



HOKKAIDO UNIVERSITY

Title	Study of localized character of 4 f electrons and ultrasonic dispersions in SmO _s 4Sb ₁₂ by high-pressure high-frequency ultrasonic measurements
Author(s)	Mombetsu, S.; Murazumi, T.; Hidaka, H. et al.
Citation	Physical Review B, 94(8), 085142 https://doi.org/10.1103/PhysRevB.94.085142
Issue Date	2016-08-25
Doc URL	https://hdl.handle.net/2115/63045
Rights	©[2016] American Physical Society
Type	journal article
File Information	PhysRevB.94.085142.pdf



Study of localized character of $4f$ electrons and ultrasonic dispersions in $\text{SmOs}_4\text{Sb}_{12}$ by high-pressure high-frequency ultrasonic measurements

S. Mombetsu, T. Murazumi, H. Hidaka, T. Yanagisawa,* and H. Amitsuka
Department of Physics, Hokkaido University, Sapporo 060-0810, Japan

P.-C. Ho
Department of Physics, California State University, Fresno, California 93740, USA

M. B. Maple
Department of Physics and Center for Advanced Nanoscience, University of California, San Diego, La Jolla, California 92093, USA
 (Received 30 March 2016; revised manuscript received 1 August 2016; published 24 August 2016)

We present high-frequency ultrasonic measurements on the filled skutterudite $\text{SmOs}_4\text{Sb}_{12}$ under hydrostatic pressure. The results clarify that the $4f$ electrons in this compound transform from delocalized at ambient pressure to localized at high pressures with a crossover pressure of approximately 0.7 GPa. This drastic change in the $4f$ electrons under pressure is apparently related to the non-Fermi-liquid state, which appears in an intermediate-pressure range of 0.5–1.5 GPa. The results of our analysis strongly suggest that the ferro-octupolar interaction becomes dominant at high pressure. Moreover, we report the pressure dependence of the ultrasonic dispersion, which is due to rattling, over a wide range of ultrasonic frequencies up to 323 MHz. The drastic change in the ultrasonic dispersions and the frequency-dependent elastic anomaly in the C_{11} mode at lower temperatures imply a possible coupling between rattling phonons and $4f$ electrons.

DOI: [10.1103/PhysRevB.94.085142](https://doi.org/10.1103/PhysRevB.94.085142)

I. INTRODUCTION

A large variety of quantum phenomena has been found in rare-earth-based intermetallic compounds, such as heavy-fermion (HF) behavior, multipolar order, and unusual types of superconductivity [1–3]. The c - f hybridization (i.e., hybridization between conduction electrons and $4f$ electrons in rare-earth ions) plays an important role in these systems. The c - f hybridization causes two competitive effects: the Kondo effect and the Ruderman-Kittel-Kasuya-Yosida (RKKY) interaction. The Kondo effect leads to itinerant $4f$ electrons that lose their local moments at low temperature. In contrast, the RKKY interaction, which is an intersite interaction between $4f$ electrons via conduction electrons, causes an ordering of local multipole moments. Especially for systems based on $\text{Ce}(4f^1)$ and $\text{Yb}(4f^{13})$, which have one electron or one hole in the $4f$ orbital, these two effects compete with one another and are well described by the Doniach phase diagram, which is a good guide for understanding this competitive feature [4]. The strength of c - f hybridization can be controlled by external parameters, such as magnetic field, chemical substitution, or pressure. For Ce compounds, when a magnetic order temperature is suppressed towards absolute zero by an external parameter, unusual physical properties, such as non-Fermi-liquid (NFL) behavior and superconductivity [5–7], have been found in the vicinity of the quantum critical point. However, the c - f hybridization and its quantum criticality for the plural $4f$ -electron systems [i.e., systems with more than two electrons in the $4f$ orbital such as $\text{Pr}(4f^2)$, $\text{Nd}(4f^3)$ and $\text{Sm}(4f^5$ or $4f^6)$] remain poorly understood.

Recently, rare-earth-based compounds with a highly symmetrical cage-like crystal structure have been attracting attention because they are prone to develop c - f hybridization and/or low-lying highly degenerate crystalline-electric-field (CEF) states, which potentially have multipolar degrees of freedom. Such states have been observed in plural $4f$ -electron systems. In Pr-based cage-structured compounds, several novel physical properties have been observed, such as the quadrupolar Kondo effect in $\text{PrV}_2\text{Al}_{20}$ (which has yet to be confirmed) [8], HF behavior, the unusual superconductivity of $\text{PrOs}_4\text{Sb}_{12}$ [9], and the unusual charge order in $\text{PrRu}_4\text{P}_{12}$ [10]. Novel physical properties are also observed in Sm-based compounds, for example, a magnetically ordered state that is unusually insensitive to external magnetic field and HF behavior in $\text{SmT}_2\text{Al}_{20}$ ($T = \text{Ti, V, Cr}$) [11,12]. The origin of the magnetically insensitive state and the role of c - f hybridization in such Sm-based compounds have not yet been clarified.

