SUBCHAPTER b

PART 310
GENERAL PROVISIONS FOR RADIATION PROTECTION

Section
310.10 Scope
310.15 Incorporations by Reference
310.20 Definitions
310.30 Exemptions
310.40 Records
310.50 Inspections
310.60 Tests
310.70 Additional Requirements
310.74 Cost Assessment
310.75 Emergency Response Cost Recovery
310.78 Deliberate Misconduct
310.80 Violations (Repealed)
310.81 Policy for Assessment of Civil Penalties
310.82 Procedures for Assessment of Civil Penalties (Repealed)
310.90 Impounding
310.100 Prohibited Uses
310.110 Communications
310.120 Plans and Specifications
310.130 The International System of Units (SI) (Repealed)
310.140 Units of Exposure and Radiation Dose
310.150 Units of Activity

310.APPENDIX A Transport Grouping of Radionuclides (Repealed)
310.APPENDIX B Tests for Special Form Licensed Material (Repealed)
310.APPENDIX C Penalty Assessment Worksheet (Repealed)

AUTHORITY: Implementing and authorized by the Radiation Protection Act of 1990 [420 ILCS 40].


Section 310.10 Scope

Except as otherwise specifically provided, this Part applies to all persons who receive, possess, use, transfer, own or acquire any source of radiation within the State of Illinois; provided, however, that nothing in this Part or 32 Ill. Adm. Code 320, 326, 330, 331, 332, 335, 340, 341, 346, 350, 351, 400, 401, 405 or 601 shall apply to any person to the extent such person is subject to regulation by the U.S. Nuclear Regulatory Commission (NRC).

AGENCY NOTE: Regulation by the State of source material, byproduct material and special nuclear material in quantities not sufficient to form a critical mass is subject to the provisions of an agreement between the State and the NRC and to 10 CFR 150 of NRC's regulations.

(Source: Amended at 35 Ill. Reg. 2908, effective February 7, 2011)

Section 310.15 Incorporations by Reference

All rules, standards and guidelines of agencies of the United States or nationally recognized organizations or associations that are incorporated by reference in this Part are incorporated as of the date specified in the reference and do not include any later amendments or editions. Copies of rules, standards and guidelines that have been incorporated by reference are available for public inspection and copying at the Illinois Emergency Management Agency, 1035 Outer Park Drive, Springfield, Illinois.

(Source: Amended at 29 Ill. Reg. 20748, effective December 16, 2005)

Section 310.20 Definitions

As used in 32 Ill. Adm. Code: Chapter II, Subchapters b and d, the following terms have the definitions set forth in this Section. Additional definitions used only in a certain Part will be found in that Part.

"Absorbed dose" means the energy imparted by ionizing radiation per unit mass of irradiated material. The units of absorbed dose are the gray (Gy) and the rad.

"Accelerator" or "particle accelerator" means any machine capable of accelerating
electrons, protons, deuterons or other charged particles in a vacuum and of discharging the resultant particulate or other radiation into a medium at energies usually in excess of 1 million electron volts (MeV).

"Accelerator-produced material" means any material made radioactive by a particle accelerator.

"Act" means the Radiation Protection Act of 1990 [420 ILCS 40].

"Activity" means the rate of disintegration (transformation) or decay of radioactive material. The units of activity are the bequerel (Bq) and the curie (Ci).

"Adult" means an individual 18 or more years of age.


"Agreement State" means any state with which the U.S. Nuclear Regulatory Commission or the U.S. Atomic Energy Commission has entered into an effective agreement under subsection 274b of the Atomic Energy Act of 1954, as amended (42 USC 2021(b) et seq.).

"Airborne radioactive material" means any radioactive material dispersed in the air in the form of dusts, fumes, particulates, mists, vapors or gases.

"Airborne radioactivity area" means any room, enclosure or operating area in which airborne radioactive material, composed wholly or partly of licensed material, exists in concentrations:

in excess of the derived air concentrations (DACs) specified in appendix B to 10 CFR 20, published at 72 Fed. Reg. 55922, October 1, 2007, exclusive of subsequent amendments or editions; or

to such a degree that an individual present in the area without respiratory protective equipment could exceed, during the hours an individual is present in a week, an intake of 0.6 percent of the annual limit on intake (ALI) or 12 DAC-hours.

"Annually" means at intervals not to exceed 1 year or once per year, at about the same time each year (plus or minus 1 month).

"As low as is reasonably achievable" or "ALARA" means making every
reasonable effort to maintain exposures to radiation as far below the dose limits in
32 Ill. Adm. Code: Chapter II, Subchapters b and d as is practical consistent with
the purpose for which the licensed or registered activity is undertaken, taking into
account the state of technology, the economics of improvements in relation to the
state of technology, the economics of improvements in relation to benefits to the
public health and safety and other societal and socioeconomic considerations, and
in relation to utilization of nuclear energy and licensed or registered sources of
radiation in the public interest.

"Background radiation" means radiation from cosmic sources, naturally occurring
radioactive materials, including radon (except as a decay product of source or
special nuclear material) and global fallout as it exists in the environment from
the testing of nuclear explosive devices, or from past nuclear accidents such as
Chernobyl that contribute to background radiation and are not under the control of
the licensee. Background radiation does not include radiation from radioactive
materials regulated by the Agency.

"Becquerel" or "Bq" means the SI unit of activity. One becquerel (Bq) is equal to
1 disintegration (transformation) per second (dps or tps).

"Bioassay" or "radiobioassay" means the determination of kinds, quantities or
concentrations and, in some cases, the locations of radioactive material in the
human body, whether by direct measurement (in vivo counting) or by analysis
and evaluation of materials excreted or removed from the human body.

"Brachytherapy" means a method of radiation therapy in which sources are used
to deliver a radiation dose at a distance of up to a few centimeters, by surface,
intracavitary, intraluminal or interstitial application.

"Brachytherapy source" means a radioactive source, a manufacturer-assembled
source train or a combination of these sources that is designed to deliver a
therapeutic dose within a distance of a few centimeters.

"By-product material" means:

any radioactive material (except special nuclear material) yielded in or
made radioactive by exposure to radiation incident to the process of
producing or utilizing special nuclear material;

the tailings or wastes produced by the extraction or concentration of
uranium or thorium from any ore processed primarily for its source
material content, including discrete surface wastes resulting from
underground solution extraction processes but not including underground ore bodies depleted by such solution extraction processes;

any discrete source of radium-226 that is produced, extracted, or converted after extraction, before, on, or after August 8, 2005, for use for a commercial, medical, or research activity;

any material that has been made radioactive by use of a particle accelerator and is produced, extracted, or converted after extraction before, on, or after August 8, 2005, for use for a commercial, medical, or research activity; and

any discrete source of naturally occurring radioactive material, other than source material, that is extracted or converted after extraction for use in commercial, medical, or research activity before, on, or after August 8, 2005, and which the U.S. Nuclear Regulatory Commission, in consultation with the Administrator of the Environmental Protection Agency, the Secretary of Energy, the Secretary of Homeland Security, and the head of any other appropriate federal agency, determines would pose a threat to the public health and safety or the common defense and security similar to the threat posed by a discrete source or radium-226. [420 ILCS 40/4(a-5)]

"Calendar quarter" means not less than 12 consecutive weeks nor more than 14 consecutive weeks. The first calendar quarter of each year shall begin in January and subsequent calendar quarters shall be so arranged such that no day is included in more than one calendar quarter and no day in any one year is omitted from inclusion within a calendar quarter. No licensee or registrant shall change the method observed by him for determining calendar quarters except at the beginning of a year.

