beta$ ratio is low, in the region of 2 to 3—more similar to late-responding normal tissues than to tumors. This essentially removes the basic rationale for a multifraction regimen of 35 or more fractions. The implication is that an external-beam regimen consisting of a smaller number of larger dose fractions, or alternatively high dose rate (HDR) brachytherapy delivered in a limited number of fractions, should result in good local tumor control without increased normal tissue damage.

Second, the outcome of several large fractionation trials, mainly involving head and neck tumors, and particularly the CHART trial, have clearly demonstrated the advantage of “acceleration,” that is, shortening the overall treatment time to improve local control. On the other hand, the demonstration that the half-times of late normal tissue repair are long severely limits the strategy of using multiple treatments per day to maintain a large number of treatments in a short overall time. The only alternative is a smaller number of larger dose fractions.

Third, the development of Intensity-modulated radiation therapy (IMRT), tomotherapy, and proton beams results in greatly improved dose distributions, with smaller volumes of normal tissues receiving high doses. This suggests the attractive possibility of increasing the dose per fraction, where the need to spare late-responding normal tissues by fractionation is reduced because of the lower dose to these tissues. The danger here, of course, is that reducing the margin around the tumor may lead to a geographical miss.

Fourth, the development of carbon ion beams has led to trials involving treatment with a small number of large dose fractions, even in some cases with a single fraction. It is not clear at present whether the apparent success of this strategy is caused by the superior dose distribution or the relatively high linear energy transfer (LET) of the radiation.

All of this means that acceleration by hypofractionation—that is, a smaller number of larger dose fractions—emerges as an interesting alternative. Disasters from the past involving unacceptable late normal tissue damage limit enthusiasm for this strategy in many quarters. All of these ideas outlined earlier, need to be approached cautiously, and the perceived benefits proven by careful clinical trials. Treatment regimens involving fewer fractions would clearly be more convenient for patients and result in significant economies in healthcare systems. However, it would be folly to espouse such strategies at the expense of either local tumor control or increased toxicity to normal tissues.
USING THE LINEAR–QUADRATIC CONCEPT TO CALCULATE EFFECTIVE DOSES IN RADIOTHERAPY

It is often useful in practice to have a simple way to compare different fractionation regimens and to assign them a numeric score. For many years, the NSD and TDF systems developed by Ellis and colleagues were used widely. They proved useful for assessing modest changes in fractionation but fell into dispute when extrapolated beyond the data range on which they were based.

The linear-quadratic model is now more widely used and has received greater acceptance. This section was suggested by Dr. Jack Fowler; the format is based on tutorials he has given at the American Society of Therapeutic Radiology and Oncology (ASTRO) and at the European Society of Therapeutic Radiology and Oncology (ESTRO) in the 1990s.

Use of the linear-quadratic model, with appropriate values for the parameters \( \alpha \) and \( \beta \), emphasizes the difference between early- and late-responding tissues and the fact that it is never possible to match two different fractionation regimens to be equivalent for both. Calculations of this sort, although a useful guide for residents in training or for research purposes, are not to be considered a substitute for clinical judgment and experience. They are presented only as examples.

**Figure 23.11** Graph illustrating that if the dose–response relationship is linear-quadratic in form for graded single doses, the effective dose-response curve for a multifraction regimen approaches an exponential function of dose for many doses. The effective dose–response relationship is a straight line from the origin through the point on the single-dose-survival curve corresponding to the daily dose fraction (typically 2 Gy). (Based on the concepts of Fowler, 1989 and Barendsen, 1982.)

**Figure 23.12** Graph illustrating the linear-quadratic nature of the radiation cell survival curve, \( S = e^{-\alpha D - \beta D^2} \), in which \( S \) is the fraction of cells surviving a dose \( D \), \( \alpha \) is the number of logs of cell kill per Gy from the linear portion of the curve, and \( \beta \) is the number of logs of cell kill per Gy\(^2\) from the quadratic component. The linear and quadratic components of cell kill are equal at a dose \( D = \alpha/\beta \).

Figure 23.11 illustrates the familiar way in which biologic effect as a function of dose varies with the number of fractions in which the radiation is delivered—always assuming that the fractions are spaced sufficiently to allow full repair of sublethal radiation damage. For a multifraction regimen, the shoulder of the curve has to be repeated many times and, as a result, the effective dose–response relationship is a straight line from the origin through the point on the single dose-survival curve for that dose fraction (typically 2 Gy). This is discussed in Chapter 3. For the linear portion of the curve, \( \alpha \) represents the loge of the cells killed per Gy. As the curve bends, the quadratic component of cell killing is represented by \( \beta \), which is the loge of the cells killed per gray squared. This is illustrated in Figure 23.12. The ratio \( \alpha/\beta \) has the dimensions of dose and is the dose at which the linear and quadratic components of cell killing are equal.

For a single acute dose \( D \), the biologic effect is given by

\[
E = \alpha D + \beta D^2
\]  
(1)

For \( n \) well-separated fractions of dose \( d \), the biologic effect is given by

\[
E = n(\alpha d + \beta d^2)
\]  
(2)

As suggested by Barendsen, this equation may be rewritten as

\[
E = (\alpha)(nd)\left(1 + \frac{d}{\alpha/\beta}\right)
\]  
(3)
but \( nd \) equals \( D \), the total dose, so

\[
E = \alpha (\text{total dose}) \ (\text{relative effectiveness})
\]

in which the quantity \( 1 + \left[ \frac{d}{\alpha/\beta} \right] \) is called relative effectiveness. If this equation is divided through by \( \alpha \), we have:

\[
\frac{E}{\alpha} = (\text{total dose}) \times (\text{relative effectiveness}) = (nd) \times \left( 1 + \frac{d}{\alpha/\beta} \right)
\]

(4)

The quantity \( \frac{E}{\alpha} \) is the biologically effective dose (BED) and is the quantity by which different fractionation regimens are intercompared. In words, the final equation is

\[
\text{BED} = (\text{total dose}) \times (\text{relative effectiveness}) = \frac{E}{\alpha} = (nd) \times \left( 1 + \frac{d}{\alpha/\beta} \right)
\]

(5)

The quantity BED was first suggested by Barendsen, but was popularized by Fowler.

**Choice of \( \alpha/\beta \)**

For calculating the examples that follow, \( \alpha/\beta \) is assumed to be 3 Gy for late-responding tissues and 10 Gy for early-responding tissues. The reader, of course, may substitute other values that seem more appropriate. It should be noted that parts of schedules can be added—that is, (partial effect)\(_1\) and (partial effect)\(_2\)—as in the concomitant boost. It also should be noted that although it is permissible to compare BEDs for late effects (in Gy\(_3\)) of one schedule with another and permissible to compare BEDs for early effects (in Gy\(_{10}\)) of one schedule with another, it is clearly not permissible or meaningful to compare early with late effects.

**Model Calculations**

1. Conventional treatment: 30 fractions of 2 Gy given one fraction per day, 5 days per week, for an overall treatment time of 6 weeks (this is written as 30F \( \times \) 2 Gy/6 weeks)

   Early effects: \( \frac{E}{\alpha} = (nd) \left( 1 + \frac{d}{\alpha/\beta} \right) \)
   \[
   = 60 \left( 1 + \frac{2}{10} \right) \\
   = 72 \text{ Gy}_{10}
   \]

   Late effects: \( \frac{E}{\alpha} = 60 \left( 1 + \frac{2}{3} \right) \)
   \[
   = 100 \text{ Gy}_{3}
   \]

   **Comment:** The subscripts to the BED are a reminder that this figure is not in gray and specify the particular values of \( \alpha/\beta \) used in the calculation.

2. Hyperfractionation: 70 fractions of 1.15 Gy given twice daily, 6 hours apart, 5 days per week, for an overall treatment time of 7 weeks; that is, 70F \( \times \) 1.15 Gy twice daily/7 weeks

   Early effects: \( \frac{E}{\alpha} = (nd) \left( 1 + \frac{d}{\alpha/\beta} \right) \)
   \[
   = 80.5 \left( 1 + \frac{1.15}{10} \right) \\
   = 89.8 \text{ Gy}_{10}
   \]

   Late effects: \( \frac{E}{\alpha} = 80.5 \left( 1 + \frac{1.15}{3} \right) \)
   \[
   = 111.4 \text{ Gy}_{3}
   \]

   **Comment:** This treatment is much “hotter,” that is, more effective than the conventional 60 Gy for both early and late effects.

3. A one-fraction-a-day control schedule frequently used to compare with hyperfractionation: 35 fractions of 2 Gy given once a day for 5 days a week, for an overall treatment time of 7 weeks; that is, 35F \( \times \) 2 Gy/7 weeks

   Early effects: \( \frac{E}{\alpha} = (nd) \left( 1 + \frac{d}{\alpha/\beta} \right) \)
   \[
   = 70 \left( 1 + \frac{2}{10} \right) \\
   = 84 \text{ Gy}_{10}
   \]

   Late effects: \( \frac{E}{\alpha} = 70 \left( 1 + \frac{2}{3} \right) \)
   \[
   = 116.7 \text{ Gy}_{3}
   \]

   **Comment:** This “control” schedule is not as effective as the hyperfractionation because it is less effective by 7% for early effects (84 vs. 89.8 Gy\(_{10}\)), but hotter for late effects by 5% (116.7 vs. 111.4 Gy\(_3\)).

4. Concomitant boost: 30 fractions of 1.8 Gy given once a day, 5 days a week, and at the same time (concomitant) a boost to a smaller field of 12 fractions of 1.5 Gy once a day; overall treatment time 6 weeks; that is, \([(30F \times 1.8 \text{ Gy}) + (12F \times 1.5 \text{ Gy})]/6 \text{ weeks}\) (this protocol is much favored at the University of Texas M.D. Anderson Hospital and Tumor Institute; by giving the boost
concomitantly, a prolongation of overall time is avoided.

Early effects: \[ \frac{E}{\alpha} = (nd) \left( 1 + \frac{d}{\alpha/\beta} \right) \]
\[ = 54 \left( 1 + \frac{1.8}{10} \right) + 18 \left( 1 + \frac{1.5}{10} \right) \]
\[ = 84.4 \text{ Gy}_{10} \]

Late effects: \[ \frac{E}{\alpha} = 54 \left( 1 + \frac{1.8}{10} \right) + 18 \left( 1 + \frac{1.5}{10} \right) \]
\[ = 113.4 \text{ Gy}_{3} \]

Comment: The Gy_{10} and Gy_{3} values should be compared with the comparable figures for the previous schedules given. The concomitant boost is hotter than the conventional schedule for both early and late effects. Compared with hyperfractionation, however, this concomitant boost is almost the same for late effects but less effective for early effects, including tumor control.

5. CHART: 36 fractions of 1.5 Gy given three fractions a day, 6 hours apart, for 12 consecutive days, with an overall treatment time of 12 days; that is, 36F \times 1.5 Gy (3F/day)/12 days.

Early effects: \[ \frac{E}{\alpha} = (nd) \left( 1 + \frac{d}{\alpha/\beta} \right) \]
\[ = 54 \left( 1 + \frac{1.5}{10} \right) \]
\[ = 62.1 \text{ Gy}_{10} \]

Late effects: \[ \frac{E}{\alpha} = 54 \left( 1 + \frac{1.5}{10} \right) \]
\[ = 81.0 \text{ Gy}_{3} \]

Comment: Direct comparison of CHART with the previous examples in terms of Gy_{10} and Gy_{3} is meaningless because CHART has an overall time of only 12 days compared with 6 or 7 weeks for the other schedules.

Allowance for Tumor Proliferation

Calculations Suggested by Fowler

The correction proposed here for tumor proliferation is a crude approximation and should not be taken too seriously. It assumes, among other things, that the rate of cellular proliferation remains constant throughout the overall treatment time.

The number of clonogens \((N)\) at time \(t\) is related to the initial number of clonogens \((N_0)\) by the expression
\[ N = N_0e^{\lambda t} \] (6)
in which \(\lambda\) is a constant related to the potential doubling time of the tumor, \(T_{pot}\), by the expression
\[ \lambda = \frac{\log_e 2}{T_{pot}} = \frac{0.693}{T_{pot}} \] (7)

The decrease in the number of clonogens because of cell killing by the fractionated radiation regimen is balanced to some extent by cell division of the surviving clonogens. The biologic effect in equation 2 now becomes
\[ E = n(\alpha d + \beta d^2) - 0.693 \frac{t}{T_{pot}} \] (8)

The BED \(E/\alpha\) becomes
\[ \frac{E}{\alpha} = (nd) \left( 1 + \frac{d}{\alpha/\beta} \right) - \frac{0.693}{\alpha} \frac{t}{T_{pot}} \] (9)
or, in words,
\[ \text{BED} = \text{(total dose)} \times \text{(relative effectiveness)} = \frac{-\log_e 2}{\alpha} \text{(no. of cell doublings)} \] (10)

The time \(t\) is the time in days available for proliferation. Rapid proliferation in tumors appears not to start up until about 21 to 28 days after treatment begins in head and neck tumors. Therefore, \(t = T - 21\) or \(t = T - 28\) is a suitable value, where \(T\) is overall time. The start-up time is called \(T_K\) for “kick-off” time, and \(t = T - T_K\).

It is now necessary to assume a value for \(\alpha\), the initial slope of the cell survival curve, as well as for \(T_{pot}\), the potential doubling time of the tumor. A reasonable value for \(\alpha\) is 0.3 ± 0.1/Gy. \(T_{pot}\) may have a value of 2 to 25 days, with a median value of about 5 days.

For typical 6-week (39-day) schedules referred to earlier, proliferation may reduce the BED by
\[ \frac{E}{\alpha} = \frac{0.693}{0.3} \times \frac{(39 - 21)}{5} = 8.3 \text{ Gy}_{10} \]

Note that because we are concerned with tumor proliferation, the reduction in BED is in Gy_{10}; that is, an early-effect \(\alpha/\beta\) value is used. By the same token, proliferation during a 7-week protocol (i.e., 46 days) would decrease the BED by:
\[ \frac{E}{\alpha} = \frac{0.693}{0.3} \times \frac{(46 - 21)}{5} = 11.6 \text{ Gy}_{10} \]

CHART calls for three fractions per day over 12 days and so rapid proliferation has not started in head and neck tumors by the time the treatment is completed in this very short schedule; that is, \(T_K\) is greater than \(T\), so \(t\) must be set at zero.
differential dose per fraction to areas of different risk on the same day, in place of the shrinking field technique. This shortens the overall treatment time and raises the question, What total dose given in 35 fractions over 7 weeks is equivalent to 50 Gy in 25 fractions over 5 weeks, or 60 Gy in 30 fractions over 6 weeks? Given that the ratio for squamous cell carcinoma of the head and neck is usually considered high, the correction factor for overall treatment time becomes more important than the correction factor for fraction size. Some groups, notably Lester Peters and colleagues at the Peter McCallum Cancer Institute in Melbourne, Australia, have taken the pragmatic approach that between 5 and 7 weeks after the start of a fractionated regimen, the dose equivalent of regeneration with protraction of treatment is about 0.5 Gy per day, rounded down to 3 Gy per week. Thus, the equivalent of 50 Gy in 5 weeks is 56 Gy in 7 weeks at 1.6 Gy per fraction, whereas the equivalent of 60 Gy in 6 weeks is 63 Gy in 7 weeks at 1.8 Gy per fraction.

Obviously, there is considerable room for debate concerning the actual numeric values chosen, but in the case of rapidly growing squamous cell tumors of the head and neck, it would be wrong to simply ignore the time factor because it is more important than the correction for fraction size. For other tumor types, the dose equivalent of regeneration is almost certainly different and in most cases, less. For some tumors, such as low to intermediate prostate cancer, it may be

Table 23.3 summarizes the effect of tumor proliferation on the BEDs characteristic of the various treatment regimens discussed earlier.

Based on the assumptions made, hyperfractionation results in the largest BED and, therefore, may be expected to result in the best tumor control, followed closely by the concomitant boost schedule. CHART is a less effective schedule based on a $T_{pot}$ of 5 days. It is necessary to assume a very fast-growing tumor, with a $T_{pot}$ of 3 days or less, for CHART to become the most effective schedule.

It must be emphasized again that calculations of this sort should be used only as a guide for residents in training because they do not, in any way, replace clinical judgment. It is useful, however, to have a yardstick by which new fractionation schemes may be judged.

**Pragmatic Approach of Peters and Colleagues**

Changing dose per fraction often results in a change in overall treatment time. This may be an important issue with the advent of IMRT, especially in the case of head and neck cancer, which may be rapidly growing. For example, with conventional treatment planning, a shrinking field technique may be used, with typically 50 Gy being given to known sites of disease and potential routes of regional spread, followed by a boost of 10 to 20 Gy using a one- or two-phase cone down, with all treatments being given in 2-Gy fractions. However, owing to the complexity of IMRT, a single plan can be utilized with

<table>
<thead>
<tr>
<th>Protocol</th>
<th>$E/\alpha$ Early, i.e., Tumor, Gy$_{10}$</th>
<th>Proliferation Correction, Gy$_{10}$</th>
<th>Corrected for Time, Gy$_{10}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Conventional protocol: 30F × 2 Gy/6 wk (39 d)</td>
<td>72</td>
<td>−8.3</td>
<td>63.7</td>
</tr>
<tr>
<td>Hyperfractionation: 70F × 1.15 Gy/7 wk (46 d)</td>
<td>89.8</td>
<td>−11.6</td>
<td>78.2</td>
</tr>
<tr>
<td>Concomitant boost: (30F × 1.8 Gy) + (12F × 1.5 Gy)/6 wk (39 d)</td>
<td>84.4</td>
<td>−8.3</td>
<td>76.1</td>
</tr>
<tr>
<td>Chart: 36F × 1.5 Gy/12 d</td>
<td>62.1</td>
<td>0</td>
<td>62.1</td>
</tr>
</tbody>
</table>

This correction for time assumes $T_p = 5$ days; $T_k = 21$ days; and $\alpha = 0.3$ In per Gy.
negligible. There is a general lack of data regarding accelerated tumor cell regeneration, regarding both its magnitude and regarding the length of the lag time before it sets in.

**SUMMARY OF PERTINENT CONCLUSIONS:**

- The “four Rs” of Radiobiology are the following:
  - Repair of sublethal damage
  - Reassortment of cells within the cell cycle
  - Repopulation
  - Reoxygenation

- The basis of conventional fractionation may be explained as follows: Dividing a dose into several fractions spares normal tissues because of the repair of sublethal damage between dose fractions and cellular repopulation. At the same time, fractionation increases tumor damage because of reoxygenation and reassortment.

- The Strandquist plot is the relation between total dose and overall treatment time. In this context, “time” includes the number of fractions. On a double log plot, the slope of the line for skin is often close to 0.33.

- The Ellis NSD system made the important contribution of separating the effects of number of fractions and overall time. The time correction was a power function ($T^{0.11}$) that is far from accurate. The system is seldom used now.

- The extra dose required to counteract proliferation in a normal tissue irradiated in a fractionated regimen is a sigmoidal function of time. No extra dose is required until some weeks into a fractionated schedule.

- The delay before an extra dose is required to counteract the effects of proliferation is much longer for late-responding tissues and is beyond the overall time for conventional radiotherapy schedules.

- Prolonging overall time within the normal radiotherapy range has little sparing effect on late reactions but a large sparing effect on early reactions.

- The dose–response relationship for late effects is more curved than for early effects. The ratio $\alpha/\beta$ is about 10 Gy for early effects and about 3 Gy for late-responding tissues. Consequently, late-responding tissues are more sensitive to changes in fractionation pattern.

- Fraction size is the dominant factor in determining late effects; overall treatment time has little influence. By contrast, fraction size and overall treatment time both determine the response of acutely responding tissues.

- Accelerated repopulation refers to the triggering of surviving cells (clonogens) to divide more rapidly as a tumor shrinks after irradiation or treatment with any cytotoxic agent.

- Accelerated repopulation starts in head and neck cancer in humans about 4 weeks after initiation of fractionated radiotherapy. About 0.6 Gy per day is needed to compensate for this repopulation.

- This phenomenon mandates that treatment be completed as soon as practical once it has started; it may be better to delay the start than to introduce interruptions during treatment.

- The basic aim of hyperfractionation is to further separate early and late effects. The overall treatment time remains conventional at 6 to 8 weeks, but because two fractions per day are used, the total number of fractions is 60 to 80. The dose must be increased because the dose per fraction is decreased. Early reactions may be increased slightly, tumor control improved, and late effects greatly reduced.

- In accelerated treatment to reduce repopulation in rapidly proliferating tumors, conventional doses and number of fractions are used, but because two doses per day are given, the overall treatment time is halved. In practice, the dose must be reduced or a rest interval allowed because acute effects become limiting.

- Hyperfractionation has been shown in randomized clinical trials of head and neck cancer to improve local tumor control and survival with no increase in acute or late effects.

- Accelerated treatment: the EORTC trial of 72 Gy in 45 fractions (three fractions per day) over 5 weeks, showed an increase in local tumor control, but no increase in survival. There was an unexpected increase in late effects, some of which were lethal. The late effects were probably “consequential”
late effects, developing out of the severe acute effects. Incomplete repair between fractions also may have been a problem because the time interval between fractions was too short.

- CHART stands for continuous hypofractionated accelerated radiation therapy. The protocol consists of 36 fractions over 12 days (three fractions per day) to a total dose of 50.4 to 54 Gy. Tumor control was maintained because of the extreme acceleration of treatment time; late effects were not increased and even may have decreased because of the low dose and acute effects were severe, but their peak occurred after completion of treatment, so patient compliance was not prejudiced. The three fractions per day, with a time interval of 6 hours between fractions, resulted in problems if the spinal cord was in the field because of the very slow repair of sub-lethal damage in this tissue.

- ARCON involves accelerated treatment to overcome tumor cell proliferation, hypofractionation to spare late-responding normal tissues, carbogen breathing to overcome chronic hypoxia, and nicotinamide to overcome acute hypoxia.

- Overall treatment time is a very important factor for fast-growing tumors. In head and neck cancer, local tumor control is decreased by about 1.4% (range of 0.4% to 2.5%) for each day that the overall treatment time is prolonged. The corresponding figure for carcinoma of the cervix is about 0.5% (range of 0.3% to 1.1%) per day. Such rapid proliferation is not seen in breast or prostate cancer.

- There is renewed interest in hypofractionation—that is, a smaller number of high dose fractions. There are several circumstances where this may be exploited: (a) for prostate cancer for which the $a/b$ ratio is closer to that for late-responding tissues, which removes the benefit of fractionation; (b) for IMRT and proton beams, where the dose distribution is so improved that the volume of normal tissue exposed to high doses is much reduced; and (c) for carbon ion beams, where the dose distribution is improved and, in addition, the radiation has a relatively high LET.

- The linear-quadratic concept may be used to calculate the biologic effectiveness of various radiotherapy protocols involving different numbers of dose fractions. The useful formula is

$$E = \alpha (td) \times \left(1 + \frac{d}{\alpha / \beta}\right)$$

- An approximate allowance can be made for tumor cell proliferation when comparing protocols involving different overall treatment times. There are two approaches considered:

1. Fowler has suggested corrections based on the $T_{pot}$ value for different tumors.
2. Peters and colleagues have suggested a pragmatic approach in the case of fast-growing squamous cell carcinomas of the head and neck, where corrections for overall time may be more important than number of fractions. They assume that between 5 and 7 weeks after the start of a fractionated regimen, the dose equivalent of regeneration with protraction of treatment is about 0.5 Gy per day, rounded down to 3 Gy per week. The correction will be different for other tumors and probably negligible for prostate cancer.

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Of patients presenting at major cancer centers in the Western world, about 1 in 10 present with a second cancer. The following are the three major reasons for this:

1. **Continued lifestyle**—For example, factors associated with the first malignancy may continue to have an effect; smoking that caused a lung cancer may result later in a head and neck cancer, or alcohol that caused esophageal cancer may also result in a cancer of the tongue.

2. **Genetic predisposition**—Some syndromes, such as Li–Fraumeni syndrome, predispose to multiple primary tumors. This involves a very small proportion of the population, but it is incredibly important to the persons affected.

3. **Treatment-related**—Radiotherapy and/or chemotherapy can induce second malignancies. In the case of radiotherapy, most second cancers not only occur within or close to the treatment field, but also occur in remote locations, especially in radiogenic organs such as the lung. This is of particular importance in young children because they are 10 to 15 times as sensitive to radiation-induced malignancies as middle-aged adults are. For a second cancer to be considered as treatment-related, the latency must be appropriate, namely, short for a leukemia and long for a solid tumor.

For any of the previously mentioned reasons, it may be necessary to consider treatment to a tumor located in any area of the body that had previously been irradiated. Decisions regarding the safety of such retreatment are complex. For example, surgery may be compromised if the tissues involved had been previously irradiated to a high dose. The discussion here, however, focuses principally on the safety of reirradiation of an area of normal tissue previously exposed to radiation. Information on this subject is far from complete, but certain general principles emerge from experimental and clinical experience. One thing is clear, namely, that if the radiation tolerance of a given organ or tissue has been exceeded by the initial treatment, to the extent that function is lost or is in the process of being lost, then subsequent retreatment cannot be contemplated with safety. By contrast, if the initial treatment is below the tolerance of the organ or tissue exposed, then some retreatment at a later date is safe, varying very much with the organ or tissue involved and depending on several other factors.

When considering retreatment with radiation, whether it is intended to be either curative or palliative, several factors must be taken into account:

1. The dose and volume treated during the initial radiotherapy and the extent to which the re-treatment fields overlap with the initial fields
2. Whether chemotherapy was added to the initial radiotherapy
3. The time interval that has elapsed since the initial radiotherapy
4. The tissues and organs involved because they differ markedly in their ability to recover
5. Highly conformal techniques, such as stereotactic radiosurgery, stereotactic body radiotherapy (SBRT), or brachytherapy, are most appropriate.
6. Whether there are alternative options to radiation that could be considered
**EARLY- AND LATE-RESPONDING TISSUES**

Early-responding tissues are usually self-renewing tissues and are characterized by a rapidly proliferating stem cell compartment that provides cells to differentiate and become the mature functioning cells. If some stem cells survive within the irradiated volume, or if undamaged stem cells can migrate into the irradiated volume from outside, the tissue architecture may be partially or completely restored. Consequently, rapidly proliferating tissues generally recover well from the initial radiotherapy and will tolerate reirradiation to almost full doses, provided sufficient time is allowed.

By contrast, most late-responding tissues are much less able to tolerate retreatment because they do not have the ability to recover from the initial damage inasmuch as they do not have a rapidly proliferating stem cell compartment. Some slowly proliferating tissues are capable of partial proliferative and functional recovery, although this takes months and some residual damage remains.

**PRECLINICAL DATA**

Experiments with rodents show that radiation-induced skin damage can recover well, with restoration of almost full radiation tolerance. As would be expected, recovery occurs quickly after low doses, but more slowly as the initial dose is increased. Using hind limb deformation as an endpoint, representing late subcutaneous fibrosis, much poorer retreatment tolerance was observed than for the early skin reaction. Preclinical data also show that retreatment with reduced doses is possible in both lung and spinal cord after an interval of 3 to 6 months. There is a plethora of data for the spinal cord in animals ranging from rodents to rhesus monkeys. The experiments with monkeys demonstrated a large capacity for the spinal cord to recover from occult radiation injury induced by a commonly prescribed elective dose (44 Gy in 2 Gy fractions) in that only 4 of 45 monkeys developed myelopathy following a retreatment with 57.2 to 66 Gy given 1 to 3 years after the initial therapy.

Other slowly proliferating organs, such as the kidney or bladder, do not appear to be capable of recovery from late functional damage even after lower subtolerance doses.

**CLINICAL STUDIES**

Clinical radiation oncologists are, quite rightly, reluctant to transfer quantitative reirradiation data from animals (especially rodents) to humans, although the general principles and lessons, particularly the big difference between different organs and tissues, are probably applicable.

Unfortunately, the clinical data available are sketchy to say the least. In most cases, the number of patients is small, collected over a long period, with changing radiotherapy techniques (dose, fractionation pattern, etc.), and with no detailed outcome of a matched control group of patients who had received the same initial therapy, but not the retreatment.

Clinical review papers are beginning to express data in terms of the biologically effective dose (BED), calculated on the basis of the linear-quadratic model, using an $\alpha/\beta$ of 10 for early-responding tissues and 3 for late-responding tissues. This is a good first step, but unconventional fractionation patterns are often used for retreatment schedules, particularly in a palliative setting, it would be necessary to convert the BED values obtained to an equivalent BED calculated for a 2 Gy per fraction schedule if different experiences are to be compared. Unfortunately, such complete data are seldom available.

**Spinal Cord**

The spinal cord is a major dose-limiting organ in the treatment of primary tumors of the spinal cord or of neoplastic disease in the direct vicinity of the cord. When initial radiation therapy is delivered in conventional 2 Gy per day fractions, the consensus is that the incidence of myelopathy is less than 1% for total doses of 50 to 55 Gy, and up to 5% for total doses of 55 to 60 Gy. The published clinical data on reirradiation suggest that there is a substantial recovery, provided the initial treatment did not exceed about 90% of the acceptable BED and that there was a time interval of a year or more between the initial and subsequent treatments. The data suggest an important threshold of a cumulative BED of 130% to 135% of the acceptable BED in a single course of therapy (calculated with an appropriate $\alpha/\beta$ ratio of about 3). The reader is referred to the appropriate review papers (Grosu et al., 2002; Nieder et al., 2000) for more details, but the overall conclusion is that it is not possible...
to suggest more detailed recommendations concerning the optimal total dose and fraction size to result in the maximal palliative effects accomplished by minimal side effects.

There are two later reports of the use of SBRT for the salvage therapy of patients with prior radiation of spinal metastases (Nelson et al., 2009; Sahgal et al., 2009). The development of this highly conformal treatment makes it possible to give a dose of approximately 24 Gy in about three fractions, without unacceptable toxicity. The authors conclude that SBRT is both safe and effective in the palliative/reirradiation setting.

In summary, the clinical data, although minimal, do not contradict the more detailed data available from primate studies, which suggest that retreatment is a real possibility in this site.

**Brain**

There are no animal data available on the reirradiation tolerance of the brain. It might be thought that information derived from studies of the spinal cord could be transferred to the brain, but the available clinical studies indicate that this is not the case, at least, in regard to the influence of the time interval between initial treatment and reirradiation.

The most recent overview of currently available clinical data on reirradiation of the brain comes from studies of the treatment of recurrent glioma (Mayer & Sminia, 2008). For conventional radiotherapy techniques, radiation-induced normal brain tissue necrosis was found when the cumulative BED (i.e., the sum of the BEDs from both the initial treatment and reirradiation) exceeded 100 Gy. In most cases, the initial treatment consisted of a standard regimen of 60 Gy in 2 Gy fractions. The BED was calculated using the linear-quadratic formalism, employing an $\alpha/\beta$ ratio of 2 Gy and normalized to 2 Gy fractions because the literature contains several reirradiation schemes with regard to total dose and fraction size. There was no correlation between the time interval between the initial and reirradiation course and the incidence of radiation-induced normal brain tissue necrosis. When more conformal techniques were employed, such as fractionated stereotactic radiosurgery, higher retreatment doses were possible without increasing the probability of normal brain tissue necrosis.

The Radiation Therapy Oncology Group Protocol 90–95 studied 156 patients with recurrent previously irradiated primary brain tumors and brain metastases, retreated with single fraction radiosurgery. They found that the maximum tolerated doses were 24 Gy, 18 Gy, and 15 Gy for tumors less than or equal to 20 mm, 21–30 mm, and 31–40 mm in diameter, respectively. Unacceptable central nervous system toxicity was more likely in patients with larger tumors, whereas local tumor control was most dependent on the type of recurrent tumor and the treatment unit used (linac versus Gamma Knife).

**Head and Neck**

Despite significant improvements in outcome for head and neck squamous cell carcinoma treated aggressively by modern techniques, the major pattern of failure continues to be locoregional recurrence. There is no consensus on the best way to handle this problem and, as a result, there is a wide variation in the approaches used. There have been several reviews in recent years (Chopra et al., 2006; DeCrevoiser et al., 1998; Kasperts et al., 2005; Lee et al., 2007; Salama et al., 2006; Wong, 2006). Radical reirradiation with doses of 50 to 60 Gy, either definitive or postoperative, does improve locoregional control and possibly improves survival, whereas lower doses are ineffective. However, reirradiation with such potentially curative doses within a few years of a similar initial treatment is associated with severe toxicity and functional sequelae, as well as a small but non-negligible incidence of treatment-related deaths. The most recent review by Sulman et al. (2009) reported on the use of intensity-modulated radiation therapy (IMRT) to retreat recurrent head and neck cancer and came to the same general conclusion except that the treatment-related morbidity, although significant, may have been less than in previously published reports using conventional techniques. In this case, the median time interval between initial radiation and retreatment was almost 4 years, and the median lifetime dose was 116.1 Gy.

**Rectum**

The addition of preoperative or postoperative chemotherapy and or radiotherapy to radical surgery for rectal carcinoma has reduced the incidence of local recurrence to approximately 10% to 15%, but this group of patients presents a substantial problem.

Mohiuddin et al. published a review of 103 patients who underwent reirradiation with concurrent 5-fluorouracil-based chemotherapy.
Overall survival was poor and there was a substantial incidence of both early and late complications. Such a wide range of doses were used, both for the initial treatment and the reirradiation, so it is difficult to draw any firm conclusions. Nevertheless, this remains an option for a particularly difficult group of patients.

**Bone Metastases**

Bone metastases are a frequent cause of morbidity in patients with malignant disease, occurring most often in patients with breast, lung, prostate, thyroid, and renal cancers, and those with multiple myeloma. Pain is the most common symptom, which can be treated successfully in most patients by external beam radiotherapy. An increasing number of patients outlive the duration of the benefits of initial palliative radiotherapy for symptomatic bone metastases, requiring reirradiation of previously treated sites.

Chow et al. (2006) summarized the studies to date, which include several retrospective and prospective trials. The available data support the reirradiation of sites of metastatic bone pain after initial irradiation, particularly when this follows an initial period of response. In other words, patients who respond to the first treatment usually respond to subsequent treatments. There is limited evidence that a small proportion of initial nonresponders would respond to reirradiation. There is no agreement over the optimal treatment schedule to be used, with reports in the literature varying from a single fraction of 4 or 8 Gy, to fractionated regimens of up to 35 Gy in four fractions. The exact mechanism by which radiation relieves bone pain is not understood. Pain relief appears to be independent of the reduction in tumor size or cell kill, as evidenced by the rapidity of onset of pain relief and the apparent absence of a dose response. Because most patients requiring treatment for painful bone metastases have a limited life expectancy, a short treatment course, with minimum inconvenience to the patient, would seem to be warranted, unless a multifraction course is clearly superior. With this in mind, Chow et al. (2006) have set up an international phase III randomized trial comparing single with multiple fractions for reirradiation of painful bone metastases.

**Breast**

Reirradiation after recurrence of breast cancer is a major problem. Harms et al. (2004) reviewed more than 250 cases, where the reirradiation was performed either with electron beams, which have a limited depth of penetration, or with continuous low dose rate (CLDR) or pulsed dose rate (PDR) brachytherapy. With both of these techniques, high doses can be applied to the chest wall, whereas deeper-seated organs (lung, heart) can be spared to a large extent.

After retreatment using electron beams, complete remissions were obtained in 41% to 74% of the patients. The brachytherapy was judged to be superior, with complete remissions in 79% to 82% of the patients, an advantage that the authors (rightly or wrongly) attributed to the protracted irradiation schedule of the CLDR/PDR, which resulted in an improved therapeutic ratio. Severe grade IV complications occurred in less than 10% of the reirradiated patients.

This is an area where the use of hyperthermia has had some modest success. Although rather dated now, Vernon et al. (1996) reported on an international collaborative group, which combined the results of five randomized controlled trials conducted in the United Kingdom, Europe, and Canada, in which external beam radiation therapy alone was compared with radiation plus hyperthermia. The greatest advantage for the combined treatment was seen in patients with recurrent lesions in previously treated areas, in whom reirradiation was limited to lower doses. In these patients, the complete response rate was almost doubled.

**Lung**

Local recurrence of lung cancer after initial external beam radiotherapy poses substantial problems for the subsequent management of the patient. Okamoto et al. (2002) is often quoted in this context, but this paper describes a very small series of only 34 patients, with a wide range of doses used for the initial treatment (30–80 Gy) and an even wider range of retreatment doses (10–70 Gy). The authors concluded that such reirradiation resulted in a symptomatic benefit in most patients, but at a cost of substantial toxicity. Most patients succumbed to their disease before the serious late effect of lung fibrosis could be expressed.

Proton therapy, characterized by a well-defined localized high dose region, would seem to be the logical way to approach reirradiation of recurrent lung cancer. Early experience from MD Anderson Cancer Center indicates that
amazingly high doses can be tolerated, but no data have been published to date. When protons are not available, a highly conformal technique such as SBRT may give improved results.

**Recurrent Vaginal Metastases**

Radiotherapy has been an accepted form of treatment for decades, but when there is recurrent or persistent disease after a full course of radiation, the best option is radical surgery, which can result in a 5-year survival rate in excess of 60% (Rubin et al., 1987). When this is not possible, reirradiation has been used in the past as an alternate, but both local control and survival rates were poor, although there was a significant rate of complications (Jones et al., 1970).

A report from China concluded that reirradiation for late recurrence in the vagina after previous radiotherapy for cervical cancer is valuable in selected cases. Early detection, so that tumor volume is small, appears to be very important, although brachytherapy was performed in most cases (Xiang et al., 1998).

**SUMMARY OF PERTINENT CONCLUSIONS**

- Of patients presenting at major cancer centers in the Western World, 10% present with a second cancer.
- If the radiation tolerance of a given organ or tissue was exceeded by the initial treatment to the extent that function is lost, or is in process of being lost, then retreatment cannot be contemplated safely.
- If the normal tissue tolerance was not exceeded by the initial treatment, some reirradiation at a later date is safe, varying very much with the tissue or organ involved and depending on other factors.
- In general, early-responding tissues recover and tolerate retreatment better than late-responding tissues, but there are exceptions.

**Animal Studies**

- Radiation-induced skin damage recovers well, with restoration of almost full radiation tolerance. Recovery is slower after larger doses.
- Poorer retreatment tolerance for fibrosis.

- Retreatment with reduced doses is possible in both lung and spinal cord. There is much data for spinal cord in both rodents and monkeys.
- Kidney and bladder are not capable of recovery from late functional damage.

**Clinical Studies**

- Most clinical studies involve small numbers of patients, variable doses, and various time intervals between the initial treatment and reirradiation.
- Reirradiation is possible in various sites with reduced doses and with a high price in terms of morbidity.
- Most data are for head and neck. Reirradiation with 50 to 60 Gy within a few years of the initial treatment improves local control and possibly survival, but with severe toxicity and functional sequelae.
- Studies using IMRT or, better still, protons to reduce the volume of normal tissue exposed, as well as hyperfractionation to spare normal tissues, may be called for in the future, but no data are available at the present time.

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The early recognition that x-rays could produce local tumor control in some patients and not in others led to the notion that other forms of ionizing radiations might be superior.

Neutrons were first introduced in a speculative way, not based on any particular hypothesis. The later use of neutrons and the introduction of protons, negative π-mesons, and heavy ions were all based clearly on a putative advantage, either of physical dose distribution or of radiobiologic properties. In the case of neutrons, they give up their energy to produce recoil protons, α-particles, and heavier nuclear fragments. Consequently, their biologic properties differ from those of x-rays: reduced oxygen enhancement ratio (OER), little or no repair of sublethal damage, and less variation of sensitivity through the cell cycle.

The use of neutrons following World War II was based squarely on the premise that hypoxic cells limit the curability of human tumors by x-ray therapy, so the lower OER characteristic of neutrons might confer an advantage. An alternative rationale for neutrons, proposed at a later date, was that their relative biologic effectiveness (RBE) is larger for slow-growing tumors, possibly giving them an advantage in a limited number of specific human tumors.

Protons have radiobiologic properties similar to those of x-rays, and their introduction into radiotherapy was based entirely on the superiority of the physical dose distribution possible with charged particles. Negative π-mesons and heavy ions were introduced with the hope of combining the radiobiologic advantages attributed to neutrons with the dose distribution advantage characteristic of protons.

Neutrons have been shown to be superior to x-rays in a limited number of situations, specifically for the treatment of prostatic cancer, salivary gland tumors, and possibly soft tissue sarcomas. Several controlled clinical trials have been performed for various cancer sites, but a small gain was apparent only in these few circumstances. On the other hand, neutrons resulted in unacceptable late normal tissue damage in many cases, and possibly a higher incidence of radiation-induced second cancers.

For almost 50 years, protons generated in facilities originally intended for physics research have found a small but important niche for the treatment of uveal melanoma and tumors such as chordomas, which are located close to the spinal cord and therefore benefit greatly from the localized dose distribution. The wider use of protons for broad beam radiotherapy is being tested now that custom-built hospital-based facilities are available, but no advantage has been demonstrated yet in controlled phase 3 clinical trials.

Negative π-mesons and heavy ions have been used to treat hundreds of patients, but prospective randomized trials have never been completed to prove their superiority over conventional x-rays. Negative π-mesons have completely disappeared.
from the scene, but high-energy carbon ions are enjoying a renaissance in Europe and Japan.

The casual reader may be content with this overview of alternative radiation modalities and may not wish to proceed further in this chapter. Interest in high-linear energy transfer (LET) radiations for radiotherapy largely has waned, but protons are very much in vogue. In this chapter, neutrons, protons, and carbon ions are considered in turn.