The filled skutterudite $\text{SmOs}_4\text{Sb}_{12}$ with cubic space group $\text{Im}\bar{3}$ (T_h^5 , No. 204) is a good candidate for studying the c - f hybridization effect of a Sm-based system. This compound has a large Sommerfeld coefficient γ of approximately $820 \text{ mJ mol}^{-1} \text{ K}^{-2}$, which indicates HF behavior at low temperature [13,14]. This value for γ remains almost constant up to 8 T [14], in contrast to typical Ce-based HF compounds [15], where the Kondo effect is weakened when subjected to an external magnetic field. The characteristic temperature T^* , below which $4f$ electrons become itinerant, is estimated to be approximately 20 K for $\text{SmOs}_4\text{Sb}_{12}$ in zero magnetic field [16]. Below T^* , this material undergoes a phase transition at $T_C \approx 2.6 \text{ K}$, accompanied by a weak spontaneous ferromagnetic (FM) moment of $0.02 \mu_B$ – $0.03 \mu_B$ per Sm ion [13,14]. Because this value is much smaller than that of the magnetic moment of a free Sm^{3+} ion ($0.71 \mu_B$), the ordered state is considered to be due to itinerant FM order. The average valence of the

*Corresponding author: tatsuya@phys.sci.hokudai.ac.jp

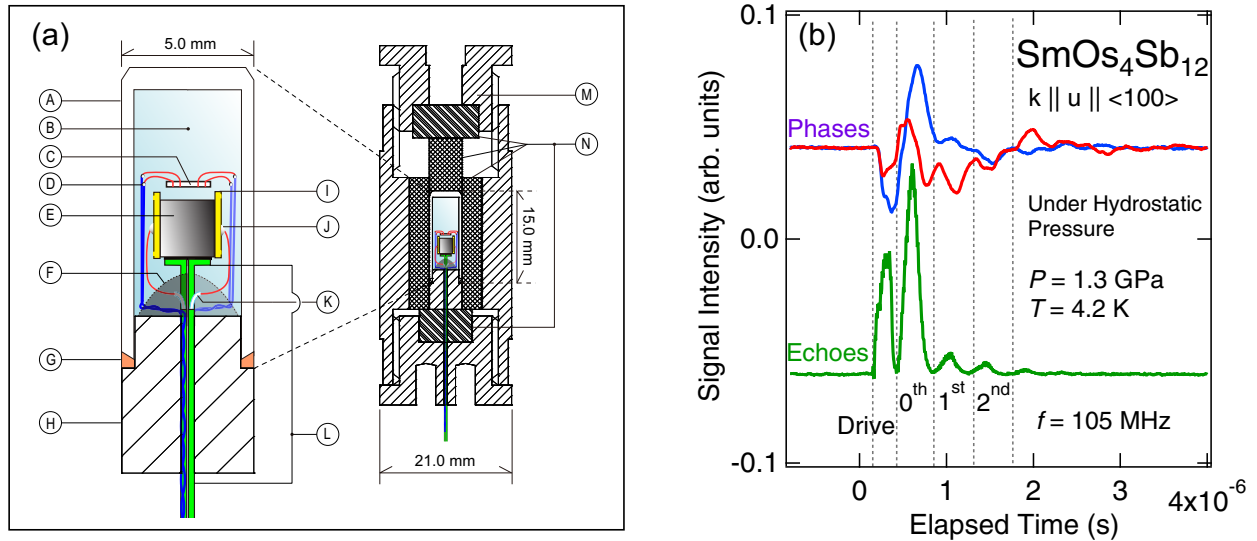


FIG. 1. (a) Schematic illustrations of hydrostatic-pressure ultrasonic measurement setup. A: Teflon capsule, B: Daphne oil 7373 for pressure medium, C: tin stick for a manometer, D: Cu and Au line, E: sample, F: Stycast 2850, G: Cu-Be sealing ring, H: body of a CuBe plug, I: LiNbO₃ transducer, J: Ag paste, K: Teflon insulation shield and inner conductor of semirigid coaxial cable, L: Cu outer shield of semirigid coaxial cable, M: Cu-Be parts (nuts and cylinder), N: NiCrAl cylinder and WC pusher. (b) Ultrasonic echo signal (green) and phase signals (blue and red) of the longitudinal C_{11} mode at 105 MHz at 1.3 GPa, 4.2 K.

Sm ion is approximately $+2.83$ at 150 K, which decreases with decreasing temperature and levels off to a constant value of approximately $+2.76$ near T^* [17]. Such anomalous coexisting of the midvalence state and heavy-fermion state in $\text{SmOs}_4\text{Sb}_{12}$ could possibly be due to the coincidence of weak c - f hybridization and the small energy difference between Sm divalent and trivalent states [18]. On the other hand, the $\log T$ behavior on the temperature dependence of the valence change indicates an unconventional Kondo effect associated with Sm $4f$ -electron charge degrees of freedom [19]. A large anharmonic vibration of a guest ion, known as rattling, has been detected in this material as well as in $\text{PrOs}_4\text{Sb}_{12}$ and other skutterudites. Reference [20] discusses the possible contribution to the HF state of charge fluctuations due to rattling of the rare-earth ion.