"Calibration" means the determination of:

the response or reading of an instrument relative to a series of known radiation values over the range of the instrument; or

the strength of a source of radiation relative to a standard.


"Chelating agent" means amine polycarboxylic acids (e.g., EDTA, DTPA), hydroxy-carboxylic acids, and polycarboxylic acids (e.g., citric acid, carbolic acid, and glucinic acid).
"Collective dose" means the sum of the individual doses received in a given period of time by a specified population from exposure to a specified source of radiation.

"Commencement of construction" means, except as specified in 32 Ill. Adm. Code 601.20, taking any action defined as "construction" or any other activity at the site of a facility subject to this Part that has a reasonable nexus to radiological health and safety.

"Committed dose equivalent" or "$H_{T,50}$" means the dose equivalent to organs or tissues of reference (T) that will be received from an intake of radioactive material by an individual during the 50-year period following the intake.

"Committed effective dose equivalent" or $H_{E,50}$ means the sum of the products of the weighting factors applicable to each of the body organs or tissues that are irradiated and the committed dose equivalent to each of these organs or tissues ($H_{E,50} = \sum w_T H_{T,50}$).

"Construction" means the installation of foundations or in-place assembly, erection, fabrication or testing for any structure, system, or component of a facility or activity subject to this Part that is related to radiological safety or security. The term "construction" does not include:

changes for temporary use of the land for public recreational purposes;

site exploration, including necessary borings to determine foundation conditions or other preconstruction monitoring to establish background information related to the suitability of the site, the environmental impacts of construction or operation, or the protection of environmental values;

preparation of the site for construction of the facility, including clearing of the site, grading, installation of drainage, erosion control and other environmental mitigation measures, and construction of temporary roads and borrow areas;

erection of fences and other access control measures that are not related to the safe use of, or security of, radiological materials subject to this Part;

excavation;
erection of support buildings (e.g., construction equipment storage sheds, warehouse and shop facilities, utilities, concrete mixing plants, docking and unloading facilities, and office buildings) for use in connection with the construction of the facility;

building of service facilities (e.g., paved roads, parking lots, railroad spurs, exterior utility and lighting systems, potable water systems, sanitary sewerage treatment facilities and transmission lines);

procurement or fabrication of components or portions of the proposed facility occurring at other than the final, in-place location at the facility; or

taking any other action that has no reasonable nexus to radiological health and safety.

"Critical group" means the group of individuals reasonably expected to receive the greatest exposure to residual radioactivity for any applicable set of circumstances.

"Curie" or "Ci" means a unit of quantity of radioactivity. One curie is that quantity of radioactive material that decays at the rate of $3.7 \times 10^{10}$ disintegrations (transformations) per second (dps or tps).

"Decommission" means to remove a facility or site safely from service and reduce residual radioactivity to a level that permits release of property for unrestricted use and termination of the license.

"Declared pregnant woman" means any woman who has voluntarily informed the licensee or registrant, in writing, of her pregnancy and the estimated date of conception. The declaration remains in effect until the declared pregnant woman withdraws the declaration in writing or is no longer pregnant.

"Dedicated check source" means a radioactive source that is used to assure the constant operation of a radiation detection or measurement device over several months or years.

"Deep dose equivalent" or "Hd" means the dose equivalent at a tissue depth of 1 centimeter (1000 milligrams per square centimeter) from external whole-body exposure.

"Densitometer" means a device that is used to provide a quantitative measurement of the optical density of x-ray film to determine the response of the film to
exposure and development.

"Depleted uranium" means the source material uranium in which the isotope uranium-235 is less than 0.711 weight percent of the total uranium present. Depleted uranium does not include special nuclear material.

"Director" means the Director of the Illinois Emergency Management Agency.

"Discrete source" means a radionuclide that has been processed so that its concentration within a material has been purposely increased for use for commercial, medical or research activities.

"Distinguishable from background" means the detectable radioactivity is statistically different from background in the vicinity of the site, or, in the case of structures, in similar materials using adequate measurement technology, survey and statistical techniques.

"Dose" or "radiation dose" means either absorbed dose, dose equivalent, effective dose equivalent, committed dose equivalent, committed effective dose equivalent, total organ dose equivalent or total effective dose equivalent.

"Dose equivalent" or "H_T" means the product of the absorbed dose in tissue, quality factor and all other necessary modifying factors (e.g., a distribution factor for nonuniform deposition) at the location of interest. The units of dose equivalent are the sievert (Sv) and the rem.

"Dose limits" or "limits" means the permissible upper bounds of radiation doses established by, or in accordance with, 32 Ill. Adm. Code: Chapter II, Subchapters b and d.

"Dosimetry processor" means an individual or an organization that processes and evaluates individual monitoring devices in order to determine the radiation dose delivered to those devices. Dosimetry processing not only includes physical or chemical processing of dosimetry in a laboratory setting to extract absorbed dose information from a storage medium as is the case with film or thermoluminescent dosimetry (TLD), but also includes the process of digitally extracting absorbed dose information locally from storage medium using approved algorithms and computer applications as is the case with direct ion storage (DIS) technology.

"Effective dose equivalent" or "H_E" means the sum of the products of the dose equivalent to each organ or tissue (H_T) and the weighting factor (W_T) applicable to each of the body organs or tissues that are irradiated (H_E = ΣW_T H_T).
"Embryo/fetus" means the developing human organism from conception until the time of birth.

"Entrance or access point" means any opening through which an individual or extremity of an individual could gain access to radiation areas or to licensed radioactive materials. This includes entry or exit portals of sufficient size to permit human entry, irrespective of their intended use.

"Exposure" means:

the quotient of \( dQ \) divided by \( dm \) where \( dQ \) is the absolute value of the total charge of the ions of one sign produced in air when all the electrons (negatrons and positrons) liberated by photons in a volume element of air having mass \( dm \) are completely stopped in air. (See Section 310.140 for SI unit coulomb per kilogram (C/kg) and the special unit roentgen (R).); or

irradiation by ionizing radiation or radioactive material.

AGENCY NOTE: The context makes clear which is the appropriate definition.

"Exposure rate" means the exposure per unit of time, such as roentgen per minute (R/min) and milliroentgen per hour (mR/h).

"External dose" means that portion of the dose equivalent received from any source of radiation outside the body.

"Extremity" means a hand, elbow, arm below the elbow, foot, knee and leg below the knee.

"Former U.S. Atomic Energy Commission (AEC) or U.S. Nuclear Regulatory Commission (NRC) licensed facilities" means nuclear reactors, nuclear fuel reprocessing plants, uranium enrichment plants, or critical mass experimental facilities where AEC or NRC licenses have been terminated.

