Low-Lying Spectra and Electromagnetic Transition Rates in ^{140—162}Gd in the Interacting Boson Model^{*}

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Abstract Spectra and electromagnetic transition for the even-even $^{140-162}$ Gd isotopes are studied in the framework of the interacting boson model. A schematic Hamiltonian can be used to describe their spectra and transition. The results show that $^{140-162}$ Gd are in the transition from the vibrational limit to rotational limit.

Key words spectra, electromagnetic transition, positive parity collective state

1 Introduction

The interacting boson model $(IBM)^{[1-3]}$ is a very effective phenomenological model for describing lowlying collective properties of nuclei. Different from the geometrical picture, it has an inherent group structure. In the first version, nucleon pairs coupled out of an entire major shell are s boson and d boson whose angular momentums are L = 0 and L = 2separately and no distinction is made between neutron boson and proton boson in this model. Scholars show strong interest in nucleons in the rare-earth region. Wen Wanxin and Gu Jingnan calculated $B(M1, 0_1^+ \rightarrow 1_M^+)$ with the theory of the neutronproton interacting boson model. The mixed symmetry states in nuclei of ¹⁵⁰⁻¹⁵⁴Gd were studied. They found that the deformation of isotope is a transition from U(5) limit to SU(3) limit or from vibration to rotation^[4]. By using the generator coordinate method approach to the dynamic group representation (DGR-DCM), Liang Shidong and Fu Deji discussed the spherical nuclei and deformation nuclei at low excitation state^[5]. The potential energy graphs of isotopes show that the potential surface of Gd isotope is a transition from spherical to axial symmetry deformation. The low-lying spectra and E2 transition rates in even-even ^{140—162}Gd are studied in the framework of the interacting boson model IBM-1. A schematic Hamiltonian is used to calculate the low-lying spectra and E2 transition rates in even-even ^{140—162}Gd, and the results show that ^{140—162}Gd are in the transition from the vibration limit to rotational limit.

2 Schematic IBM Hamiltonian

The general IBM Hamilton contains seven terms. However, for our study, we take the following schematic Hamiltonian^[6],

$$\stackrel{\wedge}{H} = \varepsilon_{\rm d} \stackrel{\wedge}{n}_{\rm d} + K \stackrel{\wedge}{Q} \cdot \stackrel{\wedge}{Q} + K_{\rm L} \stackrel{\wedge}{L} \cdot \stackrel{\wedge}{L}$$

where

$$\begin{split} & \stackrel{\wedge}{Q}_{\mu} = (\hat{s}^{+}\hat{\vec{d}} + \hat{d}^{+}\hat{s})^{2} + \chi(\hat{d}^{+}\hat{\vec{d}})^{2}_{\mu} \ , \\ & \hat{L}_{q} = \sqrt{10}(\hat{d}^{+}\hat{\vec{d}})^{(1)}_{q}, \ \ \chi = -\sqrt{7}/2 \ . \end{split}$$

The meanings of the above terms are the same as those in other papers about IBM. The Hamiltonian has three terms. The first one is the single particle

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energy, the second one is the interaction of quadruple moments and the third one is the interaction of single polarity. $\varepsilon_{\rm d}$, K, and $K_{\rm L}$ denote the intensity of interaction. Because the term $K_{\rm L}(\hat{L}\cdot\hat{L})$ is diagonal, it contributes the same to the energy levels with identical spin, and is a term adjusting energy level L. If the boson numbers are given, the Hamiltonian is principally determined by the two parameters ε_{d} and K. However, this Hamiltonian is able to give a transition from U(5) to SU(3), if $\varepsilon_d = 0$, then the Hamilton reduces to a SU(3) limit Hamiltonian. If K = 0, the Hamiltonian becomes a U(5) limit, describing the vibrational collective motion. $K_{\rm L}(\hat{L}\cdot\hat{L})$ term removes some of the degeneracy for different L values. Therefore the ratio of $k/\varepsilon_{\rm d}$ is a measure of the transition between U(5) and SU(3). The Hamiltonian is vibrational for $k/\varepsilon_{\rm d} = 0$ and the Hamiltonian is rotational for $k/\varepsilon_{\rm d} = \infty$. The Hamiltonian is in the transition between U(5) and SU(3) for the other cases. The parameters in the Hamiltonian can be determined by fitting to the experimental spectra.

3 Results and discussion

In Table 1, we give the parameters the Hamiltonian and the E2 transition operator in each nucleus studied. From Table 1, the value of ε_d increases with the increase of neutron number in the lighter even Gd isotopes. The value of ε_d decreases with the increase of neutron number in heavier even Gd isotopes. ε_d reaches its maximum and Kchanges from positive to negative near ¹⁴⁶Gd. These results show the property of energy change at excitation state to Gd isotopes

Table 1. Parameters of energy level and B (E2) operator for Gd isotopes.

