The Spin and Magnetic Moment of Ti⁴⁷ and Ti⁴⁹ and the Magnetic Moment of Ge⁷³

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The spin and magnetic moment of Ti⁴⁷ and Ti⁴⁹ have been measured by nuclear induction. The results are: $I(\text{Ti}^{47}) = 5/2$, $I(\text{Ti}^{49}) = 7/2$, $\mu(\text{Ti}^{47}) = -(0.78706 \pm 0.0001)$ nm, $\mu(\text{Ti}^{49}) = -(1.1022 \pm 0.0002)$ nm. The magnetic moment of Ge⁷³ has been measured by nuclear induction with the result: μ (Ge⁷³) = -(0.87675) ± 0.00012) nm. These results are compared to the predictions of the shell model of nuclear structure.

I. TITANIUM 47 AND 49

NUCLEAR induction signal in TiCl₄ which showed a structure has been previously reported;¹ this resonance signal was attributed to either Ti⁴⁷ or Ti⁴⁹. Using a nuclear induction spectrometer similar to those described by Proctor² and by Weaver,³ recent measurements made with separated Ti isotopes have shown that the structure of the line is not due to chemical effects as was previously thought but is actually the superposition of the close lying resonance lines of Ti⁴⁷ and Ti⁴⁹. Using two different samples of TiCl₄, one containing 63-percent Ti⁴⁷ and only 0.7percent Ti⁴⁹, the other containing only 1.2-percent Ti⁴⁷ and 77-percent Ti⁴⁹, we have observed in the same magnetic field the ratio of the nuclear magnetic resonance frequencies of Ti⁴⁷ and of Ti⁴⁹ to that of D² in a sample of D₂O containing 1 molar Mn⁺⁺ ions as a paramagnetic catalyst. Our results are

> $\nu(Ti^{47})/\nu(D^2) = 0.36721 \pm 0.00006$, (1)

$$\nu(Ti^{49})/\nu(D^2) = 0.36731 \pm 0.00006.$$
 (2)

The polarity of the Ti⁴⁷ and Ti⁴⁹ signals has been observed to be opposite to that of D^2 , indicating that their magnetic moments are negative. Using a sample of TiCl₄ containing Ti⁴⁷ and Ti⁴⁹ in their natural abundances, we have furthermore clearly resolved the two Ti resonances under conditions of higher resolution than originally used. The frequency ratios are identical to those of (1) and (2) above. We have also observed in the same magnetic field $\nu(Ti^{49})/\nu(Ti^{47}) = 1.00026$ ± 0.00002 . Thus the gyromagnetic ratios of Ti⁴⁷ and Ti⁴⁹ are remarkably close; since the spins are different, as will be shown, this is probably a coincidence.

Ti signals have also been observed in H₂TiF₆ at a frequency approximately 0.1 percent lower than in TiCl₄. This "chemical shift"⁴ has not yet been measured exactly because of the chemical instability of $H_2 TiF_6$, but its existence indicates a possible uncertainty of this order of magnitude in the magnetic moments of Ti^{47} and Ti^{49} reported below in (6) and (7).

the spins I(a) and I(b) of two nuclei a and b may be compared, in principle, by observing the heights H(a), H(b) and the widths W(a), W(b) of the nuclear induction signals for the two nuclei. It can be shown⁶ that if slow passage conditions prevail and if the radiofrequency field is small compared to its saturation value, then when the radiofrequency is the same for both nuclei,

$$R \equiv \frac{I(a)[I(a)+1]}{I(b)[I(b)+1]}$$
$$= \left[\frac{N(b)}{N(a)}\right] \left[\frac{H(a)}{H(b)}\right] \left[\frac{W(a)}{W(b)}\right]^{2} \left[\frac{\gamma(b)}{\gamma(a)}\right]^{3}, \qquad (3)$$

