Mössbauer-Effect Studies in Hafnium-Metal Single Crystals*

P. BOOLCHAND, B. L. ROBINSON, AND S. JHAT Case Western Reserve University, Cleveland, Ohio 44106 (Received 28 April 1969; revised manuscript received 10 July 1969)

The Mössbauer effect in Hf¹⁷⁸ has been studied with oriented single-crystal absorbers of hafnium metal at the temperature of liquid helium. Along the [0001] axis, only the $|\Delta m| = 1$ component of the quadrupole spectrum is observed, and along the [1010] axis only the $|\Delta m| = 1$ and $|\Delta m| = 2$ components are observed. Combining these observations with those made using a polycrystalline hafnium-metal absorber, the magnitude of the electric field gradient was found to be $V_{zz} = (0.95 \pm 0.04) \times 10^{18} \text{ V/cm}^2$, and the asymmetry parameter $\eta < 0.1$. The Debye temperature of hafnium carbide has been estimated to be (350 ± 10) °K, and the Debye temperature of hafnium metal has been found to be (227 ± 10) and (192 ± 13) °K, for γ rays propagating parallel and perpendicular to the c axis, respectively.

I. INTRODUCTION

HEN the nuclear Zeeman components of a Mössbauer line are not well resolved, the application of an external magnetic field will impose an axis of quantization; along the field direction some of the components will be suppressed and the remaining ones will be more clearly resolved. In this paper, we demonstrate that it is possible to improve the resolution of electric quadrupole spectra in the Mössbauer effect by performing polarization experiments with oriented single-crystal absorbers.1 This method suppresses some of the quadrupole components. The success of the technique lies in the strikingly different radiation patterns of the quadrupole components relative to the axis of quantization.

We have applied this technique to the study of the nuclear quadrupole interaction in hafnium metal using the 93-keV γ ray in hafnium-1782-5 with oriented single-crystal absorbers of hafnium metal. The 2-nsec 2+ first excited state of hafnium-178 6 emits a pure electric quadrupole radiation, as indicated in Fig. 1(a). In the presence of an axially symmetric quadrupole interaction (asymmetry parameter $\eta = 0$), only the $|\Delta m| = 1$ component is excited along the z axis of the electric field gradient (EFG), and only the $|\Delta m| = 1$ and $|\Delta m| = 2$ components are excited along directions perpendicular to the z axis of the EFG, as shown in Fig. 1(b). This arises from the nature of the radiation patterns of the quadrupole transition, which are shown in Fig. 1(c). The point group symmetry of hafnium metal in the (hcp) α phase⁷ (P6₃/mmc) requires the z component of the EFG to lie along the crystallographic c axis. Since V_{zz} lies along a threefold axis of rotational symmetry, it follows that the EFG should be axially symmetric.8 It may be pointed out

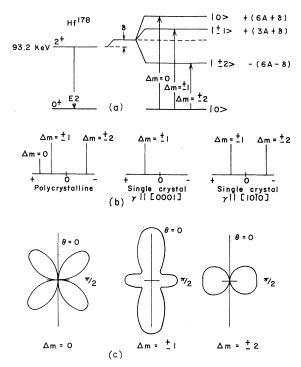


Fig. 1. (a) Quadrupole interaction in hafnium metal; (b) relative intensities of the quadrupole components of the Mössbauer line in polycrystalline hafnium metal and in oriented single crystals of hafnium metal; and (c) angular distributions of the $\Delta m = 0$, $|\Delta m| = 1$, and $|\Delta m| = 2$ quadrupole components with respect to the z axis of the EFG.

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[†] Present address: University of Cincinnati, Cincinnati, Ohio. Present address: Education Research Center, Massachusetts

Institute of Technology, Cambridge, Mass.

