Reduced transition probabilities of vibrational states in ^{154–160}Gd and ^{176–180}Hf

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The vibrational states of 154,156,158,160 Gd and 176,178,180 Hf have been studied via Coulomb excitation. Thin high-purity targets were prepared in an isotope separator. Coulomb excitation of these nuclei was studied by the scattering of 11–17 MeV α particles. The particles were detected in an Enge split-pole spectrograph. The reduced transition probabilities $B(E\lambda;0_{g,s}^{+}) \rightarrow I^{\pi}K$, were obtained for $I^{\pi}K = 2^+ 2$ states in each nucleus, and for $I^{\pi}K = 2^+ 0$ and $I^{\pi} = 3^-$ states in several nuclei. The $K^{\pi} = 2^+$ levels are reasonably constant in energy and B(E2) strength while the $K^{\pi} = 0^+$ states change rapidly in energy and excitation strength with neutron number. In both the softly deformed Gd nuclei and the more strongly deformed Hf nuclei, the lightest nucleus has the lowest energy and largest B(E2) for an $I^{\pi}K = 2^+ 0$ state, despite the difference in the sizes of the deformations at these lightest masses.

 $\begin{bmatrix} \text{NUCLEAR REACTIONS} & ^{154-160} \text{Gd}(\alpha, \alpha'), & ^{176-180} \text{Hf}(\alpha, \alpha'), & E=11-17 \text{ MeV; mea-} \\ & \text{sured } \sigma, \text{ deduced } B(E\lambda) \text{ and } M(E\lambda). \text{ Enriched targets.} \end{bmatrix}$

I. INTRODUCTION

It has been pointed out in a review¹ of the relative transition probabilities from vibrational states in gadolinium, hafnium, and neighboring nuclei that a microscopic approach is needed to explain the decay properties of these levels except in the very middle of the deformed region ($150 \leq A \leq 190$). In any microscopic calculation, the $B(E\lambda)$ strengths from the vibrational levels provide a significant test of the particular approach. The stable eveneven Gd and Hf isotopes are excellent candidates for testing microscopic calculations of deformed nuclei because (a) β - and γ -type vibrational bands occur in many of these isotopes; (b) other, $K^{\pi} = 0^{+}$, bands are reported in several of these; (c) the region of deformation they span is large since with increasing neutron number N, the Gd nuclei go from ¹⁵²Gd at the onset of deformation with N=88to the well-deformed ¹⁶⁰Gd, and the Hf nuclei go from the well-deformed ¹⁷⁴Hf to the more softly deformed ¹⁸⁰Hf. These same isotopes are also of interest because recent theoretical calculations²⁻⁴ of ground-state deformations predict the Gd nuclei to have large positive hexadecapole deformation parameters (β_4) and the Hf nuclei to have small negative values. A later paper will present E2 and

*E*4 moments from which one may extract these charge deformation parameters for most of the nuclei studied here.

In this paper we present the results of a series of measurements of the absolute $B(E\lambda)$ strengths for $I^{r}K=2^{+}2$ and $2^{+}0$, and $I^{r}=3^{-}$ vibrational states in ¹⁵⁴⁻¹⁶⁰Gd and ¹⁷⁶⁻¹⁸⁰Hf as deduced from Coulomb excitation via the (α, α') reaction. Systematic trends for the energies and collective strengths of these states are presented. The $I^{r}K=2^{+}2$ and $2^{+}0$ states in Gd and Hf show remarkable similarities with increasing neutron number despite the fact that the deformation is increasing in Gd and decreasing in Hf with N. Comparison with other reported measurements and with theoretical predictions of $B(E\lambda)$ strengths and level energies are made.

II. EXPERIMENTAL PROCEDURE

 α particles were obtained in the EN tandem Van deGraaff accelerator at the Oak Ridge National Laboratory. A 20-cm-long position-sensitive gasflow proportional counter, mounted in the focal plane of an Enge split-pole magnetic spectrograph, was used to detect the elastically and inelastically scattered α particles. Most of the targets were

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studied by observing the scattered particles at two laboratory angles, 150° and 90° . The efficiency and linearity calibrations of the detector-spectrograph system were made by measuring the yields and positions of the two main α groups in the decay of ²⁴⁴Cm as a function of magnetic field strength. This detector was found to be uniformly efficient ($\leq 1\%$ average variation) across most of its length.