1		1		
nucleus	$\varepsilon_{\rm d}/{\rm MeV}$	$K/{ m MeV}$	$K_{\rm L}/{ m MeV}$	$e_2/(eb)$
$^{140}\mathrm{Gd}$	0.482	-0.0055	0.0045	
$^{142}\mathrm{Gd}$	0.612	-0.0005	0.0005	
$^{144}\mathrm{Gd}$	0.698	0.0055	0.0055	
$^{148}\mathrm{Gd}$	0.685	0.0045	0.0000	
$^{150}\mathrm{Gd}$	0.540	0.0050	0.0025	
^{152}Gd	0.449	-0.0045	0.0025	0.1689
$^{154}\mathrm{Gd}$	0.049	-0.0115	0.0138	0.1185
$^{156}\mathrm{Gd}$	0.048	-0.0150	0.0075	0.1201
$^{158}\mathrm{Gd}$	0.048	-0.0150	0.0075	0.1155
$^{160}\mathrm{Gd}$	0.044	-0.0125	0.0075	0.1094
$^{162}\mathrm{Gd}$	0.040	-0.0095	0.0075	

and its shape. They also show the change tendency of Gd isotope from oblate to prolate. ¹⁵⁴Gd is a special nuclei because of ten times the difference between $\varepsilon_{\rm d}$, K, $K_{\rm L}$ of ¹⁵⁴Gd and those of ¹⁵²Gd. ¹⁵⁴Gd with neutron number N = 90 may be X(5) dynamical symmetry. We cannot calculate energy spectra and B(E2) of ¹⁴⁶Gd. It is probable that ¹⁴⁶Gd is a double-magic nucleus with strong shell effect. There is a little change of e_2 . By using these parameters we calculate energy level and B(E2).

3.1 Energy spectra

The comparisons between the calculated and experimental values^[7] of energy levels and B(E2) for ^{140—162}Gd are shown in Figs. 1—5 and Table 3. The theoretical energy levels accord with those of experiments, especially for the ground-state band levels in spite of high spin. γ and the first β band are the same. Because we do not consider multi-band coupling,

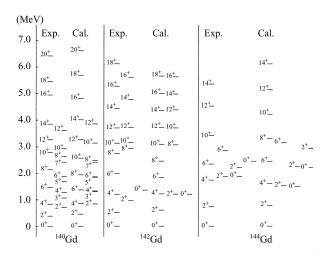


Fig. 1. Spectra for ¹⁴⁰Gd, ¹⁴²Gd and ¹⁴⁴Gd.

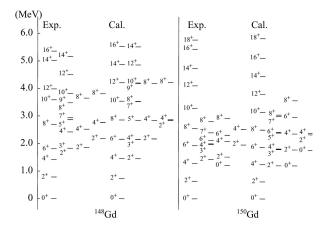


Fig. 2. Spectra for ¹⁴⁸Gd and ¹⁵⁰Gd.

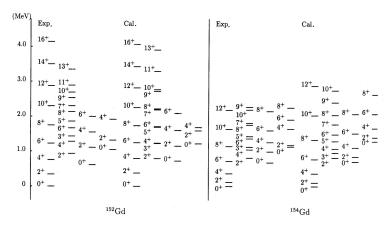


Fig. 3. Spectra for ¹⁵²Gd and ¹⁵⁴Gd.

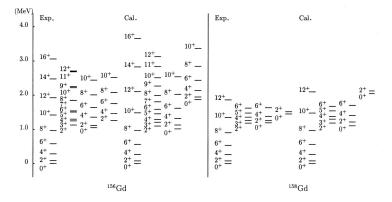


Fig. 4. Spectra for ¹⁵⁶Gd and ¹⁵⁸Gd.