"Gray" or "Gy" means the SI unit of absorbed dose. One gray is equal to an absorbed dose of 1 joule per kilogram (J/kg) (100 rad).

"Healing arts" means the art or science or group of arts or sciences dealing with the prevention and cure or alleviation of human ailments, diseases or infirmities, and has the same meaning as "medicine" when the latter term is used in its
comprehensive sense.

"High radiation area" means any area, accessible to individuals, in which radiation levels from radiation sources external to the body could result in an individual receiving a dose equivalent in excess of 0.1 rem (1 mSv) in 1 hour at 30 centimeters from the radiation source or 30 centimeters from any surface that the radiation penetrates.

"Human use" means the internal or external administration of radiation or radioactive materials to human beings.

"Individual" means any human being.

"Individual monitoring" means the assessment of:

Dose equivalent by the use of individual monitoring devices or by the use of survey data; or

Committed effective dose equivalent by bioassay or by determination of the time-weighted air concentrations to which an individual has been exposed (i.e., DAC-hours). (For the definition of DAC-hours, see 32 Ill. Adm. Code 340.30.)

"Individual monitoring devices" (personnel dosimeter or dosimeter) means devices designed to be worn by a single individual for the assessment of dose equivalent. Examples of individual monitoring devices are film badges, thermoluminescence dosimeters (TLDs), optically stimulated luminescence dosimeters (OSLs), pocket ionization chambers, personal air sampling devices and electronic dosimeters (e.g., silicon diode dosimeters).

"Inspection" means an official examination or observation including, but not limited to, tests, surveys, and monitoring to determine compliance with rules, regulations, orders, requirements, and conditions of the Agency.

"Interlock" means a device arranged or connected such that the occurrence of an event or condition is required before a second event or condition can occur or continue to occur.

"Internal dose" means that portion of the dose equivalent received from radioactive material taken into the body.

"License" means any license issued by the Agency in accordance with 32 Ill.
Adm. Code: Chapter II, Subchapters b and d.

"Licensed material" means radioactive material received, possessed, used, transferred or disposed of under a general or specific license issued by the Agency.

"Licensee" means any person who is licensed by the Agency in accordance with 32 Ill. Adm. Code: Chapter II, Subchapters b and d.

"Lost or missing source of radiation" means any licensed or registered source of radiation whose location is unknown. This definition includes, but is not limited to, radioactive material that has been shipped but has not reached its planned destination and whose location cannot be readily traced in the transportation system.

"Major processor" means a person, other than medical programs, universities, industrial radiography services, or wireline service operations, who is licensed to process, handle, or manufacture radioactive material as unsealed sources in quantities exceeding the quantities specified in appendix C to 10 CFR 20, published at 60 Fed. Reg. 20186, April 25, 1995, exclusive of subsequent amendments or editions, by a factor of at least $10^9$, or radioactive material as sealed sources in quantities exceeding the quantities specified in appendix C to 10 CFR 20 by a factor of at least $10^{10}$.

"Member of the public" means any individual, except an individual who is performing assigned duties for the licensee or registrant involving exposure to sources of radiation.

"Minor" means an individual less than 18 years of age.

"Monitoring" or "radiation monitoring" or "radiation protection monitoring" means the measurement of radiation, radioactive material concentrations, surface area activities or quantities of radioactive material and the use of the results of these measurements to evaluate potential exposures and doses.

"NARM" means any naturally occurring or accelerator-produced radioactive material. It does not include byproduct, source or special nuclear material.

"Natural radioactivity" means radioactivity of naturally occurring nuclides.

"Nuclear Regulatory Commission" or "NRC" means the U.S. Nuclear Regulatory Commission or its duly authorized representatives.
"Occupational dose" means the dose received by an individual in the course of employment in which the individual's assigned duties for the licensee or registrant involve exposure to sources of radiation. Occupational dose does not include dose received from background radiation, from any medical administration the individual has received, from exposure to individuals administered radioactive material and released as authorized by the Agency, from voluntary participation in medical research programs, or as a member of the public.

"Operator" means an individual, group of individuals, partnership, firm, corporation, association, or other entity conducting the business or activities carried on within a radiation installation. [420 ILCS 40/4(d-7)]

"Package" means the packaging, together with its radioactive contents, as presented for transport.

"Packaging" means the assembly of components necessary to ensure compliance with the packaging requirements of 32 Ill. Adm. Code 341. It may consist of one or more receptacles, absorbent materials, spacing structures, thermal insulation, radiation shielding and devices for cooling or absorbing mechanical shocks. The vehicle, tie down system and auxiliary equipment may be designated as part of the packaging.

"Particle accelerator" means any machine capable of accelerating electrons, protons, deuterons or other charged particles in a vacuum and of discharging the resultant particulate or other radiation into a medium at energies usually in excess of 1 million electron volts (MeV). For purposes of this definition, "accelerator" is an equivalent term.

"Person" means any individual, corporation, partnership, firm, association, trust, estate, public or private institution, group, agency, political subdivision of this State, any other State or political subdivision or agency thereof, and any legal successor, representative, agent, or agency of the foregoing, other than the United States Nuclear Regulatory Commission, or any successor thereto, and other than federal government agencies licensed by the United States Nuclear Regulatory Commission, or any successor thereto. "Person" also includes a federal entity (and its contractors) if the federal entity agrees to be regulated by the State or as otherwise allowed under federal law. [420 ILCS 40/4(e)]

"Personnel monitoring equipment" (see "Individual monitoring devices").

"PET" means positron emission tomography.
"Pharmacist" means an individual licensed by the State pursuant to the Pharmacy Practice Act [225 ILCS 85] to compound and dispense drugs, prescriptions, and poisons.

"Physician" means an individual licensed to practice a treatment of human ailments by virtue of the Medical Practice Act of 1987 [225 ILCS 60], the Illinois Dental Practice Act [225 ILCS 25] or the Podiatric Medical Practice Act of 1987 [225 ILCS 100], who may use radiation for therapeutic, diagnostic or other medical purposes within the limits of the individual's licensure.

"Positron emission tomography radionuclide production facility" means a facility operating a particle accelerator for the purpose of producing PET radionuclides.

"Protective apron" means any apron made of radiation attenuating materials, at least 0.25 millimeter lead equivalent, that may be used to reduce exposure to radiation.

"Qualified engineering expert" means any person qualified under the Illinois Architecture Practice Act of 1989 [225 ILCS 305], the Structural Engineering Licensing Act of 1989 [225 ILCS 340] and/or any required combination thereof.

"Quality factor" or "Q" means the modifying factor (listed in Section 310.140, Tables 1 and 2) that is used to derive dose equivalent from absorbed dose.

"Quarterly" means at intervals not to exceed 3 months.

"Rad" means the special unit of absorbed dose. One rad is equal to an absorbed dose of 100 ergs per gram or 0.01 joule per kilogram (J/kg) (0.01 Gy).

