

$N_e^-/(N_e+N_\gamma)$ for the internally converted 0.190-Mev line. This has been given by Lawson and Cork as 1.00 ± 0.30 , by Boehm and Preiswerk as 0.80 ± 0.05 , and Langer and Price⁶ as 0.80 ± 0.10 . Taking the value of 0.80 as correct the percentage K -capture is found to be 4.0 percent in good agreement with Boehm and Preiswerk. The disintegration scheme is shown in Fig. 2.

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¹ J. L. Lawson and J. M. Cork, Phys. Rev. 57, 982 (1940).

² Cork, Shreffler, and Fowler, Phys. Rev. 74, 1657 (1948).

³ F. Maienschein and J. L. Meem, Jr., Phys. Rev. 76, 899 (1949).

⁴ F. Boehm and P. Preiswerk, Helv. Phys. Acta 22, 331 (1949).

⁵ L. H. Gray, Proc. Camb. Phil. Soc. 27, 103 (1931).

⁶ L. M. Langer and H. C. Price, Jr., private communication.

Penetration and Diffusion of Hard X-Rays through Thick Barriers. IV. Multiply Scattered γ -Rays: Angular Distribution*

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IN order to gain some insight into the angular distribution of γ -rays which are scattered many times within a barrier, the following quite schematic problem has been studied theoretically: A source of monochromatic γ -rays of energy E_0 is distributed uniformly throughout an isotropic medium. Each γ -ray is emitted in exactly the same direction. What will be the distribution in angle, $N(E, \theta)$, of those photons which are degraded by Compton scattering to the energy E ? (θ is measured from the initial direction.)

This might be said to be a study of the manner in which photons "forget" their original direction as they lose their energy through Compton scattering.

Under the conditions of this problem, $N(E, \theta)$ obeys the following integral equation:

$$\begin{aligned} \mu(E)N(E, \theta) = & \int_E^{E_0} k(E', E)dE' \int_{4\pi} (1/2\pi)\delta \\ & \times (\omega \cdot \omega' - 1 + mc^2/E - mc^2/E') \times N(E', \theta') d\omega' + k(E_0, E)\delta \\ & \times (\cos\theta - 1 + mc^2/E - mc^2/E_0) \end{aligned} \quad (1)$$

where $\mu(E)$ is the probability, per unit path length, that a photon of energy E will undergo scattering, photoelectric absorption, or pair formation; $k(E', E)$ is the probability, per unit path length, that a photon of energy E' will attain the energy E through a scattering process; δ is the Dirac function and ω represents a

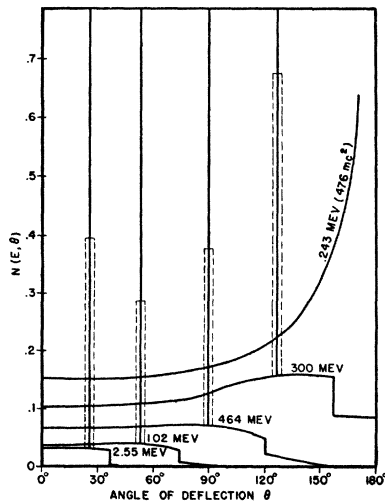


FIG. 1. Angular distribution of secondary photons for various energies of degradation. Initial energy $E_0 = 5.1$ Mev ($10 mc^2$). The dotted rectangles represent the strength of the corresponding sharp lines.

unit vector with direction coordinates (θ, ϕ) . The first term in (1) gives the rate of disappearance of photons having energy E and inclination θ . This must be balanced (under steady state conditions) by contributions coming from the source in a single scattering, represented by the third term, plus those contributions coming from the source in two or more scatterings, represented by the second term.

Using an expansion in Legendre polynomials,¹ that is,

$$N(E, \theta) = \sum_{l=0}^{\infty} [(2l+1)/4\pi] N_l(E) P_l(\cos\theta) \quad (2)$$

$$\begin{aligned} \delta(\omega \cdot \omega' - 1 + mc^2/E - mc^2/E') \\ = \sum_{l=0}^{\infty} [(2l+1)/2] P_l(1 - mc^2/E + mc^2/E') P_l(\omega \cdot \omega') \end{aligned} \quad (3)$$

we can carry out the integral over ω' . Introducing (2) and (3) into (1) we find that the coefficients $N_l(E)$ must fulfill the integral equation

$$\begin{aligned} \mu(E)N_l(E) = \int_E^{E_0} k(E', E) P_l(1 - mc^2/E + mc^2/E') N_l(E') dE' \\ + k(E_0, E) P_l(1 - mc^2/E + mc^2/E_0). \end{aligned} \quad (4)$$