- **FAST NEUTRONS**

  **Rationale**

  Neutrons were first used for cancer therapy at the Lawrence Berkeley Laboratory in California in the 1930s (Fig. 25.1). This first clinical trial of neutrons was not based on any radiobiologic rationale but was prompted largely by the availability of a new and unique beam. It is said that it received some impetus when the mother of the Lawrence brothers (E.O. Lawrence was the inventor of the cyclotron and the director of what was to become the Lawrence Berkeley Laboratory) contracted cancer, which was judged by her physician to be incurable by conventional means. She was treated with neutrons and lived for many years, although from a retrospective review of the case, it is probable that she did not have cancer in the first place. This early effort at Berkeley was hampered because the complexities of the relationship between the RBE and the dose for high-LET radiations were not understood at the time. Consequently, several patients were overdosed seriously before the trial was terminated by the entry of the United States into World War II. In reviewing their experience many years later, Dr. Robert Stone, the radiotherapist in charge of the study, concluded in his famous Janeway Lecture of 1948 that “Neutron therapy as administered by us resulted in such bad late sequelae in proportion to a few good results that it should not be continued.”

  **The Hammersmith Neutron Experience**

  The renewed interest in neutrons in the years following World War II originated at the Hammersmith Hospital in London, where neutrons were generated by the Medical Research Council’s 60-in. cyclotron. In this machine, 16-MeV deuterons incident on a beryllium target produced neutrons with a modal value of 6 MeV.

  **FIGURE 25.1** The first patient treated with neutrons in the 1930s at the Lawrence Berkeley Laboratory of the University of California. On the left is Dr. Robert Stone, the radiotherapist, and in the center is Dr. John Lawrence, the physician brother of the inventor of the cyclotron, E.O. Lawrence. (Courtesy of the University of California.)
The Hammersmith cyclotron was suggested and conceived by Gray, based on the notion that a lowered OER would be advantageous to radiotherapy. The machine suffered from the limitations of poor depth doses (equivalent to 250–kVp x-rays) and a fixed horizontal beam.

A prospective randomized clinical trial to compare neutrons with x-rays was started in 1971. Advanced tumors of the head and neck were chosen because the poor depth-dose characteristics made neutrons suitable only for treating relatively superficial lesions. The trial involved patients with tumors of the salivary glands, buccal cavity, hypopharynx, and larynx. The neutron treatments delivered in only 12 fractions were clearly superior as judged by local control of the primary tumor, but the gain was achieved at the expense of a higher complication rate.

### The United States Neutron Experience

Because of the limitations associated with low-energy cyclotrons, interest at several centers in the United States turned to the use of large cyclotrons that accelerate deuterons to energies of 22 to 50 MeV. Several such machines had been built for high-energy physics research and were converted for part-time neutron therapy. In addition, neutrons were produced at the Fermilab in Batavia, Illinois, by bombarding a beryllium target with 67-MeV protons.

All of these machines had adequate dose rates and quite good depth doses. Unfortunately, they had other disadvantages: All had fixed horizontal beams, and all were located in physics installations rather than in large, busy hospitals, so that the availability of a sufficient number of patients was a problem. Several controlled clinical trials were performed for various tumor sites, and they showed no advantage for neutrons over x-rays. Neutrons, however, appeared to be superior for salivary gland tumors, soft tissue sarcomas, and prostate cancer, but the downside was a significant increase in late normal tissue damage. It appears that high-LET radiation loses the differential sparing effect between tumors and late-responding normal tissues that is characteristic of x-rays.

Enthusiasm for neutron therapy waned just at a time when technology became available that allowed machines to be built that are suitable for clinical use. A new generation of hospital-based cyclotrons using the p⁺ → Be reaction had adequate dose rates, good percentage depth doses, and a full isocentric mount, similar to a conventional linac. A few centers operated such machines for a while, but neutrons have never become mainstream in radiotherapy.

### BORON NEUTRON CAPTURE THERAPY

The basic idea behind boron neutron capture therapy (BNCT) is elegant in its simplicity. It has appealed to physicians, and particularly to physicists, for the best part of half a century. The idea is to deliver to the cancer patient a boron-containing drug that is taken up only in tumor cells and then to expose the patient to a beam of low-energy (thermal) neutrons that themselves produce little radiobiologic effect but that interact with the boron to produce short-range, densely ionizing α-particles. Thus, the tumor is intensely irradiated, but the normal tissues are spared. There are two problems inherent in this idea that have so far proved intractable:

1. How does one find a “magic” drug that can distinguish malignant cells from normal cells? (The skeptic might add that searching for such a drug has been the holy grail of cancer research and that if one were found, the obvious strategy would be to attach an alkylating agent or an α-emitting radionuclide to it; combining its use with neutrons would be a distant third.)

2. The low-energy neutrons necessary for BNCT are poorly penetrating in tissue and consequently result in percentage depth doses that are extremely poor by today’s standards.

Several nuclides have high propensities for absorbing low-energy or thermal neutrons; that is, they have a high neutron capture cross section. Boron is the most attractive of these because it is readily available in a nonradioactive form, its chemistry is such that it can be incorporated into various compounds, and if it interacts with low-energy neutrons, it emits short-range, high-LET α-particles.

### Boron Compounds

For BNCT to be successful, the compounds used should have high specificity for malignant cells, with concomitantly low concentrations in adjacent normal tissues and in blood. In the
early days, the compounds used were not specially synthesized for BNCT, but were already available. In the brain, which is the site for which BNCT largely has been used, some selectivity is obtained because compounds do not penetrate normal brain tissue to the same degree as brain tumors in which the blood–brain barrier is absent or severely compromised.

Critical to the success of BNCT is the requirement that boron compounds be developed that target tumor versus normal cells selectively, achieve a sufficient concentration within the tumor, and produce tumor to normal tissue ratios of 3 or 4 to 1. This, of course, is a tall order.

Two classes of compounds have been proposed:

1. Low-molecular-weight agents that simulate chemical precursors required for tumor cell proliferation have the ability to traverse the cell membrane and be retained intracellularly. Two boron compounds have been identified and used clinically, known as sodium brocapitate and boronophenylalanine. Both have been used to treat brain tumors, and the latter also has been listed for cutaneous melanoma.

2. High-molecular-weight agents such as monoclonal antibodies and bispecific antibodies that contain boron have been developed. These are highly specific, but very small amounts reach brain tumors following systemic administration. Boron-containing conjugates of epidermal growth factor, the receptor for which is overexpressed on some tumors (including glioblastoma), also have been developed.

If the blood–brain barrier is disrupted temporarily, these high-molecular-weight compounds may have some utility, or direct intracerebral delivery may be required. They have not yet proved effective in clinical use.

**Neutron Sources**

During fission within the core of a nuclear reactor, neutrons are “born” that have a wide range of energies. Neutron beams can be extracted from the reactor by the application of suitable techniques and the use of appropriate moderators. Thermal neutrons, or room-temperature neutrons (0.025 eV), react best with boron to produce densely ionizing α-particles. Unfortunately, thermal neutrons are attenuated rapidly by tissue; the half-value layer is only about 1.5 cm. Consequently, it is not possible to treat to depths of more than a few centimeters without heavily irradiating surface normal tissues. Nevertheless, most clinical trials have used neutrons of this energy.

Current interest in the United States focuses on the use of epithermal neutron beams (1–10,000 eV), which have a somewhat greater depth of penetration. These can be obtained by using moderators or filters to slow the fast neutrons into the epithermal range and filtering out the residual thermal neutrons. These epithermal neutrons do not interact themselves with the boron but are degraded to become thermal neutrons in the tissue by collisions with hydrogen atoms. Even so, the peak in dose occurs at a depth of only 2 to 3 cm, with a rapid falloff beyond this depth. Thus, the very high surface doses are avoided, but the depth doses are still poor.

**Clinical Trials**

Several clinical trials have been performed over the years, beginning in the 1950s and 1960s. Results are tantalizing but never definitive. Several patients have been treated with BNCT in the United States, but the results are largely anecdotal. The concept of BNCT is as attractive as ever, but it continues to be difficult to convert to a practical treatment modality, even for shallow tumors.

**PROTONS**

Dr. Robert Wilson first proposed protons for radiotherapy as early as 1946, based on their superior physical dose distribution; their radiobiologic properties are unremarkable. The RBE of protons is indistinguishable from that of 250-kV x-rays, which means that they are about 10% more effective than megavoltage x-rays generated by a linear accelerator. The OER for protons also is indistinguishable from that for x-rays, namely about 2.5 to 3. These biologic properties are consistent with the physical characteristics of high-energy proton beams; they are sparsely ionizing, except for a very short region at the end of the particles’ range, just before they stop. In the entrance plateau, the average LET is about 0.5 keV/μm, rising to a maximum of 100 keV/μm over a few microns as the particles come to rest. This high-LET component is restricted, however, to such a tiny length of track and represents such a small proportion of the energy deposited that, for high-energy protons, it does not have any significant effect.
To treat cancer in humans with charged particles, high-energy accelerators are necessary. Protons have been used for many years to treat choroidal melanoma of the eye as an attractive alternative to enucleation. Because the eye is small, an energy of 60–70 MeV is adequate. However, to treat deep-seated solid tumors, an energy of about 200 MeV, corresponding to a range in tissue of about 26 cm, is essential. This requires a large and expensive cyclotron or synchrotron, and so it is not surprising that the number of centers where charged particle therapy is available is limited.

**Depth-Dose Patterns and the Bragg Peak**

The dose deposited by a beam of monoenergetic protons increases slowly with depth, but reaches a sharp maximum near the end of the particles’ range in the **Bragg peak**. The beam has sharp edges, with little side-scatter, and the dose falls to zero after the Bragg peak at the end of the particles’ range. The possibility of precisely confining the high-dose region to the tumor volume and minimizing the dose to surrounding normal tissue is obviously attractive to the radiotherapist. Protons come closest to realizing this dream at modest cost.

Figure 25.2 shows the depth-dose curve for the 187-MeV proton beam from the synchrocyclotron at Uppsala, Sweden. The sharply defined Bragg peak occurs at a depth in tissue that depends on the initial energy of the particles.

The early medical use of proton beams involved treatment of the pituitary, first in patients with advanced breast cancer and later in patients with diabetic retinopathy, Cushing disease, and acromegaly. Protons were used for these applications to exploit their well-defined beam, which made it possible to give a huge dose to the pituitary without causing unacceptable damage to nearby structures. These treatments were performed for many years at both Berkeley and Harvard.

The way the Bragg peak can be spread out to encompass a tumor of realistic size is illustrated in Figure 25.3. In this figure, curve A shows the narrow Bragg peak of the primary beam of the 160-MeV proton beam at the Harvard cyclotron. Beams of lower intensity and shorter range, shown in curves B, C, D, and E, are readily obtainable either by passing the beam through a rotating wheel with plastic sectors of varying thickness, or by varying the energy of the beam. The composite curve, S, which is the sum of all the individual Bragg peaks of the beams of varying range, results in a uniform dose of more than 2.8 cm. The spread-out Bragg peak (SOBP), of course, can be made narrower or broader than this, as necessary.

Many researchers consider protons to be the treatment of choice for choroidal melanoma. Figure 25.4 shows the dose distribution that is achieved at the Harvard cyclotron, which allows very high doses to be delivered to small tumors without unacceptable damage to nearby normal tissues. Protons have found a small but important place in the treatment of ocular tumors and some specialized tumors close to the spinal cord.

Broad beam radiotherapy, with the Bragg peak spread out to cover a large tumor, has been in progress at Uppsala since 1957, and a comparable US effort began at Harvard in 1961 and at Loma Linda University in 1990. Most of the proton machines used in the past were built initially for physics research and were located in physics laboratories. There is much current interest in the development of hospital-based proton facilities producing beams sufficiently penetrating to make possible the treatment of any cancer sites in the human; such facilities would include a gantry with an isocentric
FIGURE 25.3 The way the Bragg peak for a proton beam can be spread out. Curve A is the depth-dose distribution for the primary beam of 160-MeV protons at the Harvard cyclotron, which has a half-width of only 0.6 cm. Beams of lower intensity and shorter range, as illustrated by curves B, C, D, and E, can be added to give a composite curve S, which results in a uniform dose of more than 2.8 cm. The broadening of the peak is achieved by passing the beam through a rotating wheel with sectors of varying thickness. (Adapted from Koehler AM, Preston WM. Protons in radiation therapy. Comparative dose distributions for protons, photons, and electrons. *Radiology.* 1972;104:191–195, with permission.)

mount and would feed several treatment rooms. The first machine of this kind was built at Loma Linda University in California, where a 250-MeV cyclotron produces a proton beam that can be directed into any one of four treatment rooms. The layout of the facility is shown in Figure 25.5. The intention is to treat a broad spectrum of human cancers, not just the limited sites for which the dose distributions possible with protons already have proved their worth.

An even more impressive facility has been completed at the Massachusetts General Hospital...
in Boston. Although they formerly used a fixed horizontal beam from a cyclotron in the Harvard Physics Department, they now have a new facility constructed within the hospital. Similar facilities have been constructed at the MD Anderson Hospital in Houston, Texas, and at the University of Florida, whereas others are in the planning or construction stage at several locations in the United States, Europe, and Japan. Such facilities set the scene for the future. Figure 25.6 shows a

![Figure 25.5](Image)

**Figure 25.5** Model of the proton facility at Loma Linda. Protons are accelerated to energies up to 250 MeV in a large cyclotron. The protons then can be directed into any one of four treatment rooms. This arrangement minimizes “idle” time, because while one patient is being treated in one room, the next two patients can be set up in adjoining treatment rooms. This sort of facility sets the scene for the future: a large radiation therapy facility with multiple treatment rooms in the context of a cancer center. (Courtesy of Drs. James Slater and John Archambeau, Loma Linda University, Loma Linda, California.)

![Figure 25.6](Image)

**Figure 25.6** Treatment planning comparison of a seven-field intensity-modulated photon treatment plan (left) with an intensity-modulated proton (three scanned pencil beams) plan (right) for a patient with an epithelioid sarcoma involving the paravertebral tissues adjacent to the fifth through seventh vertebrae. The 30%, 70%, and 100% isodose lines are shown. The dashed line outlines the target volume. Although dose to the target volume is similar, note the significantly higher integral dose to the lungs and heart with the intensity-modulated photon plan, which may be associated with an increase in acute and late normal tissue toxicity. Treatment comparison plans were prepared by Alexei V. Trofimov, PhD, at the Northeast Proton Therapy Center using the KonRad radiation treatment planning system originally designed at Deutsches Krebsforschungszentrum, Heidelberg, Germany. For further illustrations and discussion, see Weber DC, Trofimov AV, DeLaney TF, et al. A treatment planning comparison of intensity-modulated photon and proton therapy for paraspinal sarcomas. Int J Radiat Oncol Biol Phys. 2004;58:1596–1606. (Courtesy of Dr. Thomas DeLaney, Massachusetts General Hospital, Boston.)
where a large body of radiobiologic data was accumulated and some patients were treated with various heavy ions; however, the facility closed some years ago. Interest has been rekindled largely in Europe and Japan where, at the time of writing, there are three centers using high-energy carbon ions for radiotherapy, and there is the experience of about 5,000 patients treated.

Depth-Dose Profiles

The characteristic depth-dose profile of heavy ions, which they share with protons (i.e., the increase in dose with depth of penetration), makes them attractive as a modality for the radiotherapy of deep-seated tumors.

Figure 25.7 shows the depth-dose profiles for carbon-12 (\(^{12}\text{C}\)) ions of two different energies compared with that for cobalt-60 (\(^{60}\text{Co}\)) \(\gamma\)-rays. In the case of carbon ions, dose is almost constant until close to the end of the range, when there is a sharp increase in dose in the Bragg peak. The depth at which the peak occurs depends on the energy. The peak is much too narrow to cover any tumor of realistic dimensions, and so the Bragg peak must be “spread out” to the required dimension by varying the energy of the beam. For carbon ions produced by a synchrotron, this can be achieved by varying the energy from pulse to pulse. Any required field size can be achieved from the narrow pencil beam that emerges from the synchrotron by “scanning” the beam, that is, deflecting the beam with magnetic fields so that, voxel by voxel, it conforms to the desired shape and size.

Figure 25.8 compares protons with carbon ions. For both ions, the depth-dose profile is characterized by a sharp Bragg peak.

Radiobiologic Properties

Heavy ions, including carbon ions, differ from both protons and high-energy \(\gamma\)-rays in their radiobiologic properties because of their higher LET. This is a classic case of “good news–bad news” because some differences may represent an advantage, whereas others clearly are disadvantages. The differences can be summarized as follows:

1. A lower OER. This is likely to be an advantage in that heavy ions may partly overcome the relative radioresistance of some tumors because of hypoxia. However, carbon ions are characterized by a modest reduction in
**FIGURE 25.7** Comparison of the depth-dose profiles of carbon ions of two different energies with that of cobalt-60 $\gamma$-rays. (Adapted from Kraft G. Tumor therapy with heavy charged particles. *Prog Part Nucl Phys.* 2000;45:S473–S544.)

**FIGURE 25.8** Illustrating the depth-dose profiles for protons and carbon ions compared with conventional high-energy photons. **Top panel:** Both protons and carbon ions are characterized by a sharp Bragg peak at the end of the particles’ range, but it is too sharp and too narrow to encompass a tumor of realistic size. **Bottom panel:** By using a range shifter, or by varying the energy of the beam, the Bragg peak can be spread out to encompass a tumor of any size at any depth. For both protons and carbon ions, the dose to the tumor, located in the SOBP, is higher than in the normal tissue ahead of the SOBP, but the differential is greater for carbon ions because part of the radiation dose in the SOBP has a high LET and has a correspondingly higher RBE.
OER; heavier ions would be called for if a reduced OER turns out to be the dominant advantage of heavy ions.

2. A loss of repair capacity. This is probably a major disadvantage because an important reason for the efficiency of x-rays is that they allow normal tissues to repair damage between dose fractions, more perhaps than tumors. Dose fractionation would no longer result in a differential sparing of normal tissues when heavy ions are used.

3. A smaller variation in radiosensitivity with phase of the cell cycle. This is certainly not an advantage and may be a disadvantage for heavy ions because cells in normal tissues, especially late-responding normal tissues, spend less time in the radiosensitive phases of the cell cycle (G2/M) than do cycling tumor cells. This relative sparing of normal tissues is lost if heavy ions are used, for which radiosensitivity varies less through the cell cycle.

Given the favorable depth-dose distribution and the high cell-killing effectiveness of carbon ions, short-course hypofractionated radiotherapy is currently being investigated in Europe and Japan, using only one or two large dose fractions. Time will tell if this proves to be a successful strategy.

Relative Biologic Effectiveness Considerations

For carbon ions, there is the additional factor, which may be exploited to advantage, of an increase in RBE toward the end of the particle range. The rapid change of RBE with depth is shown in the top panel of Figure 25.9. This is a classic case of “good news–bad news.” In principle, it is good news because the effectiveness of the dose is increased toward the end of the track, which is located in the tumor volume, exaggerating the change of effective dose with depth; the bad news is that one does not know the correct RBE to use, because RBE values must come from in vitro or animal experiments. Consequently, there is an element of uncertainty in these data.

Scattering and Fragmentation

Compared with protons, carbon ions have less lateral scattering, leading to sharper beam edges, but there is a “tail” of lighter fragments beyond the Bragg peak. This is illustrated in the lower panel of Figure 25.9. Carbon ions, therefore, do not share the proton advantage that dose stops sharply at the end of the range of the primary particle.

Comparison of Protons and Carbon Ions

Figure 25.8 compares protons with carbon ions. For both ions, the depth-dose profile is characterized by a sharp Bragg peak at the end of the particles range, just before it stops. However, as previously pointed out, it is too narrow to encompass a tumor of realistic size. By using a range spreader, or by varying the energy of the beam, the Bragg peak can be spread out to cover a tumor of any size at any depth (bottom panel of Fig. 25.9). The dose to the tumor located in the SOBP is higher than in the normal tissue ahead of the SOBP for both...
protons and carbon ions, but the differential is greater for carbon than for protons because of the high-LET component in the Bragg peak of the carbon ions.

**Positron Emission Tomography Verification of Treatment Plans**

There is a unique opportunity to verify the treatment plan in the case of carbon ions. The basis is as follows: When a beam of carbon ions penetrates a thick absorber, a small fraction of the ions will undergo nuclear fragmentation—that is, they will break up. A frequent process is the stripping of one or two neutrons, converting the stable carbon-12 to the positron-emitting isotopes carbon-11 and carbon-10. These isotopes travel with almost the same velocity as the main beam and stop in almost the same place. They have short half-lives, and as they annihilate, they emit γ-rays that can be detected in a conventional positron emission tomography (PET) scanner. Consequently, the location of the SOBP, and therefore the high-dose treatment volume, is visualized.

**SUMMARY OF PERTINENT CONCLUSIONS**

**Neutrons**

- Neutrons are indirectly ionizing. In tissue, they give up their energy to produce recoil protons, α-particles, and heavier nuclear fragments.
- Biologic properties of neutrons differ from those of x-rays in several ways: reduced OER, little or no repair of sublethal damage or potentially lethal damage, and less variation of sensitivity through the cell cycle.
- The rationale for the use of neutrons in radiotherapy has changed over the years. The earlier rationale was the reduced OER to overcome the problem of hypoxic cells. The revised rationale is based on a higher neutron RBE for slowly growing tumors.
- A small advantage has been demonstrated in clinical trials for neutrons in the treatment of salivary gland and prostate tumors and soft tissue sarcomas, but not for most cancer sites tested. The downside of neutrons is the unacceptable level of normal tissue damage.

- A new generation of hospital-based cyclotrons with isocentric mounts and generating neutrons by the $p^+ \rightarrow Be$ reaction became available, but by then, enthusiasm for neutrons had waned.

**Boron Neutron Capture Therapy**

- The principle of BNCT is to deliver a drug-containing boron that localizes only in tumors and then to treat with low-energy thermal neutrons that interact with boron to produce α-particles.
- Boron is a suitable substance because it has a large cross section for thermal neutrons and emits short-range, densely ionizing α-particles if bombarded by thermal neutrons. Its chemistry is such that it can be incorporated into a wide range of compounds.
- Many attempts have been made to synthesize boron-containing compounds that are selectively localized in tumors relative to normal tissues, but with limited success. They fall into two categories:
  1. Low-molecular-weight agents that simulate chemical precursors needed for tumor cell proliferation
  2. High-molecular-weight agents such as monoclonal antibodies and bispecific antibodies
- Thermal neutrons are poorly penetrating in tissue, with a half-value layer of only 1.5 cm.
- Epithermal neutrons are somewhat more penetrating. They are degraded to thermal neutrons by collisions with hydrogen atoms in tissue. The peak dose is at 2 to 3 cm, and the high surface dose is avoided.
- Results of clinical trials of the efficacy of BNCT are tantalizing but not definitive.
- The concept of BNCT is very attractive, but there are formidable practical difficulties in making it a treatment modality even for relatively shallow tumors.

**Protons**

- Protons result in excellent physical dose distributions.
- Protons have biologic properties similar to those of x-rays.
- There is an established place for protons in the treatment of choroidal melanoma or tumors close to the spinal cord in which a sharp cutoff of dose is important.
Hospital-based high-energy cyclotrons with isocentric mounts are now being used to treat a broader spectrum of cancer patients with protons. Their efficacy has yet to be proved in clinical trials, but they offer the obvious physical advantage of good dose distributions with reduced dose to normal structures.

**Carbon Ion Radiotherapy**

The use of high-energy carbon ions has experienced a renaissance in Europe and Japan.

For carbon ions, as for protons, the narrow Bragg peak must be spread out to cover a tumor of realistic dimensions by varying the energy of the beam.

Inevitably, the RBE varies across the SOBP. In theory, this could be an advantage for carbon ions because it further exaggerates tumor dose relative to normal tissue. In practice, it is a complication because one must estimate RBE values from experiments in model systems.

Hypofractionation with only one or two large dose fractions is under investigation for carbon ions.

A unique attraction of carbon ion therapy is that the target volume can be visualized by PET as some carbon-12 ions decay to radioactive carbon-11 and carbon-10.

**BIBLIOGRAPHY**


Oxygen homeostasis is deregulated in several pathologic settings. Hypoxia, or low oxygen conditions, exists in virtually all solid tumors. As tumor cells rapidly expand, their growth outpaces the ability of the existing vasculature to supply both nutrients and oxygen, resulting in tumor hypoxia. Cells experiencing hypoxia can cope with the low oxygen tension by either increasing delivery of oxygen or adapting to the low oxygen level.

**HYPOXIA-INDUCIBLE FACTOR**

Hypoxia-inducible factors (HIFs) are transcription factors that facilitate both oxygen delivery and adaptation to oxygen deprivation by regulating the expression of genes that are involved in many cellular processes including glucose uptake and metabolism, angiogenesis, erythropoiesis, cell proliferation, and apoptosis. HIFs bind to DNA as heterodimers composed of an oxygen-sensitive α-subunit and a constitutively expressed β-subunit, also known as the aryl hydrocarbon receptor nuclear translocator (ARNT). To date, three HIFs (HIF-1, -2, and -3) have been identified that regulate transcriptional programs in response to low oxygen levels.

HIF-1 was the first HIF family member to be characterized. Using DNA affinity purification, HIF-1 was identified as a hypoxic-induced factor that bound an 18-nucleotide fragment of the erythropoietin enhancer required for the hypoxic activation of erythropoietin in Hep3B cells. Structural analysis of the HIF-1α protein revealed four distinct domains, including an oxygen-dependent degradation domain (ODD) that regulated HIF-1α degradation by the ubiquitin-proteasome pathway. HIF-1 has been designated the global regulator of hypoxia-inducible gene expression because it is ubiquitously expressed and induces the expression of most hypoxia-inducible genes.

HIF-2 was the second HIF family member identified and shares approximately 50% sequence homology with HIF-1. HIF-2α is also sensitive to changes in oxygenation and possesses an ODD. Although HIF-2α heterodimerizes with ARNT like HIF-1α, the expression of HIF-2α is restricted to endothelial cells, glial cells, pneumocytes type II, interstitial cells of the pancreas and duodenum, and hepatocytes.

The third HIF family member, HIF-3α, also possesses an ODD, can dimerize with ARNT and bind to hypoxia response elements in vitro. The role of HIF-3 in the hypoxic regulation of target gene expression in vivo is not well understood.
Oxygen-Dependent Regulation of Hypoxia-Inducible Factor

The transcriptional activity of HIF is regulated through changes in protein stability and through the recruitment of transactivation coactivators (Fig. 26.1). In the presence of oxygen, HIF-1α is hydroxylated on two highly conserved proline residues: prolines 402 and 564, both found within the ODD. A family of 4-prolyl hydroxylases (PHDs), distinct from the structurally important collagen-4-prolyl hydroxylases, modifies these prolines. These enzymes catalyze a hydroxylation reaction, which uses molecular oxygen and 2-oxoglutarate as substrates and iron and ascorbate as cofactors. Hydroxylation of HIF-α is necessary for binding to the von Hippel-Lindau (VHL) tumor suppressor protein. Under normoxic conditions, HIF-α protein is rapidly turned over because of hydroxylation, binding to VHL, and destruction by the proteasome. Under hypoxic conditions, the 4-prolyl hydroxylases have reduced activity as oxygen is a requisite substrate of the hydroxylation reaction. Decreased hydroxylation of HIF-α allows it to escape recognition by VHL, resulting in protein stabilization. Once stabilized, HIF-α dimerizes with ARNT and binds to a core sequence of 5′-RCGTG-3′ in the enhancer elements of target genes to initiate gene transcription.

An additional layer of HIF regulation occurs at the level of recruitment of coactivators. A third hydroxylation of HIF-α takes place on the C-terminal transactivation domain on asparagine residue 803. This hydroxylation reaction is catalyzed by a separate asparaginyl hydroxylase, factor inhibiting HIF (FIH). Hydroxylation of
Most critical because it has potent antiangiogenic properties and is expressed in a large number of human tumors.

**Tumor Metabolism**

More than 70 years ago, cancer cells were found to shift glucose metabolism from an oxidative to glycolytic pathway. This shift has become known as the Warburg effect, and is characterized by decreased mitochondrial respiration and increased lactate production even with oxygen. Numerous studies have established that HIF-1 regulates the expression of genes involved in glycolytic metabolism including glucose transporters, glycolytic enzymes, lactate production, and pyruvate metabolism.

**Tumor Metastasis**

Metastasis is the cause of most human cancer deaths. It occurs in a series of distinct steps that include tumor cell invasion, intravasation, extravasation, and proliferation. HIF activation correlates with metastasis in multiple tumors and promotes metastasis through the transcriptional regulation of key factors such as E-cadherin, lysyl oxidase, and CXCR4 that govern cell adhesion, extracellular matrix formation, and cell migration.

**Hypoxia-Inducible Factor and Radiotherapy**

It has recently been proposed that HIF-1 plays an important role in determining tumor response to radiotherapy and tumor regrowth. Radiation results in a reoxygenation-dependent increase in HIF-1 activity by two distinct mechanisms. Tumor reoxygenation results in both HIF-1 stabilization and enhanced translation of HIF targets through the release of reactive oxygen species. The effects of HIF on tumor radiosensitivity are twofold. On one hand, HIF-1 stabilization promotes tumor vasculature radiotolerance through the release of proangiogenic cytokines such as VEGF. Paradoxically, HIF-1 can also induce tumor radiosensitivity through the induction of apoptosis. Overall, HIF-1 stabilization promotes radiotolerance because HIF-1-deficient tumors are more sensitive to radiation compared to wild-type tumors.

Studies by Ahn and Brown’s laboratory have identified an important role for bone marrow (BM)-derived cells in regenerating tumor vasculature following radiotherapy. Tumors grown...
in previously irradiated tissues exhibit decreased growth caused by insufficient neovascularization that results from radiation-induced injury to the host vasculature and connective tissue. They hypothesized that the vasculature of tumors grown in previously irradiated tissues is derived from cells from the BM. In particular, they demonstrated an important role for tissue MMP-9 and CD11b+ myelomonocytic cells from the BM to restore tumor growth in these preirradiated sites. HIF plays an important role in this recruitment of BM-derived cells by regulating several factors such as CXCR4 and SDF-1. Inhibition of HIF results in an inhibition of tumor vascularization. This study implicates vasculogenesis as an important target for adjunct therapy to radiotherapy.

UNFOLDED PROTEIN RESPONSE

Prolonged periods of hypoxia can activate non-HIF signaling pathways such as the unfolded protein response (UPR). The UPR is a cellular stress response that is induced by the accumulation of unfolded proteins in the endoplasmic reticulum (ER) in order to deal with the problems associated with the accumulation of misfolded proteins. Activation of the UPR results in a transient inhibition of protein translation to prevent further misfolded proteins from accumulating and through the induction of chaperone proteins that function in the ER to assist in folding, trafficking, and secretion of proteins (Fig. 26.2). The UPR has been well characterized in yeast and, more recently, a greater understanding has been elucidated in humans. The ER stress sensor inositol-requiring kinase 1 (IRE1) is found both in yeast and in humans. IRE1 becomes activated through homodimerization and trans-autophosphorylation. Biochemically, the endoribonuclease activity of IRE1 splices the pre-messenger ribonucleic acid (mRNA) of the X-box binding protein (XBP1) transcription factor that results in translation of XBP1. The XBP1 transcription factor has many target genes involved in ER protein maturation. Koong et al. have shown that XBP1 could be inhibited by small molecules and that this
inhibition resulted in a significant inhibition of tumor growth.

A second ER stress sensor is the protein kinase-like endoplasmic reticulum kinase (PERK). Activation of PERK prevents further translation of proteins. PERK activation occurs rapidly under severe hypoxia on the order of minutes, whereas under milder hypoxic conditions, PERK activation occurs on the order of hours. Through a combination of oligomerization and autophosphorylation, PERK becomes activated and inhibits translation by directly phosphorylating serine 51 of eIF2α, a key initiator of the mRNA translation machinery. Koumenis and colleagues have shown that activation of PERK is important to survive under hypoxic conditions as PERK-deficient cells undergo apoptosis following hypoxia, resulting in a lower survival rate compared to genetically matched cells that have the PERK pathway intact. These PERK-deficient cells are less tumorigenic, likely in part because of higher apoptosis in the hypoxic regions of the tumor. Thus, one cellular response to chronic conditions of hypoxic stress is to limit the load of misfolded proteins.

**RADIOSENSITIZING HYPOXIC CELLS**

As described in Chapter 21, a great deal of experimental work through the years had established that at least in transplanted tumors in animals, tumor control by x-rays frequently is limited by the foci of hypoxic cells that are intransigent to killing by x-rays, which may result in tumor regrowth. Among the methods suggested to overcome this problem are treatment in hyperbaric oxygen chambers and the introduction of high-linear energy transfer (LET) radiations, such as neutrons and heavy ions. Chemical sensitizers address the same problem. High-LET radiations are discussed in Chapter 7. This chapter addresses hyperbaric oxygen, chemical radiosensitizers, and the latest approach, namely, hypoxic cytotoxins.

**Hyperbaric Oxygen**

Following the identification of hypoxia as a possible source of tumor resistance, a major effort was made to solve the problem by the use of hyperbaric oxygen. Patients were sealed in chambers filled with pure oxygen raised to a pressure of three atmospheres. Churchill Davidson at St. Thomas’ Hospital in London pioneered this work, but it was taken up by researchers on both sides of the Atlantic. The clinical trials that were performed involved small numbers of patients and were difficult to interpret because unconventional fractionation schemes were used; that is, a few large fractions were used because of the time and effort involved in the technical procedures. Patient compliance was also a problem because of the feeling of claustrophobia from being sealed in a narrow tube. There was also the serious risk of fire, because tissue is highly flammable in pure oxygen (as evidenced by the accident when the crew of a space capsule died in an oxygen fire on the ground), although in practice, an accident of this nature never happened during the treatment of several thousand patients. The largest multicenter trials performed by the Medical Research Council in the United Kingdom showed a significant benefit both in local control and in survival for patients with carcinoma of the uterine cervix and advanced head and neck cancer, but not for patients with bladder cancer. The data generated a great deal of debate. An overview of the trials showed a 6.6% improvement in local control, with a suggestion, too, of an increase in late normal tissue damage. Hyperbaric oxygen fell into disuse, partly because it is cumbersome and difficult in practice, and partly because of the promise of drugs that would achieve the same result by simpler means.

The notion of improving tumor oxygenation by breathing 100% oxygen rather than air has been revived in recent years by experiments involving carbogen. If pure oxygen is breathed, it tends to lead to vasoconstriction, a closing down of some blood vessels, which, of course, defeats the object of the exercise. This is avoided if 5% carbon dioxide is added to the oxygen, a mixture called carbogen. Breathing carbogen at atmospheric pressure, then, is a relatively simple attempt to overcome chronic hypoxia, that is, diffusion-limited hypoxia. The use of carbogen in combination with nicotinamide is described subsequently in this chapter.

**Improving the Oxygen Supply to Tumors**

A group at the Princess Margaret Hospital in Toronto showed convincingly that a blood transfusion prior to radiotherapy led to a significant
improvement in local tumor control probability in patients with carcinoma of the uterine cervix. Several other studies have shown that hemoglobin levels can influence the success of radiation therapy.

Tumor oxygenation also can be improved by the use of artificial blood substances such as perfluorocarbons. Because smoking can decrease tumor oxygenation, it is clearly advisable for patients to give up smoking, at least during radiotherapy.

**Hypoxic Cell Radiosensitizers**

Spurred largely by the efforts of radiation chemists (most notably, Adams), a search was under way in the early 1960s for compounds that mimic oxygen in their ability to sensitize biologic materials to the effects of x-rays. Instead of trying to “force” oxygen into tissues by the use of high-pressure tanks, the emphasis shifted to oxygen substitutes that diffuse into poorly vascularized areas of tumors and achieve the desired effect by chemical means. The vital difference between these drugs and oxygen, on which their success depends, is that the sensitizers are not rapidly metabolized by the cells in the tumor through which they diffuse. Because of this, they can penetrate further than oxygen and reach all of the hypoxic cells in the tumor, including those most remote from a blood supply. In the early 1960s, many simple chemical compounds were found to have the ability to sensitize hypoxic microorganisms. These studies were guided by the hypothesis, now known to be correct, that sensitizing efficiency is related directly to the electron affinity of the compounds.

Adams and his colleagues listed properties that would be essential for a clinically useful hypoxic cell sensitizer. First, it has to selectively sensitize hypoxic cells at a concentration that would result in acceptable toxicity to normal tissues. Second, it should be chemically stable and not subject to rapid metabolic breakdown. Third, it must be highly soluble in water or lipids and must be capable of diffusing a considerable distance through a non-vascularized cell mass to reach the hypoxic cells, which in a tumor may be located as far as 200 \( \mu \text{m} \) from the nearest capillary. Fourth, it should be effective at the relatively low daily doses of a few grays used in conventional fractionated radiotherapy. The first candidate compound that appeared to satisfy these criteria was misonidazole.

**Misonidazole**

Figure 26.3 illustrates the numbering of the basic ring structure of the nitroimidazoles. The side chain determines position 1, and the position of the nitro group (\( \text{NO}_2 \)) leads to the classification of the drug as a 2-nitroimidazole, 4-nitroimidazole, and so on. In general, 2-nitroimidazoles have a higher electron affinity than 5-nitroimidazoles, the class that includes metronidazole, which was briefly tried as a radiosensitizer. Misonidazole is a 2-nitroimidazole; its structure is shown in Figure 26.4.

Misonidazole produces appreciable sensitization with cells in culture (Fig. 26.5). Hypoxic cells in 10 mM of misonidazole have a radiosensitivity approaching that of aerated cells. Misonidazole also has a dramatic effect on tumors in experimental animals. This is illustrated in Figure 26.6, which shows the proportion of mouse mammary tumors controlled as a function of x-ray dose delivered in a single fraction. If x-rays are used alone, the dose required to control half of the tumors is 43.8 Gy. This falls to 24.1 Gy if the radiation is delivered 30 minutes after the administration of misonidazole (1 mg/g body weight). This corresponds to an enhancement ratio of 1.8. Dramatic results, such as those shown in
The number of clinical trials, involving many different types of human tumors, in Europe and the United States. In general, the results have been disappointing. Of the 20 or so randomized prospective controlled clinical trials performed in the United States by the Radiation Therapy Oncology Group (RTOG), none yielded a statistically significant advantage for misonidazole, although a number indicated a slight benefit. The only trial that shows a clear advantage for misonidazole is the head and neck cancer trial performed in Denmark, the largest single trial performed.

Figure 26.6 in which an enhancement ratio of 1.8 was obtained, are rather misleading; they represent single-dose treatments, in contrast to the multifraction regimens common in conventional radiotherapy. Most animal tumors reoxygenate to some extent between irradiations, so in a multifraction regimen, the enhancement ratio for a hypoxic cell sensitizer is usually much less than for a single-dose treatment.

After encouraging results in laboratory studies, misonidazole was introduced into a large number of clinical trials, involving many different types of human tumors, in Europe and the United States. In general, the results have been disappointing. Of the 20 or so randomized prospective controlled clinical trials performed in the United States by the Radiation Therapy Oncology Group (RTOG), none yielded a statistically significant advantage for misonidazole, although a number indicated a slight benefit. The only trial that shows a clear advantage for misonidazole was the head and neck cancer trial performed in Denmark, the largest single trial performed.

**Figure 26.4** The structure of misonidazole, etanidazole, and nimorazole, the three compounds most widely used in clinical trials. Misonidazole and etanidazole are 2-nitroimidazoles; nimorazole is a 5-nitroimidazole. Misonidazole and etanidazole are equally active as radiosensitizers, but etanidazole is less neurotoxic because it has a shorter half-life and is hydrophilic. Nimorazole is less active but very much less toxic than either misonidazole or etanidazole, so that larger doses are tolerable.

**Figure 26.5** Survival data for aerated and hypoxic Chinese hamster cells x-irradiated in various concentrations of misonidazole (Ro-07-0582). At a concentration of 10 mM of this drug, the radiosensitivity of hypoxic cells approaches that of aerated cells. The response of aerated cells is not affected by the drug at all. (Adapted from Adams GE, Flockhart IR, Smithen CE, et al. Electron-affinic sensitization. VII. A correlation between structures, one-electron reduction potentials, and efficiencies of nitroimidazoles as hypoxic cell radiosensitizers. Radiat Res. 1976;67:9–20, with permission.)
This interesting result is shown in Figure 26.7. Other subgroups, including patients with cancer of the larynx, showed no benefit from the addition of the sensitizer.

When misonidazole was used in the clinic, the dose-limiting toxicity was found to be peripheral neuropathy that progressed to central nervous system toxicity if drug administration was not stopped. This toxicity prevented use of the drug at adequate dose levels throughout multifraction.

**FIGURE 26.6** Proportion of mouse mammary tumors controlled at 150 days as a function of x-ray dose for a single treatment. The right curve represents x-rays only; the left curve refers to x-rays delivered after the administration of 1 mg/g body weight of misonidazole (Ro-07-0582). The enhancement ratio is the ratio of x-ray doses in the absence or presence of the drug that results in the control of 50% of the tumors; it has a value of 1.8. (Adapted from Sheldon PW, Foster JL, Fowler JF. Radiosensitization of C3H mouse mammary tumours by a 2-nitroimidazole drug. *Br J Cancer*. 1974;30:560–565, with permission.)

