NUREG/CR-3569 PNL-4915

# Spectral and Dosimetric Measurements of Photon Fields at Commercial Nuclear Sites

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Pacific Northwest Laboratory Operated by Battelle Memorial Institute

Prepared for U.S. Nuclear Regulatory Commission

> 8408290171 840831 PDR NUREG CR-3569 R PDR

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Manuscript Completed: December 1983 Date Published: August 1984

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NRC FIN B2419

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#### ABSTRACT

Spectral and dosimetric measurements of photon fields were performed at seven commercial nuclear reactor sites. Revisions in 10 CFR 20 that specify exposure-to-dose conversion factors (C ) much greater than unity for photons between 40 keV and 200 keV could impact personnel monitoring practices. Monitoring at effective depths of 1 cm of tissue and shallower could underestimate doses received from high-energy photon fields (>3 MeV).

No locations with large C factors (approximately 1.5 rad/R) were found. The most significant production of low-energy photons was found to be due to photon scattering. The scatter continuum has an effective C factor of approximately 1.2 rad/R. One location was found with a nearly pure scatter spectrum. Other locations contained significant contributions from mediumenergy photons due primarily to radioactive decay of cobalt and cesium isotopes. Monitoring requirements at 0.007-cm and 1.0-cm depths in tissue were found to be adequate for estimating dose received in radiation fields containing high-energy photons. Enhanced surface doses attributed to high-energy knock-on electrons were measured in all locations monitored. Personnel monitoring techniques may provide inaccurate results in high-energy fields.

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#### ACKNOWLEDGMENTS

The authors acknowledge the help of many support groups at the Pacific Northwest Laboratory and of the health physics groups at the reactor sites. We thank Fred Eichner, Virginia Tews and Chuck Souder for the preparation and the analyses of the TL dosimeters. We thank Mark Johnson for assistance during the site measurements. We thank Marianna Cross for word processing services and Jan Baer for editing services. We also acknowledge the help of Lori Bisping and Janet Rhodes for trip scheduling and typing of progress reports. Jack Prewett and the Pacific Northwest Laboratory graphics section did an excellent job on the many line drawings and spectra. We extend a special thanks to Chris Simoner for assistance with the computer analysis of the data.

#### 1. INTRODUCTION

Current personnel monitoring practices may be inappropriate for radiation areas with significant contributions from photons with energies less than 200 keV. The use of an exposure-to-dose conversion factor of unity for all photon energies is specified in Title 10, Part 20 of the <u>Code of Federal Regulations</u> (10 CFR 20). Experimentally measured and calculated conversion factors are significantly greater than unity for energies between 40 keV and approximately 200 keV, with a maximum of 1.5 at approximately 60 keV, as listed in Table 1 (American National Standards Institute [ANSI] 1983). Revisions of 10 CFR 20 to include improved conversion factors could have a significant impact on personnel monitoring practices. In extreme cases, radiation zones may have to be monitored to determine the appropriate conversion factors.

Photon	Conversion Factor $(rem \cdot R^{-1})^{(b)}$ to Dose Equivalent					
Energy	in the ICRU Sphere at a Depth of					
(keV)	1.0 cm ("Deep")	0.3 cm	0.007 cm ("Shallow")			
15	0.28	0.67	0.90			
20	0.58	0.79	0.94			
30	1.00	1.07	1,11			
40	1.28	1.29	1.34			
50	1.46	1.46	1,50			
60	1.47	1.46	1,50			
70	1.45	1.45	1,50			
80	1.43	1.43	1,48			
90	1.41	1.41	1.45			
100	1.39	1.39	1.43			
110	1.37	1.37	1,40			
120	1.3"	1.35	1,36			
130	1.33	1.35	1.34			
140	1.32	1.32	1.32			
150	1.30	1.30	1,30			
662	1.03	1.03	1.03			

TABLE 1. Conversion Factors for Computing Dose Equivalent from Exposure for the ICRU<sup>(a)</sup> Sphere (ANSI N13.11-1983, Appendix C)

(a) ICRU = International Commission on Radiation Units and Measurements. (b) 1 rem =  $10^{-2}$  Sv; 1 R = 2.58 x  $10^{-2}$  C·kg<sup>-1</sup>.

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The presence of low-energy photons<sup>(a)</sup> in radiation areas is due not only directly to the decay of radioactive atoms, but also to photon scattering in shielding material. Shielding attenuates the primary photon field, scatters photons in all directions and shifts the spectral distribution to lower energies primarily through the Compton interaction (Fenyves and Haiman 1969). In heavily shielded areas, maximum fluxes of photons are expected between 50 keV and 150 keV, just above the rise in the photoelectric cross section for the atoms making up the shielding material.

Current personnel monitoring requirements may also be inadequate for dose estimations in fields with a significant contribution of high-energy photons, such as from  $^{16}N$  (6.1 MeV). Doses deposited by high-energy photons at tissue depths of 1 cm or shallower are significantly less than those deposited at the depth of maximum dose, which is approximately 3 cm for 6-MeV photons (Johns and Cunningham 1978). Depth-dose curves presented in Figure 1 for  $^{60}$  Co and 6-MeV photons show the reduced surface dose for the higher energies. While an "ideal" dosimeter monitoring the deposited dose at 1 cm would underestimate the actual deep dose by only approximately 20%, many practical dosimeters can underestimate the dose received by a greater amount. Dosimeters using less filtration on the deep-dose to the lens of the eye can indicate a dose of about half of the maximum received.

To help assess the impact of proposed changes in regulatory requirements for low-energy photon dosimetry and of current regulatory requirements for high-energy photon dosimetry, spectral and dosimetric measurements were performed at representative commercial nuclear sites. The spectral measurement equipment included intrinsic germanium (Ge) detectors, lithium-drifted germanium [Ge(Li)] detectors, and sodium iodide [NaI(T1)] detectors. The Ge and Ge(Li) detectors were used because of their high spectral resolution. The NaI(T1) detector measurements were performed in several locations for comparison. The dose measurement equipment consisted of ionization chambers and thermoluminescent dosimeters (TLD) embedded in a plastic material (phantom) or covered with various aluminum filters. Dose rate as a function of depth in phantom was used to assess relative dose to deep-seated organs, to estimate the spectral composition and beta-to-photon ratio, and to perform measurements using source geometries in work environments.

<sup>(</sup>a) For the purposes of this report, energies below 200 keV are referred as low energies, energies between 200 keV and 3 MeV as medium energies, and energies above 3 MeV as high energies.





Spectral measurements were performed in radiation fields of up to approximately 10 mR/h without a collimator on the detector. For fields up to 50 to 100 mR/h, a collimator was necessary to reduce the data collection rate and allow spectral data to be gathered. Ion chambers were used for greater radiation fields (above approximately 100 mR/h). The TLDs were used in areas with small and large exposure rates, overlapping with rates for both the spectral and ion chamber measurements.

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Measurements were performed at seven commercial nuclear reactor sites (Table 2). These included four pressurized-water reactors (PWRs) and three boiling-water reactors (BWRs). Measurements were performed at PWRs three times while at power (operating) and three while in refueling or maintenance

		Operating		
Site	Туре	Age, y	Status	Data
G	PWR	6	Operating	1/82*
K	PWR (twin)	1/2	Operating	3/82
		5	Shutdown	
В	PWR	8	Shutdown	4/82
			Operating	9/82
М	BWR	8	Operating	9/82
			Shutdown	3/83
0	BWR	7	Operating	11/82
P	PWR	21	Shutdown	11/82
Ν	BWR	10	Operating	3/83

#### TABLE 2. Measurement Sites

\* Equipment checkout.

conditions (shutdown). All three BWRs were visited while operating and one while shutdown. One BWR and one PWR were visited during both operating and shutdown conditions for comparison. One twin PWR was visited. Operating plant ages ranged from one-half to ten years. Shutdown plant ages ranged from 5 to 21 years. The site measurements were performed between January 1982 and March 1983. Photon spectra are compiled in Appendix A.

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Dose in rad and dose equivalent in rem are used synonymously in this report. The quality factor for photons is assumed to be unity.

#### 2. MEASUREMENT PROCEDURES AND ANALYSIS TECHNIQUES

The methods of data collection for the photon spectrometers, the tissueequivalent extrapolation chamber and the TL dosimeters are given below. Details of photon field measurement techniques are given in Appendix B.

#### 2.1 PHOTON SPECTROSCOPY

Data accumulated with the photon spectrometer system were analyzed to determine spectral composition and radioisotope identification. Figure 2 contains an example block diagram of the equipment arrangement. The Ge, Ge(Li) or NaI(T1) detectors required high voltage input. The Ge and Ge(Li) detectors also required preamp power. The signal was input to the amplifier, which was typically built into the multichannel analyzer system. Data were stored on magnetic tape cassettes, and were transferred to the analysis computer after measurements were completed.

Rzw data were in the form of pulse-height distributions. The detector did not collect the full energy for most photons because of the escape of scattered photons. The pulse-height distributions were corrected for detector efficiency and scattering losses to yield energy spectra. The analysis procedure is outlined in Figure 3. Using source geometry constraints, the detector was calibrated for efficiency and response (scattering loss) as a function of energy. During data analysis this information was used to convert the pulseheight distributions to energy spectra.



FIGURE 2. Equipment Arrangement for the Photon Spectrometer

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FIGURE 3. Block Diagram of Data Analysis Procedure for the Spectral Measurements

The efficiency curve for the intrinsic Ge detector used for most of the measurements is presented in Figure 4. The low-energy decline of efficiency is due to the thickness of the detector window. The high-energy decline is due to the increase in the penetrating power of photons at the higher energies. The efficiency was measured using radioisotope mixtures with known relative





photon intensities up to about 2 MeV. A smooth curve was drawn through the measured points using a semi-empirical model (Hajnal and Klusek 1974). The extrapolated efficiency curve above 2 MeV was checked using a <sup>16</sup>N field at a reactor site. Differences in efficiency as a function of photon entrance angle were measured to be small compared to other uncertainties of the data analysis.

The detector response function included the full-energy peak, single and double escape peaks at the higher energies, a continuum due to the loss of Compton-scattered photons, and low-energy counts due to the loss of x rays from atoms in the detector crystal. Single-energy and multiple-energy radioisotope spectra were used for input data to develop the response function. A simple parameterization for the response function was used (Seelentag and Panzer 1979). Figure 5 presents calibration data before and after the efficiency and scattering corrections. The leading edge of the Compton continuum is not fully subtracted, and remains in the corrected spectra.

Use of the simple parameterization contributed significantly to the uncertainty in the number of photons near 6 MeV versus the number at lower energies. In locations which have simple spectra dominated by 6-MeV and 511-keV photons, the undersubtraction of the leading edge of the Compton continuum added to the spectra, while the oversubtraction resulted in negative values that were set to zero. The uncertainty in the ratio of low- to highenergy photons may be as high as 50%. However, this uncertainty did not affect the results. It did not contribute significantly to uncertainties below 2 MeV.

For photon spectrometer measurements at the higher exposure rates and at locations for which directionality was considered important, a lead collimator was used. The collimator was composed of approximately 2.5 cm of lead fashioned in the shape of a cylinder with a front plate. A 3-mm-diameter hole was drilled in the center of the front plate. The collimator was mounted so that the detector could be inserted on a level surface without supporting the collimator weight. The response function of the detector with collimator was determined using isotopic sources and incorporated in the data analysis procedure. Figure 6 shows a Co source pulse-height distribution and energy distribution following analysis with collimator corrections. The residual counts at the lower energies are insignificant compared to the Co source photons.

The lower part of Figure 3 showed the procedure used to calculate effective exposure-to-dose correction, or C , factors. The corrected photon spectra represent photons entering the detector per time, or fluence. Fluence



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was converted to exposure rate for each energy interval using the curve in Figure 7 (DHEW 1970). The C factors taken from the calculations of Dimbylow and Francis (1979) for the IČRU sphere were weighted by the corresponding exposure rate to calculate effective C factors. Because factors at all energies were not available, some interpolations were required.

As indicated in Figure 3, geometry factors can significantly affect the result. Figure 8 illustrates curves for C factors for parallel incidence and isotropic incidence. For parallel incidence, all photons an assumed to travel parallel to each other with no intensity variations with distance. For isotropic incidence, all photons are generated in the space around the ICRU sphere, or on the inner surface of a large sphere concentric with the ICRU sphere. Photons travel in all directions. The shallow-depth C factors for isotropic incidence have approximately the same magnitude as the shallow and deep C factors for parallel incidence. However, the deep-depth C factors for isotropic incidence are considerably smaller than for parallel incidence. This is due to a reduction in the dose rate at 1 cm compared to the exposure rate because most of the photons have to transit a large portion of the ICRU sphere to deposit the dose.



FIGURE 7. Conversion from Photon Fluence to Exposure Rate



FIGURE 8. Curves for the Conversion of Exposure to Dose in the ICRU Sphere. Dose equivalent to tissue divided by dose to air is converted using dose per exposure to air.

Dimbylow and Francis calculated two C values: the maximum dose in the sphere at each shell depth, and the dose along the central axis of the sphere parallel to the beam direction. The central axis values approximate the worst-case example of dosimeter underresponse for high-energy photons, assuming that the dosimeter is mounted toward the source. Both quantities are listed in the data tables in Section 3 for the high-energy fields.

Correction factors other than C factors may be more useful for particular applications. The energy response of an instrument can be used to generate correction factors for instrument response to dose as easily as to generate conversion factors for exposure to dose.

#### 2 2 TISSUE-EQUIVALENT EXTRAPOLATION CHAMBER AND ION CHAMBER MEASUREMENTS

Direct measurements of effective C factors were performed using ionization chambers to measure dose in a tissue-equivalent (TE) phantom and exposure in air at the same position. The C factor was obtained by calculating the ratio of dose rate to exposure rate. This technique can determine the true C factor for a pure photon field, including all geometry effects. For each location, the C factor may change with angle due to nonuniformly distributed sources. However, given the exposure rate, the dose to the ICRU sphere or a worker would change with angle in a similar manner.

The dose measurements were performed using a parallel-plate extrapolation chamber embedded inside a TE phantom. An extrapolation chamber is an ionization chamber whose volume can be varied. Data collected for many measurement volumes were used to calculate dose for the small-volume limit. A schematic diagram is shown in Figure 9. A cylindrical plug of TE plastic was moved inside a 30-cm x 30-cm x 15-cm block of the same plastic. The front surface of the plug was coated with a conducting graphite mixture and scribed into a circular collecting electrode of 27-cm<sup>2</sup> area and an outer guard ring. A thin (approximately 7-mg/cm<sup>2</sup>) piece of TE plastic was stretched over the opening to provide a front window. The inner surface of the front window was coated with graphite to provide the high-voltage electrode. A stepping motor with remote



FIGURE 9. Equipment Arrangement for Tissue-Equivalent Extrapolation Chamber/Ion Chamber Compination

control was used to reduce the electrode separation in well-determined increments. The number of motor steps per millimeter was calibrated using a micrometer readout on the drive shaft. Voltage was applied to the electrodes, and the current going to ground from the collecting electrode was integrated using an electrometer. An ionization chamber and electrometer were used to measure exposure. Both the center of the front face of the extrapolation chamber and the center of the ionization chamber were reproducibly placed in a given location using a base plate and a mounting jig.

The composition of the TE plastic is described by Yoder et al. (1979). The low-energy response of the plastic is similar to that of tissue with trace elements. Nelson and Chilton (1983) performed Monte-Carlo calculations and found gocd agreement. Differences found between Yoder's laboratory measurements and the Monte-Carlo calculations for the TE plastic and for idealized tissue with trace elements were small compared with the uncertainties of the field measurements. Differences in response between the spherical geometry used by Dimbylow and Francis and the slab geometry of the extrapolation chamber were also small compared with the uncertainties of the field measurements. No corrections to the data were performed.

#### 2.3 THERMOLUMINESCENCE DOSIMETRY

Thermoluminescent dosimeters were used to obtain a measurement of dose at many depths simultaneously. The distribution of dose as a function of depth changes with photon energy and beta-particle energy. The phosphors used were LiF:Mg (TLD-700) and CaF.:Mn (TLD-400), which are sensitive to photon and betaparticle radiation. The<sup>2</sup>TL material was placed inside of a 20-cm x 20-cm x 15-cm Plexiglas<sup>®</sup> (methylmethacrylate) phantom or packaged in a multielement dosimeter with varying thicknesses of aluminum filters. The multielement dosimeters were used primarily to monitor for the presence of beta particles for the extrapolation chamber measurements.

The advantages of TL dosimetry are that: 1) dose rates can vary over a wide range, overlapping both the photon spectroscopy and extrapolation chamber techniques; 2) fewer assumptions of source geometry are required compared to the photon spectroscopy techniques; 3) many depths can be monitored simul-taneously; and 4) beta-particle contributions can be relatively easily identified. Disadvantages are that: 1) the data provide much less information on spectral composition, and 2) differences between the calibration and measurement geometries increase the uncertainty of the measurements.

Plexiglas is a registered trademark of Rohm and Hass Company.

The interpretation of TLDs exposed in an unknown photon field suffers because of the non-tissue-like response of the phosphors as a function of photon energy. The response of LiF:Mg increases by almost a factor of two at x-ray energies. To improve the discrimination between energies, the CaF :Mn phosphor was added to three of the phantom depth positions. As shown in<sup>2</sup> Figure 10, the response of CaF :Mn is large at x ray energies. The ratio between phosphors was used to discriminate between energies.

