Radiation Exposure of the U.S. Population from Consumer Products and Miscellaneous Sources

Recommendations of the NATIONAL COUNCIL ON RADIATION PROTECTION AND MEASUREMENTS

Issued December 30, 1987

these people would receive lung dose-equivalent rates in the range of 0.02 to 0.4  $\mu$ Sv/y (2 to 40  $\mu$ rem/y). Again, if the assumption is made that the electric power needs are met through the operation of a larger number of plants of smaller capacity, the number of people exposed would probably be larger, but their lung dose-equivalent rates would probably be correspondingly reduced. In either case, however, the dose-equivalent rates are so low as to make the contribution from this source of little significance in the assessment of the total population dose from consumer products.

3.2.6.3 Combustion of Natural Gas. Natural gas contains on the order of 370 to 740 mBq/l (10 to 20 pCi/l) of radon. This amount contributes less than 370 TBq/y ( $10^4$  Ci/y) to the worldwide inventory estimated to be 1.5 EBq (40 MCi) (NCRP, 1975). The dose equivalent to the lungs of people using natural gas in cooking ranges in the home has been calculated by Barton *et al.* (1973) and appears to be about two percent of the normal natural background dose equivalent from radon and its decay products. On that basis, the resulting effective dose equivalent rates to people in homes using natural gas for cooking would be about  $20 \mu \text{Sv/y}$  (2 mrem/y) to the alveolar region and  $90 \mu \text{Sv/y}$  (9 mrem/y) to the segmental bronchial region of the lung. This latter estimate agrees well with the  $60 \mu \text{Sv/y}$  (6 mrem/y) value reported by Johnson *et al.* (1973). According to the American Gas Association (AGA, 1974), the number of residential households using natural gas is about 41 million. If there are, on an average, 3.1 persons per household, the total number of people exposed to radon from the combustion of natural gas in ranges within homes in the United States is estimated to be about 125 million. The average annual bronchial epithelial dose equivalent to those people who are exposed would be about  $50 \mu \text{Sv}$  (5 mrem). Using a weighting factor of 0.08, the annual average effective dose equivalent to these people would be about  $4 \mu \text{Sv}$  (0.4 mrem) and the associated collective effective annual dose equivalent would be about 500 person-Sv (50.000 person-rem).

In addition, it is estimated that about 16 million people are exposed to radon as a result of the combustion of natural gas in unvented heaters in homes. The dose-equivalent rate to the bronchial epithelium for these people is estimated to be about 0.22 mSv/y (22 mrem/y) (Johnson *et al.*, 1973; Gesell, 1974). Again, using a weighting factor of 0.08, the annual average effective dose equivalent to these people would be about 18  $\mu$ Sv (1.8 mrem) and the associated collective effective annual dose equivalent to the U.S. population would be about 290 person-Sv (29,000 person-rem).

Moghissi et al. (1978) have summarized data on collective doses resulting from the use of natural gas. These estimates were based on state-by-state data provided by Johnson et al. (1973). Estimates of the annual doses to the tracheobronchial section of the lung were 8,500 organ-Sv (850,000 organ-rem) due to the operation of unvented space heaters and 18,000 organ-Sv (1,800,000 organ-rem) due to the operation of gas ranges. The estimated associated annual effective dose equivalent for people exposed to space heaters was about 50  $\mu$ Sv (5 mrem); the estimate for people exposed to gas ranges was about 10  $\mu$ Sv (1 mrem). The <sup>222</sup>Rn concentrations assumed in the natural gas were 5,550 mBq/l (150 pCi/l) for California, 3,700 mBq/l (100 pCi/l) for Texas, 1,850 mBq/l (50 pCi/l) for another 5 states, and 370 mBq/l (100 pCi/l) for all other states.

A similar analysis of the use of liquefied petroleum gas (LPG) yields annual collective doses to the tracheobronchial section of the lung of 105 organ-Sv (10,500 organ-rem) from unvented heaters, and 185 organ-Sv (18,500 organ-rem) from ranges (Gesell *et al.*, 1977). These estimates are based on an assumed 713,000 dwellings with unvented LPG heaters and 5.3 million dwellings with LPG ranges.

## 3.2.7 Glass and Ceramics

Naturally occurring radioactive materials have been used in the glass and ceramic industry for over 150 years (Jensen, 1952). Uranium compounds have been employed to produce fluorescent glassware, a variety of colored glazes, and wall tiles. More recently uranium has been incorporated into artificial teeth both for coloring and fluorescent properties. Thorium compounds have been used in wall tiles and electrical materials.

3.2.7.1 Uranium in Glassware. Sodium uranyl carbonate has been commonly employed in the production of fluorescent and iridescent glass. In particular, it was popular until the 1940's to use this material to produce dichroic properties in glass. As the concentration of uranium is increased, the glass becomes more opaque. In 1972 two manufacturers were identified as using uranium as a colorant in non-food glass products such as candlesticks and flower containers. Federal regulations allow glassware to contain up to ten percent by weight uranium or thorium, except in commercially manufactured glass brick, pane glass, ceramic tile, or other glass or ceramic used in construction. Uranium or thorium concentrations in these latter items are limited to 0.05 percent (CFR, 1986c).

