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PHYSICAL REVIEW C

VOLUME 6, NUMBER 6

DECEMBER 1972

Decay of the Five-Quasiparticle Isomeric States in $^{177}\text{Hf}^\dagger$

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(Received 11 August 1972)

The decay of 51.4-min $^{177m2}\text{Hf}$ has been studied in detail by means of Ge(Li) detectors and a Si(Li) detector. The L -subshell conversion-electron ratios for the 214.0-keV isomeric transition were measured with a double-focusing β -ray spectrometer. An $E3$ multipolarity was deduced for the isomeric transition, and thus a spin-parity of $I^\pi = \frac{37}{2}^-$ was established for the isomeric state. Coincidence measurements identify the $K^\pi = \frac{23}{2}^+$ rotational band up to the $I = \frac{31}{2}$ member. The rotational-model quantity $(g_K - g_R)^2/Q_0^2$, derived from the γ -ray intensities of the cascade and crossover transitions, was used to deduce the gyromagnetic ratios for this band.

I. INTRODUCTION

In a recent article,¹ a new high-spin isomer $^{177m2}\text{Hf}$ ($t_{1/2} = 51.6$ min) was reported. The main features of its decay were deduced from γ -ray intensity measurements. This new high-spin isomer was interpreted as being a $K^\pi = \frac{37}{2}^-$ five-quasiparticle state with a possible Nilsson configuration of $\frac{7}{2}^+ [404]_p, \frac{9}{2}^- [514]_p, \frac{7}{2}^- [514]_n, \frac{9}{2}^+ [624]_n, \frac{5}{2}^- [512]_n$.

In this paper we report a more detailed study employing high-resolution γ -ray spectroscopy, Ge(Li)-Ge(Li) γ - γ coincidences, and conversion-electron measurements. These results confirm the previously deduced decay scheme of Ref. 1 establishing the $K^\pi = \frac{23}{2}^+$ rotational band up to the $I = \frac{31}{2}$ member and the spin-parity of the isomeric state. Evidence is also presented which suggests that the $K^\pi = \frac{37}{2}^-$ isomeric state may be populating a second three-quasiparticle band in ^{177}Hf .

II. EXPERIMENTAL TECHNIQUES AND RESULTS

Sources of $^{177m2}\text{Hf}$ were produced by irradiation of isotopically enriched $^{176}\text{Yb}_2\text{O}_3$ (96%) with 46-MeV α particles. The cross section of the $^{176}\text{Yb}(\alpha, 3n)^{177m2}\text{Hf}$ reaction at this energy was estimated to be ≈ 1.5 mb. Typically, sources of 10–20 μCi of $^{177m2}\text{Hf}$ were produced for each experiment. After radiochemical separation the only observed contaminant was 23.6-h ^{173}Hf , produced from other ytterbium isotopes.

The hafnium was separated carrier free from the irradiated targets in the following manner. The target was dissolved in hot 3 N HNO_3 . Hafnium was extracted for 5 min into a few milliliters of 0.5 M TTA in xylene. The organic phase was scrubbed 3 times with 3 N HNO_3 and was then gently evaporated to dryness in a platinum crucible.

The organic portion was removed by heating it several times with 13 *N* HF. The hafnium activity was then taken up in a few drops of the HF solution, evaporated onto either a nickel or platinum backing, and flamed to red heat. The measured cross section and the observed activity of these sources indicated that the bulk ($\approx 60\%$) of the radiohafnium was reclaimed by this method. The time required for the chemical separation was about 30 min.

Sources of hafnium used for mass separation were first radiochemically separated. 5 mg of hafnium carrier was added for the TTA-xylene extraction. The hafnium fraction was then back extracted from the organic phase with 0.5 *N* HNO₃-0.5 *N* HF solution, the fluoride ions complexed with boric acid, and the hydroxide of hafnium precipitated with concentrated NH₄OH. The hydroxide precipitate was dissolved in 3 *N* HNO₃ and hafnium tetramandellate precipitated with mandelic acid. Hafnium was then converted to oxide by heating at 800°C in a muffle furnace. The hafnium oxide was introduced into the ion source of the Brookhaven electromagnetic isotope separator and converted to the volatile chlorides by passing a controlled stream of carbon tetrachloride over the heated sample. The hafnium chlorides were subsequently ionized, accelerated, and mass analyzed. The collected ¹⁷⁷Hf sample was used for γ -ray measurements.

γ -ray spectra were obtained with a 39-cm³ Ge(Li) detector which has a resolution of 0.95 keV at 122 keV and 1.86 keV at 1332 keV. Figure 1 shows a γ -ray spectrum of a typical hafnium source. Analysis of the γ -ray spectra were performed by means of a modified version of the BRUTAL² computer code. The CLSQ³ program was used for γ -ray decay curve resolutions. The energy and efficiency calibrations were determined with International Atomic Energy Agency standard sources. Internal energy calibrations were performed by using several of the γ -ray lines of ^{177m}1Hf which are known⁴⁻⁸ to within ± 0.1 keV. All of the γ -rays in Fig. 1 have been assigned to ^{177m}2Hf with 1.1-sec ^{177m}1Hf in equilibrium, except for those labeled ¹⁷³Hf. Identical lines observed in the decay of the mass separated source confirmed the assignments of these γ -rays to ¹⁷⁷Hf. Table I lists the γ -ray energies and relative intensities of ^{177m}2Hf with ^{177m}1Hf in equilibrium. The measured half-lives of the more intense γ -rays ($I_{\gamma} > 40$) averaged 51.4 ± 0.5 min.

In Table II are listed the γ -ray energies and relative intensities assigned to the decay of ^{177m}2Hf. The 214.0-, 295.1-, and 326.7-keV γ rays of ^{177m}2Hf were observed as unresolved doublets with the 214.4-, 296.5-, and 327.7-keV γ -rays of 1.1-

sec ^{177m}1Hf in equilibrium. The γ decay of 1.1-sec ^{177m}1Hf has been studied previously⁴⁻⁶ from the β^- decay of 161-day ^{177m}Lu. However, Bernthal and Rasmussen⁷ observed a 10% increase in the relative intensities over the values previously reported^{4,5} though they have not reported all the values. We therefore found it necessary to produce sources of ^{177m}Lu for measurement of the relative γ -ray intensities of ^{177m}1Hf following the β^- decay of ^{177m}Lu. A table of the γ -ray energies and relative intensities of ^{177m}Lu measured in this work is given in the Appendix. These values are in good agreement with those of Bernthal and Rasmussen.^{7,8} Contributions from 1.1-sec ^{177m}1Hf in the γ -ray multiplets of the ^{177m}2Hf sources were resolved by use of the values included in the Appendix.