The TLD were placed at eight depths, from the surface to 7 cm inside the phantom (Figure 11). They were staggered around the central axis of the phantom to reduce shielding effects by upstream dosimeters. The response of the TLD was determined for the four radiation sources listed in Table 3. The relative response per rem as a function of depth is presented in Figure 12. The estimated depth-dose response for 6-MeV photons was added for reference. The data were analyzed by performing a linear least-squares fit to the calibration data set. After the optimum combination of the calibration response

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FIGURE 11. TLD-Loaded Phantom

TABLE 3. Calibration Sources for the TLD-Loaded Phantom

Source	Туре	Energy	Comments
137 <sub>Cs</sub>	Medium-energy photon	662 keV	
MFE <sup>(a)</sup>	Low-energy photon	34 keV (effective)	NBS filter technique
H150 <sup>(a)</sup>	Low-energy photon	120 keV (effective)	NBS filter technique
90sr/90y	Beta	2.3 MeV (maximum)	

(a) Appendix of National Bureau of Standards (NBS) Special Publication 250 (1983). Support for the MFE technique has been discontinued by NBS.

functions was determined, estimates of the effective exposure-to-dose factors were made. This technique was shown to be inappropriate for areas containing large contributions of photons above 3 MeV. Even though there is a significant difference between the response to 6-MeV photons and the response to photons of other energies, the apparent presence of very high energy electrons in plant environments did not permit a similar analysis without significant ambiguities.



FIGURE 12. TL Response Per Rem Versus Depth

#### 3. SITE MEASUREMENTS

Measurements were performed at seven sites. The type and amount of data collected varied due to variations in the sites' access policies, level of contamination, equipment malfunctions, and research emphasis in areas for which additional data was required.

Delays were often experienced gaining access to the plants. Training and paperwork requirements have increased over the last few years. The time delay between arrival and the initiation of measurements varied from less than 1 day to 2-1/2 days. The greatest delays occurred when formal classroom instruction was required on radiation safety, plant security, mask fitting, and the use of a self-contained breathing apparatus (SCBA). The requirements for plant access and the use of respirator protection also varied. For example, one plant required a polygraph test, another did not allow unescorted access to any plant areas, and a third required complete medical records.

Preparation time for plant access varied with the level of contamination at the measurement locations. Typically, the measurement equipment was packaged in plastic and carried or transported by cart to the site locations where it would be used. Anticontamination dress and occasionally a respirator were required. Access to PWR containment during reactor operation required full anticontamination dress and a full-face respirator or SCBA. The number of measurements were limited by the short length of stay time due to the elevated temperatures and/or dose rates.

The data collected during each site visit is described below. The letter designations for the sites were selected to correspond with neutron and beta measurement projects reported elsewhere (Endres et al. 1983; Rathbun and Roberson 1983). The sites are listed in the approximate chronological order of data collection. When a display of site layouts was considered useful, letter designations were given to each measurement location. These designations are noted on the site layouts. Spectra for all sites are compiled in Appendix A.

#### 3.1 SITE G - OPERATING PWR

This first site visit was primarily a learning experience. The photon spectrometer measurements were not useful because dose rates exceeded maximum acceptable levels. The rates inside containment were substantially above the levels observed on previous visits. A collimator assembly for the spectrometer detector was subsequently built to allow measurements to be performed at higher rates. The extrapolation chamber experienced almost a factor of ten increase in current not associated with radiation (leakage). This was attributed to the elevated temperatures and vibration present inside containment. The addition of an insulated mounting plate and improved electronic connectors minimized this effect during future measurement trips. A return visit to Site G was planned but was cancelled due to an extended shutdown period.

The extrapolation and ion chamber measurements performed in containment are presented in Table 4. The neutron-to-gamma dose-equivalent ratio at the measurement site was 2.5. A correction for the neutron response of the detectors shifted the dose-to-exposure ratio by less than 10%. The cause of measured values greater than unity is not understood. However, the measurements have relatively large uncertainties. The elevated surface dose indicates that current monitoring requirements are adequate.

Two TLD-loaded phantom exposures were performed, one in the auxiliary building near the primary-coolant sampling station and another in containment near the primary-side piping (hot legs) of the steam generator. These data are plotted in Figure 13. Both measurements show an enhanced TLD response at the shallow depths. The measurement performed near the sampling station was analyzed for beta and photon components (Table 5). The result was a small beta component combined with a primarily medium-energy photon component. The measurement in containment showed a large surface dose characteristic of highenergy electrons (1 to 10 MeV) with a penetrating photop component. This location was expected to contain a heavy contribution from <sup>16</sup>N or other high-energy photons. The TLD phantom measurement is consistent with the combination of high-energy photons (5 to 10 MeV) with a large number of high-energy electrons created in the surrounding building and shielding materials.

Location	Exposure Rate (mR/h)	Dose Rate (mrad/h)	Effective C· Factor (rad/Ř)
Operating PWR - Containment,			
Facing Reactor			
- shallow	148	215	$1.45 \pm 0.20$
- deep	103	123	1.19 ± 0.19

TABLE 4. Extrapolation and Ion Chamber Measurements, Site G



FIGURE 13. TLD-Loaded Phantom Measurements, Site G

TABLE 5. ILD-LOAGED Phantom Measurements, SIL	ABLE !	5.	TLD-Loaded	Phantom	Measurements,	Site	G
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		Field	d Strength	(mrad/h)	for	Effective
	Shallow Dose		Photons			C_
Location	Rate (mrad/h)	34 keV	120 keV	662 keV	Beta	(rad/R)
Operating PWR						
- Auxiliary Building Sample Station	64	0	0	62	2	1.03
- Steam Generator Hot Legs	2300		(analysis	inapprop	riate)	

The surface enhancements of the dose measured inside containment do not indicate that personnel doses are being underestimated using current monitoring requirements.

#### 3.2 SITE K - SHUTDOWN AND OPERATING PWR

This twin-reactor site was visited while one of the reactors was shut down for maintenance and one was operating. The shutdown and operating units had equivalent full-power operating times of 5 years and 1/2 year, respectively. The measurement locations in containment are shown in Figures 14 and 15. Other areas visited included the auxiliary building and the contaminatedwaste storage area. The measured exposure or dose rates for each technique can vary significantly for the same location because the rates changed dramatically over a few meters when near a large centralized source.



FIGURE 14. Containment Layout at the 130-ft Elevation for the Shutdown PWR, Site K. Measurements were performed at locations C through F at the elevations shown in parentheses.



FIGURE 15. Containment Layout at the 122-ft Elevation for the Operating PWR, Site K. Measurements were performed at locations I through L and N at the elevations shown in parentheses.

Photon spectrometer measurements were performed with a 48-cm<sup>3</sup> Ge detector. The locations of the measurements are listed in Table 6. Six measurements were performed in the shutdown reactor, six measurements in the operating reactor, and one measurement in a drum storage area used for both units.

Measurements in the shutdown reactor were performed in the effluent sample room, the auxiliary building, and at two levels in containment. The exposure rates ranged from 0.3 to 12 mR/h. Three of the spectra taken are shown in Figures 16, 17, and 18. The effective C factor labelled on the figures is the greatest of the 3 maximum values calculated. The prominent characteristics are the presence of  $^{60}$ Co (5.3-y half-life),  $^{58}$ Co (71-d half-life), electron-positron annihilation radiation and a continuum between approximately 70 keV and 220 keV. The presence of the radioactive isotopes is probably due to "crud" buildup. The ratio of the number of continuum photons to source-decay photons changes by a factor of about 3 from location A (Figure 16) to location D (Figure 18), with a corresponding small increase in the C factor.

TABLE 6. Photon Spectrometer Measurements, Site K

Location	Exposure Rate (mR/h)	Effective Maximum C (rad/R) ×			Effective Central C		
		0.007 cm	0.3 cm	1.0 cm	0.3 cm	1.0 cm	2.5 cm
Shutdown PWR							
A-Effluent Sample Room	1	1.01	1.04	1.04			
B-Auxiliary Bldg. Demineralizer Room, Reactor Coolant Filter (100-ft elevation)	0.3	1.06	1.08	1.08			
C-Containment, Reactor Shield Vent Fan	1	1.02	1.05	1.05			
D-Containment, lodine Removal Fan	1	1.04	1.07	1.07			
(100-ft elevation) E-Containment, Under Reactor Coolant Pump #13 (81-ft elevation)	12	1.01	1.04	1.04			
F-Containment, Under Reactor Coolant Pump #11 (81-ft elevation)	12	1.01	1.04	1.04			
Operating PWR							
G-Effluent Sample Room	1	1.04	1.06	1.06			
H-Auxiliary Bldg. Demineralizer Room, Reactor Coolant Filter (100-ft elevation)	0.2	1.06	1.09	1.09			
I-Containment, Personnel Hatch (100-ft elevation)	0.4	1.03	1.02	1.01			
J-Containment, Near Bioshield of Steam Generator (122-ft elevation)	1.3	1.07	1.08	1.08			
K-Containment, Overlooking Reactor Cavity (122-ft elevation)	2	1.00	1.01	1.03	0.72	0.96	1.02
L-Containment, Outside Airlock (100-ft elevation)	0.9	0.93	0.97	0.95	0.70	0.98	0.99
Other							
M-Hot Drum storage Area	50	0.99	1.02	1.02			



FIGURE 16. Field Data and Corrected Spectrum, Site K, Location A (Shutdown PWR, Sample Room)



FIGURE 17. Field Data and Corrected Spectrum, Site K, Location B (Shutdown PWR, Demineralizer room)


FIGURE 18. Field Data and Corrected Spectrum, Site K, Location D (Shutdown PWR, Containment, Iodine Removal Fan)

The measurement locations in the operating reactor ranged in exposure rate from 0.2 to 2 mR/h. The pulse-height distributions (Figures 19 through 21) show radioisotope spectra from short-lived isotopes in addition to the long-lived isotopes. Prominent short-lived isotopes are <sup>133</sup>Xe (5 d) and <sup>135</sup>Xe (9 h). Xenon is a fission product that easily diffuses into the coolapt water and containment atmosphere. The long-lived isotopes present included <sup>50</sup>Co.

The relative compositions of the cobalt isotope annihilation radiations are similar for the spectra taken in the effluent sample rooms of the shutdown and operating reactors (Figures 16 and 19). However, an enhanced scatter component and the xenon radiations are prominent in the spectrum from the operating reactor. The net effect is an increase in the C factor from 1.04 to 1.06. The increase is primarily due to the increase in the scatter peak. The contribution to dose from the xenon peaks is small.

In the demineralizer rooms (Figures 17 and 20),  $^{60}$ Co is more prominent in the shutdown reactor spectrum, while annihilation radiation is prominent in the operating reactor spectrum. The operating reactor has less equivalent full-power operation (1/2 y) than the shutdown reactor (5 y), and thus has less of the "crud" buildup that contains Co. The prominence of annihilation radiation in the operating plant implies the production of short-lived positron emitters.

The spectrum shown in Figure 21 was taken overlooking the reactor cavity. The data extends up to 8 MeV. No  $^{16}$  N photons were detected. The data is consistent with a general scatter contribution, with some evidence of capture gamma rays on iron (7.6 MeV), as were detected at another operating PWR. The calculated central C factors (Table 6) imply a reduced relative dose at 0.3 cm of 0.7 and a relative dose at 1 cm of near unity.

Three measurements were performed in the shutdown reactor using the TE extrapolation chamber and the ion chamber. The measurements were performed in approximately the same locations as for the measurements performed using both the photon spectrometer and the TLD-loaded phantom. The locations, the measured exposure and dose rates, and the derived effective C factors are given in Table 7. The measurements were performed in areas with exposure rates of about 100 mR/h. Because the chamber signals were small, the measured values have large uncertainties. A measurement in the operating reactor was attempted but was unsuccessful.

All of the effective C factors derived from the extrapolation chamber measurements were near or less than unity. The values less than unity are



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FIGURE 20. Field Data and Corrected Spectrum, Site K, Location H (Operating PWR, Demineralizer Room)



FIGURE 21. Field Data and Corrected Spectrum, Site K, Location K (Operating PWR, Containment, Overlooking Reactor Cavity)

Location	Exposure Rate (mR/h)	Dose Rate (mrad/h)	Effective Exposure-to-Dose Conversion Factor
Shutdown PWR E-Drain Valve, Reactor Coolant Pump #13			
Shallow Deep	96	97 87	$1.01 \pm 0.09$ $0.91 \pm 0.15$
F-Drain Valve, Reactor Coolant Pump #11			
Shallow Deep	87	69 83	$0.79 \pm 0.12$ $0.95 \pm 0.08$
Storage Area M-Hot Drum Storage Area			
Shallow Deep	77	72 57	$0.94 \pm 0.20$ $0.74 \pm 0.16$

TABLE 7. Extrapolation Chamber and Ion Chamber Measurements, Site K

probably due to the presence of distributed sources. Corrections have been made for beta particles. The data does not indicate a significant component of low-energy photons at the higher dose rates.

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The TLD-loaded phantom measurements were performed in four locations with exposure rates ranging from 70 to 260 mrad/h. Table 8 contains the measurement locations and the results of the fitting. Locations E, F, and M are dominated by medium-energy photons, resulting in effective C factors near unity. Figure 22 contains the response of the TL dosimeters versus depth. All curves are normalized to the surface reading. Locations E and F show similar depth-dose functions characteristic of penetrating radiation. Location M has a slightly less penetrating field. Location N (steam generator) has a 7% enhancement at 1 cm compared to the surface. This is possibly caused by partially shielded beta particles from <sup>16</sup>N or high-energy electrons created by the high-energy photons.

Measurements at Site K offered a direct comparison of shutdown and operating conditions and information of the effects of plant age. The presence of xenon isotopes in the operating plant did not significantly increase the dose received. In addition, a compensating effect was the increase in the 511-keV

Surface Dose	Field S	trengths Photons	for	Effective <sup>(a)</sup> C	
Rate (mrad/h)	34 keV	120 keV	662 keV	Beta	(rað/R)
115	7	0	90	8	1.04
106	0	0	97	9	1.03
67	8	0	59	5	1.05
160	(ana	lysis ina	appropriat	e)	
	Surface Dose Rate (mrad/h) 115 106 67 160	Surface Dose Rate (mrad/h)Field S 34 keV11571060678160(anal)	Surface Dose Rate (mrad/h)Field Strengths Photons11570106006780160(analysis inational)	Field Strengths (mrad/hr)Photons34 keV 120 keV 662 keV11570901060097678059160(analysis inappropriat)	Surface Dose Rate (mrad/h) Field Strengths (mrad/hr) for Photons Field Strengths (mrad/hr) for   115 34 keV 120 keV 662 keV Beta   115 7 0 90 8   106 0 0 97 9   67 8 0 59 5   160 (analysis inappropriate) 160 160 160 160

TABLE 8. TLD-Loaded Phantom Measurements, Site K

(a) For the shallow depth (0.007 cm).

annihilation radiation for the operating plant. Differences in dosimetry between the shutdown and operating plants were mainly due to the high-energy photon fields inside containment. The effect of plant aging was to increase the amount of radioactive cobalt, lowering the effective C factors a few percent. The highest calculated C factors (1.08/1.09) occurred in the demineralizer rooms and inside containment behind the bioshield. In both cases, the low-energy scatter continuum was a dominant part of the spectrum. Other locations with large continua also had a significant high-energy tailing. The maximum effect of low-energy photons was to increase the C factors by less than 10%. The effect of operating the plant and the effect of increased age totalled only a few percent difference in C factors. The extrapolation chamber measurements indicated that the calculated C factors may be overestimated due to distributed-source effects.

The spectra taken inside containment of the operating reactor contained a sufficiently large contribution from high-energy photons to deposit less dose at 0.3-cm than at 1-cm or 2.5-cm depths. However, a TL phantom measurement did not confirm the result, possibly indicating the presence of significant beta or electron fields.



FIGURE 22. TLD-Loaded Phantom Measurements, Site K

## 3.3 SITE B - SHUTDOWN AND OPERATING PWR

The PWR at Site B was visited twice, with measurements taken once under shutdown and once under operating conditions (89% of full power). Measurements were performed at 19 locations ranging in exposure rate from 0.4 mR/h to 10.8 R/h. The measurement locations are identified alphabetically on schematic layouts of containment (Figures 23 and 24).

Spectra were taken with a 59-cm<sup>3</sup> intrinsic Ge detector (during reactor shutdown) and a 31-cm<sup>3</sup> intrinsic Ge detector (during reactor operation). Measurement locations and the calculated maximum C factors are listed in Table 9. The C values for the shutdown reactor vary between 1.02 and 1.07. A typical spectrum is presented in Figure 25. Radiation from cesium isotopes contributed significantly to the dose rates, resulting in C values near unity.



FIGURE 23. Containment Layout at the 401-ft Elevation, Site B. Measurements were performed at the locations and elevations indicated.



FIGURE 24. Containment Layout at the 336-ft Elevation, Site B. Measurements were performed at the locations indicated.

