

FIG. 1. Pulse spectra from x-rays of Te125 and I126.

spectrum measured through sufficient beryllium to absorb the beta-particles and L x-rays. For comparison, the tellurium Kx-ray spectrum resulting from the isomeric transition in Te¹²⁵ is also shown. The correspondence of the two curves proves that the I¹²⁶ radiations are tellurium K x-rays. Xenon K x-rays arising from internal conversion following β^- decay cannot contribute more than 2 percent of the x-ray intensity.

The absolute K-electron capture rate of an I^{126} sample was calculated from the x-ray pulse height distribution by the method previously described.⁶ A fluorescence yield of 0.75 was assumed.⁸ The L capture rate may be estimated⁹ to be 12 percent of the Krate. The absolute beta-disintegration rate was measured for an I¹²⁶ sample of known capture activity by correction of the measured Geiger activity for solid angle, absorption in air and counter window, back scattering, coincidence loss, and x- and gammaactivities. The ratio of the electron capture probability to the beta-decay probability for I^{126} was found to be 1.44 ± 0.15 .

Beta-ray spectrometer measurements made by Alburger demonstrated, in agreement with earlier data,¹ the existence of two beta-transitions of energies 1.24 ± 0.02 Mev (~25 percent) and 0.85 ± 0.03 Mev (~75 percent), and of conversion electrons of a 0.382 ± 0.004 -Mev gamma-ray.

Gamma-ray energies were measured with a scintillation spectrometer, in collaboration with E. der Mateosian. A 640 ± 20 -kev gamma-ray line was found, low in intensity compared with the previously known 390-kev line. The energies of the two lines were established by comparison with the gamma-rays of Cs¹³⁷ (663 kev) and of Au¹⁹⁸ (411 kev). In addition, a faint line was observed of energy corresponding to annihilation radiation.

With an I126 source, coincidences were observed between pulses from two NaI(TII) scintillation counters placed at right angles and shielded from each other by 4 mm of lead. Measurements with copper and lead absorbers showed that approximately onehalf the observed coincidence, rate represented x- γ events and one-half γ - γ events. A comparison of the coincidence rates at 180° and at 135°, when both counters were covered with 3 g/cm^2 of lead absorber, established the presence of annihilation radiation. A preliminary estimate of 2 percent positron decay may be deduced from the data. Beta-gamma coincidences were measured with the use of a Geiger counter and a NaI-scintillation counter. Comparison of the β - γ coincidence counts per β -count for I¹²⁶ and

for Au¹⁹⁸ (which has a 960-kev beta-transition followed by a 411-kev gamma-ray) and coincidence absorption measurements showed the following: (1) the 1.24-Mev betas do not coincide with gamma-rays and therefore presumably go to the ground state of Xe¹²⁶; (2) the 0.85-Mev betas are followed by a single gamma-ray transition, probably the observed 0.38-Mev gamma. The γ - γ coincidences and the 640-kev gamma-ray must therefore be in the electron capture branch. The coincidence work was done in collaboration with Dr. A. W. Sunyar.

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Angular Distributions of the Two Gamma-Rays from $Li^7(p, \gamma)Be^8$ Reaction*

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HE separate angular distributions of the 14- and 17-Mev gamma-rays from $Li^{7}(p, \gamma)Be^{8}$ reaction have been studied at two proton bombarding energies. From these measurements it has been possible to draw certain conclusions concerning the angular momentum of the energy levels involved. Earlier published experiments¹⁻⁵ have left considerable uncertainty both with regard to the angular distribution and to the nature of these levels.

The two gamma-ray lines were resolved using the gamma-ray pair spectrometer.⁶ The angular aperture of the spectrometer was limited to ± 7 degrees; then, by varying the angular position of the spectrometer with respect to the target and incident proton beam, it was possible to measure the intensity of each of the lines as a function of angle. A Pb radiator of 84 mg/cm² was used in the spectrometer.

The two proton bombarding energies were 0.50 ± 0.03 Mev and 1.15 ± 0.05 Mev. At the bombarding energy of 0.50 Mev a thick target was used, yielding mostly 440-kev resonance radiation. At the proton energy of 1.15 Mev a target thickness of 250 ± 50 kev was used. This yields mainly "nonresonant" radiation. For each proton energy, observations were made of the intensities of the two gamma-ray components at three angles of emission. Two stationary Geiger counters were used to monitor the primary beam intensity.

