Relaxing Kondo-screened Kramers doublets in CeRhSi₃

J. Pásztorová,¹ A. Howell,¹ M. Songvilay,¹ P. M. Sarte,² J. A. Rodriguez-Rivera,^{3,4} A. M. Arévalo-López,⁵ K. Schmalzl,⁶

A. Schneidewind,^{7,8} S. R. Dunsiger,^{9,10} D. K. Singh,¹¹ C. Petrovic,¹² R. Hu,¹² and C. Stock¹

¹School of Physics and Astronomy, University of Edinburgh, Edinburgh EH9 3JZ, United Kingdom

²School of Chemistry, University of Edinburgh, Edinburgh EH9 3FJ, United Kingdom

³NIST Center for Neutron Research, National Institute of Standards and Technology, 100 Bureau Drive, Gaithersburg, Maryland, 20899, USA

⁴Department of Materials Science, University of Maryland, College Park, Maryland 20742, USA

⁵Université Lille, CNRS, Centrale Lille, ENSCL, Université Artois, UMR 8181-UCCS, F-59000 Lille, France

⁶Forschungszentrum Juelich GmbH, Juelich Centre for Neutron Science at ILL, 71 avenue des Martyrs, F-38000 Grenoble, France

⁷Forschungsneutronenquell Heinz Meier-Leibnitz (FRM-II), D-85747 Garching, Germany

⁸Institut für Festkörperphysik, TU Dresden, D-1062 Dresden, Germany

⁹Physics Department, James Franck Strasse 1, Technical University of Munich, D-85748 Garching, Germany

¹⁰Centre for Molecular and Materials Science, TRIUMF, Vancouver, British Columbia, Canada V6T 2A3

¹¹Department of Physics and Astronomy, University of Missouri, Missouri 65211, USA

¹²Condensed Matter Physics, Brookhaven National Laboratory, Upton, New York, 11973, USA

(Received 19 December 2018; revised manuscript received 26 February 2019; published 25 March 2019)

CeRhSi₃ is a superconductor under pressure coexisting with a weakly antiferromagnetic phase characterized by a Bragg peak at $\vec{q}_0 = (\sim 0.2, 0, 0.5)$ [N. Aso *et al.*, J. Magn. Magn. Mater. **310**, 602 (2007)]. The compound is also a heavy-fermion material with a large specific heat coefficient $\gamma = 110 \text{ mJ mol}^{-1} \text{ K}^{-2}$ and a high Kondo temperature of $T_{K} = 50$ K, indicating CeRhSi₃ is in a strongly Kondo screened state. We apply high-resolution neutron spectroscopy to investigate the magnetic fluctuations in the normal phase, at ambient pressures, and at low temperatures. We measure a commensurate dynamic response centered around the $\vec{Q} = (0, 0, 2)$ position that gradually evolves to $H \sim 0.2$ with decreasing temperature and/or energy transfers. The response is broadened both in momentum and energy and is not reminiscent of sharp spin wave excitations found in insulating magnets where the electrons are localized. We parametrize the excitation spectrum and temperature dependence using a heuristic model utilizing the random-phase approximation to couple relaxing Ce³⁺ groundstate Kramers doublets with a Kondo-like dynamic response. With a Ruderman-Kittel-Kasuya-Yosida exchange interaction within the *ab* plane and an increasing single-site susceptibility, we can qualitatively reproduce the neutron spectroscopic results in CeRhSi₃ and, namely, the trade-off between scattering at commensurate and incommensurate positions. We suggest that the antiferromagnetic phase in CeRhSi₃ is driven by weakly correlated relaxing localized Kramers doublets and that CeRhSi3 at ambient pressures is on the border between a Rudderman-Kittel-Yosida antiferromagnetic state and a Kondo-screened phase where static magnetism is predominately absent.

DOI: 10.1103/PhysRevB.99.125144

I. INTRODUCTION

There is a growing list of heavy-fermion-based materials that show a balance between unconventional superconductivity and localized magnetism [1,2]. In particular, several Ce^{3+} compounds, such as CeCoIn₅ ($T_c = 2.3$ K), display unconventional superconducting order parameters [3,4] similar to those of cuprate- or iron-based high temperature superconductors. However, in the case of the Ce^{3+} materials, the energy scale is much more amenable to experimental techniques such as thermal and cold neutron spectroscopy [5-7]. These materials therefore provide excellent model systems to understand the balance between competing electronic and magnetic phases [8,9]. We present neutron inelastic scattering data in the low-temperature ambient and nonsuperconducting phase of CeRhSi₃ [10], illustrating the competition between commensurate magnetism and incommensurate density wave fluctuations.

CeRhSi₃ is a noncentrosymmetric heavy fermion (space group number 107, I4mm) with a tetragonal unit cell with lattice parameters a = 4.269 Å and c = 9.738 Å [11]. Magnetic Ce³⁺ ions are located at the body center and the unit cell edges. The body center Ce^{3+} position is surrounded by a layer of Rh above and by a layer of Si below, therefore breaking inversion symmetry, making CeRhSi3 noncentrosymmetric [Fig. 1(a)]. CeRhSi₃ has a comparatively high electronic specific heat coefficient of $\gamma = 110 \text{ mJ mol}^{-1} \text{ K}^{-2}$ in comparison to, for example, antiferromagnetic CeCoGe₃ with $\gamma =$ 57 mJ mol⁻¹ K⁻² along with a large Kondo temperature of $T_K = 50$ K [12]. Putting these values together to estimate the Ruderman-Kittel-Kasuya-Yosida (RKKY) temperature [given by $k_B T_{\rm RKKY} \sim 3J^2 \gamma / \pi^2$ with J estimated from the specific heat and Kondo temperature with $J = -1/\ln(3T_K\gamma/\pi^2)$] [13] gives $T_{\rm RKKY} \sim 300$ K. While $T_{\rm RKKY} > T_K$ is indicative of local magnetism [14], the combination of γ and T_K is suggestive of a strongly Kondo screened phase by



FIG. 1. (a) The *I*4mm (No. 107) crystal structure of CeRhSi₃ and (b) the crystal field scheme discussed in the text using the Stevens parameters extracted by Muro *et al.* [32].

comparing $-1/\ln(3k_BT_K\gamma\pi^{-2})$ to other cerium-based heavy fermions [13]. Furthermore, the low-temperature phase has been studied by both band structure calculations and quantum oscillation measurements reporting large electron effective mass enhancement factors of $m^*/m_e \sim 8$ [15]. Possibly on the borderline between itinerant and localized magnetism, CeRhSi₃ is weakly antiferromagnetic with a small ordered magnetic moment of ~0.1 μ_B below temperatures of $T_N =$ 1.6 K and was previously characterized by an incommensurate wave vector of (~0.2, 0, 0.5) investigated with neutron elastic scattering [16].