**FIGURE 26.7** Some results from the Danish head and neck cancer trial of misonidazole. Misonidazole produced a significant improvement of tumor control by radiotherapy only for males with tumors of the pharynx and depended on hemoglobin status. (Data from Dr. Jens Overgaard.)
regimens. The disappointing results obtained with misonidazole in the clinic must be attributed largely to the fact that doses were limited to inadequate levels because of this toxicity.

**Etanidazole and Nimorazole**

Spurred by the promise of misonidazole in the laboratory compared with its failure in the clinic, efforts were made to find a better drug. The compound chosen for the next series of clinical trials in the United States was etanidazole, a 2-nitroimidazole, the structure of which is also shown in Figure 26.4. This compound equals misonidazole as a sensitizer but is less toxic, so that doses could be increased by a factor of 3. The lower neurotoxicity is a function of a shorter half-life *in vivo* plus a lower partition coefficient, so that it penetrates poorly into nerve tissue and does not cross the blood–brain barrier. Controlled clinical trials by the RTOG in the United States and a multicenter consortium in Europe showed no benefit when etanidazole was added to conventional radiation therapy.

Nimorazole is a 5-nitroimidazole (structure also shown in Fig. 26.4); it is less effective as a radiosensitizer than either misonidazole or etanidazole, but it is much less toxic, so that very large doses could be given. In a Danish head and neck cancer trial, this compound produced a significant improvement in both locoregional control and survival compared with radiotherapy alone in patients with supraglottic and pharyngeal carcinoma. It is surprising that nimorazole has not been used elsewhere.

The development of nitroimidazoles is illustrated by the following:

- Metronidazole
- ↓
- Misonidazole: more active, toxic; benefit in subgroups
- ↓
- Etanidazole: less toxic, no benefit
- ↓
- Nimorazole: less active, much less toxic; benefit in head and neck cancer

**Overgaard’s Meta-analysis of Clinical Trials Addressing the Problem of Hypoxia**

For more than 3 decades, an enormous effort has been expended in an attempt to overcome the perceived problem of hypoxia. Dozens of clinical trials have been performed, most of which have been inconclusive or have shown results with borderline significance. Overgaard and colleagues performed a meta-analysis in which the results of all the trials were combined and analyzed together. They identified 10,602 patients treated in 82 randomized clinical trials involving hyperbaric oxygen, chemical sensitizers, carbogen breathing, or blood transfusions. Tumor sites included the bladder, uterine cervix, central nervous system, head and neck, and lung.

Overall, local tumor control was improved by 4.6%, survival by 2.8%, and the complication rate increased by only 0.6%, which was not statistically significant. The largest number of trials involved head and neck tumors, which also showed the greatest benefit. It was also concluded that the problem of hypoxia may be marginal in most adenocarcinomas and most important in squamous cell carcinomas.

**Nicotinamide and Carbogen Breathing**

Hypoxic cell radiosensitizers, such as the nitroimidazoles, were designed primarily to overcome chronic hypoxia, that is, diffusion-limited hypoxia resulting from the inability of oxygen to diffuse farther than about 100 \( \mu \text{m} \) through respiring tissue. As explained in Chapter 6, however, there is another form of hypoxia known as *acute hypoxia*, in which local regions of hypoxia are caused by the intermittent closing down of blood vessels. Nicotinamide, a vitamin B\(_3\) analogue, prevents these transient fluctuations in tumor blood flow that lead to the development of acute hypoxia, at least in mouse tumors.

A combination of nicotinamide to overcome acute hypoxia and carbogen breathing to overcome chronic hypoxia is the basis of the accelerated radiotherapy, carbogen, and nicotinamide (ARCON) trials under way at several European centers. The trials are also accelerated and hyperfractionated to avoid tumor proliferation and damage to late-responding normal tissues. Here, briefly, is a summary of the ARCON treatment:

- Accelerated, to overcome proliferation
- Hyperfractionated, to spare late-responding normal tissues
- Carbogen breathing, to overcome chronic hypoxia
Chapter 26 • The Biology and Exploitation of Tumor Hypoxia

**HYPOXIC CYTOTOXINS**

An alternative approach to designing drugs that preferentially radiosensitize hypoxic cells is to develop drugs that selectively kill hypoxic cells. It was pointed out at an early stage that the greater reductive environment of tumors might be exploited by developing drugs that are reduced preferentially to cytotoxic species in the hypoxic regions of tumors. Five classes of agents in this category are known:

1. Quinone antibiotics
2. Nitroaromatic compounds
3. Benzotriazine di-N-oxides
4. Dinitrobenzamide modified nitrogen mustard
5. 2-nitroimidazole attached to dibromo isophoramide

Mitomycin C is an example of the first class and has been used as a chemotherapy agent, active against hypoxic cells, for many years. The aerated–hypoxic differential, however, is relatively small for these compounds. Examples of the second class of compounds include dual-function agents developed previously by Adams and his group at the Medical Research Council Radiobiology Unit in England. Normal tissue toxicity prevented the trial of these compounds in the clinic. The lead compound of the third class, tirapazamine, shows highly selective toxicity toward hypoxic cells both in vitro and in vivo. The third and fourth members of this class represent a new strategy of modifying a cytotoxic chemotherapy moiety to make it hypoxia activated.

**Tirapazamine**

Figure 26.8 shows survival curves for Chinese hamster cells treated with graded concentrations of tirapazamine. Note the hypoxic/oxic cytotoxicity ratio of about 100. The hypoxic/oxic cytotoxicity ratio is not as large (about 20) in cell lines of human origin, presumably reflecting a different spectrum, or different levels, of enzymes.

The effect of x-rays plus tirapazamine is evidently much greater than additive (what would be expected from independent cell killing of the two agents). Experimentally, this was shown by treating tumors with x-rays alone, tirapazamine alone, or a combination of both agents (Fig. 26.9). The radiation schedule consisted of eight 2.5-Gy fractions designed to mimic, as far as possible, a clinical radiation therapy protocol. The combination of drug and radiation is highly effective, with the time sequence of drug before radiation slightly more effective than the reverse. In parallel experiments using the same x-ray and drug protocols, skin reactions were scored, and no radiosensitization or additive cytotoxicity was observed by the addition of the tirapazamine to the radiation treatments. This substantiates the tumor selectivity of the radiation enhancement.

Similar results were obtained in four different mouse tumors that differed significantly in their hypoxic fractions. The observed interaction between x-rays and tirapazamine results largely from the selective hypoxic toxicity of the drug. This does not totally explain the observations, however. It appears that tirapazamine can act as an aerobic radiosensitizer of cells exposed to the

**FIGURE 26.8** Dose-response curves for Chinese hamster cells exposed for 1.5 hours to graded concentrations of SR 4233 (tirapazamine) under aerated and hypoxic conditions. Cells deficient in oxygen are killed preferentially. The hypoxic cytotoxicity ratio (defined as the ratio of drug concentrations under aerated and hypoxic conditions required to produce the same cell survival) is variable between different cell lines. For the Chinese hamster cells shown, the ratio is about 100; for cells of human origin, the ratio is somewhat smaller, closer to 20. Tirapazamine is an organic nitroxide synthesized by Stanford Research International. Its structure is shown in the inset. (Courtesy of Dr. J. Martin Brown.)
drug under hypoxic conditions before or after the aerobic irradiation. This latter mechanism would be important for intermittent or acute hypoxia, that is, where a given region of a tumor cycles between aerated and hypoxic conditions.

Clinical Trials with Tirapazamine and New Bioreductive Drugs

Despite the extensive laboratory data in vitro and in vivo showing the greater-than-additive effect of the combination of x-rays with tirapazamine, little has been done clinically to combine tirapazamine with radiation therapy, except for one RTOG phase II trial. The reason may be the side effects of nausea and severe muscle cramping.

The situation adding tirapazamine to chemotherapy protocols is more advanced. One of the largest phase III trials in head and neck cancer has been completed comparing tirapazamine, cisplatin, and radiation to cisplatin and radiation for advanced squamous cell carcinoma of the head and neck (TROG 02.02, HeadSTART). In this trial, previously untreated stage III or IV squamous cell carcinomas of the head and neck were randomly assigned to receive 70 Gy of radiotherapy concurrently with cisplatin alone or cisplatin in combination with tirapazamine. A total of 861 patients were enrolled from 89 sites worldwide. The 2-year survival analysis indicated that there was no significant differences in either failure-free survival, time to locoregional progression, or quality of life.

Various explanations have been brought forth to explain this result, including failure to stratify patients based on whether their tumor was hypoxic, poor radiotherapy at some sites, and insufficient levels of hypoxia in head and neck tumors. Hay, Wilson, and Denny in Auckland have developed a series of novel tricyclic triazine-di-N-oxides (TTOs) that are related to tirapazamine, but are superior to it in regard to in vivo activity. Furthermore, because the same reductase that activates these TTOs is also responsible for activating the hypoxia marker EF5, the potential to combine the two in clinical trials and properly stratify patients could be achieved.

During the last 5 years, several other targeted therapeutics have entered clinical testing: PR-104, a novel hypoxia-activated DNA crosslinking agent, and TH-302, a hypoxia-activated dibromo isophoramide mustard. Both of them has passed phase I clinical testing and are being examined in phase II trials to determine their best clinical indication.

TARGETING TUMOR METABOLISM TO KILL HYPOXIC CELLS

Tumor cells exhibit altered metabolism that is highly glycolytic at the expense of oxidative phosphorylation. This shift from the normal oxidative metabolism gives tumor cells a growth advantage in vivo. Denko et al. have identified the HIF-1–pyruvate dehydrogenase kinase 1 (PDK1) axis as one of the major regulators responsible for this shift. HIF-1 induction of PDK1 results in phosphorylation and inactivation of pyruvate dehydrogenase (PDH), the first committed step for pyruvate entry into the mitochondria. Hypoxic tumors reduce mitochondrial function to conserve oxygen and, as a result, have a compensatory increase in glycolysis. Inhibition of PDK increases tumor hypoxia (Fig. 26.10).
Dichloroacetate (DCA) was found more than 30 years ago to be a safe inhibitor of PDKs that has been used to treat patients with inborn errors of metabolism. DCA increases mitochondrial metabolism in these patients by inhibiting the activity of PDKs. DCA in preclinical tumor models increases mitochondrial function and “reprograms” tumor metabolism to be more oxidative, like normal tissue. At the molecular level, DCA can inhibit PDH phosphorylation, reduce glucose uptake, increase mitochondrial function, and inhibit tumor growth in model tumors grown in nude mice. Figure 26.11 shows that DCA treatment can enhance the efficacy of hypoxic cytotoxins, such as tirapazamine in tumor xenografts, without causing depression of hematologic parameters. Clinical trials investigating the efficacy of DCA to alter tumor metabolism and increase tumor hypoxia have just begun, using positron emission tomography imaging of radiolabeled fluorodeoxyglucose ($^{18}$F) (FDG) and EF5 to quantify metabolic changes in the tumor.

**FIGURE 26.10** The effect of inhibiting PDK activity with dichloroacetate (DCA) on tumor hypoxia using an HIF-driven luciferase reporter gene. Treatment with DCA induces a transient increase in hypoxia as detected by increased luciferase activity. (Adapted from Cairns RA, Papandreou I, Sutphin PD, et al. Metabolic targeting of hypoxia and HIF1 in solid tumors can enhance cytotoxic chemotherapy. *Proc Natl Acad Sci USA.* 2007;104(22):9445–9450, with permission.)

**FIGURE 26.11** Demonstration of the increased efficacy of tirapazamine killing in transplanted tumors by inhibiting PDK. The graph shows a tumor growth delay experiment. The key point lies in the sequencing of PDK inhibition and tirapazamine. If tirapazamine is administered to the animal first, followed by a PDK inhibitor, little increase in cell killing beyond administering tirapazamine alone is observed. In contrast, if a PDK inhibitor is administered first, and then tirapazamine is administered subsequently, a significant increase in tumor growth delay is observed because tumor hypoxia has been increased. (Adapted from Cairns RA, Papandreou I, Sutphin PD, et al. Metabolic targeting of hypoxia and HIF1 in solid tumors can enhance cytotoxic chemotherapy. *Proc Natl Acad Sci USA.* 2007;104(22):9445–9450, with permission.)
Etanidazole is less toxic than misonidazole, and three times larger doses are tolerated. However, clinical trials in Europe and the United States showed no advantage of combined radiotherapy and etanidazole over radiotherapy alone.

Nimorazole is less active as a radiosensitizer, but so much less toxic than either misonidazole or etanidazole that much larger doses can be given. A benefit was shown for adding nimorazole to conventional radiotherapy for head and neck cancer (Danish head and neck cancer trial).

A meta-analysis of all 82 trials of hyperbaric oxygen and hypoxic cell radiosensitizers showed a 4.6% gain in local tumor control.

**Hypoxia-Inducible Factor**

- At both the molecular and cellular levels, in physiologic and pathologic settings, oxygen homeostasis is controlled primarily through the function of a heterodimeric transcription factor, the HIF-1.
- The HIF-α subunits are oxygen regulated through hydroxylation, binding to VHL and degradation in the proteasome.
- Loss of certain tumor suppressor genes such as VHL and PTEN can cause HIF to become active under normoxic conditions.
- HIF plays various critical roles in tumors, regulating metabolism, angiogenesis, and metastasis.
- HIF can also influence the response of tumors to radiotherapy by increasing the resistance of the vasculature through the increased expression of proangiogenic factors and by promoting vasculogenesis through the recruitment of BM-derived cells.

**The Unfolded Protein Response**

- The UPR responds to an accumulation of misfolded proteins by stopping protein translation through the activation of PERK and through the induction of XBP-1 and chaperone proteins that function in the ER to assist in folding, trafficking, and secretion of proteins.
- Inhibition of PERK or XBP-1 activities result in decreased tumor growth.

**Radiosensitizing Hypoxic Cells**

- Hypoxic cell radiosensitizers increase the radiosensitivity of hypoxic cells but not aerated cells. The differential effect on tumors is based on the presence of hypoxic cells in tumors but not in normal tissues.
- Sensitization is a free-radical process. The sensitizer mimics oxygen by “fixing” damage produced by free radicals.
- Misonidazole, the first compound used widely, sensitizes cells in culture as well as in both animal and human tumors.
- Doses of misonidazole that can be used clinically are limited to suboptimal levels by peripheral neuropathy. Only 1 of 30 or so clinical trials showed an advantage for misonidazole.

- Etanidazole is less toxic than misonidazole, and three times larger doses are tolerated. However, clinical trials in Europe and the United States showed no advantage of combined radiotherapy and etanidazole over radiotherapy alone.
- Nimorazole is less active as a radiosensitizer, but so much less toxic than either misonidazole or etanidazole that much larger doses can be given. A benefit was shown for adding nimorazole to conventional radiotherapy for head and neck cancer (Danish head and neck cancer trial).
- A meta-analysis of all 82 trials of hyperbaric oxygen and hypoxic cell radiosensitizers showed a 4.6% gain in local tumor control.

**BIBLIOGRAPHY**


Horsman MR, Chaplin DJ, Overgaard J. Combination of nicotinamide and hyperthermia to eliminate radioresistant...


Alice: There’s no use trying—one can’t believe impossible things.

The Queen: I dare say you haven’t had much practice. Why, sometimes I’ve believed as many as six impossible things before breakfast.

—Alice in Wonderland (Lewis Carroll)

This chapter is included after much thought and some equivocation. It was written in response to numerous requests that chemotherapeutic agents be compared and contrasted with radiation from the perspective of the experimental biologist. Many of the techniques and concepts used in chemotherapy were developed initially by radiation biologists, including quantitative tumor assay systems, the concept of cell cycle, sensitivity changes through the cell cycle, and, particularly, population kinetics. The term *growth fraction*, for example, was coined by a radiation biologist, but never assumed the importance in radiotherapy that it has in chemotherapy.

The study of chemotherapeutic agents in the laboratory, as well as in the clinic, is vastly more complicated than the study of ionizing radiations. For example, *dose* is more difficult to define or to measure, and time of drug exposure is a critical parameter. Variations in sensitivity through the cell cycle are more dramatic for chemicals than for radiation, assuming essentially an all-or-nothing effect for some agents; there are many more factors involving the microenvironment that can influence cellular response.

The term *chemotherapy* was coined by Paul Ehrlich around the turn of the 20th century to describe the use of chemicals of known composition for the treatment of parasites. Ehrlich synthesized an organic arsenic compound that was effective against trypanosome infections and rabbit syphilis. This was the first synthetic chemical effective in the treatment of parasitic disease and was rather optimistically named *salvarsan*, which roughly translates to “the savior of mankind.” The next milestone was the discovery and clinical use of penicillin in the early years of World War II. Alkylating agents had been developed as military weapons by both belligerents in World War I, but it was an explosion in Naples harbor and the exposure of seamen to these agents during World War II that led to the observation that these agents cause marrow and lymphoid hypoplasia. As a result, they were first tested in humans with Hodgkin disease in 1943 at Yale University.

It has long been known beyond doubt that a single chemotherapeutic drug, used in the appropriate schedule, can cure patients with certain rapidly proliferating cancers. The initial demonstration of this was the use of methotrexate to cure patients with choriocarcinoma and, later, the use of cyclophosphamide for Burkitt lymphoma.
The next major step forward was the use of combination chemotherapy in the treatment of acute lymphocytic leukemia in the early 1960s and, subsequently, in the treatment of Hodgkin disease, diffuse histiocytic lymphoma, and testicular cancer in the mid-1970s. These trials verified that multiple non–cross-resistant drugs with different dose-limiting normal tissue toxicities could be used effectively in combination to cure tumors that were not curable with a single agent. The principle of combination therapy was then extended to combined modality treatment, in which chemotherapy was used in conjunction with surgery or radiotherapy, or both, to cure tumors such as pediatric sarcomas.

Today, various antineoplastic agents are used routinely in clinical oncology. Drug-induced cures are claimed for choriocarcinoma, acute lymphocytic leukemia of childhood, other childhood tumors, Hodgkin disease, certain non–Hodgkin lymphomas, and some germ cell tumors of the testes. Other evidence suggests that chemotherapeutic agents given in an “adjuvant” setting for clinically inapparent micrometastatic disease may prolong disease-free survival and possibly effect cure of breast cancer and osteogenic sarcoma.

There are about 13 types of cancer for which cures are claimed by chemotherapy; this accounts for about 10% of all cancers. For comparison, the proportion of cancer patients cured by radiation therapy often is claimed to be about 12.5%. This comes from the so-called 1/2 × 1/2 × 1/2 rule; that is, one-half of all cancer patients receive radiation therapy, one-half of those treated are treated with intent to cure, and one-half of those treated definitively are cured.

The bad public image of chemotherapy relates in large part from the toxicity to normal tissue resulting from multidrug protocols used to induce remissions and achieve tumor cure. “The dose makes the poison” was the advice of Paracelsus, the 16th-century German-Swiss physician and alchemist who established the role of chemistry in medicine. In other words, anything powerful enough to help also has the power to harm. In the past, the lack of tumor-specific agents carried the burden of damage to self-renewing normal tissues, such as the gut and bone marrow. There is hope that the situation is improving with the development of targeted therapies and the new concept of synthetic lethality. (Both will be discussed later in this chapter.)

### BIOLOGIC BASIS OF CHEMOTHERAPY

Most anticancer drugs work by affecting DNA synthesis or function, and they usually do not kill resting cells unless such cells divide soon after exposure to the drug. Consequently, the effectiveness of anticancer drugs is limited by the growth fraction of the tumor—that is, by the fraction of cells in active cycle. Rapidly growing neoplasia with a short cell cycle, a large proportion of cells in S phase, and, therefore, a large growth fraction are more responsive to chemotherapy than large tumor masses in which the growth fraction is small. There is a strong tendency for growth fraction to decrease as tumor size increases, at least in experimental animal tumors.

Agents that are mainly effective during a particular phase of the cell cycle, such as S phase or M phase, are said to be **cell cycle specific**, or **phase specific**. Those whose action is independent of the position of the cell in the cycle are said to be **cell cycle nonspecific** or **phase nonspecific**. Figure 27.1 illustrates two contrasting cell cycle specific drugs. Cis-platinum compounds produce interstrand crosslinks and thus inhibit DNA synthesis; this occurs in S phase. Taxanes bind to microtubules and, by enhancing their stability and preventing disassembly, adversely affect their function. They act as mitotic inhibitors, blocking cells in the G2/M phase of the cell cycle.

Agents that are most effective against cells in S phase are relatively ineffective against cell populations that turn over slowly and have large proportions of dormant cells. On the other hand, the action of alkylating agents and other drugs interacting primarily with macromolecular DNA is largely independent of the phase of the cell cycle, and they may be effective against tumors with relatively low proliferative activity.

The other side of the coin is that the selective normal tissue toxicity of anticancer drugs is reflected in cytotoxic effects on stem cells of the intestinal epithelium or hematopoietic stem cells, which have high growth fractions.

Although many clinical oncologists claim that their thinking has been influenced by research on tumor growth kinetics, it is hard to...
fall into any of these classes. This includes the platinum compounds, procarbazine, the vinca alkaloids, the taxanes, and the newest of all, the topoisomerase inhibitors and “targeted therapy” agents that target a specific pathway that may be elevated or vulnerable in some tumor cells.

An attempt to summarize the classification of chemotherapeutic drugs is presented in Table 27.1. A few of the most commonly used agents are described briefly, with emphasis on their characteristics and mechanism of action and comments on the extent to which they interact with radiation. A thorough discussion of their clinical usefulness is outside the scope of this book.

The effectiveness of at least some chemotherapeutic agents is dependent on the presence or absence of molecular oxygen, in much the same way as x-rays. This is not surprising, at least for drugs whose action is mediated by free radicals.

**Alkylating Agents**

The alkylating agents are highly reactive compounds with the ability to substitute alkyl groups for hydrogen atoms of certain organic compounds, including DNA. There are five classes of alkylating agents:

1. Nitrogen mustard derivatives, such as cyclophosphamide, chlorambucil, and melphalan.
2. Ethylenimine derivatives, such as thiotepa.
3. Alkyl sulfonates, such as busulfan.
4. Triazine derivatives, such as dacarbazine.
5. Nitrosoureas, including bischloroethyl-nitrosourea (BCNU) (carmustine), chloroethyl-cyclohexyl-nitrosourea (CCNU) (lomustine), and methyl CCNU.

Most of these drugs contain more than one alkylating group and therefore are considered polyfunctional alkylating agents. The nitrosoureas and dacarbazine have mechanisms and cytotoxicity over and above their ability to alkylate nucleic acids. As a class, alkylating agents are considered cell cycle nonspecific.

Nitrogen mustard is the prototype for three other useful alkylating agents: cyclophosphamide, chlorambucil, and melphalan. Nitrogen mustard given intravenously interacts rapidly with cells in vivo, producing its primary effect in seconds or minutes. By contrast, cyclophosphamide (Cytoxan) is inert until it undergoes biotransformation in the liver. Disappearance
### Table 27.1 Chemotherapeutic Agents

<table>
<thead>
<tr>
<th>Compound Class</th>
<th>Chemotherapeutic Agent</th>
<th>Diseases for which Drugs Are Useful (Indications)</th>
<th>Unique/Major Toxicity</th>
<th>Targeted Pathway/ Receptor (Mechanism of Action)</th>
<th>Synergy with Radiation</th>
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<tbody>
<tr>
<td>Alkylating agents</td>
<td>Busulfan (Myleran)—BSF</td>
<td>Regular-dose therapy in chronic myelogenous leukemia (CML) (FDA approved) and polycythemia vera. High-dose therapy in bone-marrow transplant.</td>
<td>Myelosuppression pulmonary toxicity especially with total body irradiation (TBI) when used in high doses for bone marrow transplant.</td>
<td>DNA alkylation (i.e., interstrand crosslinks).</td>
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<tr>
<td></td>
<td>Carboplatin (Paraplatin)—Carbo, CBDCA</td>
<td>FDA approved for ovarian cancer, and used extensively in testicular cancer; squamous cell cancers of the head, neck, and cervix; and lung cancer.</td>
<td>Myelosuppression, especially thrombocytopenia, is dose limiting.</td>
<td>Produces intrastrand and interstrand crosslinks in DNA via association bonds with the platinum molecule, leading to DNA strand breakage during replication.</td>
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<td></td>
<td>Carmustine (BiCNU)—BCNU, Bischloronitrosourea</td>
<td>FDA approved for brain tumors, multiple myeloma, Hodgkin disease, and lymphoma.</td>
<td>Myelosuppression, especially thrombocytopenia, which is slow in onset and cumulative, is dose limiting. Interstitial lung disease, including fibrosis, is rare but can occur with any dose. Weak synergy with radiation therapy (RT).</td>
<td>Cell cycle non-specific crosslinking mechanism. Penetrates blood–brain barrier.</td>
<td>+</td>
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<tr>
<td></td>
<td>Chlorambucil (Leukeran)</td>
<td>FDA approved for chronic lymphocytic leukemia (CLL) and low-grade lymphomas. Also used for Waldenström macroglobulinemia, multiple myeloma, hairy cell leukemia, and rarely in some solid tumors.</td>
<td>Myelosuppression is dose limiting and universal, and it can be cumulative.</td>
<td>Cell cycle non-specific crosslinking mechanism.</td>
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(Continued)
### TABLE 27.1 Chemotherapeutic Agents (Continued)

<table>
<thead>
<tr>
<th>Compound Class</th>
<th>Diseases for which Drugs Are Useful (Indications)</th>
<th>Unique/Major Toxicity</th>
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<th>Synergy with Radiation</th>
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<tr>
<td>Cyclophosphamide (Cytoxan, Neosar)—CTX, CPM, Cy (prototypical alkylator)</td>
<td>FDA approved for many malignancies and used for even more. Most commonly used for breast carcinoma, non–Hodgkin lymphoma, ovarian carcinoma, and testicular cancer.</td>
<td>Myelosuppression is dose limiting, with leukopenia being most significant. Hemorrhagic cystitis common with doses more than 2 g/m².</td>
<td>Cell cycle non-specific crosslinking Prodrug activated in liver.</td>
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<tr>
<td>Dacarbazine (DTIC-Dome)—DTIC, DIC, Imidazole, Carboxamide</td>
<td>FDA approved for the treatment of malignant melanoma and Hodgkin disease, and also used for adult sarcomas and neuroblastoma.</td>
<td>Myelosuppression is dose limiting. Nausea and vomiting are severe without aggressive antiemetic therapy. Strong synergy with RT.</td>
<td>Methylates guanine bases preferentially; cell cycle non-specific.</td>
<td>+++</td>
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<tr>
<td>Ifosfamide (Ifex)</td>
<td>FDA approved for the treatment of recurrent germ cell tumors. Used for many other tumor types, including adult sarcomas, lymphoma, Hodgkin’s disease, breast cancer, and ovarian cancer.</td>
<td>Myelosuppression, hemorrhagic cystitis, and central nervous system (CNS) toxicity are all fairly common and can be dose limiting. Hemorrhagic cystitis can largely be prevented by coadministration of the uroprotective agent mesna.</td>
<td>Non-specific cell cycle alkylating agent</td>
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<tr>
<td>Lomustine (CeeNU)—CCNU</td>
<td>FDA approved for primary brain tumors and Hodgkin disease. Also used in melanoma, multiple myeloma, other lymphomas, and breast cancer.</td>
<td>Myelosuppression, especially thrombocytopenia, is dose limiting and tends to be cumulative. Pulmonary fibrosis can occur with long-term administration. Weak synergy with RT.</td>
<td>Cell cycle non-specific. Penetrates blood–brain barrier.</td>
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<tr>
<td>Drug</td>
<td>FDA Approval Details</td>
<td>Side Effects</td>
<td>Mechanism</td>
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<tr>
<td>Mechlorethamine</td>
<td>FDA approved for various hematologic malignancies and solid tumors, but generally used less in the last decade. Still used for Hodgkin disease and topically for cutaneous T-cell lymphoma.</td>
<td>This drug is a powerful vesicant, so optimal extravasation precautions and rapid infusion are a must. Myelosuppression is expected and also often dose limiting. Secondary leukemia.</td>
<td>Cell cycle non-specific alkylation agent.</td>
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<td>(Mustargen) Nitrogen mustard, HN2</td>
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<tr>
<td>Melphalan</td>
<td>Used primarily for multiple myeloma, but also FDA approved for ovarian carcinoma. Could also be useful in high-dose chemotherapy/transplant settings and in regional perfusion of extremities for melanoma and sarcoma.</td>
<td>Myelosuppression is expected and is dose limiting. Recovery can be prolonged, and effects can be cumulative. Secondary leukemia.</td>
<td>Cell cycle non-specific alkylation agent.</td>
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<tr>
<td>(Alkeran) L-PAM, L-phenylalanine mustard, L-sarcolysin</td>
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<tr>
<td>Oxaliplatin</td>
<td>FDA approved for metastatic colorectal cancer in combination with 5-fluorouracil/leucovorin. It has been used as a single agent in this disease and is being studied in other malignancies.</td>
<td>Neurotoxicity, in the form of a transient neuropathy with each dose and a persistent cumulative typical sensory polyneuropathy, is very common and dose limiting.</td>
<td>Cell cycle non-specific intrastrand and interstrand crosslinks with two strong platinum association bonds in the molecule.</td>
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<td>(Eloxatin)</td>
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<td>(Matulane) N-methylhydrazine</td>
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<td>Compound Class Chemotherapeutic Agent</td>
<td>Diseases for which Drugs Are Useful (Indications)</td>
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<tr>
<td>Temozolomide (Temodar)</td>
<td>FDA approved for treatment of recurrent high-grade astrocytomas. Used commonly for other gliomas and also for metastatic melanoma.</td>
<td>Myelosuppression is expected and dose limiting.</td>
<td>Penetrates blood–brain barrier. Oral alkylating agent. Prodrug</td>
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<tr>
<td>Atypical alkylating agents Cisplatin (Platinol)—cDDP, DDP, Cis-platinum, Cis-diammine-dichloroplatinum (II)</td>
<td>Used for almost every class of solid tumor and lymphoma. FDA approved for testicular and ovarian cancers and transitional cell carcinoma.</td>
<td>Nephrotoxicity is dose limiting for an individual dose, whereas neurotoxicity, especially painful peripheral neuropathy, is dose limiting for cumulative doses. Cumulative ototoxicity is also common. Intermediate synergy with RT.</td>
<td>Produces intrastrand and interstrand crosslinks in DNA via association bonds with the platinum molecule, leading to DNA strand breakage during replication.</td>
<td>++</td>
</tr>
<tr>
<td>Antibiotics Bleomycin (Blenoxane)—Bleo</td>
<td>FDA approved for germ cell tumors, Hodgkin disease, and squamous cell cancers. Used off-label for melanoma, ovarian cancer, and Kaposi sarcoma. Also used as a sclerosing agent for malignant pleural or pericardial effusions.</td>
<td>Pulmonary toxicity, including reversible and irreversible fibrosis, is dose limiting. Strong synergy with RT.</td>
<td>Causes DNA strand breaks directly in normal and neoplastic cells.</td>
<td>+++</td>
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<td>Drug</td>
<td>FDA Approval</td>
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<tr>
<td>Dactinomycin</td>
<td>Wilms tumor, Ewing sarcoma, rhabdomyosarcoma, uterine carcinoma, germ cell tumors, and sarcoma botryoides, also used for other sarcomas, melanoma, acute myeloid leukemia, ovarian cancer, and trophoblastic neoplasms.</td>
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<tr>
<td>Doxorubicin</td>
<td>Various cancers and used for many more. Most commonly used for breast carcinoma, adult sarcomas, pediatric solid tumors, Hodgkin disease, non–Hodgkin lymphomas, and ovarian cancer.</td>
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</table>

This drug is a moderate vesicant. Myelosuppression is dose limiting. Nausea, vomiting, skin erythema, acneiform lesions, and hyperpigmentation are common. Strong synergy with RT.

Inhibits transcription by complexing with DNA and prevents elongation by RNA polymerase.

Intercalating agent.

Doxorubicin is a potent vesicant, and extravasation precautions are a must. Myelosuppression is universal and usually dose limiting with each individual cycle. Cardiotoxicity is common and can be dose limiting, although usually subclinical. Chronic, cumulative cardiomyopathy is expected when total dose exceeds 400–500 mg/m². Strong synergy with RT. Recall skin reactions that correspond to prior RT treatment fields may develop, and can be severe. Concurrent RT or initiation of RT within 2 weeks of administration of doxorubicin should be avoided.
<table>
<thead>
<tr>
<th>Compound Class</th>
<th>Chemotherapeutic Agent</th>
<th>Diseases for which Drugs Are Useful (Indications)</th>
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</thead>
<tbody>
<tr>
<td>Mitomycin C (Mutamycin)</td>
<td>FDA approved for adenocarcinomas of the stomach and pancreas. Also used commonly in breast cancer and lung cancer.</td>
<td>Mitomycin C is a vesicant; extravasation precautions are a must. Myelosuppression is expected and is dose limiting, with a white blood cell nadir at 4 weeks and full recovery at 6–7 weeks. Strong synergy with RT.</td>
<td>Inhibits DNA and RNA synthesis.</td>
<td>++ +</td>
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<tr>
<td>Antimetabolites 5-Fluorouracil (Adrucil, Efudex)—5-FU</td>
<td>FDA approved for colon, rectum, gastric, pancreas, and breast carcinomas and used for a wide range of other neoplasms in combination regimens. Used for intrahepatic arterial infusion for liver metastases from gastrointestinal (GI) tumors and also used topically for various cutaneous neoplasms and disorders.</td>
<td>GI toxicities, primarily mucositis for bolus injection and diarrhea for prolonged infusions, are dose limiting. Rare patients with dihydropyrimidine dehydrogenase deficiency have excess GI toxicity. Dermatitis and other cutaneous toxicities, including hand–foot syndrome, are common. Intermediate synergy with RT.</td>
<td>Inhibitor of thymidylate synthase; partially cell cycle dependent.</td>
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<tr>
<td>Capecitabine (Xeloda) (antimetabolite prodrug)</td>
<td>FDA approved for metastatic breast cancer and metastatic colorectal cancer. Used also in head and neck squamous cell cancer.</td>
<td>Myelosuppression and palmar-plantar erythrodysesthesia are dose limiting. Diarrhea, fatigue, stomatitis, and hyperbilirubinemia are uncommon. Intermediate synergy with RT.</td>
<td>Prodrug of 5-Fluorouracil</td>
<td>++</td>
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<tr>
<td>Drug</td>
<td>Indications</td>
<td>Toxicities</td>
<td>Mechanism</td>
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<tr>
<td>Cytarabine (Cytosar-U)</td>
<td>Acute myelogenous leukemia (AML), acute lymphoblastic lymphoma (ALL), and non–Hodgkin lymphoma. Intrathecal use in acute leukemia.</td>
<td>Myelosuppression, often severe and prolonged, is dose limiting. Neurologic toxicity, mostly central with ataxia being predominant, is common and usually mild, but it is dose dependent and could leave permanent dysfunction. It is more common with intrathecal administration.</td>
<td>Incorporated into DNA during replication, leading to strand termination; S phase specific. Penetrates blood–brain barrier.</td>
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<tr>
<td>Fludarabine (Fludara)</td>
<td>FDA approved for the treatment of CLL. Also used for low-grade lymphomas and for AML.</td>
<td>Neurotoxicity, including cortical blindness, confusion, somnolence, coma, and demyelinating lesions, is dose limiting, but the lower doses conventionally used rarely produce these side effects.</td>
<td>Purine analog. Inhibits DNA polymerase. Only Partially cell cycle specific.</td>
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<tr>
<td>Gemcitabine (Gemzar)</td>
<td>FDA approved for advanced pancreatic adenocarcinoma, non–small cell lung cancer (NSCLC), and metastatic breast cancer; extensively used in bladder cancer also.</td>
<td>Myelosuppression, including anemia, is mild but dose limiting. Strong synergy with RT even at low doses of drug.</td>
<td>A nucleoside analogue that exhibits S-phase-specific cytotoxicity. Inhibits DNA synthesis.</td>
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<tr>
<td>Hydroxyurea (Hydrea)</td>
<td>FDA approved for CML; commonly used for other myeloproliferative disorders and also used occasionally for metastatic melanoma, refractory ovarian carcinoma, and squamous cell carcinoma of the cervix and head and neck.</td>
<td>Myelosuppression is common and dose limiting. Other toxicities include rash, headache, fever, and hyperuricemia. Nausea and vomiting are uncommon. Liver toxicity and serious neurologic toxicity are rare. Weak synergy with RT.</td>
<td>Inhibitor of ribonucleotide reductase, which converts nucleotides to the deoxyribose forms for DNA synthesis; cell cycle dependent. Penetrates blood–brain barrier.</td>
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(Continued)
### Chemotherapeutic Agents (Continued)

<table>
<thead>
<tr>
<th>Compound Class</th>
<th>Diseases for which Drugs Are Useful (Indications)</th>
<th>Unique/Major Toxicity</th>
<th>Targeted Pathway/Receptor (Mechanism of Action)</th>
<th>Synergy with Radiation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Methotrexate (Mexate, Folex, others)—MTX, Amethopterin</td>
<td>FDA approved for a wide spectrum of malignant and nonmalignant diseases. Most often used for acute leukemias, lymphomas, breast cancer, bladder cancer, squamous cell cancers, and sarcomas.</td>
<td>Myelosuppression is expected and is usually dose limiting. Renal toxicity is uncommon and usually reversible, but can be severe. Encephalopathy is rare with moderate to low-dose therapy but is more common with high doses, intrathecal administration, or concomitant CNS radiation. It can be severe and permanent. Drug should be administered prior to rather than concurrently or following brain RT when feasible to lessen risk of leukencephalopathy. Weak synergy with RT.</td>
<td>Interferes with nucleotide synthesis by inhibiting dihydrofolate reductase; cell cycle dependent. Penetrates blood-brain barrier at high intravenous doses.</td>
<td>+</td>
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<tr>
<td>Corticosteroids Prednisone (Deltasone, others)</td>
<td>FDA approved for various malignant and nonmalignant conditions. Used in oncology for lymphoid malignancies, for palliative care, and for management of side effects/toxicities.</td>
<td>Toxicity is mostly in the form of constitutional symptoms, including mood changes (depressive, anxious, or euphoric), insomnia, indigestion, enhanced appetite, weight gain, acne, and cushingoid features. Other side effects may be more serious but are less common.</td>
<td>Same as naturally occurring ones.</td>
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Hyperglycemia and increased stomach acid predisposing to ulceration occur acutely, whereas osteopenia, cataracts, skin atrophy, and adrenal insufficiency occur with prolonged use.