The data may be considered in two different ways. One may first obtain the ratio of intensity of the 17- to 14-Mev component at each angle of observation. For isotropic distributions this ratio should be independent of the angle. One may also obtain the variation of intensity of the individual gamma-ray components as a function of the angle, thus obtaining the separate angular distribution functions. Because of corrections for gamma-ray absorption in the target and the necessity to rely on the counters for long time stability, these values are less accurate. The results are given in Table I.

Within the limits of error, the resonant radiation is isotropic for both components. This is in agreement with the results of Devons and collaborators,^{1, 2} who find that under these conditions the total radiation differs from isotropy by only 4 percent and also that a pair spectrometer measurement indicates each component to be isotropic to within the accuracy of their measurements (about 20 percent). These results are in sharp disagreement with those of Nabholtz, Stoll, and Wäffler,3 who obtained the value of 2.2 for R_{90}/R_0 using photographic plates and the $C^{12}(\gamma, 3\alpha)$ reaction for energy discrimination.

TABLE I. Angular distributions of the 14-Mev and 17-Mev gamma-rays. Relative intensity = I. Ratio of relative intensity, $R = I_{17} \text{ Mev}/I_{14} \text{ Mev}$. Absolute intensity ratio, 17 Mev/14 Mev = cR.

Type of radiation	Angle in degrees	I14 Mev	I17 Mev	R	с*
$\frac{440\text{-kev resonance}}{\text{radiation.}}$ $E_p = 0.50 \text{ Mev}$	0 35 70	$\begin{array}{c} 1.00 \pm 0.07 \\ 0.97 \pm 0.07 \\ 0.98 \pm 0.07 \end{array}$	1.00 ± 0.07 1.03 ± 0.07 0.99 ± 0.07	1.00 ± 0.06 1.06 ± 0.06 1.01 ± 0.06	1.70±0.20
Nonresonant radiation. $E_p = 1.15$ Mev	0 35 75	$\substack{1.00 \pm 0.05 \\ 0.74 \pm 0.06 \\ 0.77 \pm 0.05}$	$\substack{1.00 \pm 0.05 \\ 0.69 \pm 0.06 \\ 0.56 \pm 0.05}$	$\begin{array}{c} 1.00 \pm 0.05 \\ 0.94 \pm 0.08 \\ 0.72 \pm 0.05 \end{array}$	0.62 ± 0.07

* The value of c, the ratio of absolute intensities in the forward direction, was obtained from a weighted average of the present data and earlier data of Walker and McDaniel. Corrections have been made for the variation with energy of the spectrometer detection efficiency. Resolution width, loss from vertical scattering, and pair cross section have been considered. Stated errors are standard deviations based on statistics and estimated orgidnetic errors. accidental errors.

The nonresonant radiation is strongly anisotropic, and the angular distribution is clearly different for the two gamma-ray components.

Following Devons' arguments concerning the resonance radiation, one may suggest the following level assignments. Because of the isotropy of both components under resonance excitation, the resonance level is produced by s-protons and has an angular momentum J = 1 and odd parity. Since the separate angular distributions of the two lines for nonresonant excitation are anisotropic and different from each other, it is likely, though not necessary, that the 2 lower states have different angular momenta. The long lifetime⁷ of the Be⁸ ground state would seem to favor the assignment of J=2, even, to that state and J=0, even, to the 3-Mev level. The separate angular distributions of the two components of the nonresonant radiation are not consistent with the angular distribution $(1+a\cos^2\theta)$ which would exist if p-protons alone caused this radiation. It is therefore likely that the s-protons which produce the resonant radiation, or d-protons forming some higher level, are interfering with the p-protons in producing the nonresonant radiation at these energies.

* Assisted by the ONR. ¹S. Devons and M. G. N. Hine, Proc. Roy. Soc. (London) **A199**, 56 (1949). ²S. Devons and G. R. Lindsey, Proc. Phys. Soc. (London) **A63**, 1202

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Microwave Collision Diameters and Associated Quadrupole Moments*

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IN recent papers^{1,2} Smith and Howard reported an investi-gation of the broadening of the 2.2 gation of the broadening of the 3-3 inversion line of NH₃ by other gases. From the data obtained, they calculated a collision diameter for the NH3-foreign gas collision. In the cases for which this diameter was markedly greater than that obtained from kinetic theory, the collision was ascribed to the interaction of the NH₃ dipole moment and a permanent electric quadrupole moment (averaged over the rotation)² of the foreign gas. The quadrupole moments for these gases were calculated.