Ge(Li)-Ge(Li) γ - γ coincidences were measured for the 214.0-, 254.8-, 277.3-, 295.1-, 311.5-, 326.7-, 572.4-, 606.5-, and 638.2-keV γ -rays. Table III lists the results of these measurements. Accidental coincidences were typically 3-5% of the true coincidence counting rate. Figure 2 shows the coincidence spectrum for the 214.0-217.0-keV gated region. (The gate was set on the high-energy side of the 214.0-214.4-keV doublet in order to minimize contributions from the strong 208.4-keV γ ray.) The enhancements of the 277.3-, 295.1-, 311.5-, 326.7-, 572.4-, 606.5-, and 638.2-keV γ rays clearly indicate they are in coincidence with the 214.0-keV isomeric transition. This, together with the additional coincidence information in Table III, supports the decay scheme shown in Fig. 3.

The 120.5- and 254.8-keV γ rays were not observed in coincidence with the 214.0-keV isomeric transition, although there is weak evidence that they are in coincidence with the 277 ± 4 - and 327 ± 5 -keV gated regions. In the gated region (255 ± 4 keV), the 120.5-keV γ ray was strongly enhanced and is therefore definitely in coincidence with the 254.8-keV transition. As to the other γ rays in this gated spectrum (including the 277.3- and 325.7-keV transitions), they are presumably due to the presence of the Compton distributions of higher-energy γ rays in the gate, and cannot be definitely attributed to coincidences with the 254.8-keV transition. The assignments, therefore, of the 120.5- and 254.8-keV γ rays in coincidence with the 277 ± 4 - and 327 ± 5 -keV gated regions are listed as questionable in Table III.

Conversion-electron measurements were performed with the Brookhaven double-focusing β -ray spectrometer and a Si(Li) detector of 2-mm depth. The β -ray spectrometer was only employed to measure the *L*-conversion-electron lines of the 214.0-keV isomeric transition. The spectrome-

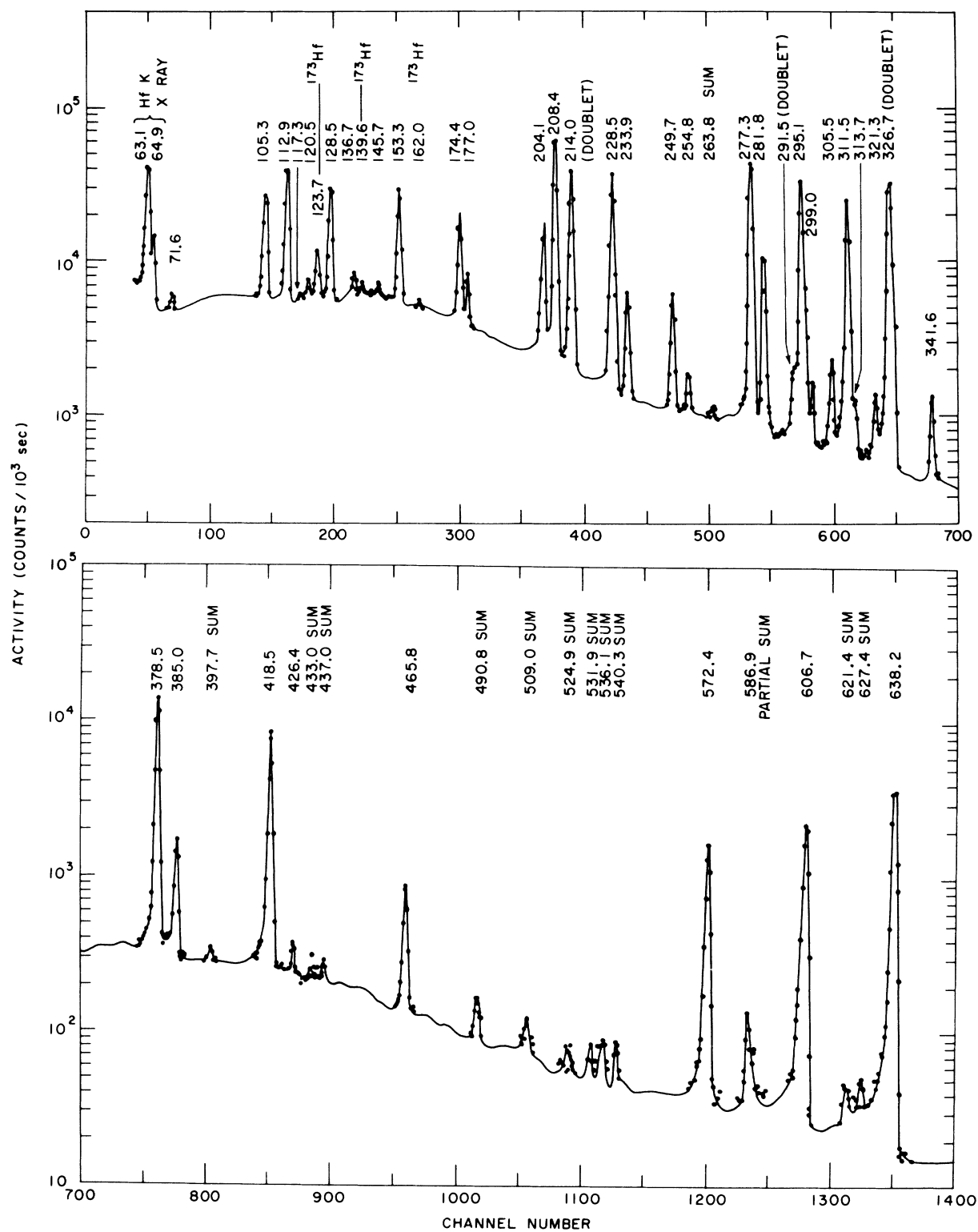


FIG. 1. Typical singles Ge(Li) γ -ray spectrum of radiochemically separated $^{177m_2}\text{Hf}$. All of the γ rays were assigned to $^{177m_2}\text{Hf}$ except those labeled ^{173}Hf .

TABLE I. γ -ray energies and intensities of $^{177m2}\text{Hf}$ and $^{177m1}\text{Hf}$ in equilibrium.