The targets were thin (~20-25 μ g/cm² in most cases) very pure (>99%) isotopes of Gd and Hf material deposited on thin (75–100 μ g/cm²) carbon or nickel foils at the ORNL 180° sector isotope separator. The ratios of the elastic peak heights to average backgrounds above the $I^{\pi} = 4^{+}$ groundband peaks were typically better than 10⁴. The ratios of the elastic peak heights to the valley depths between the elastic and 2⁺ ground-band peaks were typically better than 500. Peak widths at half-maximum were usually 18 to 25 keV. Beam energies were chosen by extrapolating the results of recent Coulomb-nuclear interference measurements on the excitations of the first 2⁺ and 4⁺ states.^{5,6} We used higher beam energies when detecting particles scattered into 90° so that about the same distances of closest approach as for the 150° experiments would be achieved. However, few studies have been made concerning Coulombnuclear excitation interferences for vibrational states. Early studies⁷ of Sm nuclei indicated that the nuclear excitation onset is the same for vibrational 2⁺ and 3⁻ states as for the ground-band 2⁺ state. A very recent study⁸ of 0s nuclei seems to yield the same conclusion for $I^{T}K = 2^{+}2$ states.

III. DATA ANALYSIS

Figures 1 and 2 show typical α -particle spectra from our measurements. Experimental ratios of inelastic-to-elastic differential scattering cross sections, $d\sigma_{2^+ \text{ or } 3^-}/d\sigma_{\text{el}} \equiv R_{\text{exp}}$, were found by normalizing the areas of the peaks due to inelastic scattering to the elastic peak, after fully resolving the latter from the $2_{g,s}^{+}$ peak. Then the R_{exp} values were compared with values (R_{calc}) calculated with the semiclassical Coulomb-excitation code of Winther and deBoer,⁹ modified¹⁰ to include E1, E3, and E4 excitations, with the matrix elements $\langle 0^{+}_{g.s.} || M(E\lambda) || I^{\pi} \rangle$ as free parameters. Baktash et al.¹¹ reported differences of ~3% (at θ_{1ab} ~170°) between semiclassical and quantum-mechanical calculations of R_{calc} for $I^{T}K = 2^{+}2$ states. This difference is in the treatment of higher-order excitation processes, such as $0 \rightarrow 2 \rightarrow 2'$. Since these are smaller at 90°, where our best data were obtained, we chose to use solely the semiclassical analysis. Reduced matrix elements connecting the $0_{g.s.}^{*}$ through $6_{g.s.}^{*}$ states with the 0^{*} and 2^{*} mem-



FIG. 1. Spectrum of elastically and inelastically scattered 15-MeV α particles from ¹⁶⁰Gd at a lab angle of 90°.

bers of excited $K^{\pi} = 0^{+}$ bands and 2^{+} and 4^{+} members of the $K^{\pi} = 2^+$ bands were used in the calculations. Interband and intraband matrix element relations¹² were deduced from the adiabatic rotor-vibrator (collective) model predictions¹³ although γ -ray branching ratio data, corrected for M1 admixtures,^{1,14} were used for interband relations when they existed. Branching ratios of γ rays from vibrational to rotational states in most nuclei in the rare-earth region deviate significantly¹ from this model because of the rotation-vibration mixing. In 156 Gd, for $\theta = 90^{\circ}$ and $E_{\alpha} = 15.3$ MeV, use of the collective model relations rather than the measured relations would decrease R_{calc} for the $I^{T}K = 2^{+}2$ state by 0.3% and for $I^{T}K = 2^{+}0$ states by about 6%. This reflects the stronger mixing with the ground band by the $K^{\pi} = 0^{+}$ band than by the $K^{\pi} = 2^{+}$ band. Baktash *et al.*¹¹ examined the



FIG. 2. Spectrum of inelastically scattered 16.9-MeV α particles from ¹⁷⁶Hf at a lab angle of 90°. Excitations of the $I^{\pi}K = 2^{+}2$, $2^{+}0$, and $3^{*}2$ states at 1341, 1227, and 1313 keV are observed.

