TDPAC STUDIES OF ELECTRIC QUADRUPOLE INTERACTION OF 133Cs IN COMPOUNDS OF BARIUM (BaCr₂O₇ AND BaCrO₄)

B. L. CHAWAT, S. N. A. JAFFREY AND J. VARMA

Department of Physics, S.B.S.H., University of Udaipur, Udaipur, India

ABSTRACT

The Electric Field Gradients (EFG) in BaCr₂O₇ and BaCrO₄ compounds were studied using time differential perturbed angular correlation technique. The observed interaction frequencies are found to be 678 ± 7 MHz and 79 ± 1.3 MHz giving the EFG in the two compounds to be $(14 \pm 1.4) \times 10^{17}$ V/cm² and $(1.6 \pm 0.03) \times 10^{17}$ V/cm² respectively.

1. INTRODUCTION

In recent years, there has been increased interest in the study of the electric field gradients in ionic compounds¹⁻⁷. Weiss⁸ has discussed various factors affecting the EFG in compounds. In the case of barium compounds like BaSO₄, BaCO₃ and Ba(NO₃)₂,

where the 6s electrons of the barium atom contribute in bonding, the major contribution to the EFG is due to ligand ions⁹. Using ¹⁸³Cs as probe Sharma et al.⁹ have measured interaction frequencies in $BaCO_3$ and $BaSO_4$ as 17×10^7 rad/sec and 11×10^7 rad/sec respectively and almost no interaction in $Ba(NO_3)_2$.