E_γ (keV) ^a	I_γ (Relative) ^b	E_γ (keV) ^a	I_γ (Relative) ^b
71.7	6.7 ± 0.7	291.4}	16.2 ± 1.9^c
105.3	$\equiv 100.0$	292.5}	
113.0	176.8 ± 12.7	295.1}	499.1 ± 38.4^c
117.2	1.4 ± 0.2	296.5}	
120.5	6.3 ± 1.0	299.0	14.9 ± 1.3
128.5	130.4 ± 10.0	305.5	15.1 ± 1.8
136.7	12.4 ± 3.4	311.5	386.8 ± 26.7
145.8	8.6 ± 0.8	313.7	9.6 ± 0.9
153.3	145.4 ± 11.2	321.3	16.0 ± 1.5
174.4	109.3 ± 8.3	326.7}	576.5 ± 43.8^c
177.0	30.2 ± 2.4	327.7}	
204.1	129.1 ± 9.9	341.6	17.8 ± 1.8
208.4	525.8 ± 40.5	378.5	239.8 ± 18.5
214.0}		385.1	26.3 ± 2.6
214.4}	323.6 ± 24.9^c	418.5	167.8 ± 13.1
228.5	318.1 ± 23.9	426.5	3.9 ± 0.5
233.9	47.1 ± 3.7	465.8	18.3 ± 1.6
249.7	53.2 ± 4.2	572.4	47.0 ± 3.8
254.8	8.8 ± 0.8	586.9	$\leq 3.6 \pm 0.5^d$
277.3	499.0 ± 39.0	606.5	75.9 ± 5.9
281.8}		638.2	133.2 ± 10.3
283.4}	124.8 ± 9.9^c		

^a Error in the γ -ray energy is ± 0.1 keV.

^b Normalized to $I_\gamma(105.3 \text{ keV}) \equiv 100.0$ units.

^c Doublet γ -ray intensities.

^d Some contribution of summing in intensity.

ter was operated at a momentum resolution of 0.2% and 1- or 2-min counts were taken starting at the lowest momentum setting. Figure 4 shows the results of the measurement. Each point has been corrected for decay and background. In addition to ≈ 7 -counts/min background of the spectrometer a source background of ≈ 200 counts/min was also observed. This source background was at-

tributed to the tails of the 277.3-keV K and 214.0-keV M lines. The uncertainties of L_I , L_{II} , and L_{III} intensities were ± 10 , ± 5 , and $\pm 5\%$, respectively. The L -subshell ratios of $L_I/L_{II}/L_{III} = 0.139/1/0.52$ measured in the present work were within experimental error of the theoretical values⁹ $L_I/L_{II}/L_{III} = 0.127/1/0.56$ for a pure $E3$ transition. (A limit of $<0.2\%$ $M4$ admixture in the 214.0-keV transition was deduced.)

Figure 5 shows the conversion-electron spectrum of $^{177m2}\text{Hf}$ measured with a Si(Li) detector. The detector had an energy resolution of 5 keV full width at half maximum. Table IV summarizes the results of the measurement. The conversion-electron intensities have been normalized to the intensity of the 214.0-keV L conversion-electron line. The γ -ray intensities (column 2) were taken from Table II, and the theoretical conversion coefficients (column 5) were obtained from the tables of Hager and Seltzer.⁹ The 212-keV electron line was observed as an unresolved multiplet of the 214.0-keV M and N lines and the 277.3-keV K line. The intensity of the 277.3-keV K line was determined by subtracting from the total line intensity the contributions of the 214.0-keV M and N lines. These contributions had been estimated from the 214.0-keV L -lines intensity and the theoretical $L/(M+N)$ ratio. $E2$ admixtures of 6–8% in the $M1$ cascade ($I \rightarrow I-1$) transitions (277.3, 295.1, 311.5, and 326.7 keV) were derived from the crossover/cascade γ -ray intensity ratios. However, owing to the large uncertainty of the K and L conversion-electron intensities, a precise measurement of the admixtures could not be made. The K/L ratios for the crossover ($I \rightarrow I-2$) transitions (572.4, 606.6, and 638.2 keV) are consistent with $E2$ multipolarity assignments.

TABLE II. γ -ray energies and intensities of $^{177m2}\text{Hf}$.

E_γ (keV)	Relative γ - ray intensity ^a	Multipolarity	Total conversion coefficient (α) ^b	Relative transi- tion intensity ^c
120.5	1.3 ± 0.3			
214.0	54.0 ± 3.5	$E3$	1.546	104.1 ± 10.0
254.8	1.8 ± 0.2			
277.3	$\equiv 100.0$	$M1 + \sim 8\% E2$	0.226	92.8
295.1	91.6 ± 7.5	$M1 + 7.4\% E2$	0.191	82.6 ± 7.9
311.5	77.5 ± 6.5	$M1 + 6.7\% E2$	0.166	68.4 ± 6.5
326.7	90.6 ± 7.8	$M1 + 6.4\% E2$	0.146	78.6 ± 7.9
572.4	9.4 ± 0.7	$E2$	0.013	7.2
606.5	15.2 ± 1.2	$E2$	0.011	11.6 ± 1.1
638.2	26.7 ± 2.0	$E2$	0.010	20.4 ± 2.0

^a Normalized to $I_\gamma(277.3 \text{ keV}) \equiv 100.0$ units.

^b Taken from Ref. 9.

^c Normalized to $T_\gamma(277.3 \text{ keV}) + T_\gamma(572.4 \text{ keV}) = 100.0$ units.

TABLE III. γ - γ coincidence results of $^{177m_2}\text{Hf}$. \times indicates observed coincidence, ? indicates doubtful coincidence.

Coin.	Gate	214	255	277	295	311	326	572	606	638
120.5			\times	\times ?			\times ?			
214.0				\times	\times	\times	\times	\times	\times	\times
254.8				\times ?			\times ?			
277.3		\times			\times	\times			\times	\times
295.1		\times		\times		\times				\times
311.5		\times		\times	\times		\times	\times		
326.7		\times		\times	\times	\times		\times	\times	
572.4		\times				\times	\times			
606.5		\times		\times			\times			
638.2		\times		\times	\times			\times		