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effect of the static moment of the $I^{\pi}K = 2^{+}2$ state on the analysis at a backward angle of ~170°. A variation of the static moment for ¹⁶⁰Gd from zero to a rotational model value increases the calculated cross section ($E_{\alpha} = 12$ MeV) of this state by ~11% at 150°, whereas for the same separation distance at 90°, the effect is ~6%. In our calculations rotational model values are assumed for the static moments of the 2⁺ states. Some recent experimental data¹⁵ indicate the validity of this assumption at least for $I^{\pi}K = 2^{+}2$ states. In general, much more experimental effort is needed to determine signs and magnitudes of static moments for $I^{\pi} = 2^{+}$ and 3⁻ states of excited bands.

Strong mixing of the ground and excited K=0 bands can affect the interband transition matrix

elements to the point that their signs may differ from the collective model predictions. Not only are the magnitudes of these matrix elements somewhat uncertain but also their signs. Kumar^{16,17} pointed out that the signs should be taken either from experiment or a microscopic theory. Lacking experimental or theoretically reliable signs, we have used those deduced from the collective model. For an estimate of the importance of this, we considered the $4_{g.s.} \rightarrow 2\beta$ element for ¹⁵⁶Gd (θ = 150° and E_{α} = 15.3 MeV). It is strongly affected by bandmixing (see Sec. IV C). A change in the sign of the matrix elements connecting the $4_{g.s.}^{+}$ state with the $I^{\pi}K = 2^{+}0$ vibrational states was made. These sign changes have a negligible (~1%) effect on R_{calc} for these $I = 2^*$ states.

TABLE I. Experimental absolute $B(E\lambda)$ strengths for ¹⁵⁴⁻¹⁶⁰Gd and ¹⁷⁶⁻¹⁸⁰Hf from (α, α') Coulomb excitation studies.

Nucleus	E (keV)	I [#] K	$B(E\lambda; 0^*g.s. \rightarrow I^{T}K)^{a}$ $(e^{2}\mathrm{b}^{\lambda})$	$\frac{B(E\lambda)}{B(E\lambda)_{\mathbf{s}_*\mathbf{p}_*}}^{\mathbf{b}}$	C measu Value	ther irements Reference
¹⁵⁴ Gd	123 815 996	2*g.s. 2*0 2*2	3.85(8) 0.015(4) 0.143(11)	157(3) 0.6(2) 5.9(5)		
¹⁵⁶ Gd	89 1129 1154 1258 1276	2*g.s. 2*0 2*2 2*0 3	4.57(5) 0.013(4) 0.120(4) <0.008 0.16(4)	183(2) 0.5(2) 4.8(2) < 0.3 16(4)		
¹⁵⁸ Gd	80 1187 1260 1517	2*g.s. 2*2 2*0 2*0	4.97(5) 0.090(10) <0.002 <0.002	196(2) 3.6(4) < 0.08 < 0.08	0.106(15)	11
¹⁶⁰ Gd	75 992 1070	2*g.s. 2*2 (2*0 or 3*?)	5.15(6) 0.101(3) 0.002(2) $e^{2}b^{2}$ or 0.01(1) $e^{2}b^{3}$ 0.127(14)	$200(2) \\ 3.9(1) \\ 0.08(8) \\ or \\ (1 \pm 1) \times 10^{-4} \\ 4.1 + 0(12)$	0.104(4)	11
¹⁷⁶ Hf	1289 88 1227 1313 1341	(3 ?) 2*g.s. 2*0 3 ⁻ ? 2*2	0.127(14) 5.19(6) 0.031(3) 0.093(29) 0.119(8)	11.9(13) 177(2) 1.0(1) 7.2(23) 4.1(3)	0.025(5) 0.075(6)	18 18
¹⁷⁸ Hf	93 1175 1277 1323 1496	2*g.s. 2*2 2*0 3-2 2*0	4.86(5) 0.115(4) 0.0018(7) 0.053(10) 0.013(2)	$163(2) \\ 3.9(1) \\ 0.07(3) \\ 4.0(8) \\ 0.44(7)$	0.100(8) ≤0.002 ≤0.010	19 19 19
¹⁸⁰ Hf	93 1201	2*g.s. 2*2	4.73(5) 0.114(7)	157(2) 3.8(3)	0.110(11)	20

