



HAL
open science

Magnetic properties and complex magnetic phase diagram in non-centrosymmetric EuRhGe₃ and EuIrGe₃ single crystals

Arvind Maurya, P. Bonville, R. Kulkarni, A. Thamizhavel, S.K. Dhar

► **To cite this version:**

Arvind Maurya, P. Bonville, R. Kulkarni, A. Thamizhavel, S.K. Dhar. Magnetic properties and complex magnetic phase diagram in non-centrosymmetric EuRhGe₃ and EuIrGe₃ single crystals. Journal of Magnetism and Magnetic Materials, 2015, 401, pp.823-831. 10.1016/j.jmmm.2015.10.134 . cea-01550709

HAL Id: cea-01550709

<https://cea.hal.science/cea-01550709>

Submitted on 29 Jun 2017

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Magnetic properties and complex magnetic phase diagram in non-centrosymmetric EuRhGe₃ and EuIrGe₃ single crystals



Arvind Maurya^a, P. Bonville^b, R. Kulkarni^a, A. Thamizhavel^a, S.K. Dhar^{a,*}

^a Department of Condensed Matter Physics and Materials Science, Tata Institute of Fundamental Research, Homi Bhabha Road, Colaba, Mumbai 400005, India

^b CEA, Centre d'Etudes de Saclay, DSM/IRAMIS/Service de Physique de l'Etat Condensé and CNRS UMR 3680, 91191 Gif-sur-Yvette, France

ARTICLE INFO

Article history:

Received 3 March 2015

Received in revised form

8 September 2015

Accepted 29 October 2015

Available online 31 October 2015

Keywords:

EuRhGe₃

EuIrGe₃

Single crystal

Magnetic phase diagram

¹⁵¹Eu Mössbauer spectra

ABSTRACT

We report the magnetic properties of two Eu based compounds, single crystalline EuIrGe₃ and EuRhGe₃, inferred from magnetisation, electrical transport, heat capacity and ¹⁵¹Eu Mössbauer spectroscopy. These previously known compounds crystallise in the non-centrosymmetric, tetragonal, *I4*mm, BaNiSn₃-type structure. Single crystals of EuIrGe₃ and EuRhGe₃ were grown using a high temperature solution growth method using In as flux. EuIrGe₃ exhibits two magnetic transition temperatures $T_{N1} = 12.4$ K, and $T_{N2} = 7.3$ K, whereas EuRhGe₃ presents a single one at $T_N = 12$ K. ¹⁵¹Eu Mössbauer spectra show evidence for a cascade of transitions from paramagnetic to incommensurate amplitude modulated followed by an equal moment phase at lower temperature in EuIrGe₃. This latter phase alone occurs in EuRhGe₃. In both compounds, the magnetisation measured up to 14 T suggests that the equal moment magnetic phase has a spiral spin arrangement. The field induced reorientations are also well documented in the magneto-transport data. A superzone gap is observed for the current density $J_{||}$ [001], which is enhanced by a transverse magnetic field. The magnetic phase diagram constructed from all the data is complex, revealing the presence of many phases in the $H - T$ space.

© 2015 Elsevier B.V. All rights reserved.

1. Introduction

The magnetic properties of several Eu-based compounds with composition EuT₃, where *T* is a *d*-block transition element and *X*=Si or Ge, have been reported in the literature [1–6]. These compounds crystallise in the non-centrosymmetric (NCS) BaNiSn₃-type structure. A transition from the paramagnetic to an incommensurate, amplitude modulated state followed by a second transition to a single moment phase in EuPtSi₃ [1] and EuNiGe₃ [5] was inferred from heat capacity and ¹⁵¹Eu Mössbauer spectroscopy. Magnetisation data on a single crystal of EuPtSi₃ showed the presence of anisotropy, probably of both crystalline and exchange origin. Only one magnetic transition is observed in EuPtGe₃ and EuPdGe₃ [2,3]. Further, magnetisation data on a single crystal of EuNiGe₃ showed that while the *ab*-plane is the hard plane, an unusual staircase-like behavior of magnetisation is observed along the *c*-axis [5]. Thus, a variety of interesting magnetic behaviours is observed in EuT₃ compounds. The magnetisation in the antiferromagnetic state shows a varying degree of anisotropy which is *a priori* surprising for a spin-only ($S=7/2$; $L=0$) ion like Eu²⁺.

The existence of iso-structural EuIrGe₃ and EuRhGe₃ is known and it was of interest to study the magnetic behaviour of these two compounds. We have probed the detailed magnetic properties of single crystalline EuIrGe₃ and EuRhGe₃ by magnetisation, resistivity and heat capacity in zero and applied fields, and ¹⁵¹Eu Mössbauer spectroscopy. While this work was in progress, the magnetic properties of single crystalline EuIrGe₃ and EuRhGe₃ have been reported, i.e. susceptibility measurements in a field of 0.1 T, zero-field electrical resistivity and heat capacity [6]. Our main observations are in agreement with the results reported in Ref. [6]; however, our more extensive data include isothermal magnetisation at selected temperatures, susceptibility measured at a number of applied fields, magnetoresistivity, construction of magnetic phase diagram, observation of the superzone gap at the antiferromagnetic transition and ¹⁵¹Eu Mössbauer spectra. In addition, we also prepared LaRhGe₃ and LaIrGe₃ as non-magnetic reference compounds and measured their heat capacity and electrical resistivity.

2. Experimental

Polycrystalline samples of EuIrGe₃ and EuRhGe₃ were first prepared by melting Eu (99.9% purity), Ir/Rh (99.99%) and Ge

* Corresponding author.

E-mail address: sudesh@tifr.res.in (S.K. Dhar).

(99.999%) in an arc furnace under an inert atmosphere of argon. The alloy buttons were flipped over three times and re-melted to ensure proper homogenisation. An excess of about 10% over the stoichiometric amount was taken for Eu, to compensate for the weight loss due to evaporation of Eu. Initially, we attempted to grow the single crystals of these compounds by using Sn as a solvent, as that choice had proved successful for EuPtSi₃ and EuPtGe₃, but it did not give the desired results. In a second attempt, charges of EuIrGe₃ and EuRhGe₃ and In (as solvent) were taken in the weight ratio 1:8, placed together in separate alumina crucibles and sealed in quartz ampoules under a partial pressure of 10⁻⁶ mbar. The sealed crucibles were placed in a box type resistive heating furnace and heated to 1100 °C at a rate of 50 °C/h. After a soaking period of 24 h, a cooling rate of 2 °C/h was employed down to 600 °C. The cooling rate was increased to 60 °C/h below 600 °C. The single crystals of EuIrGe₃ and EuRhGe₃ were separated from In-flux by centrifugation. Small traces of indium were washed away by etching the grown crystals in dilute hydrochloric acid. Polycrystalline samples of non-magnetic reference LaIrGe₃ and LaRhGe₃ were prepared by the standard technique of arc melting as described above. The magnetisation as a function of field (up to 14 T) and temperature (1.8 to 300 K) was measured using Quantum Design Magnetic properties measurement system (MPMS) and Vibration sample magnetometers (VSM). The electrical resistivity between 1.8 and 300 K and the heat capacity in zero and applied fields was measured in a Quantum Design Physical properties measurement system (PPMS) unit. ¹⁵¹Eu Mössbauer spectra were recorded at various temperatures using a constant acceleration spectrometer with a ¹⁵¹SmF₃ source. Laue diffraction patterns were recorded on a Huber Laue diffractometer fitted with an image plate, while powder-diffraction spectra were recorded on a Philips PANalytical set up using Cu-K α radiation. The crystals were cut by spark erosion electric discharge machine and oriented along the desired planes using a triple axis goniometer and Laue diffraction in the back reflection mode.