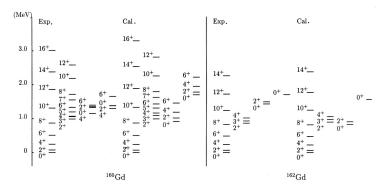


Fig. 5. Spectra for ¹⁶⁰Gd and ¹⁶²Gd.

there are differences between the theoretical and experimental values. The agreement is good for ^{140—144,148,150}Gd, but there is a backbend at the ground-state band for each of them. This result can be explained with collective backbend theory put forward by Long Guilu^[8]. Gd isotopes exhibit staggering in the excited states of ^{140—150}Gd which exist in the γ band generally. The experimental values show obvious staggering, but they are more homogeneous than those of calculation. 3^+_{γ} and 4^+_{γ} , 5^+_{γ} and 6^+_{γ} , 7^+_{γ} and 8^+_{γ} exhibit strong degeneracy. The spectra are the vibration limit with larger ε_d and smaller K, so there is the staggering. The three-body interaction^[9] between the d-boson is introduced to treat the staggering phenomenon in IBM-1. The staggering phenomenon was improved because of adding cubic term to Hamiltonian by Sevrim A and adding quadruple interaction terms of bosons to Hamilton by Liu Yuxin and Long Guilu in IBM2^[10]. For ^{144—150}Gd the two-phonon states are slightly split in energy, about 100keV, which may be understood by means of a small anharmonic term in the collective vibration. However the two-phonon states of 142,148 Gd and the three-phonon states of 150 Gd have a much larger energy splitting. It is difficult to envisage that such a large energy splitting can be caused by anharmonicties alone, which may be related to the excitation of single particle. The theoretical energy of 8_1^+ state of 144 Gd is 3.327MeV and that of 0_4^+ of 150 Gd is 1.766MeV.

For ¹⁵²Gd, the results obtained in the present work are in good agreement with those of experiments. ¹⁵²Gd exhibits strong property of vibration, which can be understood from energy spectra. We can see the differences between the theoretical and experimental data by comparing the experimental and theoretical energy spectra because there exists strong degeneracy between 3^+_{γ} and 4^+_{γ} , 5^+_{γ} and 6^+_{γ} , 7^+_{γ} and 8^+_{γ} , $9^+_{\gamma 1}$ and 10^+_{γ} in the calculated energy spectra. It can be improved by correcting Hamiltonian by adding cubic^[11] or quadruple interaction boson^[12]. The twophonon states and the three-phonon sates of ¹⁵²Gd are slightly split in energy.

For $^{154-156}$ Gd, the calculated values are in good agreement with the experimental ones, especially the ground-band, the gamma-band and the first betaband. Because multi-band coupling is not considered, it is understood that there exist differences between the calculated and the experimental values. Most calculated values of beta-band are larger. The special states, 0⁺ and 2⁺, may be mixed state or intruder state.

For the ground-band, gamma-band and first betaband of ^{158—160}Gd, the calculated values are in good agreement with the experimental ones. The calculated values of the second beta-band are larger than that of the ground-band of ¹⁶²Gd. For the groundband, gamma-band and first beta-band of ¹⁶²Gd the calculated values are in good agreement with the experimental ones. The calculated values of the second beta-band are smaller. There is no obvious staggering for the excited states of ^{158—162}Gd.

For explaining the changes of ${}^{140-162}$ Gd, the ratios $R = E(4^+_1)/E(2^+_1)$ are shown in Table 2. From Table 2, we can also see a transition from U(5) to SU(3). ^{140—152}Gd isotopes exhibit obvious properties of vibration. The properties of vibration are becoming strong with the increase of nuclei number and ^{158—162}Gd are nearly perfect rotors.

Table 2. The ratios of $E(4_1^+)/E(2_1^+)$.

A	R_{Exp}	R_{Cal}		
140	2.19	2.13		
142	2.35	2.01		
144	2.35	2.03		
148	1.81	1.98		
150	2.02	2.01		
152	2.19	2.08		
154	3.02	3.32		
156	3.24	3.33		
158	3.30	3.32		
160	3.31	3.32		
162	3.34	3.32		

3.2 E2 transition

After the determination of the spectra, the wave function is determined. The electric and magnetic transition properties can then be obtained accordingly. For example, the E2 transition operator is

$$\hat{T}(\mathbf{E}2)^2_{\mu} = e_2 \big[(\hat{s}^+ \tilde{\hat{d}} + \hat{d}^+ \hat{s})^2_{\mu} + \chi \big(\hat{d}^+ \tilde{\hat{d}} \big)^2_{\mu} \big].$$

The meanings of the above terms are the same as those in other papers about IBM. B(E2) of each nuclei are calculated by using e_2 shown in Table 1. The results are shown in Table 3.

From Table 3, we can see that the results calculated are in good agreement with those of experiment.