<sup>&</sup>lt;sup>4</sup> See Section 3.2.1.

3.2.7.2 Uranium in Glazes. Uranium in the form of oxides and as sodium uranite has been used to produce glazes of black, brown, green, and the spectrum from yellow to red. The glazes were frequently used to decorate tableware and pottery at concentrations ranging from one to twenty percent by weight. As with glassware, the restrictions on the availability of uranium during the 1940's forced manufacturers to find other coloring agents. Since the substitutes were frequently more economical, uranium has not often been used in glazes in recent years. At present no manufacturer is known to use uranium as a glaze for dinnerware. U.S. Nuclear Regulatory Commission (USNRC) exemption limits are set at a maximum of 20 percent by weight (of the glaze) for uranium compounds in glazed ceramic tableware (CFR, 1986c). The acceptability of ceramic glazes for use in food containers is subject to the food additive provisions of the Food and Drug Administration and the use of such glazes is prohibited unless specifically approved. No approvals for such applications of uranium ceramic glazes have been granted to date.

Measurements using film badges have shown that surface dose rates due to gross beta and gamma radiation from various tableware items glazed with uranium range from 5 to 200  $\mu$ Gy/h (0.5 to 20 mrad/h) (Menczer, 1965). Measured uranium enamel surface dose rates of 37  $\mu$ Gy/h (3.7 mrad/h) have been reported (USNRC, 1983). In addition to the external exposures, uranium and lead have been found on occasion to be leachable from glazed ceramics at levels of 10 to 55 ppm (concentration in the leach solution) (Kendig and Schmidt, 1972). The latter is equivalent to 630 mBq/ml (17 pCi/ml) which is in excess of the Maximum Permissible Concentration in drinking water for occupational exposures, 220 mBq/ml (6 pCi/ml) (NCRP, 1959). At the measured concentrations, the chemical toxicity of uranium is considered to be a greater hazard than its associated radiation. As a result of these measurements, the only known producer of uranium glazes for use in foodware in the U.S. has ceased operations.

3.2.7.3 Uranium in Glass Enamel. Uranium has also been used as a coloring agent in various enamel objects, including tableware and jewelry. Problems associated with the use of such items were highlighted by reports in the early 1980's of relatively high exposures associated with the use of cloisonne jewelry. A surface dose rate of 37  $\mu$ Gy/h (3.7 mrad/h) from similar sources has been reported by staff members of the U.S. Nuclear Regulatory Commission (USNRC, 1983). Such jewelry, most of which was being imported, has proven very popular in the U.S. and could be worn in direct contact with the body. Stimulated by these problems, the U.S. Nuclear Regulatory Commission in 1983 banned the use of uranium in enamel products to be sold

in the U.S. (USNRC, 1983, 1984). Products already distributed to retail outlets and consumers were not recalled because of logistical problems and the relatively low estimates of the associated radiological hazard.

3.2.7.4 Dental Products. Porcelain teeth and crowns are composed principally of feldspar minerals that contain small quantities (0.001) percent) of naturally occurring 40K. The practice of adding uranium salts was initiated at least half a century ago when it was discovered that small amounts of the element contributed a natural color and fluorescence to dentures. Restoration of natural appearance is one of the major reasons for using prostheses. Other substances have been found to imitate these characteristics over a broad range of daylight and artificial lighting conditions. The concentrations of uranium required were considered trivial and easily qualified for a licence exempt status when controls were imposed in the 1960's on the use of source material in ceramics. Under regulations of the U.S. Nuclear Regulatory Commission, neither domestic nor imported teeth and powders may contain in excess of 0.05 percent by weight of uranium (CFR, 1986a). Dental products also contain naturally occurring radioactive potassium but there are no controls over the potassium content in these products.

A study by O'Riordan and Hunt (1974) in Great Britain indicated that porcelain teeth containing 0.10 percent uranium could deliver an annual dose equivalent to the oral mucosa of almost 6 Sv (600 rem) by alpha particles and 0.028 Sv (2.8 rem) by beta particles. This estimate is in close agreement with a more recent study by Papastefanou et al. (1987) in Greece in which it was reported that uranium concentrations of 500 ppm could yield a surface dose equivalent of about 4 Sv (400 rem) per year. In a study of dental products in the U.S. (Thompson, 1976), the highest concentration observed, 0.044 percent, was calculated to deliver an annual mucosal dose equivalent of 1.3 Sv (130 rem) from alpha emissions. However, the maximum range of alpha particles in tissue is 30  $\mu$ m so that most of their energy is expended in the superficial cells overlying the sensitive basal layer. Saliva, dental pellicle, calculus, food, and tobacco residues in the mouth further reduce the intensity of the alpha flux to a level where it does not appear to present a significant hazard.

