

FIG. 1. Magnetic resonance absorption spectra of NO2 at 9360 Mc/sec. The circles are output galvanometer readings; galvanometer deflections are proportional to the first derivative of the absorption contour. The upper figure shows the three-peaked band at 5.0 mm Hg and 290°K with a modulation amplitude of 8.4 oersteds. The lower figure is for 1.3 mm Hg, 290°K, and 2.1 oersteds.

where the coupling energies A and B may depend on J and K. The magnetic moment generated by the molecular rotation is assumed to be proportional to J, the total rotational angular momentum quantum number. Using the approximation that the rotational energies are given by the symmetric top expression,^{3, 5, 6} F

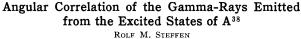
$$=0.417J(J+1)+5.87K^2$$
 cm⁻¹,

the partition function is 2700 at room temperature. It is evident that several hundred rotational levels contribute lines of low intensity, and that an observed resonance is the sum of a number of such.

Selecting the transition $\Delta J = 0$, $\Delta M_s = 1$, $\Delta M_I = \Delta M_J = 0$, and summing over J, K, M_I , and M_J (with A independent of J and K), one predicts a three-peaked band of absorption lines with the spacings between the peaks equal to A. The envelope of line intensity retains this shape at low pressure.

The general features of the partially resolved spectrum are predicted by this model with B approximately equal to A/2. The sharpness of the lines indicates that the spin-rotation term is almost independent of J and K. The coupling of the spin of the odd electron to the spin of the N¹⁴ nucleus is A = 132 Mc/sec.

* This study, together with that of reference 1, was part of a dissertation by one oi us (JGC) presented for the degree of Doctor of Philosophy in Yale University.
† Du Pont Company Fellow in Physics (1949-50); present address: University of Buffalo, Buffalo, New York.
‡ Assisted by the ONR.
‡ R. Beringer and J. G. Castle, Jr., Phys. Rev. 78, 581 (1950).
² Sce W. F. Giaque and J. D. Kemp, J. Chem. Phys. 6, 40 (1938).
³ K. B. McAriee, Phys. Rev. 78, 340 (1950).
⁴ A. K. Mann and P. Kusch, Phys. Rev. 77, 435 (1950).
⁵ G. Herzberg, Infrared and Raman Spectra of Polyatomic Molecules (D. Van Nostrand Company, Inc., New York, 1945).
⁶ Claesson, Donahue, and Schomaker, J. Chem. Phys. 16, 207 (1948).



Department of Physics, Purdue University,* Lafayette, Indiana August 17, 1950

 $\mathbf{F}_{\text{normalized}}^{\text{ROM}}$ a study of the beta-spectra¹ and the gamma-radiation² of the 38-min. Cl³⁸ it can be concluded that the two excited states of A³⁸ at 2.15 Mev and 3.75 Mev have angular momenta of $J_1=1, 2, \text{ or } 3 \text{ and } J_2=3$ respectively, the latter having a different parity than the lower excited and the ground state.

In order to obtain more information about the excited levels in A³⁸ the angular correlation of the gamma-rays successively emitted from those levels has been measured. The apparatus consisted of two sintillation counters using anthracene crystals as phosphors and RCA 5189 photo-multiplier tubes. To avoid disturbing coin-

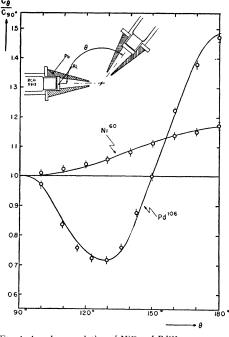


FIG. 1. Angular correlation of Ni⁶⁰ and Pd¹⁰⁶ gamma-rays.

cidence counts due to Compton scattering from one counter to the other the crystals were shielded by two lead cones and in addition covered with $\frac{1}{16}$ -in. lead foils. The angular resolution of the instrument as determined by observations on annihilation radiation was 10°30'. The resolving time of the coincidence amplifier used was 5.8×10^{-8} sec. The arrangement was tested by a careful measurement of the angular correlation of the gammaradiation emitted from the excited states of Ni⁶⁰ (decay²⁸ of Co⁶⁰⁺) and Pd¹⁰⁶ (decay^{2a} of Rh¹⁰⁶) (Fig. 1) and was found to be in perfect agreement with the results reported by Brady and Deutsch.³ The solid line represent the correlation functions