 Δn denotes phonon number of transition between different levels. Most B(E2) values shown in Table 3 are larger for $\Delta n = 1$. That means these nuclei are U(5) limit. However there exist some nuclei belonging to SU(3) limit because of smaller B(E2) values for $\Delta n = 1$. It is determined by the selection rule of $\Delta \lambda = 0$ and $\Delta \mu = 0$. Because the beta-band and gamma-band belong to the same SU(3) expression $(\lambda = 2N - 4, \ \mu = 0)$ and g-band the other SU(3)expression $(\lambda = 2N, \ \mu = 0)$, the transition between gamma-band and g-band main caused by breaking terms of SU(3) is weak. The fact that the transition probability of $B(E2, 2^+_2 \rightarrow 0^+_1)$ is not zero hints that there exist symmetry breaking terms.

Table 3. Comparison of B (E2) values in Gd nuclei^[7].

nucleus	$J_{\rm i}$	$J_{\rm f}$	$\operatorname{Exp.}/(e^2 \cdot \operatorname{fm}^4)$	$\operatorname{Cal.}/(e^2 \cdot \mathrm{fm}^4)$
¹⁵² Gd	2^+_1	0_1^+	3517	3516
Gu		0^{+}_{1}	15	20
	$\frac{-2}{2^+}$	$0^+_1 \\ 0^+_2 \\ 4^+_1$	2024	1206
	2^{+}_{2}	4^+_1	1735	830
	0^{+}_{2}	2^{+}_{1}	8672	6751
	4^+_1	2^{+}_{1}	6456	6544
	6^+_1	$\frac{-1}{4^+}$	9636	8858
	2^{+}_{2}	$\begin{array}{c} 2^+_1 \\ 2^+_1 \\ 4^+_1 \\ 2^+_1 \\ 0^+_1 \\ 0^+_2 \\ 0^+_1 \\ 0^+_1 \\ 4^+_1 \\ 4^+_1 \\ 2^+_1 \end{array}$	814	5019
$^{154}\mathrm{Gd}$	2^{+}_{1}	0^{+}_{1}	7698	7698
	2^{1}_{2}	0^{+}_{2}	4752	5853
	2^{2}_{2}	$0^{\frac{2}{+}}_{2}$	63	67
	2^{+}_{2}	0_{1}^{2}	42	1
	2^{2}_{3}	0_{1}^{+}	279	1
	2^{+}_{2}	4_{1}^{+}	961	6
	2^{2}_{3}	4_{1}^{+}	84	0
	2^{+}_{3}	2_{1}^{+}	642	2
	2^{+}_{2}	2_{1}^{+}	328	2
	0^{2}_{2}	2_{1}^{+}	2550	9
	$2^{+2}_{2} 2^{+2}_{2} 2^{+2}_{2} 0^{+1}_{2} 4^{+1}_{1} + 2^{+2}_{2} 2^{+3}_{3} 2^{+2}_{2} 2^{+3}_{3} 2^{+2}_{2} 2^{+3}_{3} 2^{+2}_{2} 2^{+3}_{3} 2^{+2}_{2} 2^{+3}_{3} 2^{+2}_{2} 2^{+3}_{3} 2^{+2}_{2} 2^{+3}_{3} 2^{+2}_$	$2^+_1 \\ 2^+_1 \\ 2^+_1 \\ 4^+_1 \\ 6^+_1$	12012	10798
	6^{+}_{1}	4^{+}_{1}	13974	11498
	8^+_1	6^{+}_{1}	15297	11438
	10^+_1	8^{+}_{1}	17651	10945
$^{156}\mathrm{Gd}$	2^+_1	0^{+}_{1}	9328	9328
Gu	2^{+}_{2}	0^{+}_{1}	31	1
	$\frac{2}{2^{+}}$	0^1_1 0^+_1	234	1
	2^+_3 2^+	0^1_1 0^+_1	15	0
	2^+_4 2^+_2 2^+_4	$0^{1}_{2^{+}}$	170	1
	$\frac{2}{2}$	2^+_1	30	0
	$\frac{2}{24}$	2^+_1		
	2^+_3	2^+_1	364	$\frac{1}{3}$
	2^+_2	$\begin{array}{c} 4_{1}^{+} \\ 4_{1}^{+} \\ 4_{1}^{+} \\ 2_{1}^{+} \end{array}$	205	
	$2^+_4 \\ 2^+_3 \\ 4^+_1 \\ 6^+_1 \\ 8^+_1$	4_{1}^{+}	214	39
	$\frac{2}{3}$	4_{1}^{+}	39	0
	4_{1}^{+}	2_{1}^{+}	13118	13121
	6_{1}^{+}	$4^+_1 \\ 6^+_1$	14715	14044
		6_{1}	15962	14086
	10^{+}_{1}	8_{1}^{+}	15662	13641
150	12^{+}_{1}	10^{+}_{1}	14964	12843
$^{158}\mathrm{Gd}$	2_{1}^{+}	0_{1}^{+}	10047	10046
	2^{+}_{2}	0_{1}^{+}	178	1
	2^{+}_{3}	0_{1}^{+}	16	0
	2_{4}^{+}	0_{1}^{+}	19	0
	2^{+}_{2}	2_{1}^{+}	299	1
	2_{4}^{+}	2_{1}^{+}	21	0
	2_{4}^{+}	4_{1}^{+}	19	0
	2_{4}^{+}	4_{2}^{+}	1218	5
	2^{+}_{2}	4_{1}^{+}	14	2
	$\begin{array}{c} 2^+_2\\ 2^+_3\\ 2^+_4\\ 2^+_2\\ 2^+_4\\ 2^+_4\\ 2^+_4\\ 2^+_2\\ 2^+_3\\ 4^+_1\\ 8^+_1\\ 10^+_1 \end{array}$	$\begin{array}{c} 0^+_1 \\ 0^+_1 \\ 2^+_1 \\ 2^+_1 \\ 4^+_1 \\ 4^+_2 \\ 4^+_1 \\ 4^+_1 \\ 2^+_1 \\ 6^+_1 \\ 8^+_1 \end{array}$	71	0
	4_{1}^{+}	2_{1}^{+}	15664	14161
	8_{1}^{+}	6_{1}^{+}	16237	15363
	10^{+}_{1}	8_{1}^{+}	16744	15014
	12^{+}_{1}	10^{+}_{1}	15730	14317
$^{160}\mathrm{Gd}$	2_{1}^{+}	0_{1}^{+}	10380	10379
	2^{+}_{2}	$\begin{array}{c} 0_1^+ \\ 0_1^+ \\ \end{array}$	196	1
	$2^+_2 \\ 2^+_2$	2^{+}_{1}	366	1
	2^{2}_{2}	$2^+_1 \\ 4^+_1$	37	3
	2	-1	- •	

