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Instructions for use

Hybridization-driven orthorhombic lattice instability in URu₂Si₂

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We have measured the elastic constant $(C_{11}-C_{12})/2$ in URu₂Si₂ by means of high-frequency ultrasonic measurements in pulsed magnetic fields $H \parallel [001]$ up to 61.8 T in a wide temperature range from 1.5 to 116 K. We found a reduction of $(C_{11}-C_{12})/2$ that appears only in the temperature and magnetic field region in which URu₂Si₂ exhibits a heavy-electron state and hidden order. This change in $(C_{11}-C_{12})/2$ appears to be a response of the 5*f* electrons to an orthorhombic and volume conservative strain field $\epsilon_{xx}-\epsilon_{yy}$ with Γ_3 symmetry. This lattice instability is likely related to a symmetry-breaking band instability that arises due to the hybridization of the localized *f* electrons with the conduction electrons and is probably linked to the hidden-order parameter of this compound.

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The heavy-fermion compound URu₂Si₂ exhibits a secondorder phase transition involving uranium's 5 *f*-electron state at $T_o = 17.5$ K, and also exhibits unconventional superconductivity at $T_c \sim 1.4$ K. Though the transition at T_o shows clear anomalies in several thermodynamic quantities,¹⁻³ the order parameter has not been identified microscopically for more than a quarter of a century; thus, the unknown transition has been called "hidden order (HO)."⁴

In high magnetic fields (using static or pulsed magnetic field) for $H \parallel [001]$ at low temperatures, URu₂Si₂ undergoes three metamagnetic transitions in the range 35 and 39 T with a collapse of the HO phase, 5-7 where the *c*-axis magnetization increases in three steps and reaches a value of $\sim 1.5 \ \mu_{\rm B}/{\rm U}$ at 60 T and 1.5 K, which is approximately half of the value expected from the localized 5 f -electron state of U³⁺ (3.62 $\mu_{\rm B}$) or U⁴⁺ (3.58 $\mu_{\rm B}$). In addition, the electrical resistivity shows an abrupt decrease above 40 T. This suggests that tuning URu_2Si_2 by means of a magnetic field decreases the hybridization between 5f and conduction electrons, and, therefore, leads to a reduced effective electron mass.^{6,7} Given this situation, it is plausible that the hybridized-electron state observed at low-temperature and in low-magnetic-field regions will change to a relatively light Fermi-liquid state when a high magnetic field is applied along the [001] axis as indicated by the disappearance of the heavy band.⁷ Thus, the collapse of the HO and the drastic change of the 5f electronic state appear to be strongly connected and suggest that the order parameter of the HO phase is veiled in the screening of the localized 5 felectronic state via their strong hybridization with conduction electrons states.

In contrast, several recent theoretical models predict that the order parameter of the HO phase is of local nature and based on higher-rank electric and magnetic multipoles.^{8–10} In particular, the $xy(x^2-y^2)$ electric hexadecapole model, which has been proposed by Kusunose and Harima and also partly related to the theory of Haue and Kotliar,¹¹ suggests that electric quadrupole moments $O_{xy}(=J_x J_y + J_y J_x)$ and $O_2^2(=J_x^2 - J_y^2)$

are induced by an in-*c*-plane magnetic field or uniaxial stress via Ginzburg-Landau coupling.

Ultrasonic measurements are a powerful tool to probe the existence of such quadrupoles, as they induce a change of the elastic constants when either temperature or magnetic field are varied. Based on a model with localized f electrons, this is generally understood as the quadrupolar susceptibility. In contrast with the hexadecapol model, our recent ultrasonic measurements of the elastic constant $(C_{11}-C_{12})/2$ in pulsed magnetic fields applied parallel to [100] and [110] have demonstrated that there is no in-plane anisotropy up to 68.7 T.¹² While higher magnetic fields or higher measurement accuracy will be required to ultimately rule out the existence of the antiferro-hexadecapolar order, this suggests that the HO is not explained by this model unless otherwise considered possible effect of itinerant f electrons.