Upon applying hydrostatic pressure to $\text{SmOs}_4\text{Sb}_{12}$, T_C increases, whereas T^* decreases [16]. These opposite tendencies imply a recovery of the localized character of $4f$ electrons when under pressure. The jump in the specific heat $\Delta C/T_C$, which occurs at T_C , becomes larger with increasing pressure, which suggests that the ordered state is stabilized under pressure. In contrast, the maximum in the ac susceptibility at T_C becomes smaller under pressure and seems to disappear at higher pressures [21,22]. These results imply that the ordered state at high pressures is not a simple FM ordered state but a higher-rank multipolar type of order. Thus, the possible multipolar contribution in $\text{SmOs}_4\text{Sb}_{12}$ should be investigated by using the appropriate probe for detecting the higher-rank multipoles.

By using ultrasonic measurements, the response of electric-quadrupole moments of localized $4f$ electrons can be obtained spectroscopically [23]. In fact, ultrasonic measurements have detected the possible effects of multipoles in numerous rare-earth-based compounds [24,25]. Recently, ultrasonic measurements of $\text{SmOs}_4\text{Sb}_{12}$ in a pulsed magnetic field have

shown clear evidence of the Γ_{67} -quartet CEF ground state [26]. These results also suggest the existence of a Kondo-like screened state and its suppression by strong magnetic fields. Furthermore, ultrasonic dispersions, i.e., frequency-dependent elastic anomalies accompanied by a peak in ultrasonic attenuation, appear near 20 and 55 K in $\text{SmOs}_4\text{Sb}_{12}$ [27]. These anomalies are considered to be resonances between thermally activated rattling and ultrasonic strain. Thus, ultrasonic measurements with c - f hybridization controlled by pressure should provide new information on multipoles, rattling, and their consequences for the exotic HF state of $\text{SmOs}_4\text{Sb}_{12}$. Here we present the results of high-frequency ultrasonic measurements of $\text{SmOs}_4\text{Sb}_{12}$ under hydrostatic pressure up to 1.3 GPa.

II. EXPERIMENTS

In this study, we used single-crystalline $\text{SmOs}_4\text{Sb}_{12}$ grown using the Sb self-flux method. The single-phase nature of the present sample was confirmed by x-ray diffraction. The crystalline quality of the specimen was also checked by electrical resistivity and magnetic susceptibility at ambient pressure, which are consistent with previous reports. A well-isolated ultrasonic echo signal with a frequency of 105 MHz (wavelength of $\approx 40 \mu\text{m}$) in the present crystal as seen in Fig. 1(b) is also a good benchmark to evidence the bulk sample's homogeneity. To generate and detect ultrasonic waves, LiNbO₃ transducers were fixed on the sample using superglue (GEL-10: Toagosei Co. Ltd.). To carry high-frequency signals under hydrostatic pressure, two ultrathin semirigid coaxial cables (SC-033/50: Coax Co. Ltd.), which comprise Cu-shield sleeves with a diameter of 0.33 mm ϕ and Au-coated W-core cables with impedance-adjusted (50Ω) diameters of 0.08 mm ϕ , were installed in the pressure cell. As shown in Fig. 1(a), the sample was mounted on

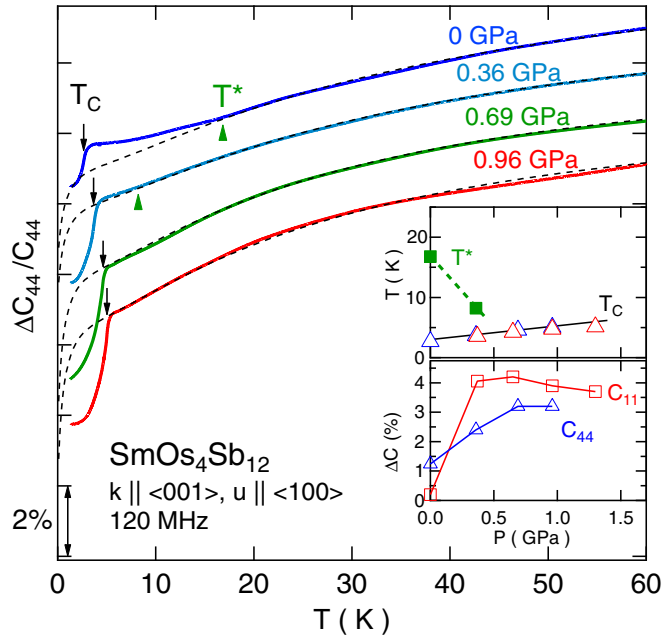


FIG. 2. Normalized elastic constant C_{44} versus temperature measured at various hydrostatic pressures (solid lines). The data are shifted vertically for clarity. Black dashed curves represent the calculated elastic constant. The upward and downward arrows indicate the characteristic temperature T^* and the ordering temperature T_C , respectively. The top inset shows a T - P phase diagram, and the bottom inset shows the jump in the elastic constants at T_C as a function of hydrostatic pressure.

the delaminated outer Cu conductors of the coaxial cables, which served as thermal anchors between the sample in the pressure cell and the thermometers outside the pressure cell. The edge of the dielectric (polytetrafluoroethylene, PTFE) and shield tubes (Cu) were carefully sealed with Stycast 2850 to prevent pressure from leaking. With this setup, we obtained clear ultrasonic-echo signals under a high pressure of 1.3 GPa at 4.2 K [see Fig. 1(b)]. Thus, this setup allowed us to make ultrasonic measurements at variable frequencies up to 323 MHz even under high pressure.