^a The uncertainties (bracketed) are one-standard-deviation values and represent variations in the last digits of the best values. For example, 0.143(11) may be written as 0.143±0.011. ^b $B(E\lambda)_{s,p} = (2\lambda+1)/4\pi [3/(\lambda+3)]^2 (0.12A^{1/3})^{2\lambda} e^{2b\lambda}$ for $I_i = 0$ and $I_f = \lambda$.

IV. RESULTS AND DISCUSSION

A. Experimental results

Table I gives energies and absolute B(E2) and B(E3) values for $I^{\pi} = 2^+$, 3⁻ states in ¹⁵⁴⁻¹⁶⁰Gd and ¹⁷⁶⁻¹⁸⁰Hf. The E2 and E4 reduced matrix elements for these nuclei are being prepared for publication, and will be compared with other results then. The B(E2) values for the $2_{g.s.}^{*}$ states do verify that the heavier Gd nuclei are more deformed than the lighter ones. The falling energies of the $2_{g.s.}^{*}$ states and the rising B(E2) values are signatures of increasing deformation. In contrast, the heavier Hf nuclei are less deformed than the lighter ones. Previous measurements¹⁸⁻²⁰ of some of the vibrational B(E2) values are also given in Table I. Uncertainties related to effects not discussed in Sec. III (e.g., those from unincorporated quantum-mechanical corrections) are not included in our guoted error limits.

1. ¹⁵⁴Gd

Because of relatively poor target quality, only the low-lying $I^{\pi}K = 2^{*0}$ and 2^{*2} states at 816 and 996 keV, respectively, could be identified; the $I^{\pi} = 3^{-}$ state at 1353 keV²¹ was not observed. Although the B(E2) strength of the 2*0 state is the largest compared with similar states in the Gd nuclei, it may be less than a single particle unit (s.p.u.). There is, however, strong mixing²² between this excited $K^{\pi} = 0^{*}$ band and the ground band.

2. 156Gd

This nucleus has three $I^{\pi} = 2^+$ states²³ within 130 keV of each other. However, only two, the K=2 state at 1154 keV and K=0 state at 1129 keV, were found to have significant B(E2) strength.

3. 158Gd

Only the $I^{\pi} = 2^{+}2$ state at 1187 keV is seen with measurable collective strength; upper limits are placed on the absolute B(E2) strengths of the two $I^{\pi}K = 2^{+}0$ states observed at 1260 and 1517 keV in decay studies,²⁴

4. 160 Gd

We observed strongly excited states at 992 and 1289 keV, and a weakly excited state at 1070 keV. All three of these states correspond to levels seen at or near these energies in other studies.^{11,21,25-27} A state at 1070 keV has previously been assigned $I^{t}K = 2^{+}0$ (Ref. 21), 3⁻¹ (Ref. 21), 3⁺(?) (Ref. 25), and 4⁺(?) (Ref. 26). However, the large direct excitation probabilities with α particles restricts the possible spin parity assignments to 2⁺ or 3⁻. We cannot choose between these two. For the same reason, we favor the $I^{\pi} = 3^{-}$ assignment²¹ rather than $I^{\pi} = 2^{-}$ (Ref. 26) for the state at 1289 keV.

5. 176Hf

Well established $I^rK = 2^*2$, 2^*0 , and 3^-2 states at 1341, 1227, and 1313 keV, respectively, are definitely excited, but there is no evidence for collectivity in the proposed²⁸ second $I^rK = 2^*0$ state at 1379 keV. Our B(E2) value for the 1227-keV state is in agreement with the recent (α, α') measurement of Hammer, Ejiri, and Hagemann¹⁸ but for the 1341-keV state their B(E2) value is two-thirds of ours. Our result is the average of data taken at 90° and 150°, which were found to be in good agreement.