	Exposure Rate	Effect	(rad/R)	um C <sub>x</sub>	Effective Central C (rad/R) x			
Location	(mR/h)	0.007 cm	0.3 cm	1.0 cm	0.3 cm	1.0 cm	2.5 cm	
Reactor Containment, PWR Shutdown								
A-Near Escape Hatch (357-ft level)	8	1.02	1.03	1.03				
B-Near Equipment Hatch (357-ft level)	5	1.06	1.07	1.07				
C-Near Elevator (357-ft level)	8	1.04	1.05	1.05				
D-Near Elevator (373-ft level)	8	1.04	1.05	1.05				
E-Near Elevator (401-ft level) F-Deck	4	1.05	1.06	1.06				
(424-ft level)	2	1.06	1.07	1.07				
G-Above Escape Hatch (373-ft level)	10	1.02	1.03	1.03				
Reactor Containment, PWR Operating								
H-Near Personnel Hatch (386-ft level)	3	0.91	0.96	0.99	0.59	0.89	0.98	
I-Near Stairway (386-ft level)	5	0.92	0.97	1.00	0.62	0.89	0.98	
E-Near Elevator (401-ft level)	7	0.84	0.90	0.95	0.38	0.80	0.94	
J-In Personnel Hatch	0.4	1.01	0.91	0.91	0.77	0.87	0.92	

TABLE 9. Photon Spectrometer Measurements, Site B

The spectra measured inside containment during reactor operation had contributions from high-energy photons. In addition to a general distribution of energies up to approximately 8 MeV, several capture gamma transitions were observed. Indicated on Figure 26 are capture gamma transitions for iron and hydrogen. The pulse-height distribution is plotted on a log scale to allow observation of the high-energy contributions. The cesium isotopes were again observed, along with positron-electron annihilation radiation and xenon isotopes. The high-energy photons contributed more to the dose rate than the medium- or low-energy photons. This resulted in maximum C factors at or below unity. Photons from <sup>10</sup>N were not observed at the locations monitored. The calculated central C factors indicated reduced doses at the 0.3-cm and 1-cm depths compared to the 2.5-cm depth.

Extrapolation chamber and ion chamber measurements for Site B are listed in Table 10. An effort was made to measure fields over a wide range of dose rate (106 mrad/h to 10.8 rad/h). Due to time constraints, it was sometimes more productive to measure several locations at one depth rather than one location at several depths. The 1-cm position was chosen because no corrections for beta dose were required.



FIGURE 25. Field Data and Corrected Spectrum, Site B, Location E (Shutdown PWR, Near Elevator, 401-ft Elevation)





Location	Exposure Rate (mR/h)	Dcse Rate (mrad/h)	Beta Dose Rate (mrad/h)	Effective Exposure-to-Dose Conversion Factor
PWR Shutdown (containment)				
K-Piping Near Steam Generator (336-ft- level basement) Shallow Deep	195	196 160	22	0.89 ± 0.07 0.82 ± 0.04
L-Piping Near Reactor Core (336-ft-level basement)	135	106		0.79 ± 0.08
M-Let-Down Heat Exchanger (336-ft level)	10,800	10,800		1.00 ± 0.05
N-Removed Nozzles (336-ft level)	900	804	-	0.89 ± 0.05
PWR Operating (containment)				
O-Upper Deck Overlooking Reactor Cavity (404-ft level)	136	223 195	0 -	$1.64 \pm 0.11$ $1.43 \pm 0.10$

TABLE 10. Expolation Chamber and Ion Chamber Measurements, Site B

All measurements for the shutdown reactor were performed at the basement level (336 ft), where sufficiently large dose rates were present. Effective  $C_x$  values were unity or less. The major contributor to dose was medium-energy photons. The values less than one were probably due to distributed sources. The source measured in the let-cown heat exchanger room (10.8 R/h) was well localized, giving the C factor expected for a Co and Cs source.

The extrapolation chamber measurement performed during reactor operation was taken from a location overlooking the reactor cavity. The neutron-to-photon ratio in dose equivalent was approximately 2:1. The correction to the dose and exposure was less than the uncertainty of the effective C measurement. The results indicate a 15% surface enhancement compared to the 1-cm

depth. There was no observed reduction of dose rate at the surface due to high-energy photons. The cause of the elevated magnitude of the measured  $C_{\chi}$  factors is not understood. The spectral measurements indicate that it cannot be explained by the presence of low-energy photons.

The TLD-loaded phantom measurements are presented in Table 11. The two measurements performed during reactor shutdown indicate a dominant contribution from medium-energy photons. Response versus depth is plotted in Figure 27. All measurements show an enhanced surface dose. The measurements performed during reactor operation are generally more penetrating.

TABLE 11. TLD-Loaded Phantom Measurements, Site B

1 1

	Field	Strengths	(mrad/h)	for	Effective
Surface Dose Rate (mrad/h)	34 keV	Photons 120 keV	662 keV	Beta	C (rað/R)
70	0	0	70	0	1.03
50	11	0	39	0	1.06
20	(ana	lysis ina	ppropriat	e)	
ar 40	(ana	lysis ina	ppropriat	e)	
	Surface Dose Rate (mrad/h) 70 50 20 ar 40	Surface Dose Rate (mrad/h) 34 keV 70 0 50 11 20 (ana ear 40 (ana	Field Strengths PhotonsSurface Dose Rate (mrad/h)Photons 34 keV 120 keV700070005011050110ar40(analysis inal analysis inal analysis inal analysis inal	Field Strengths (mrad/h)Surface Dose Rate (mrad/h)Field Strengths (mrad/h)34 keV120 keV662 keV700070501103920(analysis inappropriatar40(analysis inappropriat	Field Strengths (mrad/h) for PhotonsSurface Dose Rate (mrad/h)Field Strengths (mrad/h) for Photons34 keV120 keV662 keVBeta70007005011039050110390ar40(analysis inappropriate)

(a) Calculated for the shallow depth (7 mg/cm<sup>2</sup>).



FIGURE 27. TLD-Loaded Phantom Measurements, Site B

The calculated C factors ranged up to 1.07. The dominant contribution to the spectra was from cesium isotopes. The extrapolation chamber measurements resulted in C factors less than unity for the lower dose rates. The distribution of sources may lower the C factor below unity for most locations.

Measurements performed inside containment indicated a dominant contribution to dose from high-energy photons. However, both the extrapolation chamber and the TLD phantom measurements results in higher surface doses than depth c ses. No indications were found that per annel doses are being underestimated using current monitoring requirements

#### 3.4 SITE M - SHUTDOWN AND OPERATING BWR

The BWR at Site M was visited twice, with measurements taken under both shutdown and operating conditions (55% of full power). Measurements were performed at 29 locations ranging in exposure rate from 0.2 mR/h to 7.5 R/h.

The photon spectrometer measurements were performed with a  $31 \text{-cm}^3$  intrinsic Ge detector. The measurement locations are listed in Table 12. All measurements performed during shutdown and all those performed during operation except in the location near the turbine did not have a significant high-energy contribution. The maximum calculated C factors ranged from 1.02 to 1.16. Most values were very close to unity because the contributions to exposure were dominated by medium-energy photons. Photons from <sup>60</sup>Cc were the dominant contribution (Figure 28). The larger values were due to larger relative contributions from scattered photons. The data taken at the door of the cleanup phase separator room (Figure 29) shows an almost pure scatter spectrum. Photons from <sup>60</sup>Co scattered off a concrete wall before exiting through the door. While the spectrum peaks at about 120 keV, the higher-energy tail extends to above 500 keV. The relative importance of the medium-energy photons is enhanced by the flux-to-exposure conversion. The result is an effective C factor of 1.16.

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	Exposure Rate	Effect	ive Maxi rad/R	mum C <sub>x</sub> ,	Effective Central C <sub>x</sub> , rad/R			
Location	(mR/h)	0.007 cm	0.3 cm	1.0 cm	0.3 cm	1.0 cm	2.5 cm	
Shutdown BWR								
Reactor Building								
Refuel Pool, HEPA Filter Hose	4	0.98	1.02	1.02				
Refuel Pool, General Area(a)	0.4	1.01	1.04	1.04				
Dry Well, Valve	2	0.99	1.02	1.02				
Turbine Building								
Turbine Blade Housing Diaphram	0.2	1.02	1.05	1.05				
Operating BWR								
Reactor Building								
First Floor, Opposite Airlock	2.5	1.01	1.04	1.04				
First Floor, Outside RHR Valve Room	0.2	1.03	1.06	1.04				

TABLE 12. Photon Spectrometer Measurements, Site M

# TABLE 12. (contd)

	Exposure	Effecti	ve Maxi rad/R	mum C <sub>x</sub> ,	Effective Central C <sub>x</sub> , rad/R		
Location	(mR/h)	0.007 cm	0.3 cm	1.0 cm	0.3 cm	1.0 cm	2.5 cm
Operating BWR (contd)	1						
Reactor Building (cor	ntd)						
First Floor, Near Scram Discharge Lines	4	1.01	1.04	1.04			
Second Floor, Outside Spent Resin Room	11	1.01	1.03	1.04			
Second Floor, Near Clean-Up Phase Separator Room Door	1	1.15	1.15	1.15			
Second Floor, Near Clean-Up Phase Separator Room Do	4 oor	1.16	1.16	1.16			
Second Floor, Near Containment	5	1.01	1.04	1.04			
Third Floor, Near Jungle Room Door	4.5	1.02	1.05	1.05			
Fourth Floor, Laundry Area	1	1.02	1.05	1.05			
Fifth Floor, 0.6 m from Cavit	y 4	1.00	1.03	1.03			
Fifth Floor, 1 m from New Fue Storage	1 1.5	1.02	1.05	1.05			
Turbine Building Behind Shield Wall Near Turbine SE Corner of CO <sub>2</sub> Unit	2 14 0.3	0.83 0.81 0.96	0.88 0.86 1.01	0.94 0.93 1.01	0.38 0.32	0.82 0.79	0.99 0.92
Off-Gas Building Near Charcoal Absorbers	2	1.00	1.03	1.03			

(a) NaI(T1) data also taken at this location.





Near Clean-Up Phase Separator Room Door)

Spectra taken in the turbine room during operation show a dominant  ${}^{16}$ N peak at 6.1 MeV and the 511-keV annihilation photons. The presence of annihilation photons is expected because a major mechanism of energy loss at high energies is pair production. The  ${}^{16}$ N photons are approximately seven times more efficient in depositing dose than the 511-keV photons, and are therefore the major contributor to dose.

The effect of using the lead collimator was investigated by performing several measurements with and without the collimator. In general, the use of the collimator generated additional background counts at the low energies, which decreased the quality of the data. However, it was noted that the appearance of the spectra also changed. Figures 30 and 31 present data collected at the same location with and without using the lead collimator. The proportion of scattered photons is decreased compared to the <sup>60</sup> Co decay photons. Because the collimator allows photons to enter unattenuated only through a small opening, the conclusion is that the scatter is coming from a different direction than the <sup>60</sup> Co decay photons. If the scattered photons coming from different directions were added properly, the calculated C factor would probably be even lower. It has been observed using the extrapolation chamber that source geometry effects lower the C values in most locations.

The effect of the collimator in areas containing  $^{16}$ N-decay photons was to almost eliminate the 511-keV photons from the spectrum. This effect is in agreement with the interpretation that the 511-keV photons originate from all directions, dependent on the intensity of high-energy photons interacting in matter. The effect of this on the high-energy results is minimal, since the <sup>16</sup>N photons are the dominant component and the dosimetric measurements automatically account for geometry effects.

The results of measurements performed with the extrapolation and ion chambers are given in Table 13. Measurements performed in the high-pressure steam injection room (HPSI) and at the fuel pool heat exchanger resulted in effective C values at or below unity. The shallow-depth values were less than the 0.3-cm and the deep-depth values, indicating a nearly pure photon field consistent with the <sup>6</sup> Co energies. The deep-depth values less than unity are attributed to geometry effects. The measurement at the clean-up phase separator tank was taken with the chambers in direct view of the radiation source. The results are consistent with a dominant <sup>60</sup> Co photon field. An appropriate measurement location for the extrapolation chamber with a sufficiently large dose rate of high-energy photons was not found. The humidity and temperature were too high in the heater bay, and the dose rates were too low in the turbine room.



FIGURE 30. Field Data and Corrected Spectrum, Site M (Shutdown BWR, Refuel Pool General Area, Collimated)



FIGURE 31. Field Data and Corrected Spectrum, Site M (Shutdown BWR, Refuel Pool General Area, Uncollimated)

Location	Exposure Rate (mR/h)	Dose Rate (mrad/h)	Effective C_Factor X(rad/R)		
Shutdown BWR					
HPSI Shallow 0.3 cm Deep	2030	1770 2000 1940	$\begin{array}{c} 0.87 \pm 0.04 \\ 0.98 \pm 0.04 \\ 0.96 \pm 0.04 \end{array}$		
Operating BWR					
Fuel Pool Heat Exchanger Shallow 0.3 cm Deep	178	149 180 163	$\begin{array}{c} 0.84 \ \pm \ 0.08 \\ 1.01 \ \pm \ 0.07 \\ 0.92 \ \pm \ 0.04 \end{array}$		
Clean-Up Phase Separator Tank Deep	7350	7540	1.03 ± 0.03		

TABLE 13. Extrapolation Chamber and Ion Chamber Measurements, Site M

The TLD-loaded phantom measurements are described in Table 14. Effective C factors for locations not containing high-energy photons ranged from 1.03 to 1.15. The measurement at the clean-up phase separator door was made at the highest dose rate without opening the door and entering the room. The results are consistent with the photon spectrometer measurements. The measurement performed at the turbine blade housing diaphragm showed primarily a beta field and was not analyzed for C factor. The TLD response versus depth is presented in Figure 32 for the shutdown reactor measurements and Figure 33 for the operating reactor measurements.

The TLD phantom placed in the heater bay was directed up toward the pipes near the ceiling that carry steam to the turbine. This field was expected to contain a large contribution of  $^{16}$ N photons. A reduction in dose at the surface compared to the 2.5-cm depth was not observed. The depth-response curve has a surface enhancement of 30%.

The most significant effects of the presence of low-energy photons at Site M were found at the door to the clean-up phase separator room. The cause of the low energies was scatter from a suspected large concentration of  $^{60}$ Co in the clean-up phase separator tank. Due to the polyenergetic distribution of the scattered photons, the calculated maximum C factor calculated from the photon spectrum was 1.16. All other locations not containing energies above

	Surface Dose	Field	Strengths	for	Effective <sup>(a)</sup>	
Location	Rate (mrad/h)	34 keV	120 keV	662 keV	Beta	(rad/R)
Shutdown BWR						
Torus Drainline, HPSI	44	0	13	28	3	1.12
Turbine Blade Housing Diaphragm	162	0	0	8	154	-
Dry Well, Valve	183	0	13	162	8	1.05
Operating BWR						
Reactor Building Containment Equipment Stor- age Cage	20	0	1	18	1	1.05
Clean-Up Phase Separator Door	15	1	5	7	2	1.15
Skimmer Surge Task Room	43	0	0	38	5	1.03
Heater Bay	150					

TABLE 14. TLD-Loaded Phantom Measurements, Site M

(a) Shallow depth  $(7 \text{ mg/cm}^2)$ .

3 MeV had a large contribution from medium energy photons, primarily  ${}^{60}$ Co, which reduced the calculated C values. Large contributions to dose from high-energy photons were found. However, no surface reduction in dose was observed. Current monitoring requirements appear adequate for the high-energy photon fields encountered.

A spectrum was taken using a 2-in. x 2-in. NaI(T1) detector at the same location as one of the Ge spectra. The purpose was to demonstrate the utility of using the less-expensive detector for plant monitoring. Figure 34 presents a calibration and field measurement using a NaI(T1) detector. The resolution is poor, but the peaks are recognizable. Figure 31 showed the Ge spectrum in the same location. All of the peaks present in the Ge spectrum are present in the NaI(T1) spectrum. For spectral monitoring, the data analysis procedure would be the same for both detectors. The calibration spectrum provides efficiency and scatter signal versus energy.





FIGURE 34. Calibration and Field Data, NaI(T1) Detector

#### 3.5 SITE Q - OPERATING BWR

Measurements were performed at 12 locations including areas in the turbine building, the reactor vessel sampling station, the spent-fuel room, and the waste storage facility. Emphasis was placed on monitoring areas with dominant high-energy photon contributions. A 31-cm intrinsic Ge detector was used to measure spectra at the locations given in Table 15. The locations of the TLD phantom measurements are listed in Table 16. Extrapolation chamber measurements were not performed because the equipment was inoperable due to damage during shipment.

Measurement locations on the turbine floor of the turbine building are indicated in Figure 35. Locations included a high-rate area near the turbine (location E, approximately 900 mR/h), an area behind the bioshield (location D, 4 mR/h), and an area far from the turbine (location A, 4 mR/h). The photon spectra are dominated by high-energy photons, primarily 6.1-MeV photons from  $^{16}N$  (Figure 36). The analysis of the spectral data summarized in Table 15 includes the calculated maximum C factors and the calculated central C factors. Calculations for all turbine building locations except the laundry/ turbine loading area (location F) indicate approximately 40% and 90% of the maximum dose at 0.3 cm and 1 cm depths, respectively, for a pure photon field.

The underresponses predicted from the spectral data from the turbine building were not observed with the TLDs. The TLD phantom measurements were performed at locations C and E. The TLD phantom analysis using the linear least-squares technique was not reported because the set of basis energies were not representative of the radiations present. Figure 37 shows the depth-dose curves for these measurements. The data for both location C and location E show a surface enhancement rather than a surface underresponse. Additional measurements were made with multielement dosimeters to help characterize the surface response with location. These data, presented in Table 17, showed a surface enhancement, in approximate agreement with the TLD phantom results. Locations C and D have the greatest enhancement compared to locations E and B.

The enhancement shown in locations C and D extends to a tissue depth of approximately 1 cm. It could be the result of a range of electrons with an approximate maximum energy of 3 MeV or photons with an approximate energy of 15 keV. The observed response at depths greater than 1 cm implies penetrating photon energies, consistent with the presence of  $^{16}N$ .