where N(a), N(b) are the relative number of nuclei and $\gamma(a), \gamma(b)$ are the gyromagnetic ratios. Previous spin determinations^{6,7} have compared an unknown spin I(a)to a known spin I(b). However Eq. (3) will also uniquely determine both I(a) and I(b) even if both are unknown [except in the case I(a)=I(b)], if the additional assumption is made that the spins have quantized values. In the present case the line shape of the Ti resonances is different from that of Cl³⁵ in TiCl₄ and a comparison of I(Ti) to $I(Cl^{35})$ is not justified. The line shapes of the Ti⁴⁷ and the Ti⁴⁹ resonances, on the other hand, are identical under widely varying conditions and it is believed that application of Eq. (3) is valid. For a series of 13 measurements of the heights and widths of the Ti⁴⁷ and Ti⁴⁹ resonances in natural abundance $TiCl_4$ we find, using Eq. (3),

$$R = \frac{I(\mathrm{Ti}^{47})[I(\mathrm{Ti}^{47})+1]}{I(\mathrm{Ti}^{49})[I(\mathrm{Ti}^{49})+1]} = 0.568 \pm 0.037, \qquad (4)$$

where we have used the abundance ratio $N(Ti^{49})/$ $N(\text{Ti}^{47}) = 0.743$, which is the average of the reported values.8 Both these Ti isotopes are odd nuclei and it may be safely assumed that the spins are half odd integers. The above experimental result then establishes

According to Bloch's theory⁵ of nuclear induction,

 ¹ Jeffries, Loeliger, and Staub, Phys. Rev. 85, 478 (1952).
 ² W. G. Proctor, Phys. Rev. 79, 35 (1950).
 ³ H. E. Weaver, Phys. Rev. 89, 923 (1953).
 ⁴ W. C. Dickinson, Phys. Rev. 81, 717 (1951).
 ⁵ F. Bloch, Phys. Rev. 70, 406 (1950).

⁶ C. D. Jeffries, Phys. Rev. 90, 1130 (1953).
⁷ W. G. Proctor, Phys. Rev. 79, 35 (1950); F. Alder and F. C. Yu, Phys. Rev. 81, 1067 (1951).
⁸ A. O. Nier, Phys. Rev. 53, 282 (1938); R. F. Hibbs, Oak Ridge National Laboratory Report Y-508, 1949 (unpublished); H. C. Mattraw and C. F. Pachucki, U. S. Atomic Energy Commission Paroet AECUI 1003 (unpublished) Report AECU-1903 (unpublished).

that

$$I(\mathrm{Ti}^{47}) = 5/2$$
, and $I(\mathrm{Ti}^{49}) = 7/2$, (5)

for which the ratio R = 0.555; the nearest other possible R values are excluded: R=0.636 for $I(Ti^{47})=7/2$, $I(\text{Ti}^{49}) = 9/2$ and R = 0.440 for $I(\text{Ti}^{47}) = 7/2$, $I(\text{Ti}^{49})$ =11/2.

Using the results (1), (2) and (5) we find for the magnetic moments, without diamagnetic correction,

$$\mu(\mathrm{Ti}^{47}) = -(0.78706 \pm 0.0001) \text{ nm}, \tag{6}$$

$$\mu(\mathrm{Ti}^{49}) = -(1.1022 \pm 0.0002) \text{ nm}, \tag{7}$$

where we have used in this calculation the ratio $\mu(D^2)/\mu(H) = 0.307015$ as given by Mack⁹ and $\mu(H)$ =2.7925 as determined by Bloch and Jeffries.¹⁰

This experiment assigns an $f_{7/2}$ orbit to the odd neutron in Ti⁴⁹, as anticipated by the nuclear shell model of Mayer¹¹ and of Jensen, Haxel, and Suess.¹² For Ti⁴⁷, with I=5/2, the situation is more complicated: the single-particle model would predict an $f_{5/2}$ orbit for the odd neutron, whereas this experiment would assign a $d_{5/2}$ orbit. This "discrepancy" is quite analogous to that for the odd-proton nuclei Na²³ and Mn⁵⁵ as discussed by Mayer,¹¹ where the empirical ground level can be understood by taking into account all the protons outside the closed shells instead of just the single odd proton. A theoretical explanation has been given by Kurath¹³ and by Talmi.¹⁴ It is supposed that Mn^{55} with 25 protons has a $D_{5/2}$ ground state obtained by the jj coupling of the five protons in the incomplete $f_{7/2}$ subshell. This experiment shows that Ti⁴⁷, with 25 neutrons, behaves similarly, as anticipated by Rosenfeld¹⁵ and others. The five neutrons in the incomplete $f_{7/2}$ subshell can couple to a $D_{5/2}$ state; a simple *jj* coupling calculation of the resultant magnetic moment yields $\mu = -1.36$ nm, which is nearer to the experimentally observed value than either of the Schmidt limits.