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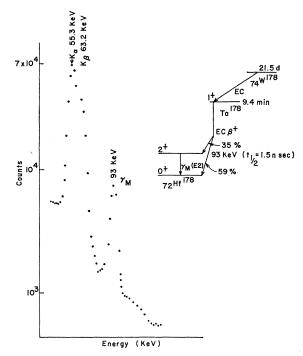


Fig. 2. Partial decay scheme of the A=178 chain: pulse-height spectrum of the K x rays and Mössbauer γ ray taken with a 25-cm³ Ge(Li) detector.

that the EFG in hafnium metal has been extensively studied through the perturbed angular correlation of the γ rays of tantalum-181 in polycrystalline hafniummetal lattice, 9-11 and the asymmetry parameter η has been found to be 0.3. However, Salomon et al. 12 studied the perturbed angular correlation of the γ rays of tantalum-181 in oriented hafnium-metal single crystals and concluded that η is zero. Apart from the inconsistency of these results, it is not clear that this method measures the EFG at a regular site in hafnium metal.

II. SOURCES AND ABSORBERS

Twenty-day $_{76}\rm W^{178}$ decays to 9.4-min $_{75}\rm Ta^{178},$ which feeds the 93-keV state of $_{74}\rm Hf^{178}$ with a 35% branch, as indicated in Fig. 2(a). Sources of W178 were made by bombarding 0.010-in. tantalum-metal foils with 43-MeV protons in the cyclotron of the Lawrence Radiation Laboratory, Berkeley. No chemical separation was needed: The tantalum-metal foil was merely etched to remove the oxidized surface layer and then annealed in vacuo for about 1 day at about 1000°C. The spectrum

of low-energy γ rays taken with a 25-cm³ Ge(Li) detector is shown in Fig. 2(b): The 93-keV γ ray is clearly resolved from the x rays and stands well above the background. Sources of W178 were also made by the bombardment of tantalum-carbide targets with 43-MeV protons and by the bombardment of enriched hafnium oxide (Hf¹⁷⁷) with 40-MeV α particles.

For the absorbers, thin platelets of hafnium metal were spark-cut along the [0001] and [1010] axes from a large single-crystal rod. 18 The single-crystal cut along the [0001] direction was etched to a thickness of $125\pm10~\mu$, and the crystal cut along the $\lceil 10\bar{1}0 \rceil$ direction was etched to a thickness of $75\pm10~\mu$. The singlecrystal absorbers could be oriented to within $\frac{1}{2}$ of arc using the back-reflection Laue technique; however, the angular resolution of the Mössbauer-effect experiments was about 2° of arc for the source-absorber distance used.

III. EXPERIMENTS AND RESULTS

The Mössbauer-effect experiments were performed with both the source and the absorber at the temperature of liquid helium in a cryostat described earlier. 14,15 The γ rays were detected in a Ge(Li) detector 3 mm thick and 4 cm² in area. In early experiments, sources of W178 in the tantalum-carbide host lattice were used. Although a large effect was observed because of the high Debye temperature of tantalum carbide, the narrowest line observed with this source and an

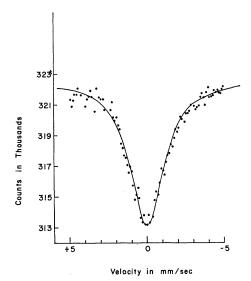


Fig. 3. Mössbauer spectrum taken with a W^{178} source in a tantalum-metal host lattice and a natural hafnium-carbide absorber, both at the temperature of liquid helium.

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absorber of hafnium carbide had a full width at halfmaximum (FWHM) of 4 mm/sec, which is twice 2Γ .

A Mössbauer spectrum taken with the source of W178 in the tantalum-metal host and a natural hafniumcarbide absorber (10 mg/cm² of Hf¹⁷⁸) is shown in Fig. 3. The FWHM is 2.6 ± 0.1 mm/sec, which is 1.3times 2Γ . The excess width is entirely attributable to the finite thickness of the absorber. Spectra were taken with different thicknesses of hafnium carbide, and from the observed excess width of the Mössbauer lines, the Debye temperature of hafnium carbide was calculated¹⁶ to be $(350\pm10)^{\circ}$ K at the temperature of liquid helium. In making this calculation, we took the total internal conversion coefficient¹⁷ to be 4.72 and the maximum resonance cross section to be 14×10^{-19} cm².