TABLE 15. Measurement Locations for Photon Spectrometer, Site Q

	Exposure Rate	Exposure Effective Max Rate rad/			Effective Central C <sub>x</sub> ,		
Location	(mR/h)	0.007 cm	0.3 cm	1.0 cm	0.3 cm	1.0 cm	2.5 cm
Turbine Building							
A-Floor 272 Near Viewing Gallery	4	0.81	0.87	0.93	0.35	0.80	0.93
B-Floor 272 Behind Stairwell	10	0.82	0.88	0.94	0.38	0.82	0.94
C-Floor 272 NW Corner of Turbine	24	0.81	0.87	0.93	0.34	0.80	0.93
D-Floor 272 SW Corner Behind Shield Wall	4	0.81	0.87	0.93	0.35	0.81	0.93
F-Floor 248, Laundry/Turbine Loading Area	0.5	1.01	1.03	1.03			
G-Floor 248, Entrance #1 to Turbine Bldg.	1.5	0.82	0.88	0.94	0.38	0.81	0.93
H-Floor 248, Entrance #2 to Turbine Bldg.	5	0.81	0.87	0.94	0.37	0.81	0.92
I-Floor 248, General Area	0.7	0.82	0.88	0.94	0.39	0.86	0.93
Other -							
J-Reactor Vessel Sampling Station	4.5	1.05	1.06	1.05			
K-Spent-Fuel Room	3.3	0.83	0.88	0.94	0.36	0.84	0.93
L-Waste Storage Area	3.0	1.04	1.05	1.05			

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Location	Shallow Dose Rate (mrad/h)	Field St 34 keV	Photons 117 keV	(mrad/h) 662 keV	) for Beta	Effective C (b) (rad/R) <sup>x</sup>
C-Turbine Bldg., Floor 272	35	(ana	alysis not	appropr	iate)	
E-Turbine Bldg., Floor 272	910	(ana	alysis not	appropr	iate)	
J-Reactor Vessel Sampling Station	16	0	2	14	0	1.07

TABLE 16. TLD-Loaded Phantom Measurements, Site Q

(a) Maximum dose at the shallow tissue depth.

(b) Shallow depth (7 mg/cm<sup>2</sup>).

The photon spectra show that the dominant dose contributor is photons from <sup>1</sup>N. Only a relatively small contribution from medium-energy photons was observed (primarily 511-keV annihilation radiation). The surface enhancement could be due to one or a combination of several mechanisms. Possibilities are: 1) very low energy photons, 2) beta particles or electrons, and 3) the geometry of incident photons or electrons. These possibilities will be discussed in order.

Photons at energies below 20 keV are present, but not in the intensities required to produce a surface enhancement. Spectral data were taken for energies down to 15 keV for several locations. The large enhancement required to input the measured surface dose was not observed.

Nitrogen-16 emits beta particles at 4 MeV to 10 MeV, along with the photons. If <sup>16</sup>N atoms were being released into the atmosphere around the turbine, the associated beta particles would give a surface enhancement. An enhancement of about a factor of two would be present without shielding. However, this cannot be the only explanation for the enhancement seen, because location D, which is behind a shielding wall and should be preferentially protected from the beta particles, also has a surface dose enhancement.





Knock-on electrons exit all material through which photons pass. In the turbine room, this material includes both the generating equipment and shielding. The maximum electron energy is the same as the energy of the photons (approximately 6 MeV in the turbine room). The mean electron energy is much lower, about one-third of the photon energy. These electrons deposit 30 to 100 times more dose per particle than do photons of the same energy. Because the pair-production reaction for photons is dominant at 6 MeV, more electrons are generated in higher-atomic-number materials. The mass attenuation coefficient is 70% higher for lead than for tissue. Rogers (1983) investigated the effects of knock-on electrons in an accelerator-produced 6- to 7-MeV beam. Measurements performed with an ion chamber with Bakelite<sup>®</sup> walls resulted in surface enhancements from +33% for a lead filter to +15% for an aluminum filter. While the differences between the geometry of an open room and that of a collimated beam could affect results significantly, the values are the right order of magnitude to explain the observed surface enhancements.

Bakelite is a registered trademark of Union Carbide Corp., Plastics Division.



(Turbine Floor 272, Behind Shield Wall)



FIGURE 37. TLD-Loaded Phantom Measurements, Site Q

TABLE 17. Relative Response for Multielement Dosimeters, Site Q

location	Exposure Rate	Relative TLD Response Versus Effective Filtration (mg/cm <sup>2</sup> )					
Location	(mR/h)	0.12	0.26	0.34	0.98	1.12	
B-Turbine Bldg., Floor 272	10	1.00	0.97	0.96	1.03	0.93	
D-Turbine Bldg., Floor 272	4	1.00	0.96	0.90	0.82	0.78	
E-Turbine Bldg., Floor 272	900	1.00	1.00	0.99	0.93	0.90	

The direction of the incident photons or electrons affects the depth-dose distribution by depositing dose with depth along a path oblique to the front surface of the phantom or dosimeter. This effect would probably be greatest for locations C and D (close to the scatter source, but shielded from the primary source) and least for location B (far from the primary and scatter sources) and location E (close to the primary source). This effect agrees with the observations of greater enhancements at locations C and D. However, because dose deposited by the scattered continuum and the 511-keV photons is small compared to the N photons, the knock-on electrons exiting all surface are probably the greatest contributors to this effect.

Locations F, J, and L had a negligible contribution from high-energy photons. These locations had a dominant contribution from <sup>6</sup>Co, resulting in effective C factors near unity. The TLD data for location J agreed with the spectrometer data.

#### 3.6 SITE P - SHUTDOWN PWR

Measurements were performed at 10 locations, including areas in the waste storage facility, the spent-fuel pit, the auxiliary building, and containment. The locations of measurements performed with the photon spectrometer (31-cm intrinsic Ge detector) are listed in Table 18 along with the exposure rates and calculated effective C factors. The TLD phantom measurements are summarized in Table 19 and Figure 38. The exposure rates in all areas monitored were relatively small (<20 mR/h). No higher rates were located in accessible areas.

The spectra show the dominance of  $^{60}$ Co and the presence of several other radionuclides emitting primarily medium-energy photons. No areas were found with dominant contributions from low-energy photons. The calculated effective C\_ factors range from 1.04 to 1.06.

The TLD phantom data also indicated that the primary spectral components are medium-energy photons, represented by the 662-keV component. The effective C values are higher for this anlaysis than for the analysis of the photon spectral data, but they are also less precise. Only a relatively small beta-field component was observed. The depth-response curves show somewhat erratic behavior, but are consistent with the above interpretation.

The data indicate that the effective  $C_x$  values are not significantly enhanced for Site P. No locations with unusual photon spectra were found.

	Exposure Rate	Effective Maximum C <sub>x</sub> (rad/R)			
Location	(mR/h)	0.007 cm	0.3 cm	1.0 cm	
Waste Storage Area					
A-Barrell Storage (outside Waste Disposal Bldg.)	3.0	1.05	1.05	1.05	
B-Compactor Area (outside Waste Disposal Bldg.)	2.0	1.06	1.05	1.05	
C-Waste Disposal Bldg.	2.7	1.06	1.06	1.06	
Spent-Fuel Pit					
D-Heat Exchanger, Spent Fuel Pit	4.4	1.05	1.05	1.05	
E-Ion Exchange Pit	2.4	1.05	1.05	1.05	
F-Fuel Transfer Shoot	3.9	1.06	1.06	1.06	
G-Spent-Fuel Pit	2.2	1.06	1.06	1.06	
Auxiliary Building	*				
H-Primary Auxiliary Bldg., General Area	0.7	1.05	1.05	1.05	
I-Open Surge Line	2.5	1.06	1.06	1.06	
Containment					
J-Above Reactor Head Area (Flooded)	2.0	1.04	1.04	1.04	

TABLE 18. Measurement Locations for Photon Spectrometer, Site P

### 3.7 SITE N - OPERATING BWR

Measurements were performed at 13 locations during operation (78% of capacity). Emphasis was placed on the presence of high-energy photons, particularly due to  $^{16}\mathrm{N}$ . Areas monitored included the heater bay, the main steam isolation valve room (MSIV), the turbine area, and the clean-up phase separator area.

	Shallow Dose Rate (mrad/h)	Field Strength <sup>(a)</sup> (mrad/h) for				(b)
Location		34 keV	Photon 117 keV	662 keV	Beta	Effective C (b) (rad/R) <sup>x</sup>
C-Waste Dis- posal Building	2.8	0	0.7	1.7	0.4	1.13
F-Fuel Transfer Shoot	16	0	6	10	0	1.16

TABLE 19. TLD-Loaded Phantom Measurements, Site P

(a) Maximum dose at the shallow tissue depth.(b) Shallow depth (0.007 cm).



FIGURE 38. TLD-Loaded Phantom Measurements, Site P

Significant contributions to dose due to <sup>16</sup>N photons were found in the heater bay, the MSIV, and near the turbine (Figure 39). The calculated central C factors predict surface underresponses based on the photon field only (Table 20). Dosimetric data were taken in these areas to measure dose at the surface compared to dose at other depths. The results of the extrapolation and ion chamber measurements made in the MSIV are contained in Table 21. The dose measured at the surface was greater than the dose at other depths. The results for the depths 0.3 cm, 1.0 cm, and 2.5 cm were essentially the same.

The TLD-loaded phantom measurements were performed in the heater bay, the MSIV, and at two locations in the turbine room (Table 22). All locations showed an enhanced dose rate at the surface (Figure 40). The measurement for the MSIV was performed in approximately the same location as the extrapolation and ion chamber measurements, with consistent results.

The TLD phantom measurements were compared to the results of the dosimetry methods used at the plant (Table 22). The exposure rate at the phantom position was checked with a survey meter. Pocket ionization chambers and personnel dosimeters (film badges) were mounted on the front face of the phantom for timed exposures. The measured surface dose rates were higher than the survey meter readings, which is consistent with enhanced surface doses. The pocket ionization chamber and the Site N personnel dosimeter overresponded by 12% and 49%, respectively. The overresponse of the film badges was probably due to the increased response at 6 MeV for the high-atomic-number material in the film emulsion and the filter packet.

These measurements do not indicate an inadequacy in the current personnel monitoring requirements. The performance of personnel dosimeters manufactured with high-atomic-number material may be poor, but they will most likely over-respond to the high-energy photon fields.


FIGURE 39. Field Data and Corrected Spectrum, Site N, Location H (Operating BWR, Turbine Room, with Collimator)

	Exposure Rate	Maximum Effective C <sub>x</sub> (rad/R)		Central Effective C <sub>x</sub> (rad/R)			
Location	(mR/h)	0.007 cm	0.3 cm	1.0 cm	0.3 cm	1.0 cm	2.5 cm
Operating BWR							
A-Heater Bay, Entrance	20	0.89	0.93	0.97	0.53	0.88	0.97
B-Heater Bay, Near Steam Lines	30	0.83	0.88	0.94	0.38	0.84	0.94
C-MSIV, Entrance Hallway	23	0.82	0.87	0.94	0.35	0.81	0.93
D-CRD Room	48	0.98	1.01	1.02			
E-Storage Room, Contaminated Pipe	60	0.99	1.02	1.02			
F-Turbine Floor, Outside Shield Wall	0.6	1.01	1.04	1.06	0.79	1.01	1.06
G-Turbine Floor, Maze Entrance to Turbine Room	6	0.81	0.87	0.93	0.33	0.79	0.93
H-Turbine Floor, Inside Turbine Room	42	0.81	0.87	0.93	0.33	0.80	0.93
I-Clean-Up Phase Separator Room Door	7	1.07	1.08	1.08			

TABLE 20. Photon Spectrometer Measurements, Site N

# TABLE 21. Extrapolation Chamber and Ion Chamber Measurements, Site N

Location	Exposure Rate	Dose Rate	Effective C_ Factor
	(mR/h)	(mrad/h)	(rad/Ř)
J-MSIV Shallow 0.3 cm Deep 2.5 cm	338	372 270 277 264	1.10 ± 0.04 0.80 ± 0.15 0.82 ± 0.06 0.78 ± 0.04

Location	Surface Dose Rate (mrad/h)	Survey Meter (mR/h)	Pocket Ionization Chamber (mR/h)	Site N Persoanel Dosimeter (mrem/h)
K-Heater Bay	106	80	130	152
J-MSIV	395	330	450	508
L-Turbine Room	84	70	83	123
M-Turbine Room	246	220	280	434

TABLE 22. TLD-Loaded Phantom Measurements, Site N



FIGURE 40. TLD-Loaded Phantom Measurements, Site N

#### 4. DISCUSSION

The discussion section has three parts covering the impact of improved C factors, high-energy photon dosimetry, and recommendations for plant monitoring procedures.

### 4.1 IMPACT OF IMPROVED C FACTORS

The areas monitored can be approximately classified into four categories: 1) radiation fields dominated by decay photons from radioactive atoms in neutron-activated or fission-product deposits; 2) radiation fields dominated by a scattered-photon continuum; 3) radiation fields containing short-lived radioactive noble gases; and 4) radiation fields dominated by high-energy photons. High-energy photons have C factors less than unity and will be discussed in section 4.2. The remaining three categories will be discussed here in turn.

Dose rates in most plant areas are dominated by lightly shielded radioactive sources in neutron-activated or fission-product deposits. The published C, values for the decay photons from these sources range from 1.00 rad/R to 1.04 rad/R (parallel geometry). Calculated C, factors for these areas are, at most, a few percent higher after the addition of contributions from the scatter continuum.

The energies comprising the scatter continuum vary with location. The maximum photon flux is at approximately 120 keV, with a half-maximum range between about 70 keV and 250 keV. A higher-energy tailing to above 500 keV skews the distribution, lowering the cumulative C factor, for most locations. The average C factor for the scatter continuum is no more than 1.2. Averaging the decay photons from long-half-life sources with the scatter continuum yields calculated C values between 1.00 and 1.10.

Only one location, near the clean-up phase separator door at Site M (BWR), was identified as being dominated by low-energy photons. The dominance of low-energy photons is present whether the plant is operating or shutdown. A similar location was monitored at Site N. However, sufficient radioactive cesium and cobalt were in the immediate vicinity to reduce the calculated C value to below 1.10.

The areas with significant radioactive ("crud") buildup will not have elevated C factors. Clean well-shielded areas are more likely to have both elevated  $C_x$  factors and low dose rates. Two locations that were monitored had

exposure rates less than 0.1 mR/h: the worker locker room at Site M and the gatehouse at Site Q. The resulting C factors for these locations were 1.17 rad/R and 1.11 rad/R, respectively. While these areas will never contribute a significant amount to occupational exposure, they do illustrate the effect for uncontaminated, highly shielded areas.

Radioactive noble gases were detected inside of containment at operating PWRs. The noble gases permeate the atmosphere, giving isotropic incidence of decay photons. For isotropic incidence, only the shallow-depth C factors are significantly greater than unity for low-energy photons. Most photons are attenuated by the ICRU sphere (or the worker) before reaching greater depths. Because the efficiency of flux-to-dose conversion favors the higher-energy photons, the presence of photons near or greater than 1 MeV overshadows the effects of the 81-keV and 249-keV photons from the xenon isotopes. Even spectra taken in personnel hatches had sufficient higher-energy contributions to keep the calculated C values near unity.

As was noted several times during the description of the site measurements, at many locations the source-decay photons and the scatter continuum originate from different locations. If the locations were all in one general direction, the shallow-dose  $C_x$  factors would be little affected. However, if they were coming from opposite directions, the  $C_x$  factors would be below unity. The extrapolation chamber measurements indicated that the most likely case is for distributed sources and for  $C_x$  factors less than unity.

Average reported doses will not be substantially affected by the use of improved C factors. The majority of worker exposures result from the higherexposure-rate locations. These locations are more likely to have dominant localized sources or distributed sources. The impact of changing the effective C values from a defined value of 1 to the actual values would not be detected among larger dosimetric uncertainties, such as changes in the response of personnel dosimeters to meet requirements in ANSI N13.11. Even dosimeters with dramatic improvements in low-energy response would not noticeably alter reported doses (<5% change). The exception would be deliberately altering the response to Cs to achieve an overall acceptable response at low energies. This change would shift the reported doses almost in direct proportion to the response shift. Such a change is potentially serious because the response of a dosimeter to Cs can be lowered by as much as 40% and still meet ANSI N13.11 criteria.

The desired accuracy of the reported dose for each worker may require special plant monitoring to be performed. The guidance provided by the International Commission on Radiation Units and Measurements (ICRU) Report 20 (ICRU 1971) and the National Council on Radiation Protection and Measurements (NCRP) Report No. 57 (NCRP 1978) is that reported doses be accurate to within 30% near maximum permissible levels. While the effects of C factors are less than 30%, it is desirable to keep each contributory effect much less than the maximum. If we chose to correct the dose if the difference were greater than one-third of the recommendation, or 10%, only one of the measurement locations investigated during this study would be identified as definitely requiring a correction. Seven other locations at four sites would be classified as marginal (calculated  $C_x > 1.08$  or TLD-measured  $C_y > 1.10$ ).

#### 4.2 HIGH-ENERGY PHOTON DOSIMETRY

Site measurements of high-energy (>3-MeV) radiation fields were performed to help determine the adequacy of current personnel monitoring requirements. The greatest depth required for monitoring is 1 cm of tissue. Because pure high-energy photon fields (3 to 10 MeV) deposit the maximum dose at depths between 2 and 4 cm of tissue, current practices could be underestimating the maximum dose received.

Measurements were performed at three operating PWRs and three operating BWRs. Both reactor types had locations with dominant high-energy photon contributions. Inside containment at operating PWRs, there were significant high-energy photons up to 8 MeV along with contributions from low- and medium-energy photons. The predicted reduction in the measured dose at 1 cm compared to the actual maximum dose was small (approximately 10%). In turbine rooms at the BWRs approximately 80% of the dose deposited was due to highenergy photons from <sup>10</sup>N. The other major contributor was 511-keV annihilation radiation. Predicted maximum reductions in dose at 1 cm were as high as 20%.