III. EXPERIMENTAL RESULTS

A. Transverse C_{44}

Figure 2 shows the relative change in the elastic constant C_{44} as a function of temperature measured at various pressures. Below 80 K, C_{44} decreases with decreasing temperature (i.e., softening) at all pressures studied. This softening can be explained by the strain-quadrupole coupling. The black dashed curves in Fig. 2 represent the calculated relative change $C_{44} = C_{44}^0 - \frac{Ng^2\chi_{\Gamma_4}}{(1+g'\chi_{\Gamma_4})}$, where C_{44}^0 , N , g , χ_{Γ_4} , and g' are the background elastic constant, the number of magnetic ions per unit volume, the quadrupole-strain coupling constant, the single-ion quadrupolar susceptibility, and the intersite quadrupole-quadrupole coupling constant, respectively [23]. In the present analysis, we assume that the mixed-valence state makes little contribution to the calculation since a small pressure dependence of the averaged valence at lowest

temperature within 0.04% up to 0.9 GPa is reported [28]. Here we discuss only the relative change in the elastic constants as a function of temperature and do not determine the absolute value of C_{44}^0 and g . The intersite quadrupole-quadrupole coupling constant g' is negative at all of the pressures, which suggests antiferro quadrupolar interaction. g' is -1.55 and -1.50 K at ambient pressure and at 0.35 GPa, respectively, and at 0.69 and 0.96 GPa, g' is -0.90 K. The value of C_{44} calculated on the basis of the localized 4f-electron model with the CEF ground state $\Gamma_{67}(0\text{ K})$ – $\Gamma_5(20\text{ K})$ reproduces well the softening above the characteristic temperature $T^* \approx 17$ K. However, below T^* , the calculated curve deviates from the measured data. This deviation can be explained by the suppression of the quadrupole moments of the 4f electrons due to the c - f hybridization that strengthens with decreasing temperature. Thus, we can treat T^* as the crossover temperature between the delocalized and localized electron state. Under a hydrostatic pressure of 0.36 GPa, the calculation can reproduce the softening of C_{44} down to 8 K. Such a decrease in T^* as determined in this study is roughly consistent with the electrical resistance and nuclear-magnetic-resonance measurements under hydrostatic pressure, which leads to an estimate of T^* that decreases slowly with increasing pressure [16]. Above 0.7 GPa, the calculation appropriately reproduces all the features of C_{44} down to T_C . Thus, we conclude that the 4f electrons recover their localized character at 0.7 GPa.

Such well-defined localized character of Sm under pressure advocated by present data is, however, inconsistent with the valence determined with x-ray absorption spectroscopy (XAS). The pressure dependence of the averaged valence determined with XAS methods shows no significant change at low temperature up to 3.5 GPa [19], even though a more significant effect of pressure on the valence would be expected if an oversized cage caused the midvalence state of Sm in the present compound. On the other hand, a relatively small change of the “temperature dependence” of the valence has been seen at around 1 GPa, at which the rattling of the Sm ion also seems to be strongly affected (as discussed later). Thus, it is obvious that ultrasonic measurement is rather sensitive for the detection of the 4f electron’s local charge degrees of freedom. At this moment, it is still controversial whether such an apparent contradiction comes from an unconventional Kondo effect due to rattling [20] or misinterpretation of either experimental results.

The controversy of the valence aside, the present results for C_{44} under pressure still provide further confirmation that the CEF ground state of the present compound is the Γ_{67} quartet because the Γ_5 CEF-ground-state model predicts that the softening of the C_{44} mode levels off below ≈ 10 K [27]. In the present analyses, however, we assume that the CEF level splitting ($\Delta \approx 20$ K at ambient pressure) remains unchanged in the hydrostatic pressure below 1 GPa for simplification. If the level splitting Δ is set as a free parameter, alternatively, the intersite quadrupole-quadrupole coupling constant needs to be changed drastically in order to reproduce the temperature and pressure dependence of C_{44} regarding the deviation at T^* . Therefore, further measurements are needed to get more information on the pressure dependence of the CEF splitting and c - f hybridization.

