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The Electric Quadrupole Moment of the 2.083 MeV State of ¹⁴⁰Ce Derived from a TDPAC-Measurement

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We studied the rotation of 329 keV-487 keV gamma-gamma angular correlation in the β^- -decay of ¹⁴⁰La (see Fig. 1) in the crystalline electric field gradient of lanthanum-magnesium double-nitrate (La₂(NO₃)₆Mg₃(NO₃)₆ · 24 H₂O) by time differential measurement. A plot of the asymmetry ratio

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$$R(t) = 2 \frac{W(180^\circ, t) - W(90^\circ, t)}{W(180^\circ, t) + W(90^\circ, t)}$$

is shown in Fig. 2. The positive sign of R(t) for $t \to 0$ is explained by a background of the prompt coincidences between the gamma lines of 487 keV and 1598 keV, since the anisotropy of a $4^+ - 2^+ - 0^+$ angular correlation is positive. Figure 2 exhibits the beginning of a very slow spin rotation. In order to derive the interaction frequency we fitted R(t) by the theoretical spin rotation function for a static polycrystalline electric interaction in an axially symmetric field gradient.



Fig. 1. Decay scheme of ¹⁴⁰La.



Fig. 2. Spin rotation observed in the 2083 keV 4⁺-state of ¹⁴⁰Ce in an environment of lanthanummagnesium double nitrate.

The result is

$$\omega_Q = \left| \frac{\mathrm{e}Q_I \langle V_{zz} \rangle}{4I(2I-1)\hbar} \right| = 1.17_{15} \mathrm{Mc/s}$$

The electric field gradient at the nuclear site of La in lanthanum-magnesium double-nitrate is derived from an NMR experiment with the stable nucleus ¹³⁹La in the same environment¹⁾ by using for the quadrupole moment of the ground state of ¹³⁹La the value:²⁾

$$Q_{7/2}(^{139}\text{La}) = +0.230_{10} \text{ b}$$

The cerium is produced by the β^{-} -decay of ¹⁴⁰La primarily in the Ce⁴⁺ charge state and we assume³⁾ that it remains in this state (with an empty 4f-shell) for the time of observation of the spin rotation.

In the evaluation of the electric quadrupole moment of the 2.083 MeV state of ¹⁴⁰Ce we took into account the small difference between the Sternheimer correction factors γ_{∞} of La³⁺ and Ce⁴⁺ and obtained:

$$Q_{4^{+}}(^{140}\text{Ce}) = 0.404_{80} \text{ b.}$$

From the measured g-factor of the same state it was

concluded³⁾ that the 4⁺ state of ¹⁴⁰Ce is predominantly the two-proton configuration (the neutron number is magic N = 82):

$$(g_{7/2}, d_{5/2})_{4^+},$$

whereas the ground state of ¹³⁹La which was used for the calibration is a $g_{7/2}$ single proton state (also with N = 82). We compared both quadrupole moments with the prediction of the shell model:

$$Q_{I}(g_{7/2})_{7/2^{+}} = -0.6667\langle j_{1}|r_{1}^{2}|j_{1}\rangle$$

$$Q_{I}(g_{7/2}, d_{5/2})_{4^{+}} = -0.2415\langle j_{1}|r_{1}^{2}|j_{1}\rangle$$

$$+ 0.1203\langle j_{2}|r_{2}^{2}|j_{2}\rangle$$

$$\approx -0.1212\langle j_{1}|r_{1}^{2}|j_{1}\rangle.$$

The shell model predicts that the quadrupole moment of the single proton state is by a factor of five larger than that of the two particle state whereas the experimental values have about the same magnitude. We take this as a remarkable experimental indication for the fact that in the actual nuclear structure many particles contribute to the nuclear spin.

As the accuracy of our experimental result is limited by the very slow rotation, we looked for another environment which produces a larger electric field gradient at the nuclear site. In recent⁵ TDPACmeasurements with antiferroelectric substances of the perovskit structure such a large electric field gradient was observed in PbTiO₃. We have therefore tried to use this lattice as an environment for the ¹⁴⁰La activity. Preliminary measurements gave indeed a faster spin rotation. The shape of R(t) showed also that the interaction is static and not time-dependent. The authors would like to express their appreciation to Prof. Dr. E. Bodenstedt for suggesting this investigation and many stimulating discussions. This work was supported by the Bundesministerium für Bildung und Wissenschaft. The numerical calculations were performed on the IBM 370/156 Computer of the GMD in Bonn.

References

- Edmonds and Lindop: Proc. Phys. Soc. 187 (1966) 721.
- Feller and Cohen: Nuclear Data Tables A5 (1969) 530.
- The main argument for this assumption is the large chemical activation energy, see also Körner *et al.*: Z. Phys. **173** (1969) 203.
- 4) Feiock and Johnson: Phys. Rev. 187 (1969) 39.
- 5) G. Schäfer, P. Herzog and B. Wolbeck: to be published in Z. Phys.

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