Dosimetric measurements were peformed at all operating reactor sites. In no location was the predicted reduction in dose at depths of 1 cm and less observed. Either no enhancement or else a large (approximately 40%) enhancement of the surface dose was measured.

Measurements performed at Site Q were designed to investigate the origin of the enhanced surface dose. The only plausible explanation was the presence of knock-on electrons created by the high-energy photons interacting with all matter in the vicinity of the measurement. Electrons generated in a tissuelike material (low atomic number) would have an intensity similar to those generated inside of the phantoms. Electrons generated by material with a higher atomic number would have a greater intensity because of the atomicnumber dependence of the pair production interaction. The observed surface enhancements were probably created by the presence of iron in the turbine areas or high-atomic-number shielding components (e.g., lead). Oblique incidence of the electrons and the 511-keV photons also increased the surface dose.

Many personnel dosimeters are designed with high-atomic-number filters to achieve responses like the 1-cm depth in tissue in a compact space. These filters are typically designed and calibrated to meet the dosimetry requirements of <sup>60</sup>Co and <sup>137</sup>Cs sources. However, because pair production is the dominant interaction at high energies, these personnel dosimeters would overrespond to 6-MeV photons. Film emulsion manufactured with high-atomic-number materials would also overrespond. This effect was observed at Site Q.

#### 4.3 RECOMMENDATIONS FOR PLANT MONITORING PROCEDURES

The greatest contribution to the difference between exposure in air and dose in phantom (or worker) is the photon backscatter created in the phantom. If a perfectly calibrated personnel dosimeter were placed on a phantom (or worker) in a low-energy photon field, it would provide an accurate dose estimate. Correction factors would only be required to compensate for a poor energy response. Both the proposed modifications to 10 CFR 20 and ANSI N13.11 (1983) encourage the use of personnel dosimeters with improved low-energy response. Exposure-rate measurements performed with icn chamber instruments require C<sub>2</sub> factors to convert to dose rates.

Correction factors other than  $C_x$  factors may be more useful for particular applications. The energy response of an instrument or dosimeter can be used to generate correction factors for response to dose as easily as to generate factors for exposure to dose.

Quick plant surveys can be performed using photon spectrometers. A Ge, Ge(Li), or NaI(Tl) detector is adequate provided that the efficiency and response function are determined. The use of a collimator will increase the measurement range. Dosimetric measurements should be performed in areas with unusual spectra or with high dose rates. The response of personnel monitoring instruments and dosimeters should be determined by direct measurement in these locations. Correction factors may be required for jobs performed in only a few locations (if any).

The potential for inaccurate dosimetry is greater for high-energy fields than for low-energy fields. Dosimeters manufactured with high-atomic-number material for element filters and film dosimeters may overrespond by as much as 60% due to differences in the cross sections for pair production. However, these dosimeters always provide conservative dose estimates.

#### 5. CONCLUSIONS

No locations containing primarily low-energy photons with large C factors (approximately 1.5 rad/R) were found during measurements at seven operating and shutdown reactors. The most significant production of low-energy photons at commercial nuclear reactors is due to scattering in shielding material. The effective C factor for the scattered-photon continuum is no more than 1.2 rad/R. Most locations have radiation fields of nearly all medium-energy photons due to radioactive decay of cobalt and/or cesium isotopes or a combination of medium-energy photons with a scatter continuum. The estimated C factors for these locations, assuming parallel incidence of the photons, ranged between 1.00 rad/R and 1.10 rad/R. Directionality measurements (made using a collimator) and dosimetric measurements indicated that most locations have distributed sources, implying effective C factors less than unity. Only one location was found with a spectrum composed almost entirely of scattered photons and a C factor of 1.16 rad/R.

Monitoring requirements at 0.007-cm and 1.0-cm depths in tissue were found to be adequate for estimating the dose received in radiation fields at commercial nuclear reactors containing high energy photons. High-energy fields were found to dominate dose received in containment of operating PWRs and in the turbine areas and heater bays of BWRs. Surface doses probably due to accompanying high-energy electrons were measured in all locations monitored. The presence of the electrons was attributed to production processes associated with the high-energy photon field interacting with all material in the vicinity. The excess surface dose observed in severa? locations was attributed to the interaction of the high-energy photons in atoms with high atomic numbers and to oblique incidence of the electrons. Personnel dosimeters using high-atomicnumber filters and film dosimeters are predicted to overrespond in areas dominated by high-energy photons.

We recommend that the response of current personnel dosimeters to highenergy photon fields be investigated. Due to the widespread use of highatomic-number materials for penetrating-radiation filters in personnel dosimeters, the overresponse problem is potentially significant. High-energy photon fields (>3 MeV) were not included in ANSI N13.11, yet are a major contributor to dose received at many locations in operating plants. Current and new dosimeter designs which meet specifications in ANSI N13.11 may respond poorly to high-energy fields.

The analysis of photon spectra performed here had considerable uncertainty for high-energy fields. The causes were the large efficiency corrections

required for the Ge detectors and the approximate scatter corrections applied. We recommend that further attention be given to data collection and analysis techniques for the high-energy fields. Improvements in the accuracy of reported doses will probably require accurate dosimetric measurements along with dosimeter redesign or the development of correction factors through accurate plant surveys.

#### 6. REFERENCES

- American National Standards Institute (ANSI). 1983. <u>American National</u> <u>Standard Criteria for Testing Personnel Dosimetry Performance</u>. ANSI N13.11-1983, New York, New York.
- Code of Federal Regulations. Title 10, Part 20, "Standards for Protection Against Radiation." U.S. Government Printing Office, Washington, D.C.
- Dimbylow, P. J., and T. M. Francis. 1979. <u>A Calculation of the Photon</u> <u>Depth-Dose Distributions in the ICRU Sphere for a Broad Parallel Beam, A</u> <u>Point Source and an Isotropic Field</u>. NRPB 92, National Radiological Protection Board, Harwell, England.
- Endres, G. W. R., et al. 1983. <u>Neutron Dosimetry at Commercial Nuclear</u> <u>Plants. Final Report of Subtask A: Reactor Containment Measurements.</u> NUREG/CR-1769, PNL-3585, Pacific Northwest Laboratory, Richland, Washington.
- Fenyves, E., and Q. Haiman. 1969. <u>The Physical Principles of Nuclear</u> Radiation Measurements. Academic Press, New York.
- Hajnal, F., and C. Klusek. 1974. "Semi-Empirical Efficiency Equations for Ge(Li) Detectors." NIM 122:559-565.
- International Commission on Radiation Units and Measurements (ICRU). 1971. Radiation Protection Instrumentation and Its Application. ICRU Report 20. ICRU Publications, Washington, D.C.
- Johns, H. E., and J. R. Cunningham. 1978. <u>The Physics of Radiology</u>. 3rd ed. Charles C. Thomas Publisher, Springfield, Illinois.
- National Council on Radiation Protection and Measurements (NCRP). 1978. Instrumentation and Monitoring Methods for Radiation Protection. NCRP Report No. 57, Washington, D.C.
- Nelson, R. F., and A. B. Chilton. 1983. Low-Energy Photon Dose Deposition in Tissue Slab and Spherical Phantoms. NUREG/CR-3425, University of Illinois.
- Rathbun, L. A., and P. L. Roberson. 1983. <u>Beta Particle Measurement and</u> <u>Dosimetry Requirements at NRC-Licensed Facilities</u>. NUREG/CR-3544, PNL-4886, Pacific Northwest Laboratory, Richland, Washington.
- Rogers, D. W. O. 1983. "A Nearly Monoenergetic 6-7 MeV Photon Calibration Source." Health Phys. 45(1):127-137.

- Seelentag, W. W., and W. Panzer. 1979. "Stripping of X-Ray Bremsstrahlung Spectra Up to 300 kVp on a Desk Type Computer." <u>Phys. Med. Biol</u>. 24(4):767-780.
- Seltzer, S. M. 1981. "Calculated Response of Intrinsic Germanium Detectors to Narrow Beams of Photons with Energies up to ~300 keV." <u>NIM 188</u>:133-151.
- U.S. Department of Health, Education, and Welfare (DHEW). 1970. <u>Radiological</u> Health Handbook. U.S. Government Printing Office, Washington, D.C.
- U.S. National Bureau of Standards (NBS). 1983. <u>Calibration and Related</u> <u>Measurement Services at the National Bureau of Standards</u>. NBS Special Publication 250, Appendix.
- Yoder, R. C., et al. 1979. <u>Confirmation of Conversion Factors Relating</u> <u>Exposure and Dose-Equivalent Index Presented in ANSI N13.11</u>. NUREG/CR-1057, PNL-3219, Pacific Northwest Laboratory, Richland, Washington.

APPENDIX A PHOTON SPECTRA

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#### APPENDIX A

#### PHOTON SPECTRA

The spectra collected during the site visits are presented here. Indexes showing the figure numbers and measurement locations for each site are given in Tables A.1 through A.6. The probable identification of radioactive-isotope photon lines are labelled on the pulse-height distributions, which were collected using germanium detectors. Photon lines not labelled were not identified. Corrected spectra are presented below the pulse-height distributions. The effective C. factors labelled on the energy spectra are the maximum calculated values assuming parallel incidence for all photons except photons from radioactive noble gases, for which isotropic incidence was assumed.

# TABLE A.1. Site K - Shutdown and Operating PWR

FIGURE	LOCATION	PAGE
Shutdown PWR		
A.1.1	A-Effluent Sample Room	A.8
A.1.2	B-Auxiliary Demineralizer Room, Reactor Coolant Filter	A.9
A.1.3	C-Containment, Reactor Shield Vent Fan	A.10
A.1.4	D-Containment, Iodine Removal Fan	A.11
A.1.5	E-Containment, Under Reactor Coolant Pump #13 (collimated)	A.12
A.1.6	F-Containment, Under Reactor Coolant Pump #11 (collimated)	A.13
Operating PWR		
A.1.7	G-Effluent Sample Room	A.14
A.1.8	H-Auxiliary Building Demineralizer Room, Reactor Coolant Filter	A.15
A.1.9	I-Containment, Personnel Hatch	A.16
A.1.10	J-Containment, Near Bioshield of Steam Generator	A.17
A.1.11	K-Containment, Overlooking Reactor Cavity	A.18
A.1.12	L-Containment, Outside Airlock	A.19
A.1.13	M-Hot Drum Storage Area (collimated)	A.20

### TABLE A.2. Site B - Shutdown and Operating PWR

FIGURE	LOCATION	PAGE
Shutdown PWR		
A.2.1	A-Near Escape Hatch, 357-ft Level	A.21
A.2.2	B-Near Equipment Hatch, 357-ft Level	A.22
A.2.3	C-Near Elevator, 357-ft Level	A.23
A.2.4	D-Near Elevator, 373-ft Level	A.24
A.2.5	E-Near Elevator, 401-ft Level	A.25
A.2.6	F-Deck, 424-ft Level	A.26
A.2.7	G-Above Escape Hatch, 373-ft Level	A.27
Operating PWR		
A.2.8	H-Near Personnel Hatch, 386-ft Level	A.28
A.2.9	I-Near Stairway, 386-ft Level	A.29
A.2.10	E-Near Elevator, 401-ft Level (collimated)	A.30
A.2.11	J-In Personnel Hatch	A.31

# TABLE A.3. Site M - Shutdown and Operating BWR

FIGURE	LOCATION	PAGE
Shutdown BWR		
A.3.1	Refuel Pool, HEPA Filter Hose (collimated)	A.32
A.3.2	Refuel Pool, General Area	A.33
A.3.3	Dry Well, Valve (collimated)	A.34
A.3.4	Turbine Blade Housing Diaphragm	A.35
Operating BWR		
A.3.5	Reactor Building, First Floor, Opposite Airlock	A.36
A.3.6	Reactor Building, First Floor, Jutside RHR Valve Room	A.37
A.3.7	Reactor Building, Near Scram Discharge Lines	A.38
A.3.8	Reactor Building, Second Floor, Outside Spent Resin Room	A.39
A.3.9	Reactor Building, Second Floor, Near Clean-Up Phase Separator Room Door	A.40
A.3.10	Reactor Building, Second Floor, Near Clean-Up Phase Separator Room Door	A.41
A.3.11	Reactor Building, Second Floor, Near Containment	A.42
A.3.12	Reactor Building, Third Floor, Near Jungle Room Door	A.43
A.3.13	Reactor Building, Fourth Floor, Laundry Area	A.44
A.3.14	Reactor Building, Fifth Floor, 0.6 m from Cavity	A.45
A.3.15	Reactor Building, Fifth Floor, 1 m from New Fuel Storage	A.46
A.3.16	Turbine Building, Behind Shield Wall	A.47
A.3.17	Turbine Building, Near Turbine (collimated)	A.48
A.3.18	Turbine Building, SE Corner of CO2 Unit	A.49
A.3.19	Off-Gas Building, Near Charcoal Absorbers	A.50

# TABLE A.4. Site Q - Operating BWR

FIGURE	LOCATION	PAGE
	Turbine Building	
A.4.1	A-Floor 272, Near Viewing Gallery	A.51
A.4.2	B-Floor 272, Behind Stairwell	A.52
A.4.3	C-Floor 272, NW Corner of Turbine (collimated)	A.53
A.4.4	D-Floor 272, SW Corner Behind Shield Wall	A.54
A.4.5	F-Floor 248, Laundry/Turbine Loading Area	A.55
A.4.6	G-Floor 248, Entrance #1 to Turbine Building	A.56
A.4.7	H-Floor 248, Entrance #2 to Turbine Building	A.57
A.4.8	I-Floor 248, General Area	A.58
	Other	
A.4.9	J-Reactor Vessel Sampling Station	A.59
A.4.10	K-Spent Fuel Room	A.60
A.4.11	L-Waste Storage Area	A.61

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TADLE A.J. SILE P - SHULDOWN	TABLE A.	5. Si	te P -	Shutdown	PWR
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FIGURE	LOCATION		
	Waste Storage Area		
A.5.1	A-Barrel Storage (Outside Waste Disposal Building)	A.62	
A.5.2	B-Compactor Area (Outside Waste Disposal Building)	A.63	
A.5.3	C-Waste Disposal Building	A.64	
	Spent Fuel Pit		
A.5.4	D-Heat Exchange, Spent Fuel Pit	A.65	
A.5.5	E-Ion Exchange Pit	A.66	
A.5.6	F-Fuel Transfer Shoot	A.67	
A.5.7	G-Spent Fuel Pit	A.68	
	Auxiliary Building		
A.5.8	H-Primary Auxiliary Building, Ceneral Area	A.69	
A.5.9	I-Open Surge Line	A.70	
	Containment		
A.5.10	J-Above Reactor Head Area (Flooded)	A.71	

# TABLE A.6. Site N - Operating BWR

FIGURE	LOCATION	PAGE
A.6.1	A-Heater Bay, Entrance	A.72
A.6.2	B-Heater Bay, Near Steam Lines (collimated)	A.73
A.6.3	C-MSIV, Entrance Hallway (collimated)	A.74
A.6.4	D-CRD Room (collimated)	A.75
A.6.5	E-Storage Room, Contaminated Pipe (collimated)	A.76
A.6.6	F-Turbine Floor, Outside Shield Wall	A.77
A.6.7	G-Turbine Floor, Maze Entrance to Turbine Room	A.78
A.6.8	H-Turbine Floor, Inside Turbine Room (collimated)	A.79
A.6.9	I-Clean-Up Phase Separation Room Door	A.80



FIGURE A.1.1. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Site K, Location A-Effluent Sample Room



FIGURE A.1.2. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Site K, Location B-Auxiliary Building Demineralizer Room, Reactor Coolant Filter



FIGURE A.1.3. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Site K, Location C-Containment, Reactor Shield Vent Fan



FIGURE A.1.4. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Site K, Location D-Containment, Iodine Removal Fan



FIGURE A.1.5. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Site K, Location E-Containment, Under Reactor Coolant Pump #13 (collimated)



FIGURE A.1.6. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Site K, Location F-Containment, Under Reactor Coolant Pump #11 (collimated)



FIGURE A.1.7. Pulse-Height and Photon Energy Distributions, Operating PWR, Site K, Location G-Effluent Sample Room



FIGURE A.1.8. Pulse-Height and Photon Energy Distributions, Operating PWR, Site K, Location H-Auxiliary Building Demineralizer Room, Reactor Coolant Filter



FIGURE A.1.9. Pulse-Height and Photon Energy Distributions, Operating PWR, Site K, Location I-Containment, Personnel Hatch



FIGURE A.1.10. Pulse-Height and Photon Energy Distributions, Operating PWR, Site K, Location J-Containment, Near Bioshield of Steam Generator



FIGURE A.1.11. Pulse-Height and Photon Energy Distributions, Operating PWR, Site K, Location K-Containment, Overlooking Reactor Cavity



FIGURE A.1.12. Pulse-Height and Photon Energy Distributions, Operating PWR, Site K, Location L-Containment, Outside Airlock



FIGURE A.1.13. Pulse-Height and Photon Energy Distributions, Site K, Operating PWR, Location M-Hot Drum Storage Area (collimated)



FIGURE A.2.1. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Site B, Location A-Near Escape Hatch, 357-ft Level



FIGURE A.2.2. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Site B, Location B-Near Equipment Hatch, 357-ft Level



FIGURE A.2.3. Pulse-Height and Photon Energy Distributions, Shutdown PWR. Site B, Location C-Near Elevator, 357-ft Level



FIGURE A.2.4. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Site B, Location D-Near Elevator, 373-ft Level