The magnetic moments of Ti⁴⁷ and Ti⁴⁹ as predicted by the scheme of Schawlow and Townes¹⁶ agree well with our measured values (6) and (7).

II. GERMANIUM 73

Using the nuclear induction spectrometer we have observed a nuclear magnetic resonance in pure GeCl₄ at

⁹ J. E. Mack, Revs. Modern Phys. 22, 64 (1950).
¹⁰ F. Bloch and C. D. Jeffries, Phys. Rev. 80, 305 (1950).
¹¹ M. G. Mayer, Phys. Rev. 78, 16 (1950).
¹² Haxel, Jensen, and Suess, Z. Physik 128, 295 (1950).
¹³ D. Kurath, Phys. Rev. 80, 98 (1950).
¹⁴ I. Talmi, Phys. Rev. 82, 101 (1950).
¹⁵ L. Rosenfeld, Physica 17, 461 (1951).
¹⁶ A. L. Schawlow and C. H. Townes, Phys. Rev. 82, 268 (1951).
The calculated value for μ(Ti⁴⁷) is in error and should be -0.80 (private communication). (private communication).

a frequency of 1.48 Mc/sec in a field of 10 000 gauss. We ascribe this resonance to Ge⁷³, the only stable odd isotope of germanium. The observed signal-to-noise ratio agrees approximately with that calculated for Ge⁷³ using the spin value of 9/2 as measured by Townes, Mays, and Dailey.¹⁷ We have been unable to observe the Cl³⁵ or Cl³⁷ resonance signals in GeCl₄, presumably because of excessive electric quadrupolar broadening of the line. However by comparing the Ge⁷³ signal to the Cl³⁵ signal in TiCl₄ we observe the magnetic moment of Ge⁷³ to be negative. This is in agreement with the prediction of the Mayer-Jensen nuclear shell model, which assigns a $g_{9/2}$ orbit to the odd neutron in Ge⁷³. In the same magnetic field the ratio of the nuclear resonance frequency of Ge⁷³ in GeCl₄ to that of Cl³⁵ in TiCl₄ is observed to be

$$\nu(\text{Ge}^{73})/\nu(\text{Cl}^{35}) = 0.35572 \pm 0.00004.$$

We have observed a slight "chemical shift"⁴ between the Cl³⁵ resonance frequency in TiCl₄ and in an aqueous solution of RbCl. The ratio of the resonance frequencies is found to be 1.00088 ± 0.000025 in these two compounds, respectively. Using this correction and the ratio $\nu(Cl^{35})/\nu(D^2)$ in RbCl+D₂O as measured by Walchli, Leyshon, and Scheitlin¹⁸ we find

$$\nu(\text{Ge}^{73})/\nu(\text{D}^2) = 0.22725 \pm 0.00003.$$
 (8)

From this we find for the nuclear magnetic moment of Ge⁷³, without diamagnetic correction,

$$\mu(\text{Ge}^{73}) = -(0.87675 \pm 0.00012) \text{ nm}, \qquad (9)$$

where we have used in this calculation the ratio $\mu(D^2)/\mu(H)$ and $\mu(H)$ as given above, and the value 9/2 for the spin of Ge⁷³ as measured by Townes et al.¹⁷ Our measured value of $\mu(Ge^{73})$ is considerably less than the predicted values.¹⁹ We have also searched for the Ge73 resonance in the powdered element without success. It should be pointed out that possible "chemical shifts"4 can make the result (9) uncertain, to about 0.1 percent.

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 ¹⁷ Townes, Mays, and Dailey, Phys. Rev. 76, 700 (1949).
 ¹⁸ Walchli, Leyshon, and Scheitlin, Phys. Rev. 85, 922 (1952).
 ¹⁹ A. L. Schawlow and C. H. Townes, Phys. Rev. 82, 268 (1951);
 J. P. Davidson, Phys. Rev. 85, 432 (1952); G. J. Bene, J. phys. et radium 13, 161 (1952).