Beta particles can penetrate in tissue to a depth of 200  $\mu$ m. The combined beta emissions of uranium and potassium-40 for the highest concentration sample observed in a study by the Bureau of Radiological Health, were calculated to deliver an annual dose equivalent of 9 mSv (0.9 rem) to the basal layer. The average concentration of uranium in U.S. dental porcelain was estimated to be 0.02 percent. This

corresponds to a uranium beta dose equivalent rate of about 5 mSv (0.5 rem) per year. The potassium-40 contribution generally ranged from 1.4 to 1.9 mSv (0.14 to 0.19 rem) per year.

As of 1971, over 19 million persons in the United States were estimated to wear full dentures and 60 million to wear crowns (DHEW, 1962, 1971). Some 90 million persons were missing at least one tooth although it is not known how many wore bridges or partial dentures. More recent published estimates are not available; however, knowledgeable sources in the dental industry indicate that 40 percent of new dental prostheses contain porcelain, and that uranium is no longer used in porcelain by domestic manufacturers (ADA, 1986). The balance of dental products are acrylics and do not contain uranium.

If it is assumed that 45 million people are wearing dental prostheses with an average concentration of 0.02 percent uranium, and that only beta dose need be considered, they will receive a dose equivalent to 7 mSv (0.7 rem) to the basal mucosa. The contribution from this source to the average annual population dose equivalent to the basal mucosa of the mouth would be estimated to be about 1.3 mSv (0.13 rem). On the basis of a weighting factor of 0.01 for the human skin, and assuming that irradiation of the basal mucosa is equivalent to irradiation of 1 percent of the skin, the weighting factor for irradiation of the basal mucosa could be estimated to be  $0.01 \times 1$  percent or  $10^{-4}$ . The resulting annual collective effective dose equivalent to the U.S. population from this source would be 31.5 person-Sv (3,150 person-rem). This dose is expected to decrease over time as porcelain without uranium displaces the old porcelain containing uranium for use in dental prostheses.

3.2.7.5 Uranium and Thorium Impurities in Ophthalmic Glass.

Ophthalmic glass is used to manufacture lenses for eyeglasses and eyepieces. At present, up to 0.05 percent by weight of source material (uranium or thorium or any combination of these materials) may be contained in any chemical mixture, compound, solution, or alloy without NRC regulation or license requirements. There is a further maximum allowable limit of 0.25 percent by weight of source material in rare earth mixtures and products (CFR, 1986c).

Pecora and Munton (1974) have reported that ophthalmic lenses, tinted by adding thorium salts, can be a source of radiation. They tested rose-tinted lenses from several manufacturers and concluded that the dose-equivalent rate to the corneal epithelium from alpha radiation was 0.1 to 0.3 mSv/h (10 to 30 mrem/h). At a depth of 0.2 cm, the beta dose-equivalent rate was calculated to range from 0.7 to 2  $\mu$ Sv/h (0.07 to 0.20 mrem/h) with a gamma dose-equivalent rate to the entire eye of 0.06 to 0.3  $\mu$ Sv/h (0.006 to 0.030 mrem/h). Another study (Yaniv, 1974) reported thorium concentrations of up to 0.14

percent by weight in some samples of ophthalmic glass, with large variations in natural thorium and uranium content for different batches of glass.

Thorium has been shown (McMillan *et al.*, 1975) to exist as an impurity in the rare earth oxides that are used in the manufacture of certain ophthalmic glasses. The thorium content was found to exceed the limit specified in federal regulations (CFR, 1986a) by as much as a factor of ten. These oxides, and their impurities, are generally thought to be the primary source of radioactivity in certain ophthalmic glasses.

Dose calculations by Tobias and Chatterjee (1974) indicate that the annual alpha-particle dose to the critical tissues of the germinal cell layer of the cornea (50  $\mu$ m), from eyeglasses containing 0.05 percent by weight of <sup>232</sup>Th in equilibrium with its decay products and worn for 16 hours a day, is 2 mGy (0.2 rad) (estimated to be accurate within a factor of two), with an approximately equal absorbed dose from beta particles. Using these data, and applying a quality factor of 20 for alpha radiation, Casarett *et al.* (1974) estimated that the dose-equivalent rate to the germinal cells of the cornea (50  $\mu$ m depth) would be approximately 40 mSv/y (4 rem/y). The dose-equivalent rate at 60  $\mu$ m tissue depth was estimated to be 10 mSv/y (1 rem/y). The beta dose-equivalent rate would be a small fraction of this, however, because of the much smaller quality factor of this radiation.