On the other hand, our results for $H \parallel [001]$ show that the change of the elastic constant $(C_{11}-C_{12})/2$ up to ~50 T is comparable to the elastic softening in the temperature dependence from 120 K to T_0 . This indicates that the lowtemperature electronic state of URu₂Si₂ exhibits a lattice instability with Γ_3 symmetry $(x^2-y^2$ -type) that is strongly related to the origin of the hybridized electronic state and the HO.¹²

In the present work, we explore a wider temperature range from 1.5 to 116 K up to 61.8 T with $H \parallel [001]$ for ultrasonic measurements on URu₂Si₂ in order to check the temperature dependence of the Γ_3 lattice instability in the high-magnetic-field region ($40 \leq H \leq 61.85$ T), where both the HO and hybridization effects are suppressed. Our results demonstrate clear evidence for an orthorhombic lattice instability that appears to be closely tied to the HO of URu₂Si₂.

A single crystal of URu_2Si_2 was grown at UC San Diego using the Czochralski method, and subsequently cut and mirror-like finishing, resulting in a length of 3.8024 mm along the [110] axis. Ultrasonic measurements were performed at the Hochfeld-Magnetlabor Dresden using the quadrature

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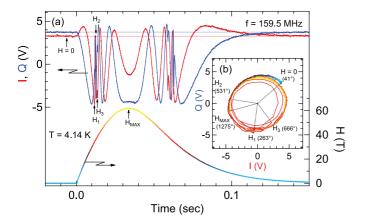


FIG. 1. (Color online) (a) Magnetic field (right axis) and output voltage of in-phase (I) and quadrature (Q) signals of the gated integrators at 4.14 K and 159.5 MHz (left axis) as a function of the elapsed time. Inset (b) shows the Lissajous curve of the I and Q signals.

procedure described in more detail in Ref. 12. For the present measurements, long-pulsed magnetic fields with a duration of \sim 150 ms and a maximum field of 61.85 T have been used. Long pulses offer the advantage that heating of the sample due to eddy currents, as well as adiabatic cooling and heating frequently observed in the vicinity of phase boundaries, when fast pulses are employed, are insignificant.

Figure 1 shows in-phase and quadrature signals of an ultrasonic wave with a frequency of 159.5 MHz for the present URu₂Si₂ sample as well as the magnitude of the pulsed magnetic field as a function of elapsed time. The arrow denoted by H_{MAX} indicates the time to reach the maximum field. In addition, H_1 , H_2 , and H_3 denote three phase boundaries observed in the high-magnetic-field phase diagram of URu₂Si₂.¹³ Notably, H_1 marks the HO transition, and H_2 and H_3 are the entry in and exit out of the so-called phase III,¹³ which is discussed as possible reentrant HO.⁵ The concentric Lissajous curve (shown in the inset of Fig. 1) demonstrates that the wave-detection system maintains good stability for the output levels and phase shift, and, in addition, that ultrasonic data obtained during magnetic-field sweeps represents true isotherms below 4.14 K.

Observed isotherms of the normalized elastic constant $(C_{11}-C_{12})/2$ are shown in Fig. 2 as a function of magnetic field $H \parallel$ [001]. Both curves for rising and falling magnetic fields are plotted and have been shifted vertically for clarity. The three phase transitions H_1 , H_2 , and H_3 discussed above are visible in our data as clear elastic anomalies for magnetic fields between 35 and 39 T and temperatures below 5.87 K as denoted by the upward facing triangles. Intriguingly, the anomaly H_1 at \sim 35 T and 1.56 K is marked by a sudden upturn, in contrast to the relatively small upturn with a precursor small dip observed at 4.14 and 5.87 K. We note that such differences are most likely due to the complex nature of the phase diagram in the vicinity of the upper phase boundary of the HO and related to the novel phases II or V reported in Refs. 5–7,13, and 14.

As apparent from Fig. 2, the magnetic field dependence of $(C_{11}-C_{12})/2$ in URu₂Si₂ looks very similar to magnetization versus *H* (applied along the *c*-axis)⁷ and thus is reminiscent

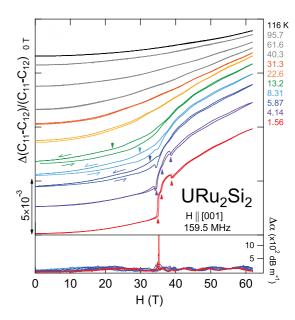


FIG. 2. (Color online) Upper panel: Elastic constant $(C_{11}-C_{12})/2$ normalized by the values at 0 T vs. the magnetic field $H \parallel [001]$ of URu₂Si₂ at various fixed temperatures between 1.56 and 116 K. Lower panel: Sound-attenuation change $\Delta \alpha$ vs. *H* for 1.56 to 13.2 K. All the data were taken with both up- and down-sweeps of the magnetic field, as indicated by arrows.