III. DISCUSSION

$K^\pi = \frac{23}{2}^+$ Rotational Band

The energy and spin-parity of the 1.1-sec $K^\pi = \frac{23}{2}^+$ isomeric state of ^{177}Hf have been well established⁴⁻⁸ from the β decay of 161-day ^{177m}Lu ($K^\pi = \frac{23}{2}^-$). This isomeric state has been assigned⁶ a Nilsson configuration of $\frac{7}{2}^+[404]_p, \frac{9}{2}^-[514]_p, \frac{7}{2}^-[514]_n$. From the decay of the 51.4-min isomeric state of ^{177}Hf , the rotational members up to $I = \frac{31}{2}$ were established¹ within the $K^\pi = \frac{23}{2}^+$ band (see Fig. 3). Substitution of the three lowest energy levels (E_I) of the $K^\pi = \frac{23}{2}^+$ band into the formula

$$E_I = E_0 + AI(I+1) + BI^2(I+1)^2 \quad (1)$$

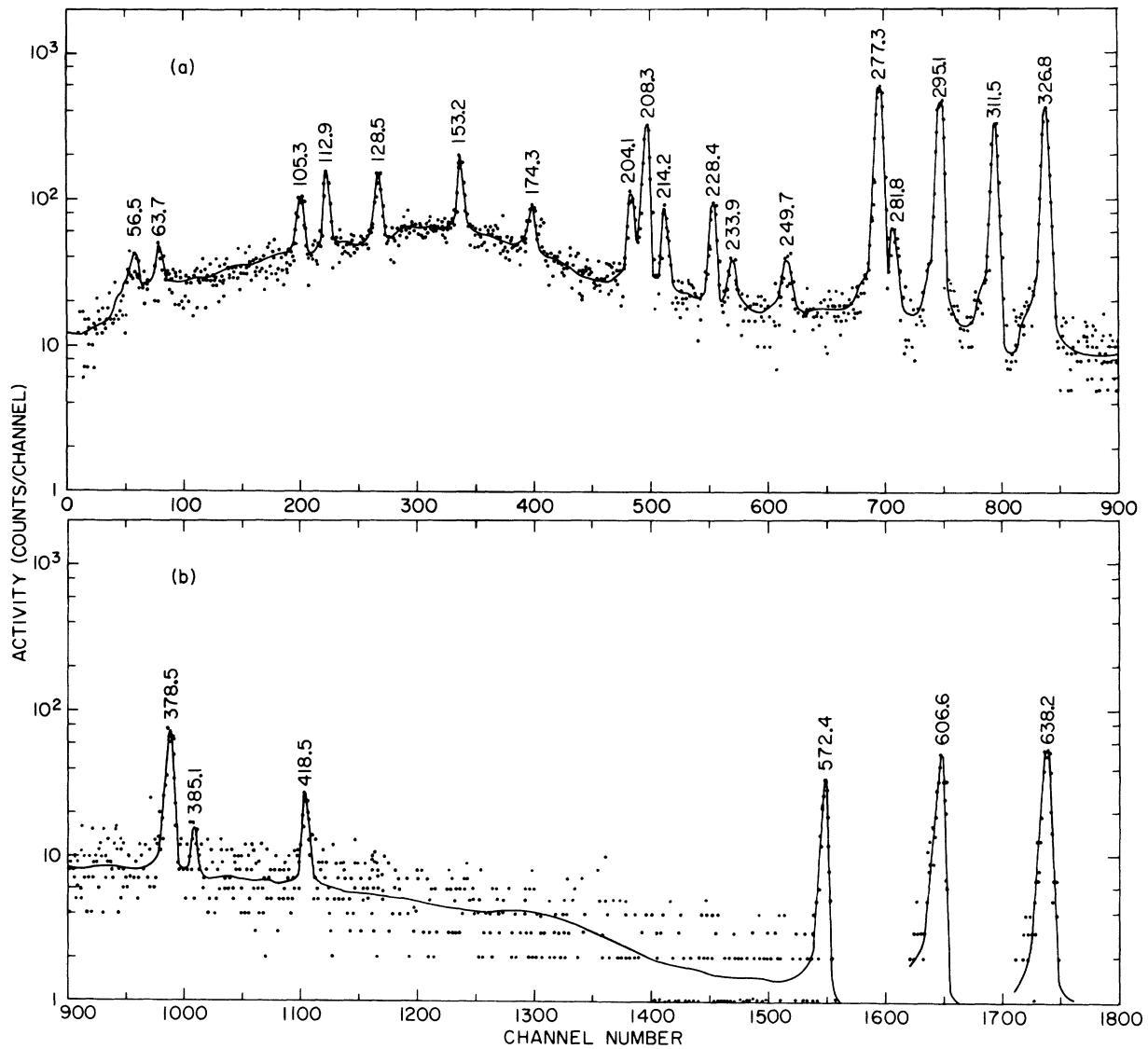


FIG. 2. Ge(Li)-Ge(Li) γ - γ coincidence spectrum gated on the 214.0-217.0-keV γ -ray region.

gave values of $A = 12.07$ keV and $B = -3.12$ eV. Energies of the remaining two band members calculated from these parameters agreed to within ± 1.6 keV with the experimental values. The parameters determined for the $\frac{7}{2}^- [514]_n$ ground-state band were $A = 12.83$ keV and $B = -6.87$ eV. It is remarkable that the parameters for these two bands are so similar.

Assuming the Alaga¹⁰ rules are valid for the $E2$ components of the cascade and crossover transitions within the $K^\pi = \frac{23}{2}^+$ band, one can calculate the $E2/M1$ mixing ratios (δ^2) of the cascade transitions and the gyromagnetic factors, g_K and g_R , within the framework of the rotational model. The $E2/M1$ mixing ratio (δ^2) can be derived from the following formula

$$1/\delta^2 = \left(\frac{E_\gamma}{E_{\gamma'}} \right)^5 \frac{[(2I+1)(I-2+K)(I-1-K)]}{(2K)^2(2I-1)\lambda} - 1, \quad (2)$$

where λ is the experimental crossover/cascade ratio with γ and γ' denoting the crossover ($I \rightarrow I-2$) and cascade ($I \rightarrow I-1$) transitions, respectively. The rotational-model quantity $(g_K - g_R)^2/Q_0^2$ can then be obtained from the following formula:

$$\frac{1}{\delta^2} = \frac{2.87 \times 10^5 (2I+2)(2I-2)}{E_{\gamma'}^2} \frac{(g_K - g_R)^2}{Q_0^2}, \quad (3)$$

where $E_{\gamma'}$ is in keV. In Table V are listed the experimental crossover/cascade ratios (λ), the $E2/M1$ mixing ratios (δ^2), and the quantity $(g_K - g_R)^2/Q_0^2$ for the three cascade transitions within the $K^\pi = \frac{23}{2}^+$ band. The values of $(g_K - g_R)^2/Q_0^2$ are constant within the experimental error with an averaged value of $(5.52 \pm 0.61) \times 10^{-3}$. Inserting the value¹¹ $Q_0 = 6.74$ b determined for the ground state, one obtains $g_K - g_R = 0.50 \pm 0.03$.