For $l=0$, this equation coincides with Eq. (1) in the accompanying paper.²

We solve Eq. (1) in the following manner: We obtain the contributions to $N(E, \theta)$ which are scattered only one time or only two times through iteration of (1). We subtract these same contributions from Eqs. (4), likewise through iteration. Orders of scattering higher than the second can contain at most a discontinuity in the derivative $dN/d\theta$; consequently the part of the sum (2) representing these higher orders of scattering will converge reasonably well. Actually, we found that using values of l up to four we obtain sufficiently good convergence at all except the highest energies.

We did some numerical work for purposes of orientation choosing $E_0 = 10 mc^2$; and for materials we used aluminum and lead as representative light and heavy elements. Figures 1 and 2 show the results of our calculations for various values of E in the case of aluminum. In Fig. 1, the vertical straight lines indicate the portion of the radiation of each energy which arises from single scattering and whose angular distribution is peaked like a δ -function. The dotted rectangles give a measure of the intensity of this component. The other discontinuity in each curve is at the maximum angle obtainable by a photon going from E_0 to E in two scatterings. These last appear also in Fig. 2.

The high orders of scattering become increasingly effective as the energy decreases, as evidenced by the increase in the number of photons at larger angles than the "two-scattering" cut-off. It will be noticed that at $E = 0.476 mc^2$, which corresponds to a single scattering through 180° , this multiply scattered beam tends to ∞ at the backward pole. This is because those photons attaining $0.476 mc^2$ in two scatterings can reach this direction no matter what the intermediate energy is. At still lower energies than

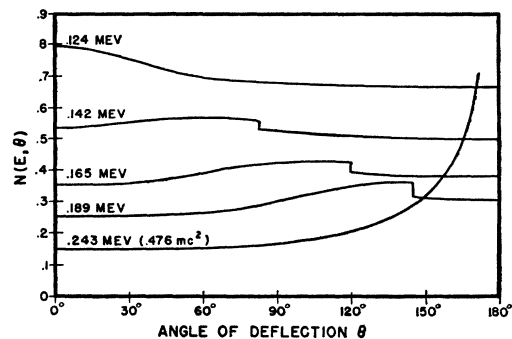


FIG. 2. Angular distribution of secondary photons for various energies of degradation. Initial energy $E_0 = 5.1$ Mev ($10 mc^2$).

$E=0.476$ mc² there is no longer a singly scattered beam; the solution has the appearance of a wave running across as the energy decreases. Not all the departure from isotropy at these lower energies is due to the beam which is scattered twice—there is still a slight “wave” in existence when the second scattered beam is eliminated. This is largely due to photons which are scattered every time except once through angles close to 0° or to 180° and thus have some remembrance of their original direction.

The solution in the case of lead has the same general features, except that the low orders of scattering are much more important, due to the increased competition which pair production and photoelectric effect give to Compton scattering. Also, instead of a build-up in the total density of photons at lower energies, there is a sharp decrease, due to photoelectric effect.

A more detailed account of this work will appear in the Journal of Research of the National Bureau of Standards.

We should like to thank Dr. U. Fano for suggesting this problem and for many valuable discussions during the course of the work.

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¹ See e.g. M. Verde and G. C. Wick, Phys. Rev. 71, 852 (1947).

² P. R. Karr and J. C. Lamkin, Phys. Rev. 76, 1843 (1949). This is paper III of the series; papers I and II are quoted there.

Radioactive Lanthanum 140

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THE single natural isotope of lanthanum of mass 139 should, on neutron capture in the pile, yield the radioactive isotope of mass 140. This emitter has been previously studied by many investigators.¹ Because of the difficulty of chemical purification, impurities are likely to be present. A highly purified sample separated by Dr. G. E. Boyd was kindly made available and irradiated in the Oak Ridge pile.

This specimen shows a remarkably pure decay through eight octaves with a half-life of 41.4 hours. In photographic beta-spectrometers, many electron conversion lines are observed, about 16 in all, with energies less than 500 kev. These are shown in Table I, together with their proposed interpretation and the values of the resultant probable 12 gamma-rays. The $K-L$ differences were observed are characteristic of cerium, indicating that gamma-emission follows the loss of a beta-particle from the lanthanum nucleus.