of magnetoelastic coupling. We can estimate the effect of the magnetostriction on $(C_{11}-C_{12})/2$ using the recently reported thermal-expansion data obtained in static magnetic fields up to 45 T.¹³ Using the Ehrenfest relation,¹⁵ and the pressure dependence of the *c*-axis at H_1 ,¹⁶ we find that the change of volume at H_1 is of the order of $\sim 10^{-4}$. The change of the [110] axis will be less or of the same order, and accordingly $(C_{11}-C_{12})/2$ will be affected with a factor of 2 in the phase comparative method. In turn, we conclude that the dramatic change of $\Delta(C_{11}-C_{12})/(C_{11}-C_{12})$ of 6×10^{-3} observed in our measurements near H_1 to H_3 is an order of magnitude larger than the estimated influence of the magnetostriction and, therefore, suggests the presence of an additional effect. For completeness, we note that our argument is only complete in the case where the CEF effect of the quadrupole moment O_2^2 is also negligible, e.g., not in the case of the Γ_5 doublet ground state. On the other hand, clear hystereses is observed below \sim 38 T for 5.87, 8.31, and 13.2 K. The hysteresis suggests that we do not have isothermal conditions, at least in this temperature range during up and down sweeps, probably due to strong magnetocaloric effects of this non-Fermi-liquid region relative to other temperature and magnetic field regions. Indeed, these regions may correspond to soft quantum fluctuations arising in the vicinity of the quantum critical end point associated with the destruction of the HO as suggested in Ref. 17.

Figure 3 represents the elastic constant change versus temperature at several fixed magnetic fields, which was obtained by converting isotherms of $(C_{11}-C_{12})/2$ recorded as a function of magnetic field (Fig. 2). Here, the data sets at zero magnetic field and 17.8 T were measured as function of temperature at constant fields using a standard phase-comparator method and served as a basis to extract the temperature dependence

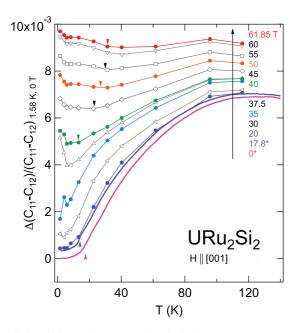


FIG. 3. (Color online) Normalized elastic constant $(C_{11}-C_{12})/2$ vs. temperature at various magnetic fields $H \parallel [001]$ from 0 to 61.85 T. The curves indicated by a magnetic field with an asterisk, 0 and 17.8 T, have been measured with the conventional method as a basis for the conversion from isotherms to curves at constant fields. Upward and downward facing triangles indicate the HO transition and local minimum, respectively.

of $(C_{11}-C_{12})/2$ from Fig. 2. The softening of 7×10^{-3} in the relative change of $(C_{11}-C_{12})/2$ below 120 K at zero magnetic field is gradually suppressed with increasing magnetic fields. Above 40 T, $(C_{11}-C_{12})/2$ shows a minimum, indicated by downward facing triangles, which shifts to higher temperatures with increasing magnetic field.

Figures 2 and 3 are compiled into a three-dimensional plot in Fig. 4. It is clearly seen that the softening of $(C_{11}-C_{12})/2$, i.e., the Γ_3 lattice instability, is enhanced in the red-colored region in and around the HO phase, where the strong hybridization is also developed. Conversely, the lattice instability disappears at high temperatures and high magnetic fields, where the HO collapses, as expected from previous work.⁷ Our present results obtained over a wide temperature and magnetic-field range, therefore, strongly support that the view that the hybridized electronic state entails the Γ_3 lattice instability.

The open (white) and closed (black) circles on the surface plot in Fig. 4 indicate the hidden-order transition defined by other experiments and local minima found in the temperature dependence of $(C_{11}-C_{12})/2$, respectively. The dotted curve and the white line are projections of the position of the local minima on the *H*-*T* phase diagram illustrated onto the basal plane of the plot. We note that the *H*-*T* phase diagram obtained in this work is similar to the phase diagram presented in Ref. 7 that was constructed via high-magnetic-field measurements of the electrical resistivity and magnetization.