"Radiation" or "ionizing radiation" means gamma rays and x-rays, alpha and beta particles, high-speed electrons, neutrons, protons, and other nuclear particles, or electromagnetic radiations capable of producing ions directly or indirectly in their passage through matter; but does not include sound or radio waves, or visible infrared or ultraviolet light. [420 ILCS 40/4(f)]

"Radiation area" means an area, accessible to individuals, in which radiation levels could result in an individual receiving a dose equivalent in excess of 0.05 mSv (0.005 rem) in 1 hour at 30 centimeters from the source of radiation or from any surface that the radiation penetrates.

"Radiation dose" (see "Dose").
"Radiation emergency" means the uncontrolled release of radioactive material from a radiation installation which poses a potential threat to the public health, welfare and safety. [420 ILCS 40/4(f-5)]

"Radiation Installation" is any location or facility where radiation machines are used or where radioactive material is produced, transported, stored, disposed or used for any purpose [420 ILCS 40/4(g)], except when the radioactive materials or facility are subject to regulation by the NRC.

"Radiation machine" means any device that produces radiation when in use [420 ILCS 40/4(h)], except those that produce radiation only from radioactive materials.

"Radiation safety officer" means an individual who has the knowledge and responsibility to apply appropriate radiation protection regulations and has been assigned that responsibility by the licensee or registrant.

"Radioactive material" means any solid, liquid, or gaseous substance which emits radiation spontaneously. [420 ILCS 40/4(i)] It includes material defined as "byproduct material" in the Act.

"Radioactivity" means the disintegration (transformation) of unstable atomic nuclei by the emission of radiation.

"Radiobioassay" (see "Bioassay").

"Registrant" means any person who is registered with the Agency and is legally obligated to register with the Agency pursuant to the Radiation Protection Act of 1990 [420 ILCS 40] and 32 Ill. Adm. Code 320.10.

"Registration" means registration with the Agency in accordance with 32 Ill. Adm. Code 320.10.

"Regulations of the U.S. Department of Transportation" or "regulations of USDOT" means the regulations in 49 CFR 100-189, revised November 14, 2014, exclusive of subsequent amendments or editions.

"Rem" means the special unit of any of the quantities expressed as dose equivalent. The dose equivalent in rem is equal to the absorbed dose in rad multiplied by the quality factor (1 rem = 0.01 Sv).
"Research and development" means:

theoretical analysis, exploration, or experimentation; or

the extension of investigative findings and theories of a scientific or technical nature into practical application for experimental and demonstration purposes, including the experimental production and testing of models, devices, equipment, materials, and processes.

Research and development does not include the internal or external administration of radiation or radioactive material to human beings.

"Residual radioactivity" means radioactivity in structures, materials, soils, groundwater and other media at a site resulting from activities under the licensee's control. This includes radioactivity from all licensed and unlicensed sources used by the licensee, but excludes background radiation. It also includes radioactive materials remaining at the site as a result of routine or accidental releases of radioactive material at the site and previous burials at the site, even if those burials were made in accordance with the provisions of 32 Ill. Adm. Code 340 or the equivalent provisions of 10 CFR 20.

"Restricted area" means any area access to which is limited by the licensee or registrant for purposes of protecting individuals against undue risks from exposure to sources of radiation. Restricted area shall not include areas used for residential quarters, but separate rooms in a residential building may be set apart as a restricted area.

"Roentgen" means the special unit of exposure. One roentgen (R) equals 2.58 x 10^-4 coulombs per kilogram (C/kg). (See "Exposure" and Section 310.140.)

"Sealed source" means any device containing radioactive material to be used as a source of radiation which has been constructed in such a manner as to prevent the escape of any radioactive material.

"Sealed source and device registry" means the national registry that contains all the registration certificates generated by the Agency, U.S. Nuclear Regulatory Commission or an Agreement State that summarize the radiation safety information for the sealed sources and devices and describe the licensing and use conditions approved for the product.

"Sensitometer" means a device that is used to test the setup and stability of film processing procedures and equipment by providing a standard pattern of light
exposure of x-ray film.

"Shallow dose equivalent" or "Hs", which applies to the external exposure of the skin or an extremity, means the dose equivalent at a tissue depth of 0.007 centimeter (7 milligrams per square centimeter).

"SI" means the abbreviation for the International System of Units.

"Sievert" or "Sv" means the SI unit of any of the quantities expressed as dose equivalent. The dose equivalent in sievert is equal to the absorbed dose in grays multiplied by the quality factor (1 Sv = 100 rem).

"Source material" means:

- uranium or thorium, or any combination thereof, in any physical or chemical form; or
- ores that contain by weight one-twentieth of one percent (0.05 percent) or more of uranium, thorium or any combination thereof.

Source material does not include special nuclear material.

"Source of radiation" means any radioactive material or any device or equipment emitting, or capable of producing, radiation.

"Special form radioactive material" means radioactive material that satisfies the following conditions:

- It is either a single solid piece or is contained in a sealed capsule that can be opened only by destroying the capsule;
- The piece or capsule has at least one dimension not less than 5 millimeters (0.197 inch); and
- It satisfies the test requirements specified in 10 CFR 71.75 and 71.77, published at 60 Fed. Reg. 50264, September 28, 1995, exclusive of subsequent amendments or editions, except that special form radioactive material designed or constructed prior to July 1, 1985 need only meet the requirements of 10 CFR 71.75 and 71.77 in effect on June 30, 1983.

"Special nuclear material" means:
plutonium, uranium 233, uranium enriched in the isotope 233 or in the isotope 235 and any other material which the Agency declares by order to be special nuclear material after the United States Nuclear Regulatory Commission, or any successor thereto, has determined the material to be such, but does not include source material; or

any material artificially enriched by any of the foregoing, but does not include source material.  [420 ILCS 40/4(1)]

"Special nuclear material in quantities not sufficient to form a critical mass" means uranium enriched in the isotope U-235 in quantities not exceeding 350 grams of contained U-235; U-233 in quantities not exceeding 200 grams; plutonium in quantities not exceeding 200 grams; or any combination of them, except source material, in accordance with the following formula: For each kind of special nuclear material, determine the ratio between the quantity of that special nuclear material and the quantity specified above for the same kind of special nuclear material. The sum of such ratios for all of the kinds of special nuclear material in combination shall not exceed one. For example, the following quantities in combination would not exceed the limitation and are within the formula:

\[
\frac{175 \text{ (grams contained U-235)}}{350} + \frac{50 \text{ (grams U-233)}}{200} + \frac{50 \text{ (grams Pu)}}{200} = 1
\]

"Survey" means an evaluation of the radiological conditions and potential hazards incident to the production, use, transfer, release, disposal, or presence of sources of radiation. Such an evaluation includes, but is not limited to, measurements or calculations of levels of radiation, or concentrations or quantities of radioactive material present.

"Test" means the process of verifying compliance with an applicable regulation.

"Total effective dose equivalent" or "TEDE" means the sum of the deep dose equivalent for external exposures and the committed effective dose equivalent for internal exposures.

"Total organ dose equivalent" or "TODE" means the sum of the deep dose equivalent and the committed dose equivalent to the organ receiving the highest dose as described in 32 Ill. Adm. Code 340.1160(a)(6).