6. ¹⁷⁸Hf

Two $I''K = 2^{*0}$ states²⁹ have been observed here, one at 1277 keV and the other at 1496 keV. The latter has a B(E2) value more typical of a vibrational state. Earlier $(\alpha, \alpha'\gamma)$ Coulomb excitation results¹⁹ for these states and the $I'' = 2^{*}$ state at 1175 keV are in good agreement with our more accurate B(E2) values.

7. ¹⁸⁰Hf

Our study has shown that only the $I^{*}K = 2^{+}2$ state at 1201 keV is collective, thus supporting the work of Varnell, Hamilton, and Robinson.²⁰

B. Collective strength and energy level trends

The trends of the B(E2) strengths (in single particle units) and of the energies of $I^{\pi} = 2^+$ vibrational states are shown in Fig. 3 for ¹⁵⁴⁻¹⁶⁰Gd. Figure 4 shows these systematics for ¹⁷⁴⁻¹⁸⁰Hf, with the data for ¹⁷⁴Hf taken from Ref. 30. In both the Gd and Hf nuclei the $I^{T}K = 2^{+}2$ states are nearly constant in energy and decrease slowly in collectivity as Nincreases. On the other hand, the most collective $I^{T}K = 2^{+}0$ states change rapidly in energy. The $2^{+}2$ and $2^{+}0$ states with the largest B(E2) values are in the most neutron deficient isotopes of Gd and Hf. The 2*0 state then lies the lowest in energy. That the same trends are seen in both Gd and Hf nuclei is surprising in that for increasing N, the Gd nuclei are more deformed but the Hf nuclei are less deformed.

In a macroscopic sense, the increase of a vibrational energy and corresponding decrease in its B(E2) strength indicate an increasing stiffness to this vibration since the level energy E and the B(E2) value are related in the collective model to



FIG. 3. Absolute B(E2) strength (in single particle units) and level energy systematics for $^{154-156}$ Gd for 2^{+} states of bands with K values given in the figure.

the vibrational stiffness parameter C according to

$$E \propto \left(\frac{C}{B}\right)^{1/2}$$
 and $B(E2) \propto (BC)^{-1/2}$,

where B is a mass parameter, which includes moment-of-inertia effects. Figures 5 and 6 show plots of ratios of energies and B(E2) values of the vibrational states to the corresponding values for the ground-band 2⁺ state. Such ratios should minimize the ground-state deformation effects (by near cancellation of the mass parameter) and emphasize



FIG. 4. Absolute B(E2) strength (in single particle units) and level energy systematics for ¹⁷⁴⁻¹⁸⁰Hf for 2⁺ states of bands with K values given in the figure. The data for ¹⁷⁴Hf are taken from Ref. 30.



FIG. 5. Ratios of level energies and absolute B(E2) strengths of vibrational states to those of the $2_{g,s}^{+}$ states in ¹⁵⁴⁻¹⁶⁰Gd. The B(E2) ratios for $K^{\pi} = 0^{+}$ states in ^{158,160}Gd and the 1258-keV state in ¹⁵⁶Gd are extremely small and are omitted.

the stiffness as a function of neutron number. Our data presented in this manner show the similarity of the Gd and Hf nuclei with the Gd nuclei being slightly more sensitive to the neutron number than the Hf nuclei.



FIG. 6. Ratios of level energies and absolute B(E2) strengths of vibrational states to those of the $2^+_{g,s,}$ states in ¹⁷⁴⁻¹⁸⁰Hf. The data for ¹⁷⁴Hf are taken from Ref. 30. The B(E2) ratio for the 1277-keV $K^{\pi} = 0^+$ state in ¹⁷⁸Hf is extremely small and is omitted.