FIGURE A.2.5. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Site B, Location E-Near Elevator, 401-ft Level



FIGURE A.2.6. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Site B, Location F-Deck, 424-ft Level



FIGURE A.2.7. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Site B, Location G-Above Escape Hatch, 373-ft Level



FIGURE A.2.8. Pulse-Height and Photon Energy Distributions, Operating PWR, Site B, Location H-Near Personnel Hatch, 386-ft Level



FIGURE A.2.9. Pulse-Height and Photon Energy Distributions, Operating PWR, Site B, Location I-Near Stairway, 386-ft Level



FIGURE A.2.10. Pulse-Height and Photon Energy Distributions, Operating PWR, Site B, Location E-Near Elevator, 401-ft Level (collimated)



FIGURE A.2.11. Pulse-Height and Photon Energy Distributions, Operating PWR, Site B, Location J-In Personnel Hatch



FIGURE A.3.1. Pulse-Height and Photon Energy Distributions, Shutdown BWR, Site M, Refuel Pool, HEPA Filter Hose (collimated)



FIGURE A.3.2. Pulse-Height and Photon Energy Distributions, Shutdown BWR, Site M, Refuel Pool, General Area

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FIGURE A.3.3. Pulse-Height and Photon Energy Distributions, Shutdown BWR, Site M, Dry Well, Valve (collimated)



FIGURE A.3.4. Pulse-Height and Photon Energy Distributions, Shutdown BWR, Site M, Turbine Blade Housing Diaphragm



FIGURE A.3.5. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Reactor Building, First Floor, Opposite Airlock



FIGURE A.3.6. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Reactor Building, First Floor, Outside RHR Valve Room



FIGURE A.3.7. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Reactor Building, Near Scram Discharge Lines



GURE A.3.8. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Reactor Building, Second Floor, Outside Spent Resin Room



FIGURE A.3.9. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Reactor Building, Second Floor, Near Clean-Up Phase Separator Room Door



FIGURE A.3.10. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Reactor Building, Second Floor, Near Clean-Up Phase Separator Room Door



FIGURE A.3.11. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Reactor Building, Second Floor, Near Containment



FIGURE A.3.12. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Reactor Building, Third Floor, Near Jungle Room Door



FIGURE A.3.13. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Reactor Building, Fourth Floor, Laundry Area



FIGURE A.3.14. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Reactor Building, Fifth Floor, 0.6 m from Cavity



FIGURE A.3.15. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Reactor Building, Fifth Floor, 1 m from New Fuel Storage



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FIGURE A.3.16. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Turbine Building, Behind Shield Wall



FIGURE A.3.17. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Turbine Building, Near Turbine (collimated)



FIGURE A.3.18. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Turbine Building, SE Corner of CO<sub>2</sub> Unit



FIGURE A.3.19. Pulse-Height and Photon Energy Distributions, Operating BWR, Site M, Off-Gas Building, Near Charcoal Absorbers



FIGURE A.4.1. Pulse-Height and Photon Energy Distributions, Operating BWR, Turbine Building, Site Q, Location A-Floor 272, Near Viewing Gallery



FIGURE A.4.2. Pulse-Height and Photon Energy Distributions, Operating BWR, Turbine Building, Site Q, Location B-Floor 272, Behind Stairwell



FIGURE A.4.3. Pulse-Height and Photon Energy Distributions, Operating BWR, Turbine Building, Site Q, Location C-Floor 272, NW Corner of Turbine (collimated)



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FIGURE A.4.4. Pulse-Height and Photon Energy Distributions, Operating BWR, Turbine Building, Site Q, Location D-Floor 272, SW Corner Behind Shield Wall



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FIGURE A.4.5. Pulse-Height and Photon Energy Distributions, Operating BWR, Turbine Building, Site Q, Location F-Floor 248, Laundry/Turbine Loading Area

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FIGURE A.4.6. Pulse-Height and Photon Energy Distributions, Operating BWR, Turbine Building, Site Q, Location G-Floor 248, Entrance #1 to Turbine Building



FIGURE A.4.7. Pulse-Height and Photon Energy Distributions, Operating BWR, Turbine Building, Site Q, Location H-Floor 248, Entrance #2 to Turbine Building



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FIGURE A.4.8. Pulse-Height and Photon Energy Distributions, Operating BWR, Turbine Building, Site Q, Location I-Floor 248, General Area



FIGURE A.4.9. Pulse-Height and Photon Energy Distributions, Operating BWR, Site Q, Location J-Reactor Vessel Sampling Station



FIGURE A.4.10. Pulse-Height and Photon Energy Distributions, Operating BWR, Site Q, Location K-Spent Fuel Room

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FIGURE A.4.11. Pulse-Height and Photon Energy Distributions, Operating BWR, Site Q, Location L-Waste Storage Area



FIGURE A.5.1. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Waste Storage Area, Site P, Location A-Barrel Storage (Outside Waste Disposal Building)



FIGURE A.5.2. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Waste Storage Area, Site P, Location B-Compactor Area (Outside Waste Disposal Building)



FIGURE A.5.3. Pulse-Height and Photon Energy Distributions, Shutdown FWR, Waste Storage Area, Site P, Location C-Waste Disposal Building



FIGURE A.5.4. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Spent Fuel Pit, Site P, Location D-Heat Exchange







FIGURE A.5.6. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Spent Fuel Pit, Site P, Location F-Fuel Transfer Shoot



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FIGURE A.5.8. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Auxiliary Building, Sice P, Location H-Primary Auxiliary Building, General Area



FIGURE A.5.9. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Auxiliary Building, Site P, Location I-Open Surge Line



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A.5.10. Pulse-Height and Photon Energy Distributions, Shutdown PWR, Containment, Site P, Location J-Above Reactor Head Area (Flooded)



Pulse-Height and Photon Energy Distributions, Operating BWR, Site N, Location A-Heater Bay, Entrance



FIGURE A.6.2. Pulse-Height and Photon Energy Distributions, Operating BWR, Site N, Location B-Heater Bay, Near Steam Lines (collimated)



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FIGURE A.6.3. Pulse-Height and Photon Energy Distributions, Operating BWR, Site N, Location C-MSIV, Entrance Hallway (collimated)



FIGURE A.6.4. Pulse-Height and Photon Energy Distributions, Operating BWR, Site N, Location D-CRD Room (collimated)



FIGURE A.6.5. Pulse-Height and Photon Energy Distributions, Operating BWR, Site N, Location E-Storage Room, Contaminated Pipe (collimated)







FIGURE A.6.7. Pulse-Height and Photon Energy Distributions, Operating BWR, Site N, Location G-Turbine Floor, Maze Entrance to Turbine Room



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FIGURE A.6.8. Pulse-Height and Photon Energy Distributions, Operating BWR, Site N, Location H-Turbine Floor, Inside Turbine Room (collimated)





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APPENDIX B

PHOTON FIELD MEASUREMENT TECHNIQUES

#### APPENDIX B

#### PHOTON FIELD MEASUREMENT TECHNIQUES

The preparation time required to conduct photon field surveys in occupational environments can be considerable. A large part of this time is devoted to studying the literature and preparing the computer software for data analysis. The techniques used for this study are described below to help others reduce their initial time investment. Familiarity with the discussions in Section 2 is assumed.

Described here are example data analysis formulae and computer software. (Instructions for the operation of equipment are typically provided by the manufacturer.) An HP-1000 computer with an enhanced version of ANSI FORTRAN X3.9-1966 was used for the programs discussed here and reproduced at the end of this appendix. The computer programs can be adapted to many micro- or larger computers with Fortran capability. The data file handling procedures will probably require modifications.

#### **B.1 PHOTON SPECTROSCOPY**

The photon spectrometer manufacturer's demonstrations and literature show methods of collecting data in the form of pulse-height distributions, typically recorded by a multichannel analyzer. To conduct an effective spectral monitoring program, the capability of electronically transferring the data into an analysis computer should be available. Input parameters for the analysis software are obtained from detector calibration spectra.

#### B.1.1 Efficiency Calibration

The detector can be calibrated using simple sources (e.g., <sup>109</sup>Cd, <sup>137</sup>Cs, and <sup>co)</sup> for response function characteristics and using calibrated multinuclide sources for efficiency measurements. Multinuclide sources are commercially available. With additional effort, the efficiency calibration can be performed with individually calibrated sources. The sources are positioned at a distance from the detector to simulate parallel incidence of photons. The calibration sources are moved or the detector is rotated to determine the efficiency at other entrance angles. If a collimator is used, the aperture is placed along the axis of the detector, and the calibration is performed with the source along the same axis. Detector efficiency at each source photon energy is calculated by dividing the number of counts in the full-energy peak by the number of incident photons calculated using the source calibration data and the source-detector geometry. The semi-empirical formula given in Equation 8 of Hajnal and Klusek (1974) can be used to parameterize the detector efficiency  $(\Sigma_m)$  for germanium detectors. The formula is:

$$\varepsilon_{m} = e^{-\mu} A^{W} e^{-\mu A_{1}} \left\{ \frac{1 - e^{-\mu A_{2}}}{\mu_{1}} \right\} \left\{ A_{3^{\tau}} + (B.1) \right\}$$

$$\left[A_{4^{\sigma}a} + A_{5^{\sigma}s} \frac{1 - e^{-\mu'A_{6}}}{\mu'} \left(\tau' + A_{7^{\sigma}} e^{-E_{2}A_{8}}\right)\right] e^{-E_{1}A_{9}} + A_{10^{\kappa}} e^{-T_{2}A_{11}}$$

where 
$$E_1 = E\sigma_s/\sigma$$
  
 $E_2 = E_1\sigma_s'/\sigma'$   
and  $\ln Q = \sum_{i=1}^{N} a_i [\ln E]^{i-1}$ 

for  $Q = \tau$ ,  $\sigma_{coh}$ ,  $\sigma_{inc}$ ,  $\kappa$  for aluminum and  $\tau$ ,  $\sigma_{coh}$ ,  $\sigma_{inc}$ ,  $\kappa$  for germanium and  $E_1$ . The parameters are identified in Table B.1 along with the corresponding variable names used in Subroutine EFF (Program TEST). Many of the parameters are fixed using the detector characteristics. Hajnal and Klusek (1974) explain the meaning of the parameters in detail. Other parameters are varied until an acceptable fit is found. The a coefficients are tabulated in Table 1 of Hajnal and Klusek and in Function F of Program TEST. Program TEST, which is used to calculate the efficiency curves, calls Subroutine EFF to calculate Equation B.1. Subroutine EFF requires Function F to calculate the 'Q' parameters. Detector 'ID' = 1 was used to generate the curve in Figure 4 (page 7) of this report.

The highest energy available from the calibration sources used in this work was 1836 keV from <sup>88</sup>Y. The extrapolation to 6 MeV using the semiempirical model was checked using <sup>16</sup>N fields at a reactor site. Two locations with radiation fields almost totally due to <sup>16</sup>N decay were used. Results for the Ge detector are compared to results for a NaI detector and TL dosimeter measurements. The measured dose rates at depths greater than 3 cm were used to estimate the five ce of 6-MeV photons. The Ge efficiency was low, but the resulting error was smaller than that caused by the use of the simple formula for unfolding the spectra.

## TABLE B.1. Efficiency Formula Parameters

Symbol	Unit, Name	FORTRAN Label
τ	barn, photoelectric cross section (Ge)	F(5,E)
σ <sub>a</sub>	barn, Compton energy absorption cross section	SA
σ <sub>s</sub>	barn, Compton scattering cross section (Ge)	SS
σ	barn, average Compton collision cross section (Ge)	S
к	barn, pair-production cross section (Ge)	F(8,E)
μ1	barn, $\mu_1 = \tau + \sigma + \kappa$ (Ge)	XM1
μ	cm <sup>2</sup> g <sup>-1</sup> , mass attenuation coefficient (Ge)	XM
μ <sub>A</sub>	cm <sup>2</sup> g <sup>-1</sup> , mass attenuation coefficient (A1)	ХМА
E	MeV, primary photon energy	E
E <sub>1</sub> , E <sub>2</sub>	MeV, the average once- or twice-Compton- scattered photon energy (Ge)	E1, E2
T <sub>2</sub>	MeV, average energy of electron/positron (Ge)	T2
W	gcm <sup>-2</sup> , window thickness (A1)	A(12)
Ai	Parameters of the semi-empirical Equation (8)	A(1)-A(11)
$\Sigma_{m}(E)$	Semi-empirical efficiency at energy E .	EFM
Σ <sub>T</sub>	Total efficiency without scattering loss	EFT

Primed quantities refer to Compton-scattered photons and are denoted by the FORTRAN label followed by a 'P'.

# B.1.2 Conversion to Energy Distributions

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The efficiency equation (Equation B.1) and the simple model of Seelentag and Panzer (1979) are used to unfold the pulse-height distributions to yield energy spectra. Program HILO is used to calculate the necessary corrections using Subroutine EFF and the parameters developed with Program TEST.

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The parameterization for Compton scattering (see Figure 4 of Seelentag and Panzer) is:

$$H(E,E_{i}) = \begin{cases} \frac{3h(E_{i})}{2} (1 - E/E_{c}) & 0 < E < E_{c}/3 \\ h(E_{i}) & E_{c}/3 < E < E_{c} \\ \frac{h(E_{i})}{2} (1 - 3/4 (E/E_{c})) & E_{c} < E < 4 E_{c}/3 \end{cases}$$

where E is the energy channel,  $E_{\rm C}$  is the Compton edge and h is the amplitude. The data are analyzed by correcting the highest energy bin, subtracting the scattered radiation from all lower bins, and repeating the procedure for all lower bins in order. Dividing the data into bins of 20- to 50-keV increments simplifies the analysis. The corrected number of counts in energy bin E is

$$N_{c}(E) = \frac{N(E) - \frac{6}{7} \sum_{E_{i} > E} H(E, E_{i})}{\sum_{m}(E)}$$

where  $h(E_i) = N_c(E_i)(\Sigma_T - \Sigma_m(E_i))$ and N(E) is the original number of counts in energy bin E. These formulae are coded in lines 137 to 175 of Program HILO.

Corrections for pair production effects are required at high energies. The single- and double-escape peaks are subtracted from the spectra using the number of counts in the full-energy peak and the average probability of recording a 511-keV photon that was created inside the detector by the annihilation of a positron and an electron. The 511-keV detection probability factor is determined for each detector using average ratios of the fullenergy, single-escape and double-escape peaks. Lines 179 to 200 of Program HILO accomplish this.

Calculations using a collimator can be performed by establishing the difference in detector response functions with and without the collimator. The difference is parameterized and stored in Subroutine LEAD. Each collimatordetector combination may require different variable values. The modifications required for Program HILO are given in Table B.2.

TABLE B.2.	Modifications	to	Program	HILO	for	the	Use	of	а	Collin	mator
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Locat	ion	Change				
After Li	ne 11	insert COMMON COL (2100)				
After Li	ne 126	insert CALL LEAD (PEAK(1), PEAK(2))				
After Li	ne 151	<pre>insert COMPT(II) = COMPT(II)*(1.+COL(II))</pre>				
After Li	ine 161	<pre>insert COMPT(II) = COMPT(II)*(1.+COL(II))</pre>				
After Li	ine 171	<pre>insert COMPT(II) = COMPT(II)*(1.+COL(II))</pre>				

Although artifacts in the spectra caused by the approximate formulae used are easily seen, these techniques were determined to be adequate for the purpose. Improved formulae for computer corrections at low energies (<300 keV) are found in the literature (Seltzer 1981). Improved methods at high energies would have to be developed.

#### B.1.3 Calculation of Dosimetry Correction Factors

ed.

As indicated in Figure 3 (page 6) of this report, the spectra are used to generate dosimetry correction factors using a conversion from fluence to exposure and a conversion from exposure to dose ( $C_x$ ). Program CX uses the following formula to calculate effective  $C_x$  factors (( $C_x^{Eff}$ ).

 $c_{x,k}^{Eff} = \frac{\sum_{E_i}^{C_{x,k}(E_i)N(E_i)/F(E_i)}}{\sum_{E_i}^{N(E_i)/F(E_i)}}$ 

The 'k' index refers to the type and tissue depth of the C factor and  $F(E_i)$  is the photon flux per Roentgen. The C factors are listed in the DATA statement for Program CX and correspond to the energies in keV in the DATA statement for E. Eight sets of C factors are listed consecutively for the maximum and central values at 2.5 cm, 1 cm, 0.3 cm, and 0.007 cm.

Correction factors for an instrument or dosimeter that does not have a flat energy response can be calculated by changing the DATA statement for Program CX. The energy response of the instrument or dosimeter must be known.

#### **B.2 THERMOLUMINESCENCE DOSIMETRY**

Thermoluminescent dosimeters can be used to determine the approximate spectral composition of photon and electron fields. The advantages of this technique are that monitoring can be performed fairly inexpensively and will not be limited to a small range of dose rates. The TL material is packaged so that relative magnitudes of dose due to specific energy ranges can be determined. Many (approximately 10) elements are needed to achieve the required sensitivity. Computer analysis of the data can be performed in a manner similar to the TLD-phantom analysis discussed in Subsection 2.3 (page 14). Program POLY uses least-square fitting techniques to determine the optimum linear combination of calibration responses to describe the field data. The multielement TL dosimeter is calibrated using the photon and beta-particle energies that represent the occupational environment. The number of calibration types and energies must be less than the number of elements in the dosimeter. It is important to design the multielement dosimeter to respond uniquely to each energy of interest.

Some of the calculations in Program POLY apply specifically to the analysis of the TLD-phantom data. For example, inverse-distance-squared corrections are made to remove geometry-phantom effects for the least-square fitting. For a different design, the data statement in Subroutine GCORR can be changed.