"Unrefined and unprocessed ore" means ore in its natural form prior to any processing, such as grinding, roasting or beneficiating, or any refining.
Processing does not include sieving or encapsulation of ore or preparation of samples for laboratory analysis.

"Unrestricted area" means any area access to which is not controlled by the licensee or registrant for purposes of protection of individuals from exposure to radiation and radioactive material, and any area used for residential quarters.

AGENCY NOTE: Licensees or registrants may control access to certain areas for purposes other than radiation protection, but such action does not affect whether the areas are unrestricted areas as defined in this Part.

"Uranium fuel cycle" means the operations of milling of uranium ore, chemical conversion of uranium, isotopic enrichment of uranium, fabrication of uranium fuel, generation of electricity by a light-water-cooled nuclear power plant using uranium fuel and reprocessing of spent uranium fuel to the extent that these activities directly support the production of electrical power for public use. Uranium fuel cycle does not include mining operations, operations at waste disposal sites, transportation of radioactive material in support of these operations and the reuse of recovered non-uranium special nuclear and byproduct materials from the cycle.

"U.S. Department of Energy" means the agency created by the Department of Energy Organization Act (established by P.L. 95-91, 91 Stat. 565, 42 USC 7101 et seq.), to the extent that the Department of Energy, or its duly authorized representatives, exercises functions formerly vested in the U.S. Atomic Energy Commission, its Chairman, members, officers and components and transferred to the U.S. Energy Research and Development Administration and to the Administrator thereof pursuant to sections 104(b), (c) and (d) of the Energy Reorganization Act of 1974 (P.L. 93-438, 88 Stat. 1233 at 1237, 42 USC 5814) and retransferred to the Secretary of Energy pursuant to section 301(a) of the Department of Energy Organization Act (P.L. 95-91, 91 Stat. 565 at 577-578, 42 USC 7151).

"Very high radiation area" means an area, accessible to individuals, in which radiation levels from radiation sources external to the body could result in an individual receiving an absorbed dose in excess of 500 rads (5 grays) in 1 hour at 1 meter from a radiation source or 1 meter from any surface that the radiation penetrates.

AGENCY NOTE: For very high doses received at high dose rates, units of absorbed dose (e.g., gray and rad) are appropriate rather than units of dose equivalent (e.g., sievert and rem).

"Waste" means those low-level radioactive wastes containing source, special
nuclear or byproduct material that are acceptable for disposal in a land disposal facility. For the purposes of this definition, low-level radioactive waste means radioactive waste not classified as high-level radioactive waste, transuranic waste, spent nuclear fuel, or byproduct material as defined in Section 4(a-5)(2) of the Act.

"Waste handling licensee" means a person licensed by the Agency, the U.S. Nuclear Regulatory Commission or an Agreement State to receive radioactive wastes for storage or treatment, or both storage and treatment, prior to disposal as well as any person licensed to receive radioactive waste for disposal away from the point of generation.

"Week" means 7 consecutive days starting on Sunday.

"Whole body" means, for purposes of external exposure, head, trunk (including male gonads), arms above the elbow or legs above the knee.

"Worker" means any individual engaged in work under a license or registration issued by the Agency and controlled by a licensee or registrant, but does not include the licensee or registrant.

"Working level" or "WL" means any combination of short-lived radon daughters in 1 liter of air that will result in the ultimate emission of $1.3 \times 10^5$ MeV of potential alpha particle energy. The short-lived radon daughters are for:

- radon-222: polonium-218, lead-214, bismuth-214 and polonium-214; and
- radon-220: polonium-216, lead-212, bismuth-212 and polonium-212.

"Working level month" or "WLM" means an exposure to 1 working level (WL) for 170 hours. (2,000 working hours per year divided by 12 months per year is approximately equal to 170 hours per month.)

"Year" means the period of time beginning in January used to determine compliance with the provisions of 32 Ill. Adm. Code: Chapter II, Subchapters b and d. The licensee or registrant may change the starting date of the year used to determine compliance by the licensee or registrant provided that the decision to make the change is made not later than December 31 of the previous year. If a licensee or registrant changes a year, the licensee or registrant shall assure that no day is omitted or duplicated in consecutive years.

(Source: Amended at 45 Ill. Reg. 9911, effective July 22, 2021)
Section 310.30  Exemptions

a) General Provisions − The Agency may, upon application or upon its own initiative, grant such exemptions or exceptions from the requirements of 32 Ill. Adm. Code: Chapter II, Subchapters b and d as it determines are authorized by law and will not result in undue hazard to public health and safety or property.

b) U. S. Department of Energy Contractors and U. S. Nuclear Regulatory Commission Contractors − Any U. S. Department of Energy contractor or subcontractor and any U. S. Nuclear Regulatory Commission contractor or subcontractor of the following categories operating within this State is exempt from 32 Ill. Adm. Code: Chapter II, Subchapters b and d to the extent that such contractor or subcontractor under contract receives, possesses, uses, transfers or acquires sources of radiation:

1) Prime contractors performing work for the Department of Energy at U. S. Government-owned or controlled sites, including the transportation of sources of radiation to or from such sites and the performance of contract services during temporary interruptions of such transportation;

2) Prime contractors of the Department of Energy performing research in, or development, manufacture, storage, testing or transportation of, atomic weapons or components thereof;

3) Prime contractors of the Department of Energy using or operating nuclear reactors or other nuclear devices in a United States Government-owned vehicle or vessel; and

4) Any other prime contractor or subcontractor of the Department of Energy or of the Nuclear Regulatory Commission when the State and the Nuclear Regulatory Commission jointly determine:

   A) that, under the terms of the contract or subcontract, there is adequate assurance that the work thereunder can be accomplished without undue risk to the public health and safety; and

   B) that the exemption of such contractor or subcontractor is otherwise appropriate.

c) Federal governmental agencies or contractors of such agencies providing training for State and local governmental entities within this State are exempt from the fee
requirements in 32 Ill. Adm. Code 331 provided they meet the notification

(Source: Amended at 45 Ill. Reg. 9911, effective July 22, 2021)

Section 310.40  Records

Each licensee and registrant shall maintain records showing the receipt, transfer, use, storage and
disposal of all sources of radiation. Additional record requirements are specified elsewhere in 32
Ill. Adm. Code: Ch. II, Subchapters b and d. Each record required by this Part shall be legible
throughout the specified retention period. The record shall be the original or a reproduced copy
or a microform provided that the copy or microform is authenticated by authorized personnel.
The microform shall be capable of producing a clear copy throughout the required retention
period. Records may be stored in electronic media with the capability for producing legible,
accurate and complete records during the required retention period. Records such as letters,
drawings and specifications shall include all pertinent information such as stamps, initials and
signatures. The licensee or registrant shall maintain adequate safeguards against tampering with
and loss of records.

(Source: Amended at 35 Ill. Reg. 2908, effective February 7, 2011)

Section 310.50  Inspections

a) Each person shall afford the Agency at all reasonable times opportunity to inspect
radiation installations and sources of radiation and the premises and facilities in
which those radiation installations and sources of radiation are used or stored.
(See 420 ILCS 40/27)

b) Each person shall make available to the Agency for inspection, upon reasonable
notice, records maintained pursuant to 32 Ill. Adm. Code: Chapter II,
Subchapters b and d.