3. Results and discussion

3.1. Structure

Well faceted crystals having a platelet geometry and typical dimensions of ~5 mm × 5 mm × 1 mm were obtained after centrifuging out the In solvent. The composition of the crystals was confirmed using electron dispersive analysis by x-rays (EDAX). The powder x-ray diffraction spectra, obtained by crushing a few single crystals to powder, could be indexed to tetragonal BaNiSn₃-type structure.

The lattice parameters obtained by the Rietveld analysis of the powder diffraction spectra using FullProf software package [7] are listed in Table 1 and are in good agreement with the previously reported values [6,8].

3.2. Magnetisation

The inverse susceptibility, χ^{-1} , of EuIrGe₃, between 1.8 and 300 K, with the magnetic field (0.1 T) applied parallel to [100] and

Table 1
Lattice parameters *a* and *c*, and unit cell volume *V* of EuRhGe₃ and EuIrGe₃ obtained from the Rietveld refinement of x-ray powder diffraction pattern.

Compound	<i>a</i> (Å)	<i>c</i> (Å)	<i>V</i> (Å ³)
EuRhGe ₃	4.407(3)	10.068(7)	195.57(7)
EuIrGe ₃	4.430(0)	10.041(6)	197.06(5)

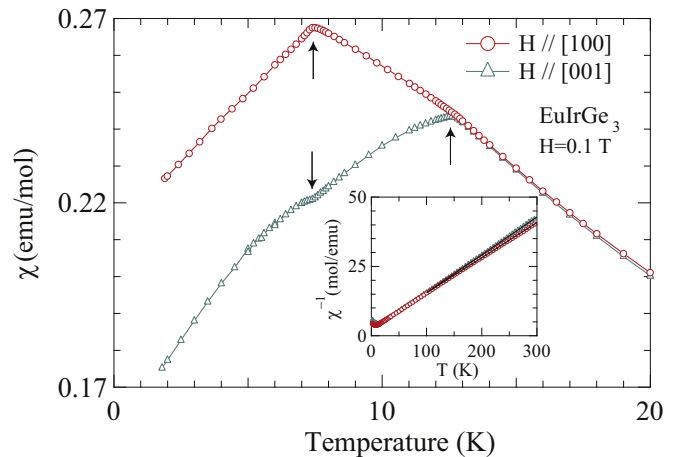


Fig. 1. Magnetic susceptibility and inverse magnetic susceptibility (in inset) of EuIrGe₃ at field 0.1 T along [100] and [001].

[001] directions is shown in the inset of Fig. 1. The fit of χ^{-1} to the Curie–Weiss expression $\chi(T) = \mu_{eff}^2 / [8(T - \theta_p)]$ between 100 and 300 K provides the following parameters: $\mu_{eff} = 7.94$ and $7.65 \mu_B$, and $\theta_p = -22.1$ and -13.7 K for $H \parallel [100]$ and $[001]$, respectively. The value of μ_{eff} along [100] matches exactly with the Hund's rule derived value. An antiferromagnetic interaction between the divalent Eu ions is inferred from the negative values of θ_p along the two directions. The susceptibility below 20 K is shown in the main panel of Fig. 1. There is a cusp at $T_{N1} = 12.4$ K typical of antiferromagnetic ordering, followed by a mild shoulder at 7.3 K for $H \parallel [001]$.

On the other hand for $H \parallel [100]$ there is a very subtle change of slope observed at 12.4 K and a very clear cusp is observed at 7.3 K. The main features of the susceptibility in the magnetically ordered state are in good agreement with those reported in Ref. [6]. For a collinear bipartite antiferromagnet, the susceptibility below T_N is temperature independent along the magnetic hard axis while it gradually decreases to zero along the easy axis as the temperature is lowered to zero. In the present case, χ decreases along both [001] and [100] indicating a magnetic configuration which is different from simple collinear bipartite antiferromagnetic. This is confirmed by the isothermal magnetisation data, to be described below, which suggest a spiral spin arrangement. T_{N1} is comparable to θ_p for $H \parallel [001]$ but it is smaller than θ_p for $H \parallel [100]$. We mention here that in Ref. [6] a single θ_p value of -17 K is reported.

$M(T)/H$ at selected fields, along [100] and [001] was measured in various fields ranging from 0.5 to 6 T and the data below 15 K are shown in Fig. 2. Additional features appear as the field is increased above 0.1 T.

At 0.5 T, M/H ($H \parallel [100]$) shows a knee near 5.5 K, which shifts to lower temperatures as the field is increased and either vanishes at $H = 3$ T or occurs below 1.8 K. The anomalies at 12.4 K and 7.3 K also shift slightly to lower temperatures with increasing field. For $H \parallel [001]$, the magnetisation between 2.5 and 4 T shows a prominent upturn at low temperatures, indicating a field-induced change in the direction of the magnetic moments. The mild shoulder at 7.3 K shifts to higher temperatures with increasing field while the peak at 12.4 K (in 0.1 T field) shifts to lower temperatures. The two appear to merge in a field of 6 T. At high fields, ~3 T and above, the magnetisation M/H at temperatures approaching 1.8 K is comparable for both the directions. These plots show that the configuration of the Eu moments in the magnetically ordered regime is modified by the applied field. The critical points derived from $M/H(T, H)$ have been included while constructing the $H - T$ phase diagram in Fig. 12.

The magnetisation at selected temperatures for $H \parallel [100]$ and

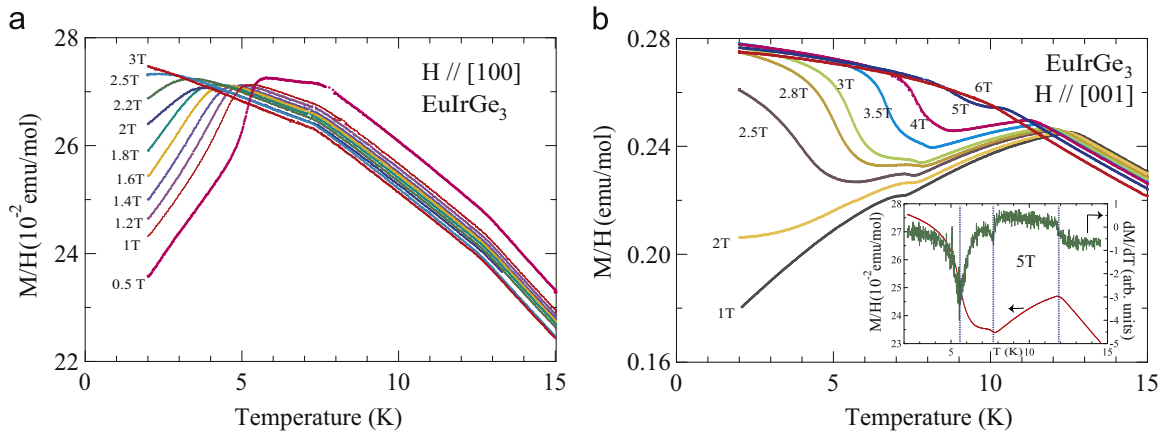


Fig. 2. Temperature dependence of M/H at selected fields along [100] (left panel) and [001] (right panel) for EuIrGe_3 . Inset in (b) shows $M/H(T)$ at 5 T on the left scale and its derivative on the right scale with vertical dotted line at critical points.