Above 40 T, where the hybridization of f and conduction electrons is reduced, we are able to reproduce the temperature and magnetic field dependence of $(C_{11}-C_{12})/2$ employing the theory for a quadrupolar susceptibility based on a localized 5f electron with appropriate CEF scheme. By using the

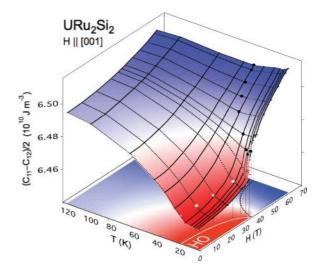


FIG. 4. (Color online) Three-dimensional plots of the elastic constant $(C_{11}-C_{12})/2$ vs. temperature and magnetic field along the *c* axis of URu₂Si₂. The white and black circles indicate the hidden-order transition and local minima found in the temperature dependence of $(C_{11}-C_{12})/2$, respectively. The bottom of the box shows the magnetic field-temperature phase diagram of URu₂Si₂ (See text for details).

following CEF scheme, $\Gamma_1^{(1)}$ - $\Gamma_2(60 \text{ K})$ - Γ_3 (178 K), which we have also employed in our previous work,¹² the features of $(C_{11}-C_{12})/2$ are qualitatively explained with the exception of the upturn below the local minimum. Since the position of the local minimum in the temperature dependence of the elastic constant (as shown in Fig. 3) may depend on multiple sources such as the gradient of the phonon-background, and the onset of the enhanced magnetization, the reason for this upturn is currently unclear. Further measurements in pulsed magnetic fields using different ultrasonic modes and magnetic-field directions will help to understand the details of the temperature dependence of $(C_{11}-C_{12})/2$ at the high-magnetic-field region. In particular, performing measurements on the nonmagnetic reference sample ThRu₂Si₂ to obtain the phonon background will facilitate this analysis.

Next, we consider the possible origin of the Γ_3 -lattice instability that we observe below 40 T. We expect this orthorhombic instability of URu₂Si₂ is caused by hybridization of the 5f electrons with the conduction electrons, since the non-5 f reference compound ThRu₂Si₂ does not show a softening in the $(C_{11}-C_{12})/2$ mode at zero magnetic field.¹⁸ In addition, the softening appears to be sensitive to chemically induced pressure and/or carrier doping as a small amount of Rh substitution (7%) significantly reduces the magnitude of the softening of $(C_{11}-C_{12})/2$ below 120 K.¹⁹ This suggests that for the magnetic-field region below 40 T, the so-called band Jahn-Teller (BJT) effect would be an explanation for the observed elastic response in contrast to the localized picture that we used at high magnetic fields. Here, the energy gain achieved through the formation of a hybridized band leads to this Jahn-Teller-type deformation.

The effect of the BJT distortion on the elastic constants was first discussed in the ultrasonic study of CeAg, LaAg, and their alloys with In.^{20,21} In the LnAg (Ln = La, Ce) compounds, the deformation-potential coupling effect was confirmed by

band-structure calculations and structural analysis of microscopic measurements. At present, microscopic measurements, such as ²⁹Si-NMR or high-precision x-ray and neutron scattering,^{22–24} provide no evidence for a structural symmetry breaking in URu₂Si₂. Since the elastic softening of URu₂Si₂ $(\sim 0.7\%)$ is much smaller compared to those of the BJT compounds (\sim 15–50%), we expect that a putative band deformation would be of a staggered type, instead of the uniform type observed in the BJT compounds.^{20,25} There is, however, no standard formulation of the BJT effect for such a staggered potential deformation. Such a Fermi-surface instability (i.e., evoked by Fermi-surface nesting along the *a* axis with $Q \neq 0$, predicted theoretically by Oppeneer et al.²⁷ and confirmed experimentally by Kawasaki et al.26) has an indirect effect on physical quantities at the Brillouin-zone center (Q = 0) for bulk measurements, in general. Thus, exotic effects such as a higher-order coupling or mode-mode-coupling of two different fluctuation mechanisms may need to be considered to interpret the present results.