(Source: Amended at 45 Ill. Reg. 9911, effective July 22, 2021)

Section 310.60  Tests

Each licensee and registrant shall perform upon instructions from the Agency, or shall permit the
Agency to perform, such reasonable tests as the Agency deems appropriate or necessary
including, but not limited to tests of:

a) sources of radiation;
b) installations in which sources of radiation are used or stored;

c) radiation detection and monitoring instruments; and

d) other equipment and devices used in connection with utilization or storage of licensed or registered sources of radiation.

(Source: Amended at 29 Ill. Reg. 20748, effective December 16, 2005)

Section 310.70 Additional Requirements

a) The Agency is authorized to inspect and investigate the premises and operations and personnel of any radiation installation, whether or not the installation is required to be registered or licensed by the Agency, for the purpose of studying and evaluating the health hazards caused by the use and operation of machines and material.

b) The Agency may impose additional requirements upon any licensee or registrant if the Agency deems these requirements to be necessary to minimize the danger to public health and safety or the environment.

c) Nothing in 32 Ill. Adm. Code: Chapter II, Subchapters b and d relieves the licensee or registrant from complying with other applicable Federal, State or local requirements governing any toxic, hazardous, medical or any other property of these materials or products containing these materials.

(Source: Amended at 29 Ill. Reg. 20748, effective December 16, 2005)

Section 310.74 Cost Assessment

The Agency has authority under the Radiation Protection Act of 1990 [420 ILCS 40] to take actions necessary to abate violations of the Act or any rules or regulations promulgated under the Act and may provide that all or a portion of the cost of such actions be assessed to operators of radiation installations or other persons responsible for the violation or contamination. [420 ILCS 40/36]

a) The Agency may assess all or a portion of the costs incurred to abate violations to responsible operators of radiation installations or other responsible persons. Costs that are assessed shall be based on the Agency's actual response costs, including, but not limited to:

1) Time required by the Agency professional staff to coordinate response;
2) Time spent traveling and providing administrative support;

3) Performance or oversight of decontamination activities at properties contaminated with radioactive material;

4) Performance or oversight of confirmatory environmental monitoring;

5) Performance or oversight of treatment, storage, transfer and disposal of sources of radiation;

6) Equipment and supplies; and

7) Contractual support, if any, incurred by the Agency.

AGENCY NOTE: These support service costs may include, but are not limited to, rental of specialized equipment, acquisition of additional professional expertise not available within the Agency and laboratory fees charged to the Agency.

b) Any party affected by an order of the Agency assessing cost shall have the right to a hearing before the Agency in accordance with 32 Ill. Adm. Code 200.

(Source: Amended at 29 Ill. Reg. 20748, effective December 16, 2005)

Section 310.75 Emergency Response Cost Recovery

The Agency has authority under the Radiation Protection Act of 1990 [420 ILCS 40] to respond to conditions that constitute an immediate threat to health and to assess the costs of its response against the person or persons responsible for the creation or continuation of the threat. If the Agency is unable to determine who is responsible for the creation or continuation of the threat, the costs shall be assessed against the owner of the property and shall constitute a lien against the property until paid [420 ILCS 40/38(b)].

a) Costs that are assessed shall be based on:

1) The Agency's actual response costs, including, but not limited to:
   A) Time required by Agency professional staff to coordinate response;
   B) Time spent traveling and providing administrative support;
C) Performance or oversight of decontamination activities at properties contaminated with radioactive material;

D) Performance or oversight of confirmatory environmental monitoring;

E) Performance or oversight of treatment, storage and disposal of sources of radiation;

F) Equipment and supplies; and

G) Contractual support, if any, incurred by the Agency.

AGENCY NOTE: These support service costs may include, but are not limited to, rental of specialized equipment, acquisition of additional professional expertise not available within the Agency and laboratory fees charged to the Agency.

2) Costs incurred by other units of government while assisting the Agency, including agencies of the federal government, provided the costs are submitted as follows:

A) Unless otherwise notified by the Agency, the request for reimbursement must be received by the Agency within 45 days after the assistance is rendered to the Agency or 45 days after the costs are determined, whichever is later, but in any case, not later than one year after the assistance is rendered;

B) The request shall be in writing and shall include documentation justifying costs to be reimbursed; and

C) Reimbursable costs may include, but are not limited to, items specified in subsection (a)(1) of this Section.

b) All reimbursable costs described in a reimbursement request by a governmental unit are subject to approval by the Director of the Agency. The Agency may request additional information in support of the requested reimbursement.

c) If a request by a governmental unit for costs is denied, or denied in part, the Agency shall notify the requesting governmental unit of the decision within 30 days after the date the request was submitted.
d) Each bill for emergency response costs assessed under this Section shall identify the items claimed and the costs related to each. Payment is due to the Agency within 45 days after receipt of the bill.

e) After all emergency response costs have been paid by the responsible parties, the Agency shall pay governmental units based on approved requests.

f) Any person assessed costs under this Section shall have the right to a hearing before the Agency provided a written request for a hearing is served on the Agency within 10 days after notice of the assessment. In the absence of receipt of a request for a hearing, the affected party shall be deemed to have waived the right to a hearing [420 ILCS 40/38(b)]. Hearings shall be conducted in accordance with 32 Ill. Adm. Code 200.

(Source: Amended at 29 Ill. Reg. 20748, effective December 16, 2005)

Section 310.78 Deliberate Misconduct

a) Any licensee, registrant, applicant for a license or certificate of registration, employee of a licensee, registrant or applicant, or any contractor (including a supplier or consultant), subcontractor, employee of a contractor or subcontractor of any licensee or registrant or applicant for a license or certificate of registration, who knowingly provides to any licensee, applicant, registrant, contractor or subcontractor any components, equipment, materials or other goods or services that relate to a licensee's, registrant's or applicant's activities in this Part shall not:

1) Engage in deliberate misconduct that causes, or would have caused if not detected, a licensee, registrant or applicant to be in violation of any statute, regulation, limitation on any license issued by the Agency, or order; or

2) Deliberately submit to an Agency licensee, an applicant, or a licensee's, certificate holder's or applicant's contractor or subcontractor, information that the person submitting the information knows to be incomplete or inaccurate in some respect material to the Agency.

b) A person who violates subsection (a)(1) or (a)(2) of this Section may be subject to enforcement action as provided in Section 36, 39, or 45 of the Radiation Protection Act of 1990.

c) For the purposes of subsection (a)(1), deliberate misconduct by a person means an intentional act or omission that the person knows:
1) Would cause a licensee, registrant or applicant to be in violation of any regulation, statute or order, or any term, condition or limitation of any license issued by the Agency; or

2) Constitutes a violation of a requirement, procedure, instruction, contract, purchase order or policy of a licensee, registrant, applicant, contractor or subcontractor.