[001] in applied fields up to 14 T is shown in Figs. 3a and b, respectively. In the inset of Fig. 3a the data at 1.8 K along the two directions are plotted. Above 3 T and between 3 and 8 T, the magnetisations at 1.8 K along the two directions virtually overlap (in conformity with the $M - T$ data discussed above) and then slightly bifurcate at higher fields. Whereas the magnetisation for $H \parallel [100]$ is linear with field up to about 10 T, the magnetisation along [001] shows a clear dip around 2 T at 1.8 K, which shifts to higher field as temperature increases. Such a behaviour reminds that of a planar spiral magnetic structure with sizeable anisotropy [9]: for a field perpendicular to the spiral plane, a conical structure progressively develops and the magnetisation is linear with the field; for a small field within the spiral plane, anisotropy prevents the conical structure to appear and the moments “lag” in their plane until a threshold field is reached, recovering the conical arrangement. Therefore, the susceptibility in this latter case is lower than for the perpendicular case, yielding both the observed dip in the $M(H)$ curve and the lower M/H values reported above. For EuIrGe_3 , this suggests that the plane of the spiral is perpendicular to [100] and contains [001].

At 1.8 K, the magnetisation at 14 T along [001] ($6.3 \mu_B/\text{Eu}$) is slightly larger than along [100] ($6.15 \mu_B/\text{Eu}$), and the saturation value of $7 \mu_B$ is not reached along both directions. Thus the spin-flip field at 1.8 K is nearly independent of the direction (like in EuPtGe_3) and should be close to 16 T.

In contrast to EuIrGe_3 , the Rh analog shows only one anti-ferromagnetic transition which occurs close to $T_N = 12$ K in applied field of 0.1 T (see, Fig. 4), in accordance with Ref. [6]. Fig. 4 shows the susceptibility data below 20 K for $H \parallel [100]$, [110] and [001]. It is noticed that the response in the ab -plane is isotropic. Above 100 K, the inverse susceptibility fits well to the Curie–Weiss law with the values: $\mu_{\text{eff}} = 7.56$ and $7.78 \mu_B$, $\theta_p = -7$ and -11 K for $H \parallel [100]$ and [001], respectively. It is likely that the value of μ_{eff} is slightly lower due to the presence of tiny inclusions of In metal in the crystal which get incorporated during the crystal growth. As a result the amount of Eu used in the calculation of μ_{eff} is actually slightly overestimated. We infer the presence of In from the slight dip in the susceptibility measured in an applied field of 0.005 T near the superconducting transition temperature of In. The dip vanishes when the applied field is increased to 0.05 T.

At higher fields applied parallel to [100] the transition shifts to lower temperatures reaching 9.4 K in 10 T (see, Fig. 5a). An additional peak which occurs close to 11 K in 1 T gradually shifts to 2.8 K in 8 T. Between 4 and 6 T, the magnetisation shows additional features below 5 K which are depicted in Fig. 5b. The field induced changes in the magnetic configuration have a first order character as indicated by the hysteresis observed at 5 T (see inset

of Fig. 5a). The data depicted in Figs. 5a and 5b indicate a field induced complex phase diagram. On the other hand, for fields along the [001] direction the character of the plots remains unaltered except that T_N decreases with field (see Fig. 5c). While our observations do not provide us the actual configuration of the magnetic moments, they appear to suggest a non-collinear anti-ferromagnetic structure which evolves in a complex fashion for field applied in the ab -plane.

The main panel of Fig. 6a shows the isothermal magnetisation plots at 2 K for $H \parallel [100]$, [110] and [001]. The magnetisation along [100] at various temperatures is plotted in Fig. 6b; the inset shows the magnetisation at 2 and 5 K for $H \parallel [001]$.

Along [001] the magnetisation increases nearly linearly with the field reaching a value of $6.2 \mu_B/\text{Eu}$ ion at 14 T. There is hardly any change in the magnetisation at 2 and 5 K (Fig. 6b inset). By contrast, for $H \parallel [100]$ and [110], one sees a deviation from linearity as a large dip up to 5 T. This behaviour could be due to a spiral spin arrangement, like in EuIrGe_3 , but with the plane of the spiral perpendicular to [001]. Furthermore, the structure observed in the dip near 3.5 T could point to a still more complex moment arrangement. Above 10 T, the ab -plane isotropy of the magnetisation is slightly violated. At the highest field (14 T) the magnetisations along the three directions are nearly identical. A derivative plot of magnetisation for $H \parallel [100]$, dM/dH , shown in the inset of Fig. 6a shows three peaks which are a signature of the field induced changes in the magnetic configuration. The magnetisation shows hysteresis around 5 T at 2 K (see, upper inset of Fig. 6a) which correlates nicely with the data depicted in the inset of Fig. 5a. The temperature dependence of $M(H)$ along [100] is shown in main panel of Fig. 6b.

3.3. Heat capacity

The specific heat measured down to 100 mK in zero field (Fig. 7a) confirms the presence of two transitions in EuIrGe_3 , at $T_{N1} = 12.4$ and $T_{N2} = 7.2$ K, in close correspondence with the low-field magnetisation data presented above. The magnitude of the jump in the heat capacity at T_{N1} , ~ 5 J/mol K, which is far below the mean-field value of 20.14 J/mol K for a mol of spin $S = 7/2$. This suggests that the transition at T_{N1} is from paramagnetic to amplitude modulated anti-ferromagnetic configuration. At T_{N2} the transition from this intermediate state to an equal moment configuration takes place, as confirmed by ^{151}Eu Mössbauer spectra (see below).