We now consider differences between our result and the theoretical explanation of recent magnetic-torque measurements,²⁸ which have been interpreted as spontaneous fourfold rotational symmetry breaking in the tetragonal basal plane that takes place in the HO phase. Here, Thalmeier et al. and Ikeda *et al.* have proposed Γ_5^+ (E⁺)-type quadrupole and Γ_5^- (E⁻)-type dotriacontapole order parameters, respectively, to explain the torque-measurement results (here the sign of +and - indicate the parity of the time-reversal symmetry).^{10,29} Here, the elastic constant C_{44} corresponds to E^+ (Γ_5^+)symmetry and is expected to show characteristic softening above T_{o} and a relatively large anomaly at T_{o} compared to that of $(C_{11}-C_{12})/2$ with B_{2g} (Γ_3)-symmetry via electron-phonon interaction, even if the order parameter exhibits nematicity.³⁰ However, no distinct change of C_{44} has been observed at around T_0 ,³¹ except for a tiny kink, which could be caused by a thermal expansion effect on the sound velocity measurement, and thus, the Γ_3 lattice instability is highlighted. One simple interpretation for these discrepancies may be the anisotropic Jahn-Teller-type coupling between lattice and the electron systems, i.e., the electron-phonon coupling constant could be "extremely" weak for transverse modes but finite only for the $(C_{11}$ - C_{12} /2 mode. Otherwise, we have to simply conclude in agreement with NMR and x-ray diffraction^{22,24} that there is no lattice symmetry breaking of Γ_5 and Γ_4 symmetry in the HO phase.

It has also been pointed out that the formation of micronsize domains may wash out measurements of bulk properties in URu₂Si₂.²⁸ However, in case the continuum approximation is effective, that is when the ultrasonic wavelength in the solid sample is much longer than a force range due to the internal stresses, which occurs when a body is deformed by the elastic-wave propagation,³² and also the wavelength is shorter than the domain size or at least comparable with that, ultrasonic measurements will not be influenced by the existence of domains. For the present experiment this condition is fulfilled as the wavelength of the ultrasound waves is estimated $\lambda \sim 11$ –15 μm for $v \sim 2511 \text{ ms}^{-1}$ and f = 225– 159 MHz. Moreover, no unnecessary reflection or absorption of the ultrasound has been observed, and ultrasonic echoes have been well separated. On the other hand, a recent report of thermal-expansion measurements³³ suggests that multiple domains may orient to form an orthorhombic monodomain state when small in-plane uniaxial stress is applied. Therefore, the effects of uniaxial pressure on the elastic response will need to be probed to exclude any influence from domains entirely.

In conclusion, we have investigated the lattice instability existing within the HO and the associated hybridized electron state in URu₂Si₂. Our results demonstrate that the orthorhombic instability related to the electronic hybridization arising in the HO state x^2-y^2 -type (Γ_3) symmetry in stark contrast to the Γ_5 - or Γ_4 -type HO parameters that have been recently proposed on the basis of theory.

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- ¹T. T. M. Palstra, A. A. Menovsky, J. van den Berg, A. J. Dirkmaat, P. H. Kes, G. J. Nieuwenhuys, and J. A. Mydosh, Phys. Rev. Lett. **55**, 2727 (1985).
- ²M. B. Maple, J. W. Chen, Y. Dalichaouch, T. Kohara, C. Rossel, M. S. Torikachvili, M. W. McElfresh, and J. D. Thompson, Phys. Rev. Lett. **56**, 185 (1986).
- ³W. Schlabitz, J. Baumann, R. Pollit, U. Rauchschwalbe, H. M. Mayer, U. Ahlheim, and C. D. Bredl, Z. Phys. B **62**, 171 (1986).
- ⁴N. Shah, P. Chandra, P. Coleman, and J. A. Mydosh, Phys. Rev. B **61**, 564 (2000).

- ⁵N. Harrison, M. Jaime, and J. A. Mydosh, Phys. Rev. Lett. **90**, 096402 (2003).
- ⁶K. H. Kim, N. Harrison, M. Jaime, G. S. Boebinger, and J. A. Mydosh, Phys. Rev. Lett. **91**, 256401 (2003).
- ⁷G. W. Scheerer, W. Knafo, D. Aoki, G. Ballon, A. Mari, D. Vignolles, and J. Flouquet, Phys. Rev. B **85**, 094402 (2012).
- ⁸H. Harima, K. Miyake, and J. Flouquet, J. Phys. Soc. Jpn. **79**, 033705 (2010).
- ⁹H. Kusunose and H. Harima, J. Phys. Soc. Jpn. **80**, 084702 (2011).
- ¹⁰P. Thalmeier and T. Takimoto, Phys. Rev. B **83**, 165110 (2011).
- ¹¹K. Haule and G. Kotliar, Nat. Phys. 5, 796 (2009).