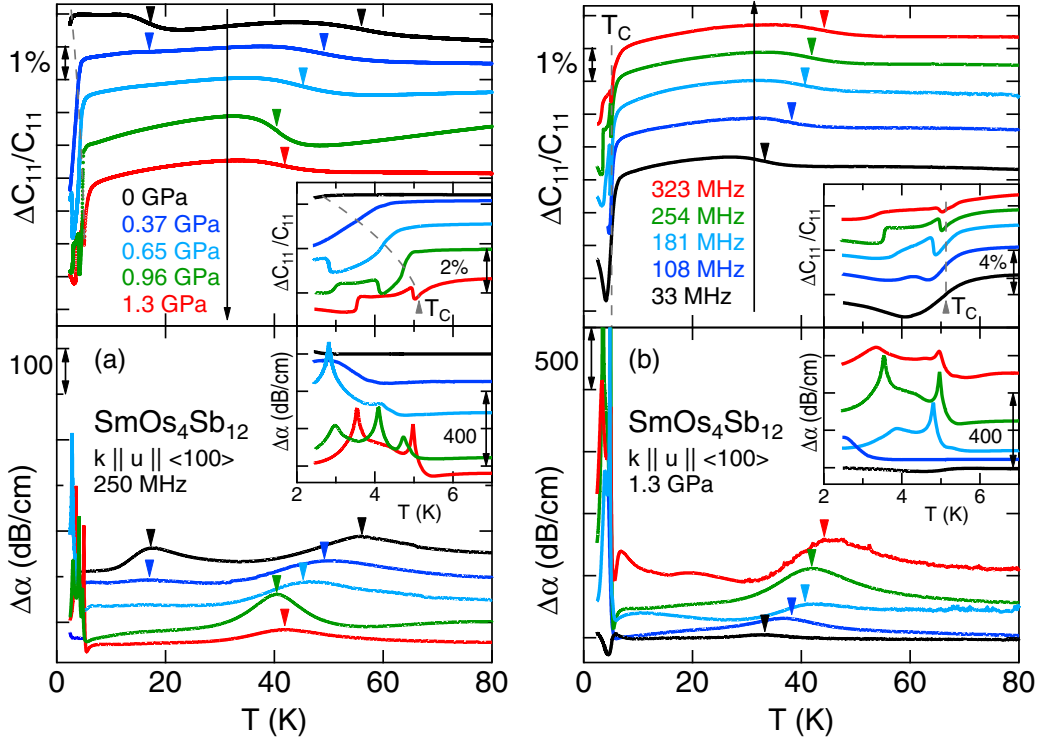


FIG. 3. Normalized elastic constant of C_{11} vs temperature (a) measured at 250 MHz and under various hydrostatic pressures and (b) measured at 1.3 GPa and at various frequencies. The downward arrowheads indicate the temperatures at which the resonance relation, $\omega\tau \sim 1$, is satisfied. The insets give enlarged views of the low-temperature range. The bottom panels show the corresponding ultrasonic attenuations. The data have been shifted vertically for clarity.

B. Longitudinal C_{11}

Figure 3(a) shows the relative change in the elastic constant C_{11} as a function of temperature measured at 250 MHz and under various hydrostatic pressures. At ambient pressure, C_{11} reveals shoulderlike anomalies near 15 and 55 K which are ultrasonic dispersions (UDs) and an abrupt decrease near T_C due to magnetic ordering, as described in Ref. [27]. The elastic anomalies of the UD reveal a clear dependence on pressure. Between these UD, C_{11} softens, which is also due to the strain-quadrupole coupling. This softening must become less strong below the crossover temperature $T^* \approx 17$ K in the same manner as C_{44} , although the presence of the UD near 15 K makes it difficult to distinguish the crossover phenomena in the C_{11} mode. Upon applying pressure, the softening of C_{11} becomes larger and continuous down to T_C , which is again explained by a recovery of the localized character of the $4f$ electrons, as discussed for C_{44} . The relatively small upturn near 15 K at ambient pressure becomes smaller once pressure is applied and seems to disappear near 0.65 GPa [see inset of Fig. 3(a)]. Below T_C , C_{11} reveals another steplike elastic anomaly for pressures exceeding 0.65 GPa. The anomaly and its dependence on frequency are discussed in detail below [Fig. 3(b)].

The insets in Fig. 2 show a phase diagram and the jump in the elastic constants ΔC of C_{11} and C_{44} at T_C as functions of pressure. The anomalies at T_C of both elastic constants become larger at higher pressure, which is also explained by a recovery of the localized character of $4f$ electrons. This drastic change might have the same origin as the disappearance of the nuclear

quadrupole resonance signal over a wide range at 0.4 GPa [29]. Moreover, the peak in ac susceptibility at T_C increases rapidly after approximately 0.5 GPa is applied [22]. These results imply that the ordered state changes drastically from itinerant FM order to localized FM order after a pressure greater than 0.4 GPa is applied. As shown by the phase diagram in the inset in Fig. 2, the pressure at which T^* drops below T_C can be estimated by extrapolation to be 0.7 GPa. Note that this pressure is close to the pressure region where the exponent n of the temperature dependence of electrical resistivity is a minimum [16]. Thus, these results strongly suggest that the NFL characteristic is related to the change in the localized character of the $4f$ electrons.

IV. DISCUSSION

A. Mean-field analysis of quadrupole susceptibility

We now discuss the ordered state in $\text{SmOs}_4\text{Sb}_{12}$ at high pressure, where the $4f$ electrons are well localized, based on a comparison of the experimental results with a calculation based on the mean-field approximation. Because weak FM ordering appears at ambient pressure, we begin the calculation by considering a primary FM order. We also consider the intersite interaction of the Γ_4 magnetic-octupole moments $T_i^\beta = (\bar{J}_i J_j^2 - \bar{J}_k^2 J_i)$, where $(i, j, k) = (x, y, z), (z, x, y)$, or (y, z, x) and the bar represents normalized symmetrization. The Hamiltonian is

$$H = H_{\text{CEF}} - K_{\text{DD}} \langle \mathbf{J} \rangle \mathbf{J} - K_{\text{OO}} \langle T^\beta \rangle T^\beta - g_{\Gamma} O_{\Gamma} \varepsilon_{\Gamma}, \quad (1)$$

