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Measurement of the Magnetic Moments of the Microsecond-isomers in ⁷³As and ²⁰⁶Pb ⁺

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The isomeric state of $^{73}\mathrm{As}$ (426 keV, 5.8 $\mu\mathrm{sec}$) was produced and aligned by the reaction $^{71}\mathrm{Ga}\left(\alpha,2\mathrm{n}\right)$ in a liquid metal target. The anisotropies of the depopulating $\gamma\text{-rays}$ were measured vs. the dc magnetic field applied perpendicular to the beam-detector plane. From the attenuation and rotation, we obtain $g=+1.03\pm0.11$. Limits of $\delta^2<10^{-2}$ can be set to the multipole admixtures in both $\gamma\text{-transitions}$. The same technique was applied to the 123 $\mu\mathrm{sec}$ isomer of $^{206}\mathrm{Pb}$ (2200 keV) produced by $^{204}\mathrm{Hg}\left(\alpha,2\mathrm{n}\right)$. After estimating the relaxation time, $g=-\left(0.035\pm0.020\right)$ is obtained.

Considerable alignment is given to the low-energy, high-spin states populated by (α,xn) reactions. Without perturbation, the time integrated angular distribution of gamma radiation emitted by such states is described as $W(\Theta) = 1 + \sum A_k P_k(\cos\Theta)$, $k = 2, 4, \ldots$. With $A_k = B_k F_k$, the coefficients B_k depend upon the degree of alignment, and the F_k are determined by the multipole character of the transition and by the spins involved. If one applies a static magnetic field H_\perp perpendicular to the beam-detector plane, the angular distribution becomes 1

$$W(H_{\perp}, \Theta) = 1 + \sum \frac{b_k/b_0}{1 + (k\omega_{\perp}\tau)^2} \left[\cos k \Theta - k\omega_{\perp}\tau \sin k \Theta\right];$$

$$k = 2, 4, \dots, \tag{1}$$

where $b_0 = 1 + (1/4) A_2 + (9/64) A_4 \dots,$ $b_2 = (3/4) A_2 + (5/16) A_4 \dots,$ $b_4 = (35/64) A_4 \dots,$

and
$$\omega_{\perp} = g \,\mu_n \, H_{\perp}/\hbar$$
, $g = \mu_I/I \,\mu_n$. (2)

It is to be noted that a) no static interaction other than the one given by H_{\perp} must be acting during the time of measurement; b) the time constant τ with which the anisotropic emission of the γ -rays decays has to be known.

The experiments were performed at the 88" cyclotron at Berkeley using the reactions $^{71}\text{Ga}(\alpha,2n)\,^{73}\text{As}$ or $^{204}\text{Hg}(\alpha,2n)\,^{206}\text{Pb}$ and isotopically enriched liquid metal targets. The Ge(Li) detectors and the electronics used were very similar to the setup described in 2 . The variable field H_{\perp} was known to about $\pm\,2\%$.

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- ¹ R. M. Steffen, Adv. Phys. 4, 293 [1955].

The ⁷³As isomer decays by two γ -rays in cascade, 360 keV $(9/2^+ \rightarrow 5/2^-, E2)$ and 66 keV $(5/2^- \rightarrow 3/2^-, M1)$. From an angular distribution measurement $(H_{\perp} = 0)$, and from the data of Fig. 1 we get

$$A_2(360) = +0.34(4);$$

 $A_4(360) = -0.05(5);$
 $A_2(66) = -0.20(5).$ (3)

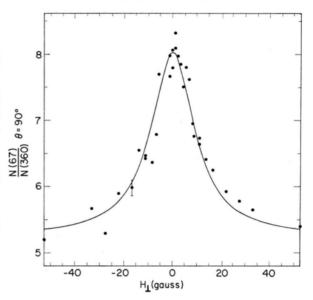


Fig. 1. Ratio of γ -ray intensities in the 90° detector vs. magnetic field H_{\perp} for the ⁷³As 6 μ sec isomer. The statistical error is indicated only once. The curve is from a least squares fit.

