

Spin determination and role of pairing correlations in the SD bands of Hg isotopes

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Introduction

Since the discovery of first high-spin superdeformed (SD) band in ^{152}Dy by Twin *et al.* [1], many SD bands have been studied considerably in $A \sim 190, 150, 130$ and 80 mass regions. With the advancement in the γ -ray detectors, rich variety of data on SD bands is provided which mainly consists of the intraband γ -transition energies. One of the most crucial challenges in the study of superdeformation is the accurate determination of spins, parities and excitation energies of SD bands. The unavailability of discrete γ -transitions, directly linking SD states to normal deformed states makes the spin assignment of SD bands tedious. The availability of band-head spins is vital for understanding the behavior of SD bands. The moment of inertia (MoI) also proves to be a crucial quantity to study the SD rotational bands. Two types of MoI are available viz. kinematic MoI ($\mathfrak{S}^{(1)}$) and dynamic MoI ($\mathfrak{S}^{(2)}$). The spin independence of $\mathfrak{S}^{(2)}$ makes it the most efficient parameter to explore the SD bands. A characteristic feature of the SD bands near $A \sim 190$ region is the smooth and large increase of the dynamic MoI $\mathfrak{S}^{(2)}$ with rotational frequency $\hbar\omega$. The SD bands near $A \sim 190$ and 130 region display pronounced variations in $\mathfrak{S}^{(2)}$. The smooth rise in $\mathfrak{S}^{(2)}$ for SD bands in $A \sim 190$ region was attributed to the gradual alignment of higher-N orbitals in the presence of pairing correlations [2]. Dadwal and Mittal investigated the identical SD bands in $A \sim 190$ region and the role of pairing correlations was studied [3]. Re-

cently, the flat SD bands in the $A \sim 190$ were investigated and the unusual behavior of $\mathfrak{S}^{(2)}$ was explored [4]. Many theoretical models are available to deduce the reliable spins of the SD bands. In the present analysis, we have deduced the band-head spins of the SD bands in Hg isotopes using the exponential model with pairing attenuation.

Formalism

The exponential model with pairing attenuation was given by Sood and Jain [5] based on the considerations of single particle level densities. Moretto [6] identified the dependence of pairing gap on the angular momentum I as

$$\Delta(I) = \Delta_0 \left(1 - \frac{I}{I_c}\right)^{1/\nu}. \quad (1)$$

Sood and Jain [5] included the explicit spin dependence of pairing gap in addition to the the exponential dependence of moment of inertia on pairing gap and obtained the rotational energy expression as

$$E(I) = \frac{\hbar^2}{2\mathfrak{S}_{(Exp\sigma)}} I(I+1) e^{[\Delta_0(1 - \frac{I}{I_c})^{1/2}]}, \quad (2)$$

where $\mathfrak{S}_{(Exp\sigma)}$ and Δ_0 are the parameters and I_c is adopted to be the value where the moment of inertia achieves a maximum value.

$$E_\gamma = E(I) - E(I-2). \quad (3)$$

The γ -ray transition energy E_γ obtained from the exponential model can be expressed using Eq. 2 and Eq. 3, in the following form

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$$E_\gamma = \frac{\hbar^2}{2\mathfrak{S}_{(Exp0)}} \times (I(I+1)e^{\Delta_0\sqrt{1-\frac{I}{I_c}}} - (I-2)(I-1)e^{\Delta_0\sqrt{1-\frac{(I-2)}{I_c}}}) \quad (4)$$

The band-head moment of inertia (\mathfrak{S}_0) are calculated using

$$\frac{2\mathfrak{S}_0}{\hbar^2} = \frac{2\mathfrak{S}_{(Exp0)}}{\hbar^2} e^{(-\Delta_0)}. \quad (5)$$

Results and Discussion

The observed transition energies [7, 8] of the SD bands in the Hg isotopes have been fitted to the exponential model. The calculated and experimental transition energies agree extremely well whenever the correct band-head spin is assigned and the root-mean-square deviation is found to be minimum. By using the assigned band-head spins and the intraband γ -transitions, the corresponding fitting parameters viz. the band-head MoI (\mathfrak{S}_0) and the pairing gap parameter (Δ_0) are calculated. The fitting parameters along with the deduced spins using the exponential model have been listed in Table I.

We observe that the band-head spins deduced using the exponential model agree well with those listed in Refs. [7, 9] except for $^{191}\text{Hg}(1)$, $^{193}\text{Hg}(5,6)$ and $^{195}\text{Hg}(3)$. The nearly identical intraband γ -transition energies obtained for the pairs $\{^{191}\text{Hg}(2), ^{193}\text{Hg}(2)\}$ and $\{^{191}\text{Hg}(3), ^{193}\text{Hg}(3)\}$ proves that they are identical SD band pairs. The $\mathfrak{S}^{(2)}$ are calculated after firmly assigning the band-head spins and its variation with respect to the rotational frequencies ($\hbar\omega$) is studied, and a comparison with the experiment is made.

Conclusion

In the present analysis, by using the exponential model with pairing attenuation, the band-head spins have been assigned for the SD bands in Hg isotopes and the variation of $\mathfrak{S}^{(2)}$ with the rotational frequencies has been investigated. The exponential model proves

TABLE I: The calculated band-head spins and the corresponding fitting parameters for the SD bands in Hg isotopes using the exponential model.

SD Band	E_γ^{exp} (keV)	Ref. [7]	Exponential model		
		I_0 (\hbar)	I_0 (\hbar)	\mathfrak{S}_0 ($\hbar^2\text{MeV}^{-1}$)	Δ_0
$^{191}\text{Hg}(1)$	310.9	15.5	12.5	86.43	0.37
$^{191}\text{Hg}(2)$	252.4	10.5	10.5	91.76	0.30
$^{191}\text{Hg}(3)$	272.0	11.5	11.5	91.42	0.34
$^{191}\text{Hg}(4)$	280.9	12.5	10.5	81.90	0.39
$^{192}\text{Hg}(1)$	214.4	8	8	85.98	0.43
$^{192}\text{Hg}(2)$	241.2	10	10	90.59	0.43
$^{193}\text{Hg}(1)$	233.2	9.5	9.5	90.33	0.41
$^{193}\text{Hg}(2)$	254.0	10.5	10.5	91.28	0.31
$^{193}\text{Hg}(3)$	233.5	9.5	9.5	90.89	0.34
$^{193}\text{Hg}(4)$	254.0	10.5	10.5	90.98	0.34
$^{193}\text{Hg}(5)$	291.0	13.5	11.5	87.00	0.36
$^{193}\text{Hg}(6)$	240.5	10.5	9.5	89.42	0.24
$^{194}\text{Hg}(2)$	200.8	8	8	91.31	0.34
$^{195}\text{Hg}(1)$	294.0	12.5	12.5	90.38	0.40
$^{195}\text{Hg}(2)$	273.9	11.5	11.5	90.66	0.39
$^{195}\text{Hg}(3)$	244.0	10.5	9.5	87.26	0.36
$^{195}\text{Hg}(4)$	341.9	15.5	15.5	95.53	0.24

to be a powerful method to deduce the band-head spins and to study the general behavior of SD bands in Hg isotopes.

Acknowledgments

Monica Karday would like to acknowledge the MHRD, Govt. of India for the assistantship at NITJ.

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