Under pressures greater than ~12 kbar, CeRhSi₃ is found to be superconducting, with $T_c \sim 1$ K [12,17] and unusually large critical fields along the *c* axis [18–21]. The superconducting phase is more peculiar as muon spectroscopy finds antiferromagnetism to persist into this phase, being completely suppressed only at pressures of 23.6 kbar following a second-order transition indicative of a quantum critical point [22]. This is consistent with ac susceptibility measurements under pressure [23]. The nature of this critical point has been further explored by resistivity measurements under pressure and applied magnetic field [24] but remains unclear due to the lack of a high-pressure and high-field Fermi liquid phase. However, we note that recent penetration depth measurements report magnetism and superconductivity coexisting up to the largest pressures measured [25].

The noncentrosymmetric crystal structure combined with the strongly correlated electronic nature evidenced by heat capacity allows the possibility of novel unconventional electronic orders [26,27]. In centrosymmetric superconductors, spin-up and spin-down bands are degenerate in energy; however, when the crystal structure is noncentrosymmetric, spinorbit coupling can split these two bands, and the orbital and spin wave functions cannot be treated independently [28]. Magnetic fluctuations may therefore play a key role in superconductivity [29–31], and this has been further implicated by the transport and spectroscopic measurements outlined above, which seem to suggest that the Néel and superconducting temperatures are correlated.

We characterize the magnetic fluctuations in the lowtemperature Kondo-screened, normal state and at ambient

TABLE I. Stevens coefficients taken from Ref. [32].

Coefficient	Value			
B_{2}^{0}	- 0.151 meV			
B_4^0	0.0329 meV			
B_4^4	0.409 meV			

conditions in CeRhSi₃ by applying neutron inelastic scattering. This paper is divided into six sections, including this introduction and a final summary and conclusions component. We first discuss the single-ion crystal field theory to motivate our spectroscopic experiments. The experiments and materials are described in the next section, along with how we utilized the multidetector array on Multi Axis Crystal Spectrometer (MACS) to correct for the background. We then present our data illustrating the competing commensurate and incommensurate responses in CeRhSi₃ and discuss the results in terms of a heuristic model based on weakly coupled Kramers doublets.

II. SINGLE-ION CRYSTAL FIELD THEORY

As a starting point towards understanding the neutron scattering cross section characterizing the magnetic excitations and static order in CeRhSi₃, we review the crystal field theory results and also calculate matrix elements relevant for the discussion of the static and dynamic magnetism in this heavy-fermion compound. In the discussion Sec. V, we apply this analysis to investigate the possibility that the thermally isolated ground-state doublet can be considered in terms of projecting onto a $j_{\text{eff}} = 1/2$ angular momentum operator. For Ce³⁺ with j = 5/2 in a local $C_{4\nu}$ environment, as is the case for CeRhSi₃ in Fig. 1(a), the crystal field scheme should consist of three Kramers doublets. The corresponding Hamiltonian takes the following form:

$$\mathcal{H}_{CEF} = B_2^0 O_2^0 + B_4^0 O_4^0 + B_4^4 O_4^4, \tag{1}$$

where O_n^m are the Stevens operators based on angular momentum operators acting on the $|j, m\rangle$ basis and B_n^m are the Stevens parameters [33].

To obtain an idea of how much neutron scattering intensity should reside in the different crystal field doublets and in particular the intensity in the inelastic and elastic channels, we have taken the Stevens parameters extracted from susceptibility measurements given in Table I.

The resulting crystal field scheme is illustrated in Fig. 1(b) and is found to be in good agreement with the energy values of the crystal field excitations extracted from neutron inelastic scattering experiments. Substituting these coefficients into \mathcal{H}_{CEF} , we can obtain the eigenstates and calculate the matrix elements characterizing the elastic and inelastic neutron cross sections. We then obtain the following cross sections for neutron scattering exciting dynamics within the ground-state doublet (denoted as $I_{\text{inelastic}}$) and from the elastic cross section (I_{elastic}) [34,35]:

$$\begin{split} I_{\text{inelastic}} \propto \sum_{i=x,y} |\langle -|J^i| + \rangle|^2 \mu_B^2 &= 2.3 \mu_B^2 \\ I_{\text{elastic}} \propto g_J^2 |\langle 0|J^z|0\rangle|^2 \mu_B^2 \sim 10^{-3} \mu_B^2, \end{split}$$

with $g_J = 6/7$ for Ce³⁺. The above calculation suggests, based on the single-ion crystal field Hamiltonian, a large dynamic cross section in the neutron scattering response and comparatively little intensity in the elastic channel, a result consistent with initial powder work with neutrons [36]. A similar situation exists with YbRh₂Si₂ with a small ordered moment in comparison with the spectral weight in the inelastic channel [37]. The crystal field prediction of a low static ordered magnetic moment at low temperature is consistent with the small entropy gain measured from heat capacity (only $0.08R\ln 2$), which is one of the smallest values found for Ce³⁺-based materials [12]. The small static magnetic moment may therefore possibly be indicative of the underlying crystalline electric field rather than quantum criticality [38], which would mark the boundary between an itinerant phase and one where the mass of electrons diverges being characterized by more localized magnetism [14]. Such a critical point has also been suggested to host a Griffiths phase [39]. The lack of quantum criticality has been supported by transport measurements under pressure [40], and the presence of low-energy spin fluctuations has been implicated as the origin of the linear resistivity [29].

III. EXPERIMENTAL DETAILS

Having outlined the background and crystal field theory describing the magnetic excitations, we now discuss the experimental techniques and results obtained with neutron scattering.

A. Materials and scattering experiments

Material preparation and sample mounting. Single crystals of CeRhSi₃ were synthesized using a flux technique. Given the relatively small sample sizes for neutron scattering, an array of 2 g of single crystals was aligned on a series of aluminum plates as displayed in Figs. 2(c) and 2(d) with the rocking scan shown in Fig. 2(a), indicating a mosaic of 4°. The individual single crystals were secured to the plates using hydrogen-free Fomblin oil and were covered with aluminum foil as shown in Figs. 2(c) and 2(d). The aluminum plates were shaped in circles with two pins so that the sample could be aligned and swapped between the *H0L* and *HHL* scattering planes. Given the sample mounting, our sample was, on average, centrosymmetric as the mounting does not distinguish between $\pm c$ of the individual crystals.