The Yaniv (1974) study concluded that the radiation dose rates from the ophthalmic glass could be reduced significantly with better quality control of the rare earth and zirconium oxides. Another problem revealed by this study was that the observed radiation is not directly related to the source material content of the glass, due to widely varying daughter-parent equilibrium conditions. The radiation emissions are, in fact, mainly due to the short-lived decay products of <sup>232</sup>Th and <sup>238</sup>U, which can be present in glass even after the parent radionuclides are removed. Thus, control of source material content is not sufficient to eliminate radioactive material from glass. Yaniv recommended that new regulations for ophthalmic glass be established on the basis of emission rates rather than on the abundance by percent of weight of the parent nuclide. The Optical Manufacturers Association, with the assistance of the U.S. Nuclear Regulatory Commission and other governmental agencies, has established a voluntary radiological standard for ophthalmic glass (OMA, 1975).

In 1977, about 96,000,000 persons in the U.S. wore eyeglasses (BOC, 1979). Currently, it is estimated that about half of the eyeglasses in use in the U.S. contain plastic lenses which do not contain radioactive material. The same is true for plastic contact lenses (Buckley *et al.*,

1980). As a result, the current estimate of the number of people who are wearing eyeglasses with glass lenses in the U.S. totals about 50,000,000. Assuming an annual dose equivalent of 40 mSv (4 rem) to the cornea at 50 micrometers depth, and assuming a tissue weighting factor of  $\leq 10^{-4}$ , the annual collective effective dose equivalent to the U.S. population would be about  $\leq 200$  person-Sv ( $\leq 20,000$  person-rem).<sup>5</sup>

## 3.2.8 Thorium Products

Thorium is used in optical glass, gas mantles, tungsten welding electrodes, and in various metal alloys. Approximately 250,000 pounds of thorium were used for these purposes within the United States during 1972.

3.2.8.1 Thoriated Optical Glass. Thorium is added to optical instrument glass in concentrations up to 30 percent by weight to provide certain optical properties. Specifically, glasses having an index of refraction greater than 1.65 or with a product of Abbe number and index of refraction greater than 70 are often made with glasses of high thorium content.

The most abundant isotope of natural thorium, <sup>232</sup>Th, is the very longlived parent of the "thorium series." The presence of the decay products in equilibrium with the parent produces alpha-emission rates six times the parent emission rate. Along with the alpha emissions, a very significant beta- and gamma-emission rate also exists.

The use of thorium in optical glass raises few problems unless the glass is used for an eyepiece in an optical instrument. The alpha and beta radiations are easily stopped by almost any lens enclosure. However, the direct exposure of the eye to a lens at a close distance for long periods of time can deliver a significant dose to the outer tissues of the eye. The use of these special optical glasses near the eye is not authorized under the exemptions issued by the U.S. Nuclear Regulatory Commission (CFR, 1986c). However, cases have been reported of eyepieces containing large quantities of thorium (McMillan and Horne, 1973). These thoriated eyepieces were without labels or specifications indicating that thoriated glass was used. The extent to which these lenses are used (deliberately or inadvertently) for eyepieces in optical instruments is not known.

Casarett et al. (1974) have calculated the dose-equivalent rate delivered to the critical tissue (50 to 60  $\mu$ m depth) of the eye. They assumed that the eyepiece of an instrument that was used for 20 hours per week by a professional user was made from glass with 16 percent <sup>232</sup>Th in equilibrium with its daughters. By also assuming an air gap distance of 0.1 cm between the lens and the outer surface of the lacrimal layer of the eye, they calculated the annual absorbed dose to the critical tissue of the eye at 50 and 60  $\mu$ m depth to be 0.44 and 0.18 Gy (44 and 18 rad), respectively. With the assumption that the quality factor for alpha particles is 20, the corresponding dose-equivalent rates to the eye are 8.8 and 3.0 Sv/y (880 and 360 rem/y), respectively. Eyepieces made from glass having 0.05 percent thorium may be expected to yield a dose-equivalent rate of approximately 30 mSv/y (3 rem/y) at 50  $\mu$ m depth.

McMillan and Horne (1973), after having discovered that these lenses were being used as eyepieces, made similar calculations. Their calculated results agree with those of Casarett et~al. to within 20 to 40 percent. They also confirmed by laboratory measurement the accuracy of their calculated fluence rates at the surface of the glass. One measurement of a lens containing 18 percent thorium by weight yielded a dose rate at the surface of 10  $\mu$ G/h (1 mrad/h). The number of such lenses in use is unknown although it is probably small. An estimate of the average annual population dose equivalent is not possible.

3.2.8.2 Gas Mantles. The mantles in gas-lanterns and gas yard lights consist almost entirely of the oxides of thorium (95 percent), magnesium, aluminum, cerium, beryllium, and silicon. These mantles are the major incandescent element in such lanterns. During their initial curing, they release about 50 percent of their stable beryllium oxide and many of the decay products of thorium into the atmosphere (Griggs, 1973). Although beryllium probably represents the greater health hazard, because of its chemical toxicity to the lungs, alpha and gamma radiation associated with the thorium oxide can also result in radiation exposure to the user.