(Source: Amended at 45 Ill. Reg. 9911, effective July 22, 2021)

Section 310.80 Violations (Repealed)

(Source: Repealed at 45 Ill. Reg. 9911, effective July 22, 2021)

Section 310.81 Policy for Assessment of Civil Penalties

a) Civil penalties shall be assessed in accordance with the provisions of this Section.

b) A civil penalty will be assessed whenever the Agency, based on consideration of the factors set forth in subsection (c), determines that a civil penalty is appropriate and issues a Preliminary Order and Notice of Opportunity for Hearing, in accordance with 32 Ill. Adm. Code 200.

c) Factors to be Considered in Assessing Civil Penalties

1) The Agency shall consider the factors contained in subsection (c)(2) to determine whether a penalty should be assessed, as provided in subsection (d), and the amount of the penalty. However, if the Agency has by rule established the amount to be assessed for a particular violation, the Agency shall assess the penalty as specified in that rule without regard to the factors contained in subsection (c)(2).

AGENCY NOTE: For an example of a rule that establishes the amount of the civil penalty to be assessed, see 32 Ill. Adm. Code 401.170, which specifies the civil penalties to be assessed for violations of the Agency's radiologic technologist accreditation requirements.

2) The factors to be considered by the Agency are:

A) History of Previous Violations. The Agency shall consider the person's history of previous violations of the Radiation Protection Act of 1990, the Agency's rules promulgated under that Act (Title
32, Chapter II, Subchapters b and d) and licenses issued pursuant to the Act. Each prior violation will be considered without regard to whether it led to a civil penalty assessment. A prior violation shall not be considered, however, if the notice or order relating to the prior violation is the subject of pending administrative or judicial review, or if the time to request such review or to appeal any administrative or judicial decision relating to the prior violation has not expired. The Agency shall not consider a prior violation if a Preliminary or Final Order pertaining to that prior violation has been vacated. The Agency shall not consider previous violations that occurred more than 6 years prior to the issuance of the Preliminary Order or other action taken by the Agency for those violations.

B) Severity of the Violation. The Agency shall consider the severity of the violation, including, but not limited to, actual or potential contamination of the environment resulting from the violation and any actual or potential hazard to the health or safety of the public or to workers, resulting from the violation. When evaluating the severity of the violation, the Agency may also consider the impact that the violation has on the Agency's ability to determine compliance with requirements established by statute, regulation or license condition.

C) Culpability. The Agency shall consider whether the person to whom the Preliminary Order was issued was negligent in causing, allowing, or failing to correct the violation, condition, or practice which was cited in the Preliminary Order. The Agency shall also consider:

i) whether the violation was intentional or inadvertent;

ii) whether the violation was allowed to continue once identified;

iii) whether actions were taken to correct or mitigate the violation and the timeliness of those actions; and

iv) whether the violation was voluntarily reported to the Agency.

d) Determination of the Amount of Penalty; Assessment of Separate Violations for
Each Day

1) The Agency may assess a civil penalty not to exceed $10,000 per violation for each day the violation continues. If the Agency's rules (Title 32, Chapter II, Subchapters b and d) specify the amount of the civil penalty to be assessed for a particular violation, the Agency shall assess the civil penalty in that amount so specified, without consideration of the factors listed in subsection (c) of this Section.

2) When determining the amount of penalty, the Agency shall consider each day of a continuing violation to be a separate violation. Accordingly, the Agency may assess a separate penalty, in accordance with this Section, for each day that a violation continues.

(Source: Amended at 45 Ill. Reg. 9911, effective July 22, 2021)

Section 310.82 Procedures for Assessment of Civil Penalties (Repealed)

(Source: Repealed at 45 Ill. Reg. 9911, effective July 22, 2021)

Section 310.90 Impounding

a) Authority of Department in cases constituting an immediate threat to health. Notwithstanding any other provision of the Act, whenever the Department finds that a condition exists which constitutes an immediate threat to health due to the violation of any provisions of this Act or any code, rule, regulation or order promulgated under this Act and requiring immediate action to protect the public health or welfare, it may issue an order reciting the existence of such an immediate threat and the findings of the Department pertaining thereto. The Department may summarily cause the abatement of such violation or may direct the Attorney General to obtain an injunction against such violator. [420 ILCS 40/38]

b) Such order shall be effective immediately but shall include notice of the time and place of a public hearing before the Department to be held within 30 days of the date of such order to assure the justification of such order. On the basis of such hearing the Department shall continue such order in effect, revoke it or modify it. Any party affected by an order of the Department shall have the right to waive the public hearing proceedings. [420 ILCS 40/38]

(Source: Amended at 23 Ill. Reg. 14454, effective January 1, 2000)
Section 310.100  Prohibited Uses

a) Hand-held fluoroscopic screens shall not be used with x-ray equipment.

b) Shoe-fitting fluoroscopic devices shall not be used.

(Source: Amended at 17 Ill. Reg. 18472, effective January 1, 1994)

Section 310.110  Communications

All communications and reports concerning these regulations and any applications filed thereunder may be submitted to the Agency as follows:

a) By mail addressed: IEMA-Division of Nuclear Safety, 1035 Outer Park Drive, Springfield, IL 62704;

b) By hand delivery to the Agency's offices at 1035 Outer Park Drive, Springfield, IL 62704; or

c) Where practicable, by electronic submission. Electronic submission shall be made in a manner that enables the Agency to receive, read, authenticate, distribute, and archive the submission, and process and retrieve it a single page at a time. Guidance on making electronic submissions can be obtained by contacting the appropriate Agency program.

(Source: Amended at 45 Ill. Reg. 9911, effective July 22, 2021)

Section 310.120  Plans and Specifications

The Director may require the user of any new or altered radiation installation to prepare plans and specifications of the proposed installation and submit them to the Agency for review and approval prior to starting construction or operation.

(Source: Amended at 29 Ill. Reg. 20748, effective December 16, 2005)

Section 310.130  The International System of Units (SI) (Repealed)

(Source: Repealed at 17 Ill. Reg. 18472, effective January 1, 1994)

Section 310.140  Units of Exposure and Radiation Dose

a) As used in 32 Ill. Adm. Code: Chapter II, Subchapters b and d, the unit of exposure is the coulomb per kilogram (C/kg) or roentgen (R). One roentgen (R) is
equal to $2.58 \times 10^{-4}$ C/kg.

b) As used in 32 Ill. Adm. Code: Chapter II, Subchapters b and d, the units of radiation dose are:

1) "Gray" (Gy) is the SI unit of absorbed dose. One Gy is equal to an absorbed dose of 1 joule per kilogram (J/kg). (1 Gy = 100 rad).

2) "Rad" is the special unit of absorbed dose. One rad is equal to an absorbed dose of 100 ergs per gram or 0.01 joule per kilogram (J/kg). (1 rad = 0.01 Gy).

3) "Rem" is the special unit of any of the quantities expressed as dose equivalent. The dose equivalent in rem is equal to the absorbed dose in rad multiplied by the quality factor (1 rem = 0.01 Sv).