The heat capacity was also measured in applied fields of 8 and 14 T with $H \parallel [100]$. At 8 T, the two peaks at 7.2 and 12.4 K (zero field) have shifted slightly lower in temperature to 6.1 and 11 K,

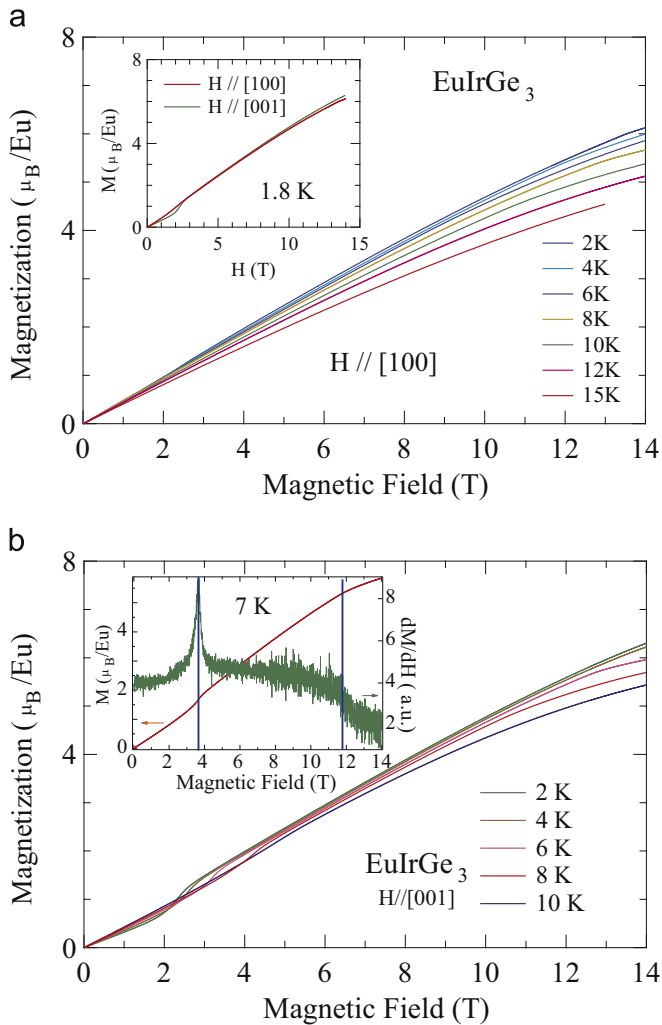


Fig. 3. Field dependence of magnetisation at selected temperatures in EuIrGe_3 along (a) [100] and (b) [001]. Inset in (a) represents nearly isotropic magnetisation plots in EuIrGe_3 and in (b) a representative derivative of magnetisation plot along [001] at 7 K (right scale) along with the $M(H)$ data (left scale) is shown. (For interpretation of the references to color in this figure caption, the reader is referred to the web version of this paper.)

respectively, in correspondence with the magnetisation data shown in Fig. 2. The jump in the heat capacity is slightly reduced. At 14 T, there is only one peak at 7 K, with appreciable reduction in

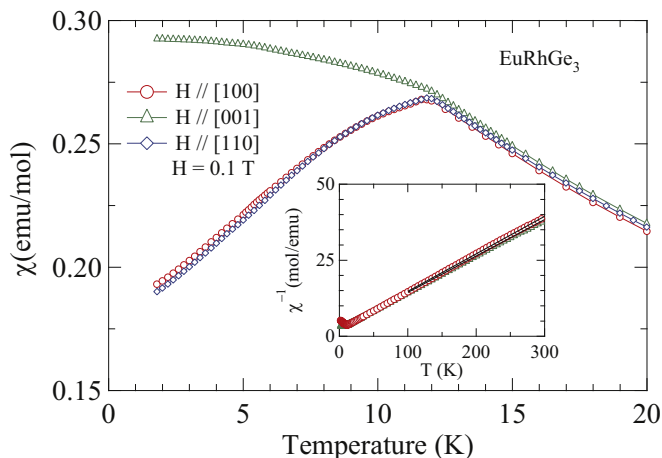


Fig. 4. Anisotropic magnetic susceptibility and inverse magnetic susceptibility (in inset) as a function of temperature at magnetic field 0.1 T of EuRhGe_3 .

peak height. For $H \parallel [001]$, at 5 T both peaks come closer but are still well resolved; at higher fields we observe only a single peak in agreement with magnetic phase diagram in Fig. 12b.

The heat capacity of the iso-structural LaIrGe_3 is also plotted in Fig. 7a, and the 4f contribution to the heat capacity, C_{4f} , and entropy S_{4f} were calculated under the assumption that the phonon heat capacities of LaIrGe_3 and EuIrGe_3 are identical, after normalisation due to the slightly different atomic masses of La and Eu. The entropy attains the value of $R \ln 8$ (for Eu^{2+} ions, $S=7/2$ and $L=0$) near 18 K but keeps on increasing at higher temperatures, indicating a poor validity of the assumption of identical phonon spectra in LaIrGe_3 and EuIrGe_3 at least at higher temperatures. A similar situation was earlier encountered in EuPtSi_3 [11]. The C/T vs. T^2 plot of LaIrGe_3 is linear below 8 K, characterized by $\gamma = 4.0 \text{ mJ/mol K}^2$ and $\beta = 0.349 \text{ mJ/mol K}^4$. A Debye temperature, θ_D , of 303 K is inferred from β .

The heat capacity data of EuRhGe_3 measured in zero and 8 T ($H \parallel [100]$) are plotted in Fig. 7b. The relatively sharp jump in the heat capacity near 12 K in zero field is in excellent agreement with the magnetisation data discussed above. Surprisingly, the jump in the heat capacity at T_N in EuRhGe_3 is about 13 J/mol K, which is far below the value for a transition to an equal moment antiferromagnetic state (20.14 J/mol K) in the mean field model [10]. It may be noted that the shape of the heat capacity variation below T_N is rather unusual and similar to some cases of amplitude modulated moment state described in Fig. 5 in Ref. [10]. In applied fields the magnetic transition shifts to lower temperatures and additional peaks, marked by downward arrows in Fig. 7 for 5 T data are observed, in conformity with the in-field magnetisation data (Fig. 5) described above. The heat capacity of non-magnetic, reference analogue LaRhGe_3 is also plotted. The entropy S_{4f} was estimated using the method mentioned above. It again exceeds the maximum value of $R \ln 8$ like in the Ir compound. For LaRhGe_3 , $\gamma = 6.7 \text{ mJ/mol K}^2$ and $\beta = 0.376 \text{ mJ/mol K}^4$. A Debye temperature, θ_D , of 296 K is inferred from β , which is close to that of the Ir compound.

3.4. Electrical resistivity

The electrical resistivity of EuIrGe_3 and EuRhGe_3 with the current density J parallel to [100] and [001], respectively, is shown in Fig. 8. The resistivity shows anomalies for EuIrGe_3 at the two transitions T_{N1} and T_{N2} along both directions though with slightly different characteristics. Along [100] the resistivity decreases faster at each transition due to the rapid reduction in the spin-disorder scattering. Above T_{N1} the resistivity monotonically increases up to the room temperature. On the other hand, for $J \parallel [001]$ at T_{N1} there is a slight upturn on cooling. The upturn at T_{N1} is suggestive of a gap-opening at the Fermi surface along [001] direction with AFM order, often referred to as superzone gap. Many other rare earth intermetallics have been found to show this kind of behaviour [11–13].