Recent reports indicate that about 25 million thorium gas mantles are distributed annually in the U.S. (O'Donnell and Etnier, 1981 and Buckley et al., 1980). Approximately 85 percent are used in portable lanterns (e.g., camping), 8 percent in residential and commercial outdoor lights, 5 percent in residential indoor lights and 2 percent in recreational vehicles. The collective dose rate estimates for transportation, distribution, and use of these devices are about 9 person-Sv/y (900 person-rem/y) (Buckley et al., 1980) and 40 person-Sv/y (4,000 person-rem/y) (O'Donnell and Etnier, 1981). The major differences are due to an estimate by O'Donnell and Etnier of a dose of 10 person-

<sup>&</sup>lt;sup>5</sup> There are no reported radiation induced cancers of the cornea. Based on this observation, a weighting factor of  $\leq 10^{-4}$  has been considered appropriate for estimating the effective dose equivalent due to exposures of this organ.

## References

- ABELSON, P. H. (1973). "Consumer product safety," Science, 179, 17.
- ACOG (1984). American College of Obstetricians and Gynecologists. Testimony before the House of Representatives, Subcommittee on Health and Safety, Committee on Education and Labor, May 8, 1984.
- ADA (1986). American Dental Association. Private communications with knowledgeable sources as referenced by the American Dental Association.
- AGA (1974). American Gas Association. Gas Facts—A Statistical Record of the Gas Utility Industry, 1973 (American Gas Association, Arlington, Virginia).
- AMA (1949). American Medical Association. "Shoe-fitting fluoroscopes," J. Am. Med. Assoc. 139, 1004.
- ANSI (1972). American National Standards Institute. Safety Requirements for X-radiation Limits for AC High-Voltage Power Vacuum Interrupters Used in Power Switchgear, Report ANSI-C37.85 (American National Standards Institute, New York).
- AUXIER, J. A. (1973). Contribution of Natural Terrestrial Sources to the Total Radiation Dose to Man, Report ORNL-TM-4323 (Oak Ridge National Laboratory, Oak Ridge, Tennessee).
- Baptiste, M. S., Rothenberg, R., Nasca, P. C., Janerich, D. T., Stutzman, C. D., Rimawi, K., O'Brien, W. and Matuszek, J. (1984). "Health effects associated with exposure to radioactively contaminated gold rings," J. Am. Acad. Dermatol. 10, 1019.
- Barker, R. F., Hopkins, D. R. and Tse, A. N. (1974). "Radiation dose to population (crew and passengers) resulting from the transportation of radioactive material by passenger aircraft in the United States," Paper IAEA-SM-184-15, presented at a seminar on Radiological Safety Evaluation of Population Doses and Application of Safety Standards to Man and the Environment (Por Toroz, Yugoslavia).
- Barton, C. J., Moore, R. E. and Rohwer, P. S. (1973). Contribution of Radon in Natural Gas to the Natural Radioactivity Dose in Homes, Report ORNL-TM-4154 (Oak Ridge National Laboratory, Oak Ridge, Tennessee).
- BECKER, S. (1970). "An investigation of x-radiation from color television receivers in Suffolk County, New York," Radiol. Health Data Rep. 11, 179.
- BECKER, S. (1971). "Results of a follow-up radiation survey on color television sets, Suffolk County, New York," Radiol. Health Data Rep. 12, 457.
- BEE, R., GORDON, J., KWAN, Q., VIERZBA, E. AND WALLO, A. (1985). The Feasibility of Detecting the Import of Unauthorized Radioactive Materials into the United States, NRC Report NUREG/CR-4357 (Nuclear Regulatory Commission, Washington, DC).
- Belanger, R., Buckley, D. W. and Swenson, J. B. (1979). Environmental