Fig. 1 shows the change of the γ -intensities with H_{\perp} . From this measurement and one with one of the detectors at 54°, we obtain through Eq. (1)

$$g = +1.03(11),$$
 (4)

using the average ^{3, 4} $T_{1/2} = 5.8(5)$ µsec. This gives the sign for and agrees with the more precise value

$$|g| = 1.146(7)$$

obtained later 5.

An estimate of the possible combinations of F_2 , F_4 can be derived for a particular gamma transition from the observed coefficients A_2 and A_4 , if no perturbations occur in the decaying state. This is because for k=2 and k=4, the factors 6 α_k describing the loss of alignment during the neutron- and γ -emission turn out to be approximately related as $\alpha_4 \approx (\alpha_2)^3$, when $\alpha_2 > 0.5$. Together with the limits on the mixing parameters δ

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- ³ R. W. HAYWARD and D. D. HOPPES, Phys. Rev. **101**, 93 [1956].
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- ⁶ T. YAMAZAKI, Nucl. Data A 3, 1 [1967].

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given in Ref. ³ and Ref. ⁷, we thus obtain the region of possible values δ (360) and δ (66) which is displayed in Fig. 2.

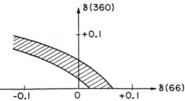


Fig. 2. Mixing ratios $\delta(360)$ and $\delta(66)$ for the two γ -rays depopulating the 6 μ sec isomer in ⁷³As $(9/2^+ \rightarrow 5/2^-, \text{ predominantly E2}$ and $5/2^- \rightarrow 3/2^-, \text{ predominantly M1}, \text{ respectively}).$

For the isomeric state in ^{206}Pb , the change of the γ -ray intensities with H_{\perp} is displayed in Fig. 3. A least squares fit of Eq. (1) to these data, neglecting the A_4 and higher terms, gave

$$g \cdot \tau = -(3.0 \pm 1.0) \cdot 10^{-6} \text{ sec.}$$
 (5)

For each of the γ -transitions, the value of A_2 is reduced to about 0.5 of the value for maximum alignment and lowest possible multipole character in all transitions.

For ^{206}Pb , relaxation in the isomeric state may reduce the time during which the interaction $g~\mu_n~H_\perp$ is effectively observed below the nuclear lifetime 8

$$\tau = 177.9(1.6) \mu sec.$$

To estimate this effect, we assume an effective relaxation time T_r for the P_2 term; the coefficients A_2 are then reduced by the factor $T_r/(\tau+T_r)$, and so is the g factor entering Eq. (2). We find magnetic relaxation negligible (see 9), $T_r \gtrsim \tau$, and

$$g = -0.035(20). (6)$$

The g factor was calculated for a $p_{1/2}i_{13/2}$ neutron hole configuration as g=-0.06, using the g factors of the $^{207}\mathrm{Pb}$ and $^{197}\mathrm{Hg}$ ground states (-0.20 with the Schmidt values). The use of more accurate wave functions 10 does not improve the agreement with experiment, giving g=-0.08.

M. TATCHER and H. LINDEMAN, Nucl. Instr. Methods 61, 58 [1968]. The magnetic moment of the ²⁰⁶Pb-isomer is being determined more precisely by MAIER et al. using the stroboscopic observation of the Larmor precession ¹¹.

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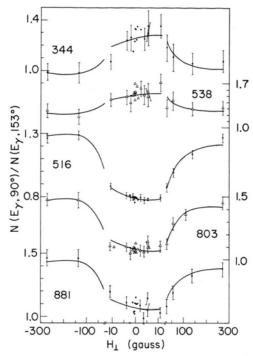


Fig. 3. Ratio of γ -ray intensities vs. magnetic field for the $^{206}{\rm Pb}$ 123 $\mu{\rm sec}$ isomer. The γ -energies (in keV) are given between the data and the ordinate scale applicable. Note the changes in abscissa scale. The points in the region $H_{\perp}\approx 0$ have nearly the same errors as the points farther out. The curves are from a least squares fit.

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