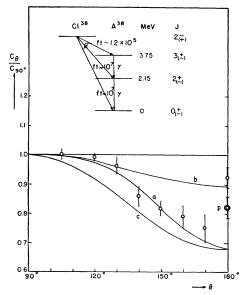


FIG. 2. Angular correlation of gamma-rays emitted from the A³⁸ nucleus. FIG. 2. Angular correlation of gamma-rays emitted from the A* nucleus. Curve a: Correlation function for quadrupole-quadrupole transition between states of J=3, 2, and 0. Curve b: Correlation function for quad-rupole-dipole transition and J=3, 1, 0 and dipole-quadrupole J=3, 2, 0. Curve c: Correlation function Q-D, J=1, 1, 0; D-Q, J=1, 2, 0; and D-D, J=1, 1, 0.

 $f(\theta) = 1 + 0.125 \cos^2\theta + 0.042 \cos^4\theta$ (Ni⁶⁰) and $f(\theta) = 1 - 1.5 \cos^2\theta$ $+2\cos^4\theta(\mathrm{Pd}^{106})$, corrected for the finite angular resolution of the instrument. The above angular correlation of the Ni⁶⁰ gamma-rays is to be expected for a quadrupole-quadrupole transition between states of angular momenta J=4, 2, and 0. No theoretical explanations for the angular distribution of the Pd106 gamma-rays has been given yet.5,6

The angular correlation of the gamma-rays of A³⁸ has been measured using sources of Cl³⁸ produced by bombarding LiCl with 10-Mev deuterons in the Purdue cyclotron. The results are shown in Fig. 2. The measured points follow, within the experimental error, the angular correlation function $f(\theta) = 1 - \frac{1}{3}\cos^4\theta$. For comparison all the other known correlation functions⁴ for quadrupole or dipole transitions are plotted which give smaller emission probabilities at 180° than at 90°. The deviation from the theoretical curve near 180° may be partly due to internal pair production chiefly of the 2.15-Mev gamma-transition. Nevertheless the correction applied (point P in Fig. 2) using Rose's calculations⁷ of the coefficient of internal pair production assuming electric quadrupole transition (see discussion below) is not sufficient to bring the point down to the expected place. Part of this effect is probably due to positron annihilation produced by pair formation of the gammarays in the absorbers, which are necessary to stop the rather energetic beta-particles from Cl38.

The correlation function $1-\frac{1}{3}\cos^4\theta$ is characteristic for two quadrupole quanta and angular momenta J=3, 2, 0 respectively of the A³⁸ levels involved. The parity of the two excited states must be equal since otherwise electric dipole radiation would be possible.

Measurements of Myers and Wattenberg² indicating that the direct transition from the second excited level to the ground state of A³⁸ occurs in less than 3×10^{-4} of the disintegrations make the assignment of the same parity of the second excited level and the ground state of A³⁸ necessary. It must therefore be assumed that both transitions are by electric quadrupole radiation and that all three states of A³⁸ involved have the same parity. Even if mixtures of different multipole radiations are admitted giving different correlation functions due to interference effects8 no other assignments can be found which are compatible with selection rules and the abundance of the cross-over transitions.