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<tr>
<th>Enzymes</th>
<th>FDA approved for ALL; also used in AML, late-stage CML, CLL, and non–Hodgkin lymphomas.</th>
<th>Hypersensitivity can be life threatening, requiring anaphylaxis precautions and a 2-unit test dose. Coagulopathy is common and requires monitoring. Lethargy, somnolence, fatigue, depression, and confusion are seen, as are pancreatitis and fever.</th>
<th>Catalyzes the hydrolysis of amino acid asparagine, which is an essential amino acid required by rapidly proliferating cells, to aspartic acid.</th>
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<tr>
<td>I-Asparaginase (Elspar)—Colaspase</td>
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<td>Steroidal progestational agents</td>
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### TABLE 27.1 Chemothapeutic Agents (Continued)

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<tr>
<th>Compound Class Chemotherapeutic Agent</th>
<th>Diseases for which Drugs Are Useful (Indications)</th>
<th>Unique/Major Toxicity</th>
<th>Targeted Pathway/Receptor (Mechanism of Action)</th>
<th>Synergy with Radiation</th>
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<tr>
<td><strong>Targeted therapy</strong>&lt;br&gt;Bevacizumab (Avastan)</td>
<td>FDA approved for combination with 5-FU for first- and second-line treatment of metastatic colorectal cancer; for combination with carboplatin and paclitaxel; for unresectable, locally advanced, recurrent, or metastatic non-squamous, non-small cell lung cancer; and for combination with interferon alpha for metastatic renal cell cancer.</td>
<td>Most toxicities related to inhibition of angiogenesis. GI perforation is the major life-threatening side effect. Treatment can also inhibit wound healing and may prevent surgical incisions to close, also leading to fatality in some instances. Bleeding in stomach, brain, nose, and vagina. Common side effects are nosebleeds, high blood pressure, headache, and inflammation.</td>
<td>Inhibits the function of vascular endothelial growth factor. Causes regression of existing tumor vessels. Makes existing tumor vessels more functional in supplying oxygen and chemotherapy. Inhibits new tumor vessel growth.</td>
<td>++</td>
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<tr>
<td><strong>Cetuximab</strong>&lt;br&gt;(Erbitux)</td>
<td>Colorectal cancer, lung cancer, head and neck cancer.</td>
<td>Self-limiting sterile, nonsuppurative acne-like skin rash is common. Resolves with cessation of drug. Patients exhibiting a grade 2 rash or above have better survival. Strong synergy with RT.</td>
<td>Blocks EGFR receptor dimerization and tyrosine kinase phosphorylation, which inhibits tyrosine kinase pathway signal transduction.</td>
<td>+++</td>
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<tr>
<td>Imatinib mesylate (Gleevec)</td>
<td>FDA approved for treatment of CML in the frontline setting, in accelerated phase, and in blast crisis. It is also approved for treatment of recurrent inoperable or metastatic gastrointestinal stromal tumors.</td>
<td>There is no definite dose-limiting toxicity of imatinib. Myelosuppression is significant in CML but mild in gastrointestinal stromal tumors. Hepatotoxicity is common but usually mild. Liver function tests should be monitored closely during therapy. Fluid retention is common but usually mild, as are nausea, vomiting, and diarrhea. Rash and fever are uncommon.</td>
<td>Specific receptor tyrosine kinase inhibitor, which selectively inhibits the tyrosine kinases of the bcr-abl, c-kit, and platelet-derived growth factor (PDGF) receptors.</td>
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<tr>
<td>Poly adenosine diphosphate-ribose polymerase (PARP) inhibitors</td>
<td>Soon to be FDA approved for BRCA-mutation-positive breast, ovarian, and prostate cancer. Doubling of progression-free survival for patients with triple negative breast cancer when inhibitors are combined with gemcitabine and carboplatin.</td>
<td>Dizziness, nausea, vomiting, diarrhea, lymphopenia, anemia, and fatigue.</td>
<td>PARP inhibitors block DNA base excision, leading to the collapse of replication forks and generation of DNA double-strand breaks. Cells possessing BRCA mutations cannot repair PARP inhibitor-induced DNA double-strand breaks by homologous recombination.</td>
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<tr>
<td>Rituximab (Rituxan)</td>
<td>FDA approved for relapsed or refractory low-grade or follicular, CD20-positive, B-cell lymphomas.</td>
<td>Fever, chills, and malaise are common during administration, even with premedication with acetaminophen and diphenhydramine. Other infusion-related symptoms include nausea, vomiting, flushing, urticaria, angioedema, hypotension, dyspnea, bronchospasm, fatigue, headache, rhinitis, and pain at disease sites. These symptoms are generally self-limited, improve with slowing of the infusion, and resolve after infusion. Short-lived myelosuppression, abdominal pain, and myalgia are uncommon. Arrhythmias and angina pectoris are rare.</td>
<td>Directed against the B-cell surface antigen CD20.</td>
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<tr>
<td><strong>Trastuzumab</strong> (Herceptin)</td>
<td>FDA approved for HER2/neu overexpressing metastatic or locally advanced breast cancer; has shown clinical benefit as a single agent and in conjunction with paclitaxel-based chemotherapy.</td>
<td>Common toxicities include acute fever, chills, nausea, vomiting, and headache. Trastuzumab seems to worsen leukopenia, anemia, and diarrhea when given with chemotherapy compared with chemotherapy alone. Also, trastuzumab could have uncommon acute cardiotoxicity, which might add to the more common anthracycline-induced cardiotoxicity; therefore, the use of trastuzumab with doxorubicin is not indicated by the FDA.</td>
<td>Directed against the HER2/neu growth factor receptor overexpressed on many invasive breast carcinomas; mechanism of action for clinical activity in breast cancer is unknown, but could be complement-mediated cell lysis, antibody-dependent cellular cytotoxicity, or induction of apoptosis.</td>
<td></td>
</tr>
<tr>
<td><strong>Taxanes</strong></td>
<td><strong>Docetaxel</strong> (Taxotere)</td>
<td>FDA approved for metastatic breast cancer and first- and second-line NSCLC. Clinical experience increasing in ovarian cancer and other epithelial neoplasms.</td>
<td>Myelosuppression is universal and dose limiting. Alopecia is also universal. Edema and fluid accumulation, including pleural effusions and ascites, are common and can be dose limiting. Fluid accumulation is partially preventable with corticosteroid treatment before and after each cycle of docetaxel. Mild sensory or sensorimotor neuropathy is common.</td>
<td>Inhibits the mitotic spindle apparatus by stabilizing tubulin polymers, leading to death of mitotic cells.</td>
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<tr>
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<tr>
<td>Paclitaxel (Taxol, Onxol)</td>
<td>FDA approved for salvage therapy in ovarian cancer and for breast cancer in both the metastatic and adjuvant setting. Used also in lung cancer, head and neck cancers, and bladder cancer.</td>
<td>Paclitaxel is an irritant or mild vesicant when extravasated into subcutaneous tissue. Myelosuppression, predominantly neutropenia, is expected and is dose limiting. Shorter infusions of the same dose produce less neutropenia. Mucositis is with very common, particularly with longer infusions. Peripheral neuropathy is common, usually mild, and increases with cumulative dose. Acute neuromyopathy is also common and occurs for several days after each dose. Hypersensitivity reactions to paclitaxel, including urticaria, wheezing, chest pain, dyspnea, and hypotension, are common but are reduced in frequency and severity by premedication with corticosteroids and H1 and H2 histamine receptor blockers (recommended regimen is dexamethasone 20 mg PO 12 and 6 hours prior to paclitaxel and diphenhydramine 50 mg and cimetidine 300 mg IV 30 minutes prior to paclitaxel). Weak synergy with RT.</td>
<td>Inhibits depolymerization of tubulin in the spindle apparatus, thereby inducing apoptosis in dividing cells.</td>
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Paclitaxel is an irritant or mild vesicant when extravasated into subcutaneous tissue. Myelosuppression, predominantly neutropenia, is expected and is dose limiting. Shorter infusions of the same dose produce less neutropenia. Mucositis is with very common, particularly with longer infusions. Peripheral neuropathy is common, usually mild, and increases with cumulative dose. Acute neuromyopathy is also common and occurs for several days after each dose. Hypersensitivity reactions to paclitaxel, including urticaria, wheezing, chest pain, dyspnea, and hypotension, are common but are reduced in frequency and severity by premedication with corticosteroids and H1 and H2 histamine receptor blockers (recommended regimen is dexamethasone 20 mg PO 12 and 6 hours prior to paclitaxel and diphenhydramine 50 mg and cimetidine 300 mg IV 30 minutes prior to paclitaxel). Weak synergy with RT.
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<tr>
<th><strong>Topoisomerase inhibitors</strong></th>
<th><strong>Etoposide</strong> (Vespid)—VP-16, Epipodophyllotoxin; also available as Etoposide Phosphate (Etopophos)</th>
<th>FDA approved for germ cell tumors and small cell lung cancer (SCLC). Also used for lymphomas, AML, brain tumors, NSCLC, and as high-dose therapy in the transplant setting for breast cancer, ovarian cancer, and lymphomas.</th>
<th>Myelosuppression, primarily leukopenia, is universal and dose limiting. Nausea and vomiting are common with PO administration but rare when the drug is given IV. Stomatitis and diarrhea are rare with normal doses but common with high doses. Secondary AML has been reported after etoposide.</th>
<th>Partially cell cycle specific. Topoisomerase II inhibitor.</th>
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<tr>
<td><strong>Irinotecan</strong> (Camptosar)—CPT-11</td>
<td>Irinotecan is FDA approved for refractory or recurrent metastatic colon cancer, and it has now been used in other malignancies, including lung cancer, ovarian cancer, and lymphoma.</td>
<td>Myelosuppression, primarily neutropenia, is common and dose limiting. Diarrhea is also common and can be dose limiting.</td>
<td>Partly cell cycle specific. Topoisomerase I inhibitor.</td>
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<tr>
<td><strong>Vinca alkaloids</strong></td>
<td><strong>Vinblastine</strong> (Velban, Velsar, others)—VLB, Vinca-leukoblastine</td>
<td>FDA approved for multiple hematologic and solid neoplasms. Most often used for Hodgkin disease, non-Hodgkin lymphoma, germ cell tumors, and breast cancer.</td>
<td>Vinblastine is a soft tissue vesicant, requiring extravasation precautions during administration. Myelosuppression, especially leukopenia, is expected and dose limiting.</td>
<td>Inhibitor of tubulin polymerization and thereby mitosis; G2 phase specific</td>
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<tr>
<td><strong>Vincristine</strong> (Oncovin, Vincasar)—Leurocristine, VCR</td>
<td>FDA approved for Hodgkin disease and other lymphomas, acute leukemias, rhabdomyosarcoma, neuroblastoma, and Wilms tumor. Used for many other neoplasms as well.</td>
<td>Vincristine is a vesicant and should be administered with extravasation precautions. Neurotoxicity is dose limiting in the form of peripheral neuropathy, which is related to total cumulative dose.</td>
<td>Inhibitor of tubulin polymerization and thereby mitosis; G2 phase specific.</td>
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### TABLE 27.1 Chemotherapeutic Agents (Continued)

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<tr>
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<tr>
<td>Vinorelbine (Navelbine)—5’-noranhydrovinblastine, NVB</td>
<td>FDA approved for the treatment of relapsed metastatic breast cancer and for NSCLC as a single agent or combined with a platinating agent.</td>
<td>Vinorelbine is a mild vesicant, requiring extravasation precautions. Myelosuppression, mostly leukopenia, is expected and dose limiting. Neurotoxicity in the form of neuropathy is less common and milder than that seen with vincristine. Tumor pain during administration has been reported.</td>
<td>Inhibitor of tubulin polymerization and thereby mitosis; $G_2$ phase specific.</td>
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<tr>
<td>Luteinizing hormone-releasing hormone (LHRH) Goserelin acetate (Zoladex)</td>
<td>FDA approved for advanced prostate cancer and used also in metastatic breast cancer.</td>
<td>Toxicity is mild. Endocrine side effects are most prominent and include hot flashes, diminished libido, impotence, gynecomastia, amenorrhea, and breakthrough vaginal bleeding.</td>
<td>Inhibits pituitary–gonadal axis function; causes steroid hormone withdrawal from dependent tissues, including prostate cancer and breast cancer cells.</td>
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<tr>
<td>Leuprolide acetate (Leupron)—Luprorelin Acetate</td>
<td>FDA approved for the treatment of hormone-dependent advanced prostate cancer. Also used for breast cancer and endometriosis.</td>
<td>Usually well tolerated, but side effects can affect many systems, including endocrine (hot flashes, impotence, gynecomastia, breast tenderness, diminished libido, amenorrhea, atrophic vaginitis, and increased cholesterol).</td>
<td>Gonadotropin-releasing hormone agonist, which shuts down the pituitary release of gonadotropins, resulting in a dramatic decrease in gonadal estrogens and androgens, and growth inhibition of hormone-dependent neoplasms.</td>
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<td><strong>Nonsteroidal antiandrogens</strong></td>
<td><strong>Tamoxifen</strong></td>
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<tr>
<td><strong>Bicalutamide (Casodex)</strong></td>
<td><strong>FDA approved for stage D2 prostate cancer, in combination with an LHRH agonist agent.</strong></td>
<td><strong>FDA approved for the treatment of breast cancer, generally in postmenopausal patients or those with estrogen receptor-positive tumors. The same dose has been approved for chemoprevention of breast cancer in high-risk patients. Higher doses are used for melanoma and pancreatic cancer.</strong></td>
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<td></td>
<td><strong>Constitutional symptoms predominate, including hot flashes, decreased libido, depression, weight gain, edema, gynecomastia, early disease-site pain (flare reaction), and constipation.</strong></td>
<td><strong>Tamoxifen is usually very well tolerated. Constitutional symptoms are most prevalent and usually dose limiting. Hot flashes, sweating, mood changes, weight gain or loss, and stomach upset are most common.</strong></td>
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| **Nonsteroidal aromatase inhibitors** | **FDA approved as adjuvant therapy for breast cancer and for treatment of postmenopausal women with breast carcinoma who have progressed on tamoxifen therapy.** | **The drug is very well tolerated. Asthenia, headache, and hot flashes occur in less than 15% of women. Thrombophlebitis has been reported.** |
|-----------------------------|---------------|
| **Anastrozole (Arimidex)**  | **Blocks estrogen production selectively.** | |

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<tr>
<th><strong>Cytoprotectants</strong></th>
<th><strong>FDA approved for pretreatment with cisplatin. Useful as a bone marrow, kidney, and nerve cytoprotectant. Useful with other alkylators. Also FDA approved as a radiation protectant to reduce xerostomia.</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Amifostine (Ethyol)—WR-2721, Ethiofos</strong></td>
<td><strong>Transient hypotension is dose limiting. Nausea, vomiting, and somnolence are common.</strong></td>
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+ indicates the degree of synergy with radiation.

Compiled by Christopher Schultz, MD, 2005.
Drugs acting by intercalation appear to augment the cytotoxic effects of bleomycin, as do x-rays and chemicals that generate superoxide radicals. Bleomycin is considered cell cycle nonspecific. It is more damaging to nonproliferating cells than to most proliferating cells.

Mitomycin C (Mutamycin) is an extremely toxic antitumor antibiotic. Unlike most other antibiotics, it is activated in vivo to a bifunctional or trifunctional alkylating agent. It is cell cycle nonspecific and is considerably more toxic to hypoxic than to aerated cells. Mitomycin C is usually administered intravenously; it is cleared rapidly from the plasma, with a half-life of 10 to 15 minutes, primarily by metabolism in the liver. It does not appear to cross the blood–brain barrier. The major toxicity of mitomycin C is myelosuppression.

Antimetabolites
The antimetabolites are analogues of normal metabolites required for cell function and replication. They may interact with enzymes and damage cells by any of these modes of action:

1. **Substituting** for a metabolite normally incorporated into a key molecule
2. **Competing** successfully with a normal metabolite for occupation of the catalytic site of a key enzyme
3. **Competing** with a normal metabolite that acts at an enzyme regulatory site, thereby altering the catalytic rate of the enzyme

Methotrexate is a folic acid antagonist. It works by competing for the folate-binding site of the enzyme dihydrofolate reductase. This results in decreased synthesis of thymidine and purine nucleotides. The cytotoxicity of methotrexate can be reversed by leucovorin, which is converted readily to other forms of reduced folate within the cell and which then can act as methyl donors for various biochemical reactions. The use of high-dose methotrexate with leucovorin rescue is based on the pharmacology of the two drugs, with the possibility of a differential effect between tumors and normal tissues in their ability to transport the two drugs across cell membranes. How true this differential effect turns out to be is another matter.

5-Fluorouracil
5-Fluorouracil (5-FU) is a structural analogue of the DNA precursor thymine. It works primarily as an irreversible inhibitor of the enzyme thymidylate kinase of injected cyclophosphamide from the plasma is biexponential, with an average half-life of 4 to 6.5 hours. Like all useful alkylating agents, cyclophosphamide produces toxicity in rapidly proliferating normal tissues. Chlorambucil (Leukeran) is an aromatic derivative of nitrogen mustard and is the slowest acting alkylating agent in general use. Melphalan (Alkeran, L-PAM) is a phenylalanine derivative of nitrogen mustard.

The nitrosoureas are a group of lipophilic alkylating agents that undergo extensive biotransformation in vivo, leading to various biologic effects, including alkylation, carbamylation, and inhibition of DNA repair. The multiple mechanisms of action may explain why the nitrosoureas generally lack cross-resistance with other alkylating agents. These compounds are very lipid soluble and readily cross the blood–brain barrier. They disappear from plasma rapidly, but their metabolites may persist for days.

**Antibiotics**
The clinically useful antibiotics are natural products of various strains of the soil fungus Streptomyces. They produce their tumoricidal effects by directly binding to DNA, and so their major inhibiting effects are on DNA and RNA synthesis. As a class, these drugs behave as cell cycle–nonspecific agents. Doxorubicin (Adriamycin) and daunorubicin (also known as daunomycin) are closely related anthracycline antibiotics. After intravenous injection, both drugs undergo extensive bioreduction in the liver to active and inactive metabolites, are bound extensively in tissues, and persist in plasma for prolonged periods. Neither drug crosses the blood–brain barrier to any appreciable extent. Both doxorubicin and daunorubicin are highly toxic drugs, producing various severe reactions; the major limiting toxicity, however, is cardiac damage.

Dactinomycin (actinomycin D) inhibits DNA-primed RNA synthesis by intercalating with the guanine residues of DNA; at higher concentrations, it also inhibits DNA synthesis. The net effect is cell cycle–nonspecific cytotoxicity. Dactinomycin must be administered intravenously. It is important longer plasma half-life is about 36 hours, and the drug is extensively bound to tissues.

Bleomycin sulfate (Blenoxane) affects cells by directly binding to DNA, resulting in reduced synthesis of DNA, RNA, and proteins. It also can lead to single-strand DNA breaks.
huge spreading yew tree, dating from medieval times when the archer's weapon, the long bow, was fabricated from the wood of the yew—but the tree is always found in the walled-in church cemetery, because the leaves or bark are toxic to browsing livestock. This toxicity is caused by a class of compounds known as taxanes. For those who see chemotherapy for cancer as simply the latest round in the never-ending battle between plants and animals, taxanes do indeed provide the perfect example.

**Paclitaxel** is the prototype of a new class of antineoplastic agents, the taxanes that targets the microtubules. It is a natural product, isolated from the bark of the western yew, *Taxus brevifolia*. Docetaxel is a largely synthetic derivative.

**Taxanes** are potent microtubule-stabilizing agents and promoters of microtubule assembly. This is in contrast to agents such as the vinca alkaloids and colchicine that bind to tubulin, the subunit of microtubules, and inhibit microtubule formation. The taxanes block or prolong the transit time of cells in the G2/M phase of the cell cycle. The inability of these cells to pass through the G2 and M phases of the cycle results from the inability of these cells to form a competent mitotic spindle or to disassociate a drug-treated spindle.

In addition to multiple *in vitro* studies from the early 1970s on, human studies have demonstrated the ability of taxanes to increase the mitotic index in various normal tissues *in vivo*, whereas the two taxanes in clinical use, Taxol (paclitaxel) and Taxotere (docetaxel), have demonstrated significant levels of activity in a broad range of human tumors. The taxanes are of particular interest to radiobiologists because of the way in which they interact with radiation (described later).

**Miscellaneous Agents**

**Procarbazine**

Procarbazine is a hydrazine derivative that must undergo biotransformation before it can exert its cytotoxic effects. The precise mechanism of action is not clear, because it interferes with various biochemical processes. Procarbazine is well absorbed from the gastrointestinal tract and is cleared from the plasma with a half-life of about 10 minutes. The drug freely crosses the blood–brain barrier. It is used primarily in the treatment of advanced Hodgkin disease.
Targeted Therapy

Achieving tumor response with traditional chemotherapeutic agents, such as the alkylating agents, cis-platinum, and 5-FU, for example, inevitably involved substantial toxicity to self-renewal normal tissues, such as the gastrointestinal tract and the blood-forming organs. There is some promise that the situation is much more favorable in the case of the new generation of targeted therapeutic agents that, in combination with radiotherapy, enhance tumor control with little, if any, increase in normal tissue toxicity. Cetuximab (Erbitux) might be regarded as a model for this class of pathway-targeting agents. Promising but still early results for bevacizumab await further clinical testing.

Cetuximab (Erbitux)

There appears to be a strong correlation between both locoregional control and overall survival and the level of epidermal growth factor receptor (EGFR) expression in advanced head and neck squamous carcinoma. Because high levels of EGFR predict for a poor outcome, EGFR is an attractive target for cancer therapy.

Cetuximab (Erbitux) is a recombinant human/mouse chimeric monoclonal antibody produced in mammalian cell culture. Cetuximab specifically binds to EGFR, blocking phosphorylation and activation of receptor-associated kinases, resulting in inhibition of cell growth, induction of apoptosis, and decreased matrix metalloproteinase and vascular endothelial growth factor (VEGF).

Studies using cetuximab with a xenograft tumor in mice indicate that although the antibody alone produces a modest tumor growth delay, there is some evidence that cis-platinum is more toxic to hypoxic than to aerated cells—that is, that it is a hypoxic cell radiosensitizer, although not as powerful in this regard as the nitroimidazoles.

Hydroxyurea

Hydroxyurea was first synthesized as long ago as 1869 and was found to be bone marrow suppressive in 1928. It was not used in the treatment of cancer until the 1960s. It acts as an inhibitor of ribonucleotide reductase, an enzyme essential to DNA synthesis, and is consequently specifically cytotoxic to cells in the S phase of the cell cycle. In experimental biology, hydroxyurea is used to synchronize cells, because in addition to killing S phase cells, it causes survivors to pile up at a block at the G1/S interface. Clinically, hydroxyurea is used primarily in the treatment of chronic myeloid leukemia.

Cis-Platinum

Structurally, cis-platinum (cis-dichlorodiammine-platinum) is an inorganic complex formed by an atom of platinum surrounded by chlorine and ammonium ions in the cis position of the horizontal plane. Cis-platinum bears a resemblance to the bifunctional alkylating agents based on nitrogen mustard. It inhibits DNA synthesis to a greater extent than it does the synthesis of RNA or protein. It binds to DNA, causing both interstrand and intrastrand crosslinking.

Cis-platinum is cell cycle nonspecific. Its isomer, trans-platinum, is much less cytotoxic, presumably because of the different way that it crosslinks to DNA. There is some evidence that cis-platinum is more toxic to hypoxic than to aerated cells—that is, that it is a hypoxic cell radiosensitizer, although not as powerful in this regard as the nitroimidazoles.

Topoisomerase Inhibitors

DNA topoisomerases are nuclear enzymes that reduce twisting and supercoiling that occur in selected regions of DNA as a result of transcription, replication, and repair recombination. Little is known of the actual mechanism by which they kill cells.

Top I inhibitors, thus far consisting primarily of camptothecin analogues, interact with the enzyme–DNA complex and prevent the resealing of DNA strand breaks, leading to cell death in cells actively replicating.

Top II inhibitors prevent religation of DNA cleaved by top II and lead to cell death. Etoposide, a semisynthetic podophyllotoxin derivative, is a prime example.
FIGURE 27.2  A: Five-year survival data for locoregionally advanced head and neck cancer patients treated with radiotherapy alone or the combination of radiotherapy and cetuximab. The addition of cetuximab increased survival by 9%. B: Five-year survival for locoregionally advanced head and neck cancer based on whether patients had an acne-like rash of grade 2–4. Patients with a rash above grade 2 had 2.5 times longer survival than those who had a grade 1 rash or no rash. (Adapted from Bonner JA, Harari PM, Giralt J, et al. Radiotherapy plus cetuximab for locoregionally advanced head and neck cancer: 5-year survival data from a phase 3 randomised trial, and relation between cetuximab-induced rash and survival. Lancet Oncol. 2010;11:21–28, with permission.)
Bevacizumab (Avastin)
In 1971, Judah Folkman proposed that the formation of new blood vessels (angiogenesis) was an essential process for tumor growth and metastasis. This concept has been widely confirmed by other investigators and launched the era of “antiangiogenic agents.” The most commercially successful of these agents is bevacizumab, a monoclonal antibody that neutralizes the activity of the proangiogenic mitogen VEGF.

The first clinical trials with bevacizumab in the treatment of recurrent and metastatic breast cancer resulted in failure to demonstrate a survival benefit. The major insight in the use of antiangiogenic agents was found to be in combination with cytotoxic chemotherapy. Currently, bevacizumab is approved with chemotherapy for metastatic colorectal cancer, lung cancer, and renal cancer. The increased survival of patients treated with the combination of antiangiogenic agents and chemotherapy compared to antiangiogenic therapy alone brought about the vascular normalization hypothesis that proposes that agents such as bevacizumab eliminate the malformed and inadequate vessels in a tumor, reducing tumor size and making it better perfused and more responsive to cytotoxic chemotherapy and radiotherapy (Fig. 27.3).

More recently, Willet and his colleagues have reported the results of some early phase I clinical trials to determine whether inhibition of VEGF is safe and enhances the effect of chemoradiotherapy with locally advanced rectal cancer. The combination of bevacizumab, 5-FU, and external beam irradiation resulted in decreased tumor vasculature density and decreased interstitial fluid pressure. These results are consistent with the inhibition of angiogenesis and a vascular normalizing effect of bevacizumab. These results strongly suggest that antiangiogenic agents can be safely and effectively combined with radiotherapy.

Poly Adenosine Diphosphate-Ribose Polymerase Inhibitors
The enzyme poly adenosine diphosphate-ribose polymerase 1 (PARP-1) plays an essential role in the base excision repair pathway, in particular,
in the recruiting of x-ray cross complementing factor 1 (XRCC1), which acts as a scaffold protein to coordinate the repair of damaged bases (see Chapter 2). Inhibition of PARP results in decreased based excision repair activity and accumulation of single-strand breaks that will lead to the collapse of replication forks in the S phase of the cell cycle and the generation of DNA double-strand breaks (DSBs) and cell lethality. If tumors are deficient in the repair of DNA DSBs by homologous recombination in the S phase of the cell cycle because of mutations in BRCA1 or BRCA2, then PARP inhibition will result in cell lethality. This so-called synthetic lethal interaction between PARP inhibition and BRCA deficiency has served as the basis for several clinical trials where PARP inhibitors are administered alone or in combination with cytotoxic chemotherapy agents to patients with BRCA mutations in breast, ovarian or prostate cancer, or to patients with triple negative breast cancer. The scientific rationale behind these trials is that inhibition of PARP alone or mutations in BRCA genes alone have little effect on cell viability. However, only when the two are combined do you have cell killing. This type of interaction has been termed synthetic lethality because of the requirement that two pathways need to be inhibited that typically are unaffected and lead to very few viable progeny (Fig. 27.4). Not surprisingly, most studies to date have focused on PARP-1 inhibitors as a monotherapy or administered with some forms of cytotoxic chemotherapy. However, inhibition of PARP renders cells sensitive to radiotherapy and even more so in a BRCA mutation containing tumors. Thus, although PARP inhibitors are quite new, the combination of impressive phase I and II clinical results and potential synergism with radiotherapy will most probably increase their use with radiotherapy in the future.

**Dose–Response Relationships**

Dose–response relationships have been produced for a wide range of chemotherapeutic agents using techniques developed initially for radiation, although much less effort has been expended on fitting data to models than has been the case for ionizing radiations. From even a cursory examination of the data, however, it is evident that—with some clear exceptions—the shape of the survival curve is unremarkable and reminiscent of that of survival curves for ionizing radiations. If surviving fraction is plotted on a log scale against drug dose on a linear scale, the dose-response curve has an initial shoulder followed by a region that becomes steeper and straighter (Fig. 27.5). The antibiotics doxorubicin (Adriamycin), bleomycin, and dactinomycin (actinomycin D) are clear exceptions. For these agents, the dose-response curve appears to have no shoulder, and the curve is concave upward. A dose-response curve with this shape is usually associated with a variation of sensitivity within the population (i.e., nonuniform sensitivity of cells). This has never been demonstrated experimentally, however. For example, synchronously dividing cells likewise show the same upwardly concave dose-response curve. This shape therefore must remain unexplained at the present time.

Dose–response curves indicate that, at best, anticancer drugs kill by first-order kinetics; that is, a given dose of the drug kills a constant fraction of a population of cells, regardless of its size. This assumes, of course, that the growth fraction and the proportion of sensitive to resistant cells remains the same. This leads to the conclusion that the chance of eradicating a cancer is greatest if the population size is small, or that there is an inverse relationship between curability and the tumor cell burden at the initiation of chemotherapy. This conclusion has been arrived at from long and bitter clinical experience, but, in fact, it is an inevitable consequence of the shape of the simplest dose–response relationship.

**FIGURE 27.4** Schematic representation of the synthetic lethality principle. Synthetic lethality occurs when individual loss of two genes (A or B) is viable but is lethal when in combination (A and B). This principle is exemplified with genes involved in DNA repair pathways, PARP (BER) and BRCA1 (HR). (From Hammond EM, Pires I, Giaccia AJ. DNA damage and repair. In: Leibel SA, Phillips TL, eds. *Textbook of Radiation Oncology*. 3rd ed. Elsevier; in press, with permission.)
FIGURE 27.5  Dose–response relationships in vitro for six commonly used chemotherapeutic agents. Note the diverse shapes. Many have shapes similar to survival curves for x-rays, except that drug concentration replaces absorbed dose. The antibiotics bleomycin (graph B), Adriamycin (doxorubicin) (graph E), and actinomycin D (graph A) have dose–response relationships that are concave upward. A: Dose–response relationship for dividing CHO cells treated for 1 hour with graded doses of actinomycin D. (Adapted from Barranco SC, Fluorney DR. Modification of the response to actinomycin-D-induced sublethal damage by simultaneous recovery from potentially lethal damage in mammalian cells. Cancer Res. 1976;36:1634–1640, with permission.) B: Dose–response relationship for plateau-phase CHO cells treated for 1 hour with graded doses of bleomycin. (Adapted from Barranco SC, Novak JKJ, Humphrey RM. Response of mammalian cells following treatment with bleomycin and 1,3-bis(2-chloroethyl)-1-nitrosourea during plateau phase. Cancer Res. 1973;33:691–694, with permission.) C: Dose–response relationship for CHO cells treated for 1 hour with graded doses of CCNU. (Adapted from Barranco SC. In vitro responses of mammalian cells to drug-induced potentially lethal and sublethal damage. Cancer Treat Rep. 1976;60:1799–1810, with permission.) D: Dose–response relationship for human lung cancer cells exposed for 1 hour to graded doses of melphalan. (Unpublished data, courtesy of Dr. Laurie Roizin-Towle.) E: Dose–response relationship for V79 Chinese hamster cells exposed for 1 hour to graded doses of Adriamycin (doxorubicin). (Adapted from Belli JA, Piro AJ. The interaction between radiation and Adriamycin damage in mammalian cells. Cancer Res. 1975;37:1624–1630, with permission.) F: Dose–response relationship for V79 Chinese hamster cells exposed for 1 hour to graded doses of cis-platinum (cisplatin). (Unpublished data, courtesy of Dr. Laurie Roizin-Towle.)
demonstrated by an increase in survival if a dose of radiation (or other cytotoxic agent) is divided into two or more fractions separated in time. There is a tendency for the extent of sublethal damage repair to correlate with the shoulder of the acute dose-response curve, but this is not always true. Repair of potentially lethal damage is manifest as an increase in survival if cells are held in a nonproliferative state for some time after treatment.

Similar studies have been performed with various chemotherapeutic agents. The results are not as clear-cut as for radiation, and there is much greater variability between different cell lines. Potentially lethal damage repair is a significant factor in the antibiotics bleomycin and doxorubicin. Data for bleomycin are shown in Figure 27.7. Potentially lethal damage repair is also seen after treatment with dactinomycin. Sublethal damage repair is essentially absent with all of these drugs.

Another characteristic of chemotherapy agents is that the sensitivity to cell killing varies enormously among cell types. Radiosensitivity varies, too, of course, but not to the same extent as chemosensitivity. Dose-response curves for several cell lines exposed to paclitaxel are shown in Figure 27.6, illustrating the wide range of sensitivities.

**SUBLETHAL AND POTENTIALLY LEthal DAMAGE REPAIR**

Studies with radiation led to the concepts of sublethal damage repair and potentially lethal damage repair, which are discussed in some detail in Chapter 5. These are still largely operational terms, although a notable exception is mitomycin C in which the gene for repair of DNA damage has been identified and cloned. Sublethal damage repair is demonstrated by an increase in survival if a dose of radiation (or other cytotoxic agent) is divided into two or more fractions separated in time. There is a tendency for the extent of sublethal damage repair to correlate with the shoulder of the acute dose-response curve, but this is not always true. Repair of potentially lethal damage is manifest as an increase in survival if cells are held in a nonproliferative state for some time after treatment.

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Potentially lethal damage repair is a significant factor in the antibiotics bleomycin and doxorubicin. Data for bleomycin are shown in Figure 27.7. Potentially lethal damage repair is also seen after treatment with dactinomycin. Sublethal damage repair is essentially absent with all of these drugs.
No potentially lethal or sublethal damage repair is seen with nitrosourea, even though the dose-response curves for single doses have substantial shoulders. The breakdown products of the nitrosoureas are known to inhibit DNA repair, and this may be a contributing factor.

Studies of repair of sublethal damage with drugs are complicated, because if a split-dose study is performed, decisions must be made about the equivalence of drug concentration and time. It is frequently assumed that biologic response is determined by an integral dose (i.e., the product of concentration and time), but this has not been checked and confirmed in all cases. There appears to be no correlation between the existence of a shoulder on the dose-response curve for single doses and the appearance of sublethal damage repair, as evidenced by an increase in survival in a split-dose experiment. It is possible that a shoulder in a survival curve does not have the same meaning for chemically induced damage as it does for radiation-induced damage. A shoulder on a survival curve for a chemotherapeutic agent may reflect more about drug concentrations and the time required for entry of the drug into the cells and interaction with a target molecule than it does about the accumulation and repair of sublethal damage.

### THE OXYGEN EFFECT AND CHEMOTHERAPEUTIC AGENTS

The importance of the oxygen effect for cell killing by radiation is discussed in Chapter 6. It has been known for more than half a century that the presence or absence of molecular oxygen has a dramatic influence on the proportion of cells surviving a given dose of x-rays. Only in more recent years has the influence of oxygen on the cytotoxicity resulting from chemotherapeutic agents been studied. It is certainly more complicated than for ionizing radiations.

Some agents, such as bleomycin, are more toxic to oxygenated cells than to chronically hypoxic cells. Dose-response curves for cells exposed to graded concentrations of bleomycin in the presence or absence of oxygen are shown in Figure 27.8.

**FIGURE 27.8** Molecular oxygen can be either a sensitizer or a protector, depending on the particular chemotherapeutic agent. **A:** Survival curves for EMT6 cells treated for 1 hour with graded doses of mitomycin C under aerated or hypoxic conditions. In the absence of oxygen, the cells are substantially more sensitive. (Data from Teicher BA, Lazo JS, Sartorelli AC. Classification of antineoplastic agents by their selective toxicities toward oxygenated and hypoxic tumor cells. *Cancer Res.* 1981;41:73–81.) **B:** Survival curves for Chinese hamster cells in culture exposed for 4 hours to graded doses of bleomycin under aerated or hypoxic conditions. In the absence of molecular oxygen, the cells are more resistant. (Adapted from Roizin-Towle L, Hall EJ. Studies with bleomycin and misomnida zole on aerated and hypoxic cells. *Br J Cancer.* 1978;37:254–260, with permission.)
macromolecular synthesis and inducing genes that promote angiogenesis and tissue remodeling. This is discussed in some detail in Chapter 26. The ability of malignant cells to survive fluctuations in oxygen tension is important in the context of cancer therapy, because it is well documented that tumors that are hypoxic are aggressive and respond poorly to all forms of treatment. The shortage of oxygen in hypoxic cells renders them refractory to killing by ionizing radiations that require oxygen to “fix” the damage produced in DNA by the hydroxyl radicals; it is equally true that cells may be refractory to killing by “radiation-mimetic drugs” that operate via a free radical mechanism. Cessation of cell division and loss of apoptotic potential (cell suicide), however, in hypoxic cells are likely to be more important reasons why hypoxic cells are resistant to chemotherapy. Hypoxic cells located remote from capillaries are least likely to be killed by radiation but most likely to be killed by a hypoxic cytotoxin. This is illustrated in Figure 27.9.

The most effective way to partly circumvent the problem introduced by hypoxia in reducing the efficacy of chemotherapy is to introduce a bioreductive drug or hypoxic cytotoxin that is most effective at reduced oxygen concentrations. This is described in more detail in Chapter 26. Mitomycin C and its derivatives have been used for this purpose for many years.
**DRUG RESISTANCE AND CANCER STEM CELLS**

The biggest single problem in chemotherapy is drug resistance, which either may be evident from the outset or may develop during prolonged exposure to a cytostatic drug. Cells resistant to the drug take over, and the tumor as a whole becomes unresponsive. The development of resistance can be demonstrated readily for cells in culture. Figure 27.10 shows a substantial resistance to doxorubicin developing as cells are grown continuously in a low concentration of the drug for a period of weeks.

Underlying this problem of drug resistance are genetic changes that can sometimes be seen in chromosome preparations. Figure 27.11 shows two illustrations involving gene amplification or the presence of multiple minute chromosome fragments.

Drug resistance is an important factor that occurs readily—a phenomenon quite alien to the radiobiologist. Radiation-resistant cells can be produced and isolated, but it is a difficult and time-consuming process. For instance, cells continuously irradiated at low dose rates occasionally do spawn radioresistant clones. By contrast, resistance to chemotherapeutic agents is acquired quickly, uniformly, and inevitably.

If a resistant clone can arise by a chance mutation of a gene responsible for one of the important steps in drug action, then the probability of it occurring would be expected to increase rapidly as the tumor increases in size. The average mutation rate for mammalian genes is about $10^{-7}$ to $10^{-8}$ per division, so that in a tumor containing $10^{10}$ cells that go through many divisions, a mutation is almost certain to occur, especially in the presence of a powerful mutagen, which most chemotherapeutic agents are.

Another contributing factor to chemotherapy resistance is cancer stem cells. The current model of cancer stem cells proposes that although these cells represent only a small subpopulation of the tumor, they are the most important because they possess the potential to give rise to both stem cells and differentiated non-stem cells. Therapeutically, stem cells have been...
FIGURE 27.10 Change in survival response to doxorubicin (Adriamycin) of Chinese hamster cells grown in culture and exposed continuously to a low concentration of the drug (0.05 μg/ml) for prolonged periods, namely, 1, 17, 46, or 74 weeks. The closed circles show the survival response for the parent cell line. A dramatic resistance to the drug develops by 17 weeks; that is, prolonged exposure at a low concentration renders the cells resistant to subsequent high concentrations. (Adapted from Belli JA. Radiation response and Adriamycin resistance in mammalian cells in culture. Front Radiat Ther Oncol. 1979;13:9–20, with permission.)

FIGURE 27.11 Most forms of drug resistance probably have a genetic basis. A few extreme examples can be seen in chromosome changes. A: The arrow indicates an elongated chromosome, which on banding shows the features of an extended homogeneously staining region. This karyotype was observed in the human breast cancer cell line (MCF-7), which is resistant to methotrexate. (From Cowan KH, Goldsmith ME, Levine R, et al. Dihydrofolate reductase gene amplification and possible rearrangement in estrogen-responsive methotrexate-resistant human breast cancer cells. J Biol Chem. 1982;257:15079–15086, with permission.) B: Small-cell lung carcinoma line derived from a patient treated with methotrexate. These cells are very resistant to the drug and contain numerous double minute chromosomes. A pair is indicated by arrows. (From Curt GA, Carney DN, Cowan KH, et al. Unstable methotrexate resistance in human small-cell carcinoma associated with double minute chromosomes. N Engl J Med. 1983;308:199–202, with permission.)
reported to be resistant to ionizing radiation and chemotherapy. Resistance to radiation has been processed to be mediated through decreased reactive oxygen species that are scavenged before they give rise to DNA strand breaks. In contrast, chemotherapy resistance is thought to be associated with members of the adenosine triphosphate-binding cassette (ABC) transporter that transports drugs across the membrane. High levels of free radical scavengers in cancer stem cells will also make them refractory to certain chemotherapy agents such as bleomycin and doxorubicin. In addition to intrinsic mechanisms of resistance, cancer stem cells may also reside in specialized niches that are hypoxic, or impede chemotherapy activity.

The usual strategy to overcome the problem of induced resistance is to use a battery of different drugs, applied sequentially and cyclically, that produce their cytotoxicity by diverse mechanisms. By this strategy, cells that develop resistance to drug A are killed by drug B, and so on.

The bigger problem is pleiotropic resistance, the phenomenon by which the development of resistance to one drug results in cross-resistance to other drugs, even those with different mechanisms of action. There are four interesting points to be made:

1. Multidrug resistance in tumor cells is caused by extrusion of the drugs; that is, cells pump the drugs out as fast as they get in. This is mediated by increased expression of the product of the multiple drug resistance gene (mdr), a p-glycoprotein expressed in the cell membrane. This membrane protein is a polypeptide of 1,280 amino acids composed of two similar domains, each containing six potential transmembrane segments and two putative adenosine triphosphate-binding regions. Its structure is similar to that of various transporters of ions, amino acids, peptides, or proteins in bacterial, yeast, and animal cells. Indeed, it has been reported that the mdr gene in human tumor cells shows considerable homology to the gene in yeast that extrudes an attractant that is important in the reproductive cycle. The mdr gene has been mapped to human chromosome 7. Resistance by this means can be reversed by calcium channel-blocking drugs, such as verapamil. This has been shown to be an important mechanism of resistance to doxorubicin in Chinese hamster ovary (CHO) cells in culture, and there appears to be expression of this same gene for resistance in cells from some human solid tumors that have acquired resistance.

2. Glutathione is a naturally occurring thiol in all cells. Elevated levels of glutathione have been observed in resistant cells, especially those made resistant by treatment with melphalan. Drugs are available that block the synthesis of glutathione and that can be used to lower the levels of this compound in tumors and normal tissues. The best known example is buthionine sulfoximine. Use of buthionine sulfoximine has been shown to reduce cross-resistance, particularly between melphalan and cis-platinum in tumor-bearing mice. The use of buthionine sulfoximine would not be advisable in combination with doxorubicin or cis-platinum, because an increase in specific normal tissue toxicity (lung or kidney, respectively) would be expected.

3. A marked increase in DNA repair has been noted in some cells resistant to melphalan or cis-platinum. In principle, drugs could be used to block repair; aphidicolin has been used in experimental systems, but no suitable drugs are available for clinical use.

4. A debatable issue is whether cells that have acquired resistance to chemotherapeutic agents are also resistant to radiation. The consensus is that they are not. There may be some data from clinical experience to suggest that they are, but the laboratory data show rather clearly that acquiring resistance to a drug does not necessarily result in radioresistance. This is illustrated in Figure 27.12, in which cells that have acquired extreme resistance to melphalan show a normal response to radiation. Radioresistance and chemoresistance may occur together, but radiation rarely induces chemoresistance and vice versa.