The electrical resistivity of EuRhGe_3 has some similarities with that of EuIrGe_3 described above. For $J \parallel [100]$ the resistivity decreases at the single antiferromagnetic transition, while for $J \parallel [001]$ the initial increase at $T_N (= 12 \text{ K})$ again indicates the opening of a superzone gap like in EuIrGe_3 .

We have fitted the Bloch Grüneisen expression to our $\rho(T)$ data in the paramagnetic region given by following expression:

$$\rho(T) = A + B \left(\frac{T}{\theta_R} \right)^5 \int_0^{\theta_R/T} \frac{x^5}{(e^x - 1)(1 - e^{-x})} dx \quad (1)$$

where θ_R is the Debye temperature determined from the $\rho(T)$ data, A is the temperature independent part of resistivity comprising of electron scattering caused by crystal imperfections and spin

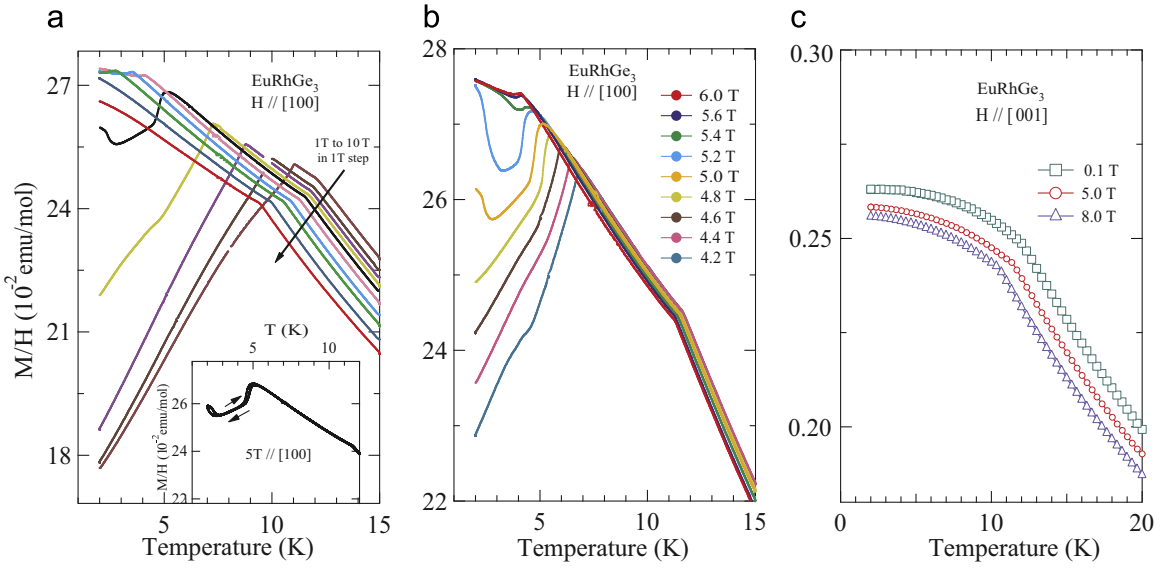


Fig. 5. $M(T)/H$ at selected fields along [100] (a and b) and [001] (c) for EuRhGe_3 . Inset in (a) shows the hysteresis in the data taken in the warming and cooling cycle at 5 T.

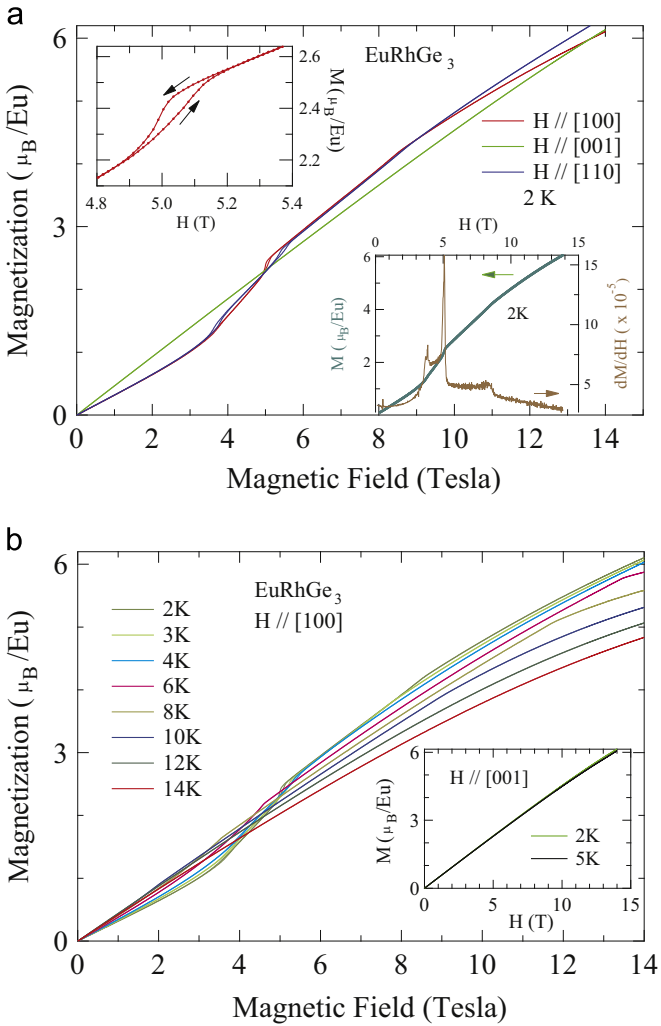


Fig. 6. (a) Isothermal magnetisation $M(H)$ at 2 K of EuRhGe_3 along the principal crystallographic directions. Bottom inset shows $M(H)$ at 2 K on left scale and its derivative on right scale revealing field induced spin reorientations. (b) Temperature evolution of $M(H)$ of EuRhGe_3 along [100] in main panel and along [001] in the inset. (For interpretation of the references to color in this figure caption, the reader is referred to the web version of this paper.)