Heat capacity measurements were performed on a (3.2 ± 0.5) -mg sample, using a physical property measurement system (Quantum Design) in a temperature range between 1.2 and 3 K. A relaxation method with a 2τ fitting procedure was used. The data displayed in Fig. 2(b) show a transition near 2 K, as expected from magnetic ordering.

Neutron spectroscopy. Attempts to search for a static magnetic Bragg peak using the D23 diffractometer [Institut Laue-Langevin (ILL), France] were not successful in resolving a temperature-dependent signal from the background. This may be expected based on the crystal field analysis discussed above, which suggests a comparatively weak static moment in comparison to the inelastic scattering. We therefore focused our measurements on studying the dynamics with spectroscopy.



FIG. 2. (a) The rocking curve through the (002) Bragg peak with a full width at half maximum of 4.0° at room temperature. (b) Heat capacity data as a function of temperature for CeRhSi₃, confirming a transition at ~1.8 K. (c) and (d) The sample mount used for the neutron inelastic scattering studies on CeRhSi₃ discussed in the text.

Initial triple-axis measurements were carried out on the PANDA spectrometer (FRM2), where it was established that the magnetic scattering was highly extended in momentum space. This was further confirmed by experiments on the Spin Polarized Inelastic Neutron Spectrometer (NIST). Given the need for measurements of a broad range in momentum space, the MACS cold triple-axis spectrometer (NIST) was utilized. Cooled filters of beryllium and beryllium oxide were placed before the monochromator and after the sample, respectively. With a fixed $E_f = 3.7$ meV, this configuration afforded energy transfers up to 1.3 meV. The use of a double-filter configuration was found to be necessary to keep higher-energy neutrons from scattering off the monochromator onto the sample. In such a situation, without the first beryllium filter, it was found that the sample mount and Fomblin oil gave a large background, making extraction of the magnetic signal difficult and unreliable.

Motivated by the observation of a commensurate response which may be indicative of ferromagnetic interactions, we measured the neutron inelastic response under a magnetic field. We note that in YbRh₂Si₂, the incommensurate response was found to be strongly affected by an applied field [7]. Measurements with a magnetic field were done using the IN12 cold triple-axis spectrometer at the ILL with $E_f = 3.5$ meV and using a 6-T cryomagnet with a ³He insert so that temperatures as low as 0.5 K could be reached.

B. Background subtraction on MACS

Given our complex sample-mounting scheme shown in Fig. 2, background scattering of neutrons from the Fomblin



FIG. 3. The method used to determine the background. (a) The smoothed intensity. (b) The remaining data after removing the main magnetic signal, Bragg peaks, and central beam. (c) The background signal generated by finding the radial average of (b). (d) The magnetic signal found by removing the background in (c) from the smoothed data in (a).

grease and also the aluminum sample holder was an issue. We outline here how the magnetic inelastic signal was isolated from the background using the detector arrangement on the MACS cold triple-axis spectrometer (NIST). The wide coverage of MACS allowed for a simultaneous measurement of both the magnetic inelastic signal and also a large background region where a comparatively strongly temperature dependent magnetic signal was not observed.

The methodology used to calculate the background for any given temperature and energy is illustrated in Fig. 3. Figure 3(a) shows the smoothed intensity including background and magnetic scattering, adding equivalent data at $\pm L$ positions. The intensity consists of a main magnetic region surrounded on both sides by a weaker scattering signal independent of the sample rotation angle, which appears as a ring of intensity in $|\vec{Q}|$. This background also includes Bragg peaks that occur near the (1, 0, 1) and (-1, 0, 1) positions. The relative strength of this background varied with temperature and energy. Therefore, the background needed to be approximated at each temperature and energy.

To subtract this background, for a given temperature and also energy transfer, we removed strips that contained the magnetic signal and the Bragg peaks, as shown in Fig. 3(b). The remaining data were then used to determine a radially averaged background, as shown in Fig. 3(c). This averaging did not consider small-angle scattering from the main beam indicated by the bright region about the origin in Fig. 3. Subtracting Fig. 3(c) from Fig. 3(a) returns the magnetic inelastic signal shown in Fig. 3(d). A similar background subtraction procedure was applied in a recent study on MACS investigating CeCo($In_{1-x}Hg_x$)₅ (x = 0.01) [41].

IV. EXPERIMENTAL RESULTS

Having discussed the single-ion properties of CeRhSi3 and the experimental details, we discuss the results obtained for the dynamic neutron response. We first show the temperature dependence of the correlated low-energy magnetic scattering. Motivated by previous neutron scattering reports of a weak, low-temperature magnetic Bragg peak at ($\sim 0.2, 0, 0.5$) [16], we focused our measurements on the (H, 0, L) scattering plane. Figure 4 shows the temperature dependence of the correlated magnetic scattering near $\vec{Q} = (0, 0, 2)$ taken on MACS with the sample oriented in the (H0L) plane and with the background subtracted following the methodology discussed above. Figures 4(a)-4(c) illustrate low-temperature constant E = 0.5 meV slices at T = 0.3, 6, and 10 K. The results show enhanced but momentum-broadened scattering near $\vec{Q} = (0, 0, 2)$ for these three temperatures. We note that the commensurate scattering at $\vec{Q} = (0, 0, 2)$ is strongly contaminated by nuclear elastic scattering given that (0,0,2) is an allowed crystallographic Bragg peak. The



FIG. 4. Constant E = 0.5 meV slices taken on MACS at (a) T = 0.3 K, (b) 6 K, and (c) 10 K. Slices along the $(H, 0, 1.5 \pm 0.2)$ direction are shown in (d)–(f). The lines are fits to Gaussians, with (d) illustrating two symmetrically displaced peaks and (e) showing a fit to two peaks and also a single peak. At high temperatures of 10 K in (f), the correlated scattering is well described by a single peak centered at the H = 0 position.

background-corrected scattering near (0,0,2) in Fig. 4(a) consists of both a sharp component originating from scattering from the nuclear peak and also scattering elongated along the *L* direction and extending over the entire Brillouin zone. The momentum-broadened rods of scattering are elongated along *L*, indicative of weak correlations along *c*. With increasing temperature, shown in Figs. 4(b) and 4(c), the scattering decreases in intensity, confirming the magnetic origin.