The evolving story of drug resistance has an impact on the development and screening of new drugs. In the past, the initial screening for new agents consisted of fast-growing, highly drug-sensitive mouse tumors. Tests against specific patterns or types of drug resistance were not included. The screening systems, therefore, were weighted heavily in favor of producing more of the same types of drugs. This has changed, and
Chapter 27 • Chemotherapeutic Agents from the Perspective of the Radiation Biologist

Surviving fraction

\[
\begin{array}{c|c|c|c|c|c|c|c|c|c|c}
\text{Melphalan (\mu g/ml)} & 0 & 4 & 8 & 12 & 16 & 20 \\
\hline
\text{AB} & \bullet & \circ & \circ & \circ & \circ & \circ \\
\text{C5} & \circ & \circ & \circ & \circ & \circ & \circ \\
\end{array}
\]

**FIGURE 27.12** Chinese hamster pleiotropic multidrug-resistant cells are not necessarily resistant to radiation. The parental ovary cell line is designated AB. The multidrug-resistant cell line C5 was isolated by Dr. Victor Ling by exposing the parental line to the mutagen ethyl methane sulfonate, after which surviving cells were grown for an extended period in increasing concentrations of colchicine. A clone was isolated that is resistant to colchicine and to various chemotherapeutic agents. **A:** C5 cells are resistant to melphalan, compared with the parental line (AB). They are also resistant to other agents, such as daunorubicin. **B:** The radiation responses of the parental and the chemotherapy-resistant cell lines are virtually indistinguishable. (Adapted from Mitchell JB, Gamson J, Russo A, et al. Chinese hamster pleiotropic multidrug resistant cells are not radioresistant. *Natl Cancer Inst Monogr.* 1988;6:187–191, with permission.)

The screening of new drugs for activity is performed using a battery of cells of human origin cultured in vitro.

### COMPARISON OF CHEMOTHERAPEUTIC AGENTS WITH RADIATION

The title of this chapter includes the words “from the perspective of the radiation biologist.” This limited and specialized viewpoint must be kept in mind in what follows. Several important differences are evident in the response of cells to chemotherapeutic agents versus ionizing radiation:

1. There is a much greater variation of sensitivity to chemotherapeutic agents than there is to radiation. In the case of x-rays, the variation of D₀ from the most sensitive to the most resistant known mammalian cells may be a factor of about 4. By contrast, the response of various cell lines to a given chemotherapeutic agent may differ by orders of magnitude. A particular cell line may be exquisitely sensitive to one drug and extremely resistant to another. A different cell line may have a different order of sensitivity to various drugs, as well as a quite different absolute sensitivity. Different clones derived from a common stock may exhibit quite different sensitivity to a given agent. This variability is shown in Figure 27.13, which gives the response of one cell line to nine different cytotoxic agents, and in Figure 27.14, which shows the widely different response to Me CCNU of three permanent clones derived from a common astrocytoma cell line.

2. The sensitivity of a given cell line to a given drug may be manipulated to a much greater extent than for radiation.

3. Repair of sublethal and potentially lethal damage is more variable and less predictable for drugs than for radiation.

4. The oxygen effect is more complex for drugs than for ionizing radiations. For radiation, the presence or absence of molecular oxygen has an important influence on the proportion of cells surviving a given dose of low-linear energy transfer (LET) radiation, in which about two-thirds of the damage is caused by indirect action (i.e., mediated by free radicals). As the LET of the radiation increases and the balance shifts from indirect to direct action, the importance of oxygen decreases. For very high-LET radiations (above about 200 keV/\mu m), the biologic effect for a given dose is independent of the presence or absence of molecular oxygen. Under no circumstances is oxygen protective in the case of ionizing radiations.
For drugs in which the biologic effect involves free radicals, the presence or absence of oxygen is important in the same way as for low-LET ionizing radiations. The new factor in the case of drugs is that there is a whole class of antineoplastic agents that undergo bioreduction in the absence of oxygen, so that they are more effective in hypoxic cells. There is no parallel for ionizing radiations.

Other agents do not depend primarily on free radicals for their biologic effects, nor do they undergo bioreduction under hypoxic conditions; consequently, the effect of a given treatment is independent of the presence or absence of molecular oxygen, a property these agents have in common with very densely ionizing radiations.

5. Resistance to drugs develops more quickly and more regularly than it does to radiation.

Acquired resistance to drugs does not necessarily involve resistance to x-rays as well.

6. Drug resistance may be caused by changes in thiol levels or by molecular changes observable at the chromosome level that result in the activation of a gene that functions to pump the drug out of the cells.

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**ADJUNCT USE OF CHEMOTHERAPEUTIC AGENTS WITH RADIATION**

The initial rationale for the combination of radiation and chemotherapeutic agents was what is usually known as “spatial cooperation” (Fig. 27.15). Radiation may be more effective for controlling the localized primary tumor, because it can be aimed and large doses are given, but it is ineffective against disseminated disease.
Chemotherapy, on the other hand, may be able to cope with micrometastases, whereas it could not control the larger primary tumor. In other situations, chemotherapy is the primary treatment modality, and radiation is used only to treat “sanctuary” sites not reached by the drug.

Although spatial cooperation was the original rationale, it is no longer the only one. Radiation and chemotherapeutic agents are combined in an attempt to achieve better local control. A specific example is the integrated use of taxanes and x-rays for the treatment of breast cancer. This is based on laboratory experiments in vitro. The initial hypothesis for the mechanism of interaction is that cell cycle alterations induced by paclitaxel leave cells in a state in which they are more sensitive to radiation. This is illustrated in Figure 27.16, which shows survival curves for astrocytoma cells.
of human origin cultured in vitro and exposed to graded doses of γ-rays, either alone or following a 24-hour treatment with 10-nM paclitaxel. This drug treatment alone kills about 95% of the cells and accumulates the survivors in the radiosensitive G2/M phase of the cell cycle (Fig. 27.16, inset).

Inasmuch as the cell killing resulting from the combination of drug and radiation is greater than the sum of the two separately, the interaction may be described as “synergistic.” For this to occur, the drug must accumulate some cells in G2/M, which is a radiosensitive phase of the cycle, without killing them. Although similar results have been obtained for several cell lines, there are a significant number of reports in the literature in which the interaction between radiation and paclitaxel was shown to be purely additive. Additivity is observed in any cell line in which the G2/M-arrested cells are doomed to die as a consequence of the drug alone. Even an additive interaction, however, may be therapeutically useful. For example, in animal tumors in vivo, the paclitaxel-induced death of G2/M cells can lead to tumor shrinkage and reoxygenation, which result in an enhanced response to subsequent irradiation.

Although the initial rationale for the combination of taxanes and radiation was based on classic radiobiologic considerations regarding the cell cycle dependence of radiosensitivity, there are other possible explanations for a synergistic effect if the two agents are used together. For example, the contribution of p53 to paclitaxel-dependent cytotoxicity is in marked contrast with its role for DNA-damaging agents such as radiation. Cells with a mutant p53 phenotype are more sensitive to paclitaxel treatment than are nontransformed or wild-type p53 cells. This finding explains the effectiveness of paclitaxel as a cytotoxic agent alone, as well as for the efficacy of its combination with radiation. In this theory, paclitaxel and radiation act as non-cross-resistant agents; radiation is more effective against wild-type p53 cells in a tumor population, and paclitaxel is relatively more cytotoxic to cells with a mutant p53 phenotype.

In general, a therapeutic gain requires differential effects between tumor and normal tissue. One or more of the following tumor characteristics may be exploited to achieve this difference:

1. Genetic instability of tumor cells
2. Rapid proliferation of some tumor cells
3. Cell age distribution of tumor cell populations
4. Hypoxia (characteristic of larger tumors)
5. pH (often low in tumors)
6. Elevation of specific pathways in tumors (for example, EGFR)

The goal of combining the two modalities of radiation and cytotoxic drugs is to increase local tumor control, relapse-free survival, and overall survival, and to alter the pattern of relapse. Three obvious strategies are (1) to start chemotherapy after the completion of the local treatment (adjuvant chemotherapy), (2) to start chemotherapy before the local treatment (induction chemotherapy), or (3) to give chemotherapy during local treatment (concurrent chemotherapy). Most cytotoxic agents do not provide enough differential sensitization of tumors compared with normal tissues; consequently, the probability of dose-limiting or life-threatening toxicity to critical tissues may be increased.

Integrating surgery, chemotherapy, and radiotherapy has been found to increase materially the cure rate of a dozen or more experimental animal tumors. Emerging principles include the following:

1. The lower the initial body burden of the tumor, the better.
2. A maximally effective drug regimen should begin as soon as possible after surgery, with maximum practicable doses.
3. Concurrent chemotherapy may lead to improved local control, but at the price of increased local toxicity.

Improving local control can improve overall survival rates and avoid uncontrolled local growth and the possible need for mutilating surgery. For example, induction chemotherapy prior to radiation therapy has a proven benefit in larynx preservation. For this reason alone, aggressive use of combined modalities to improve local control is warranted.

It has often been said that “you can only kill the sensitive cells once.” There is no point in using a battery of different agents that all target the same subpopulation of sensitive cells. Rather, the heterogeneity of the tumor cell population should be acknowledged and exploited. One agent should be effective against cycling cells, another against resting cells, and a third perhaps against hypoxic cells; in other words, the strategy should be to combine agents that specifically attack the different subpopulations in the tumor.

A problem to be avoided is the triggering of “accelerated repopulation.” This refers to triggering surviving clonogens to divide and repopulate even
more rapidly as a tumor shrinks after treatment with a cytotoxic agent. It is a phenomenon described in Chapter 22. It may be one of the reasons why radiotherapy after induction chemotherapy has shown disappointing results.

**ASSAYS FOR SENSITIVITY OF INDIVIDUAL TUMORS**

A great deal of effort has been expended to develop ways to assess which agents are likely to be effective for a particular tumor. The long-term goal would be to mimic the testing of a bacterial infection for sensitivity to a wide range of antibiotic drugs to select the one most suitable and effective.

One approach is to take biopsy specimens from a tumor in a patient, grow the cells *in vitro*, and subject the cells to a battery of chemotherapeutic agents in the petri dish. This approach has the advantages of not being too expensive to be practical and of providing answers quickly enough to influence the treatment and modify the protocol of the patient from whom the cells were taken. It suffers, of course, from the obvious disadvantages of focusing attention solely on the question of inherent cellular sensitivity and not addressing the questions of drug access, hypoxia, or any of the more complex factors involved as determinants of overall tumor response.

A different approach is to grow cells from human tumors as xenografts in immune-suppressed mice. This is a difficult and limited technique beset with problems, some of which are discussed in Chapter 21. Human tumor cell, however, maintain many characteristics of the clinical response of the donor tumors. Indeed, there is a good correlation between clinical remission in donor patients and growth delay in xenografts established from transplanted cells. The method of establishing xenografts and then performing the necessary growth delay experiments is sufficiently slow and time consuming that the technique cannot be expected ever to provide realistic input into deciding treatment strategy in individual patients, although it can provide guidance on the sensitivity of broad categories of human tumors to a battery of chemotherapeutic agents.

**SECOND MALIGNANCIES**

Late effects are the key to the acceptance of combined treatments. The induction of second malignancies is one of the unfortunate late effects of treatment with radiation or cytotoxic drugs. In a large series of 3,000 patients with Hodgkin disease treated with a combination of radiotherapy and chemotherapy, 114 developed second malignancies. The greatest relative risk was leukemia, but the greatest in number were solid tumors.

Radiation is a relatively weak carcinogen; chemotherapeutic agents vary widely. There is a choice of many chemotherapeutic agents, and the variable potential for producing a second malignancy must be a factor influencing the choice of drug in patients who are likely to be long-term survivors.

Table 27.3 compares radiation, bleomycin, ultraviolet radiation, and benzopyrene in terms

<table>
<thead>
<tr>
<th>Agent</th>
<th>D$_{37}$</th>
<th>DNA Lesion</th>
<th>Number of Lesions per Cell per D$_{37}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>X-rays</td>
<td>1 Gy</td>
<td>SSB</td>
<td>1,000</td>
</tr>
<tr>
<td></td>
<td></td>
<td>DSB</td>
<td>40</td>
</tr>
<tr>
<td>Bleomycin</td>
<td>5.5 µg × 1 h</td>
<td>SSB</td>
<td>150</td>
</tr>
<tr>
<td></td>
<td></td>
<td>DSB</td>
<td>30</td>
</tr>
<tr>
<td>Ultraviolet light</td>
<td>10 J/m$^2$</td>
<td>TT dimer</td>
<td>1,000,000</td>
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<tr>
<td></td>
<td></td>
<td>SSB</td>
<td>100</td>
</tr>
<tr>
<td>Benzopyrene</td>
<td>—</td>
<td>Adduct</td>
<td>100,000</td>
</tr>
</tbody>
</table>

SSB, single-strand break; DSB, double-strand break; TT, thymine-thymine.

Courtesy of Dr. John Ward, University of California at San Francisco.
Many classically used chemotherapeutic agents fall into one of several classes:

- Alkylating agents, which are highly active, with the ability to substitute alkyl groups for hydrogen atoms in DNA, include nitrogen mustard derivatives, cyclophosphamide, chlorambucil, melphalan, and the nitrosoureas (BCNU and CCNU).

- Antibiotics, which bind to DNA and inhibit DNA and RNA synthesis, include dactinomycin, doxorubicin, daunorubicin, and bleomycin.

- Antimetabolites, which are analogues of the normal metabolites required for cell function and replication, include methotrexate, 5-FU, cytarabine, and 5-azacytidine.

- Many of the newer and widely used drugs do not fall into any of these classes, including the vinca alkaloids, the taxanes, procarbazine, hydroxyurea, platinum complexes, topoisomerase inhibitors, and “targeted therapy” agents that target a specific pathway that may be elevated in some tumors (Erbitux, Herceptin, Gleevec, and Rituxan).

- Dose–response relationships for many chemotherapeutic agents resemble those for radiation, with drug concentration replacing absorbed dose; that is, there is an initial shoulder followed by an exponential relationship between surviving fraction and dose. The exceptions are doxorubicin, bleomycin, dactinomycin, and taxanes, which have dose–response curves that are concave upward.

- At best, traditional anticancer drugs kill cells by first-order kinetics; that is, a given dose kills a constant fraction of cells. Consequently, the chance of eradicating a cancer is greatest if the population size is small (i.e., there is an inverse relationship between curability and tumor cell burden) at the initiation of chemotherapy.

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- Studies of sublethal damage and potentially lethal damage are more confusing and less clear-cut for drugs than for radiation.

- Potentially lethal damage repair is a significant factor for bleomycin and doxorubicin, but sublethal damage repair is essentially absent. Neither potentially lethal nor sublethal damage repair is reported for the nitrosoureas.
The oxygen effect is more complex for drugs than for radiation.

Some drugs (e.g., bleomycin) are more toxic to aerated than to hypoxic cells. For these drugs, free radicals are involved in the mechanism of cell killing, as is the case for x-rays.

Some drugs (such as mitomycin C) are more toxic to hypoxic than to aerated cells because they undergo bioreduction. This applies also to tirapazamine, as discussed in Chapter 26.

Other drugs (including 5-FU, methotrexate, cis-platinum, and the nitrosoureas) appear to be equally cytotoxic to aerated and hypoxic cells.

The effectiveness of chemotherapeutic agents decreases with distance from a capillary because the drug concentration falls off because of metabolism and because cells are not proliferating because they are hypoxic.

The addition of a hypoxic cytotoxin to other chemotherapeutic agents may alleviate this problem. As the effectiveness of conventional agents falls off with distance from a capillary, hypoxic cytotoxins become more effective because they only function in low oxygen concentrations.

Drug resistance is the biggest single problem in chemotherapy. For example, cells exposed continuously to low levels of doxorubicin become very resistant to subsequent treatments with this drug.

The usual strategy to overcome resistance is to use a battery of drugs that produce cytotoxicity by diverse mechanisms.

Pleiotropic resistance occurs if the development of resistance to one drug results in cross-resistance to other drugs with a different mechanism of action.

Underlying acquired resistance are genetic changes.

Resistance may be associated with the following: decreased drug accumulation and the expression of p-glycoproteins in the cell membrane from gene amplification; elevated levels of glutathione; marked increase in DNA repair.

RADIORESISTANCE AND CHEMORESISTANCE

The adjunct use of chemotherapy with radiation may involve sequential or concurrent treatments.

The extent to which chemotherapeutic agents show synergy with radiation varies widely. Some interact strongly (e.g., doxorubicin, gemcitabine), others less so, and some not at all. See Table 27.1.

Some agents penetrate the blood–brain barrier (e.g., BCNU, temozolomide, cytosine arabinoside, hydroxurea), others only at high intravenous doses (e.g., methotrexate), whereas many do not cross the blood–brain barrier at all. See Table 27.1.

A therapeutic gain requires a differential between tumor and normal tissue. This may be achieved by exploiting one or more of the following tumor characteristics: genetic instability; rapid proliferation; cell age distribution; hypoxia; pH; elevated specific pathways (e.g., EGFR).

Sensitive cells can be killed only once. Tumor heterogeneity should be exploited by using a combination of drugs effective against different cell subpopulations.

Sensitivity of individual tumors to chemotherapeutic agents with or without radiation may be assessed by the following: *in vitro* clonogenic assays; xenografts in nude mice; micronuclei in treated cells.

**BIBLIOGRAPHY**


The medical manipulation of body temperature for treatment of disease has a long and fascinating history. Ancient references to the specific use of hyperthermia (or induced elevation of temperature above normal either locally, in part of the body, or of the whole individual) are found within the medical history of cultures from around the world. Modern interest in the cellular effects of heat energy by radiation oncologists was initially piqued by studies in the 1970s and 1980s demonstrating the highly quantifiable, time-dependent cytotoxic effects of “heat-shock” range temperatures (usually 41° to 45° C) on cells, similar to the highly reproducible and predictable effects of radiation. Moreover, hyperthermia in this temperature range was revealed to have clear additive, as well as synergistic, radiosensitizing properties. Although an early phase III trial in the United States in the 1980s testing the benefit of adding hyperthermia to radiation in patients with superficial tumors failed to show benefit, subsequent analysis of this trial revealed significant flaws in design, including the fact that many patients’ tumors were suboptimally heated at several of the participating centers. Indeed, since that time, multiple randomized trials examining the use of hyperthermia in combination with radiation and/or chemotherapy for a range of solid tumors, including deep tumors, have now been completed, demonstrating clear and significant clinical benefits. As a result, today, in some countries such as the Netherlands and Germany, hyperthermia is a fully approved and a reimbursable part of cancer care. The improved ability to control and measure thermal dose, in combination with a more complete recognition of the underlying biologic contribution of hyperthermia in radiochemosensitization, is currently driving a vigorous research effort, which will certainly further optimize the use of hyperthermia as an adjuvant for various oncologic and nononcologic disease targets.

For example, it is now appreciated that heating tumors in situ (and at temperatures as low as 39° to 42° C) can activate vascular, metabolic, and immunologic parameters of the tumor microenvironment, which may play an additional role in radiochemosensitization beyond hyperthermia-induced cell killing of tumor cells. Moreover, new functions for the focused delivery of heat in oncology are being identified, including as a highly specific means by which to concentrate cytotoxic drugs at the site of tumor using thermosensitive liposomes and other nanoparticles.
New image-guided approaches are now rapidly expanding the medical applications of thermal medicine. This includes the use of radiofrequency (RF) or ultrasound ablation technologies capable of focusing heat within tumors in situ to achieve very high (50° to 60° C) tumor temperatures over a very short interval, effectively instantly killing most tumor tissues without harming surrounding normal tissues. New nanoparticle technology including ferromagnetic fluids, nanotubes, and gold-coated particles are leading to completely new strategies for local heating of cells and tumors. In addition, this chapter also summarizes several recently completed clinical applications of hyperthermia with radiation and/or chemotherapy.

- CONTRIBUTION OF INDIRECT AND DIRECT EFFECTS OF HYPERThERmIA TO RADIATION SENSITIZATION

**Early Emphasis on the Direct Cytotoxic Effects of “Heat Shock” Temperatures**

The research, which led directly to initial clinical trials during the 1980s and 1990s that aimed to locally heat patients’ tumors to temperatures to approximately 43° C, focused on the observation that tumor cell damage from focused heat energy that could add to, or even synergize with, the antitumor effects of radiation without the systemic toxicity typical of chemical radiation sensitizers. When cells are heated in vitro to a sufficient temperature and for a long enough duration, they die in a predictable, exponential manner, and the rate of killing increases with temperature. Figure 28.1 shows a series of survival curves for cells exposed for various periods to a range of temperatures from 41.5° to 46.5° C. The cell survival curves for heat are similar in shape to those obtained for x-rays (i.e., an initial shoulder followed by an exponential region), except that the time of exposure to the elevated temperature replaces the absorbed dose of x-rays. For lower temperatures, the picture is complicated, because the survival curves flatten out after a protracted exposure to hyperthermia, possibly indicating the development of “thermotolerance” or an acquired resistance or tolerance to the elevated temperature. The similarity in the shape of the cell survival curves for heat energy and x-rays is misleading. It is important, therefore, not to draw conclusions for heat based on the interpretation of radiation dose-response curves, because the amount of energy involved in cell inactivation is a thousand times greater for heat than for x-rays. This reflects the different mechanisms involved in cell killing by heat and x-rays. Families of survival curves similar to those in Figure 28.1 have been obtained for many different cell types, and it is clear that cells differ somewhat in their sensitivity.

**FIGURE 28.1** Survival curves for mammalian cells in culture (Chinese hamster ovary line) heated at different temperatures for varying lengths of time. (Adapted from Dewey WC, Hopwood LE, Sapareto LA, et al. Cellular responses to combinations of hyperthermia and radiation. Radiology. 1997;123:463–474, with permission.)
to heat-induced cellular damage in this temperature range. However, as with ionizing radiation, there is no consistent difference between normal and malignant cells heated in vitro.

Survival data for Chinese hamster ovary cells exposed to various levels of hyperthermia (taken from Fig. 28.1) are replotted in Figure 28.2, with $1/D_0$ on the ordinate and $1/T$ on the abscissa. $T$ is the absolute temperature; $D_0$ is the reciprocal of the slope of the exponential region of the survival curve (i.e., the time at a given temperature that is necessary to reduce the fraction of surviving cells to 37% of their former value). This type of presentation is known as an Arrhenius plot; its slope gives the activation energy of the chemical process involved in the cell killing. The obvious change in slope, which is typical of Arrhenius plots, occurs at a temperature referred to as the “breakpoint” and is about 43°C for human cells. Above this temperature, an increase of 1°C doubles the rate of cell killing, and below the breakpoint, the rate of cell killing by heat drops by a factor of 2 to 4 for each drop of 1°C. The difference in activation energy above and below this temperature may reflect different mechanisms of cell killing (i.e., different targets for cytotoxicity above and below 43°C) or may reflect the development of thermotolerance within cells. Importantly, the slopes of Arrhenius plots derived from many in vitro and in vivo studies are nearly identical, and this consistency has provided long-standing usefulness of the Arrhenius analysis as a basis for assessing thermal dose in clinical hyperthermia applications up until the present time.

The similarity of the activation energy for protein denaturation to the activation energy for heat cytotoxicity, calculated from the Arrhenius analysis, led to the hypothesis that the target for heat cell killing in this temperature range resides in cellular proteins. Specifically, the heat of inactivation for cell killing and thermal damage is similar to the energy needed for protein denaturation (130–170 kcal/mol). Additional evidence for proteins as being the primary target for thermal cell killing is the importance of heat shock proteins (HSPs) in protecting thermotolerant cells from thermal damage. One of the primary functions of HSPs is to stabilize and refold other proteins that have been denatured or damaged, an event that signals their enhanced production. Structural chromosomal proteins, nuclear matrix, mitochondria and cytoskeleton, repair enzymes, transport proteins, and other membrane components are

**FIGURE 28.2** An Arrhenius plot for heat inactivation of mammalian cells in culture. The reciprocals of the $D_0$ values obtained for Figure 28.1 are plotted versus the reciprocal of the absolute temperature. (Adapted from Dewey WC, Hopwood LE, Sapareto LA, et al. Cellular responses to combinations of hyperthermia and radiation. *Radiology.* 1997;123:463–474, with permission.)
all included among targets that have been shown to be denatured in a time-dependent manner by temperatures between 42° and 45° C. Enzymes in the respiratory chain are more heat sensitive than enzymes in the glycolytic pathway.

Protein damage represents an important mechanistic difference between the manner in which heat and ionizing radiation lead to cellular damage, and this suggested that the two therapies could result in at least an additive ability to kill tumor cells if used together. There are other differences between irradiation and heating, suggesting that they could have additive effects. After irradiation, cells die only in attempting the next or a subsequent mitosis (except in very unusual circumstances), whereas heat-induced damage is expressed early, causing cells to die by various necrotic or apoptotic mechanisms, depending on the temperature. In addition, heat affects differentiating as well as dividing cells. Heat-induced chromosomal aberrations can occur if cells are in S phase during the time of heating. Although there is a familiar delay in x-ray response related to the time it takes for stem cells to progress through the process of differentiation and become functional, this delay is absent in the case of heat-induced cell death because all cells are affected, differentiated, or dividing; the thermal damage to the tissue is expressed immediately if the temperature is high enough. The DNA repair process itself is heat sensitive and this may be one of the mechanisms that lead to heat-induced radiosensitization and chemosensitization. Finally, although hypoxia renders cells resistant to radiation, it does not appear to affect sensitivity to heat.

**Thermal Enhancement Ratio**

The extent of the interaction of heat and radiation can be expressed in terms of the thermal enhancement ratio (TER), defined as the ratio of doses of x-rays required to produce a given level of biologic damage with and without the application of heat. The TER has been measured for various normal tissues, including skin, cartilage, and intestinal epithelium. The data form a consistent pattern of increasing TER with increasing temperature, up to a value of about 2 for a 1-hour heat treatment (HT) at 43° C. The TER is more difficult to measure in transplanted tumors in laboratory animals, because the direct cytotoxic effect of the heat tends to dominate. Heat often can control experimental tumors with acceptable damage to normal tissues, because cell killing by heat is strongly enhanced by nutritional deprivation and increased acidity, conditions that are typical of the poorly vascularized parts of solid tumors. Thus, a moderate HT, which can be tolerated by well-vascularized normal tissues, destroys a large proportion of the cells of many solid tumors in experimental animals. In those cases in which thermal radiosensitization has been studied, typical TER values are 1.4 at 41° C, 2.7 at 42.5° C, and 4.3 at 43° C, with heat applied for 1 hour. TERs have been developed for canine and human tumors however in studies by Gillette et al. and Overgaard et al. These estimates came from retrospective analyses of dose-effect relationships with and without hyperthermia. In canine oral squamous cell carcinomas, it was estimated to be approximately 1.15, when hyperthermia was administered twice a week during a course of fractionated radiotherapy. Importantly, the use of hyperthermia increased the slope of the TCD50 curve, such that the doses required to achieve a high level of local tumor control were maintained well below that which caused normal tissue damage (defined here as bone necrosis). TERs have also been estimated to be in the range of 1.5 for several superficial human tumor types.

**Heat and the Therapeutic Gain Factor**

The therapeutic gain factor can be defined as the ratio of the TER in the tumor to the TER in normal tissues. There is no advantage to using heat plus lower doses of x-rays if there is no therapeutic gain compared with the use of higher doses of x-rays alone. The question of a therapeutic gain factor is complicated in the case of heat, because the tumor and normal tissues are not necessarily at the same temperature. If the statement is made that heat preferentially damages tumor cells compared with normal tissue, it is implied that they are both at the same temperature. In a practical situation, however, this is not always the case. For example, if a poorly vascularized tumor is treated with microwaves, it may reach a higher temperature than the surrounding normal tissue, because less heat is carried away by the flow of blood. In addition, the overlying skin can be cooled actively by a draft of air or even a cold-water pack. In these circumstances, the normal tissues may be at a significantly lower temperature than the tumor, which therefore exaggerates the differential response in a favorable direction.
Factors that Modify Direct Cellular Damage from Hyperthermia

Because the earliest rationale for using heat as a radiation sensitizer involved maximizing direct heat-induced protein denaturation and cellular damage, there has been considerable attention by researchers given to factors that could modify tumor cell thermal sensitivity. For example, modification of cell membrane lipid content or use of membrane active agents, such as alcohols, was found to sensitize cells to heat killing, and the sensitization is probably related to destabilization of the membrane as it relates to lipid–protein interactions. Cells in an acid pH environment appear to be more sensitive to killing by heat. The pH dependence of cytotoxicity at elevated temperatures, however, is affected by pH history. Cells can adapt to pH changes and avoid the heat sensitivity shown for low pH.

Cells deficient in nutrients are quite heat sensitive. This can be demonstrated with cells in culture in which sensitivity to heat increases progressively as cells have their energy supply compromised, either by depriving them of glucose or by the use of a drug that uncouples oxidative phosphorylation. These conclusions about pH and nutrients, obtained under controlled conditions with cells in culture, led to the speculation that cells in tumors that are nutritionally deprived and at acid pH because of their location remote from a blood capillary may be particularly sensitive to heat. Because of their environment, it is likely that these cells are out of cycle and possibly hypoxic also. In this context, heat and x-rays would be predicted to be complementary in their action, because the cells that are most resistant to x-rays (out of cycle and hypoxic because of their remoteness from a capillary) show enhanced sensitivity to heat. A further enhancing factor that contributes to the complementary interaction of hyperthermia and radiation is that regions of a tumor in which the vasculature is poorly developed and potentially hypoxic tend to be at an elevated temperature because the cooling effect of blood flow is reduced.

Increased Recognition of the Importance of Indirect Effects of Hyperthermia: Heat and the Tumor Microenvironment

Some studies in mice had shown that local hyperthermia (using temperatures of 43° to 44° C) could naturally damage tumor vascular endothelium as well as tumor cells. This was predicted to lead to an increased accumulation of heat (because the damaged blood vessels would carry less heat away) while the pH, pO2, and nutrient status of the cells would decline, leading to enhanced tumor cell sensitivity to heat-induced cell killing. As a result of these studies, it was suggested that care was needed in combining heat and radiation, since tumor perfusion would be compromised by heating. For these reasons, several protocols have used hyperthermia in clinical applications after the first dose of radiation.

However, temperatures that can be achieved experimentally in mouse tumors are significantly higher than those which typically occur in human tumors under clinical conditions using locally applied, external heat energy; whether tumor vasculature in human tumors is damaged by the more modest temperatures reached under clinical conditions is still open to question. Further, it is important to also note that in animal studies in which more moderate temperatures are used (41° and 41.5° C), hyperthermia has been shown to promote tumor reoxygenation, with the degree of reoxygenation correlating with the level of the radiosensitivity of the tumor. Importantly, these data have been confirmed in a clinical study of patients with soft tissue sarcomas and breast cancer. Brizel and colleagues showed that one HT led to reoxygenation of human soft tissue sarcomas within 24 to 48 hours, whereas there was no measurable reoxygenation during a prior week of standard radiation therapy (RT). Jones and colleagues reported that mild hyperthermia (41° and 41.5° C at 90% of the measured points for 1 hour) significantly increased the pO2 in hypoxic, but not normoxic, human breast cancers. Such increases in tumor oxygenation could significantly improve tumor response to radiotherapy and is likely to be the primary important effect of local/regional or whole body forms of clinical hyperthermia. If this assumption is correct, then there should be much more attention devoted to optimal scheduling of hyperthermia in future trials to maximize oxygen-dependent radiosensitization. Indeed, instead of aiming to kill cells, mild hyperthermia may have the ability to rapidly stimulate global changes in the tumor microenvironment that collectively could enhance RT; this includes effects on pH, oxygen concentration, metabolism, protein/gene expression, and vascular perfusion.
It is well documented in human physiology that in normal tissues, a modest increase in temperature prompts highly sensitive thermoregulatory homeostatic mechanisms that increase heat dissipation in the vasculature. However, the ability to optimally thermoregulate is itself a highly temperature-dependent phenomenon. Optimal thermoregulation in terms of vascular effects occurs at mild to moderate hyperthermia and begins to fail above 41° and 42° C, resulting in heatstroke symptoms. While at temperatures of 43° C or higher, direct vascular and cellular damage can occur, and blood flow can be impaired. In cooler regions, the largest contributing factor to radiation sensitization by heating appears to be its ability to stimulate vascular activity, which, in turn, enhances delivery of additional oxygenated blood into tumors, sensitizing the tumor to radiation.

Despite an incomplete understanding of the actual basis of radiation sensitization by hyperthermia, an overestimation of the role of thermotolerance and suboptimal sequencing of hyperthermia and radiation fractions, multiple trials have been completed, revealing a critical role for hyperthermia in enhancing RT. A very active, preclinical, and clinical research effort is under way at the present time to develop the optimal dose and scheduling protocols and to design new clinical trials based on a more complete understanding of the physiologic effects of hyperthermia.

Thermotolerance

Exploration of the changing view of the role of “thermotolerance” in hyperthermia serves as a useful example demonstrating how our understanding of the radiosensitizing effects of hyperthermia have evolved over the past 2 to 3 decades. Thermotolerance, or induced thermal resistance, is usually described as the development of a transient and nonhereditary resistance to subsequent heating by an initial HT. Specifically, in vitro clonogenic assays of thermal cytotoxicity reveal that although one dose of heat kills a substantial fraction of cells, subsequent daily treatments are comparatively ineffective. Thermotolerance can begin to occur a few hours after the first treatment and may take as much as a week to decay. This would suggest that thermotolerance would be a significant problem clinically when heat is combined with fractionated radiotherapy, and most clinical trials were designed to minimize thermotolerance by keeping at least several days between hyperthermia fractions. However, it is now clear that in actual tumors in situ, an increase in blood flow and consequent increase in oxygenation by heating may significantly enhance the response of tumors to radiotherapy to a greater degree than any inhibition of thermal killing related to the occurrence of thermotolerance. Further, heat-induced radiosensitization is not subject to thermotolerance. Thus, although fears regarding thermotolerance had a historical importance regarding scheduling of hyperthermia for patients (schedules that may have turned out to be suboptimal in terms of achieving maximal benefits of hyperthermia), it is now clear that under normal heating conditions used in the clinic, thermotolerance may not be induced to any significant degree. Newer trials that are being designed to use more frequent applications of heating are likely to result in improved clinical benefit.

Another related area of research that has undergone considerable evolution in terms of understanding radiation sensitization by hyperthermia involves HSPs. It has been known for a long time that if cells are exposed to heat, proteins of a defined molecular weights or families are observed to be strongly induced above normal levels of expression. It was shown long ago that the appearance of these HSPs coincides with the development of thermotolerance, and their disappearance coincides with the decay of thermotolerance. Later molecular studies that specifically altered expression of certain HSPs demonstrated that they clearly exert a significant protective effect against potentially lethal heat exposures. Indeed, it has been long understood that HSP families are ubiquitous in evolution and are found in all living organisms. Similarly, a thermotolerance phenomenon also appears to be a conserved response in evolution resulting from their increased expression following exposure to certain stress conditions. However, although thermotolerance was one of the earliest functions attributed to HSPs, it is recognized today that these proteins play essential roles in numerous macromolecular processes under normal physiologic conditions as well as under stress conditions. In fact, there is now considerable research exploring induction of HSPs by hyperthermia treatments because of their likely role in promoting antitumor immune responses.
IMMUNOLOGIC EFFECTS OF HYPERThERMIA

One of the most recent directions in the field of thermal medicine is the exploration of whether thermal stimulation of the antitumor immune response contributes to improved local control or long-term survival of patients receiving treatments of hyperthermia. Historically, some of the earliest attempts to use immunotherapy in cancer are linked to the medical origins of hyperthermic oncology, because it was reported that cancer patients who experienced the highest fevers following injection with infectious agents (i.e., Coley toxins) in an attempt to treat tumors may have also experienced the longest survival. Although there are major and important differences between hyperthermia and fever, both involve an upward shift in body temperature, a phenomenon that has been strongly correlated with improved survival following infection in multiple species. One possibility that is being considered is that clinical hyperthermia may trigger some of the same thermally sensitive targets in the immune system that have evolved over millions of years to respond to fever. An exciting possibility is that strategically applied HTs may serve as an adjuvant, helping to overcome at least some of the tumor escape mechanisms used by tumors to overcome immune recognition and destruction.

Although much more work is needed, current evidence that supports this possibility includes (1) enhanced immunogenicity and HSP expression seen after tumor cells are heated, (2) thermally enhanced immune effector cell activation and function, and (3) thermally enhanced vascular perfusion and delivery or trafficking of immune effector cells to tumors (Fig. 28.3). Current research goals in this rapidly growing field include definition of the role of thermal stress-induced HSPs in mediating enhanced T lymphocyte responses, and establishment of heating protocols in which thermally regulated immune effector mechanisms could be maximally

![Mild hyperthermia diagram](https://example.com/mildhyperthermia.png)

**FIGURE 28.3** Mild hyperthermia can affect multiple aspects of the antitumor immune system (From Peer, Grimm MJ, Zynda ER, et al. Diverse immune mechanisms may contribute to the survival benefit seen in cancer patients receiving hyperthermia. *Immunol Res*. 2010;46:137–154, with permission.)
solve the problem of how to normalize the time–temperature data, which was seen to vary significantly from patient to patient. Sapareto and Dewey proposed the concept of “cumulative equivalent minutes” (CEM) at 43°C. More specifically, the measure of thermal dose stated as CEM 43°C T90 refers to the number of CEM at 43°C exceeded by 90% of the monitored points within the tumor. For this analysis, 43°C was chosen because it represents the breakpoint temperature for most human cells as judged from the Arrhenius plot. How does this idea work in practice? Because it is generally agreed that the effects of a 1°C rise of temperature is equivalent to a reduction of time by a factor of 2, consequently, above this transition temperature,

\[
\frac{t_2}{t_1} = 2^{T_1 - T_2}
\]

in which \(t_1\) and \(t_2\) are the heating times at temperatures \(T_1\) and \(T_2\), respectively, to produce equal biologic effect. For temperatures lower than the transition temperature, an increase in temperature by 1°C requires that time be decreased by a factor of 4 to 6:

\[
\frac{t_1}{t_2} = (4 \text{ to } 6)^{T_1 - T_2}
\]

The CEM 43°C T90, or the thermal dose, may be calculated from one of the other of these expressions or a combination of both: That is, the heat dose associated with a changing temperature may be calculated as the sum of equivalent heating times at 43°C for each temperature. Thus,

\[
\text{CEM 43°C} = tR^{(43-T)}
\]

where CEM 43°C refers to the cumulative equivalent minutes at 43°C (the temperature suggested for normalization), \(t\) is the time of treatment, \(T\) is the average temperature during desired interval of heating, and \(R\) is a constant. When above the breakpoint, \(R\) is 0.5. When below the breakpoint, \(R\) is 0.25. For a complex time–temperature history, the heating profile is broken into short intervals of time “t” length (typically 1 to 2 minutes), where the temperature remains relatively constant. CEM 43°C is calculated for each interval and summed to give a final CEM 43°C for the entire HT:

\[
\text{CEM 43°C} = \sum tR^{(43-T_{avg})}
\]

CEM 43°C, also known as the “thermal iso-effect dose formulation, has now been used to...
in addition to the ability to monitor resultant tumor temperatures over time, varied enormously at the various centers that participated in this trial. Despite this early setback, basic, translational, and clinical research in the field of thermal medicine continued, as well as research on the physics and engineering principles associated with optimal heat delivery to tumors. Further, significant progress in the ability to measure accurately the resultant tumor temperature over time was achieved. Although much more work is needed, it is indeed remarkable that there are now at least 10 randomized clinical trials that demonstrate clear benefits to hyperthermia as an adjuvant to radiation for the treatment of various types of cancers, including superficial and deep tumors, and involving both palliative and potentially curable settings. Moreover, several additional phase III trials provide strong evidence for combining hyperthermia with chemoradiation, or chemotherapy. Today, in some countries such as the Netherlands and Germany, hyperthermia is a fully approved and a reimbursable part of cancer care. Much can be learned from these trials by those interested in radiation oncology, and several of the most important phase III trials showing clear enhancement of radiation by hyperthermia are presented here.

Carcinoma of the cervix: A trial that contributed significantly to the fact that hyperthermia is added as part of standard care for some cancers in parts of Europe is a phase III trial done by a Dutch group, which tested the use of radiation alone versus radiation and hyperthermia in patients with locally advanced pelvic tumors (cervical carcinoma, as well as bladder and rectal tumors were included) in a randomized study that involved 361 patients. In this study, HT were given once weekly for five treatments. It is also important to note here that although some had considered that hyperthermia would only be useful for superficial tumors, the heating technology used here resulted in adequate deep heating capable of heating pelvic tumors. Although improved responses were seen in all tumor types, most of the benefit appeared to occur in patients with cervical cancer. The complete response (CR) rate following RT + HT was 83% compared with 57% after RT alone. Three-year survival was 27% in the RT-alone group of

- PHASE III CLINICAL TRIALS TESTING BENEFIT OF HYPERTERMIA FOR ENHANCING RADIATION THERAPY

Early studies showed that the simultaneous application of heat and ionizing radiation maximizes thermoradiosensitization; simultaneous thermoradiotherapy should cause more cell killing with mild temperatures than does sequential delivery. However, because of logistical and practical issues, most clinical trials have sequenced heat before or after radiation, not at the same time.