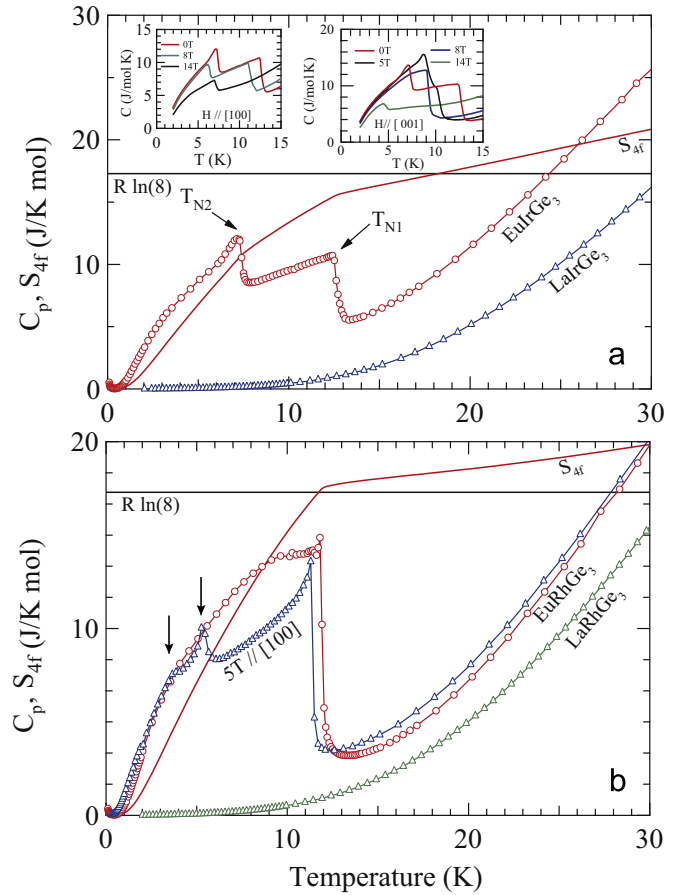


Fig. 7. Heat capacity ($C_p(T)$) and calculated magnetic entropy (S_{4f}) of (a) EuRhGe_3 and (b) EuRhGe_3 , respectively. $C_p(T)$ of corresponding La-analogues are also shown in the main panels. Insets in (a) show the $C_p(T)$ curves when field is applied along [100] (left) and [001] (right). In (b), blue trace represents data taken at 5 T field applied parallel to a -axis, capturing the field induced phase transitions indicated by arrows. (For interpretation of the references to color in this figure caption, the reader is referred to the web version of this paper.)

disorder in the paramagnetic state and B is a material dependent prefactor. Parameters determined from the fit are listed in Table 2. It may be noted that the magnitude of θ_R is different from θ_D . This

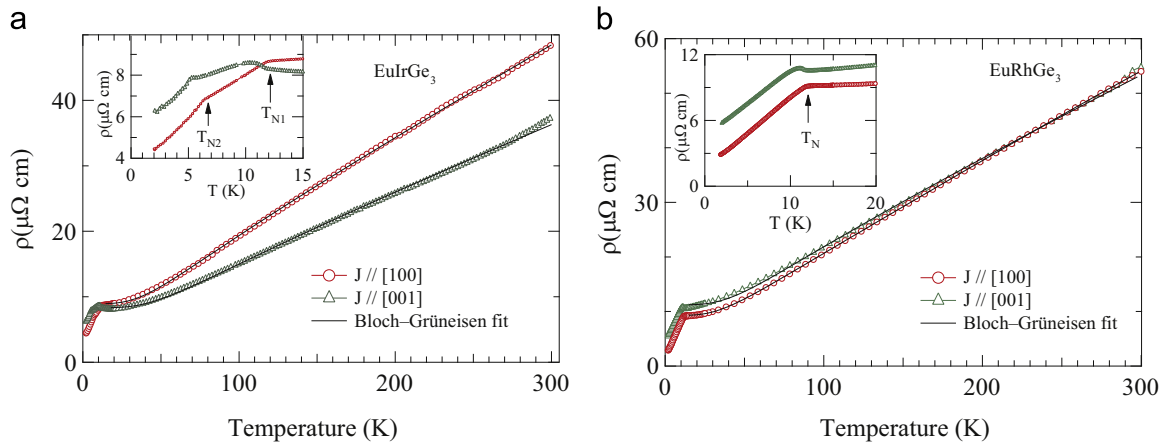


Fig. 8. Variation of electrical resistivity ($\rho(T)$) for current density $J_{||}$ [100] and [001] for (a) EuIrGe_3 and (b) EuRhGe_3 . Insets show low temperature data on an expanded scale where the magnetic transitions are marked by arrows.

is not unusual as θ_R considers only the longitudinal lattice vibrations.

The transverse magnetic field dependence of electrical resistivity under different configurations is shown in Figs. 9 and 10 for EuIrGe_3 and EuRhGe_3 , respectively. The main features in the $\rho(T)$ data of both compounds are in excellent correspondence with the magnetic susceptibility data. The upturn in the resistivity at T_N becomes more prominent as the field is increased (see, Figs. 9a and 10a), suggesting an enhancement of the superzone gap in the two compounds.

The magnetoresistivity MR , defined as $MR(H) = (\rho(H) - \rho(H=0)) \times 100/\rho(H=0)$, of EuIrGe_3 for different transverse configurations is shown in Figs. 11(a–c). For $H_{||}$ [001], [010] and $J_{||}$ [100] at $T=2$ K, the MR is positive, increases rapidly with field and shows a minor anomaly near 2 T which corresponds well with the spin-flop like transition seen in the magnetisation. The positive MR is typically expected in an antiferromagnet as the field disrupts the antiferromagnetic ordered state. The positive MR peaks near 12 T and then decreases slightly, indicating the proximity of the spin-flip field around 14 T. As the temperature is raised (see, Fig. 10c) the minor anomaly shifts to higher fields tracking the corresponding increase of the spin-flop like transition field in the magnetisation, and the magnitude of positive MR decreases due to the increase of temperature. At 6 K the MR becomes negative for $H > 10$ T, and the crossover field value decreases with further increase of temperature. At 15 K the MR in the paramagnetic state is negative at all fields, most likely due to the ordering effect of the field on the fluctuating moments.

The MR for $H_{||}$ [010] shows qualitatively similar field dependence (see, Fig. 10b) as described above for $H_{||}$ [001]. The anomalies observed in MR data are included in phase diagram corresponding well to the magnetisation data.

The MR of EuRhGe_3 at selected temperatures is shown in Figs. 11(d–f). Fig. 11d shows a comparison between MR data taken at 2 K in different configurations. The nature of MR curves is dependent upon the field direction as well as the direction of current

density J . For $H_{||}$ [100] and $J_{||}$ [001] the MR at 2 K is positive and initially increases with field. It shows anomalies at 5 and 8.5 T which mirror the anomalies seen in the magnetisation at these fields (Fig. 5b). Above 8.5 T, the MR is still positive but begins to decline in its absolute values most likely due to the increasing alignment of the moments along the field direction as the spin-flip field is approached. As the temperature is increased, the two anomalies approach each other, shifting in opposite directions, and above ~ 4 –5 K they apparently merge and then the single anomaly shifts to lower fields with increasing temperature (see Fig. 11e). At 6 K, the MR shows an anomaly in 13–14 T range which shifts to 11–12 and 8–9 T intervals at 8 and 10 K, respectively. This feature matches well with the magnetisation plots measured at these temperatures (indicated by arrows in Fig. 6b). In the paramagnetic region (15 and 20 K) the MR is negative. Lastly, the MR data for $H_{||}$ [001] and $J_{||}$ [100] are shown in Fig. 11f. It may be recalled that [001] is relatively the hard-axis of magnetisation in EuRhGe_3 . The MR up to 5 K is positive, increasing with field and showing a slight decline above 12 T. At 8 and 10 K the decline in MR is marked by a sharp knee at ~ 11.5 and ~ 9 T, respectively, which are phenomenologically similar to the one's seen in Fig. 11e and occur at similar values of fields as well, and may have a similar origin. Again, the MR in the paramagnetic region is negative.