Figures 4(d)-4(f) illustrate cuts through these rods of scattering near $\vec{Q} = (0, 0, 2)$ integrating along *L* in the $(H, 0, 1.5 \pm 0.2)$ direction. At the low temperatures (T = 0.3 K) shown in Fig. 4(d), two clear peaks can be seen to be symmetrically displaced along *H*, located at $H_0 = 0.14 \pm 0.03$. On heating, the structure disappears until 10 K, where the scattering is well described by a commensurate peak centered at H = 0. The low-energy magnetic scattering in CeRhSi₃ therefore crosses over from incommensurate scattering along *H* to a commensurate response with increasing temperature.

Having established the presence of momentum-broadened correlations along L which weaken with increasing temperature as expected for magnetic scattering, we now show the energy dependence at low temperatures. Figure 5 shows a series of constant-energy slices taken at T = 0.5 K using the MACS spectrometer. At low-energy transfers of E = 0.5 meV [Fig. 5(a)], extended scattering along L which peaks near $H = 0.14 \pm 0.03$ real lattice unit (r.l.u.) is found. However, as shown in Figs. 5(b) and 5(c), on increasing the energy



FIG. 5. T = 0.5 K constant-energy cuts at (a) E = 0.5 meV, (b) 0.75 meV, and (c) 1.0 meV. The correlated magnetic scattering weakens and also shifts spectral weight to the commensurate position with increasing energy transfer.

transfer, the scattering becomes weaker and also centered around the commensurate $\vec{Q} = (0, 0, 2)$ position. For energy transfers at E = 1.0 meV, shown in Fig. 5(c), the scattering remains highly extended in momentum yet more centered at the commensurate position. The magnetic scattering therefore crosses over from an incommensurate response to a commensurate response through increasing either the energy transfer or temperature.

The experimental data were primarily taken in the (H, 0, L) plane, motivated by the discovery of a weak magnetic Bragg peak in this scattering plane. However, in most tetragonal heavy-fermion materials, like the Ce(Rh,Co)In₅ family, the scattering is located near the antiferromagnetic (0.5, 0, 5) position [7,37,42]. In Fig. 6, the neutron responses in both



FIG. 6. A comparison of the magnetic scattering in the (a)–(c) (H, 0, L) and (d)–(f) (H, H, L) scattering planes.

the (H, 0, L) and (H, H, L) scattering planes are compared for a series of energy transfers. The scattering in both planes is qualitatively similar, with the magnetic scattering being highly extended along the L direction. At low temperatures and energy transfers, in the (H, H, L) plane, as illustrated in Fig. 6(d), the scattering is also suggestive of being extended along the $[1, \overline{1}, 0]$ direction. With increasing energy transfer, the scattering broadens and weakens considerably near the (0,0,2) position in both scattering planes.

The presence of magnetic scattering near the commensurate (002) position is suggestive of an underlying ferromagnetic response as observed in YbRh₂Si₂ [37]. While ferromagnetism in Ce-based compounds is rare, it has been reported in CeRuPO [43] and also CeSb₂ [44]. However, given the body-centered nature of the crystallographic structure, it is also consistent with an antiferromagnetic interaction between nearest Ce^{3+} neighbors along c. To test for this hypothesis, we have applied a vertical magnetic field in order to check for a strong response of the magnetic fluctuations. These experiments were performed on IN12 and are summarized in Fig. 7, which compares a constant (H, 0, 1.5) scan at 0 and 6 T at low temperatures. While the data are statistics limited, we do not observe a strong or significant response of magnetic fluctuations to a magnetic field aligned within the *a-b* plane. As an additional check for a possible ferromagnetic response, the elastic (0,0,2) nuclear Bragg peak was also studied as a function of magnetic field, with no measurable change observed. Based on this and the lack of a strong change with applied field in the inelastic channel, we conclude that the underlying interaction is dominated by antiferromagnetic interactions.



FIG. 7. Constant E = 0.5 meV scans at 0 and 6 T taken on the IN12 spectrometer (ILL) with $E_f = 3.5$ meV. A temperatureindependent background has been subtracted from the data. No strong or significant response of the magnetic fluctuations to field is observed.

V. HEURISTIC DESCRIPTION

Having presented the experimental results, we now attempt to understand the interplay between the commensurate magnetic fluctuations at high temperatures and energies and the crossover at low energies and temperatures to an incommensurate response. We first discuss the results in terms of overdamped $j_{eff} = 1/2$ spin waves and show that the local crystal field symmetry is not consistent with such a description. We then discuss the fluctuations in terms of weakly coupled Kramers doublets through the use of the randomphase approximation (RPA). We find that this latter approach allows us to reproduce, qualitatively, the interplay between commensurate and incommensurate responses.

A. $j_{\text{eff}} = 1/2$ ground state and excitations?

The crystal field scheme in Fig. 1 shows that the groundstate doublet is separated from the next highest energy doublet by ~19 meV = 220 K. Given the energy scale separating these two excitations, at low temperatures we would expect this doublet to be well separated from the higher energy and then to possibly behave as a $j_{eff} = 1/2$ magnet similar to the spin-orbit split levels in Co²⁺ [45] or 4*d* or 5*d* transition-metal ions [45–50]. In this section, we investigate this point by studying the lowest-energy doublets and the angular momentum operators in this two-dimensional subspace.

To investigate this point, we have transformed the coordinates of the angular momentum operators from a $|j, m\rangle$ basis to a basis corresponding to the eigenstates of the crystal field Hamiltonian \mathcal{H}_{CEF} (denoted as the $|CEF\rangle$ basis). This corresponds to writing a transformation matrix with the columns

the eigenstates in the $|j, m\rangle$ basis as follows:

$$\mathcal{C} = \begin{bmatrix} 0.605 & 0 & 0 & 0 & 0.796 & 0 \\ 0 & 0.796 & 0 & 0 & 0 & -0.605 \\ 0 & 0 & 1.000 & 0 & 0 \\ 0 & 0 & 0 & 1.000 & 0 & 0 \\ -0.796 & 0 & 0 & 0 & 0.605 & 0 \\ 0 & -0.605 & 0 & 0 & 0 & -0.796 \end{bmatrix}$$

Using this matrix, we can then transform any operator (denoted as A) from the $|j, m\rangle$ to the $|CEF\rangle$ basis and vice versa using

$$A_{|CEF\rangle} = \mathcal{C}^{-1} A_{|j,m\rangle} \mathcal{C}.$$
 (2)

 \mathcal{H}_{CEF} after this transformation takes the following diagonal form, with the elements being the energy eigenvalues:

				$\mathcal{H}^{CEF}_{ CEF}$	$\langle \sigma_{\gamma} \rangle = C^{-1}$	${}^{1}\mathcal{H}^{CEF}_{ j,m\rangle}\mathcal{C}$,
	[−13.97	0	0	0	0	ך 0	
=	0	-13.97	0	0	0	0	
	0	0	5.16	0	0	0	
	0	0	0	5.16	0	0	•
	0	0	0	0	8.82	0	
	0	0	0	0	0	8.82	

Using the transformation C, we can transform the angular momentum operators between the different bases, for example, with J^z taking the following form:

				$J^z_{ C }$	$\langle EF \rangle = C^{2}$	$^{-1}J^{z}_{ j,m\rangle}\mathcal{C} =$:
Γ	- 0.034	0	0	0	1.928	ך 0	
	0	0.034	0	0	0	-1.928	
	0	0	0.5	0	0	0	
	0	0	0	-0.5	0	0	•
	1.928	0	0	0	1.034	0	
L	0	-1.928	0	0	0	-1.034	

To determine if the ground-state doublet can be projected onto a $j_{\text{eff}} = 1/2$ angular momentum operator, we consider the upper 2 × 2 matrix (denoted in bold) which operates on this subspace. Calculating this for the \tilde{A}^x , \tilde{A}^y , and \tilde{A}^z operators we get

$$\tilde{A}^{x} = \begin{bmatrix} 0 & 1.078\\ 1.078 & 0 \end{bmatrix},$$
$$\tilde{A}^{y} = \begin{bmatrix} 0 & -1.078i\\ 1.078i & 0 \end{bmatrix},$$
$$\tilde{A}^{z} = \begin{bmatrix} -0.034 & 0\\ 0 & 0.034 \end{bmatrix},$$

These 2 × 2 matrices do not follow the canonical commutation relations summarized by $\vec{J} \times \vec{J} = i\vec{J}$ as satisfied by the Pauli spin matrices which belong to the SU(2) group. We therefore conclude that the ground-state crystal field doublet of CeRhSi₃, while isolated at low temperatures from higherenergy crystal field doublets, cannot be projected onto a $j_{\text{eff}} = 1/2$ angular momentum operator. Therefore, $\tilde{A} \neq \alpha J$, where J is an angular momentum operator and α is a scaler projection factor. From a symmetry perspective, this result is not surprising given the highly anisotropic crystalline electric field. We will discuss this result below in parametrizing the low-energy and temperature-dependent spin fluctuations in CeRhSi₃.

B. Correlated relaxing Kondo sites

The magnetic scattering illustrated in Figs. 4 and 5 is highly extended in both momentum and energy, displaying little $\vec{Q} - \omega$ structure like that reported in other more localized Ce³⁺-based systems such as CeRhIn₅ [42,51] or discussed in the context of temporally sharp spin excitations in CeCoIn₅ [52]. Therefore, parametrization of the excitations involving crystal fields or well-defined harmonic magnons is not appropriate to describe the low-energy magnetic dynamics in CeRhSi₃ at ambient pressures. Furthermore, the crystal field analysis discussed above illustrates that the ground state cannot be projected onto a $j_{eff} = \frac{1}{2}$ ground state, and therefore, damped spin wave theory involving Heisenberg coupling of $j_{\rm eff} = \frac{1}{2}$ spins cannot be applied. Any heuristic model must be able to describe the temperature and energy dependence of the magnetic excitations discussed in the previous sections. In particular, the results outlined above indicate two competing effects, with one described by a commensurate wave vector near $\vec{Q} = (0, 0, 2)$ and another associated with the low-temperature incommensurate wave order appearing near $H \sim 0.2$. In this section, we investigate whether a weakly correlated lattice of Kondo sites can consistently describe the commensurate scattering and the extended nature of it in momentum. We emphasize that this model is not unique, and similar incommensurate to commensurate magnetic responses have been observed in a number of materials, notably in, for example, the cuprates and pnictides with models based on band structure or stripes [53,54].

The magnetic neutron cross section is proportional to $S(\vec{Q}, \omega)$, which is related to the susceptibility by

$$S(\vec{Q},\omega) \propto [n(\omega)+1]\chi''(\vec{Q},\omega),$$

where $[n(\omega) + 1]$ is the Bose factor and $\chi''(\vec{Q}, \omega)$ is the imaginary part of the susceptibility. To model the energy-broadened commensurate component, we note that the extended nature of the neutron cross section in momentum indicates weak correlations between Ce³⁺ sites. We therefore consider a single-site susceptibility describing fluctuations within the ground state of the localized Kramers doublet with a characteristic energy scale Γ set by the relaxational timescale τ via $\Gamma \sim \frac{1}{\tau}$. This results in a single-site susceptibility given by

$$\chi_0(\omega) = \frac{X_0\Gamma}{\Gamma - i\omega},$$

where X_0 is the temperature-dependent single-site static susceptibility. Similar single-site susceptibilities have been used to describe the paramagnetic scattering in uranium-based heavy fermions [55–57] and also in the paramagnetic phase

of the cuprates [58,59]. In this case, given that the dominant single-ion cross section discussed above is for transitions between members of the ground-state doublet, this susceptibility can be interpreted physically as transitions between the members of the lowest-energy Kramers doublet with a timescale set by $\tau \propto \frac{1}{\Gamma}$. We therefore call this single-ion response a relaxing Kramers doublet.

To parametrize the coupling of these relaxing Kramers doublets on different Ce^{3+} sites, we apply the RPA, which gives a final susceptibility with the following form:

$$\chi(\vec{Q},\omega) = \frac{\chi_0(\omega)}{1 - \chi_0(\omega)J(\vec{Q})}$$

where $J(\vec{Q})$ is the Fourier transform of the coupling between the localized magnetic sites, each consisting of a low-energy two-level energy scheme. We consider the following heuristic form for a two-dimensional magnet with only nearestneighbor coupling along the *a* and *b* axes for simplicity:

$$J(\vec{Q}) = \sum_{r} J_{r} e^{i\vec{Q}\cdot\vec{r}} = 2J_{\rm RKKY}[\cos(2\pi H) + \cos(2\pi K)].$$
 (3)

Putting this all together to calculate the imaginary part of the susceptibility proportional to the neutron cross section gives

$$\chi''(\vec{Q},\omega) = \frac{X_0 \Gamma \omega}{\omega^2 + \Gamma^2 [1 - X_0 J(\vec{Q})]^2}.$$
 (4)

This relation highlights the fact that the cross section is peaked in momentum when $1 - X_0 J(\vec{Q})$ is a minimum. We note that this form of χ'' is an odd function in energy required for detailed balance [60]. Given the scattering is confined along the (0, 0, L) direction and extended along L, indicative of short-range correlations, we have considered the cross section from in-plane fluctuations and weakly correlated Ce³⁺ spins along c. The neutron cross section can therefore be written as

$$S(\vec{Q},\omega) \propto [n(\omega)+1]\chi''(\vec{Q},\omega) \left[1 - \left(\frac{Q_{ab}}{Q}\right)^2\right] \cdots \times [1 + \alpha \cos(\vec{Q} \cdot \vec{c})].$$
(5)

The parameter α denotes the strength of nearest-neighbor correlations along *c*. We note the distinction between coupling, described by $J(\vec{Q})$, and correlations parameterized by the susceptibility and α .