Although an early Radiation Therapy Oncology Group phase III trial that included patients with various types of superficial tumors yielded well-publicized disappointing results in the late 1980s (showing only a modest benefit of addition of hyperthermia and only in patients with tumors that were less than 3 cm), it was quickly realized that the ability to actually heat tumors,
Superficial malignancies: 
Recurrent chest wall breast cancer: Managing and treating recurrences of breast cancer in the chest wall present a significant challenge to both patients and their clinicians. However, in this setting, hyperthermia appears to contribute significant benefits and new clinical trials are definitely warranted as soon as possible. Five separate phase III trials have been conducted, which were eventually combined as an international collaborative study. In this study, patients were randomized to either RT alone or RT with HT, and a significant improvement in CR rate was seen for patients receiving HT + RT compared with RT alone. It is notable that the greatest effect was observed in patients with lesions in areas that had previously been irradiated, thus limiting the ability of those patients to receive optimal subsequent irradiation. Related to this trial, data from a seven-institution retrospective review of treatment for chest wall recurrence was presented at the American Society for Therapeutic Radiology and Oncology 2006 meeting. Although these data are retrospective, the complete response rate was 67% in those treated with hyperthermia and radiation versus 31% treated with radiation alone.

Superficial malignancies: An important single-institution prospective randomized trial of radiation and hyperthermia for various superficial tumors (most of which were recurrent chest wall breast tumors) was conducted by Jones et al. and included patients who were candidates for local RT for a superficial lesion less than 3 cm in thickness. In this study, the investigators decided to first determine whether a tumor was “heatable” prior to randomization by conducting a test hyperthermia treatment and measuring whether the tumor could be heated to a prescribed temperature ahead of time. All patients whose tumors were judged to be heatable were later randomized to receive radiation alone (with no additional heat) or radiation combined with hyperthermia. The complete response rate in the hyperthermia/radiation group was 66% versus only 42% in the radiation-alone group. Again, previously irradiated patients had the greatest benefit, enjoying a 68.2% response rate in the hyperthermia/radiation group versus 23.5% in the radiation-alone group (Fig. 28.4). This was the first study to capitalize on the idea that for hyperthermia to act as a radiation sensitizer, the tumor must be able to be heated, and it will surely help to define the need for improved pretreatment identification of the patients most likely to benefit from hyperthermia in the future. Moreover, as discussed previously, this trial provides strong evidence that although heating is designed to bring the tumor to at least 43°C, the resultant CEM 43°C T90 is significantly lower, suggesting that mild hyperthermia can have significant radiosensitizing effects, which may not be caused by thermal cytotoxicity.

A second phase III trial, which has been completed, that prospectively evaluated heatability prior to randomization is an important canine trial conducted by Thrall et al., conducted on pet dogs with spontaneous soft tissue sarcomas; again, the data from this trial confirm the benefit of the heatability of tumors with respect to clinical benefits.

Head and neck cancer: Head and neck cancers are significantly more likely to respond better if hyperthermia is added to RT. In studies conducted by Valdagni et al., patients in stage III receiving RT + HT had a 58% CR compared with 20% in the RT group. Similarly, patients with stage IV disease achieved a CR of 38% compared with 7% for those receiving RT alone. Despite the difficulties encountered when trying to heat tumors in the head and neck region, this trial evaluated both the primary site as well as neck nodes. In another trial by this group in which only
an important research area for extending hyperthermia treatment to the brain.

**Clinical Trials Assessing the Benefit of Hyperthermia in Combination with Chemotherapeutic Agents**

As with RT, there are now several very compelling clinical demonstrations of the benefit of adding hyperthermia to chemotherapy protocols, and understanding the underlying rationale is a topic of much current research. In several fundamental ways, enhancement of efficacy of both radiation and chemotherapy by heating is likely to involve overlapping mechanisms. *In vitro* data reveal the potential for enhanced chemosensitization for several chemotherapeutic agents by increased temperature, even of only 1° to 2° C. Agents that have been shown to synergize with hyperthermia include melphalan, cisplatin and related compounds, anthracyclines, bleomycin, mitomycin C, nitrosoureas, and nitrogen mustards. Moreover, hypoxic cell sensitizers have

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**FIGURE 28.4** Results of a randomized phase III trial comparing efficacy of delivering a low versus high cumulative thermal dose in combination with radiotherapy for superficial tumors (primarily chest wall recurrences of breast cancer). **A:** Clinical trial design. **B:** Duration of local control in patients with superficial tumors treated with a low versus high cumulative thermal dose combined with RT. The difference in CR rate was significant as well as duration of local control following administration of a higher cumulative thermal dose. (From Jones EL, Oleson JR, Prosnitz LR, et al. Randomized trial of hyperthermia and radiation for superficial tumors. *J Clin Oncol.* 2005;23:3079–3085, with permission.)
been shown to synergize with heat. Table 28.1 lists some drugs that are potentiated by heat and some that are not; however, more work is needed in this area, particularly on assessment of the efficacy of a wider range of temperatures in vivo. There may be several different mechanisms that underlie the interaction of heat with chemotherapeutic drugs in vivo. These include (1) increased drug uptake and/or retention in cells, (2) increased DNA damage and inhibition of repair processes, (3) increased oxygen radical formation, and (4) increased vascular delivery and tumor penetration. It is also clear that the extent of hypoxia and the pH of the tumor may affect the interaction between heat and chemotherapy. Whatever the mechanisms involved, the enhancement by hyperthermia of the efficacy of several different major classes of chemotherapeutic drugs may prove very useful in the chemotherapies of solid tumors. Further, there is a clear rationale for testing, as soon as possible, other types of cancer therapies that depend on vascular delivery and/or cellular sensitivity or uptake of drug, such as small molecule inhibitors and monoclonal antibodies. Several newer and informative clinical trials are presented here.

### Table 28.1 Interaction of Heat and Chemotherapeutic Agents

<table>
<thead>
<tr>
<th>Effect</th>
<th>Drug</th>
</tr>
</thead>
<tbody>
<tr>
<td>Potentiated by heat</td>
<td>Melphalan</td>
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<tr>
<td></td>
<td>Cyclophosphamide</td>
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<tr>
<td></td>
<td>BCNU</td>
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<tr>
<td></td>
<td><em>Cis</em>-DDP</td>
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<td></td>
<td>Mitomycin C</td>
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<td></td>
<td>Bleomycin</td>
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<td></td>
<td>Vincristine</td>
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<tr>
<td>Unaffected by heat</td>
<td>Hydroxyurea</td>
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<tr>
<td></td>
<td>Methotrexate</td>
</tr>
<tr>
<td></td>
<td>Vinblastine</td>
</tr>
<tr>
<td>Complex interaction</td>
<td>Doxorubicin</td>
</tr>
</tbody>
</table>

BCNU, bischloroethylnitrosourea; *Cis*-DDP, *cis*-diamminedichloroplatinum.


**Esophageal cancer**: Two randomized studies by Sugimachi et al. have now demonstrated a significant clinical advantage by adding hyperthermia to CT alone or to chemoradiotherapy during neoadjuvant treatment of squamous cell carcinoma of the esophagus. In one study, patients were randomized to preoperative hyperthermia, RT, and chemotherapy (bleomycin), compared with radiation and chemotherapy alone. Clinical complete responses as well as pathologic responses were significantly improved in the trimodality arm. In a follow-up study, patients were treated with CT alone (bleomycin and cisplatin) or combined with hyperthermia (i.e., no RT was given in this trial). A significant improvement in histopathologic response was noted in the group receiving hyperthermia (19% versus 41%).

**High-risk sarcoma**: In new clinical data recently reported by Issels and colleagues in a randomized phase III trial involving 341 patients with localized high-risk soft tissue sarcoma (the largest phase III trial for any evaluation of therapy for sarcoma), it was revealed that heating the tumor and the surrounding area (i.e., deep regional or part-body heating at a range between 40° to 43° C) during chemotherapy can significantly improve tumor control, including those in the extremities (Fig. 28.5). Specifically, regional hyperthermia during induction is associated with a 42% reduction in risk of local progression or death compared with chemotherapy alone. Further, there was a 30% improvement in disease-free survival and an improved treatment response rate in the combined heat–chemotherapy group. In this impressive study, patients were randomized to neoadjuvant chemotherapy (etoposide, ifosfamide, and doxorubicin) alone or combined with regional hyperthermia achieved by using RF for 60 minutes on days 1 and 4 of each chemotherapy cycle during induction and postinduction therapy. Importantly, this trial demonstrates the fact that a nonuniform heating field can be very successful (the temperature across the tumor varied over 1° to 3° C), which should simplify the design of heating equipment for future clinical applications. Although determination of whether there are long-term survival benefits from the addition of hyperthermia must await future analysis, recently...
FIGURE 28.5 Results of a phase III trial in patients with high-risk sarcoma. (A) Schematic of the trial design. Patients with previous surgery were also assigned to receive the complete induction and postinduction therapy for randomization. EIA, etoposide + ifosfamide + doxorubicin; RHT, regional hyperthermia. (B) Kaplan-Meier estimates of local progression-free survival, (C) disease-free survival, (D) overall survival in all patients randomly allocated treatment, and (E) overall survival in the per-protocol-induction population. (From Issels RD, Lindner LH, Verweij J, et al. Neo-adjuvant chemotherapy alone or with regional hyperthermia for localised high-risk soft-tissue sarcoma: a randomised phase 3 multicentre study. Lancet Oncol. 2010;11:561–570, with permission.)
completed interim analysis does suggest the possibility of improved long-term survival. Importantly, a new phase III trial that will include the comparison of presurgery and postsurgery radiation along with hyperthermia has been initiated. Because radiotherapy is most often performed in the United States for patients with high-risk sarcoma, a demonstration of the benefits of hyperthermia in this setting should help to increase interest in evaluating the benefits of hyperthermia as an adjuvant for other disease settings, in addition to sarcoma.

**Development and Evaluation of “Thermosensitive” Liposomes for Improved Tumor Targeting of Chemotherapy**

The addition of local hyperthermia to systemically delivered chemotherapy may have the obvious advantage of “targeting” and localizing the principal effect of the drug, allowing greater tumor cell killing for a given systemic toxicity. This would help to overcome one of the principal problems and limitations of chemotherapy. This possibility is being realized in exciting new preclinical and clinical research by Dewhirst and colleagues in which hyperthermia is being used to direct the release of chemotherapeutic drugs at the site of tumors through the use of thermosensitive liposomes. Liposomes consist of a lipid membrane that can be filled with a cytotoxic chemotherapeutic agent such as doxorubicin. Long circulating liposomes that are covered with polyethylene glycol reduce recognition by the reticuloendothelial system and can extravasate from tumor blood vessels and accumulate in the extravascular space. This is termed the enhanced permeability and retention (EPR) effect. Whereas this effect can passively lead to enhanced drug accumulation in tumors by virtue of the presence of endothelial cell membrane pores, hyperthermia can increase the size of these pores and enhance drug delivery by 4 to 5 fold over what can be achieved from just the EPR effect. Further enhancement of drug delivery can be achieved by using thermally sensitive liposomes that are engineered to melt at mild temperatures (e.g., 41° to 42° C). These formulations yield further improvement in drug delivery by an additional factor of 4 to 5 versus nonthermally sensitive liposomes. Indeed, these latter formulations have been shown to enhance doxorubicin delivery by 25 to 30 fold, compared with free drug, resulting in impressive antitumor effects in tumors that were completely refractory to free drug. It is speculated that the combination of hyperthermia-enhanced liposomal extravasation and intravascular-triggered drug release results in enhanced delivery to tumor cells and a large therapeutic effect. The mechanism is illustrated in Figure 28.6. More recent studies by this group...

**FIGURE 28.6** Characteristics of thermally sensitive liposome. **A:** The liposome contains a mixture of phospholipids including a small percentage of single-chain fatty acid. At body temperature, the liposome is in a frozen or gel state. The single-chain fatty acid tends to concentrate between plates of frozen lipid. When heated, the grain boundaries melt first, the single-chain fatty acids line the grain boundaries, creating pores in the lipid membrane. This permits very rapid release of drug (doxorubicin in this example) from the liposome (<20s). **B:** In a heated tumor, this feature promotes deposition of high amounts of drug because of intravascular release, which drives the drug far into the interstitial tissue, down its concentration gradient. (Panel B – from Gong G, et al. Efficacy of Liposomes and Hyperthermia in a Human Tumor Xenograft Model: Importance of Triggered Drug Release. Cancer Res 2000;60:6950-6957, with permission.)
are helping to establish more details by which li-posomally encapsulated doxorubicin may work in combination with hyperthermia. For example, it has been determined that although extracellu-lar pH or *in vitro* sensitivity to doxorubicin does not appear to predict efficacy *in vivo*, tumor cell doubling time, vascular perfusion, and extent of hypoxia do appear to correlate. Clinical validation of this strategy is now well under way for patients with recurrent chest wall cancers and hepatocellular carcinomas. In the latter setting, thermosensitive liposomes are being combined with high temperature RF ablation. Here, the strategy is designed to increase the zone of tumor destruction particularly at the tumor margins but capitalizing on the ability of the mild hyper-thermia that occurs in this region to release drug from the liposomes.

**Isolated Limb Perfusion, Limb Infusion, and Hyperthermic Intraperitoneal Chemotherapy**

Mild hyperthermia is also finding a very useful role in important surgical modalities that employ chemotherapy: isolated limb perfusion (ILP), isolated limb infusion (ILI), and intraperitoneal chemotherapy. ILP (Fig. 28.7) is a technique that aims to avoid amputation of limbs containing bulky tumors or small multiple tumors (e.g., in transit metastases of melanoma) by achieving regional drug concentrations 15 to 25 times higher than systemic administration of the drug, which helps to reduce systemic side effects while exposing tumors to highest possible concentrations of drugs. ILP is generally achieved by clamping and cannulation of major blood vessels, connecting an oxygenated extracorporeal circuit, ligating collateral vessels, and applying a tourniquet. Melphalan (L-phenylalanine mustard) has become the most commonly used drug for this procedure; however, the addition of tumor necrosis factor-alpha (TNF-α) has been demonstrated to be critical for clinical efficacy because of its ability to enhance tumor vascular uptake of drug and has helped lead to approval for this therapy in Europe. ILP also incorporates very mild hyperthermia (39° to 40° C) because it has become clear that keeping the limb slightly warm during the treatment can stimulate vascular function and help with drug delivery to the interior of large tumors. However, if the temperature is too warm (i.e., above 41° C), there is increased local toxicity limiting the benefits of ILP. Mild hyperthermia is also employed in a surgical modification of ILP termed ILL, a simpler procedure that does not require arterial and venous catheters to be inserted through the surgical incision, thus resulting in lower morbidity.

**Hyperthermic Intraperitoneal Chemotherapy**

Many patients are at high risk for tumor growth in the abdomen from microscopic residual disease that remains following surgery, or have tumors that involve the ovaries or abdominal organs, such as the pancreas, or have cancer cells in the ascites fluid. In these cases, the intraoperative use of chemotherapy is being combined with hyperthermia by heating of the chemotherapy solution prior to its addition to the abdominal cavity, where organs are exposed to the heated solution (usually at approximately 40° C) for various periods. Heat is thought to potentiate the effect of chemotherapy
by reducing tumor cell resistance. Several new clinical trials are demonstrating improved survival and improvements in overall quality of life when hyperthermic intraperitoneal chemotherapy is used in comparison to conventional chemotherapy. However, additional work is needed to clarify the exact benefit of hyperthermia in this setting.

Systemic or Whole Body Applications

In addition to deep regional, or part-body hyperthermia studies, other phase I and phase II clinical studies from Bull and colleagues are demonstrating the feasibility and safety of using mild temperature whole body treatments using temperatures near 39° to 40° C in combination with chemotherapy. Although more work is needed on the development and testing of systemic heat therapies, it is likely that the mechanisms of action of whole body and part-body hyperthermia protocols overlap in terms of their effects on chemotherapeutic efficacy. Furthermore, the whole body or systemic approach may have the advantage of allowing treatment of widely disseminated tumors in metastatic patients, with the same HT.

MAJOR METHODS OF TUMOR HEATING

Most of the original laboratory research that led to clinical applications of local hyperthermia used cells cultured in vitro usually heated by hot-water baths. Indeed, the simplest and most reliable way to heat cells in a petri dish, or to heat a tumor grown in the leg of a mouse or rat, is to immerse the target in a thermostatically controlled bath of water. This approach has been used most often in the preclinical research in hyperthermia over the past 3 decades. Water temperature can be controlled within a fraction of a degree, and temperature measurement involves no problem. However, even in this simplest of situations for rodent tumor heating, the tumor may not be at the same temperature as the skin (or water bath) because the heated tumor is constantly drained and resupplied by cooler blood coming from the nonheated regions of the body. Moreover, in water-bath applications, rodents usually need to be anesthetized, and this will significantly limit the role of neurovascular regulation in response to tissue heating and will create abnormal temperature gradients because of significant heat loss during general anesthesia. Because the core temperature of mice is very easily and quickly adjusted (because of their large surface to volume ratio), another convenient method developed by Repasky and colleagues simply uses warm air incubators to systematically change the temperature of mice bearing tumors. This protocol does not require anesthetization and results in rapid core body temperature increases that is likely to incur additional thermoregulatory vascular responses. Another protocol used recently by Bull and colleagues to mimic systemic heating involves anesthetized rats that are largely immersed in warm water. The “optimal” heating protocol for rodent tumors, which can best mimic the situation that may occur in patients, should be dependent on the question being addressed and on the type of tumor or heating method being studied. Because of fundamental differences in thermoregulatory mechanisms between rodents and humans, it is also likely that no rodent model is completely suitable for mimicking human hyperthermia treatments, which makes the clinical studies done on larger canine models by Thrall et al. especially valuable. In any case, unlike the relatively simple situation of heating cultured cells (or even rodents bearing tumors), the design of equipment to heat patients’ tumors accurately, without damaging surrounding normal tissues, has required significant engineering efforts, which continue to advance at the current time.

For localized hyperthermia in human tumors to be achieved by microwaves, RF-induced currents, or ultrasound, there have been some serious technical problems that have had to be overcome. In the case of microwaves, good localization can be achieved at shallow depths, but at greater tumor depths, even if the frequency is lowered to allow deeper penetration, the localization is much poorer and surface heating limits therapy. If ultrasound is used, the presence of bone or air cavities causes distortions of the heating pattern, but adequate penetration and good uniform temperature distributions can be achieved in soft tissues, particularly with ultrasound in focused arrays. In practice, then, superficial tumors such as recurrent chest wall nodules can be treated adequately with microwaves or plane wave ultrasound, and it should theoretically be possible to heat deep-seated tumors below the diaphragm with focused ultrasound, regional microwave devices, or interstitial techniques.

In all cases, however, methods of heating pose a complex problem, although significant progress has been made as a result of the clever application
of focused arrays. For example, the use of multi-phased arrays allows uniformity of temperature to the edge of the field. The picture is complicated, and it is unlikely that one simple answer to all the complex problems will be found. This is a highly active area at the present time, and the following summary of available heating systems is likely to change in the near future as new breakthroughs occur. Because of space, the summary provided here is limited to devices that are already in use clinically or that have the potential to become commercially available in the near future. Indeed, there have been a great number of other systems developed and tested over the last 3 decades, but few had been produced commercially in part because of the U.S. Food and Drug Administration classification of hyperthermia devices.

**PROGRESS IN CLINICAL THERMOMETRY**

Probably no area in thermal engineering research is more important than the need to accurately assess the resultant temperature of tumors or normal tissues following heating. This is essential for accurate comparison of clinical trial data using different heating equipment or different tumor targets.

**Invasive Thermometry Methods**

Direct measurement of temperatures during clinical hyperthermia with invasive sensors was for many years the only method of treatment monitoring, control, and thermal dose calculation. Sensors should be able to measure temperatures to an accuracy and precision of about 0.1° C in a water bath and of about 0.2° C in vivo in intense electromagnetic (EM) or ultrasound fields.

There are three basic kinds of invasive thermometers: (1) electrically conducting, (2) minimally conducting, and (3) nonconducting (optical) sensors. Thermistors and thermocouple sensors with metallic leads are examples of the first kind that are not suitable for EM hyperthermia. A commonly used minimally conducting sensor is the high-resistivity thermistor with carbon-impregnated plastic leads. The high-resistivity material allows accurate measurement in strong EM environment. Nonconducting optical sensors use materials at the tip of an optical fiber whose optical properties are a known function of temperature. The second and third kinds of sensors are used in EM hyperthermia.

Incorrect temperature measurements are possible because of direct heating of either the sensor or the catheter in which the sensor is placed. Such errors have been eliminated for EM systems by the use of high-resistive thermistors and optical sensors. For ultrasound hyperthermia, however, artifacts of several degrees are possible because acoustic waves can be absorbed by the catheter or sensor-coating material, and/or because of frictional (viscous) heating of sensors. In addition, thermal conduction heat transfer along metallic leads can affect the accuracy of the measurement. In general, thin needle thermocouple sensors and the measurement of temperature during short interruptions of ultrasound power minimize these ultrasound artifacts. A sufficient number of thermometry sensors are important for accurate evaluation of the quality of hyperthermia treatment and calculation of the thermal dose delivered. In general, the more locations monitored during treatment, the better the evaluation and thermal dose estimate. To this end, multisensor probes, thermal mapping techniques, and quality assurance guidelines have been implemented. Moreover, advanced heating systems demand extensive thermometry for effective utilization of temperature feedback power control.

However, an important clinical reality is that only a limited number of invasive catheters are desired or feasible. For further improvements in hyperthermia delivery, monitoring, control, assessment, and quality, a more complete knowledge of the temperature field in humans during treatment is necessary. This has motivated a great deal of research in noninvasive thermometry methods.

**Progress toward Clinically Achievable Noninvasive Thermometry**

One of the most sought-after breakthroughs in the hyperthermia field is achieving control and measurement of temperature in living tissue in patients, particularly a noninvasive technology that could provide detailed temperature distributions during treatment. Several noninvasive thermal measurement approaches are under investigation, including infrared thermography and thermal monitoring sheet fiberoptic arrays (surface distributions only), electrical impedance tomography, microwave tomography, microwave radiometry, ultrasonic temperature estimation techniques, and magnetic resonance thermal imaging (MRTI). The most promising approach to date
Currently being used in combination with radiation and chemotherapy applications, thermal tumor ablation is becoming a major clinically used form of hyperthermia in which tumors (liver, kidney, lung, bone, adrenal gland, or prostate) are heated for short intervals at high temperatures between 50° C and 100° C to thermally “ablate” nodules. At 50° C, it takes a few minutes to kill cells; above 60° C, it takes only seconds. Moreover, there are now clinical trials in place in which thermal ablation is being combined with more traditional applications of hyperthermia to improve drug delivery through the use of thermosensitive liposomes at the margins of ablated tumors. Ablative heating is produced by needle-type RF or microwave applicators (~1.5 mm diameter) inserted into the tumor under imaging guidance (CT or ultrasound), and RF electric current or microwaves heat tissue surrounding the applicator. The zone of tissue coagulation grows as heat is thermally conducted from the hot region close to the applicator into the tissue. Blood flow ceases inside the coagulation zone, reducing effects of perfusion compared to mild hyperthermia, and final coagulation zones are ~3 to 5 cm in diameter, depending on device.

Tumor ablation can be performed minimally invasively by an interventional radiologist or during surgery (laparoscopy or open surgery [Fig. 28.8]). Long-term results from primary and metastatic liver cancer show promising results.

**Thermal Ablation Applications in Thermal Medicine**

Although the major goal of this chapter is to focus on hyperthermia treatments that are currently being used in combination with radiation and chemotherapy applications, thermal tumor ablation is becoming a major clinically used form of hyperthermia in which tumors (liver, kidney, lung, bone, adrenal gland, or prostate) are heated for short intervals at high temperatures between 50° C and 100° C to thermally “ablate” nodules. At 50° C, it takes a few minutes to kill cells; above 60° C, it takes only seconds. Moreover, there are now clinical trials in place in which thermal ablation is being combined with more traditional applications of hyperthermia to improve drug delivery through the use of thermosensitive liposomes at the margins of ablated tumors. Ablative heating is produced by needle-type RF or microwave applicators (~1.5 mm diameter) inserted into the tumor under imaging guidance (CT or ultrasound), and RF electric current or microwaves heat tissue surrounding the applicator. The zone of tissue coagulation grows as heat is thermally conducted from the hot region close to the applicator into the tissue. Blood flow ceases inside the coagulation zone, reducing effects of perfusion compared to mild hyperthermia, and final coagulation zones are ~3 to 5 cm in diameter, depending on device.

Tumor ablation can be performed minimally invasively by an interventional radiologist or during surgery (laparoscopy or open surgery [Fig. 28.8]). Long-term results from primary and metastatic liver cancer show promising results.
thermotolerance plays a significant role in modulating resistance to heating in vivo.

Hyperthermia at a sufficient temperature and duration of heating induces quantifiable damaging effects in both the nucleus and the cytoplasm of cells. The Arrhenius plot continues to be useful in assessing effects of thermal damage in tissue. The mechanisms of thermal damage may be different above and below the break temperature in the Arrhenius plot.

Thermal ablation technology using temperatures between 50° and 60° C has significantly advanced the concept of using thermal energy to kill the largest number of tumor cells in the shortest period, whereas lower tumor temperatures may be most effective for simultaneously changing several critical parameters within the tumor microenvironment including vascular perfusion, oxygenation state, pH, and metabolism. Each of these represents important targets that can greatly affect the efficacy of radiation and chemotherapy. Immune sensitization by heat may also contribute significantly to the adjuvant potential of various forms of hyperthermia and may lead to novel combinations of heat and immunotherapy.

Several phase III randomized trials now demonstrate a clear and significant benefit for the addition of hyperthermia to standard RT and/or chemotherapy. The cancer types tested include cervical cancer, superficial localized breast cancer, recurrent or metastatic malignant melanoma, nodal metastases from head and neck cancer, glioma, esophageal cancer, and high-risk sarcoma. Although results from recent trials involving hyperthermia are very encouraging, it is clear that many more trials should be conducted. Further improvements in clinical outcome could be seen with trial designs that are based on a more complete understanding of the biologic effects of heating and on tumor-type specific optimization of heating technology and thermometry.

Prospective control of thermal dose reveals that if a tumor is not sufficiently “heatable,” the patient may not benefit from hyperthermia. Thus, as is true for several other forms of cancer therapies, identification of biomarkers of heating efficacy will

SUMMARY OF PERTINENT CONCLUSIONS

- Survival curves for heat are similar in shape to those for radiation, except that time at the elevated temperature replaces absorbed dose. No consistent difference in inherent sensitivity exists between normal and malignant cells. The age-response function for heat complements that for x-rays. S phase cells that are resistant to x-rays are sensitive to heat.
- Cells at low pH and nutritionally deprived (more likely to be in tumors) are more sensitive to heat, although cells can adapt to pH changes and lose their sensitivity to heat.
- Hypoxia does not protect cells from heat as it does from x-rays. Hyperthermia may be very effective at reversing hypoxia through its effects on vascular perfusion. This may be an important mechanism of radiosensitization by hyperthermia.
- Thermotolerance is the induced resistance to a second heat exposure by prior heating and may be monitored by the appearance of HSPs. It is increasingly unlikely that
help in improved selection of patients who can most benefit by the addition of hyperthermia as an adjuvant.

- The TER is the ratio of radiation doses with and without heat to produce the same biologic effects. TER values of 2 to 4 can be obtained in tumors and normal tissues in experimental animals and has been reported to be between 1.15 and 1.5 in clinical analyses. With improved methods of heating and temperature monitoring, it is likely that TER values in the clinic will be larger in the future.

- Temperature measurement in vivo is difficult but improving. Progress toward noninvasive thermometry includes the use of magnetic resonance imaging.

- Novel applications of heat include enhanced targeted delivery of chemotherapy by thermosensitive liposomes and in combination with gene therapy applications. Increased use of ILP and intraperitoneal chemotherapy for specific types of tumors are continuing to add data regarding the usefulness of hyperthermia in combination with chemotherapy.

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**Biologic changes and action level:** Concentration of radon in a house above which it is recommended that some action be taken to reduce it. Currently, the action level is 148 Bq/m³.

**Absorbed dose (D):** The energy imparted per unit mass by ionizing radiation to matter at a specific point. The SI unit of absorbed dose is joule per kilogram (J/kg). The special name for this unit is gray (Gy). The previously used special unit of absorbed dose, the rad, was defined to be an energy absorption of 100 erg/g. Thus, 1 Gy = 100 rad.

**Absorption:** Removal of γ-rays from a beam.

**Accelerated fractionation:** A treatment method used in addition to radiation or chemotherapy. It utilizes a schedule in which the average rate of dose delivery exceeds the equivalent of 10 Gy per week in 2-Gy fractions.

**Accelerated proliferation:** Increase in the clonogen proliferation rate after treatment with radiation or chemotherapy, relative to its pretreatment value.

**Accelerator (linear):** A machine, often called a linac, that produces high-energy X-rays for the treatment of cancer.

**Acidic fibroblast growth factor (FGF):** A mitogen for many types of cells of mesodermal origin, including endothelial cells, chondrocytes, and fibroblasts. In presence of heparin, induces blood vessel growth.

**Action level:** Concentration of radon in a house above which it is recommended that some action be taken to reduce that radon level; currently, the action level is 148 Bq/m³ in the United States.

**Activation:** The process of making a material radioactive by bombardment with neutrons, protons, or other nuclear particles. See activation analysis.

**Activation analysis:** Identifying and measuring chemical elements in a sample of material. First, the sample is made radioactive by bombardment with neutrons, charged particles, or γ-rays. Then the newly formed radioactive atoms in the sample give off characteristic nuclear radiations, such as γ-rays, which tell the types and quantities of atoms present. Activation analysis is usually more sensitive than chemical analysis.

**Activity:** Quantity of a radionuclide that describes the rate at which decays occur in an amount of a radionuclide. The SI unit of radioactivity is the becquerel (Bq). One becquerel corresponds to one disintegration of a radionuclide per second.

**Acute hypoxia:** Low oxygen concentration in regions of a tumor associated with changes in blood flow through vessels, for example, by transient closing of vessels. Also called perfusion limited hypoxia.

**Acute radiation syndrome (ARS):** Biologic changes and symptoms, including death, that occur within weeks after a high-intensity total body irradiation.

**Acyclovir:** A nucleoside analogue of guanosine that blocks DNA replication when incorporated into an elongating polynucleotide. Used under the trade name Zovirax as a treatment for herpes.

**Additive:** A situation in which the effect of a combination is the sum of the effects of the separate treatments (i.e., independent cell kill).

**Adenoma:** A benign tumor of epithelial origin, such as a polyp of the colon.

**Adenovirus:** One of a group of viruses responsible for upper respiratory and other infections in humans, other mammals, and birds. Adenoviruses are useful as a vector in gene therapy because they infect both dividing and non-dividing cells, but the downside to their use is that they evoke an immune response, which makes their repeated use difficult.

**Adjuvant therapy:** A treatment method used in addition to the primary therapy. Radiation therapy is often used as an adjuvant to surgery or chemotherapy.

**Agreement state:** Any state with which the Nuclear Regulatory Commission has entered into an effective licensing agreement to enable the state to regulate source, special nuclear, and by-product materials.

**ALARA (as low as reasonably achievable):** The principle of limiting the radiation dose of exposed persons to levels as low as are reasonably achievable, economic and social factors being taken into account.

**Alleles:** Alternate forms of a gene or DNA sequence on the two homologous chromosomes of a pair.

**Alopecia:** Hair loss.

**Alpha/beta ratio (α/β ratio):** The ratio of the parameters α and β in the linear-quadratic model; used to quantify the fractionation sensitivity of tissues.

**Alpha fetoprotein (AFP):** A 70-kDa glycoprotein synthesized in embryonic development by the yolk sac. High levels of this protein in the amniotic fluid are associated with neural tube defects such as spina bifida. Lower than normal levels may be associated with Down syndrome.

**Alpha particle (α-particle):** A positively charged particle emitted by radioactive materials. It consists of two neutrons and two protons bound together; hence, it is identical with the nucleus of a helium atom. It is the least penetrative of the three common types of radiation—α, β, and γ—and it is stopped by a sheet of paper. It is not dangerous to plants, animals, or humans unless the α-emitting substance has entered the organisms. α-particles are ejected from a nucleus during the decay of some radioactive elements; for example, an α-particle is emitted if either of the radon progeny polonium-218 or polonium-214 decays.

**Alpha ray (α-ray):** A stream of α-particles. Used loosely as a synonym for α-particles.

**Alu sequence:** An interspersed DNA sequence of approximately 300 bp found in the genome of primates that is cleaved by the restriction enzyme Alu I. Alu sequences are
A mode of rapid cell death after irradiation in which the cell nucleus displays characteristic densely staining globules and at least some of the DNA is subsequently broken down into internucleosomal units. Some- times postulated to be a “programmed,” and therefore potentially controllable, process. Plays an important part in embryogenesis and in tissue regeneration following an insult and can eliminate cells whose DNA has been damaged and not repaired with a high fidelity.

ARCON therapy: The use of accelerated radiotherapy in conjunction with carbogen and nicotinamide.

Arrhenius plot: Probably the most common life–stress relationship when the stress is thermal. In hyperthermia research, a plot of 1/D0 versus 1/T, where D0 is the reciprocal of the slope of the cell survival curve and T is the absolute temperature. Named after the Swedish physical chemist.

asexual reproduction: Production of offspring in the absence of any sexual process.

atomic mass: See atomic weight.

atomic mass unit (amu): One-sixteenth the mass of a neutral atom of the most abundant isotope of oxygen, 16O.

atomic number (Z): The number of protons in the nucleus of an atom and also its positive charge.

atomic weight (at. wt.): The mass of an atom relative to other atoms. The present-day basis for the scale of atomic weights is oxygen; the most common isotope of this element arbitrarily has been assigned an atomic weight of 16. The unit of the scale is 1/16 the weight of the 16O atom, or roughly the mass of one proton or one neutron. The atomic weight of any element is approximately equal to the total number of protons and neutrons in its nucleus. Compare atomic number.

autoimmune disease: The production of antibodies that results from an immune response to one’s own molecules, cells, or tissues. Such a response results from the inability of the immune system to distinguish self from nonself. Diseases such as arthritis, scleroderma, systemic lupus erythematosus, and perhaps diabetes are considered autoimmune diseases.

autoradiography: Use of a photographic emulsion to detect the distribution of a radioactive label in a tissue specimen.

autosomes: Chromosomes other than the sex chromosomes. In humans, there are 22 pairs of autosomes.

B lymphocyte: A white blood cell responsible for production of antibodies involved in the humoral immune response.

background radiation: The radiation in the natural environment, including cosmic rays and radiation from the naturally radioactive elements, both outside and inside the bodies of humans and animals. Also called natural radiation. The term also may mean radiation that is unrelated to a specific experiment.

bacteriophage: A virus that infects bacteria. Altered forms are used as vectors for cloning DNA.

bacterium: A single-celled prokaryotic organism.

basal cells: Cells at the base of the wall of the lung airways. These cells divide to replenish the other cells in the lung wall and often are considered the key cells that, if damaged, can lead to lung cancer.

base pair: A pair of complementary nitrogenous bases in a DNA molecule—adenine–thymine and guanine–cytosine. Also, the unit of measurement for DNA sequences.

basic fibroblast growth factor (FGF): A mitogen for many types of cells of mesodermal or neuroectodermal origin. Has various angiogenic properties.

B-DNA: See double helix.

beam: A stream of particles or electromagnetic radiation moving in a single direction.

becquerel (Bq): Unit of radioactivity, corresponding to one radioactive disintegration per second. See activity.
benign: Describing a slow-growing, not malignant, tumor that does not spread to other parts of the body. If completely removed, benign lesions do not tend to recur. Incompletely removed tumors may recur but do not spread. Although benign, these tumors may cause permanent damage to some structures in the brain.

BER: Base excision repair, a pathway for repairing damage to DNA bases.

beta particle (β-particle): An elementary particle emitted from a nucleus during radioactive decay, with a single electrical charge and a mass equal to 1/1,837 that of a proton. A negatively charged β-particle is identical to an electron. A positively charged β-particle is called a positron. β-radiation may cause skin burns, and β-emitters are harmful if they enter the body. β-particles, however, are stopped easily by a thin sheet of metal.

BeV: One billion electron volts. Also written as GeV.

bilateral: On both sides of the body.

bioassay: A technique used to identify, quantify, and/or specify the location of radionuclides in the body by direct or indirect analysis of tissues or excretions from the body.

biochemistry: Chemical reactions that sustain life.

biologic half-life: See half-life, biologic.

biologic therapy: Treatment to stimulate or restore the ability of the immune system to fight infection and disease. Also called immunotherapy.

biologically effective dose (BED): In fractionated radiotherapy, the quantity by which different fractionation regimens are compared.

biopsy: The removal of a small portion of a tumor to allow a pathologist to examine it under a microscope and provide a diagnosis of tumor type.

biotechnology: Commercial or industrial processes that use biologic organisms or products.

blob: A concentration of about 12 ion pairs in a region about 7 nm in diameter.

blood count: The number of red blood cells, white blood cells, and platelets in a sample of blood.

BNCT: Boron neutron capture therapy.

body burden: The amount of radioactive material present in a human or an animal. See background radiation, whole-body counter.

bone marrow: Spongy tissue in the cavities of large bones where the body’s blood cells are produced.

bone seeker: A radioisotope that tends to accumulate in the bones; for example, the strontium-90 isotope, which behaves chemically like calcium.

brachytherapy: Internal radiation treatment achieved by implanting radioactive material directly into the tumor or very close to it. Sometimes called internal radiation therapy.

Bragg peak: Region of maximum dose deposition near to the end of the track of a charged particle, such as a proton or carbon ion. This allows precise spatial definitions of dose distributions for radiotherapy with charged particles.

breeder reactor: A nuclear reactor that produces fissionable material as well as consuming it, especially one that creates more than it consumes. The new fissionable material is obtained during the production or the use of source material or fissionable material. By-product material includes fission products and many other radioisotopes produced in nuclear reactors.

bystander effect: Induction of biologic effects in cells that are not directly traversed by a charged particle but are in close proximity to cells that are.

cancer: A general name for more than 100 diseases in which abnormal cells grow out of control; a malignant tumor.

cancer stem cell: Cells within a tumor that have the capacity to self-renew.

carcinogen: A physical or chemical agent capable of causing cancer, such as radon progeny, cigarette smoke, or asbestos.

carcinogenesis: Process that leads to the formation of a cancer. It involves several stages (resulting from successive alterations of the genome). The first stage is initiation (which may, for instance, be caused by the mutation of a proto-oncogene into an oncogene). For a normal cell to be “transformed” (i.e., for it to become preneoplastic), its genome has to undergo several modifications: appearance of an oncogene, inactivation of both copies of a suppressor gene, immortalization (i.e., acquisition of an unlimited capacity to proliferate), changes affecting the apoptosis system, and so on. A transformed cell can give rise to an invasive cancer at the end of the second stage, known as promotion, which is associated with the proliferation of the descendants of the initiated cell and the escape of one of them from the control of the normal surrounding cells and of the body.

carcinogenic: Having the potential (as, for example, tobacco smoke or alcohol) to contribute to the development of cancer (same as oncogenic).

carcinoma: A malignant tumor derived from epithelial tissue, which forms the skin and outer cell layers of internal organs.

cardiac catheterization: Passage of a small catheter through a vessel in an arm, leg, or neck and into the heart, permitting the securing of blood samples, determination of intracardiac pressure, detection of cardiac anomalies, and the injection of contrast media for imaging.

CAT scan: Computerized axial tomography, often called a CT scan, which provides three-dimensional x-ray images of some part of the body. It is useful for diagnosing cancer and for planning radiation therapy treatments.

catalyst: A substance that promotes a chemical reaction by lowering the activation energy of a chemical reaction, but itself remains unaltered at the end of the reaction.

cathode: An opacification in the normally transparent lens of the eye.

cathode: Negative side of the x-ray tube; contains the filament and focusing cup.

CD11b+: A cell surface glycoprotein found on monocytes, granulocytes, NK cells and some peripheral blood lymphocytes. Usually CD11b is found as a heterodimer with beta 2 integrin.
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**chain reaction:** A reaction that stimulates its own repetition.

**cervix:** The lower part of the uterus, which projects out into the vagina.

**c-erbB2 (HER2/neu):** A gene closely related to the epidermal growth factor receptor, which is amplified in various cancers, including that of the breast.

**Cdk1:** Kinase that associates with cyclin B to regulate entry into mitosis. Complex is activated by Cdc25-mediated dephosphorylation. Also associates with cyclin A during M phase.

**Cdk2:** Kinase that associates with cyclin E during the G1/S transition and cyclin A during S phase. Inhibited by p21 and p27.

**Cdk4:** Kinase that associates with cyclin D1. Complex can phosphorylate pRb to allow cells to progress through G1. Inhibited by p16, p21, and p27.

**Cdk5s (cyclin-dependent kinases):** Proteins that complex with their cyclin regulatory subunits to phosphorylate proteins necessary for progression through the cell cycle.

**cDNA (copy DNA):** DNA synthesized from an RNA template using reverse transcriptase.

**cDNA library:** A library composed of complementary copies of cellular mRNAs (i.e., the exons without the introns).

**cell cycle:** Sum of the phases of growth of an individual cell type; divided into G1 (gap 1), S (DNA synthesis), G2 (gap 2), and M (mitosis); the cycle of cellular events from one mitosis to the next.

**cell cycle checkpoint:** Control mechanism to verify that each phase of the cell cycle has been accurately completed before progression to the next phase.

**cell cycle time:** The time between one mitosis and the next.

**cell loss factor (Φ):** The rate of cell loss from a tumor as a proportion of the rate at which cells are being added to the tumor by mitosis.

**cells:** The body’s tiny functioning units, which can be observed under a microscope. Each cell plays a specialized role in the body. Groups of cells are organized together to form tissues. Tissues are organized to form organs in the body.

**cellular oncogene (proto-oncogene):** A normal gene that if mutated or improperly expressed, contributes to the development of cancer.