- Assessment of Ionization Chamber Smoke Detectors Containing Am-241, NRC Report NUREG/CR-1156 (Nuclear Regulatory Commission, Washington, DC).
- Belli, M., Salvadori, P., Sgrilla, E. and Susanne, A. (1978). "Public health aspects in the use of radium-226 and americium-241 in lightning rods," page 441 in *Radioactivity in Consumer Products*, NRC Report NUREG/CP-0001 (Nuclear Regulatory Commission, Washington, DC).
- BOC (1979). Bureau of the Census. Statistical Abstracts, 1979 (Government Printing Office, Washington, DC).
- BOGGS, R. F., SCHMIDT, G. D. AND WILLIAMS, K. D. (1969). "Radiological health aspects of spent radon seeds," Radiol Health Data Rep., 10, 185.
- Boice, J. D. and Blot, W. J. (1986). Private communication. (National Cancer Institute, Bethesda, Maryland).
- Bradley, F. J. (1986). Private communication.
- Braestrup, C. B. and Mooney, R. T. (1959). "X-ray emission from television sets," Science 130, 1071.
- BRH (1970). Bureau of Radiological Health. *Electronic Product Radiation and the Health Physicist*, BRH/DEP Report 70-26 (Center for Devices and Radiological Health, Rockville, Maryland).
- BRH (1972). Bureau of Radiological Health. 1971 Annual Report on the Administration of the Radiation Control for Health and Safety Act of 1968, U.S. House of Representatives Document No. 92-334 (Center for Devices and Radiological Health, Rockville, Maryland).
- BRH (1981). Bureau of Radiological Health. An Evaluation of Radiation Emission from Video Display Terminals, BRH Report FDA 81-8153 (Center for Devices and Radiological Health, Rockville, Maryland).
- Buckley, D. W., Belanger, R., Martin, P. E., Nicholaw, K. M. and Swenson, J. B. (1980). *Environmental Assessment of Consumer Products Containing Radioactive Material*, NRC Report NUREG/CR-1775 (Nuclear Regulatory Commission, Washington, DC).
- CASARETT, G. W., TOBIAS, C. A., GOLDMAN, M., APPLETON, B., ARCHER, V. E. AND COGAN, D. G. (1974). Radiobiological Evaluation of Thorium in Optical and Ophthalmic Glass (Directorate of Regulatory Standards, U.S. Atomic Energy Commission, Washington, DC).
- CDRH (1986a). Center for Devices and Radiological Health. Unpublished information based on national surveys and internal laboratory testing results (Center for Devices and Radiological Health, Rockville, Maryland).
- CDRH (1986b). Center for Devices and Radiological Health. Unpublished information based on state and industry contacts (Center for Devices and Radiological Health, Rockville, Maryland).
- CFR (1970). Code of Federal Regulations. *Performance Standards for Television Receivers*, 42 CFR 78.210 (now 21 CFR 1020.10) (Government Printing Office, Washington, DC).
- CFR (1973). Code of Federal Regulations. Regulations for Aircraft Security, 14 CFR 121.538 (Government Printing Office, Washington, DC).
- CFR (1974). Code of Federal Regulations. Federal Performance Standards for Cabinet X-ray Systems, 21 CFR 1020.40 (Government Printing Office,

73

- Washington, DC).
- CFR (1981). Code of Federal Regulations. Airplane Operator Security, 14 CFR 108.17 (Government Printing Office, Washington, DC)
- CFR (1986a). Code of Federal Regulations. Certain Items Containing Byproduct Material, 10 CFR 30.15 and Exempt Quantities, 10 CFR 30.18 (Government Printing Office, Washington, DC).
- CFR (1986b). Code of Federal Regulations. Gas and Aerosol Detectors Containing Byproduct Material, 10 CFR 32.26-32.29 (Government Printing Office, Washington, DC).
- CFR (1986c). Code of Federal Regulations. *Unimportant Quantities of Source Material*, 10 CFR 40.13 (Government Printing Office, Washington, DC).
- COHEN, B. S., EISENBUD, M. AND HARLEY, N. H. (1980). "Measurement of the α-radioactivity on the mucosal surface of the human bronchial tree," Health Phys. 39, 619.
- CORBETT, J. O. (1983). "The radiation dose from coal burning: A review of pathways and data," Radiat. Prot. Dosimetry, 4, 1, 5-19.
- COTHERN, C. R., LAPPENBUSCH, W. L. AND MICHEL, J. (1986). "Drinking-water contribution to natural background radiation," Health Phys. 50, 33.
- CPRDH (1971). Commonwealth of Puerto Rico Departartment of Health. "Results of a survey of x-radiation from color television receivers in the metropolitan area of San Juan, Puerto Rico, 1969–1970," Radiol. Health Data Rep. 12, 547.
- CPSC (1985). Consumer Product Safety Commission. Natural Sample Survey of Unreported Residential Fires, Contract Report No. C83-1239 by Audits and Surveys (Consumer Product Safety Commission, Washington, DC).
- CRCPD (1978). Conference of Radiation Control Program Directors. *Natural Radioactivity Contaminator Problems*, published as report EPA-520/4-77, 015 (U.S. Environmental Protection Agency, Washington, DC).
- CRCPD (1982a). Conference of Radiation Control Program Directors. Suggested State Regulations for Control of Radiation, published as Department of Health and Human Services, publication FDA 83-8203 (Government Printing Office, Washington, DC).
- CRCPD (1982b). Conference of Radiation Control Program Directors. Suggested State Regulations for Control of Radiation, Vol. 1, Ionizing Radiation, Sec. C.4(c)(2), Self-Luminous Products Containing Radioactive Material (ii) Radium-226, HHS Publication FDA 83-8203 (Center for Devices and Radiological Health, Rockville, Maryland).
- CRCPD (1982c). Conference of Radiation Control Program Directors. Suggested State Regulations for Control of Radiation, Vol. 1, Ionizing Radiation, Sec. C.4(c)(3), Gas and Aerosol Detectors Containing Radioactive Materials. HHS Publication FDA 83-89203 (Government Printing Office, Washington, DC).
- Cross, F. T. (1984). "Radioactivity in Cigarette Smoke Issue," Health Phys. 46, 205.
- Cross, F. T., Harley, N. H. and Hofmann, W. (1985). "Health effects and risks from <sup>222</sup>Rn in drinking water," Health Phys. **48**, 659.
- DasGupta, A. K. and Fujimoto, K. R. (1970). "Radiation hazard assessment of privately owned colour television receivers in Canada," page 221 in