#### PROGRAM TEST

```
1=00004 15 DN CR00026 USING 00018 BLKS R-0000
16
           FTN4X .L PROGRAM TEST
0001
2889
             C--THIS PROGRAM CALCULATES DETECTOR EFFICIENCIES FOR GE OR GELI
C--DEFECTORS.
0004
0006
                           DIMENSION EN(7). EE(40). EF(40)
DA(A EN/59,5.88.03.122.1.511..661.6.1173.2.2754.1/
WRITE(1.'("ENTER DETECTOR NUMBER OR CHOOSE FARAMETERS (ID=5)"))
WRITE(1.'("ID = "))
READ(1.'(II)')ID
DO 16 I=1.7
E-EH(1)(1000
0008
0009
0010
0012
                           DU 10 1-1.7
E=EM(I)/1000.
CALL EFF(E.ID.EFM.EFT)
WRITE(1.'(3F10.4)') E.EFM.EFT
WRITE(6.'(3F10.4)') E.EFM.EFT
0014
0016
0017
0018
0019
                          WRITE(6.'(3+10.4)') E.EFM.EFT

CUNIINUE

E=0.020

DU 3( I=1.40

E=E*1.189207

CALL EFF(E.ID.EFM.EFT)

EE(I)=E

EF(I)=EFM

CONTINUE

WRITE(6.'(4(2E10.4.5X))')(EE(I).EF(I).I=1.40)

SIDE

END

SUBMOUTTING EFE(E.I.) EED EET EE CO.00)
               10
30
 0024
0026
             SUBROUTINE EFF(E.1), EFN.EFT.PE.CO.PP)

C...DETECTOR EFFICIENCY BASED ON HAJNAL AND KLUSEK.NIM 122.559(1974).

DIMENSION A(12), AA(12,4)
0020
0029
                         DIMENSION A(12).AA(12.4)

DATA AA/.4.15...96..30..56.3.25.8.27.7.57.1.15..111.1.19..3.

X.1.20...9..45..3.5.2..9.9.5.1.5..1..6..14.

X.3.226. 42.23..05180..1743..1521. 5.158.90.11. 9.507. 1.473.

X7.458. 2.993..1370/

DD 4 I=1.9

X=F(I.E)

CONTINUE

LF(I.E)=D1-D50 T0 7
0032
0034
 0036
00370039
               4
                            IF(10.EQ.-1)G0 TO 7
IF(10.NE.5)G0 TO 6
0040
0041
0042
0043
                           ID=-1
WRITE(1,'("SPECIFY DETECTOR PARAMETERS(12)")')
READ(1,'(F10,3)') A
WRITE(1,'("RUNNING")')
G0 TU 7
 0044
0045
0045
0046
0047
                           CONTINUE
DO 5 I=1,12
A(I)=AA(I.ID)
CONTINUE
1048
              6
0050
             うう
0052
           C....ABSORBING MULTIPLIERS
0054
                           XM1=+(5.E)++(6.E)++(7.E)++(8.E)
XMA=(+(1.E)++(2.E)+F(3.E)++(4.E))*.02232
XM=XM1*.008297
EX = 1.-EXP(-XH*A(2))
0056
0057
0058
0058
0060
0061
0062
0063
                           AT = LXP(-XMA*A(1?))*EXP(-XM*A(1))
EF1-AT*EX
           6 ...
           C....PHOTOELECTRIC
C.....PHOTOELECTRIC
C.....PE=A(3) ¥F(5.E) ¥EX/XM1
0064
0065
1066
            6 ....
           C...COMPTON
6600
0069
0070
0071
                         B-F(7,E)
E1=F(9,E)
E2=F(9,E1)
S5=54E1/E
0072
0074
0074
0075
0075
0075
0077
0078
                          SA=S-35
XM1P=F(S.E1)+F(6.E1)+F(7.E1)+F(8.E1)
XMF=XD1P*,008297
SP= F(7.E1)
```

#### PROGRAM TEST (continued)

.

#### PROGRAM HILO

' T4 1=00004 15 ON CR00026 USING 00036 BLKS R-0000 0001 FTN4X.L #FILES(2,2) #EMA(LARGE.0) 0004 PROGRAM HILD() ANALYZE A HI-LO GANMA SPECTRUM 1005 0006 0007 0008 C--THIS PROGRAM CONVERTS PULSE HEIGHT DISTRIBUTION TO AN ENERGY SPECTRUM. INTEGER NAME(6), ITITLE(36), CLEAR(2), LCRSR, LOLK, UNLOCK INTEGER ICHAN(8200) DIMENSION FX(5), CKEV(5,2), PEAK(4), AA(4,12) COMMON/LARGE/DATA(8200), ICHAN, COMPT(8200) 0009 DATA CLEAR/155508,155128/ DATA LCRSR/155078/ DATA LCRSR/155678/ DATA LOCK/155548/ DATA UNLOCK/155558/ 0012 0013 0016 0017 0018 0019 C--GET THE DATA FILE NAME WRITE(1,'(2A2,/,T25,"HI-LO GAMMA SPECTRUM ANALISIS",/)') CLEAR WRITE(1,'(A2)') LOCK WRITE(1,'("ENTER FILE NAME (SIX CHARACTERS MAX): ")') READ(1,'(6A2)') NAME OPEN(80,FILE=NAME,IOSTAT=IOS.ERR=200) 100 C--READ THE TITLE OF THE FILE READ(80.'(36A2)') ITITLE WRITE(1.'(2A2.36A2./)') CLEAR,ITITLE WRITE(1.'("IS THIS THE FILE YOU WHNT??")') READ(1,'(A2)') IANS IF(IANS.NE.2HNO) GOTO 110 CLOSE(80) WRITE(1.'("TRY AGAIN?")') READ(1.'(A2)') IANS IF(IANS.NE.2HNO) GOTO 100 WRITE(1.'("A2)') UNLOCK STOP 0028 00323003400035 0036 0037 0038 6339 C--READ THE DATA FROM THE FILE AND TOTAL THE COUNIS 0040 6---0041 0042 0043 WRITE(1, /(2A2,/,6A2,/,36A2,/,A2)/) CLEAR,NAHE.ITITLE.LOCK WRITE(1, /("GETTING DATA FROM FILE: "6A2)/) NAME TOTAL=0 110 TOTAL=0 DO 120 I=1.8500 READ(80.\*.END-130) ICHAN(I).DATA(I) WRITE(1.'(I4.A2."")') ICHAN(I).LCR5H IF(1.LE.2) GOTO 120 TOTAL=TOTAL+DATA(I) 0044 1045 0046 0047 1048 CONTINUE I=I=1 WRITE(1,'("DATA POINTS:".IS./."TOTAL COUNTS:".F8.2./)') 1.TOTAL CLOSE(80) 0049 0050 0051 0052 120 C---GET NAME OF OUTPUT STORAGE FILE WRITE(1.'("DATA STORAGE FILE NAME: ")') READ(1.'(6A2)') NAME OPEN(81.FILE-NAME.IDSTAT=10S.ERR-140) WRITE(81.'(36A2)') ITITLE DATA(1)=0. DATA(2)=0. 6--140 0060 0061 0062 0063 DATA(2)=0C--CHOOSE DETECTOR 0064 C----WR(1E(1,155) 155 FORMAT("DETECTOR: 1, 31cc"/.T11"2. 48cc"/.T11"3. 59cc"/.T11. +"4. TEST"./.T9."77") 0065 0066 READ(1. (11)) 100 0068 0069 0070 0071 0072 0073 0073 C--- LALCULATE KeV/CHANNEL FOR SPECTRUM (T-MA+b) 256 WRITE(1." ("ENTER CHANNEL NUMBER . KEY IN CHANNEL (-99=END INPUT)") ... KR=0 MAX=5 0076 0077 0078 SUMA=0. SUMA2=0. SUMY-0.

#### PROGRAM HILO (continued)

SUMXY=0. DO 157 K=1, MAX READ(1,\*) CKEV(K.1).CKEV(K.2) 1F(CKEV(K.1),EQ.-99) GUTO 158 SUNX=SUNX+CKEV(K.1) SUNX2=SUNX2+(CKEV(K.1)\*\*2) SUMX2=SUMX2+(CKEV(K.1)\*\*2) SUMY=SUM1+CKEV(K.2) SUMX1=SUMX1+CKEV(K.1)\*CKEV(K.2) 157 CONTINUE 158 K=K-1 K=K-1 A1=(SUMXY-((SUMX\*SUMY)/K))/(SUMX2-(SUMX\*\*2)/K) AD=(SUMY/K)-(A1\*(SUMX/K)) WKITE(1.?("Y="F5.2" X+"F5.2)?) A1.A0 C CALCULATE THE NUMBER OF CHANNELS/BLOCK FOR GIVEN KEV/BLOCK WRITE(1./("TOTAL REV/BLOCK: ")') READ(1.\*) KEVB ICHBK=IFIX(KEVB/A1) WRITE(1./(F7.2." KeV/BLOCK = "IS" CHANNELS -- OK \_")') KEVB.ICHBK READ(1./(A2)') IANS IF(IANS.NE.2HNO) GOTO 160 WRITE(1./(TRY AGAIN? ")') READ(1./(A2)') IANS IF(IANS.NE.2HNO) GOTO 159 GOTO 300 0101 0102 0103 C CALCULATE THE NUMBER OF BLOCKS/SPECTRUM 0105 0106 0107 166 NBK-I/ICHBK 0110 0111 0111 C CALCULATE THE AVERAGE COUNTS AND CENTER KEV IN THE HIGHEST BLUCK N=1 B00 IB-NBK-N SUM-0. FIRST=I-(N¥ICHBK)+1 LAST=FIRST+ICHBK-1 D0 900 K=FIRST.LAST SUN=SUM+DATA(K) 900 CONTINUE 900 CONTINUE 0114 0115 0116 0118 0119 CONTINUE AVE=SUM/ICHBK PEAK(1)=((FIRST\*A1+A0)+(LAST\*A1+A0))/2, PEAK(2)=(FIRST+LAST)/2, E=PEAK(1)/1000. CALL EFF(E.1D0.EFM.EFT.PE.CO.PP) WRITE(1.'("FIRST\_LAST\_SUM\_AVE.CHANNEL")') WRITE(1.'(216.2X.F7.2.2X.F7.2)') FIRST.LAST.SUM.AVE 0120 0121 01223 01223 01223 01225 01225 01227 C--CALCULATE THE COMPTON EDGE 0128 0129 0130 0131 0132 01334 01334 0135 PEAK(3)=PEAK(1)-PEAK(1)/(1.+(2.\*PEAK(1)/511.)) PEAK(4)=(PEAK(3)-A0)/A1 WRITE(1.'("PEAK ENERGY\*PEAK CHANNEL\*EDGE ENERGY\*EDGE CHAMNEL")) WRITE(1.'(4(F7.3.5X))') PEAK C--SUBTRACT COMPTON BACKGROUND DO 186 IK=1.8200 COMPT(IK)=0. COMPT(IK)=0. COMTINUE INEXT-IFIX(PEAK(4)) P=PEAK(1) IF(KR.GT.0)GO TO 188 KR=1 WRITE(1,'("CHOOSE COMPTON CORRECTION COEFFICIENT(DEF=1.)")') READ(1.\*) CCOEFF IF(CCOEFF.LE.0.001) CCOEFF=1.0 H=SUM\*(EFT/EFN-1.)/PEAK(4)\*.86\*CCUEFF WRITE(1.'("AVERAGE H=".F7.1)') H N1=INEXT/3. DO 190 LI=N1.INEXT 0142 0143 0144 0145 0145 0147 0149 0150 0151 DO 190 LIENA, INEXT DATA(II)=DATA(II)-(COMFI(II)) IF(DATA(II).LT.0) DATA(II)=0 CONTINUE H2=H/2. N2=4.\*INEXT/3. SLUFE=(H2=0)/(INEXT-N2) WRITE(1.'("FIRST SLOPE="F7,3)') SLOPE 015/

## PROGRAM HILO (continued)

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LOS . NAME

### SUBROUTINE LEAD

13	T = 0 (	0004 15 ON CR00026 USING 00006 BLKS R=0000
1001 0003 0003 0004 0005 0004 0005 0007 0008 0007 0008 0009 0010	C{	SUBROUTINE LEAD(E,C) SUBROUTINE TO SUBTRACT LEAD COLLINATOR SCATTER (DHMON COL(2100) DIMENSION MU(12).EMU(12).RANGE(40) REAL MU.MUCO DATA MU/S.2.,95.,22.,12.,086.,069.,06.,051.,045.,042.,043.,047/ DATA MU/S.2.,95.,22.,12.,086.,069.,06.,051.,045.,042.,043.,047/ DATA MU/S.2.,95.,22.,12.,086.,069.,06.,051.,045.,042.,043.,047/ DATA MU/S.2.,95.,22.,12.,086.,069.,1200.,1400.,1750.,2500.,5500. X 6500.,10000./ DATA RANGE/1.9.2.1.1.5.1.6.1.6.1.5.1.8.2.2.2.2.2.2.2.4.2.3. X 2.1.2.4.2.3.2.1.2.0.2.1.1.9.2.0.1.8.1.8.1.7.1.2.1.6.1.6.1.4.1.4. X 1.3.1.25.1.25.1.25.1.3.1.3.1.3.1.25.1.25
0012 0013 0014 0015	CD4	ALCULATE NET TRANSMISSION THRU COLLINATOR - 31.86/CM##2 ATA IS NORMALIZED TO CU-50
001478 00119 00119 00022345 000222345 000222345 000222345 000222345 000222345 000222345 000222345 00023312334	10 20 35 40	$\begin{array}{l} \text{MUCD-0.06} \\ \text{T-31.8} \\ 18F = 3\theta \\ \text{DD 10 I=1.10} \\ \text{IF(E.L1.EMU(I)) GO TO 20} \\ \text{CONTINUE} \\ \text{NET-MU(I)-MUCO} \\ \text{Ab=EXP(-NET*T)} \\ \text{DO 40 I=1.2100} \\ \text{EC=1/C*E} \\ \text{ED=EC/INF} \\ \text{J=IFIX(ED)} \\ \text{IF(J.GT.38) GO TO 35} \\ \text{COL(I)=AB*(RANGE(J)-1.)} \\ \text{GO 10 40} \\ \text{COL(I)=AB*0.3} \\ \text{CONTINUE} \\ \text{RETUKN} \\ \text{ENU} \end{array}$

## PROGRAM CX

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Ti	T=00004 IS ON CR00026 USING 00024 BLKS R=0000
0001 0002 0003	FIN4X.L \$FILES(2.2) \$EMA(LARGE.0) PROGRAM CX().CALCULATES EFFECTIVE CX VALUES FOR THE ICRU SPHERE
0006	CDATA INTERPOLATED FROM DIMBYLOW AND FRANCIS, 1979, A CALCULATION OF THE CPHOTON DEPTH-DOSE DISTRIBUTIONS IN THE ICRU SPHERE FOR A BROAD PARALLEL CBEAM.A POINT SOURCE AND AN ISOTROPIC FIELD.NAT RAD PROT BOARD, NRPE 92, CHARWELL, ENGLAND.
$\begin{array}{c} 3001123456789012334567890142345678901423456789014233456789014223345678901423345678901400000000000000000000000000000000000$	INTEGER NAME(6).ITITLE(36).CSCRN(2).LCRSR.LOCK.UNLOCK INTEGER ICHAN(8200) DIMENSION F(5).CKEV(5,2).E(31),FL(31),CX(31.B).C(8).CT(8) COMMON/LARGE/DATA(8200).ICHAN.COMPT(2100) DATA LCRSR/15550B.IS5128/ DATA LCRSR/15555B/ DATA LUX/15555B/ DATA UNLOCK/15555B/ DATA (V1)5555B/ DATA (V1)55555B/ DATA (V1)55555B/ DATA (V1)55555B/ DATA (V1)55555B/ DATA (V1)555555555555555555555555555555555555
0050	CGET THE DATA FILE NAME AND LU
0052	WRITE(1.'(2A2./.T25."SPECTRUM ANALYSIS PROGRAM"./)') (SCRN WRITE(1.'(A2)') LOCK WRITE(1.'(A2)') LOCK WRITE(1.'(GACA)') NAME (SIX CHARACTERS MAX): ")') READ(1.'(GAC)') NAME CPEN(B0.FILE=NAME.IOSTAT=LOS.ERR=200)
0058	CREAD THE TITLE OF THE FILE
0050 0051 0053 0053 00553 00555 00555	READ(80,'(36A2)') ITITLE WRITE(1.'(2A2.35A2./)') CSCRN.ITITLE WRITE(1.'("15 THIS THE FILE YOU WANT?? ")') READ(1.'(A2)') IANS IF(IANS.NE.2HNO) GOTO 110 CLOSE(80) GOTO 100
0068	CREAD THE DATA FROM THE FILF
0089	110 WRITE(1, '("SEITING DATA FROM FILE: "642)') MANE
1072 073 0074 0075	120 1=1,8500 READ(80,*.END=130) ICHAN(1).DATA(1) WRITE(1./(I4.SX.F7.0.A2."")) ICHAN(1).DATA(1).LCRSR 130 I=1-1
0077	CLUSE (B0)