Assuming the contributions of the three particles of the $K^\pi = \frac{23}{2}^+$ state are additive, one obtains

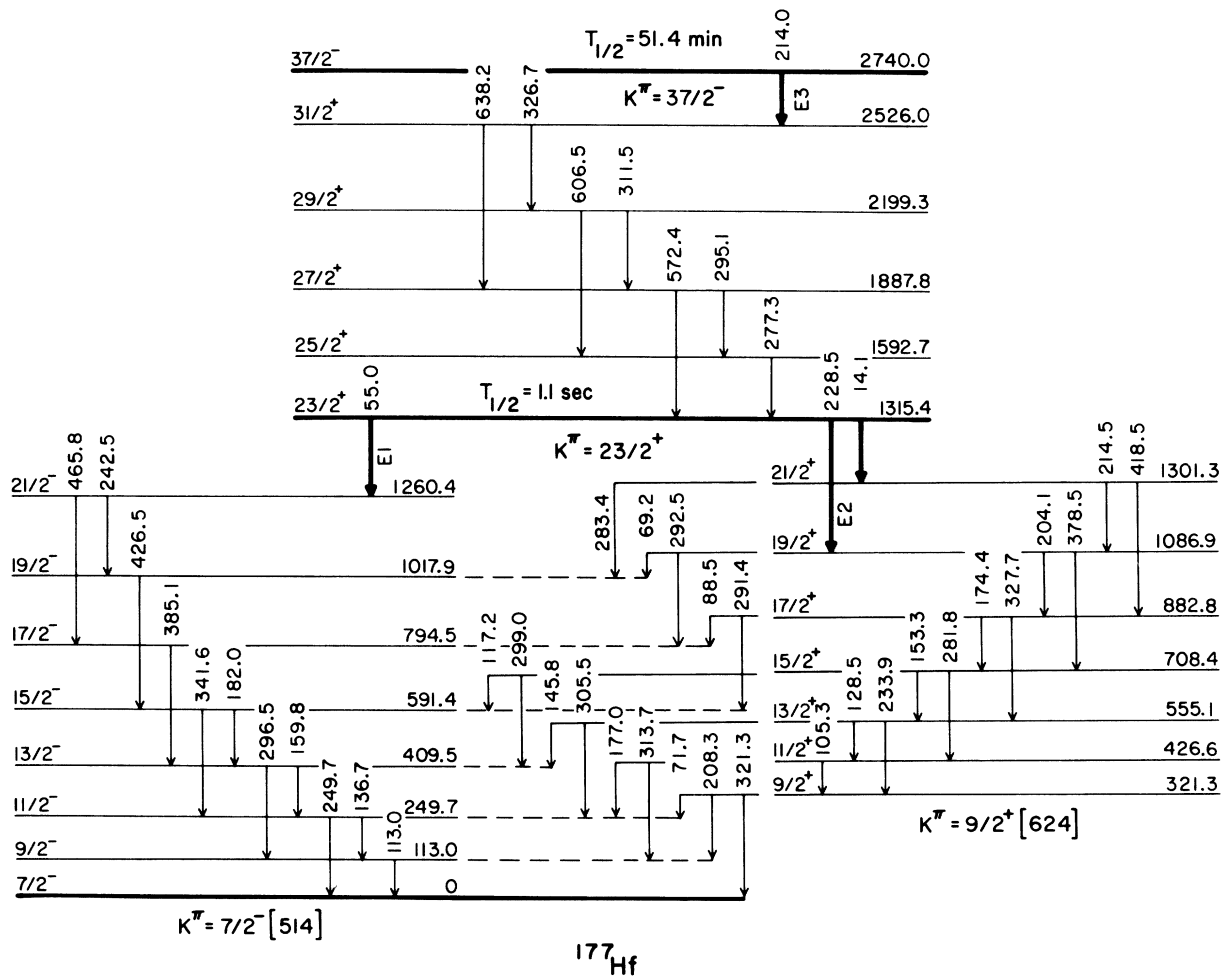


FIG. 3. Proposed decay scheme of 51.4-min $^{177m2}\text{Hf}$ ($K^\pi = \frac{37}{2}^-$). The decay of 1.1-sec $^{177m1}\text{Hf}$ ($K^\pi = \frac{23}{2}^+$) was previously (Refs. 4-8) well established.

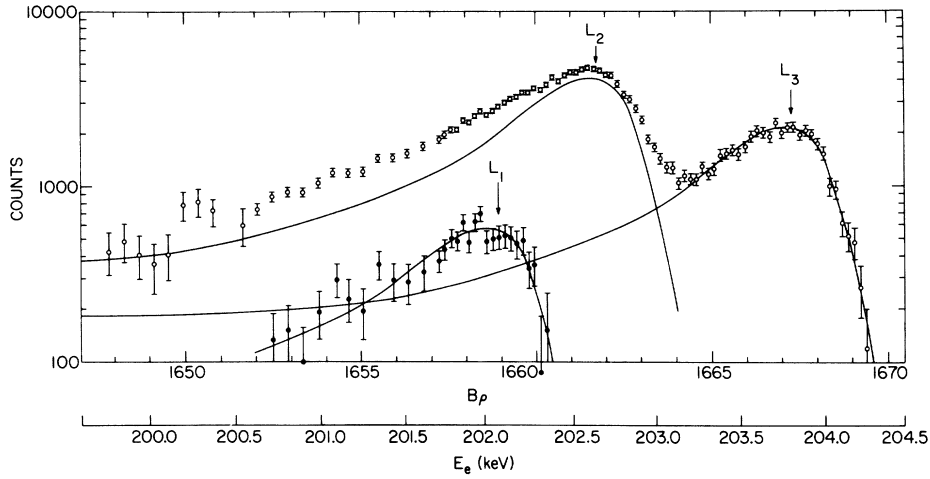


FIG. 4. L -conversion-electron spectrum of the 214.0-keV isomeric transition of $^{177m2}\text{Hf}$. The spectrum was obtained with a double-focusing β -ray spectrometer.