4) "Sievert" (Sv) is the SI unit of any of the quantities expressed as dose equivalent. The dose equivalent in sievert is equal to the absorbed dose in gray multiplied by the quality factor (1 Sv = 100 rem).

c) As used in 32 Ill. Adm. Code: Chapter II, Subchapters b and d, the quality factors for converting absorbed dose to dose equivalent are as follows:

<table>
<thead>
<tr>
<th>Type of Radiation</th>
<th>Quality Factor (Q)</th>
<th>Absorbed Dose Equal to a Unit Dose Equivalenta</th>
</tr>
</thead>
<tbody>
<tr>
<td>X, gamma or beta radiation and high-speed electrons</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Alpha particles, multiple-charged particles, fission fragments and heavy particles of unknown charge</td>
<td>20</td>
<td>0.05</td>
</tr>
<tr>
<td>Neutrons of unknown energy</td>
<td>10</td>
<td>0.1</td>
</tr>
<tr>
<td>High-energy protons</td>
<td>10</td>
<td>0.1</td>
</tr>
</tbody>
</table>

*a Absorbed dose in gray equal to 1 Sv or the absorbed dose in rad equal to 1 rem.
d) If it is more convenient to measure the neutron fluence rate than to determine the neutron dose equivalent rate in rem per hour or sievert per hour, as provided in subsection (c) of this Section, 0.01 Sv (1 rem) of neutron radiation of unknown energies may, for purposes of 32 Ill. Adm. Code: Chapter II, Subchapters b and d, be assumed to result from a total fluence of 25 million neutrons per square centimeter incident upon the body. If sufficient information exists to estimate the approximate energy distribution of the neutrons, the licensee may convert a measured tissue dose in gray (rad) to dose equivalent in sievert (rem) by using the fluence rate per unit dose equivalent or the appropriate Q value shown below.

<table>
<thead>
<tr>
<th>Neutron Energy (MeV)</th>
<th>Quality Factor(^a) (Q)</th>
<th>Fluence per Unit Dose Equivalent(^b) (neutrons cm(^{-2}) Sv(^{-1}))</th>
<th>Fluence per Unit Dose Equivalent(^b) (neutrons cm(^{-2}) rem(^{-1}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>2.5 E(^{-8}) (thermal)</td>
<td>2</td>
<td>980 E(^{8})</td>
<td>980 E(^{6})</td>
</tr>
<tr>
<td>1 E(^{-7})</td>
<td>2</td>
<td>980 E(^{6})</td>
<td>980 E(^{6})</td>
</tr>
<tr>
<td>1 E(^{-6})</td>
<td>2</td>
<td>810 E(^{8})</td>
<td>810 E(^{6})</td>
</tr>
<tr>
<td>1 E(^{-5})</td>
<td>2</td>
<td>810 E(^{6})</td>
<td>810 E(^{6})</td>
</tr>
<tr>
<td>1 E(^{-4})</td>
<td>2</td>
<td>840 E(^{6})</td>
<td>840 E(^{6})</td>
</tr>
<tr>
<td>1 E(^{-3})</td>
<td>2</td>
<td>980 E(^{6})</td>
<td>980 E(^{6})</td>
</tr>
<tr>
<td>1 E(^{-2})</td>
<td>2.5</td>
<td>1010 E(^{8})</td>
<td>1010 E(^{6})</td>
</tr>
<tr>
<td>1 E(^{-1})</td>
<td>7.5</td>
<td>170 E(^{8})</td>
<td>170 E(^{6})</td>
</tr>
<tr>
<td>5 E(^{-1})</td>
<td>11</td>
<td>39 E(^{8})</td>
<td>39 E(^{6})</td>
</tr>
<tr>
<td>1</td>
<td>11</td>
<td>27 E(^{8})</td>
<td>27 E(^{6})</td>
</tr>
<tr>
<td>2.5</td>
<td>9</td>
<td>29 E(^{8})</td>
<td>29 E(^{6})</td>
</tr>
<tr>
<td>5</td>
<td>8</td>
<td>23 E(^{8})</td>
<td>23 E(^{6})</td>
</tr>
<tr>
<td>7</td>
<td>7</td>
<td>24 E(^{8})</td>
<td>24 E(^{6})</td>
</tr>
<tr>
<td>10</td>
<td>6.5</td>
<td>24 E(^{8})</td>
<td>24 E(^{6})</td>
</tr>
<tr>
<td>14</td>
<td>7.5</td>
<td>17 E(^{8})</td>
<td>17 E(^{6})</td>
</tr>
<tr>
<td>20</td>
<td>8</td>
<td>16 E(^{8})</td>
<td>16 E(^{6})</td>
</tr>
<tr>
<td>40</td>
<td>7</td>
<td>14 E(^{8})</td>
<td>14 E(^{6})</td>
</tr>
<tr>
<td>60</td>
<td>5.5</td>
<td>16 E(^{8})</td>
<td>16 E(^{6})</td>
</tr>
<tr>
<td>1 E(^2)</td>
<td>4</td>
<td>20 E(^{8})</td>
<td>20 E(^{6})</td>
</tr>
<tr>
<td>2 E(^2)</td>
<td>3.5</td>
<td>19 E(^{8})</td>
<td>19 E(^{6})</td>
</tr>
<tr>
<td>3 E(^2)</td>
<td>3.5</td>
<td>16 E(^{8})</td>
<td>16 E(^{6})</td>
</tr>
<tr>
<td>4 E(^2)</td>
<td>3.5</td>
<td>14 E(^{8})</td>
<td>14 E(^{6})</td>
</tr>
</tbody>
</table>

\(^a\) Value of quality factor (Q) at the point where the dose equivalent is maximum in a 30-centimeter diameter cylinder tissue-equivalent phantom.

\(^b\) Monoenergetic neutrons incident normally on a 30-centimeter diameter cylinder tissue-equivalent phantom.
Section 310.150 Units of Activity

For the purposes of 32 Ill. Adm. Code: Chapter II, Subchapters b and d, activity is expressed in the SI unit of becquerel (Bq) or in the special unit of curie (Ci), or their multiples, or disintegrations (transformations) per unit of time (dps, dpm, tps or tpm).

a) One becquerel (Bq) = 1 disintegration (transformation) per second (dps or tps).

b) One curie (Ci) = \(3.7 \times 10^{10}\) disintegrations (transformations) per second (dps or tps) = \(3.7 \times 10^{10}\) becquerel (Bq) = \(2.22 \times 10^{12}\) disintegrations (transformations) per minute (dpm or tpm).

(Source: Amended at 23 Ill. Reg. 14454, effective January 1, 2000)

(Source: Added at 17 Ill. Reg. 18472, effective January 1, 1994)
Section 310.APPENDIX A  Transport Grouping of Radionuclides (Repealed)

(Source: Repealed at 10 Ill. Reg. 17259, effective September 25, 1986)
Section 310.APPENDIX B  Tests for Special Form Licensed Material (Repealed)

(Source: Repealed at 10 Ill. Reg. 17259, effective September 25, 1986)

Section 310.APPENDIX C  Penalty Assessment Worksheet (Repealed)

(Source: Repealed at 17 Ill. Reg. 18472, effective January 1, 1994)