- Electronic Product Radiation and the Health Physicist, BRH/DEP Report 70-26 (Center for Devices and Radiological Health, Rockville, Maryland).
- DELHOVE, J. (1970). A Study of the Radiological Aspects of the Manufacture and Use of Radioactive Lightning Conductors, Euratom Report EUR-4292 (In French) (National Technical Information Service, Springfield, Virginia).
- DHEW (1962). Department of Health, Education and Welfare. Decayed, Missing, and Filled Teeth in Adults, United States, Vital and Health Statistics Series 11, No. 23 (Health Resources Administration, Department of Health, Education and Welfare, Washington, DC).
- DHEW (1971). Department of Health, Education, and Welfare, *Edentulous Persons-United States-1971*, Vital and Health Statistics, Series 11, No. 23 (Health Resources Administration, Department of Health, Education, and Welfare, Washington, DC).
- DILLINGHAM, W. E. AND STALKER, W. W. (1951). "Kentucky study reveals most x-ray shoe-fitting machines are defective," Ind. Health Monthly II, No. 5.
- DOT (1980). Department of Transportation. Federal Register, 45 FR 20101, March 27, 1980.
- DUPREE, W. G. AND CORSENTINO, J. S. (1975). *United States Energy Through the Year 2000*, (Revised), Report of the Bureau of Mines (U.S. Department of Interior, Washington, DC).
- EIA (1971). Electronic Industries Association. "Evaluation of television contribution to the annual genetically significant radiation dose to the population," Radiol. Health Data Rep. 12, 363.
- EPA (1977a). United States Environmental Protection Agency. Radiological Surveys of Idaho Phosphate Ore Processing—The Thermal Process Plant, Technical Note ORP/LV-77-3 (United States Environmental Protection Agency, Las Vegas, Nevada).
- EPA (1977b). United States Environmental Protection Agency. *Radiation Protection Activities*, EPA Report EPA-520/4-78-003 (United States Environmental Protection Agency, Washington, DC).
- EPA (1978). United States Environmental Protection Agency. Radiological Surveys of Idaho Phosphate Ore Processing—The Wet Process Plant, Technical Note ORP/LV-78-1 (United States Environmental Protection Agency, Washington, DC).
- EPA (1984). United States Environmental Protection Agency. Background Information Document (Integrated Risk Assessment); Final Rule for Radionuclides, EPA Report EPA 520/1-84-002-2 Vol. II (United States Environmental Protection Agency, Washington, DC).
- EPRI (1987). Electric Power Research Institute. Classification of Fly Ash for Use in Cement and Concrete, Report EPRI CS-5116 (Electric Power Research Institute, Palo Alto, California).
- FITZSIMMONS, C. K., McNelis, C. K. and Wruble, D. T. (1972). "Tritium activity in urine from luminous dial wristwatches," Health Phys. 22, 514.
- Fornes, E. and Ortiz, P. (1978). "Radioactive lightning rods, static eliminators and other radioactive devices," page 462 in *Radioactivity in Consumer Products*, NRC Report NUREG/CP-0001 (Nuclear Regulatory Commission, Washington, DC).