Figure 8(a) illustrates the X_0 and J phase diagram, with the colors indicating where in H the term in the susceptibility $[1 - X_0 J(\vec{Q})]$ is a minimum and therefore where the neutron scattering cross section is maximum. The value of Γ was chosen to be 0.5 meV to match the energy range where strong magnetic fluctuations are observed on MACS. The plot is done for X_0 and J and shows that for small values of the susceptibility X_0 , a maximum in the scattering cross section occurs at the commensurate value $H \sim 0$, where for larger values of either X_0 or J, the cross section is maximum at H = 0.25 when $2\pi H = 1/2$. This is further illustrated in Figs. 8(b) and 8(c), which plot constant-energy slices for the two extreme cases. Figure 8(b) shows the scattering cross section for the case of both large susceptibility X_0 and also large correlations along c. The intensity profiles display peaks displaced along H near H = 0.25 and also L = 0.5. When the



FIG. 8. (a) The location in *H* of the maximum scattering intensity as a function of X_0 and exchange constant *J* between different Ce³⁺ ions. (b) A simulated constant E = 0.5 meV slice taken with large X_0 and interlayer correlations α . (c) A simulated constantenergy slice with weak X_0 and interlayer correlations. The calculations were done with $\Gamma = 0.5$ meV.

susceptibility X_0 is reduced, the cross section becomes peaked around the commensurate position, as illustrated in Fig. 8(c).

This simple and heuristic model of correlated and relaxing localized Kramers doublets captures the main qualitative results of our experiments. The calculations with $\Gamma = 0.5$ meV provide an estimate for the exchange constant of ~1 meV (assuming $X_0 \sim 1\mu_B^2/\text{meV}$), which should be compared with an estimate of ~10 meV extracted from the Kondo temperature and specific heat coefficient. Our choice of X_0 is consistent with Q = 0 susceptibility measurements, which indicate a low-temperature susceptibility of the order of ~10⁻² emu/mol [32]. Our calculated value is sensitive to the choice of Γ and also X_0 but, nevertheless, is in reasonable agreement with the value estimated based on Ref. [13].

The experimental results shown above exhibit a trade-off in intensity between incommensurate magnetic scattering peaking near (~0.2,0,1.5) and also the commensurate (0,0,2) position. The maximum in the cross section shifts with increasing temperature or energy transfer. In both cases, our model would imply that such a change in either energy or temperature coincides with a reduction in susceptibility X_0 . We note that neutron diffraction results have reported a Bragg peak near $\vec{q_0} = (\sim 0.22, 0, 0.5)$. The displacement of the wave vector from the 0.25 position along *H* may be accounted for by incorporating farther-neighbor exchange constants, resulting in a more complex form of $J(\vec{Q})$ discussed above.

VI. CONCLUSIONS AND DISCUSSION

The momentum- and energy-broadened response reported here for CeRhSi₃ differs from dispersing spin waves measured in insulating magnets and also metallic compounds with strongly localized moments. In this context CeRhSi₃ differs from other noncentrosymmetric systems like CePt₃Si, which displays momentum dispersing spin wave excitations, albeit damped [61]. The energy- and momentum-broadened spin response at low energies also contrasts with the localized systems with the same crystal structures, such as CeCoGe₃ [62] and CeIrGe₃ [63], and is suggestive of highly itinerant *f* electrons. This latter point is consistent with quantum oscillation measurements, which fail to measure a divergence in effective mass in the magnetic phase as a function of pressure, which would indicate a crossover to strongly localized moments [64].

The results presented here also differ from muon results which suggest static magnetism on the megahertz timescale. We observe relaxational dynamics with a frequency scale on the order of terahertz, and this may, indeed, be outside the time window of muons. The pressure dependence of the magnetic ordering [22] has been interpreted in terms of the Doniach phase diagram [13,65], which captures the competition between RKKY-driven antiferromagnetism and a Kondo-screened state [66-68] separated by a critical exchange constant value. Our results point to CeRhSi3 being on the borderline between these different phases, with relaxational dynamics reminiscent of a screened antiferromagnetic phase without long-range order. However, our results are consistent with a weak RKKY coupling between sites and also a growing susceptibility which drives the incommensurate order. This might point to the existence of well-defined spin excitations at low energies in more localized variants such

PHYSICAL REVIEW B 99, 125144 (2019)

as CeCoGe₃, where much higher pressures (~4 GPa) are required to achieve superconductivity [69] and where thermodynamic measurements suggest a weaker Kondo effect than in CeRhSi₃ [11]. We would expect a decrease in J_{RKKY} with pressure as Kondo screening becomes more prevalent. This is consistent with thermoelectric measurements under pressure [70] and also resistivity on CeRhSi₃ [71].

In summary, we have measured the low-energy magnetic response in CeRhSi₃ at ambient pressures and at low temperatures. A momentum- and energy-broadened response was observed which crosses over from commensurate at high energies to an incommensurate wave vector along H at low-energy transfers and low temperatures. The response is not consistent with dispersing or damped spin excitations as observed in other Ce-based non-centrosymmetric superconductors. We found a heuristic model based on weakly correlated relaxing and strongly Kondo screened localized Kramers doublets qualitatively accounts for this behavior. We suggest that CeRhSi₃ is on the border between a Rudderman-Kittel-Yosida antiferromagnetic phase with strong and underdamped magnetic correlations and a fully Kondo screened phase where no correlations exist.