A Mössbauer spectrum taken with the polycrystalline absorber of hafnium metal is given in Fig. 4(c). Spectra made with the polycrystalline metallic absorbers were fitted by the method of least squares to three Lorentzians, with no restrictions on line positions or intensities. The interval between the $\Delta m = 0$ and $|\Delta m| = 1$ components, 0.85±0.11 mm/sec, was very nearly equal to one-third the interval between the $|\Delta m| = 1$ and $|\Delta m| = 2$ components, 2.2 ± 0.1 mm/sec, from which it is concluded that $\eta < 0.10$, which is consistent with the expectation from the crystal symmetry that η be zero.

The Mössbauer spectrum taken with a hafniummetal single crystal oriented so that the γ rays were transmitted parallel to the c axis shows a single line [Fig. 4(a)], while the spectrum taken with the γ rays transmitted perpendicular to the c axis and parallel to the $\lceil 10\overline{1}0 \rceil$ axis shows only two lines $\lceil \text{Fig. 4(b)} \rceil$. It is clear that these spectra confirm the predicted suppression of the quadrupole components. It can be noted that the positions of the $|\Delta m| = 1$ and $|\Delta m| = 2$ lines in the spectrum made with the polycrystalline absorber coincided with those in the spectra made with the single-crystal absorbers. From the interval observed in Fig. 4(b), the quadrupole interaction in hafnium metal at the temperature of liquid helium is found to be $e^2qQ = -1.83 \pm 0.06$ µeV. Using the electronically measured mean life⁶ of the 93-keV state of Hf¹⁷⁸ and the theory of transition rates among the rotational levels of deformed nuclei, 17,18 we obtain

$$\begin{array}{l} B(E2)_{\rm ex} \! = \! 4.08 \! \times \! 10^{-61} \, E_{\gamma}^{-5} \tau^{-1} \! \big[1 \! + \! \alpha_T \big]^{\! -1} \\ = \! Q_0^2 \! 5 / 16 \pi \end{array}$$

and

$$Q(I,K) = Q_0[3K^2 - I(I+1)]/(I+1)(2I+3)$$
,

for $B(E2)_{ex}$ in cm⁴ with E_{γ} in MeV and τ in sec, where α_T is the total internal conversion coefficient and the

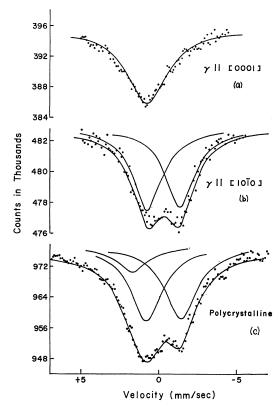


Fig. 4. Mössbauer spectra with hafnium-metal absorbers; (a) oriented single-crystal absorber with γ rays propagating parallel to the c axis; (b) oriented single-crystal absorber with γ rays propagating perpendicular to the c axis; and (c) polycrystalline absorber.

electric quadrupole moment of the 93-keV state was calculated to be $Q = -1.93 \pm 0.05$ b. The value of the EFG is then $V_{zz} = +(0.95\pm0.04)\times10^{18} \text{ V/cm}^2$.

The room-temperature perturbed angular correlation studies of Marklund et al. 11 and of Sommerfeldt et al. 10 gave $\eta = 0.3$ as mentioned above, and their results for the EFG are considerably smaller than ours. Although it seems likely that the magnitude of the EFG should decrease as the temperature increases, it is difficult to imagine that the asymmetry parameter, a quantity intimately tied to the structural symmetry of the lattice, would change with temperature and develop a nonvanishing value.

Gerdau et al.3 obtained a Mössbauer spectrum with a polycrystalline hafnium absorber and a supposed singleline source. They fitted their spectrum to three Lorentzians assuming $\eta = 0$, and they obtained a value of -1.752 ± 0.024 µeV for the quadrupole interaction and a value of $+(0.90\pm0.03)\times10^{18}$ V/cm² for the EFG. By a similar procedure, Snyder et al.5 obtained a value of -1.84 ± 0.01 µeV for the quadrupole interaction energy in hafnium metal.

Finally, it may be pointed out that the three components of the Mössbauer spectrum taken with the

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