**centimorgan:** A unit of distance between genes on chromosomes. One centimorgan represents a value of 1% crossing over between two genes.

**central nervous system (CNS):** The brain and spinal cord.

**centriole:** A cytoplasmic organelle composed of nine groups of microtubules, generally arranged in triplets. Centrioles function in the generation of cilia and flagella and serve as foci for the spindles in cell division.

**centromere:** The chromosome constriction to which the spindle fiber attaches. The position of the centromere determines whether chromosomes are metacentric (X-shaped; e.g., chromosomes 1, 3, 16, 19, 20) or acrocentric (inverted V-shaped; e.g., chromosomes 13–15, 21, 22, Y). During mitosis, the identical chromatids of each chromosome are separated by shortening of the spindle fibers attached to opposite poles of the dividing cell.

**c-Erb-B2 (HER2/neu):** A gene closely related to the epidermal growth factor receptor, which is amplified in various cancers, including that of the breast.

**chronic:** Persisting for a long time.

**chronic hypoxia:** Persistent low oxygen concentrations, such as exists in viable tumor cells close to regions of necrosis.

**chronic hypoxia:** Persistent low oxygen concentrations, such as exists in viable tumor cells close to regions of necrosis.

**c-fos:** An early-response gene induced by mitogenic stimuli and stress responses. Forms complexes in the nucleus that act as transcription factors. Recognizes AP-1 sites if complexed with c-jun.

**chain reaction:** A reaction that stimulates its own repetition. In a fission chain reaction, a fissionable nucleus absorbs a neutron and fissions, releasing additional neutrons. These, in turn, can be absorbed by other fissionable nuclei, releasing still more neutrons. A fission chain reaction is self-sustaining if the number of neutrons released in a given time equals or exceeds the number of neutrons lost by absorption in nonfissioning material or by escape from the system.

**characteristic x-rays:** X-rays produced following ionization of inner-shell electrons; characteristic of the target element.

**charged particle:** An ion; an elementary particle that carries a positive or negative electrical charge.

**CHART:** Continuous hyperfractionated accelerated radiotherapy, delivering 54 Gy in 36 fractions with 3 fractions on 12 consecutive days.

**chemotherapist:** A physician who specializes in the use of drugs to treat cancer, now called a medical oncologist.

**chemotherapy:** A treatment for cancers that involves administering chemicals toxic to malignant cells.

**chromatid:** Each of the two progeny strands of a duplicated chromosome joined at the centromere during mitosis and meiosis.

**chromatin:** The complex of DNA, RNA, histones, and non-histone proteins that make up chromosomes.

**chromatography:** Technique for the separation of a mixture of solubilized molecules by their differential migration over a substrate.

**chromosomal aberration:** Any change resulting in the duplication, deletion, or rearrangement of chromosomal material.

**chromosomal instability:** An effect of irradiation in which chromosomal aberrations continue to appear through many cell generations.

**chromosomal mapping:** See chromosomal aberration.

**chromosomal polymorphism:** Alternate structures or arrangements of a chromosome that are carried by members of a population.

**chromosome:** In prokaryotes, an intact DNA molecule containing the genome; in eukaryotes, a DNA molecule complexed with RNA and proteins to form a threadlike structure containing genetic information arranged in a linear sequence.

**chromosome banding:** Technique for the differential staining of mitotic or meiotic chromosomes to produce a characteristic banding pattern or selective staining of certain chromosomal regions such as centromeres, the nucleolus organizer regions, and GC- or AT-rich regions.

**chromosome map:** A diagram showing the location of genes on chromosomes.

**clinical trials:** Medical research studies conducted with volunteers. Each study is designed to answer scientific questions and to find better ways to prevent or treat disease.

**clone:** Genetically identical cells or organisms all derived from a single ancestor by asexual or parasexual methods; for example, a DNA segment that has been enzymatically inserted into a plasmid or chromosome of a phage or bacterium and replicated to form many copies.
cloned library: A collection of cloned DNA molecules representing all or part of an individual's genome.

clonogenic cells: Cells that have the capacity to produce an expanding family of descendants (usually at least 50). Also called colony-forming cells or clonogens.

c-e-myc: An early-response gene induced by mitogenic stimuli, as well as TGF-β. Is highly overexpressed as a result of translocations in Burkitt lymphomas. Gene is amplified in certain cancers, as is its relative, N-myc.

cobalt-60: A radioactive substance used as a radiation source to treat cancer.

code: See genetic code.
codon: A group of three nucleotides that specifies the addition of one of the 20 amino acids during translation of mRNA into a polypeptide.
colchicine: An alkaloid compound that inhibits spindle formation during cell division. Used in the preparation of karyotypes to collect a large population of cells inhibited at the metaphase stage of mitosis.
collective dose: Usually refers to the collective effective dose obtained by multiplying the average effective dose by the number of persons exposed to that given dose. To be expressed in person-Sv. The old unit was the man-rem.
collective effective dose: (person-Sv) The sum of all the individual effective doses in the population of concern.

colon: Large intestine.

colony: A group of identical cells derived from a single ancestor cell.

combination chemotherapy: The use of more than one drug to treat cancer.

committed effective dose: Effective dose due to absorbed doses in the specified organs or tissues integrated over 50 years following an intake of a radionuclide by ingestion, inhalation, or dermal absorption.

complementarity: Chemical affinity between nitrogenous bases as a result of hydrogen bonding. Responsible for the base pairing between the strands of the DNA double helix.

complementary nucleotides: Members of the pairs adenine–thymine, adenine–uracil, and guanine–cytosine, which have the ability to hydrogen-bond to each other.

complementation: Identification of whether a (radiosensitive) phenotype in different mutants is caused by the same gene. Studied by means of cell fusion.

complementation test: A genetic test to determine whether two mutations occur within the same gene. If both mutations are introduced into a cell simultaneously and produce a wild-type phenotype (i.e., they complement each other), they are often nonallelic. If a mutant phenotype is produced, the mutations are noncomplementing and are often allelic.

Compton effect: Scattering of x-rays resulting in ionization and loss of energy. The energy lost by the photon is given to the ejected electron as kinetic energy.

concomitant boost: The practice of adding an extra dose to the specified organs or tissues integrated over a very low-power level, permitting study of the assembly's components for various fissionable materials in different geometric arrangements.

cosmic rays: Radiation of many sorts, but mostly protons and heavier atomic nuclei with very high energies originating outside the earth's atmosphere. Cosmic radiation is part of the natural background radiation. Some cosmic rays are more energetic than any human-made forms of radiation.

cosmid: A vector designed to allow cloning of large segments of foreign DNA (25,000–45,000 bp). Cosmids are hybrids composed of the cos sites of lambda inserted into a plasmid. In cloning, the recombinant DNA molecules are packaged into phage protein coats, and after infection of bacterial cells, the recombinant molecule replicates and can be maintained as a plasmid.

covalent bond: A nonionic chemical bond formed by the sharing of electrons.

critical: Capable of sustaining a chain reaction. See criticality.

critical assembly: An assembly of sufficient fissionable material and moderator to sustain a fission chain reaction at a very low-power level, permitting study of the assembly's components for various fissionable materials in different geometric arrangements.

critical mass: The smallest mass of fissionable material that supports a self-sustaining chain reaction under stated conditions.

criticality: The state of a nuclear reactor if it is sustaining a chain reaction.

cross section (σ): A measure of the probability that a nuclear reaction will occur. Usually measured in barns, it is the apparent or effective area presented by a target nucleus or particle to an oncoming particle or other nuclear radiation, such as a photon of γ-radiation.

crossing over: The exchange of chromosomal material (parts of chromosomes) between homologous chromosomes by breakage and reunion. The exchange of material between nonsister chromatids during meiosis is the basis of genetic recombination.

CT scan (CAT scan, CT x-ray): A three-dimensional x-ray. CT stands for computerized tomography.
cure: An outcome of treatment that leaves the patient disease free, with no likelihood of recurrence.
curie (Ci): Old unit of radioactivity, corresponding to 3.7 × 1010 radioactive disintegrations per second. Now replaced by the becquerel.

cXCR4: Receptor for the chemokine stromal derived factor-1 (SDF-1) that is involved in regulating chemotaxis. Plays an important role in vascularization.
cyclic adenosine monophosphate (cAMP): An important regulatory molecule in both prokaryotic and eukaryotic organisms.
cyclin A: Regulatory protein that associates with Cdk2 during the S phase of the cell cycle.
cyclin B: Regulatory protein that associates with Cdk2 to regulate entry into mitosis.
cyclin D: One of three cyclin D family members (also D2, D3). Associates with Cdk4 or Cdk6 to regulate through the G1 phase of the cell cycle. Is induced by various mitogens.
cyclin E: Protein that associates with Cdk2 during late G1 phase and is thought to regulate entry into S phase.
cyclins: Proteins that complex with CdkS to regulate progression through the cell cycle.
cyst: A cavity, usually filled with a liquid, sometimes associated with benign or malignant tumors.
cytokines: Study that relates the appearance and behavior of chromosomes to genetic phenomena.
cytogenetics: Procedures for determining the nucleotide sequence of a DNA fragment. Identified by Southern hybridization or by polymerase chain reaction.
dna: DNA is the genetic material of most organisms and consists of two polynucleotide strands of DNA molecules. Are held together by hydrogen bonds between adenine-thymine and cytosine-guanine base pairs.
dna fingerprint: A unique pattern of DNA fragments identified by Southern hybridization or by polymerase chain reaction.
dna polymerase: An enzyme that catalyzes the synthesis of DNA from deoxyribonucleotides and a template DNA molecule.
dna polymorphism: One or two or more alternate forms of a chromosomal locus that differ in nucleotide sequence or have variable numbers of repeated nucleotide units.
dna sequencing: Procedures for determining the nucleotide sequence of a DNA fragment.
dNase (deoxyribonuclease): An enzyme that degrades or breaks down DNA into fragments or constitutive nucleotides.
dominant: Expression of a trait in the heterozygous condition.
dominant gene: A gene whose phenotype is expressed if it is present in a single copy.
dominant-oncogene: A gene that stimulates cell proliferation and contributes to oncogenesis when present in a single copy.
dose: General term for the quantity of radiation. See absorbed dose, equivalent dose, effective dose, collective dose.
dose limit: A limit on dose that is applied for exposure of individuals to prevent the occurrence of deterministic effects and to limit the probability of stochastic effects.
dose rate: The radiation dose delivered per unit time and measured, for instance, in grays per hour.

dose-modifying factor (DMF): A ratio indicating the dose without to the dose with the agent for the same level of effect. (Similarly, dose-reduction factor or sensitizer enhancement ratio.)

dose-rate effect: Decreasing radiation response with decreasing radiation dose rate.

dosimetrist: A person who plans and calculates the proper radiation dose for treatment.

double helix: The model for DNA structure proposed by James Watson and Francis Crick, involving two antiparallel, hydrogen-bonded polynucleotide chains wound into a right-handed helical configuration, with 10 base pairs per full turn of the double helix. Often called B-DNA.

double minutes: Chromosome fragments that are the result of chromosome duplication and in which oncogenes may be amplified. A common cytologic feature of tumor cells.

double trouble: The situation in which a hot spot within a treatment field receives not only a higher dose but also a higher dose per fraction, which means that the biologic effectiveness of the dose is also greater.

doubling dose: Applied to heritable effects, the dose required to double the spontaneous mutation incidence; put another way, the dose required to produce an incidence of mutations equal to the spontaneous rate.

doubling time: The time it takes for a cell population or tumor volume to double its size.

Drosophila melanogaster: The fruit fly, whose common use in genetic studies was introduced by Thomas Hunt Morgan in the early 1900s.

early-response gene: A gene such as c-fos, c-jun, or c-myc, whose mRNA levels are induced dramatically following mitogenic stimuli or stress.

early responses: Radiation-induced normal tissue damage that is expressed in weeks to a few months after exposure. Generally caused by damage to parenchymal cells. The $\alpha/\beta$ ratio tends to be large.

ED$_{50}$ (effect dose 50%): Radiation dose that produces a specified effect in the normal tissues of 50% of animals.

edema: Abnormal accumulation of fluid (e.g., pulmonary edema, a buildup of fluid in the lungs).

EF5: This molecule is a fluorinated derivative of the 2-nitroimidazole etamidazole and is used to determine oxygen levels in normal and tumor tissue. It binds rather selectively and irreversibly to cellular macromolecules in hypoxic cells. A radioactive version has been generated for PET imaging is currently undergoing testing.

effective dose: The radiation dose allowing for the fact that some types of radiation are more damaging than others, and some parts of the body are more sensitive to radiation than others. It is defined as the sum over specified tissues of the products of the equivalent dose in a tissue and the weighting factor for that tissue.

eIF2a: Eukaryotic translation initiation factor 2 alpha is a protein that catalyzes the formation of the preinitiation complex involved in protein synthesis.

electromagnetic radiation: Radiation consisting of associated and interacting electrical and magnetic waves that travel at the speed of light, such as light, radio waves, $\gamma$-rays, and x-rays. All electromagnetic radiation can be transmitted through a vacuum.

electron (e): An elementary particle with a unit negative electrical charge and a mass 1/1,837 that of the proton. Electrons surround the positively charged nucleus and determine the chemical properties of the atom. Positive electrons, or positrons, also exist.

electronvolt (eV): The amount of energy gained by a particle of charge e $(-1.6 \times 10^{-19}$ C) if it is accelerated by a potential difference of 1 V. 1 eV $= 1.6 \times 10^{-19}$ J.

electrophoresis: The technique of separating charged molecules in a matrix to which an electrical field is applied.

electroporation: A process whereby high-voltage pulses of electricity are used to open pores in the cell membrane, through which foreign DNA can pass.

Ellis formula: The relation between dose, overall time, and number of fractions in radiotherapy (NSD).

endonuclease: An enzyme that cleaves at internal sites in the substrate molecule.

epidermal growth factor (EGF): Protein that promotes growth of epidermal cells and can stimulate or inhibit the proliferation and differentiation of various cells.

epithelium: A thin layer of cells in the skin, mucous membrane, or any duct that replaces worn-out cells by cell division...

equivalent dose: A quantity used for radiation protection purposes that takes into account the different probability of effects that occur with the same absorbed dose delivered by radiations with different radiation weighting factor values. It is defined as the product of the average absorbed dose in a specified organ or tissue and the radiation weighting factor values. If dose is in grays, equivalent dose is in sieverts.

erb-B (EGFR/HER1): Membrane receptor that binds epidermal growth hormone.

error-free repair: DNA repair whereby the molecule is reconstituted with a high fidelity (i.e., without loss of information).

erythropoietin: Cytokine that stimulates late erythroid progenitors to form small colonies of erythrocytes.

Escherichia coli (E. coli): A bacterium found in the human colon that is used widely as a host for molecular cloning experiments.

ethidium bromide: A fluorescent dye that is used to stain DNA and RNA and that intercalates between nucleotides and fluoresces if exposed to ultraviolet light.

eukaryote: An organism whose cells possess a nucleus and other membrane-bounded vesicles, including all members of the protist, fungi, plant, and animal kingdoms.

evolution: The origin of plants and animals from preexisting types. Descent with modifications.

excision repair: Repair of DNA lesions by removal of a polymucleotide segment and its replacement with a newly synthesized, corrected segment.

exon: DNA segment of a gene that is transcribed and translated into protein.

exonuclease: An enzyme that breaks down nucleic acid molecules by breaking phosphodiester bonds at the 3’ or 5’ terminal nucleotides.

exponential growth: Growth according to an exponential equation.

exponential survival curve: A survival curve without a threshold or shoulder region that is a straight line on a semilogarithmic plot.
exposure (X): (Often used in its more general sense and not as the specially defined radiation quantity). A measure of the quantity of x- or γ-radiation based on its ability to ionize the air through which it passes. The SI unit of exposure is coulomb per kilogram (C/kg). The previously used special unit of exposure was the röntgen (R).

expression library: A library of cDNAs whose encoded proteins are expressed by specialized vectors.

expression vector: A plasmid or phage-carrying promoter region designed to cause expression of cloned DNA sequences.

external radiotherapy: Radiation therapy that uses a machine located outside the body to aim high-energy rays at cancer cells. Sometimes called external beam radiotherapy.

extrapolated total dose (ETD): Calculated isoeffect dose when the dose rate is very low or when fraction size is very small.

extrapolation number: A parameter in the multitarget equation: the point on the survival scale to which the straight part of the curve back-extrapolates.

fallout: The radioactive material falling from the atmosphere to Earth’s surface after a nuclear event, such as a weapon’s test or accident.

familial trait: A trait transmitted through and expressed by members of a family.

fast neutrons: Neutrons with energy greater than approximately 100,000 eV. Compare thermal neutrons.

fibroblast: A precursor cell of connective tissue that is relatively easy to maintain in cell culture.

field: A term used in radiation oncology to describe or define an area through which x-rays are directed toward the tumor.

field-size effect: The dependence of normal tissue damage on the size of the irradiated area; also known as volume effect.

film badge: An assembly containing a packet of unexposed photographic film and various filters (absorbers); when the film is developed, the dose and type of radiation to which the wearer was exposed can be estimated.

fingerprint: The pattern of ridges and whorls on the tip of a finger. The pattern obtained by enzymatically cleaving a protein or nucleic acid and subjecting the digest to two-dimensional chromatography or electrophoresis.

FISH (fluorescence in situ hybridization): The process whereby fluorescent dyes are attached to specific regions of the genome, thus aiding the identification of chromosomal damage.

fission: The splitting of a heavy nucleus into two approximately equal parts (which are nuclei of lighter elements), accompanied by the release of a relatively large amount of energy and generally one or more neutrons. Fission can occur spontaneously, but usually it is caused by nuclear absorption of γ-rays, neutrons, or other particles.

fission products: The nuclei (fission fragments) formed by the fission of heavy elements plus the nuclei formed by the fission fragments’ radioactive decay.

flanking region: The DNA sequences extending on either side of a specific locus or gene.

gene: The fundamental physical unit of heredity whose existence can be confirmed by allelic variants and that occupies a specific chromosomal locus; a DNA sequence coding for a single polypeptide.

gene amplification: The presence of multiple copies of a gene and one of the mechanisms by which proto-oncogenes are activated to result in neoplasia.

gene expression: The process of producing a protein from its DNA- and mRNA-coding sequences.

gene mutation: See point mutation.

gene pool: The total of all genes possessed by reproductive members of a population.

generic burden: Average number of recessive lethal genes carried in the heterozygous condition by an individual in a population. Also called genetic load.
granulocyte colony-stimulating factor (G-CSF): Cytokine that stimulates differentiation of progenitors into granulocytes.

growth delay: Extra time required for an irradiated tumor to reach a given size, compared with an unirradiated control.

growth factor: A serum protein that stimulates cell division when it binds to its cell surface receptor.

growth factor receptor: A membrane-spanning protein that selectively binds its growth factor and then transduces a signal for cell division to other molecules in the cytoplasm and nucleus.

growth fraction: Proportion of viable cells in active cell division.

growth hormone (GH): Secreted by the anterior pituitary gland, a hormone that acts mainly on the growth of bone and muscles. Can be secrected by lymphocytes in response to phorbol ester treatment and may be involved in lymphocyte growth. Also known as somatotropin.

genetic code: The three-letter code that translates nucleic acid sequence into protein sequence.

genetic counseling: Analysis of risk for genetic defects in a family and the presentation of options available to avoid or ameliorate possible risks.

genetic disease: A disease that has its origin in changes to the genetic material, DNA. Usually refers to diseases that are inherited in a mendelian fashion, although noninherited forms of cancer also result from DNA mutation.

genetic drift: Random variation in gene frequency from generation to generation, most often observed in small populations.

genetic effects of radiation: Radiation effects that can be transferred from parent to offspring; any radiation-caused changes in the genetic material of sex cells. Now called heritable effects.

genetic engineering: The manipulation of an organism’s genetic endowment by introducing or eliminating specific genes. A broad definition of genetic engineering also includes selective breeding and other means of artificial selection.

genetic polymorphism: The stable coexistence of two or more discontinuous genotypes in a population. If the frequencies of two alleles are carried to equilibrium, the condition is called balanced polymorphism.

genetically significant dose: The dose that, if given to every member of a population, should produce the same heritable harm as the actual doses received by the individuals. Expressed in sievers, this dose takes into account the childbearing potential of those receiving the dose.

genetics: The branch of biology that deals with heredity and the expression of inherited traits.

gene: The genetic complement contained in the chromosomes of a given organism, usually the haploid chromosome state.

gray (Gy): The special name for the SI unit of absorbed dose, kerma, and specific energy imparted equal to 1 J/kg. The previous unit of absorbed dose, rad, has been replaced by the gray. One gray equals 100 rad.

ground state: The state of a nucleus, an atom, or a molecule at its lowest (normal) energy level.

geneva (G): The time required for a biologic system, such as a human or an animal, to eliminate by natural processes half the amount of a substance, such as a radioactive material, that has entered it.

haploid: Having only one of each type of chromosome, as is usually the case in gametes (oocytes and spermatozoa).

heat shock: A transient response following exposure of cells or organisms to elevated temperatures. The response involves activation of a small number of loci, inactivation of previously active loci, and selective translation of heat-shock mRNA. Appears to be a nearly universal phenomenon, observed in organisms ranging from bacteria to humans.

heavy water (D2O): Water containing significantly more than the natural proportion (1 in 6,500) of heavy hydrogen (deuterium) atoms to ordinary hydrogen atoms. Heavy water is a moderator in some nuclear reactors because it slows down neutrons effectively and also has a low cross section for absorption of neutrons.

gene: An enzyme that participates in DNA replication by unwinding the double helix near the replication fork.

genes: The branch of biology that deals with heredity and the expression of inherited traits.

genetherapy: The study of blood and its disorders.

hemizygous: Describing a condition in which a gene is present in a single dose. Usually applied to genes on the X chromosome in heterosexual males.

hemoglobin (HB): An iron-containing, conjugated respiratory protein occurring chiefly in the red blood cells of vertebrates. Carries oxygen to the tissues.

hemophilia: A sex-linked trait in humans associated with defective blood-clotting mechanisms.

herpesvirus: One of a group of viruses causing herpes in man and other primates. These viruses are potentially useful as vectors in gene therapy because they are large and can accommodate a large insert, but they are seldom used in practice because they are potentially more pathogenic and difficult to control.

heterochromatin: The heavily staining, late-replicating regions of chromosomes that are condensed in interphase. Thought to be devoid of structural genes.
**Glossary**

**heterozygote:** An individual with different alleles at one or more loci. Such individuals produce unlike gametes and therefore do not breed true.

**hierarchical tissues:** Cell populations comprising a lineage of stem cells, proliferating cells, and mature cells. The mature cells do not divide.

**high-dose-rate (HDR) remote brachytherapy:** A type of internal radiation in which each treatment is given in a few minutes with the radioactive source in place. The source of radioactivity is removed between treatments.

**histones:** Proteins complexed with DNA in the nucleus. They are rich in the basic amino acids arginine and lysine and function in the coiling of DNA to form nucleosomes.

**HLA:** Cell surface proteins, produced by histocompatibility loci, that are involved in the acceptance or rejection of tissue and organ grafts and transplants.

**homogeneously staining region:** Segment of mammalian chromosomes that stains lightly with Giemsa following exposure of cells to a selective agent. These regions arise in conjunction with gene amplification and are regarded as the structural locus for the amplified gene.

**homologous chromosomes:** Chromosomes that have the same linear arrangement of genes; a pair of matching chromosomes in a diploid organism.

**homologous recombination (HR):** DNA repair pathway for double-strand breaks by using as a template an undamaged homologous DNA sequence, usually from the sister chromatid.

**homologue:** Any member of a set of genes or DNA sequences from different organisms whose nucleotide sequences show a high degree of one-to-one correspondence.

**homozygote:** An individual with identical alleles at one or more loci. Such individuals produce identical gametes and therefore breed true.

**hormesis:** The phenomenon whereby a physical or chemical agent has one effect at high doses and the reverse effect at low doses. It probably results from the activation of defense mechanisms. Hormesis is observed with several drug molecules that are toxic at high doses, but which can have a beneficial protective effect at low doses.

**hormones:** Factors synthesized in endocrine glands that if released, act to regulate and modulate the functions of multicellular organisms.

**Human Genome Project:** A project coordinated by the National Institutes of Health and the Department of Energy to determine the entire nucleotide sequence of the human chromosomes.

**hybrid:** The offspring of two parents differing in at least one genetic characteristic.

**hybridization:** The hydrogen bonding of complementary DNA or RNA sequences to form a duplex molecule.

**hybridoma:** A somatic cell hybrid produced by the fusion of an antibody-producing cell and a cancer cell, specifically a myeloma. The cancer cell contributes the ability to divide indefinitely, and the antibody cell confers the ability to synthesize large amounts of a single antibody.

**hydrogen bond:** An electrostatic attraction between a hydrogen atom bonded to a strongly electronegative atom such as oxygen or nitrogen and another atom that is electronegative or contains an unshared electron pair.

**hyperbaric oxygen (HBO):** The use of high oxygen pressures (two or three atmospheres) to enhance oxygen availability in radiotherapy.

**hyperdiploid:** Additional chromosomes; the modal number is 47 or more.

**hyperfractionated radiation:** Division of the total dose of radiation into smaller doses usually given more than once a day.

**hyperthermia:** The use of heat to treat cancer.

**hypodiploid:** Loss of chromosomes with modal number 45 or less.

**hypo fractionation:** The use of dose fractions substantially larger than the conventional level of ~2 Gy.

**hypopharynx:** Part of the lower throat beside and behind the larynx (voice box).

**hypoxia:** Low oxygen tension; usually the very low levels required to make cells maximally radiosensitive. Sometimes used to mean anoxia (literally, the complete absence of oxygen).

**hypoxic cell cytotoxic:** Any agent, typically a bioreductive drug, that preferentially kills hypoxic cells.

**hypoxic fraction:** The fraction of hypoxic cells in a tumor that are resistant to radiation, but still viable.

**ICRP:** International Commission on Radiological Protection.

**identical twins:** See monozygotic twins.

**IFNs:** Cytokine that is produced in response to viral infection and that protects cells from viruses and causes growth arrest of normal and tumor cells.

**IFNβ:** Cytokine that is produced in response to viral infection and that protects cells from viruses and causes growth arrest of normal and tumor cells.

**IFNγ:** Cytokine that is produced in response to viral infection and that activates macrophages, protects cells from viruses, and causes growth arrest of normal and tumor cells.

**IL-1:** Cytokine involved in the regulation of immune and inflammatory responses.

**IL-2:** Cytokine that stimulates growth of T cells. Also stimulates B cell growth and differentiation, generation of lymphokine-activated killer cells, activation of macrophages, and production of other cytokines.

**IL-3:** Cytokine that induces the proliferation of hematopoietic cells, particularly erythroid and myeloid cells. Produced by activated T cells and mast cells.

**IL-4:** Cytokine that has general effects on hematopoietic cells, including the activation, growth, and differentiation of B cells. Also induces growth of mast cells and T cells.

**IL-5:** Cytokine that induces eosinophil differentiation, as well as B cell activation, growth, and differentiation.

**IL-6:** Cytokine that regulates B cell differentiation, T cell activation, killer cell induction, and other physiologic responses. Induced by cytokines, ultraviolet irradiation, and other stimuli. Many effects are similar to IL-1 and TNF.

**IL-7:** Cytokine that promotes the growth of B and T cell progenitors.

**IL-8:** One of an extended family of cytokines that act as chemotactants for neutrophils, T cells, and basophils.

**IL-9:** Cytokine that stimulates proliferation of T cells and enhances mast cell activity and growth.

**IL-10:** Cytokine that enhances IL-2 and IL-4 proliferative response of T cells. Also acts to inhibit cytokine production by various different cells.

**IL-11:** Cytokine that together with other cytokines stimulates the growth of various hematopoietic progenitors.
IL-12: Cytokine that promotes growth of T and natural killer cells and enhances cytotoxic T cell responses.

immortalization: Term used to describe the change as cells from somatic tissues that are only able to perform a limited number of cell divisions undergo a “crisis” and become capable of unlimited cell divisions.

immortalizing oncogene: A gene that upon transfection enables a primary cell to grow indefinitely in culture.

immune system: The body’s defense system, which protects it from foreign substances, such as bacteria and viruses that are harmful to it.

immunoglobulin: The class of serum proteins having the properties of antibodies.

implant: A quantity of radioactive material placed in or near a cancer.

IMRT: Intensity-modulated radiation therapy. A technique using nonuniform radiation beams to achieve high conformity treatment plans.

in situ hybridization: A technique for the cytologic localization of RNA sequences complementary to a given nucleic acid or polyribonucleotide.

in vitro: Literally, in glass; outside the living organism; occurring in an artificial environment.

in vivo: Literally, in the living; occurring within the living body of an organism.

inbreeding: Mating between closely related organisms.

incomplete repair: Increased damage from fractionated radiotherapy if the time interval between doses is too short to allow complete recovery of sublethal damage.

indirect action: Damage to DNA by free radicals formed through the ionization of nearby water molecules.

inducible response: A response to irradiation that is modified by a small dose of radiation given shortly before.

infusion: Slow or prolonged intravenous delivery of a drug or fluid.

initial slope: The steepness of the initial part of the oxic cell survival curve.

initiating agent: Something that causes initial “latent” damage to the DNA. The cell requires more damage from a second “promoting agent” before the damage is expressed as cancer. Radiation usually is considered an initiating agent.

initiation codon: The mRNA sequence AUG, which codes for methionine and initiates translation.

intercalating agent: A compound that inserts between bases in a DNA molecule, disrupting the alignment and pairing of bases in the complementary strands (e.g., acridine dyes).

interferon: One of a family of proteins that act to inhibit viral replication in higher organisms. Some interferons may have anticancer properties.

interphase: That portion of the cell cycle between divisions.

interphase death: The death of irradiated cells before they reach mitosis.

interstitial implant: The placement of fine tubes in a gridlike pattern through tissues containing a cancer; these tubes are filled later with radioactive sources for brachytherapy.

intracavitary implant: The placement of a small tube within a body cavity, such as the bronchus or vagina; this tube later is filled with radioactive sources for brachytherapy.

intraoperative radiation: A type of radiation used to deliver a large dose of radiation therapy to the tumor bed and surrounding tissue at the time of surgery.

intravenous (IV): Into a vein.

intron: A portion of DNA that is located between coding regions in a gene and which is transcribed, but which does not appear in the mRNA product.

inverse square law: A physical law stating that the intensity of x- or y-radiation from a point source emitting uniformly in all directions is inversely proportional to the square of the distance from the source.

inversion: Two breaks occurring in the same chromosome with rotation of the intervening segment. If both breaks are on the same side of the centromere, it is called a pericentric inversion. If they are on opposite sides, it is called a paracentric inversion.

ion: An electrically charged atom or group of atoms.

ion pair: A closely associated positive ion and negative ion (usually an electron) having charges of the same magnitude and from a neutral atom or molecule by radiation.

ionization: The process of adding one or more electrons to, or removing one or more electrons from, atoms or molecules, thereby creating ions. High temperatures, electrical discharges, or nuclear reactions can cause ionization.

ionization chamber: A device for detection of ionizing radiation or for measurement of radiation dose and dose rate.

ionizing radiation: Any radiation displacing electrons from atoms or molecules, thereby producing ions. Examples of ionizing radiation are α-, β-, and γ-radiation, x-rays, and shortwave ultraviolet light. Ionizing radiation may produce severe skin or tissue damage. See radiation burn, radiation illness.

ipsilateral: On the same side of the body (opposite of contralateral).

irradiation: Exposure to radiation, as in a nuclear reactor.

isobars: Atoms having the same number of nucleons but different numbers of protons and neutrons.

isochromosome: A chromosome that consists of identical copies of one chromosome arm with loss of the other arm. Thus, an isochromosome for the long arm of chromosome 17 (q17;p10) contains two copies of the long arm (separated by the centromere) with loss of the short arm of the chromosome.

isoeffect plots: Graphs of the total dose for a given effect (e.g., ED50) plotted, for instance, against dose per fraction or dose rate.

isomers: Atoms having the same number of protons and neutrons but a different nuclear energy state.

isotopes: Forms of a chemical element that have the same number of protons and electrons but differ in the number of neutrons contained in the atomic nucleus. Unstable isotopes undergo a transition to a more stable form with the release of radioactivity.

isotonic: Having equal intensity in all directions.

Karnofsky score: A measure of the patient’s overall physical health following treatment, judged by his or her level of activity.

karyotype: Arrangement of chromosomes from a particular cell according to a well-established system such that the largest chromosomes are first and the smallest ones are
LD50/30: The time between an injury occurring and late responses:

labeling index (LI): Proportion or percentage of cells in the late responses:

kilovolt (kV): A unit of length consisting of 1,000 nucleotides.
kilobase (kb): The sum of the initial kinetic energies of all the kilovolt (kV): A unit of electrical potential difference equal to 1,000 V.
lowering radiation: All radiation coming from within the source assembly except for the useful beam.

lethal dose (LD): A dose of ionizing radiation sufficient cause death. Median lethal dose (MLD, or LD 50) is the dose required to kill, within a specified period of time, half the individuals in a large group of organisms similarly exposed. The LD 50 for humans is about 4 Gy.

lethal gene: A gene whose expression results in death. The process in gametogenesis or sporogenesis during which one replication of the chromosomes is followed by two nuclear divisions to produce four haploid cells.

leukemia: A malignant cancer of the blood-forming tissues.

leukocyte: White blood cell.

lifetime risk: The risk of dying of some particular cause over the whole of a person’s life.

ligase: An enzyme that catalyzes a reaction that links two DNA molecules by the formation of a phosphodiester bond.

ligation: The process of joining two or more DNA fragments.

linear accelerator: A machine creating high-energy radiation to treat cancer. See accelerator.

linear energy transfer (LET): The rate of energy loss along the track of an ionizing particle, usually expressed in keV/μm.

linear-quadratic (LQ) model: Model in which the effect (E) is a linear-quadratic function of dose (d): $E = ad + βd^2$. For cell survival: $S = \exp(-ad - βd^2)$.

linkage: Condition in which two or more nonallelic genes tend to be inherited together. Linked genes have their loci along the same chromosome, do not assort independently, but can be separated by crossing over.

local invasion: The spread of cancer from an original site to the surrounding tissues.

local tumor control: The complete remission of a tumor without later regrowth. This requires that all cancer stem cells are inactivated.

localized tumors: Tumors that are contained in one particular site and have not spread.

locally multiply damaged site (LMDS): Any of various complex lesions, including base damage as well as double-strand breaks, produced by “spurs” and “blobs” from a high-LET track.

log-phase culture: A cell culture growing exponentially.

long terminal repeat (LTR): Sequence of several hundred base pairs found at the ends of retroviral DNAs.

lymph node: A collection of lymphocytes within a capsule and connected to other lymph nodes by fine lymphatic vessels; a common site for certain cancer cells to grow after traveling along lymphatic vessels.

lymphatic system: A network of fine lymphatic vessels that collects tissue fluids from all over the body and returns these fluids to the blood. Accumulations of lymphocytes, called lymph nodes, are situated along the course of lymphatic vessels.

lymphocyte: A type of white blood cell that helps protect the body against invading organisms by producing antibodies and regulating the immune system response.

lymphoma: A type of cancer beginning in an altered lymphocyte. There are two broad categories of lymphomas: Hodgkin disease and non–Hodgkin lymphoma.

lysis: The destruction of the cell membrane.

macrophage: A type of white blood cell assisting in the body’s fight against bacteria and infection by engulfing and destroying invading organisms.

macrophage colony-stimulating factor (M-CSF): Cytokine that stimulates formation of macrophages from pluripotent hematopoietic cells.

magnetic resonance imaging (MRI): A method of taking pictures of body tissue using magnetic fields.

mammogram: An x-ray of the breast used to detect cancer, sometimes before it can be detected by palpation. Women older than 50 years are advised to have a mammogram yearly; women in their 40s, every 2 years.

mapping: Determining the physical location of a gene or genetic marker on a chromosome.

mass number (A): The sum of the neutrons and protons in a nucleus. It is the nearest whole number to an isotope’s atomic weight. For instance, the mass number of the uranium-235 isotope is 235. Compare atomic number.

mean: Arithmetic average.

median: The value in a group of numbers below and above which there are an equal number of data points or measurements.

medical oncologist: A doctor specializing in using chemotherapy to treat cancer.

mega: A prefix that multiplies a basic unit by 1 million (10^6).

meiosis: The process in gametogenesis or sporogenesis during which one replication of the chromosomes is followed by two nuclear divisions to produce four haploid cells.

melanoma: A type of cancer that begins in the pigment-containing cells of a skin mole or the lining of the eye.

mendelian: Referring to diseases that are caused by mutations in single genes located on either the autosomes or...
the sex chromosomes and that show a simple, predictable pattern of inheritance.

**malignant:** A type of brain tumor that is relatively common and usually benign.

**messenger RNA (mRNA):** The class of RNA molecules that copies the genetic information from DNA in the nucleus and carries it to ribosomes in the cytoplasm and is translated into the amino acid sequence of a polypeptide.

**metacentric chromosome:** A chromosome with a centrally located centromere, producing chromosome arms of equal lengths.

**metaphase:** The stage of cell division in which the condensed chromosomes lie in a central plane between the two poles of the cell and in which the chromosomes become attached to the spindle fibers.

**metastasis:** The ability of cancerous cells to invade surrounding tissues, enter the circulatory system, and establish new malignancies in body tissues distant from the site of the original tumor.

**metastatic cancer:** An advanced stage of cancer in which cells from the original (primary) site have spread (metastasized) to other organs.

**methionine:** The amino acid encoded by the sequence AUG.

**methylate:** The addition of one or more methyl groups (CH3) to a molecule.

**MeV:** One million (10^6) electron volts.

**micro (μ):** A prefix that divides a basic unit by 1 million (10^-6).

**microarray:** An array of DNA spots of known sequence, usually on a glass slide, used to quantify amounts of genomic DNA or mRNA (reverse transcribed into cDNA or cRNA) present in cells or tissues. Capable of monitoring expression of all known genes and their variants. Also known as “gene expression microarrays” or “chips.”

**microinjection:** Introducing DNA into a cell using a fine micropipette.

**micrometer (μm):** A unit of length equal to 1 × 10^-6 m. Previously called a micron.

**micron:** See micrometer.

**migration coefficient:** An expression of the proportion of migrant genes entering the population per generation.

**millimeter (mm):** A unit of length equal to one thousandth of a meter.

**minimal medium:** A medium containing only those nutrients that support the growth and reproduction of wild-type strains of an organism.

**miRNA:** MicroRNAs, small (19 to 22 nucleotides) single-stranded noncoding RNAs expressed in cells that can regulate gene expression.

**mismatch repair (MMR):** DNA repair pathway for repairing or replacing mismatched bases in DNA.

**misrepair (error-prone repair):** Reconstitution with a loss of information (e.g., deletion caused by the loss of a fragment of the molecule or mutation or translocation).

**missense mutation:** A mutation that alters a codon to that of another amino acid, causing an altered translation product to be made.

**mitochondrion:** Found in the cells of eukaryotes, a cytoplasmic, self-reproducing organelle that is the site of ATP synthesis.

**mitogen:** A substance (e.g., phytohemagglutinin) that stimulates mitosis in nondividing cells.

**mitogen-activated protein (MAP) kinase:** A family of two protein kinases of 42 and 44 kDa (ERK1 and ERK2) and 38 kDa that act to induce certain early-response genes.

**mitosis:** The replication of a cell to form two progeny cells with identical sets of chromosomes.

**mitotic death:** Cell death associated with a postirradiation mitosis.

**mitotic delay:** Delay of entry into mitosis, or accumulation in G2, as a result of treatment.

**mitotic index (MI):** Proportion or percentage of cells in mitosis at any given time.

**MMP-9:** Matrix metalloproteinase 9 encodes a 92 kDa type IV collagenase. This enzyme is involved in breaking down the extracellular matrix and has been implicated in the metastatic process of tumor cells.

**molecular imaging:** Medical imaging visualizing the spatial distribution of molecular targets, signaling pathways, or cellular phenotypes. Examples include PET and SPECT.

**molecule:** A group of atoms held together by chemical forces. The atoms of the molecule may be identical, as in H2, S2, and S8, or different, as in H2O and CO2. A molecule is the smallest unit of matter that can exist by itself and retain all its chemical properties.

**monoclonal antibodies:** Immunoglobulin molecules of single-epitope specificity that are secreted by a clone of B cells.

**mosaicism:** Describing an aneuploid condition in which one member of a chromosome pair is missing; having a chromosome number of 2n - 1.

**monozygotic twins:** Twins produced from a single fertilization event; the first division of the zygote produces two cells, each of which develops into an embryo. Also known as identical twins.

**morbidity:** Sickness, side effects, and symptoms of a treatment or disease.

**mRNA:** See messenger RNA.

**mtDNA:** Mitochondrial DNA.

**mucositis:** Inflammation of the lining of areas such as the mouth.

**multifactorial:** Referring to diseases known to have a genetic component but whose transmission patterns cannot be described as simple mendelian.

**multitarget equation:** Model that assumes the presence of a number of critical targets in a cell, all of which require inactivation to kill the cell. Survival is given by S = 1 - (1 - exp [D/D0])n.

**mutagen:** Any agent that causes an increase in the rate of mutation.

**mutant:** A cell or organism carrying an altered or mutant gene.

**mutation:** A relatively stable change in the DNA of the cell nucleus. Mutations in the germ cells of the body (ova and sperm) may lead to inherited effects in the offspring. Mutations in the somatic cells of the body may lead to effects in the individual (e.g., cancer).