C. Interpretations of $I^{\pi}K = 2^+2$ and 2^+0 vibrational states

If collective, the $I^{\pi} = 2^+$ states may be interpreted as γ vibrational when K=2, or β vibrational when K=0. Our results show that the γ vibrational mode is present in each nucleus studied here. The $I^{r}K$ = 2^{+0} states in ^{154,156}Gd and ¹⁷⁴⁻¹⁷⁸Hf with the largest B(E2) values do have strengths typical of β vibrational states. The B(E2) values are only ~1 s.p.u. and one might incorrectly conclude that these 2* states are not members of a collective band. However, the collective $K^{\pi} = 0^{+}$ vibrational band may mix strongly with the $K^{\pi} = 0^{*}$ ground band. As a result of the mixing, the collective E2 strength is redistributed between the two K=0 bands. To first order, the $0_{g,s_{\bullet}}^{+} \rightarrow 2_{\beta}^{+}$ matrix element is decreased, the $2^+_{g,s} - 0^+_{\beta}$ and $4^+_{g,s} - 2^+_{\beta}$ elements are increased. Then, in a first-order perturbation treatment of this mixing, ²² the intrinsic value of $B(E2; 0_{g,s,-}^* - 2_{\beta}^*)$ will be multiplied by the factor $(1 - 6Z_{\beta})^2$. The value of $B(E2; 0^+_{\beta} - 2^+_{g_{\alpha}, g_{\alpha}})$ will be multiplied by $(1 + 6Z_{\beta})^2$. For ¹⁵⁴Gd, where $Z_{\beta} \approx 0.05$ (Ref. 22), the measured $B(E2; 0^+_{g,s,} - 2^+_{\beta})$ is thus 49% of the intrinsic value. The intrinsic $4^+_{g,s} \rightarrow 2^+_{\beta}$ element is multiplied by $(1+14Z_{\beta})$. With $Z_{\beta} \approx 0.05$, the measured value of $B(E2; 4^{*}_{g,s_{\bullet}} \rightarrow 2^{*}_{\beta})$ is 289% of its intrinsic value. It is difficult then to conclude the collectivity of a K = 0 band simply by measuring $B(E2;0_{c_{s_{s}}}^{+}-2_{\beta}^{+})$. The complementary measurement of $B(E2; 0_{\beta}^{+} - \hat{2}_{R,s_{*}}^{+})$ can lead to a better assessment of the collectivity. Mixing of the γ and ground bands produces a smaller effect on $B(E2; 0_{g.s.}^{+} \rightarrow 2_{\gamma}^{+})$. Here, the factor has the form $(1 - Z_{\gamma})^2$. For ¹⁵⁴Gd, $Z_{\gamma} \approx 0.09$ (Ref. 22), and the reduction is only 17%. The weaker $I^{T}K = 2^{+}0$ states are experimentally strikingly different from the stronger $I^{T}K = 2^{+}0$ states. Studies of the (p, t) reaction on Gd (Ref. 31) and Hf (Ref. 32) nuclei show that the 0^+ and 2^+ members of the band usually associated with the β vibration are more strongly excited than the second excited K = 0 band members. Such is true for the (d, d') reaction²¹ also. Directional correlation studies show that the $2^+0 - 2^+_{g,s}$ transitions from the second K=0 band, such as in ¹⁵⁶Gd (Ref. 33) and 178 Hf (Ref. 34), are mostly M1 in character. It is interesting that for ¹⁷⁸Hf the transition from the 1277-keV 2* state to the 2* ground state has an 80% M1 admixture³⁴ and that E2 γ -ray branching ratios from this state do not agree¹ with the collective model even with perturbational corrections. On the other hand, the transition from the more collective 2* state at 1496 keV to the 2* ground state has also a large (62%) M1 admixture³⁴ but there the E2 branching ratios do agree¹ when a perturbational correction is applied. Such M1admixtures were originally thought³⁵ to be responsible for bandmixing anomalies in the β bands but

this was subsequently ruled out (see Ref. 1). Other interpretations besides that of a collective β vibration have been given to such bands. A single twoquasiparticle amplitude may be dominant, possibly as in the cases of the $K^{\pi} = 0^{+}$ bandheads at 1168 keV in ¹⁵⁶Gd (Ref. 36) and at 1199 keV in ¹⁷⁸Hf (Ref. 34). The 1715-keV 0⁺ state in ¹⁵⁶Gd may be a two- β phonon ($n_{\beta} = 2, n_{\gamma} = 0$) bandhead.³⁷ Also, theoretical calculations³⁸ of potential energy surfaces in this region indicate that it is possible to have a K = 0band built on a minimum corresponding to an oblate deformation, whereas the normal β vibrations occur about prolate shapes.