for the intrinsic g factor the expression

$$g_K = \frac{1}{K} (\Omega_1 g_{\Omega 1} + \Omega_2 g_{\Omega 2} + \Omega_3 g_{\Omega 3}), \quad (4)$$

where Ωg_{Ω} for the individual particles are given^{12, 13} by

$$\Omega g_{\Omega} = \Omega g_i + (g_s - g_i) \langle S_z \rangle. \quad (5)$$

The calculated g_K value is relatively insensitive to the assumed g_s factor and the deformation. We have used a value of $g_s^{\text{eff}} = (0.512 \pm 0.032)g_s^{\text{free}}$ (where g_s^{free} is 5.585 for protons and -3.826 for neutrons) which was experimentally derived¹⁴ for the $\frac{7}{2}^- [514]_n$, ^{177}Hf ground state. The expectation values of the z component of the spin ($\langle S_z \rangle$) were calculated using Nilsson¹² wave functions with a deformation¹⁵ of $\eta = 4.88$. The calculated values of $\langle S_z \rangle$ were -0.374 , -0.455 , and 0.451 for the $\frac{7}{2}^- [514]_n$, $\frac{9}{2}^- [514]_p$, and $\frac{7}{2}^+ [404]_p$ orbitals, respectively. Using the above values in Eqs. (4) and (5) we calculate $g_K = 0.76$ for the $K^\pi = \frac{23}{2}^+$ state. The collective gyromagnetic factor (g_R) can now be determined using our experimental value of $g_K - g_R = 0.50 \pm 0.03$, yielding $g_R = 0.26 \pm 0.03$. This value of g_R is in agreement with the predicted¹⁵ values, $g_R = 0.24$ and 0.28 , and the experimental value¹⁴ for the ^{177}Hf ground state, $g_R = 0.253 \pm 0.013$.

$$K^\pi = \frac{37}{2}^- \text{ Isomeric State}$$

A spin-parity of $\frac{37}{2}^-$ was assigned to the isomeric state based on the data presented above, with the $K^\pi = \frac{23}{2}^+$ rotational band established up to the $I^\pi = \frac{31}{2}^+$ rotational level by an $E3$ transition. A state with $K^\pi = \frac{37}{2}^-$ can only be explained if one assumes it to be a five-quasiparticle state. No combinations of three-quasiparticle Nilsson orbitals are available at this excitation energy that could produce this high spin. The lowest-energy five-quasiparticle state expected in ^{177}Hf is the $K^\pi = \frac{37}{2}^-$ state with a Nilsson configuration of $\frac{7}{2}^+ [404]_p$, $\frac{9}{2}^- [514]_p$, $\frac{7}{2}^- [514]_n$, $\frac{9}{2}^+ [624]_n$, $\frac{5}{2}^- [512]_n$.

Formation of the high-spin state can be viewed as a coupling of the $\frac{7}{2}^- [514]_n$ ground state of ^{177}Hf with protons from a broken pair in the $\frac{7}{2}^+ [404]_p$ and $\frac{9}{2}^- [514]_p$ orbitals and neutrons from a broken

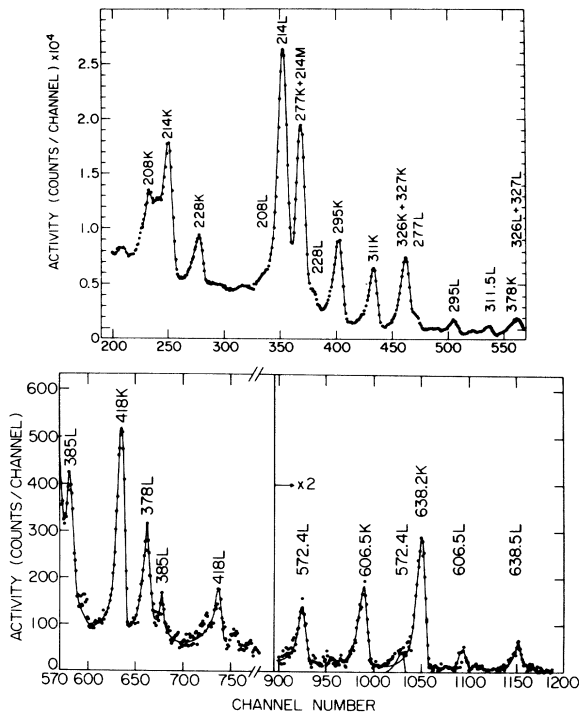


FIG. 5. Conversion-electron spectrum of $^{177m2}\text{Hf}$ taken with a 2-mm-depth Si(Li) detector which had a resolution of 5 keV full width at half maximum.

pair in the $\frac{5}{2}^- [512]_n$ and $\frac{9}{2}^+ [624]_n$ orbitals. The expected energy above the ground state is

$$E(\frac{37}{2}^-) \approx 2\Delta_p + 2\Delta_n, \quad (6)$$

where Δ_p and Δ_n are the proton and neutron pairing energies respectively, obtained from cranking-model calculations.¹⁵ The pairing energies (Δ_p and Δ_n) can be identified with the odd-even mass differences (P_p and P_n). Using the values $\Delta_p = P_p = 780$ keV and $\Delta_n = P_n = 720$ keV given by Prior, Bohem, and Nilsson,¹⁵ one estimates the five-quasiparticle energy at ≈ 3.0 MeV. A closer approximation can be obtained by using the three-quasiparticle energy of the $K^\pi = \frac{23}{2}^+$ state and the odd-even mass difference for the broken neutron pair,

$$E(\frac{37}{2}^-) \approx E(\frac{23}{2}^+) + 2\Delta_n, \quad (7)$$

which yields 2755 keV above the ground state. This is to be compared with the value of $E(\frac{37}{2}^-) = 2740.0$ keV in Fig. 3.

The hindrance factor¹⁶ (H) calculated for the K -forbidden $E3$ isomeric transition is 2.41×10^5 , or a factor of 22 for each degree of forbiddenness, $\nu = \Delta K - L = 4$. Such a hindrance factor would be expected for a transition between states with relatively pure K quantum numbers. However, it should be noted that the K -forbiddenness factor ($f \approx 22$) for the 214.0-keV transition is about 3 to 5 times smaller than for other $E3$ K -forbidden transitions in this mass region. For example, $f(E3) \approx 62$ and ≈ 96 , respectively, for ^{177m}Lu (K^π

$= \frac{23}{2}^-$) and $^{179m2}\text{Hf}$ ($K^\pi = \frac{25}{2}^-$).¹⁷ The decrease in the K -forbidden factor for $^{177m2}\text{Hf}$ may be an indication that either: (a) The initial state or the final state may be admixed with other five- or three-quasiparticle states, respectively, or (b) due to the similarity of the initial and final state configurations the decay of the two additional neutrons is somewhat more favored over the configurations involved in the decay of ^{177m}Lu and $^{179m2}\text{Hf}$.