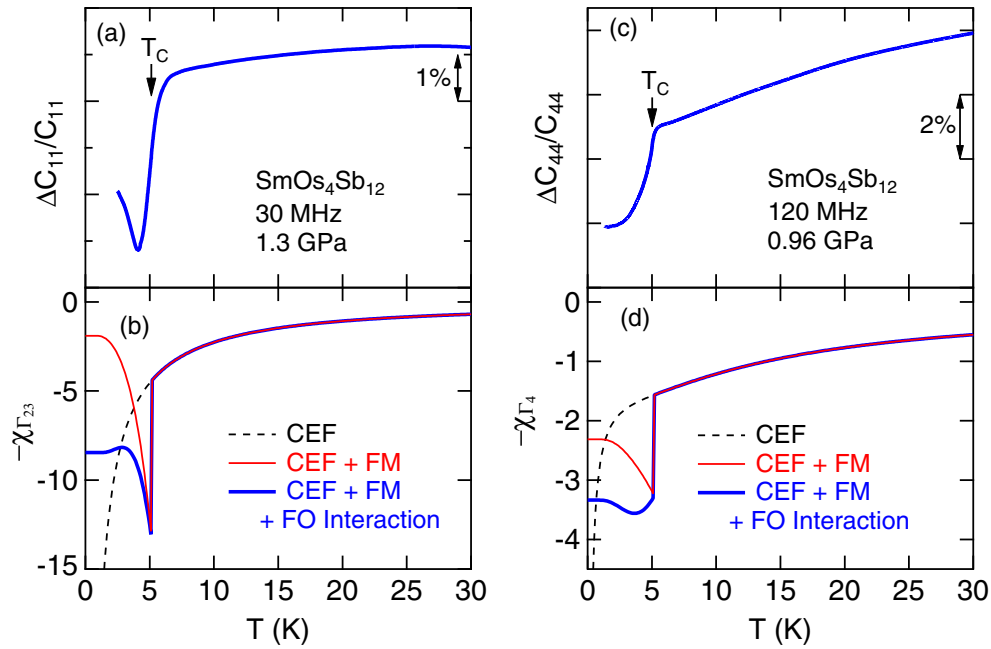


FIG. 4. (a) Temperature dependence of C_{11} , which corresponds to the response of the quadrupole O_2^0 with Γ_{23} symmetry, at 1.3 GPa, (b) quadrupolar susceptibility $-\chi_{\Gamma_{23}}$ calculated by applying a mean-field approximation (see text), (c) temperature dependence of C_{44} , which corresponds to the response of the quadrupole O_{yz} with Γ_4 symmetry, at 0.96 GPa, and (d) quadrupolar susceptibility $-\chi_{\Gamma_4}$ calculated by applying the mean-field approximation.

where the first term corresponds to the CEF effect and the coefficients K_{DD} , K_{OO} , and g_{Γ} are the magnetic dipole-dipole coupling constant, magnetic octupole-octupole coupling constant, and the strain-quadrupole coupling constant with symmetry $\Gamma = \Gamma_{23}$ or Γ_4 , respectively. In this calculation, we assume $K_{DD} = 2.2$ and $K_{OO} = 0.03$. Because the Γ_{67} -quartet ground state requires the magnetic easy axis for $\langle 100 \rangle$, we assume FM ordering along $\langle 100 \rangle$. In this calculation, we also consider three possible domain states by averaging over them. Figure 4 shows the experimental results obtained at the highest pressures and the calculated quadrupolar susceptibilities. The quadrupolar susceptibility $-\chi_{\Gamma_{23}}$ calculated by assuming primary FM ordering agrees qualitatively with the experimentally obtained C_{11} . However, the same calculation for the C_{44} mode gives a minimum C_{44} at T_C , and at low temperatures, C_{44} increases with decreasing temperature, which is inconsistent with the experimental result. As shown in Fig. 4, if a ferro-type octupole intersite interaction is included in the calculation, the calculated quadrupolar susceptibility reproduces the leveling-off feature of C_{44} at low temperatures.

Therefore, we conclude that the octupole-octupole interaction plays a role in the ordered state in this material. Considering the results of the specific heat and ac susceptibility under pressure [22], as discussed above, we strongly expect this interaction to become dominant and, at higher pressure, the ordered state to change into an octupolar ordered state.

B. Pressure dependence of the ultrasonic dispersions

Next, we describe how the UD depend on pressure. Figure 3(b) shows how C_{11} and the ultrasonic attenuation $\Delta\alpha$ depend on temperature at 1.3 GPa and at various frequencies. As can be seen, a clear UD also appears under high pressures.