3.5. Magnetic phase diagram

From $M(T, H)$ and $\rho(T, H)$ data we have constructed the magnetic phase diagrams of EuRhGe_3 and EuIrGe_3 shown in Fig. 12. The conclusions derived from these two sets of data correspond very well with each other. AF1, AF2, ... denote phases specified by different antiferromagnetic (or spiral/conical) configurations. In EuRhGe_3 , at low fields a second transition appears which shifts to lower temperatures with increasing magnetic field until the occurrence of a tricritical point at (5 K, 4.9 T), followed by another tricritical point at (4 K, 5.4 T). Dotted lines are plausible extrapolations.

The magnetic phase diagram in EuIrGe_3 also shows a similar degree of complexity. Here the red symbols have been used for points determined from $H_{||}$ [001] and blue symbols for $H_{||}$ [100] data, respectively for the M vs. T , M vs. H , R vs. H and R vs. T experiments. For $H_{||}$ [001], on increasing the field, T_{N1} and T_{N2} come closer and merge together accompanied with the appearance of other field induced transitions forming a closed dome centered around 10 K. This dome corresponds to the incommensurate modulated phase. Above 5 T we could observe only one transition along the c -axis. On the other hand, for field parallel to the a -axis

Table 2

Parameters derived from the Bloch Grüneisen fit to the $\rho(T)$ data of EuRhGe_3 and EuIrGe_3 .

	EuRhGe_3		EuIrGe_3	
	A ($\mu\Omega\text{cm}$)	θ_R (K)	A ($\mu\Omega\text{cm}$)	θ_R (K)
$J_{ }$ [100]	9.5	246	8.9	224
$J_{ }$ [001]	11.2	244	8.3	270

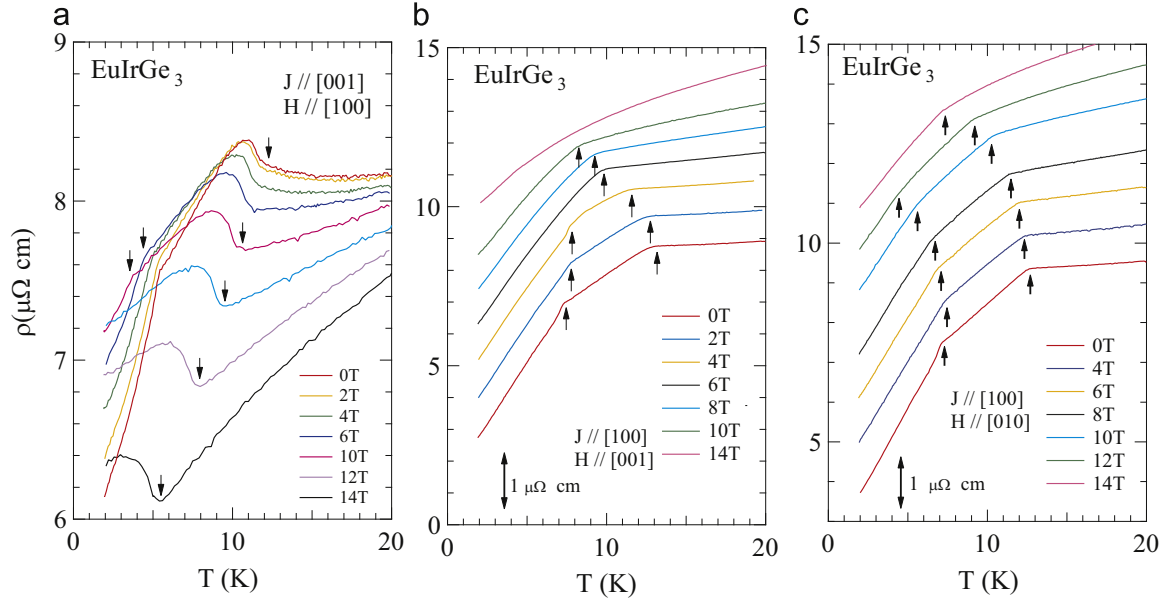


Fig. 9. $\rho(T)$ of EuIrGe_3 for different configurations of current and applied magnetic fields. In b and c, the traces for non-zero fields have been moved in vertical direction for clarity. Arrows indicate magnetic phase transitions. (For interpretation of the references to color in this figure caption, the reader is referred to the web version of this paper.)

T_{N1} and T_{N2} are suppressed, apparently merging at around 14.2 T at absolute zero. Interestingly, it looks that the high field phase for $H \parallel [001]$ seems to converge at the same point at 0 K as the merging point of two phase lines for $H \parallel [100]$. An additional low field phase line nearly parallel to the temperature axis, having opposite curvatures for a and c axes but merging with each other at around 2 K is observed. It may be noted that the phase boundary between PM and AF1 phase is also a demarcation line for the superzone gap in both EuRhGe_3 and EuIrGe_3 .

3.6. ^{151}Eu Mössbauer spectra

The ^{151}Eu Mössbauer spectra at 4.2, 8 and 12 K in EuIrGe_3 are shown in Fig. 13. The 4.2 K spectrum is a hyperfine field pattern with the single value $H_{hf} = 28.9(2)$ T, and it shows an isomer shift of $-10.6(1)$ mm/s with respect to the SmF_3 γ -ray source. Both these values are characteristic of divalent Eu, and the single hyperfine field indicates an equal moment magnetic ordering. At 8 K,

just above T_{N2} , the spectrum has changed shape and cannot be fitted using a single hyperfine field. Its shape is typical of an incommensurate magnetic structure (see for instance Ref. [5]), and it was fitted to a lineshape where the hyperfine field distribution is decomposed into its Fourier components:

$$H_{hf}(kx) = \sum_n m_{2n+1} \sin[(2n+1)kx], \quad (2)$$

where n is an integer running from zero to a maximum value n_{max} , and kx runs from 0 to 2π , x being the distance along the propagation vector \mathbf{k} . A good fit was obtained with $n_{max} = 4$, i.e. with 5 Fourier components, yielding a modulation with a maximum hyperfine field value of 29 T, equal to the 4.2 K single value. At 12 K, just below T_{N1} , the spectrum is broad and featureless, and can be fitted to a superposition of an incommensurate pattern, with maximum hyperfine field 17 T, and of a single narrow line characteristic of the paramagnetic phase, as observed at 15 K (spectrum not shown). The coexistence of the two phases close to T_{N1}