**\$**-1

\* - 3 - 4 <sup>(</sup>

1

Qa -

# PROGRAM CX (continued)

\*

0079	CCA	ILCULATE CX VALUES
000823	8	R1=0. D0 8 K=1.8 L1(K)=0. D0 30 J=3.2048
0086	10	ENELCHARUSS DO 10 I=1,31 IF(EN.LT.E(I))GO TO 20 CONTINUE
0089	50	IG=I IF(IG, LT, 2)G0 TO 30 EI=(E(IG)-EN)/(E(IG)-E(IG-1)) E==(EI(IG)-EN)/(E(IG)-E(IG-1)) E==(EI(IG)-EN)/(E(IG-1)) E==(EI(IG)-EN)/(E(IG)-E) + EI(IG)
0093	25	DO 25 K=1.8 C(K)=CX(IG,K)-(CX(IG,K)-CX(IG-1,K))*EI R= DATA(J)/F
0096 0097 0098 0099	28	RT=RT+R DO 28 K=1.8 CT(K)=CT(K)+R*C(K) LOPTINUE
6100 0101 0102	32	DO 32 K=1,8 C(K)=CT(K)/RT WRITE(1,'("CX FACTORS FOR THE ICRU SPHERE, PLANE PARALLEL INCIDENCE WRITE(1,'("CX FACTORS FOR THE ICRU SPHERE, PLANE PARALLEL INCIDENCE
0103 0105 0105 0106 0107 0108		WRITE(1, '(/"MAXIMUM VALUES: 2.5CM = ",F5.2./.16X."DEEP = ", XF5.2./.16X."D.3CM = ",F5.2./.16X."SHALL = ",F5.2.//"CENTRAL VALUES X: 2.5CM = ",F5.2./.16X."DEEP = ",F5.2./.16X."O.3CM = ",F5.2./. X:6X."SHALL = ",F5.2./. WRITE(1, '(/"TOTAL EXPOSURE = ",E10.4." 10**-8 R*CM**2")')RT
0107 0110 0111 0112 0113 0114	200	G0 10 300 WRITE(1,'("READ ERROR "14" ON FILE "6A2" OPEN = ")') 105.NAME CLUSE(80) WRITE(1,'(/.T25."PROGRAM IS FINISHED")') WRITE(1,'(A2)') UNLOCK STOP
0115		END

9

### PROGRAM POLY

15	T = (	10004 15 ON CR00026 USING 00042 BLKS R=0000
00 1	FTN	ALL AN
00.3	Se Li	PROGRAM POLY( ). PHANTOM DATA ANALYSIS
00 4 00 5 00 5 00 9 00 9 00 9 00 10	1000000	THIS PROGRAM ANALYSES DATA FROM THE PLEXIGLASS ILD PHANIOM. FLD DATA IS ENTERED ALONG WITH THE CALIBRATION AND BACKGROUND: IF THE GEOMETRY IS A POINT SOURCE, AN INVERSE R2 CORRECTION IS APPLIED. THE PROGRAM THEN FITS EXISTING CALIBRATIONS FOR THE PHANTOM TO THE DATA. TO DETERMINE THE RELATIVE CONTRIBUTIONS OF THE COMPONENTS.
1012	č	NOTE: AN OPTION ALLOWS OMISSION OF ANY TWO COMPONENTS.
0013 0014 0015 0016 0047	C	REAL X.D(11).SklP(11) REAL T(11,5).R(11).CXS(5).CXD(5).C(11.11).A(6.7).FRAC(5).DIST INTEGER EN(6.3).TITTLE.LENT.GEOM.NAME(3).ANS.MUDE.SLAL.TIPE.INUM INTEGER SCAL1
0019	C.	DATA D/0.0,0.48.0.95.1.43.1.91.3.02.5.08.7.14.0.48.1.91.7.14/
0500	Ç.	ENTER DATA
0022		10 FORMAT ("1" /// TIS. "PLEXIGLASS TED PHANTOM DATA ANALYSIS" ///)
0024 0025 0025 0027 0027	5	MODE=2 1DEFAULT VALUE 10 FORMAT (//,T7,"PHANTOM DATA OR MODIFIED THI DOSIMETER?"./. + T7." (ENTER 2 FOR PHANTOM, 1 FUR HODIFIED THI) READ (1.'(11)') MODE TNUM=0
0029	5	IF (MODE.EQ.2) WRITE (1,520) 20 FORMAT (77,17,"TLD-200 DALY (ENTER 1)"
0031		+ "OR BUTH TLD-700 AND TLD-400 (ENIER 2) 7 ")
0033		IF (TYPE, EQ. 2) TNUM=TNUM+3 111 READINGS IF CaF:Mn INCLUDED
0035	6	MODE=MODE+2 INUMBER OF ENTRIES PER LINE
0037		11 FORMAT (//, T7. "15 DATA IN A FILE? ")
0038	0.13	NEAD (1.12) ANS 12 FORMAT (A1)
0040	c	IF (ANS.NE.IHT) GD TO 19
0042	č	DATA IN A FILE
0044		13 FORMAT (///T7."ENTER 6 CHARACTER FILE NAME ")
0045		READ (1.14) (NAME(1),1=1.3) 14 FORMAT (3A2)
0047		OPEN (89.FILE=NAME) READ (89.12) ANS III NORE FILE BEADING
0049		DU 16 J=1, INUN
0031		$\frac{(K = A)}{16} (HY, *) (T(J, K), K = 1, MUDE)$
0052		CLOSE (89) G0 TO 28
0022	E	DATA FROM TERMINAL
0056	(* L)	19 WRITE (1,20)
0038		+ "DEPIH, ". 7. T7. "(NOTE: VALUES OF ZERO WILL BE IGNORED)". 7)
0059		DU 2/ J=1.INUM WRITE (1.25) J
0061		25 FORMAT (17, "POSITION ".11,") ")
0063	C '	
0065		AVENAGE VALUES FOR EACH PUSITION 28 DO 40 J=1.INUM
0065		5Um=0. CNI=1.0#MODE
0068		DO 30 K=1.MODE
0020		IF ([(],K),EQ.0.) ERT=CN1-1.0 FIGNORE ZERO VALUES
0072		IF (UNI,EQ.0.) CNT=CNT+1.0 IINO DIVISION OF (ENO)
0073		R(J)=5UM/CNT
0025		WRITE (1.50)
0077		READ (1.*) BKG. BKG4
A		
#### PROGRAM POLY (continued)

```
WRITE (1.'(////T7."BACKGROUND SUBTRACTED VALUES".//))
DO 60 J=1.INUM
IF(J,GI,B)BK=HKG4
R(J)=R(J)-BK
IF (R(J).LT.0.0) R(J)=0.
WRITE (1.*) R(J)
0079
0080
0081
0082
0083
0084
0085
                   60 CONTINUE
            ć
                          CORRECT FOR SOURCE GEOMEIRY
0008
            £
                   WRITE (1.62)

62 FORMAT (7/7.17."OPTIONAL GEOMETRY CORRECTION:".

+ 77.19."1. NO CURRECTION (UNIFORM FIELD)".

+ 77.19."2. 1/R CORRECTION (LINE SOURCE)".

+ 71.2."3. 1/R**2 CURRELIUN (POINT SOURCE)".

+ 71.7."ENTER OPTION DESIRED (1.2 OR 3) ")
008900900091
0092
0094
                   * //,T/."ENTER OPTION DESIRED (1.2 OR 3) ")
READ(1.64) GEOM
64 FORMAT(I1)
IF (GLOM.EQ.0)GEOM=1 |DEFAULT
IF (GEOM.EQ.1) GO TO 68
WRITE (1.66)
66 FORMAT (//,T7."ENTER DISTANCE FROM SOURCE TO PHANTOM (CM)
READ (1.*) DIST
CALL GLORR(R.DIST.GEOM.INUM)
68 CONTINUE
0096
0049
0100
                                                                                                                                                                 0101
0102
0103
0103
0104
0105
0105
0107
0108
             00
                   NORMALIZE TO 0.5 CM AND DISPLAY TLD RESULTS
WRITE (1,'(///)')
DO 72 J=1.TNUM
X=R(J)/R(2)
WRITE (1.*) D(J),R(J).X
72 CONTINUE
0108
0119
0111
0111
0111
0113
             00
                          REAF CALIBRATION VALUES FROM FILE
            0
            C OPTION TO IGNORE A CALIBRATION
WRITE (1,1070)
1070 FORMAT (77.17."ENTER CALIBRATIONS TO BE SKIPPED (78 FOR NONE)
READ (1."(242)') SCAL.SCAL1
0114
0116
                                                                                                                                                                       " )
0118
            π.
                          IF (NODE.EQ.3) [HEN
OPEN (12.FILE='TCALIB',
ELSE
OPEN (12.FILE='PCAL')
ENVIE
0120
                   READ (12,70) TITTLE
70 FORMAT (AL)
ICNT=0
DU 90 I=1.5
0124
                                                                                                                    IFILE HEADING
8139
                   DU 90 1=1.5

READ (12.80 END=100) (EN(I.L).L=1.3) IREAD CALIBRATION SOURCE

80 FURMAT (3A2)

1F ((EN(I.1).EQ.SCAL).OR.(EN(I.1).EQ.SCAL)) THEN LOPTION TO SKIP CALIB

READ (12.*) LS.CD

READ (12.*) (SKIP(J).J=1.TNUM)

ELE
0128
0130
0132
                          ELSE
ELSE
READ (12.*) CXS(I).CXD(I)
READ (12.*) CC(ICNI.J).J=1.INUM)
0134
                                                                                                                    ISHALLOW AND DEEP CX
                                                                                                                          TILD READINGS FUR IN UR 1 RAD
0130
0137
0138
0139
                   DO 68 L-1.3
88 EN(ICNT.L)=EN(I.L)
ENDIF
                 90 LONTINUE
100 CONTINUE
0140
0141
0142
0143
            8
                          DETERMINE CROSS PRODUCTS
AND SET UP MATKIX FOR SIMULTANEOUS EQUATION SOUUTION
BY GAUSSIAN ELIMINATION.
1144
0145
0146
              č
             č
                            THE INPUT MATKIX HAS DIMENSIONS TENT BY ICNI+1. WHERE
TENT IS THE NUMBER OF CALIBRATIONS
                          THE
0148
             €.
0149
             ĩ.
                          N=ICNI+1
SEI MATRIX 10 ZERO
             C
                 DO 120 I=1.ICNI
DO 110 J=1.0
110 A(I.J)=0.0
120 CONTINUE
0152
0153
0154
0155
0139
                          CALCULATE CR055 PRODUCTS
D0 180 1=1.1CMT
D0 160 J=1.1CMT
             6
                                                                 i J-COLUMN NUMBER
0150
```

### PROGRAM POLY (continued)

DO 140 L=1, INUM I = DEPTIA(I,J)= A(I,J) + C(I,L)\*C(J,L) 140 CONTINUE 160 CONTINUE 180 CUNTINUE 0159 I L=DEPTH (POSITION IN PHANION) 9160 0161 0162 0163 0164 C D0 200 I=1.ICNT D0 190 L=1.TNUN 190 A(I.N)=A(I.N)+R(L)\*C(I.L) 200 CUNTINUE 0166 ILAST COLUMN OF MAIRIX 0168 01669012345 C C PRINT INTERMEDIATE VALUES DO 205 L=1,ICN1 WRITE (1.\*) (A(L.LL).LL=1.N) 205 CUN(INUE CLOSE (12) CC 0176 0176 0177 0178 0179 0180 SOLVE FOR COEFFICIENTS CALL MATRX(ICNT.A) ANSWERS ARE NOW IN LAST COLUMN OF A CC DISPLAY RESULTS DISPLAT RESULTS WRITE (1, (///)) 100 230 L=1, ICNT 230 WRITE (1, 240) (EN(L,J), J=1.3), A(L,N) 240 FURNAT (17.3A2.2X.F6.3./) 0181 1182 0183 0184 END f, 0186 SUBROUTINE GCORR(R.DIST.GEOM.TNUM) REAL R(11).DIST.D(11) INTEGER GEOM.EXP.TNUM DATA D/0.0.0.48.0.95.1.43.1.91.3.02.5.08.7.14.0.48.1.91.7.14/ EXP=GEUM-1 D0 10 I=1.TNUM R(I)=R(I)\*(((DIST+D(I))/(DIST))\*\*EXP) 10 CONTINUE IF (TNUM.EQ.1) WRITE (1.20) (R(I).I=1.8) IF (TNUM.EQ.1) WRITE (1.25) (R(I).I=1.1) 20 FORMAT (//.T7."CORRECTED READINGS: "./.T9.8(F5.0.1X)) FORMAT (//.17."CORRECTED READINGS: "./.T9.11(F5.0.1X)) RETURN END C 0188 0189 0190 0191 END CC SUBROUTINE MATRX(M.A) REAL ALG.7 INTEGER N, 0204 0205 0209 0209 0210 0211 GODDARD COMPUTER SCIENCE INSTITUTE C LCC SUBROUTINE NAME,
 LALLING NAME,
 LALLING NAME,
 SIMUL FANEOUS EQUATION SOLVER
 STATUS/CHANGE LEVEL,
 FROGRAMMED BY,
 ADAPTED TO HP FORMAT
 SIMUL FANEOUS EQUATION SOLVER
 SIMUL FANEOUS EQUATION SOLVER 0212 G 0213 000 0215 0216 0217 0216 0219 SUPPORTED IN PART BY GRANT NO. FR00197 BY THE SPECIAL RESEARCH RESOURCES BRANCH, NIH-USPHS, 0 02250 £ SUPPORTED IN PART BY GRANT N., FR00197 BT THE SPECIAL RESEARCH RESOURCES BRANCH, NIH-USPHS. 02224567 02290 0 ٤. 5 00000 C 5 0236 0237 0238 The array A contains the input coefficients and is passed in ConnoN. The number of equations (i) is passed to the subroutine by the call statement- CALL SSIME(M) (M (ar=2c). The resultant G LC.

# PROGRAM POLY (continued)

39 C 40 C		values are found in the (M+1)th elements of the array H: i.e., variable $i = (i, M+1)$ , variable $2 = (2, M+1)$ , variable $M = (M, M+1)$ .
42 č		
43		N=n+1
44		11=1
15	10	13=11
46		5UD-6B5(6(11,11))
17		DU301=11.M
18		IF (SUN-ABS(A(1.11)))20.30.30
99	50	13=1
2Q	70	ラリロテクビカとの(1,11)/
21	39	
24		
20	40	
27		
23	E 0.	
20	20	H13.37-00H
21	60	
28	20	
77	10	H(1,11)-H(1,11/)H(11,11)
60	80	
01		
02	0.0	
0.3	70	D01003-13.M
64	+ 0.0	$\Delta (T + 1) = \Delta (T + T) = \Delta (T + T) + \Delta (T - T)$
03	100	TELLO / HALL OF HALL FIRMAN
29	110	
60	***	
40		DOLOTION M
20		
21	120	A(I,II) = A(I,II) - A(I,J) * A(J,II)
25		IE(11-H)10, 80, 10
22	130	D0150(=1.M
74	100	J2=M I
75		13-12+1
76		A(13,N) = A(13,N)/A(13,13)
77		(F(J2))40,160,140
78	1.40	001501=1.12
599	150	A(J,N) = A(J,N) - A(I3,N) * A(J,I3)
0.0	160	RETURN
10.1	200	ENI

2

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BIBLIOGRAPHIC DATA SHEET	1. REPORT NUME NUREG/CR PNI - 4915	ER (Assigned by DDC) -3569	
4 TITLE AND SUBTITLE (A dd Volume No. 1/ appropriate) Spectral and Dosimetric Measurements of Photon F	ields at	2. (Leave Disn's)	
Commercial Nuclear Sites	3. RECIPIENT'S A	CCESSION NO.	
PL Roberson RA Fox KL Holbrook <u>GWR Endres</u> DL Haggard LA Rathbun	5. DATE REPORT MONTH December	COMPLETED VEAR 1983	
Pacific Northwest Laboratory	August	ISSUED	
Richland, WA 99352	8. (Leave blank)		
12. SPONSORING ORGANIZATION NAME AND MAILING ADDRESS Unclude Zip C Division of Facility Operations Office of Nuclear Regulatory Research U.S. Nuclear Regulatory Commission Washington, DC 20555	10. PROJECT, TASK	. B2419	
13. TYPE OF REPORT PER Research Report	O COVERED (Inclusive dates)		
15. SUPPLEMENTARY NOTES	14 /Leave ofank)		
Commercial nuclear reactor sites. Revisions to 1 conversion factors (C <sub>X</sub> ) much greater than unity f could impact personnel monitoring practices. Mon tissue and shallower could underestimate doses re (>3 MeV). No locations with large C <sub>X</sub> factors (approxim significant production of low-energy photons was The scatter continuum has an effective C <sub>X</sub> factor tion was found with a nearly pure scatter spectru cant contributions from medium-energy photons due cobalt and cesium isotopes. Monitoring requireme tissue were found to be adequate for estimating d taining high-energy photons. Enhanced surface do electrons were measured in all locations monitor may provide inaccurate results in high-energy fie	O CFR 20 that specif or photons between 4 itoring at effective ceived from high-ene ately 1.5 rad/R) wer found to be due to p of approximately 1.2 n. Other locations primarily to radioa nts at 0.007-cm and ose received in radi ses attributed to hi ed. Personnel monit lds.	y exposure-to-dose o keV and 200 keV e depihs of 1 cm of ergy photon fields re found. The most hoton scattering. rad/R. One loca- contained signifi- ctive decay of 1.0-cm depths in ation fields con- gh-energy knock-on oring techniques	
Photons, Gamma ray, Dosimetry Radiation measurement, Gamma Spectrometers	ESCRIPTOR		
175. IDENTIFIERS OPEN ENCED TERMS			
8 AVAILABILITY STATEMENT	Unclassified	21 NO OF PAGES	
nlimited 20	"Unclassified """	22 PRICE	

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