ACKNOWLEDGMENTS

The work here was funded by the EPSRC, the Carnegie Trust for the Universities of Scotland, and the STFC. Access to MACS was provided by the Center for High Resolution Neutron Scattering, a partnership between the National Institute of Standards and Technology and the National Science Foundation under Agreement No. DMR-1508249. Work at Brookhaven National Laboratory was supported by US DOE, Office of Science, Office of Basic Energy Sciences (DOE-BES), under Contract No. DE-SC0012704

- [1] G. R. Stewart, Rev. Mod. Phys. 56, 755 (1984).
- [2] C. Pfleiderer, Rev. Mod. Phys. 81, 1551 (2009).
- [3] K. Izawa, H. Yamaguchi, Y. Matsuda, H. Shishido, R. Settai, and Y. Onuki, Phys. Rev. Lett. 87, 057002 (2001).
- [4] N. D. Mathur, F. M. Grosche, S. R. Julian, I. R. Walker, D. M. Freye, R. K. W. Haselwimmer, and G. G. Lonzarich, Nature (London) **394**, 39 (1998).
- [5] O. Stockert, J. Arndt, E. Faulhaber, C. Geibel, H. S. Jeevan, S. Kirchner, M. Loewenhaupt, K. Schmalzl, W. Schmidt, Q. Si, and F. Steglich, Nat. Phys. 7, 119 (2011).
- [6] C. Stock, C. Broholm, J. Hudis, H. J. Kang, and C. Petrovic, Phys. Rev. Lett. 100, 087001 (2008).
- [7] C. Stock, C. Broholm, Y. Zhao, F. Demmel, H. J. Kang, K. C. Rule, and C. Petrovic, Phys. Rev. Lett. 109, 167207 (2012).
- [8] P. Coleman, Phys. Rev. B 35, 5072 (1987).
- [9] S. Burdin, A. Georges, and D. R. Grempel, Phys. Rev. Lett. 85, 1048 (2000).
- [10] T. Kawai, M. Nakashima, Y. Okuda, H. Shishido, T. Shimoda, T. D. Matsuda, Y. Haga, T. Takeuchi, M. Hedo, Y. Uwatoko, R. Settai, and Y. Onuki, J. Phys. Soc. Jpn. 76, 166 (2007).
- [11] Y. Muro, D. Eom, N. Takeda, and M. Ishikawa, J. Phys. Jpn. 67, 3601 (1998).

- [12] N. Kimura, Y. Muro, and H. Aoki, J. Phys. Jpn. 76, 051010 (2007).
- [13] Y. F. Yang, Z. Fisk, H. O. Lee, J. D. Thompson, and D. Pines, Nature (London) 454, 611 (2008).
- [14] P. Coleman, C. Pepin, Q. Si, and R. Ramazashvili, J. Phys.: Condens. Matter 13, R723 (2001).
- [15] T. Terashima, M. Kimata, S. Uji, T. Sugawara, N. Kimura, H. Aoki, and H. Harima, Phys. Rev. B 78, 205107 (2008).
- [16] N. Aso, H. Miyano, H. Yoshizawa, N. Kimura, T. Komatsubara, and H. Aoki, J. Magn. Magn. Mater. 310, 602 (2007).
- [17] N. Kimura, K. Ito, K. Saitoh, Y. Umeda, H. Aoki, and T. Terashima, Phys. Rev. Lett. 95, 247004 (2005).
- [18] N. Kimura, T. Sugawara, and H. Aoki, J. Phys. Jpn. 80, SA019 (2011).
- [19] Y. Tada, N. Kawakami, and S. Fujimoto, Phys. Rev. Lett. 101, 267006 (2008).
- [20] N. Kimura, K. Ito, H. Aoki, S. Uji, and T. Terashima, Phys. Rev. Lett. 98, 197001 (2007).
- [21] T. Sugawara, N. Kimura, H. Aoki, F. Lévy, I. Sheikin, and T. Terashima, J. Phys. Soc. Jpn. 79, 063701 (2010).
- [22] N. Egetenmeyer, J. L. Gavilano, A. Maisuradze, S. Gerber, D. E. MacLaughlin, G. Seyfarth, D. Andreica, A. Desilets-Benoit, A. D. Bianchi, C. Baines, R. Khasanov,

Z. Fisk, and M. Kenzelmann, Phys. Rev. Lett. **108**, 177204 (2012).

- [23] T. Sugawara, N. Kimura, and H. Aoki, J. Phys. Soc. Jpn. 80, SA067 (2011).
- [24] H. Iida, T. Sugawara, H. Aoki, and N. Kimura, Phys. Status Solidi B 250, 502 (2013).
- [25] J. F. Landaeta, D. Subero, D. Catalá, S. V. Taylor, N. Kimura, R. Settai, Y. Ōnuki, M. Sigrist, and I. Bonalde, Phys. Rev. B 97, 104513 (2018).
- [26] Y. Yanase and M. Sigrist, J. Phys. Soc. Jpn. 76, 124709 (2007).
- [27] P. Y. Chang, O. Erten, and P. Coleman, Nat. Phys. 13, 794 (2017).
- [28] L. P. Gor'kov and E. I. Rashba, Phys. Rev. Lett. 87, 037004 (2001).
- [29] Y. Tada, N. Kawakami, and S. Fujimoto, Phys. Rev. B 81, 104506 (2010).
- [30] Y. Tada, S. Fujimoto, and N. Kawakami, J. Phys. Chem. Solids 69, 3341 (2008).
- [31] Y. Tada, N. Kawakami, and S. Fujimoto, J. Phys. Soc. Jpn. 77, 054707 (2008).
- [32] Y. Muro, M. Ishikawa, K. Hirota, Z. Hiroi, N. Takeda, N. Kimura, and H. Aoki, J. Phys. Jpn. 76, 033706 (2007).
- [33] M. T. Hutchings, Solid State Phys. 16, 227 (1964).
- [34] R. J. Birgeneau, E. Bucher, L. Passell, and K. C. Turberfield, Phys. Rev. B 4, 718 (1971).
- [35] K. C. Turberfield, L. Passell, R. J. Birgeneau, and E. Bucher, J. Appl. Phys. 42, 1746 (1971).
- [36] A. Krimmel, M. Reehuis, and A. Loidl, Appl. Phys. A 74, s695 (2002).
- [37] C. Stock, C. Broholm, F. Demmel, J. Van Duijn, J. W. Taylor, H. J. Kang, R. Hu, and C. Petrovic, Phys. Rev. Lett. 109, 127201 (2012).
- [38] Q. M. Si and F. Steglich, Science 329, 1161 (2010).
- [39] A. H. Castro Neto, G. Castilla, and B. A. Jones, Phys. Rev. Lett. 81, 3531 (1998).
- [40] T. Sugawara, H. Iida, H. Aoki, and N. Kimura, J. Phys. Soc. Jpn. 81, 054711 (2012).
- [41] C. Stock, J. A. Rodriguez-Rivera, K. Schmalzl, F. Demmel, D. K. Singh, F. Ronning, J. D. Thompson, and E. D. Bauer, Phys. Rev. Lett. **121**, 037003 (2018).
- [42] C. Stock, J. A. Rodriguez-Rivera, K. Schmalzl, E. E. Rodriguez, A. Stunault, and C. Petrovic, Phys. Rev. Lett. 114, 247005 (2015).
- [43] C. Krellner, N. S. Kini, E. M. Bruning, K. Koch, H. Rosner, M. Nicklas, M. Baenitz, and C. Geibel, Phys. Rev. B 76, 104418 (2007).
- [44] P. C. Canfield, J. D. Thompson, and Z. Fisk, J. Appl. Phys. 70, 5992 (1991).
- [45] P. M. Sarte, R. A. Cowley, E. E. Rodriguez, E. Pachoud, D. Le, V. García-Sakai, J. W. Taylor, C. D. Frost, D. Prabhakaran, C. MacEwen, A. Kitada, A. J. Browne, M. Songvilay, Z. Yamani, W. J. L. Buyers, J. P. Attfield, and C. Stock, Phys. Rev. B 98, 024415 (2018).
- [46] G. L. Stamokostas and G. A. Fiete, Phys. Rev. B 97, 085150 (2018).
- [47] R. A. Cowley, W. J. L. Buyers, C. Stock, Z. Yamani, C. Frost, J. W. Taylor, and D. Prabhakaran, Phys. Rev. B 88, 205117 (2013).