Anderson³ has since shown that the interaction of the quadrupole moment of NH₃, calculated from its structure, and the dipole induced in the foreign gas gives a collision diameter in good agreement with those molecules whose experimental collision cross sections approached kinetic theory values. However, for N₂ and others this diameter is too small, and the assignment of a permanent electric quadrupole to these molecules seems justified. TABLE I. Collision diameters (b) of NH₂ with various colliding molecules.

Gas	b×10 ⁸ (cm) (micro- wave)	b×10 ⁸ (cm) (kinetic theory)	Polariza- tion (α) $\times 10^{24}$ s	b×10 ⁸ (cm) (Anderson)	Quadrupole moment (Q) $\times 10^{16} (\text{cm}^2)$
C ₂ H ₂	8.79		3.33	3.92	1.1
C ₂ H ₄	6.67	4.79 ^b	4.27	4.09	0.48
C2H6	5.64	4.86 ^b	4.53	4.15	(0.28)
CaHa	9.76	4.50°	10.32	4.85	1.3
N ₂ O	7.32	4.35d	2.99	3.89	0.59
NO	5.64	3.90d	1.72	3.52	0.28
cõ	5.97	3.96 ^d	1.95	3.59	0.34

^a H. A. Stuart, Molekulstruktur (Verlag, Julius Springer, Berlin, 1934).
 ^b E. H. Kennard, Kinetic Theory of Gases (McGraw-Hill Book Company, Inc., New York, 1938), p. 149.
 ^c Landolt-Bornstein, Phys. Chem. Tab., Eg. I (a), p. 105.
 ^d L. Loeb, Kinetic Theory of Gases (McGraw-Hill Book Company, Inc., New York, 1934), p. 551

New York, 1934), p. 651.

The NH₃ 3-3 line breadth has been measured for mixtures of NH3 with several more gases, using the same experimental technique as in reference 1. The results obtained are listed in Table I. For the gases investigated, either the dipole moment or its average over a collision was zero, and the collision may be described by an interaction other than dipole-dipole.

The diameters calculated from Anderson's formula are quite a bit lower than those found from broadening data, and except for C_2H_6 a quadrupole moment has been given. For C_2H_6 the microwave diameter is sufficiently close to the kinetic theory diameter that the quadrupole moment is given only as an upper limit.

For the C₂H₂, C₂H₄, C₂H₆ series the microwave diameter and associated quadrupole moment are seen to increase with the order of the $C-\bar{C}$ bond, indicating an increase in the asymmetry of the charge distribution. The moments obtained for NO and CO are very close to that for N₂ found by Smith and Howard, while the quadrupole moment for N2O agrees very well with their values for COS, CS₂, and CO₂.

Further work in the interpretation of these data is in progress.

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¹ R. R. Howard and W. V. Smith, Phys. Rev. **79**, 128 (1950).
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Survey Experiment on Elastic Scattering*

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SING cameras similar to those described by Fulbright and Bush¹ two survey experiments on the elastic scattering of protons have been performed.

In one, the angular dependence of the differential elastic cross section in the range 26° to 106° was measured at 15° intervals for the elements W, Pd, Ni, and Al. The proton beam energy, as determined from the magnetic field strength and the camera geometry, was 18.6 ± 0.4 Mev. The total energy spread at the halfmaximum intensity of the incident beam was less than 1 Mev. The circulating beam was scattered by wires 0.001 in. in diameter.

The camera used in this experiment had three slits. In a single run the relative values of the elastic scattering at three angles could be obtained. The measurements were extended to other angles by overlapping sets of observations.

The relative values of the cross sections were then determined by counting the number of tracks per unit area in the elastic lines which appear on the developed film.

The results of this experiment are shown in Fig. 1. We have plotted $\ln(\sigma_o/\sigma_R)$ vs θ , where σ_o is the observed cross section, σ_R is that expected for Rutherford scattering, and θ is the angle

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