D. Application of a pairing-plus-quadrupole model to ^{154, 156}Gd

For γ vibrations, the pairing-plus-quadrupole (PPQ) interaction calculations of Bes *et al.*³⁹ attempted to provide an in-depth view of this mode in the rare-earth region. However, B(E2)strengths are overestimated by factors of 2 or 3. Much better agreement with experiment has been obtained either by the more exact PPQ calculations of Kumar for ^{150,152}Sm (Ref. 40), or by a variable moment of inertia approach⁴¹ for ¹⁵²Sm. Gupta *et al.*^{42,43} have extended the PPQ calculations to the more deformed ^{154,156}Gd. Tables II and III present our experimental values of the reduced *E2* matrix elements $\langle 0^*_{\underline{K},\underline{s}}, ||M(E2)||I^{\underline{r}} = 2^*, K = 0$ or $2\rangle$, which are square roots of the B(E2) values of Table I. Experimental magnitudes of

$$\langle 2_{g,s}^{*} \text{ or } 4_{g,s}^{*} ||M(E2)||I^{T} = 2^{*}, K = 0 \text{ or } 2 \rangle$$

were obtained by combining our B(E2) values with B(E2) ratio data¹ from γ -ray studies. These are compared with PPQ predictions as well as with predictions of the collective model, using the

TABLE II. E2 matrix elements in ¹⁵⁴Gd compared with pairing-plus-quadrupole and collective models.

	$M(E2; IK \rightarrow I'K')$ (eb)				
$IK \rightarrow I'K'$	<i>E_{I'K'}</i> (keV)	Exp (magnitude)	PPQ ^a	Collective model ^b	
$0g \rightarrow 2g$	123	1.96(2)	1.96		
$0g \rightarrow 2\gamma$	996	0.38(6)	0.37		
$0g \rightarrow 2\beta$	815	0.12(7)	0.14		
$2g \rightarrow 2\gamma$		0.46(9)	-0.54	_0.68(10)	
$2g \rightarrow 2\beta$		0.35(18)	0.41	0.15(8)	
$4g \rightarrow 2\gamma$		0.22(3)	-0.14	-0.10(2)	
$4g \rightarrow 2\beta$		0.6(4)	-0.88	-0.20(10)	

^aPredictions of the pairing-plus-quadrupole model calculations of Gupta *et al.* (Refs. 42 and 43).

^bThese values are obtained using the collective model (Ref. 13) relations (Ref. 12) between B(E2) values and our experimental $0g \rightarrow I'K'$ matrix elements. The signs result from assuming that PPQ theory has correctly predicted the signs of the $0g \rightarrow I'K'$ matrix elements.

	$M(E2; IK \rightarrow I'K')$ (eb)				
	E ['K'	Exp		Collective	
$IK \rightarrow I'K'$	(keV)	(magnitude)	PPQ ^a	model ^b	
$0g \rightarrow 2g$	89	2.14(2)	-2.09		
$0g \rightarrow 2\gamma$	1154	0.35(3)	-0.38		
$0g \rightarrow 2\beta$	1129	0.12(7)	-0.14		
$0g \rightarrow 2o$	1258	< 0.09	-0.08		
$2g \rightarrow 2\gamma$		0.42(3)	-0.47	-0.41(3)	
$2g \rightarrow 2\beta$		0.28(2)	0.40	0.14(8)	
$2g \rightarrow 2o$		< 0.07 °	0.09	< 0.11	
$4g \rightarrow 2\gamma$		0.12(2)	-0.21	-0.09(1)	
$4g \rightarrow 2\beta$		0.31(19)	-0.58	-0.18(11)	
$4g \rightarrow 2o$		< 0.34	-0.15	_<0.14	

TABLE III. Matrix elements in ¹⁵⁶Gd compared with pairing-plus-quadrupole and collective models.

^a Predictions of the pairing-plus-quadrupole model calculations of Gupta *et al.* (Refs. 42 and 43).