- [48] J. Sakurai, W. J. L. Buyers, R. A. Cowley, and G. Dolling, Phys. Rev. 167, 510 (1968).
- [49] J. Kanamori, Prog. Theor. Phys. 17, 177 (1957).
- [50] J. Kanamori, Prog. Theor. Phys. 17, 197 (1957).
- [51] P. Das, S. Z. Lin, N. J. Ghimire, K. Huang, F. Ronning, E. D. Bauer, J. D. Thompson, C. D. Batista, G. Ehlers, and M. Janoschek, Phys. Rev. Lett. **113**, 246403 (2014).
- [52] Y. Song, J. V. Dyke, I. K. Lum, B. D. White, S. Jang, D. Yazici, L. Shu, A. Schneidewind, P. Cermak, Y. Qiu, M. B. Maple, D. K. Morr, and P. Dai, Nat. Commun. 7, 12774 (2016).
- [53] C. Stock, R. A. Cowley, W. J. L. Buyers, C. D. Frost, J. W. Taylor, D. Peets, R. Liang, D. Bonn, and W. N. Hardy, Phys. Rev. B 82, 174505 (2010).
- [54] C. Stock, E. E. Rodriguez, O. Sobolev, J. A. Rodriguez-Rivera, R. A. Ewings, J. W. Taylor, A. D. Christianson, and M. A. Green, Phys. Rev. B 90, 12113 (2014).
- [55] C. Broholm, J. K. Kjems, G. Aeppli, Z. Fisk, J. L. Smith, S. M. Shapiro, G. Shirane, and H. R. Ott, Phys. Rev. Lett. 58, 917 (1987).
- [56] C. Stock, D. A. Sokolov, P. Bourges, P. H. Tobash, K. Gofryk, F. Ronning, E. D. Bauer, K. C. Rule, and A. D. Huxley, Phys. Rev. Lett. 107, 187202 (2011).
- [57] A. Schröder, G. Aeppli, E. Bucher, R. Ramazashvili, and P. Coleman, Phys. Rev. Lett. 80, 5623 (1998).
- [58] Z. Yamani, W. J. L. Buyers, F. Wang, Y. J. Kim, J. H. Chung, S. Chang, P. M. Gehring, G. Gasparovic, C. Stock, C. L. Broholm, J. C. Baglo, R. Liang, D. A. Bonn, and W. N. Hardy, Phys. Rev. B 91, 134427 (2015).
- [59] C. Stock, W. J. L. Buyers, Z. Yamani, Z. Tun, R. J. Birgeneau, R. Liang, D. Bonn, and W. N. Hardy, Phys. Rev. B 77, 104513 (2008).
- [60] G. Shirane, S. M. Shapiro, and J. M. Tranquada, *Neutron Scattering with a Triple-Axis Spectrometer* (Cambridge University Press, Cambridge, 2004).
- [61] B. Fåk, S. Raymond, D. Braithwaite, G. Lapertot, and J.-M. Mignot, Phys. Rev. B 78, 184518 (2008).
- [62] M. Smidman, D. T. Adroja, A. D. Hillier, L. C. Chapon, J. W. Taylor, V. K. Anand, R. P. Singh, M. R. Lees, E. A. Goremychkin, M. M. Koza, V. V. Krishnamurthy, D. M. Paul, and G. Balakrishnan, Phys. Rev. B 88, 134416 (2013).
- [63] V. K. Anand, A. D. Hillier, D. T. Adroja, D. D. Khalyavin, P. Manuel, G. Andre, S. Rols, and M. M. Koza, Phys. Rev. B 97, 184422 (2018).
- [64] T. Terashima, Y. Takahide, T. Matsumoto, S. Uji, N. Kimura, H. Aoki, and H. Harima, Phys. Rev. B 76, 054506 (2007).
- [65] S. Doniach, Phys. B (Amsterdam, Neth.) 91, 231 (1977).
- [66] P. Coleman, Phys. Rev. B 28, 5255 (1983).
- [67] A. Auerbach and K. Levin, Phys. Rev. Lett. 57, 877 (1986).
- [68] C. Lacroix and M. Cyrot, Phys. Rev. B 20, 1969 (1979).
- [69] R. Settai, I. Sugitani, Y. Okuda, A. Thamizhavel, M. Nakashima, Y. Onuki, and H. Harima, J. Magn. Magn. Mater. 310, 844 (2007).
- [70] H. A. Tanaka, N. Aso, M. Hedo, T. Nakama, Y. Takaesu, H. Iida, N. Kimura, and H. Aoki, J. Korean Phys. Soc. 62, 2016 (2013).
- [71] M. B. T. Tchokonte, P. V. Plessis, and A. M. Strydom, Solid State Commun. 117, 321 (2001).