According to the measurements of Langer¹ the beta-transition from Cl³⁸ to the ground state of A³⁸ having zero spin is once-forbidden involving a spin change of two units and change of parity suggesting a spin of 2 for Cl³⁸ and odd parity if we assume an even parity of the ground state of A³⁸. The intermediate group of electrons has a *ft*-value of $\sim 10^7$ indicating a once-forbidden transition having a spin change of 0 or 1. This indicates an angular momentum of 1, 2, or 3 and even parity in agreement with the results of our correlation measurements $(J_1=2)$. The low energy betaspectrum is allowed according to its *ft*-value of 1.2×10^5 . The selection rules for allowed beta-transitions ($\Delta J = 0, 1, n_0$) give an angular momentum $J_2=2, 3$, or 4 of the second excited state of A^{38} and odd parity. The angular momentum 3 is also found from our measurements, but an odd parity of this level is not compatible with our results, which indicate the same parity for all levels of A³⁸. The interpretation of the angular correlation would suggest therefore that the low energy spectrum in spite of its small ft-value of 1.2×10^5 is once-forbidden. More direct information concerning the parities of the excited levels of A38 will be obtained by polarization-correlations experiments which are intended to be performed in this laboratory

We wish to thank Dr. D. J. Tendam for his help in bombarding the samples in the Purdue cyclotron.

- * Supported by the ONR.
 ¹ L. M. Langer, Phys. Rev. 77, 50 (1950).
 ² V. Myers and A. Wattenberg, Phys. Rev. 75, 992 (1949).
 ² Obtained from the Isotope Division, AEC, Oak Ridge, Tennessee.
 ³ E. L. Brady and M. Deutsch, Phys. Rev. 78, 558 (1950).
 ⁴ D. R. Hamilton, Phys. Rev. 58, 122 (1940). D. L. Falkoff, thesis,
 ⁴ D. S. Ling and D. L. Falkoff, Phys. Rev. 76, 430 (1949).
 ⁶ A. Spiers, Phys. Rev. 76, 575 (1950).
 ⁷ M. E. Rose, Phys. Rev. 76, 678 (1949).
 ⁸ D. S. Ling and D. L. Falkoff, Phys. Rev. 74, 1224 (1948).

Erratum: On the Primary Cosmic-Ray Spectrum

[Phys. Rev. 78, 819 (1950)] J. A. VAN ALLEN AND S. F. SINGER Applied Physics Laboratory, Johns Hopkins University, Silver Spring, Maryland

N the course of publication certain errors occurred in this Letter to the Editor which requires correction. The units of geomagnetic cut-off, as given in terms of the momentum/charge ratio, pc/Ze, are properly billion volts (Bv), and not billion electron volts (Bev) as printed, and should be so understood throughout, including Tables I and II. Line 13 of the text should read "... charged component to geomagnetic latitude." The heading of column 3 of Table II should be "pc/Ze."

Apparent Error in the Measured Mass of S³²

A. S. PENFOLD Belatron Group, University of Saskatchewan, Saskatoon, Saskatchewan, Canada August 14, 1950

X/HILE attempting to calculate the threshold energy for the reaction $S^{32}(\gamma, d)P^{30}$ an apparent anomaly between the measured masses of S32 and Si30 was encountered.

Discussing mass defects in light nuclei, Rosenfeld¹ states, "In the interval $A = 29 \cdots 34$ there is a large unexplained discrepancy between the values derived from nuclear reaction data and those based on the mass spectrograph measurement of S32." The latter mass was used in Rosenfeld's table of mass defects.

This discrepancy can seemingly be attributed to an error in the measured mass of S22. The following calculations are offered in support of this statement.

The reaction energies of five relevant nuclear reactions are given in the isotopic report of Mattauch and Flammersfeld² and in a recent paper on thresholds.3 These values, together with the reaction energy for the reaction $S^{32}(\gamma, d) P^{30}$ which was measured in this laboratory, are listed in Table I.

Seaborg's tables⁴ give 3.85 and 3.50 Mev for the β -decay energies of S³¹ and P³⁰. Both values were determined by magnetic spectrometer measurements.

Figure 1 illustrates the situation. The reactions considered are shown by solid arrows and the β -decays by dotted arrows.

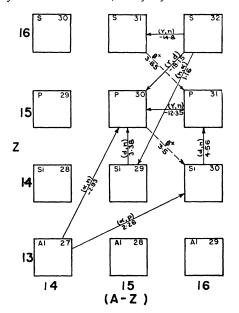


FIG. 1. Sketch showing the reactions, reaction energies, β -decays, and decay energies used in the calculations to determine the mass of S³².