The 120.5- and 254.8-keV γ -rays were observed to decay with the same half-life of 51.4-min $^{177m2}\text{Hf}$ but were not observed in coincidence with the 214.0-keV isomeric transition. However, they were found to be in coincidence with each other. One can exclude the possibility that they belong to a weak branch in the decay of the $K^\pi = \frac{23}{2}^+$ isomeric state, since its decay has been well studied⁴⁻⁷ (see Appendix). One can also exclude the possibility of contamination, since these γ rays were observed in the γ -ray spectrum of mass-separated $^{177m2}\text{Hf}$. We believe that these weak transitions indicate that the $K^\pi = \frac{37}{2}^-$ five-quasiparticle state may be populating a second three-quasiparticle band in ^{177}Hf . The 120.5- and 254.8-keV γ -rays have been observed¹⁸ with in-beam spectroscopy through the $^{176}\text{Yb}(\alpha, 3n)^{177}\text{Hf}$ reaction and were found to have lifetimes of 10–20 nsec. Owing to the low intensity of these γ rays we could not establish other members in this decay branch. There are only two other Nilsson orbital configurations available for construction of other three-quasipar-

TABLE IV. Relative conversion-electron intensities of $^{177m2}\text{Hf}$.

E_γ (keV)	I_γ^a	Multipole order	Conversion line	α_{Theory}^b	$I_{\gamma\alpha_{\text{th}}}^c$	$I_{\text{ce(Exp.)}}^c$
214.0	54.0 ± 3.5	$E3$	K	4.298×10^{-1}	51.7	52.6 ± 1.7
			L	8.320×10^{-1}	$\equiv 100.0$	$\equiv 100.0$
277.3	$\equiv 100.0$	$M1/E2$	K	1.971×10^{-1}	43.9	45.3 ± 2.1^d
			L	3.022×10^{-2}	6.7	7.2 ± 0.3
295.1	91.6 ± 7.5	$M1/E2$	K	1.666×10^{-1}	34.0	34.7 ± 1.5
			L	2.550×10^{-2}	5.2	5.3 ± 0.3
311.5	77.5 ± 6.5	$M1/E2$	K	1.440×10^{-1}	24.8	28.1 ± 1.3
			L	2.201×10^{-2}	3.8	4.5 ± 0.3
326.7	90.6 ± 7.8	$M1/E2$	K	1.267×10^{-1}	25.5	29.4 ± 1.3
			L	1.933×10^{-2}	3.9	4.7 ± 0.4
572.4	9.4 ± 0.7	$E2$	K	1.099×10^{-2}	0.230	0.220 ± 0.015
			L	2.266×10^{-3}	0.047	0.045 ± 0.005
606.6	15.2 ± 1.2	$E2$	K	9.651×10^{-3}	0.327	0.345 ± 0.021
			L	1.928×10^{-3}	0.065	0.070 ± 0.007
638.2	26.7 ± 2.0	$E2$	K	8.622×10^{-3}	0.512	0.562 ± 0.042
			L	1.679×10^{-3}	0.100	0.114 ± 0.010

^a See Table II.

^b Theoretical conversion coefficient (α) taken from the tables of Hager and Seltzer (Ref. 9). The conversion coefficients listed for the $M1/E2$ admixed transitions are the values for pure $M1$ transitions.

^c The relative conversion-electron intensities are normalized to I_{ce} (214-keV L line) $\equiv 100.0$ units.

^d See text for explanation of intensity.

TABLE V. The g factors and branching ratios for the $K^\pi = \frac{23}{2}^+$ rotational band in ^{177}Hf . λ is the experimental ratio between crossover ($I \rightarrow I-2$) and cascade ($I \rightarrow I-1$) transitions, δ^2 the $E2/M1$ mixing ratios of the ($I \rightarrow I-1$) transitions, and $(g_K - g_R)^2/Q_0^2$ a rotational model parameter.

I	λ	δ^2	$10^3(g_K - g_R)^2/Q_0^2$
$\frac{27}{2}$	0.103 ± 0.007	0.080 ± 0.008	5.23 ± 0.63
$\frac{29}{2}$	0.196 ± 0.017	0.072 ± 0.007	5.63 ± 0.51
$\frac{31}{2}$	0.294 ± 0.027	0.068 ± 0.007	5.70 ± 0.68

ticle states within ≈ 0.5 -MeV excitation energy of the $K^\pi = \frac{23}{2}^+$ state, namely,

$$\left(\frac{7}{2}^+[404]_p, \frac{9}{2}^-[514]_p, \frac{9}{2}^+[624]_n\right)K^\pi = \frac{25}{2}^-$$

and

$$\left(\frac{7}{2}^-[514]_n, \frac{9}{2}^+[624]_n, \frac{5}{2}^-[512]_n\right)K^\pi = \frac{21}{2}^+.$$

The estimated band-head energies for the $K^\pi = \frac{25}{2}^-$ and $\frac{21}{2}^+$ three-quasiparticle states were calculated as ≈ 1640 and ≈ 1440 keV, above the ground state, respectively, by use of an equation similar to Eq. (7).

ACKNOWLEDGMENTS

The authors would like to acknowledge the thoughtful advice and encouragement of Dr. J. B. Cumming and Dr. S. Katcoff throughout this investigation. The authors would also like to express their appreciation to Dr. A. W. Sunyar for the use of his Si(Li) detector system and his invaluable assistance in the conversion-electron measurements. We also acknowledge the many valuable discussions during the course of this work with our colleagues here at Brookhaven National Laboratory, in particular Dr. G. S. Goldhaber and Dr. M. L. Perlman, and with Professor G. T. Emery at Indiana University. Assistance provided by the operators of the Brookhaven cyclotron and High Flux Beam Reactor facilities in obtaining irradiations is gratefully acknowledged.

TABLE VI. γ -ray energies and relative intensities of ^{177m}Lu .