^b These values are obtained using the collective model (Ref. 13) relations (Ref. 12) between B(E2) values and our experimental $0g \rightarrow I'K'$ matrix elements. The signs result from assuming that PPQ theory has correctly predicted the signs of the $0g \rightarrow I'K'$ matrix elements.

^cIn Ref. 14 the branching ratios were taken from Ref. 1 and are not corrected for the 87.4% M1 admixture in the 2_g-2_o transition. This result is corrected.

"Alaga rules"¹²; that is, ratios of the appropriate Clebsch-Gordan coefficients normalized to the measured $0^*_{g_*s_*} \rightarrow 2^*$ matrix elements. Overall agreement with PPQ theory is very good, even to predicting the small matrix element to the 2⁺ state at 1278 keV in ¹⁵⁶Gd.

E. Results and discussion for 3⁻ states

Several $I^{\pi} = 3^{-}$ octupole vibrational states were observed and the measured B(E3) strengths are presented in Table IV. Also presented are energies, proposed K values, and the predictions⁴⁴ of the microscopic calculations of Neergard and Vogel.^{45,46}

Our experimental results verify several predictions of the theory. To be noted is the very good agreement between theory and experiment in both absolute strength and energy level position. The inclusion of the Coriolis interaction is responsible for this agreement, as also is the case for the actinides.⁴⁶ For example, in ¹⁵⁶Gd, unperturbed random-phase approximation calculations yield B(E3) values for K=0, 1, 2, and 3 of 5.6, 6.1, 5.7, and 0.0 $e^{2}b^{3} \times 10^{-2}$ with energies of 1420, 1370, 1570, and 1840 keV, respectively.⁴⁵ But, as is observed experimentally, the collectivity concentrates in the state with the lowest energy. In addition, the measurements seem to verify the prediction that the collectivity decreases with increasing mass.

TABLE IV. Comparisons of measurements and micro-
scopic calculations of $B(E3; 0^+g.s. \rightarrow 3^-)$ for the one-octu-
pole-phonon 3 ⁻ states. The label α is the K quantum num-
ber corresponding to the largest component of the Cori-
olis-coupled wave function since K is not a "good" quan-
tum number for the mixed octupole states.

		Experiment			Theory ^a		
		E	B(E3)		E	B(E3)	
Nucleus	I [¶] K	(keV)	$(e^{2}b^{3} \times 10^{-2})$	α	(keV)	$(e^{2}b^{3} \times 10^{-2})$	
¹⁵⁶ Gd	3-(0)	1276	16(4)	0	1848	1.1	
				1	1309	15.2	
				2	1664	1.0	
				3	1818	0.0	
¹⁶⁰ Gd	(3-?)	1289	12.7(14)	0	1855	0.0	
				1	1607	0.9	
	,			2	1434	11.3	
				3	1882	0.0	
¹⁷⁶ Hf	3-2	1313	9.3(29)	0	1962	0.1	
				1	1642	0.9	
				2	1294	8.1	
				3	1890	4.9	
¹⁷⁸ Hf	3-2	1323	5.3(1)	0	2117	0.5	
				1	1549	0.0	
				2	1411	9.8	
				3	1806	4.4	

^aReferences 44 and 45.

V. CONCLUSION

We have used the (α, α') reaction to Coulombexcited $I^{\pi} = 2^{*}$ and 3⁻ vibrational states in even-even ¹⁵⁴⁻¹⁶⁰Gd and ¹⁷⁶⁻¹⁸⁰Hf nuclei to obtain precise values of the absolute B(E2) and B(E3) strengths.

For both the Gd and Hf nuclei, the 2^* states behave with marked similarity in trends with increasing mass of both level energy and E2 collectivity despite the opposite trend in ground state deformation. The $I^{*}=2^*$ members of second K=0excited bands have small, if any, B(E2) strength.

A pairing-plus-quadrupole model applied to the $I^{\pi} = 2^{+}$ states in ¹⁵⁴⁻¹⁵⁶Gd works very well and suggests further applicability of this model to well-deformed nuclei. For octupole-vibrational states, the theory of Neergard and Vogel works well if the Coriolis interaction is included.

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