- FR (1975). Federal Register. Proposed exemption for "Persons using spark gap irradiators containing cobalt-60," FR 40, No. 207 (Government Printing Office, Washington, DC).
- GESELL, T. F. (1974). "Estimation of the dose equivalent to the U.S. population from radon in liquified petroleum gas," page 347 in *Proceedings of a Symposium on Population Exposures, Eighth Midyear Topical Symposium of the Health Physics Society, Knoxville, Tennessee*, Report CONF-741018 (National Technical Information Service, U.S. Department of Commerce, Springfield, Virginia).
- GESELL, T. G., JOHNSON, R. H. AND BERNHARDT, D. E. (1977). Assessment of Potential Radiological Population Health Effects from Radon in Liquified Petroleum Gas, EPA Report EPA 520/1-75-002 (Environmental Protection Agency, Washington, DC).
- GORDON, J. A. (1968). Interim Report of the Study of Public Health Aspects of Fossil Fuel and Nuclear Power Plants (Southeastern Radiological Health Laboratory Report, Montgomery, Alabama).
- GRIGGS, K. (1973). "Toxic metal fumes from mantle-type camp lanterns," Science 181, 842.
- GUIMOND, R. J. AND WINDHAM, S. T. (1975). "Radioactivity Distribution in Phosphate Products, By-Products, Effluents, and Wastes," Technical Note ORP/CSD—75-3 (U.S. Environmental Protection Agency, Washington, DC).
- GUIMOND, R. J. (1978). "The radiological aspects of fertilizer utilization," pages 380-393 in *Radioactivity in Consumer Products*, NRC Report NUREG/CP-0003 (Nuclear Regulatory Commission, Washington, DC).
- GREENHOUSE, N. A. AND PETERSON, G. C. (1972). "An evaluation of radiation hazards from high voltage vacuum interrupters," Health Phys. 23, 85.
- HALPERIN, J. A. AND HESLEP, J. M. (1966). "Radium in military surplus commodities," Public Health Rep. 81, 1057.
- HAYWOOD, F. F., WARD, D. R., LITTLETON, C. P., POSTON, J. W., ROBINSON,
  E. M. AND BURSON, Z. G. (1970). "HPRR and accelerator operations," page
  187 in Health Physics Division Annual Report for the Period Ending July
  31, 1970, Report ORNL-4584 (Oak Ridge National Laboratory, Oak Ridge,
  Tennessee).
- HESS, C. T., MICHEL, J., HORTON, T. R., PRICHARD, H. M. AND CONIGLIO, W. A. (1985). "The occurrence of radioactivity in public water supplies in the United States," Health Phys. 48, 553.
- HHS (1986). U.S. Department of Health and Human Services. Smoking and Health—A National Status Report. Report to the Congress, DHHS Publication No. (CDC) 87-8396 (Public Health Service, Centers for Disease Control, Atlanta, Georgia).
- IAEA (1967). International Atomic Energy Agency. Radiation Protection Standards for Radioluminous Timepieces—Recommendations of the European Nuclear Energy Agency and the International Atomic Energy Agency, IAEA Safety Series No. 23 (International Atomic Energy Agency, Vienna).
- ICRP (1955). International Commission on Radiological Protection. Recommendations of the International Commission on Radiological Protection,

- Published as Supplement No. 6, Br. J. Radiol. (British Institute of Radiology, London).
- ICRP (1960). International Commission on Radiological Protection. Article 19, page 14 in Report of Committee III on Protection Against X-Rays Up to Energies of 3 MeV and Beta- and Gamma-Rays from Sealed Sources, ICRP Publication 3 (Pergamon Press, New York).
- ICRP (1977). International Commission on Radiological Protection. Recommendations of the International Commission on Radiological Protection, ICRP Publication 26, Annals of the ICRP, 1, 3 (Pergamon Press, New York).
- ICRP (1981). International Commission on Radiological Protection. *Limits for Inhalation of Radon Daughters by Workers*, ICRP Publication 32, Annals of the ICRP, 6, 1, 1 (Pergamon Press, New York).
- JENSEN, D. (1952). "Uranium glass and its fluorescence," Rocks and Minerals, 27, 230.
- JOHNSON, R. H., BERNHARDT, D. E., NELSON, N. S. AND CALLEY, H. W., JR. (1973). Assessment of Potential Radiological Health Effects from Radon in Natural Gas, EPA Report EPA-520/1-73-004 (U.S. Environmental Protection Agency, Washington, DC).
- JONES, D., BJARNGARD, B., SIMON, N. AND HARLEY, J. (1968). "The application of phosphor-teflon thermoluminescent dosimeters to the measurement of dose-rate from radioactive jewelry," Brit. J. Radiol., 41, 944.
- Keating, L. (1986). Private communication.
- KENDIG, J. F. AND SCHMIDT, G. D. (1972). Radiological Hazards Evaluation: Bureau of Foods Pottery Sample No. 093-549E Mango/Red, Staff Report, Radioactive Materials Branch (Center for Devices and Radiological Health, Rockville, Maryland).
- KLEIN, H. F., ROBINSON, E. W., GRAGG, R. L. AND ROLOFSON, J. W. (1970). "Evaluation of emissions from radium dial watches," Radiol. Health Data Rep. 11, 7.
- KRUGER, P. (1962). "In ruthenium metal—Ru<sup>106</sup> concentrated from fallout," Nucleonics 20, 102.
- Lewinsky, B. (1985). "Th-232 in T.V. camera lenses," Radiat. Prot. Dosimetry, 11, 65.
- LITTLE, J. B., RADFORD, E. B., McCombs, H. L. and Hunt, V. R. (1965). "Distribution of polonium in pulmonary tissues of cigarette smokers," N. Engl. J. Med. **273**, 1343.
- Lubenau, J. O. and Nussbaumer, D. (1986). "Radioactive contamination of manufactured products," Health Phys. 51, 409-425.
- MARTELL, E. A. (1974). "Radioactivity of tobacco trichromes and insoluble cigarette smoke particles," Nature **249**, 216.
- Martin, J. E. (1974). "Comparative population radiation dose commitments of nuclear and fossil fuel electric power cycles," page 317 in *Proceedings of a Symposium on Population Exposures, Eighth Midyear Topical Symposium of the Health Physics Society, Knoxville, Tennessee*, Report CONF-741018 (National Technical Information Service, Springfield, Virginia).
- MARTIN, J. E., HARWARD, E. D. AND OAKLEY, D. T. (1969). "Comparison of