E_γ (keV) ^a	I_γ (Relative) ^b	E_γ (keV) ^a	I_γ (Relative) ^b
71.7	7.4 ± 0.7	249.7	51.3 ± 2.5
105.3	$\equiv 100.0$	268.8	28.3 ± 1.5
113.0	179.1 ± 8.4	281.8	116.7 ± 5.3
115.9	5.0 ± 0.5	283.4	4.3 ± 0.6
121.6	48.7 ± 2.9	291.4	14.9 ± 1.3 ^c
128.5	126.9 ± 5.7	292.5	
136.7	11.4 ± 1.1	296.5	44.5 ± 2.7
145.8	7.5 ± 0.8	299.1	14.3 ± 1.0
147.1	30.2 ± 2.3	305.5	14.5 ± 1.2
153.3	150.0 ± 7.1	313.7	11.5 ± 0.8
159.8	5.0 ± 0.6	319.0	85.7 ± 4.7
171.8	41.0 ± 2.2	321.3	11.6 ± 0.9
174.4	105.3 ± 5.3	327.7	145.9 ± 6.4
177.0	28.9 ± 1.8	341.6	14.9 ± 1.3
181.9	0.8 ± 0.2	367.4	24.8 ± 1.6
195.6	7.2 ± 0.7	378.5	232.2 ± 10.7
204.1	119.0 ± 5.6	385.1	24.5 ± 1.6
208.4	510.0 ± 22.4	413.7	137.5 ± 7.0
214.4	54.6 ± 3.1	418.5	167.0 ± 8.4
218.1	25.1 ± 3.0	426.5	3.4 ± 0.4
228.5	310.0 ± 12.6	465.9	19.4 ± 1.3
233.9	47.1 ± 2.3		

^a Error in the γ -ray energies is ± 0.1 keV.

^b Normalized to $I_\gamma(105.3 \text{ keV}) = 100.0$ units.

^c Doublet γ -ray intensity.

APPENDIX

Sources of 161-day ^{177m}Lu were produced by 5-h irradiation of 3 mg of 75% enriched $^{176}\text{Yb}_2\text{O}_3$ in the Brookhaven High Flux Beam Reactor ($\phi_n \approx 2 \times 10^{14}$ neutrons $\text{sec}^{-1} \text{cm}^{-2}$). The 6.7-day ^{177}Lu decayed during the 120-day prior to mass separation. The source strength after mass separation was 9×10^5 dis/min. The γ -ray energies and relative intensities of ^{177m}Lu are listed in Table VI. These measurements were used to determine contributions of ^{177m}Hf to the unresolved multiplets in $^{177m2}\text{Hf}$.

[†]Research supported by Advanced Research Projects Agency (ARPA order No. 1590) and the U. S. Atomic Energy Commission.

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PHYSICAL REVIEW C

VOLUME 6, NUMBER 6

DECEMBER 1972

Nuclear Orientation Study of the Decays of ^{126, 127, 128}Sb[†]

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(Received 10 July 1972)

Angular distributions have been measured for a number of γ rays emitted by ^{126, 127, 128}Sb polarized at low temperatures in iron. From the 0–90° anisotropies of the angular distributions, taking the proposed spin values $J=8$ (^{126, 128}Sb) and $J=\frac{1}{2}$ (¹²⁷Sb), the ground-state magnetic moments have been deduced to be: $|\mu(^{126}\text{Sb})| = 1.28 \pm 0.07 \mu_N$; $|\mu(^{127}\text{Sb})| = 2.59 \pm 0.12 \mu_N$; $|\mu(^{128}\text{Sb})| = 1.31 \pm 0.19 \mu_N$. Based on the anisotropies of the ¹²⁷Te γ rays the following ¹²⁷Te spin assignments are favored (level energies in keV): $\frac{3}{2}^+$ (0), $\frac{1}{2}^+$ (61), $\frac{11}{2}^-$ (88), $\frac{9}{2}^-$ (341), $\frac{5}{2}^+$ (473), $\frac{7}{2}^-$ (632), $\frac{7}{2}^+$ (686), $\frac{5}{2}^+$ (784), $\frac{9}{2}^-$ (786), $\frac{7}{2}^+$ (924), $\frac{5}{2}^+$ or $\frac{9}{2}^+$ (1077), $\frac{5}{2}^+$ (1142), $\frac{5}{2}^+$ (1290). Based on these spin assignments, $E2/M1$ mixing ratios have been deduced for a number of ¹²⁷Te γ rays.

I. INTRODUCTION

In a number of recent publications,¹⁻³ we have reported investigations of the angular distribution of γ rays emitted by ^{122, 124, 125}Sb nuclei polarized in Fe at low temperatures. Among the parameters which were deduced from these measurements were the angular momentum multipolarities of the β - and γ -radiation fields and the Sb ground-state magnetic moments. We report here similar investigations of the decays of ^{126, 127, 128}Sb.

II. DECAY SCHEMES

The decays of ^{126, 127, 128}Sb to levels of ^{126, 127, 128}Te are illustrated in Figs. 1, 2, and 3, respectively. β and γ radiations and conversion electrons emitted in the decay of the 12.4-day ¹²⁶Sb have been investigated by Orth, Dropesky, and Freeman,⁴ and spin assignments for the higher-lying ¹²⁶Te states have been proposed by Kiselev *et al.*⁵; the decay scheme of Fig. 1 is based primarily on the latter work. β and γ radiations from the decay of 3.9-day ¹²⁷Sb were studied by Ragaini, Gordon, and Walters (RGW)⁶ and by Takemoto, Iwashita, and Kageyama.⁷ The measurement of RGW⁶ proposed several spin assignments; the level scheme of Fig. 2 is taken primarily from that work, with additional spin and parity assignments based on

the present work included. β and γ spectra from the 8.6-h ¹²⁸Sb decay were investigated by Kiselev *et al.*,^{5, 8} and γ -ray coincidence and conversion-electron studies were done by Kerek.⁹

Additional investigations of the ¹²⁶Te levels have been done through the (d, p) reaction by Graue *et al.*¹⁰ and through the ($\alpha, 2n$) reaction by Kerek.¹¹ The ¹²⁷Te levels have also been studied by Graue *et al.*¹² through the (d, p) reaction. The reaction data are consistent with the level identifications proposed in the above decay studies.

III. EXPERIMENTAL DETAILS

A. Sample Preparation

The Sb isotopes were produced by α -induced fission of U. Following chemical separation of the Sb, a small amount of ¹²⁵Sb was added; the well-understood decay of ¹²⁵Sb³ permits its use as thermometer and also as a verification of the success of the Sb diffusion into the Fe. A quantity of this solution was placed on the surface of a 0.1-mm 99.99%-pure Fe foil, and the Sb was allowed to diffuse into the Fe by annealing at 1100°C for 2 h in a H₂-Ar atmosphere. During annealing, ⁵⁴Mn and ⁵⁷Co activities, which had previously been adsorbed onto the walls of the annealing crucible, diffused into the foil. The low-energy γ