The UD that appears at higher temperatures, which we call UD(high), shifts to lower temperatures upon increasing the pressure to 0.96 GPa. As seen in Ref. [27], the UD in ROs_4Sb_{12} systems appears at higher temperatures as the atomic number increases. In other words, as the ionic mass increases and the radius of the R ion decreases, the temperature at which the UD appears shifts toward higher temperatures, whereas the size of the atomic cage does not change in ROs_4Sb_{12} systems [30]. By applying pressure, the atomic cage gets smaller, and the relative size of the guest ion increases. Thus, we expect that, under pressure, the UD shifts to lower temperatures, as observed below 0.96 GPa. However, at 1.3 GPa, the elastic anomaly of UD(high) at 250 MHz is broader and appears at higher temperatures. This result cannot be explained on the basis of the analogy above and suggests a possible relationship with the other parameters, such as the recovery of the localized character and the change of the mixed-valence state. As previously noted, the temperature dependence of the averaged valence is reported to change upon applying pressure, and the temperature at which the averaged valence starts to decrease has a minimum around 1 GPa [28]. The ion size also strongly depends on the valence. Thus, UD(high) is affected by such a change in the valence state.

The UD that appears at lower temperatures, which we call UD(low), gradually disappears with increasing pressure. Interestingly, this UD(low) seems to disappear at the pressure where T^* drops below T_C . This result implies a possible relationship between the UD(low) and the c - f hybridization. To verify the veracity of this relationship, further ultrasonic measurements of C_{11} at high pressure and high magnetic fields are needed because a recovery of the localized character of 4*f* electrons was also indicated by our ultrasonic measurements above 20 T at ambient pressure [26].

Below T_C , we report in the insets in Figs. 3(a) and 3(b), the two frequency-dependent steplike elastic anomalies at high pressures, which are accompanied by sharp peaks in ultrasonic absorption. In contrast, C_{44} does not exhibit this type of anomaly in the region of pressure investigated in this work (i.e., these anomalies are mode selective, as are the UD). Thus, these anomalies seem to have the same origin as the UD. Assuming the resonance relation $\omega\tau \sim 1$ is satisfied at the temperature where the absorption peaks, the relaxation time τ between these two anomalies seems to diverge below T_C . This result suggests that some phonon modes freeze out in this range of temperature and pressure, which couples to Γ_1 and/or Γ_{23} strain waves. We also observed a similar dynamical elastic response at 0.96 and 0.65 GPa. Interestingly, this anomaly seems to appear under high pressure where the UD(low) disappears. These experimental results for the frequency-dependent and mode-selective elastic anomalies lead us to the following simple conclusion: the anharmonic potential of the guest Sm ion, which may be causing the UD(low), might change as the character of the $4f$ electrons changes from itinerant to localized upon applying pressure. The origin and quadripartite relationship between c - f hybridization, valence fluctuations, local charge fluctuations due to rattling, and multipolar interactions remain questions for discussion. Further measurements at higher pressure and over a wide range of temperature and frequency are required to elucidate these remaining issues.

V. CONCLUSIONS

In summary, we reported high-frequency and high-pressure ultrasonic measurements of the filled skutterudite $\text{SmOs}_4\text{Sb}_{12}$. We detected a signature of the crossover from the itinerant $4f$ -electron region to the localized $4f$ -electron region near 0.7 GPa. We also confirm the Γ_{67} CEF ground state and the presence of octupolar degrees of freedom of the $4f$ electrons in this material. We also measured how the UD due to rattling depend on ultrasonic frequency and pressures. The dependence of UD on pressure implies that a relationship exists between the local charge fluctuation due to off-center degrees of freedom regarding rattling and the exotic $4f$ -electronic states of this compound.

ACKNOWLEDGMENTS

This research was supported by JSPS KAKENHI Grants No. 26400342, No. 15K05882, and No. 15K21732 (J-Physics) and the Strategic Young Researcher Overseas Visits Program for Accelerating Brain Circulation from the Japan Society for the Promotion of Science and was also supported by the U.S. DOE (Grant No. DEFG02-04-ER46105) and the U.S. NSF (Grant No. DMR-1506677). We would like to thank S. Kasai (Coax Co. Ltd.) for his technical assistance with the ultrathin coaxial cable.