For decay from 2_3^+ to 4_1^+ , the transition between states of $\Delta n = 1$ can be described by vibration and the transition from beta-band to g-band the rotation. The B(E2) value of ¹⁵⁴Gd, ¹⁵⁶Gd, ¹⁵⁸Gd from 2_3^+ to 4_1^+ is nearly zero. That means these nuclei exhibit strong rotation property.

For further explaining the shape changes of $^{152-158}$ Gd, we analyze g-band, B (E2: $L + 2 \rightarrow L$), values. Results are shown in Fig. 6. From Fig. 6, we can see that the g-band value increases with the increase of angular momentum and it begins to decrease at 1/4 maximum angular momentum. It is similar to SU(3) property^[13]. That means $^{156-158}$ Gd exhibit transition from U(5) to SU(3).

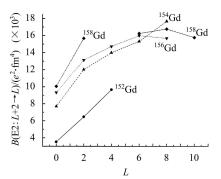


Fig. 6. The values of B (E2: $L + 2 \rightarrow L$) of the g-band as function of spin L in ¹⁵⁴⁻¹⁵⁸Gd isotopes.

In the U(5) limit, the $\Delta n_{\rm d} = 1$ selection rule allows a decay from the 0_2^+ state to the 2_1^+ state that can be observed in ¹⁵²Gd, ¹⁵⁴Gd and the decay is strong. In the O (6) limit, the $\Delta \sigma = 0$ and $\Delta \tau = 1$ selection rule allows a decay from the 0_2^+ state to the 2_2^+ state and the decay from the 0_2^+ state to the 2_1^+ state is forbidden. For all Gd isotopes, the decay from 0_2^+ to 2_2^+ state is not observed. That means ^{140—162}Gd are the U(5) to SU(3) transitional nuclei.

4 Conclusion

We have made a detailed study of the energy levels and E2 transition in Gd isotopes in the framework of the interacting boson model. The results indicate that $^{140-162}$ Gd are the U(5) to SU(3) transitional nuclei. Meanwhile, the discrepancy between the calculated and the experimental data is found, which means that other factors must be introduced into Hamiltonian, such as pair interacting, isospin effect,

high angular momentum boson, IBM-2 and so on.

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^{140—162}Gd核的低能谱和电磁跃迁的相互作用玻色子模型*

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摘要 采用相互作用玻色子模型研究了^{140—162}Gd偶偶核的低能谱和电磁跃迁,应用一个 $U(5) \rightarrow SU(3)$ 的简化 哈密顿量很好地描述它们的低能谱和电磁跃迁过渡.结果表明^{140—162}Gd同位素核基本上属于 $U(5) \rightarrow SU(3)$ 的 过渡核.

关键词 能谱 电磁跃迁 低能正宇称集体态

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