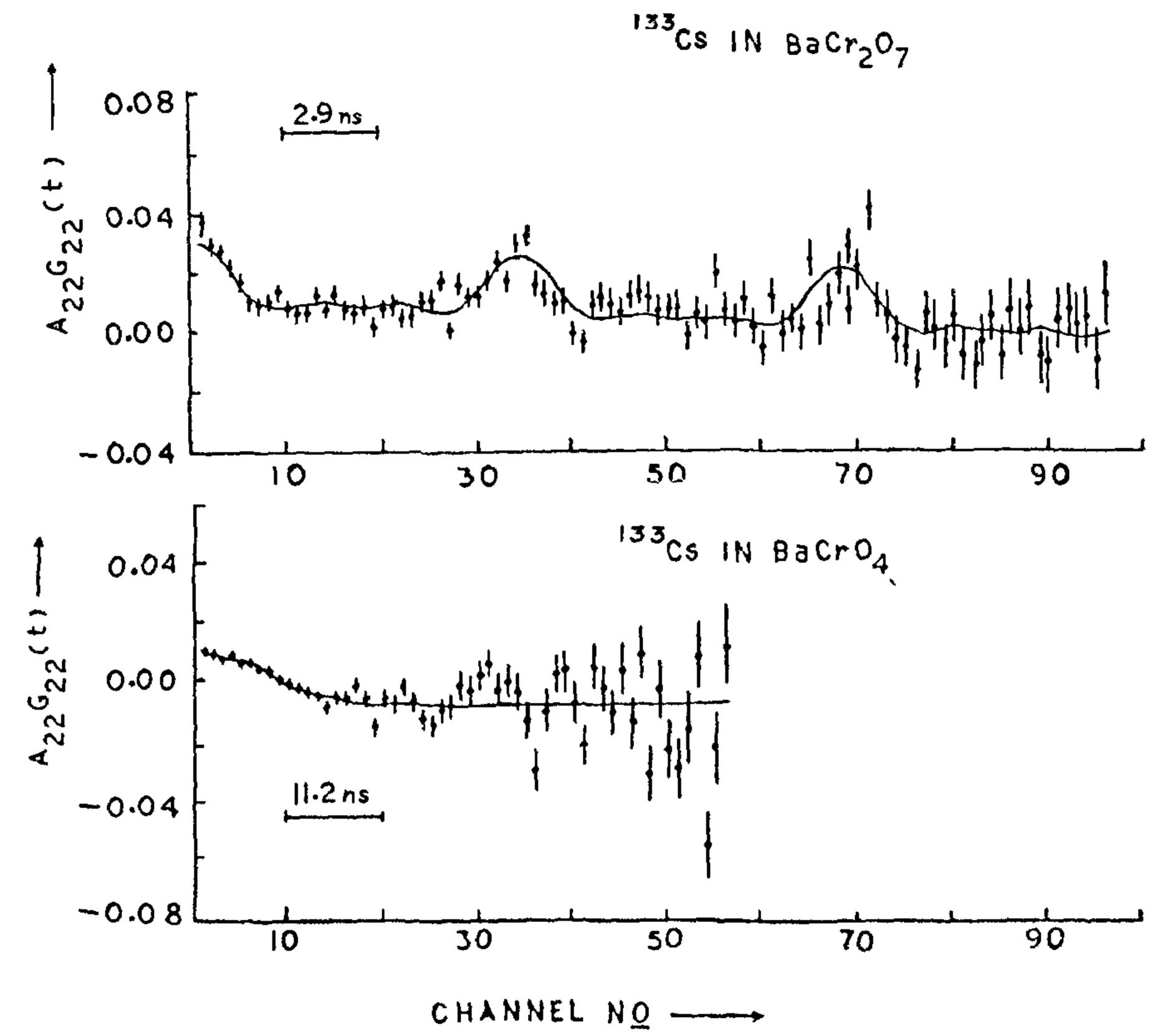


Fig. 1. TDPAC spectrum of barium dichromate and barium chromate at room temperature.

In the present work the quadrupole interaction of the 81 keV (5/2) level of 131 Cs in monoclinic barium-dichiomate (BaCr₂O₇) and rhombohedral barium chromate (BaCrO₄) is studied using time differential perturbed angular correlation (TDPAC) technique. The radioactive 131 Ba decays to 133 Cs via the 356-81 keV gamma ray cascade with intermediate state having half life of 6.35 ns; electric quadrupole moment 2.0 barns as an upper limit⁹⁻¹⁰. The Sternheimer antishielding factor γ_{∞} for 133 Cs is reported to be -157^{11} .

2. EXPERIMENTAL

The compounds BaCr₂O₇ and BaCrO₄ were prepared by chemical processing of radioactive BaCl₂ with potassium—dichromate and potassium chromate in acidic solutions respectively.

The TDPAC measurements were carried out with powder samples at room temperature using a four counter set up with ORTEC TAC and ND 1100 MCA. The time resolution of the system was found to be 2.3 ns (FWHM). After subtracting the chance coincidences, the ratio $A_{22}G_{22}$ (t) was calculated as defined below:

$$A_{22}G_{22}(t) \approx \frac{2 \left[W(180^{\circ}, t) - W(90^{\circ}, t)\right]}{\left[W(180^{\circ}, t) + 2W(90^{\circ}, t)\right]}$$

where W (180°, t) and W (90°, t) are counting rates of coincidences at 180° and 90° respectively and 't' is the time delay between two gamma-ray photons. The data shown in Fig. 1 were computer analysed using standard function for randomly oriented EFG for spin $(5/2)^{12}$, keeping the asymmetry parameter η as floating.

3. RESULTS AND DISCUSSION

The computer fit of the data gave the interaction frequency v_a in barium dichromate to be 678 \pm 7 MHz with a spread of 3% and the asymmetry parameter $\eta = 0$. Combining this with the reported quadrupole moment of the ¹³³Cs excited state, the EFG was obtained to be $(14 \pm 1.4) \times 10^{17} \text{ V/cm}^2$. In the

case of barium chromate the interaction frequency was found to be 79 ± 1.3 MHz but with a large spread of 61%, a fact which does not allow us to derive any meaningful result for the EFG in this case. The value of A_2^{eff} at time t=0 was found to be less than that expected for the cascade and could perhaps be explained by the time dependant electric quadrupole interactions during the charge neutralisation process of the ion $(\tau \text{ ion } \simeq 10^{-8} \text{ sec})$.

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