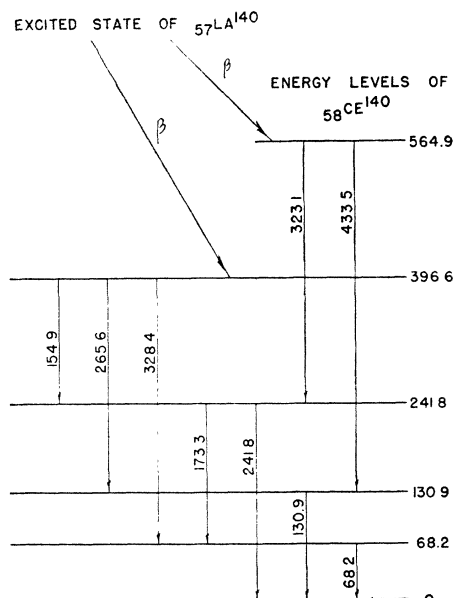


FIG. 1. A possible level scheme associated with $^{57}\text{La}^{140}$.

The observed gamma-rays may be fitted into a level scheme as shown in Fig. 1. The sums of all possible combinations agree as expected to within about 0.2 kev. The gamma-rays observed at 109.1 and 487.1 kev do not, however, fit into the level diagram. The only positions they could occupy and remain consistent with the scheme would be either above or below the group of levels shown in Fig. 1. The electron line at 483.2 kev may also be interpreted as a K line, in which case the gamma-energy is 523.5 kev. This value happens to be almost exactly the sum of three observed gamma-rays of 109.1, 173.3, and 241.8 kev. The presence of these two anomalies indicates that some ultimate revision of the proposed level scheme may be necessary, to include also the higher energy gamma-rays.

TABLE I. Conversion electron and probable gamma-ray energies.

Electron energy	Probable identification	Energy total	Gamma-energy
27.9 kev	K_1	68.2 kev	68.2 kev
68.7	K_2	109.0	109.1
90.7	K_3	131.0	130.9
103.0	L_2	109.2	
107.9	M_2	109.3	
114.6	K_4	154.9	154.9
124.5	L_3	130.7	
133.0	K_5	173.3	173.3
201.5	K_6	241.8	241.8
225.3	K_7	265.6	265.6
282.8	K_8	323.1	323.1
288.2	K_9	328.5	328.4
322.0	L_9	328.2	
393.2	K_{10}	433.5	433.5
446.8	K_{11}	487.1	487.1
483.2	K_{12} or L_{11}	523.5	523.5
		489.4	

The beta-emission had been previously shown to be complex. Higher energy gamma-radiation is present which by absorption in lead had an energy of 1.88 Mev. The long-lived 140-day activity previously reported does not appear to be present in this specimen.

This investigation was made possible by the joint support of the ONR and the AEC.

¹ Weimar, Pool, and Kurbatov, Phys. Rev. 63, 67 (1943); R. Osborne and M. Peacock, Phys. Rev. 69, 679 (1946); C. Mandeville and M. Scherb, Phys. Rev. 73, 1434 (1948); Mitchell, Langer, and Brown, Phys. Rev. 71, 140 (1947); Cork, Shreffler, and Fowler, Phys. Rev. 74, 240 (1948).

Ferroelectricity in the Ilmenite Structure

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SINGLE crystals of LiTaO_3 and LiCbO_3 have been found to be strongly ferroelectric as is indicated by the existence of a saturation polarization in their dielectric hysteresis loop. The saturation value increases rapidly with temperature as shown in Fig. 1. LiCbO_3 exhibits a similar behavior at higher temperatures.

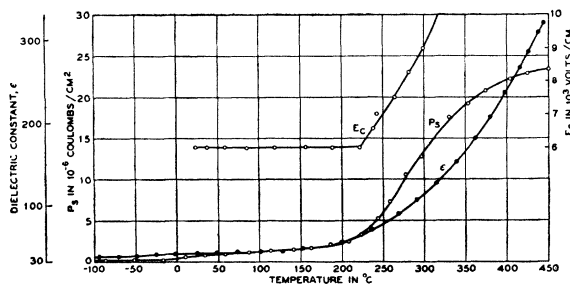


FIG. 1. Saturation values in a single crystal of LiTaO_3 as a function of temperature.

LiCbO_3 , with which LiTaO_3 is isomorphous, was described by Zachariasen¹ as belonging to the ilmenite structure, which is centro-symmetric. The existence of a spontaneous polarization, however, indicates the absence of a center of symmetry. A further