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- ¹²T. Yanagisawa, S. Mombetsu, H. Hidaka, H. Amitsuka, M. Akatsu, S. Yasin, S. Zherlitsyn, J. Wosnitza, K. Huang, and M. B. Maple, J. Phys. Soc. Jpn. 82, 013601 (2013).
- ¹³V. F. Correa, S. Francoual, M. Jaime, N. Harrison, T. P. Murphy, E. C. Palm, S. W. Tozer, A. H. Lacerda, P. A. Sharma, and J. A. Mydosh, Phys. Rev. Lett. **109**, 246405 (2012).
- ¹⁴A. Suslov, J. B. Ketterson, D. G. Hinks, D. F. Agterberg, and B. K. Sarma, Phys. Rev. B 68, 020406(R) (2003).
- ¹⁵W. Knafo, C. Meingast, K. Grube, S. Drobnik, P. Popovich, P. Schweiss, P. Adelmann, Th. Wolf, and H. v. Löhneysen, Phys. Rev. Lett. **99**, 137206 (2007).
- ¹⁶T. Inoue, K. Kindo, H. Okuni, K. Sugiyama, Y. Haga, E. Yamamoto, T. C. Kobayashi, Y. Uwatoko, and Y. Ōnuki, Physica B **294-295**, 271 (2001).
- ¹⁷M. Garst and A. Rosch, Phys. Rev. B 72, 205129 (2005).
- ¹⁸T. Yanagisawa, H. Saito, Y. Watanabe, Y. Shimizu, H. Hidaka, and H. Amitsuka, J. Phys. Conf. Ser. **391**, 012079 (2012).
- ¹⁹T. Yanagisawa (unpublished).
- ²⁰M. Niksch, B. Lüthi, and J. Kübler, Z. Phys. B **68**, 291 (1987).
- ²¹K. Knorr, B. Renker, W. Assmus, B. Lüthi, R. Takke, and H. J. Lauter, Z. Phys. B **39**, 151 (1980).
- ²²S. Takagi, S. Ishihara, S. Saitoh, H. Sasaki, H. Tanida, M. Yokoyama, and H. Amitsuka, J. Phys. Soc. Jpn. **76**, 033708 (2007).

- ²³H. Amitsuka, T. Inami, M. Yokoyama, S. Takayama, Y. Ikeda, I. Kawasaki, Y. Homma, H. Hidaka, and T. Yanagisawa, J. Phys.: Conf. Ser. **200**, 012007 (2010).
- ²⁴C. Tabata, private communications.
- ²⁵M. Yoshizawa, D. Kimura, T. Chiba, Simayi, Y. Nakanishi, K. Kihou, C.-H. Lee, A. Iyo, H. Eisaki, M. Nakajima, and S. Uchida, J. Phys. Soc. Jpn. **81**, 024604 (2012).
- ²⁶I. Kawasaki, S. I. Fujimori, Y. Takeda, T. Okane, A. Yasui, Y. Saitoh, H. Yamagami, Y. Haga, E. Yamamoto, and Y. Ōnuki, Phys. Rev. B 83, 235121 (2011).
- ²⁷P. M. Oppeneer, S. Elgazzar, J. Rusz, Q. Feng, T. Durakiewicz, and J. A. Mydosh, Phys. Rev. B 84, 241102 (2011).
- ²⁸R. Okazaki, T. Shibauchi, H. J. Shi, Y. Haga, T. D. Matsuda, E. Yamamoto, Y. Ōnuki, H. Ikeda, and Y. Matsuda, Science **331**, 439 (2011).
- ²⁹H. Ikeda, M. Suzuki, R. Arita, T. Takimoto, T. Shibauchi, and Y. Matsuda, Nature Phys. 8, 528 (2012).
- ³⁰S. Fujimoto, Phys. Rev. Lett. **106**, 196407 (2011).
- ³¹K. Kuwahara, H. Amitsuka, T. Sakakimara, O. Suzuki, S. Nakamura, T. Goto, M. Mahalik, A. A. Menovsky, A. de Visser, and J. J. M. Franse, J. Phys. Soc. Jpn. **66**, 3251 (1997).
- ³²B. Lüthi, *Physical Acoustics in the Solid State* (Springer, Berlin/Heiderberg/New York, 2005).
- ³³S. Kambe, D. Aoki, B. Salce, F. Bourdarot, D. Braithwaite, J. Flouquet, and J.-P. Brison, Phys. Rev. B **87**, 115123 (2013).