-
- [1] F. Steglich, J. Aarts, C. D. Bredl, W. Lieke, D. Meschede, W. Franz, and H. Schäfer, *Phys. Rev. Lett.* **43**, 1892 (1979).
 - [2] R. Shiina, H. Shiba, and P. Thalmeier, *J. Phys. Soc. Jpn.* **66**, 1741 (1997).
 - [3] Y. Ōnuki, R. Settai, K. Sugiyama, T. Takeuchi, T. C. Kobayashi, Y. Haga, and E. Yamamoto, *J. Phys. Soc. Jpn.* **73**, 769 (2004).
 - [4] S. Doniach, in *Valence Instabilities and Related Narrow Band Phenomena*, edited by R. D. Parks (Plenum, New York, 1977), p. 169.
 - [5] D. Jaccard, K. Behnia, and J. Sierro, *Phys. Lett. A* **163**, 475 (1992).
 - [6] E. Vargoz, P. Link, and D. Jaccard, *Phys. B (Amsterdam, Neth.)* **230-232**, 182 (1997).
 - [7] H. Hegger, C. Petrovic, E. G. Moshopoulou, M. F. Hundley, J. L. Sarrao, Z. Fisk, and J. D. Thompson, *Phys. Rev. Lett.* **84**, 4986 (2000).
 - [8] A. Sakai, K. Kuga, and S. Nakatsuji, *J. Phys. Soc. Jpn.* **80**, 063701 (2011).
 - [9] E. D. Bauer, N. A. Frederick, P.-C. Ho, V. S. Zapf, and M. B. Maple, *Phys. Rev. B* **65**, 100506(R) (2002).
 - [10] C. Sekine, T. Uchiumi, I. Shirotni, and T. Yagi, *Phys. Rev. Lett.* **79**, 3218 (1997).
 - [11] R. Higashinaka, T. Maruyama, A. Nakama, R. Miyazaki, Y. Aoki, and H. Sato, *J. Phys. Soc. Jpn.* **80**, 093703 (2011).
 - [12] A. Sakai and S. Nakatsuji, *Phys. Rev. B* **84**, 201106(R) (2011).
 - [13] W. M. Yuhasz, N. A. Frederick, P.-C. Ho, N. P. Butch, B. J. Taylor, T. A. Sayles, M. B. Maple, J. B. Betts, A. H. Lacerda, P. Rogl, and G. Giester, *Phys. Rev. B* **71**, 104402 (2005).
 - [14] S. Sanada, Y. Aoki, H. Aoki, A. Tsuchiya, D. Kikuchi, H. Sugawara, and H. Sato, *J. Phys. Soc. Jpn.* **74**, 246 (2005).
 - [15] G. R. Stewart, B. Andraka, C. Quitmann, B. Treadway, Y. Shapira, and E. J. McNiff, Jr., *Phys. Rev. B* **37**, 3344 (1988).
 - [16] H. Kotegawa, H. Hidaka, T. C. Kobayashi, D. Kikuchi, H. Sugawara, and H. Sato, *Phys. Rev. Lett.* **99**, 156408 (2007).
 - [17] M. Mizumaki, S. Tsutsui, H. Tanida, T. Uruga, D. Kikuchi, H. Sugawara, and H. Sato, *J. Phys. Soc. Jpn.* **76**, 053706 (2007).
 - [18] A. Yamasaki, S. Imada, H. Higashimichi, H. Fujiwara, T. Saita, T. Miyamachi, A. Sekiyama, H. Sugawara, D. Kikuchi, H. Sato, A. Higashiya, M. Yabashi, K. Tamasaku, D. Miwa, T. Ishikawa, and S. Suga, *Phys. Rev. Lett.* **98**, 156402 (2007).
 - [19] K. Fushiya, R. Miyazaki, R. Higashinaka, A. Yamada, M. Mizumaki, S. Tsutsui, K. Nitta, T. Uruga, B. Suemitsu, H. Sato, and Y. Aoki, *Phys. Rev. B* **92**, 075118 (2015).
 - [20] K. Hattori, Y. Hirayama, and K. Miyake, *J. Phys. Soc. Jpn.* **74**, 3306 (2005).
 - [21] H. Kotegawa, H. Hidaka, Y. Shimaoka, T. Miki, T. C. Kobayashi, D. Kikuchi, H. Sugawara, and H. Sato, *J. Phys. Soc. Jpn.* **74**, 2173 (2005).
 - [22] T. C. Kobayashi, K. Miyazawa, K. Tabira, Y. Ikeda, S. Araki, H. Kotegawa, H. Sugawara, D. Kikuchi, and H. Sato, *J. Phys. Soc. Jpn.* **80**, SA034 (2011).
 - [23] T. Goto and B. Lüthi, *Adv. Phys.* **52**, 67 (2003).
 - [24] M. Yoshizawa, Y. Nakanishi, M. Oikawa, C. Sekine, I. Shirotni, S. R. Saha, H. Sugawara, and H. Sato, *J. Phys. Soc. Jpn.* **74**, 2141 (2005).
 - [25] I. Ishii, H. Muneshige, Y. Suetomi, T. K. Fujita, T. Onimaru, K. T. Matsumoto, T. Takabatake, K. Araki, M. Akatsu, Y. Nemoto, T. Goto, and T. Suzuki, *J. Phys. Soc. Jpn.* **80**, 093601 (2011).

- [26] S. Mombetsu, T. Yanagisawa, H. Hidaka, H. Amitsuka, S. Yasin, S. Zherlitsyn, J. Wosnitzer, P.-C. Ho, and M. B. Maple, *J. Phys. Soc. Jpn.* **85**, 043704 (2016).
- [27] T. Yanagisawa, H. Saito, H. Hidaka, H. Amitsuka, K. Araki, M. Akatsu, Y. Nemoto, T. Goto, P.-C. Ho, R. E. Baumbach, and M. B. Maple, *J. Phys. Soc. Jpn.* **80**, 043601 (2011).
- [28] S. Tsutsui, N. Kawamura, M. Mizumaki, N. Ishimatsu, H. Maruyama, H. Sugawara, and H. Sato, *J. Phys. Soc. Jpn.* **82**, 023707 (2013).
- [29] H. Kotegawa, K. Tabira, Y. Irie, H. Hidaka, T. C. Kobayashi, D. Kikuchi, K. Tanaka, S. Tatsuoka, H. Sugawara, and H. Sato, *J. Phys. Soc. Jpn.* **77**(Suppl. A), 90 (2008).
- [30] D. J. Braun and W. Jeitschko, *J. Less Common Met.* **72**, 147 (1980).