**mutation component (MC):** This allows for the observation that only a proportion of mutations lead to a disease.

**mutation rate:** The frequency with which mutations take place at a given locus or in a population.

**myeloma:** A tumor of the cells of the bone marrow.

**nano (n):** A prefix that divides a basic unit by 1 billion (10^-9).

**nanometer (nm):** A unit of length equal to 1 × 10^-9 m.
nasopharynx: Part of the breathing passage behind the nasal cavity.
natural radioactivity: See background radiation.
natural selection: Differential reproduction of some members of a species resulting from variable fitness conferred by genotypic differences.
natural uranium: Uranium as found in nature, containing 0.7% of the isotope $^{235}\text{U}$, 99.3% of $^{238}\text{U}$, and a trace of $^{234}\text{U}$.
negligible individual dose: Level of effective dose to an individual per source or practice that may be ignored. Defined by National Council on Radiation Protection and Measurements as 0.01mSv.
neutrino ($\nu$): An electrically neutral elementary particle with a negligible mass. It interacts very weakly with matter and hence is difficult to detect. It is produced in many nuclear reactions (e.g., in $\beta$ decay) and has high penetrating power. Neutrinos from the sun usually pass right through the earth.
neutron ($n$): An uncharged elementary particle that has a mass slightly greater than that of the proton and that is found in the nucleus of every atom heavier than hydrogen. A free neutron is unstable and decays with a half-life of about 13 minutes into an electron, a proton, and a neutrino. Neutrons sustain the fission chain reaction in a nuclear reactor.
NHEJ: Nonhomologous end-joining. A DNA repair pathway for repairing double-strand DNA breaks without using any homologous sequence as a template.
nondisjunction: An accident of cell division in which the homologous chromosomes (in meiosis) or the sister chromatids (in mitosis) fail to separate and migrate to opposite poles; responsible for defects such as monosomy and trisomy.
nonsense mutation: A mutation that alters a codon to one that encodes no amino acid; for example, UAG (amber stop codon), UAA (ochre codon), or UGA (opal codon). Leads to premature termination during the translation of mRNA.
nonstochastic effect: An effect, the severity of which increases with increasing dose, after a threshold region. Now called deterministic effect.
normal distribution: A probability function that approximates the distribution of random variables. The normal curve, also known as a Gaussian or bell-shaped curve, is the graphic display of the normal distribution.
northern blotting: A procedure in which RNA fragments are transferred from an agarose gel to a nitrocellulose filter from which the RNA is then hybridized to a radioactive probe.
NSD: Nominal standard dose in the Ellis formula.
nuclear fuel cycle: Activities associated with production, utilization, and disposition of fuel for nuclear reactors and related by-products.
nuclear reactor: A structure in which nuclear fission may be sustained in a self-supporting chain reaction. In thermal reactors, the fission is produced by fission neutrons, and in fast reactors by fast neutrons.
nuclease: An enzyme that breaks bonds in nucleic acid molecules.
nucleic acid: A class of organic acids that play a role in protein synthesis, in the transmission of heritable traits, and in the control of cellular activities.
nucleon: Proton or neutron.
nucleoside: A purine or pyrimidine base covalently linked to a ribose or deoxyribose sugar molecule.
nucleotide: A building block of DNA and RNA, consisting of a nitrogenous base, a five-carbon sugar, and a phosphate group.
nucleotide pair: The pair of nucleotides (A and T or G and C) in opposite strands of the DNA molecule that are hydrogen-bonded to each other.
nucleus (of a cell): The membrane-bound region of a eukaryotic cell that contains the chromosomes.
nucleus (of an atom): The small, positively charged core of an atom. It is only about 1/10,000 the diameter of the atom but contains nearly all the atom’s mass. All nuclei contain both protons and neutrons, except the nucleus of ordinary hydrogen, which consists of a single proton.
nucleide: A general term applicable to all atomic forms of the elements. The term is often used incorrectly as a synonym for isotope, which properly has a more limited definition. Whereas isotopes are the various forms of a single element (hence are a family of nuclides) and all have the same atomic number and number of protons, nuclides comprise all the isotopic forms of all the elements. Nuclides are distinguished by their atomic number, atomic mass, and energy state.
occupationally exposed: Exposed to radiation as a direct result of occupational duties.
oligonucleotide: A DNA polymer composed of only a few nucleotides.
oncogene: A gene that contributes to cancer formation when mutated or inappropriately expressed.
oncogenic: Having the potential to cause cancer (same as carcinogenic).
oncologist: A physician specializing in the study and treatment of cancer.
oncology: The study of cancer.
orbit: A four-letter word.
organ or tissue weighting factor (Wt): A factor that indicates the ratio of the risk of stochastic effects attributable to irradiation of a given organ or tissue to the total risk if the whole body is uniformly irradiated. Organs that have a large tissue weighting factor are those susceptible to radiation-induced carcinogenesis (such as the breast or thyroid).
oxidative stress: Formation of reactive oxygen species (ROS) in and outside cells, such as those resulting from the lysis of water molecules induced by ionizing radiation. This stress not only can activate several enzyme systems, but can also modify the transcription of genes. These reactions are known collectively as oxidative stress.
oxrogen enhancement ratio (OER): The ratio of the radiation dose given under anoxic conditions to produce a given effect relative to the radiation dose given under fully oxygenated conditions to produce the same effect.
p15: G1 inhibitor induced in epithelial cells by TGF-β. Inhibits cyclin D1–Cycl4 and cyclin D1–Cdk6 complexes.
p16: G1 inhibitor of epithelial cells. Inhibits cyclin D1–Cycl4 and cyclin D1–Cdk6 complexes. Gene is deleted in familial melanomas and other tumor types.
**p21 (WAF1):** Inhibitor of Cdc2, Cdk4, and Cdk6. Induced through p53 pathway.

**p27:** Cell cycle inhibitor induced in epithelial cells by TGF-β. Inhibits cyclin E–Cdk2 complex.

**p53:** Considered the guardian of the genome. Mediates cellular responses to DNA-damaging agents such as ionizing radiation at the G1 checkpoint; induces p21. The gene p53 on chromosome 17 is mutated in colon, breast, esophageal, and other various human cancers. Binds DNA; can act as transcription factor.

**palindrome:** A word, number, verse, or sentence that reads the same backward or forward. In nucleic acids, a sequence in which the base pairs read the same on complementary strands (5’ → 3’). For example: 5’ GAAATTC 3’, 3’ CTTAGG 5’. These often occur as sites for restriction endonuclease recognition and cutting.

**palliative care:** Treatment to relieve, rather than cure, symptoms caused by cancer. Palliative care can help people live more comfortably.

**palpate:** To examine by carefully feeling with the fingers.

**paracentric inversion:** A chromosomal inversion that does not include the centromere.

**parent:** Radioactive atom that disintegrates to a different atom, its progeny.

**particle:** A minute constituent of matter, generally one with a measurable mass. The primary particles involved in radioactivity are α-particles, β-particles, neutrons, and protons.

**pathologist:** A specialist who attempts to describe the nature of a disease by analyzing samples obtained from tissues, organs, or body fluids.

**pathology:** The study of diseased tissues, both by gross and by microscopic examination of tissues removed during surgery and postmortem.

**pedigree:** In human genetics, a diagram showing the ancestral relationships of a given genotype manifest in a specific mutant phenotype associated with a trait.

**penetration:** The frequency (expressed as a percentage) with which individuals of a given genotype manifest at least some degree of a specific mutant phenotype associated with a trait.

**peptide bond:** The covalent bond between the amino group of one amino acid and the carboxyl group of another amino acid.

**pericentric inversion:** A chromosomal inversion that involves both arms of the chromosome and thus involves the centromere.

**Person-sievert:** The unit of collective dose.

**phage:** See bacteriophage.

**pharynx:** Medical term for the throat from the nasal and oral cavities above to the larynx and esophagus below.

**phosphodiester bond:** In nucleic acids, the covalent bond between a phosphate group and adjacent nucleotides, extending from the 5’ carbon of one pentose (ribose or deoxyribose) to the 3’ carbon of the pentose in the neighboring nucleotide. Phosphodiester bonds form the backbone of nucleic acid molecules.

**photodynamic therapy:** Cancer treatment using light to activate a photosensitizing agent, thereby releasing cytotoxic free radicals.

**photoelectric effect:** Absorption of an x-ray by ionization.

**photon:** The carrier of a quantum of electromagnetic energy. Photons have an effective momentum but no mass or electrical charge.

**pico:** A prefix that divides a basic unit by 1 trillion (10^{-12}).

**picocuries per liter (pCi/L):** A unit of measurement of the activity concentration of a radioactive material; measures, for example, how many radioactive disintegrations of radon occur every second in a liter of air.

**plaque:** A clear spot on a lawn of bacteria or cultured cells on which cells have been lysed by viral infection and replication.

**plasmid:** A circular DNA molecule, capable of autonomous replication, which typically carries one or more genes encoding antibiotic resistance proteins.

**plateau-phase cultures:** Cell cultures grown to confluence so that proliferation is markedly reduced (also called stationary phase).

**platelet-derived growth factor (PDGF):** A protein that induces growth fibroblasts and is involved in wound healing. Also acts on some epithelial and endothelial cells and on mesenchymal cells.

**platelets:** Special blood cells that help stop bleeding.

**plating efficiency:** The proportion or percentage of in vitro-plated cells that form colonies.

**pleiotropy:** Condition in which a single mutation simultaneously affects several characters.

**ploidy:** Relates to the number of sets of chromosomes in a cell. Diploid cells have two sets of chromosomes, a chromosome complement twice that found in the gametes. Tetraploid cells have four sets of chromosomes.

**point mutation:** A mutation that can be mapped to a single locus. At the molecular level, a mutation that results in the substitution of one nucleotide for another.

**polar body:** A cell that is produced at either the first or second meiotic division in females and which contains almost no cytoplasm as a result of an unequal cytokinesis.

**polyacrylamide gel electrophoresis:** Electrophoresis through a matrix composed of a synthetic polymer, used to separate small DNA or RNA molecules (up to 1,000 nucleotides) or proteins.

**polyclonal antibodies:** A mixture of immunoglobulin molecules secreted against a specific antigen, each recognizing a different epitope.

**polymerase:** An enzyme that catalyzes the addition of multiple subunits to a substrate molecule.

**polymerase chain reaction (PCR):** A procedure that enzymatically amplifies a DNA sequence through repeated replication by DNA polymerase.

**polymorphism:** The existence of two or more discontinuous, segregating phenotypes in a population.

**polypeptide:** A molecule made up of amino acids joined by covalent peptide bonds. This term is used to denote the amino acid chain before it assumes its functional three-dimensional configuration.

**polyploid:** A cell or individual having more than two sets of chromosomes.

**population:** A local group of individuals that belong to the same species and that are actually or potentially interbreeding.

**positron (β^+):** An elementary particle with the mass of an electron but charged positively. It is the “antineutron.” It is emitted in some radioactive disintegrations and is formed by the interaction of high-energy γ-rays with matter.
positron emission tomography (PET): An imaging technique using radionuclides that emit positrons whose annihilation photons are imaged in coincidence to form tomographic views of the body.

potential doubling time ($T_{\text{pot}}$): Tumor doubling time, taking into account the cell cycle time and the growth fraction, but ignoring cell loss.

potentially lethal damage (PLD): Cellular damage that is repaired during the interval between treatment and assay, especially under suboptimal growth conditions.

pRb: A protein of ~110 kDa that regulates cell cycle progression through G1. Phosphorylated by cyclin D–Cdk complexes to release G1 transcription factors. Inactivated in hereditary retinoblastoma and sporadic tumors of the bone, breast, esophagus, and other tissues.

precursor: In a radioactive decay chain, a member of the decay chain that occurs before a particular atom in question.

pre-mRNA: The initial mRNA transcript prior to any mRNA processing.

primary cell: A cell or cell line that is taken directly from a living organism and that is not immortalized.

primary tumor: The place in which a cancer originates, which is referred to regardless of the site of its eventual spread. Thus, prostate cancer that spreads to the bone is still prostate cancer and is not referred to as bone cancer.

primer: A short DNA or RNA fragment annealed to single-stranded DNA.

primordial: Existing at the beginning of the universe or at the beginning of the earth.

prion: An infectious pathogenic agent devoid of nucleic acid and composed mainly of a protein, PrP, with a molecular weight of 27,000 to 30,000 Da. Prions are known to cause scrapie, a degenerative neurologic disease in sheep, and are thought to cause similar diseases in humans, such as kuru and Creutzfeldt-Jakob disease.

probability: Ratio of the frequency of a given event to the frequency of all possible events.

probe: A single-stranded DNA (or RNA) that has been labeled radioactively and is used to identify complementary sequences.

prodromal phase: Signs and symptoms in the first 48 hours following irradiation.

progeny: Formerly called a “daughter” in a radioactive decay chain.

prognosis: The predicted or likely outcome.

programmed cell death: Cell death that occurs as the result of an active process carried out by molecules in the cell, for example, apoptosis.

prokaryotes: Organisms lacking nuclear membranes, meiosis, and mitosis. Bacteria and blue-green algae are examples of prokaryotic organisms.

promoter: A region of DNA extending 150 to 300 bp upstream from the transcription start site that contains binding sites for RNA polymerase and several proteins that regulate the rate of transcription of the adjacent gene.

promoter site: Region having a regulatory function and to which RNA polymerase binds prior to the initiation of transcription.

promoting agent: Something that acts on earlier cellular damage caused by an initiating agent; can cause the earlier damage to be expressed as cancer. Tobacco smoke usually is considered a promoting agent.

prophylactic: Preventive measure or medication.

prostate: A gland at the base of the bladder in males for the production of seminal fluids. Cancer of this gland is common in elderly men.

protein: A molecule composed of one or more polypeptides, each composed of amino acids covalently linked together.

protein kinase: An enzyme that adds phosphate groups to a protein molecule at serine, threonine, or tyrosine residues.

protein kinase C (PKC): A family of protein kinases involved in mitogenic signaling. Activated by second messengers, including diacylglycerol and Ca$^{2+}$ (some isoforms). Can be activated directly by the phorbol ester class of tumor promoters. Can induce early-response genes through ras.

proteomics: Study of the proteins expressed in cells including structure and function.

protocol: A standardized combination of therapies developed specifically for particular tumors.

proton: An elementary particle that is a component of all nuclei and that has a single positive electrical charge and a mass approximately 1,837 times that of the electron. The nucleus of an ordinary or light hydrogen atom. The atomic number of an atom is equal to the number of protons in its nucleus.

proto-oncogene: A gene generally active in the embryo and fetus and during proliferation processes. A mutation can result in the permanent activation of a proto-oncogene, which then becomes an oncogene.

PTEN: PTEN stands for Phosphatase and Tensin Homolog. The gene is commonly found mutated in many different cancers and is classified as a tumor suppressor gene. The gene encodes a phosphatidylinositol-3,4,5-trisphosphate 3-phosphatase that dephosphorylates phosphoinositide substrates and is critical in the regulation of the Akt signaling pathway.

pulsed-field gel electrophoresis (PFGE): Process whereby current is alternated between pairs of electrodes set at angles to one another to separate very large DNA molecules of up to 10 million nucleotides.

purines: Organic bases with carbon and nitrogen atoms in two interlocking rings; components of nucleic acids and other biologically active substances.

pyrimidines: Nitrogenous bases composed of a single ring of carbon and nitrogen atoms; components of nucleic acids.

quality factor: The factor by which absorbed dose is multiplied to obtain a quantity that expresses on a common scale, biological effects (risks) from exposure to different radiations. Largely replaced by radiation weighting factor.

quasi-threshold dose ($D_0$): Point of extrapolation of the exponential portion of a multitarget survival curve to the level of zero survival: $D_s = D_0 \ln(n)$.

rad: The old unit of absorbed dose, equivalent to an energy absorption of $10^{-2}$ J/kg. Replaced by the gray (Gy). See absorbed dose.

radiation: The emission and propagation of energy through matter or space by means of electromagnetic disturbances (x-rays) that display both wavelike and particle-like behavior; in this context the “particles” are known as photons. Also, the energy so propagated. The term has been extended to include streams of fast-moving particles, such as α- and β-particles, free neutrons, and cosmic radiation. Nuclear radiation is that emitted from atomic nuclei in various nuclear reactions, including α-, β-, and γ-ray radiation and neutrons.
radiation (ionizing): Any electromagnetic or particulate radiation capable of producing ions, directly or indirectly, by interaction with matter. Examples are x-rays, photons, charged atomic particles and other ions, and neutrons.
radiation absorbed dose: See Gray.
radiation accidents: Accidents resulting in the spread of radioactive material or in the exposure of individuals to radiation.
radiation burn: Radiation damage to the skin. β-burns result from skin contact with or exposure to emitters of β-particles. See beta particle, ionizing radiation.
radiation chemistry: The branch of chemistry that is concerned with the chemical effects, including decomposition, of energetic radiation or particles on matter.
radiation detriment: Measure of stochastic effects from exposure to ionizing radiation that takes into account the probability of fatal cancers, probability of heritable effects in future generations, probability of nonfatal cancers weighted by the lethality fraction, and the relative years of life lost per fatal health effect.
radiation dose: The amount of radiation absorbed by an irradiated object. This unit is the gray (Gy), defined to be 1 J/kg.
radiation illness: An acute organic disorder that follows exposure to relatively large doses of ionizing radiation. It is characterized by nausea, vomiting, diarrhea, blood cell changes, and (in later stages) hemorrhage and loss of hair.
radiation protection: Legislation and regulations to protect the public and workers against radiation. Also, measures to reduce exposure to radiation.
radiation quality: Relative penetrability of an x-ray beam to reduce exposure to radiation.
radiation shielding: Reduction of radiation by interposing a shield of absorbing material between any radioactive source and a person, laboratory area, or radiation-sensitive device.
radiation sterilization: Using radiation to make a plant or animal sterile. Also, radiation to kill all forms of life, especially bacteria, in food or surgical equipment.
radiation warning symbol: An officially prescribed symbol, a magenta trefoil on a yellow background, that should be displayed whenever a radiation hazard exists.
radiation weighted dose: New name proposed by ICRP for equivalent dose.
radiation weighting factor (W R): A factor used for radiation protection purposes that accounts for differences in biologic effectiveness between different radiations. The radiation weighting factor is independent of the tissue weighting factor.
radiative decay: See decay, radioactive.
radiative half-life: Time for a radioisotope to decay to one-half its activity.
radiative isotope: One of the forms of an element, differing in atomic weight and possessing an unstable nucleus that emits ionizing radiation.
radiactive series: A succession of nuclides, each in turn transforming by radioactive disintegration into the next nuclide until a stable nuclide is reached. The first member is called the parent, the intermediate members are called progeny, and the final, stable member is called the end product.
radiative tracer: A small quantity of a radioactive isotope, either with or without a carrier, used to follow biologic, chemical, or other processes by detection, determination, or localization of the radioactivity.
radiative waste: Unwanted radioactive materials in any form. Often categorized in the nuclear power industry into low-level, intermediate-level, and high-level waste.
radiative activity: A property of all unstable elements that regularly decay to an altered state by releasing energy in the form of photons (γ-rays) or particles (e.g., electrons, α-particles).
radiobiology: The body of knowledge and the study of the principles, mechanisms, and effects of ionizing radiation on living matter.
radiogenic: Of radioactive origin; produced by radioactive transformation.
radiography: The production of images on film or other media by the action of x-rays transmitted through a patient.
radioisootope: A radioactive isotope; an unstable isotope of an element that decays or disintegrates spontaneously, emitting radiation. More than 1,300 natural and artificial radioisotopes have been identified.
radiologist: A physician with special training in reading diagnostic x-rays and performing specialized x-ray procedures.
radiology: The science that investigates all forms of ionizing radiation in the diagnosis and treatment of disease.
radiouclide: A radioactive nuclide.
radioprotector: A chemical compound that reduces the biologic consequences of radiation.
radioreistance: A relative resistance of cells, tissues, organs, or organisms to the harmful action of radiation.
radiosensitivity: A general term indicating the overall level of clinical response to radiotherapy.
radiosensitivity: (1) A relative susceptibility of cells, tissues, organs, or organisms to the effects of radiation. (2) The radiation dose required to produce a defined level of cell inactivation, usually indicated by the surviving fraction at 2 Gy (i.e., SF2) or by the parameters of the linear-quadratic or multitarget equations.
radiosensitizer: In general, any agent that increases the sensitivity of cells to radiation. Most commonly applied to electron-affinic chemicals that mimic oxygen in fixing free radical damage.
radium (Ra): A radioactive metallic element with atomic number 88. As found in nature, the most common isotope has an atomic weight of 226. It occurs in minute quantities associated with uranium in pitchblende, carnotite, and other minerals. The uranium decays to radium in a series of α- and β-emissions. By virtue of being an α- and γ-emitter, radium is used as a source of luminescence and as a radiation source in medicine and radiography.
radium (Rn): Colorless, odorless, naturally occurring radioactive gas; a radioactive element and the heaviest gas,
known. Its atomic number is 86 and its atomic weight varies from 200 to 226. Radon-222 is the progeny of radium in the uranium radioactive series.

radon daughter: Any atom that is below radon-222 in the uranium decay chain; often specifically refers to polonium-218 and polonium-214, as these have the most biologic significance; now referred to as radon progeny.

raf: A protein kinase that is activated by GTP-bound ras. Acts to transduce mitogenic signaling by phosphorylation of MAP kinases.

ran: A family of 21-kDa proteins (H-, K-, and N-ran) found to be activated by point mutations at codons 12, 13, and 61 in various tumors. Involved in mitogenic signaling, coupling growth signals from growth factor receptors to raf activation, and downstream stimulation of early-response genes. Binds GTP in its activated state. Is found at the inner face of the cell membrane.

RBE: See relative biologic effectiveness.

reading frame: A series of triplet codons beginning from a specific nucleotide.

reassortment (redistribution): Return toward a more even cell age distribution, following the selective killing of cells in certain phases of the cell cycle.

recessive: Term describing an allele that is not expressed in the heterozygous condition.

recessive gene: Gene whose phenotype is expressed only when both copies of the gene are mutated or missing.

recessive-acting oncogene (anti-oncogene): A single copy of this gene is sufficient to suppress cell proliferation; the loss of both copies of the gene contributes to cancer formation.

reciprocal translocation: A chromosomal aberration in which nonhomologous chromosomes exchange parts.

recombinant DNA: The process of cutting and recombining DNA fragments as a means to isolate genes or to alter their structure and function.

recovery: An increase in cell survival as a function of time during or after irradiation. See repair.

recurrence: The return of a cancer after all detectable traces had been removed by primary therapy; recurrences may be local (near the primary site) or distant (metastatic).

regression rate: The rate at which the tumor volume shrinks during or after treatment with radiation or a chemotherapy agent.

relapse: Recurrence of a disease following treatment.

relative biologic effectiveness (RBE): A factor used to compare the biologic effectiveness of different types of ionizing radiation. It is the inverse ratio of the amount of absorbed radiation required to produce a given effect to a standard (or reference) radiation required to produce the same effect.

relative risk: Situation in which the risk of a disease resulting from some injury is expressed as some percentage increase of the normal rate of occurrence of that disease; in contrast to an absolute risk in which the risk of a disease resulting from an injury does not depend on the normal rate of occurrence of that disease.

rem: Old unit of equivalent or effective dose. It is the product of absorbed dose (in rad) and the radiation weighting factor. One rem is one hundredth of a sievert.

remediation: Reducing a home's indoor radon level.

remission, complete: Condition in which no cancerous cells can be detected and the patient appears to be free from disease.
of electrical charge (either positive or negative) in 1 cm³ of dry air under standard conditions.

**rRNA**: The RNA molecules that are the structural components of the ribosomal subunits. In prokaryotes, these are the 16S, 23S, and 5S molecules; in eukaryotes, they are the 18S, 28S, and 5S molecules.

**Saccharomyces cerevisiae**: Brewer’s yeast.

**sarcoma**: A type of cancer derived from connective bone or fat tissue. Examples include fibrosarcoma, osteogenic sarcoma, and liposarcoma.

**satellite DNA**: DNA that forms a minor band if genomic DNA is centrifuged in a cesium salt gradient. This DNA usually consists of a short sequence repeated many times in the genome.

**scan**: A diagnostic test usually involving the movement of a detector to produce a picture. Examples include ultrasound, nuclear medicine, computer-assisted tomographic, and magnetic resonance scans.

**scattered radiation**: Radiation that, during passage through matter, is changed in direction (the change is usually accompanied by a decrease in energy).

**SCE**: Sister chromatid exchange.

**Scheimpflug imaging system**: An imaging system that gives an objective and quantitative assessment of the severity of an ocular cataract. Named after the Austrian army officer Theodor Scheimpflug (1865–1911).

**SDF-1**: The ligand for the chemokine receptor CXCR4. Plays important roles in chemotaxis and vascularization.

**secondary cancer**: Cancer arising from a primary cancer; metastatic cancer.

**segregation**: The separation of homologous chromosomes into different gametes during meiosis.

**selectable marker**: A gene whose expression makes it possible to identify cells that have been transformed or transfected with a vector containing the marker gene. It is usually a gene for resistance to an antibiotic.

**selection**: The force that brings about changes in the frequency of alleles and genotypes in populations through differential reproduction.

**semiconservative replication**: A model of DNA replication in which a double-stranded molecule replicates in such a way that the progeny molecules are composed of one parental (old) and one newly synthesized strand.

**senescence**: Permanent arrest of cell division associated with aging, differentiation, or cell damage.

**sensitizing agent**: A substance that increases the biologic effectiveness of a given dose of radiation.

**sex chromosome**: A chromosome, such as the X or Y in humans, that is involved in sex determination.

**sex linkage**: The pattern of inheritance resulting from genes located on the X chromosome.

**sexual reproduction**: Reproduction through the fusion of gametes, which are the haploid products of meiosis. 

**SF₂**: Surviving fraction at 2 Gy.

**sickle cell anemia**: A genetic disease in humans caused by an autosomal recessive gene, usually fatal in the homozygous condition. Caused by an alteration in the amino acid sequence of the β chain of globin.

**side effects**: Symptoms directly related to treatment, such as the side effect of nausea resulting from radiation treatment over the stomach. Side effects are considered acute if they occur during treatment and subside when treatment is complete. Those symptoms persisting over a longer period are considered chronic.

**sievert (Sv)**: Unit of equivalent dose or effective dose. It is equal to the dose in gray multiplied by a weighting factor. One sievert equals 100 rem.

**SINES**: Short interspersed repetitive sequences found in the genomes of higher organisms, such as the 300-bp Alu sequence.

**single photon emission computed tomography (SPECT)**: An imaging technique in which one or more gamma cameras sample a region of the body from several angles, producing tomographic images (slices) of the body.

**sister chromatid exchange (SCE)**: A crossing-over event that can occur in meiotic and mitotic cells. Involves reciprocal exchange of chromosomal material between sister chromatids (joined by a common centromere). Such exchanges can be detected cytologically after BrdUrd incorporation into the replicating chromosomes.

**slow repair**: Long-term recovery that takes place on a time scale of weeks to months.

**solid tumor**: A cancer originating in an organ or tissue other than bone marrow or the lymph system.

**somatic**: Pertaining to the body; pertaining to all cells except the germ cells.

**somatic cell**: Any cell other than a germ cell that composes the body of an organism and that possesses a set of multiploid chromosomes.

**somatic effects of radiation**: Effects of radiation limited to the exposed individual, as distinguished from genetic effects, which also affect subsequent, unexposed generations. Large radiation doses can be fatal. Smaller doses may make the individual noticeably ill, may produce temporary changes in blood cell levels detectable only in the laboratory, or may produce no detectable effects.

**somatic mutation**: A mutational event occurring in a somatic cell. Such mutations are not heritable.

**SOS response**: The induction of enzymes to repair damaged DNA in Escherichia coli. The response involves activation of an enzyme that cleaves a repressor, activating a series of genes involved in DNA repair.

**Southern blotting**: A procedure in which DNA restriction fragments are transferred from an agarose gel to a nitrocellulose filter, where the denatured DNA is then hybridized to a radioactive probe.

**spatial cooperation**: The use of radiotherapy and chemotherapy to treat disease in different anatomic sites.

**species**: A group of actually or potentially interbreeding individuals that are morphologically distinguishable.

**specific activity**: The radioactivity of a radioisotope of an element per unit weight of the element in a sample; the activity per unit mass of a pure radionuclide; the activity per unit weight of any sample of radioactive material.

**spheroid**: Clump of cells grown together in tissue culture suspension.

**spindle fibers**: Cytoplasmic fibrils formed during cell division that are involved in the separation of chromatids at anaphase and their movement toward opposite poles in the cell.
**split-dose (SLD) recovery:** Decrease in radiation effect if a single radiation dose is split into two fractions separated by times up to a few hours. Also called Elkind recovery or recovery from sublethal damage.

**spontaneous mutation:** A mutation that is not induced by a mutagenic agent.

**spore:** A unicellular body or cell that is encased in a protective coat and that is produced by some bacteria, plants, and invertebrates. Capable of survival in unfavorable environmental conditions and can give rise to a new individual upon germination. In plants, spores are the haploid products of meiosis.

**spur:** A concentration of about three ion pairs in a volume about 4 nm in diameter. See locally multiply damaged site.

**SSBR:** Single-strand break repair. DNA repair pathway for repairing a break in only one of the DNA strands.

**stable isotope:** An isotope that does not undergo radioactive decay. Compare radioisotope.

**stage:** The anatomic extent of a cancer. Cancer may exist in the organ of origin and extend locally, or it may spread to regional tissues, then to local lymph nodes, and then to distant areas as metastases.

**standard deviation:** The radiation dose that results in a 50% probability of tumor control.

**standard error:** A quantitative measure of the amount of variation in a sample of measurements from a population.

**stathmokinetic method:** Study of cell proliferation using agents that block cells in mitosis.

**stem cells:** Cells capable of self-renewal and of differentiation to produce all the various types of specialized cells in a lineage.

**sterility:** The condition of being unable to reproduce; the condition of being free from contaminating microorganisms.

**sticky end:** A single-stranded nucleotide sequence produced if a restriction endonuclease cleaves off-center in its recognition sequence.

**stochastic effects:** Effects, the probability of which, rather than their severity, is a function of radiation dose without threshold. (More generally, stochastic means random in nature.)

**strain:** A group with common ancestry that has physiologic or morphologic characteristics of interest for genetic study or domestication.

**stringency:** Reaction conditions, such as temperature, salt, and pH, that dictate the annealing of single-stranded DNA/DNA, DNA/RNA, and RNA/RNA hybrids. At high stringency, duplexes form only between strands with perfect one-to-one complementarity; lower stringency allows annealing between strands with less than a perfect match between bases.

**sublethal damage (SLD):** Nonlethal cellular injury that can be repaired or accumulated with further dose to become lethal.

**submetacentric chromosome:** A chromosome with the centromere placed so that one arm of the chromosome is slightly longer than the other.

**suppressor genes:** Genes that oppose the continuous proliferation of cells. Also known as tumor suppressor genes.

**supra-additivity (synergism):** A biologic effect caused by a combination of effects that is greater than would be expected from the addition of the effects of the component agents.

**target cell:** A stem cell whose death contributes to a reduction in growth or tissue function.

**target theory:** (1) A theory based on the idea that death of a cell is caused by the inactivation of specific targets within the cell. (2) The idea that the shoulder on cell survival curves is a result of the number of unrepaird lesions per cell.

**targeted agents:** Small molecules or antibodies that inhibit cellular pathways that are specific to cancer cells, or substantially overexpressed in malignant compared with normal cells.

**targeted radiotherapy:** Treatment of disseminated cancer by means of drugs that localize in tumors and carry therapeutic amounts of radioactivity.

**TBI:** Total body irradiation.

**TCD50:** The radiation dose that results in a 50% probability of tumor control.

**TCP:** Tumor control probability.

**telangiectasia:** Pathologically dilated capillaries, observed in tissues as a late effect of radiation.

**telomeres:** Long arrays of TTAGGG repeats that cap and protect the ends of chromosomes. Each time a normal somatic cell divides, the terminal end of the telomere is lost.

**telophase:** The stage of cell division in which the progeny chromosomes reach the opposite poles of the cell and re-form nuclei. Telophase ends with the completion of cytokinesis.

**temperature-sensitive mutation:** A conditional mutation that produces a mutant phenotype at one temperature range and a wild-type phenotype at another temperature range.

**template:** An RNA or single-stranded DNA molecule on which a complementary nucleotide strand is synthesized.

**teratocarcinoma:** Embryonal tumors that arise in the yolk sac or gonads and are able to undergo differentiation into various cell types. These tumors are used to investigate the regulatory mechanisms underlying development.

**termination (stop) codon:** Any of three mRNA sequences (UGA, UAG, UAA) that do not code for an amino acid and thus signal the end of protein synthesis.

**therapeutic gain factor:** In hyperthermia, the ratio of the thermal enhancement ratio in the tumor to the thermal enhancement ratio in normal tissue. For high-linear energy transfer radiations, the therapeutic gain factor is the ratio of the relative biologic effectiveness in the tumor to the relative biologic effectiveness in normal tissue.
therapeutic index (therapeutic ratio): Tumor response for a fixed level of normal-tissue damage.

thermal dose: A function of temperature and heating time that is thought to relate well to biologic effect. It is defined to be the cumulative equivalent minutes at 43° C.

thermal enhancement ratio (TER): The ratio of radiation doses, with and without heat, to produce the same biologic effects.

thermal neutrons: Neutrons in thermal equilibrium with their surrounding medium. Thermal neutrons are those that have been slowed down by a moderator to an average speed of about 2,200 m/s at room temperature from the much higher initial speeds that they had when expelled by fission.

thermoluminescent dosimeter (TLD): A dosimeter containing a crystalline solid for measuring radiation dose, plus filters (absorbers) to help characterize the types of radiation encountered. If heated, TLD crystals that have been exposed to ionizing radiation give off light proportional to the energy they received from the radiation.

thermotolerance: The induced resistance to a second heat exposure by prior heating.

thorium series: Radioactive decay chain starting with thorium-232; one member of the chain is radon-220. This chain is of much less significance than the uranium series, and other agents.

threshold: A level (e.g., of radiation dose) below which there is no observable effect. There is no threshold for induction of cancer by radiation: All levels of radiation are considered harmful.

threshold dose: The minimum dose of radiation that produces a detectable biologic effect.

thymidine kinase (tk): An enzyme that allows a cell to use an alternate metabolic pathway for incorporating thymidine into DNA. Used as a selectable marker to identify transfected eukaryotic cells.

thymine dimer: A pair of adjacent thymine bases in a single polynucleotide strand between which chemical bonds have formed. This lesion, usually the result of damage caused by exposure to ultraviolet light, inhibits DNA replication unless repaired by the appropriate enzyme system.

time-dose relationships: The dependence of isoeffective radiation dose on the duration (and number of fractions) in radiotherapy.

tissue weighting factor: A factor that indicates the ratio of the risk of stochastic effects attributable to irradiation of a given organ or tissue to the total risk when the whole body is uniformly irradiated.

tolerance: The maximum radiation dose or intensity of fractionated radiotherapy that the therapist judges to be acceptable, usually expressed in dose units. Actual values depend on fractionation, field size, concomitant treatments, and so on.

topoisomerase: A class of enzymes that converts DNA from one topologic form to another. During DNA replication, these enzymes facilitate the unwinding of the double-helical structure of DNA.

totipotent: Referring to the ability of a cell or embryo part to give rise to all adult structures. This capacity usually is restricted progressively during development.

trait: Any detectable phenotypic variation of a particular inherited character.

transcription: Transfer of genetic information from DNA by the synthesis of an RNA molecule copied from a DNA template.

transfection: The uptake and expression of foreign DNA by cultured eukaryotic cells.

transformation: In higher eukaryotes, the conversion of cultured cells to a malignant phenotype. In prokaryotes, the natural or induced uptake and expression of a foreign DNA sequence.

transforming growth factor alpha (TGF-α): Functional and structural analogue of epidermal growth factor. Induces the growth of epithelial cells as well as fibroblasts and keratinocytes. May be involved in tumor-associated neovascularization.

transforming growth factor beta (TGF-β): A cytokine that regulates many of the biologic processes essential for embryonic development and tissue homeostasis and which therefore plays a role in the healing of a tissue and carcinogenesis. The effects of TGF-β may differ, depending on the tissue involved. For instance, TGF-β inhibits proliferation of epithelial cells, but stimulates that of fibroblasts.

transforming oncogene: A gene that upon transfection converts a previously immortalized cell to the malignant phenotype.

transgenic: A vertebrate organism in which a foreign DNA gene (a transgene) is stably incorporated into its genome early in embryonic development. The transgene is present in both somatic and germ cells, is expressed in one or more tissues, and is inherited by offspring in a mendelian fashion.

transient hypoxia: Low oxygen concentrations associated with the transient closing and opening of blood vessels. Sometimes called acute or cyclic hypoxia.

translation: The process of converting the genetic information of mRNA on ribosomes into polypeptides.

translocation: The movement or reciprocal exchange of large chromosomal segments, typically between two different chromosomes.

trisomy: The condition in which a cell or organism possesses two copies of each chromosome, except for one, which is present in three copies. The general form for trisomy is therefore 2n + 1.

tritium (H, T): A radioactive isotope of hydrogen with two neutrons and one proton in the nucleus. It is human-made and heavier than deuterium (heavy hydrogen). Tritium is used in industrial thickness gauges and as a label in chemical and biologic experiments. Its nucleus is a triton.

tRNA (transfer RNA): A small RNA molecule that contains a three-base segment (anticodon) that recognizes a codon in mRNA, a binding site for a specific amino acid, and recognition sites for interaction with the ribosome and the enzyme that links it to its specific amino acid.

tumor: An abnormal growth of cells or tissues. Tumors may be benign (noncancerous) or malignant (cancerous).

tumor bed effect (TBE): Slower rate of tumor growth after irradiation, resulting from stromal injury in the irradiated “vascular bed.”

tumor cord: Sleeve of viable tumor growing around a blood capillary.

tumor necrosis factor (TNF): Two proteins, TNF-α and TNF-β, involved in immune response control and inflammation. Induced by cytokines, ultraviolet radiations, and other agents.
**UNSCCER:** United Nations Scientific Commission on Effects of Atomic Radiation.

**uranium (U):** A radioactive element with atomic number 92 and, as found in natural ores, an average atomic weight of approximately 238. The two principal natural isotopes are $^{235}\text{U}$ (0.7% of natural uranium), which is fissionable, and $^{238}\text{U}$ (99.3% of natural uranium), which is fertile. Natural uranium also includes a minute amount of $^{234}\text{U}$.

**uranium series (sequence):** The series of nuclides resulting from the radioactive decay of the uranium isotope $^{238}\text{U}$. The end product of the series is the lead isotope $^{206}\text{Pb}$.

**variance:** A statistical measure of the variation of values from a central value, calculated as the square of the standard deviation.

**vascular targeted therapies:** Treatments designed to specifically target tumor vasculature, including angiogenesis inhibitors.

**vector:** An autonomously replicating DNA molecule into which foreign DNA fragments are inserted and then propagated in a host cell.

**viability:** The measure of the number of individuals in a given phenotypic class that survive, relative to another class (usually wild-type).

**viral oncogene:** A viral gene that contributes to malignancies in vertebrate hosts.

**virulent phage:** A bacteriophage that infects and lyses the host bacterial cell.

**virus:** An infectious particle that is composed of a protein capsule and a nucleic acid core and that is dependent on a host organism for replication.

**volume effect:** Dependence of radiation damage to normal tissues on the volume of tissue irradiated.

**volume-doubling time:** Time taken for a tumor to double in volume.

**waste, radioactive:** Equipment and materials (from nuclear operations) that are radioactive and have no further use. Wastes are generally classified as high level: radioactivity concentrations of hundreds to thousands of curies per gallon or cubic foot; low level: in the range of 1 μCi (microcurie) per gallon or cubic foot; or intermediate: between these extremes.

**wavelength:** Distance between similar points on a sine wave; length of one cycle.

**Western blot:** Similar to a Southern blot (for DNA) or a northern blot (for RNA), except that protein is used.

**white blood cells:** The blood cells that fight infection.

**whole-body counter:** A device used to identify and measure the radioactivity in the body (body burden) of humans and animals. It uses heavy shielding (to keep out background radiation), ultrasensitive scintillation detectors, and electronic equipment.

**wild type:** The most commonly observed phenotype or genotype, designated as the norm or standard.

**X chromosome:** The female sex chromosome.

**Xenografts:** Transplants between species; usually applied to the transplantation of human tumors into immune-deficient mice and rats.

**xerostomia:** Dryness of the mouth caused by malfunctioning salivary glands.

**X-linkage:** See sex linkage.

**x-ray:** A penetrating form of electromagnetic radiation emitted either if the inner orbital electrons of an excited atom return to their normal state (these are characteristic x-rays) or if a metal target is bombarded with high-speed electrons. X-rays are always extranuclear in origin.

**x-ray crystallography:** A technique to determine the three-dimensional structure of molecules through diffraction patterns produced by x-ray scattering by crystals of the molecule under study.

**yeast artificial chromosome (YAC):** A vector that is used to clone DNA fragments of up to 400,000 bp and that contains the minimum chromosomal sequences needed to replicate in yeast.

**Z:** The symbol for atomic number; the number of protons in the nucleus.

**zinc fingers:** A structural motif of DNA-binding proteins in which fingerlike loops of amino acids are stabilized by interactions with zinc atoms.

**zygote:** The diploid cell produced by the fusion of haploid gametic nuclei.
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