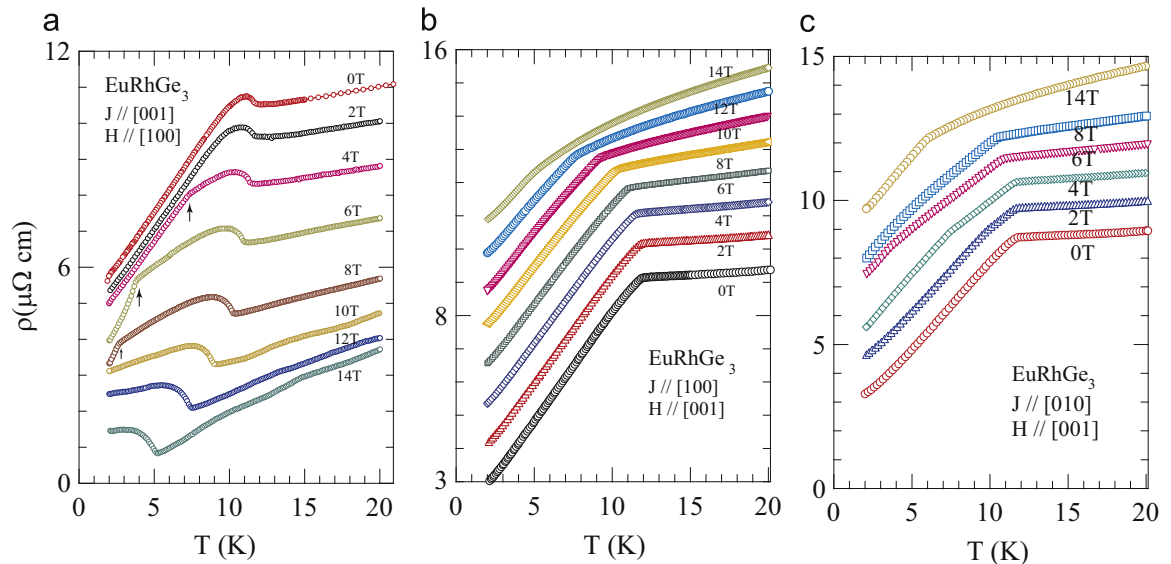


Fig. 10. $\rho(T)$ of EuRhGe_3 for different configurations of current and applied magnetic fields. $\rho(T)$ curves other than 0 T have been shifted vertically for clarity.

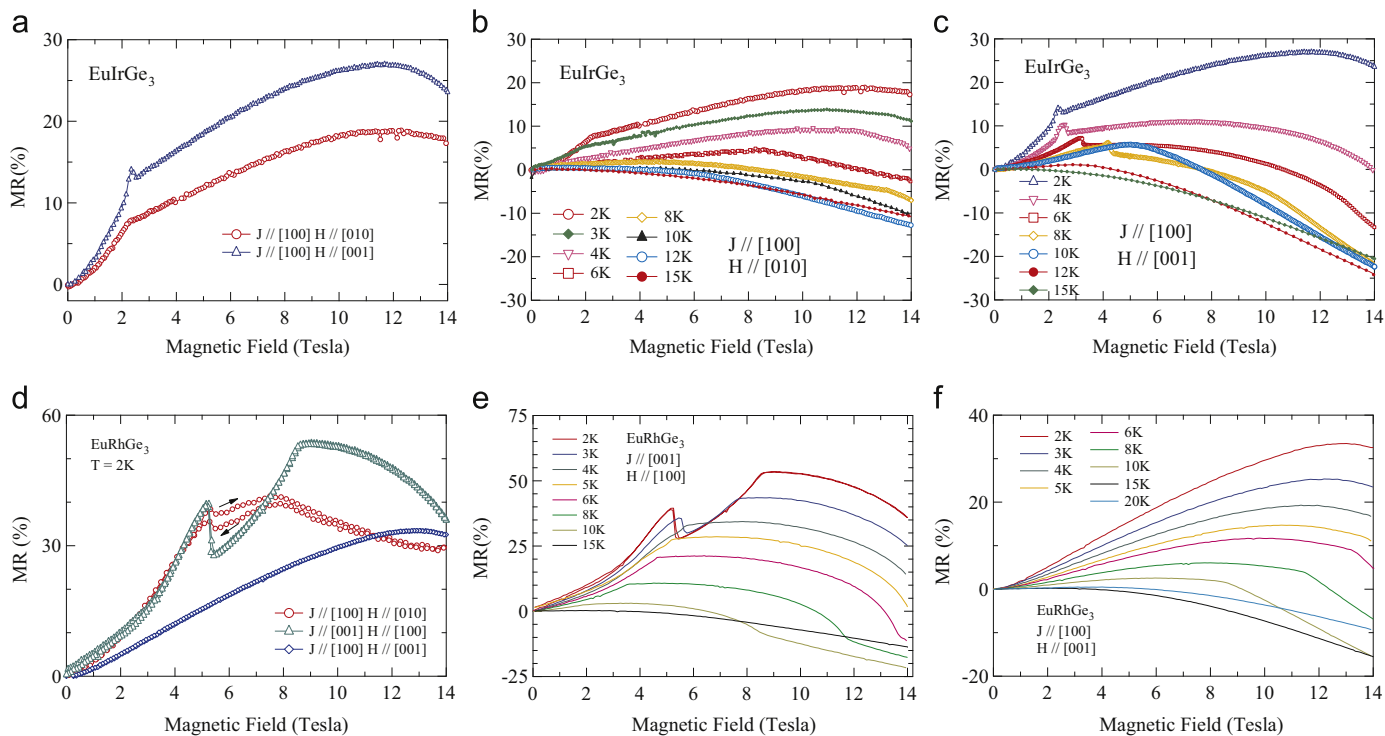


Fig. 11. The variation of magnetoresistance with field at selected temperatures in EuIrGe₃ (upper panels) and EuRhGe₃ (lower panels).

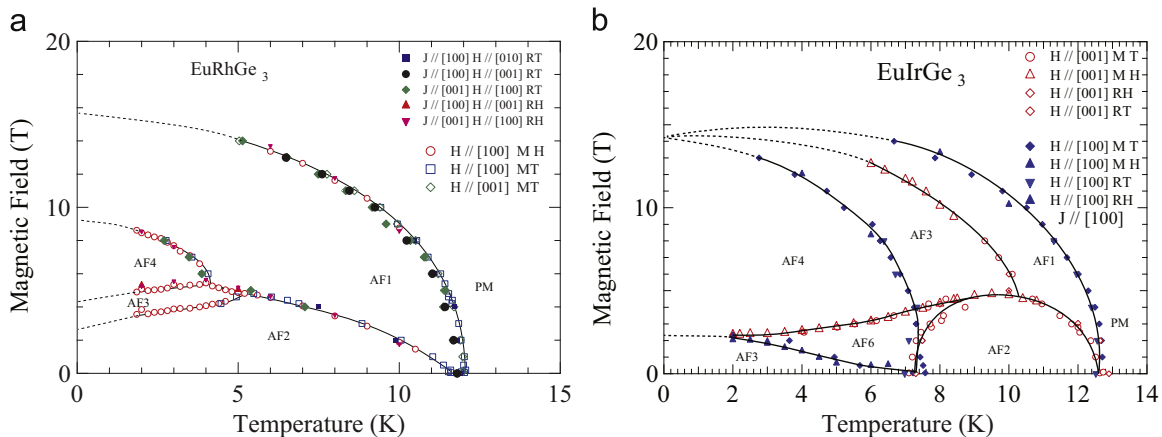


Fig. 12. H – T phase diagram of (a) EuRhGe₃ and (b) EuIrGe₃ constructed from critical points in $M(T, H)$ and $\rho(T, H)$ data. Solid lines are guide to the eye and dotted lines are extrapolation.

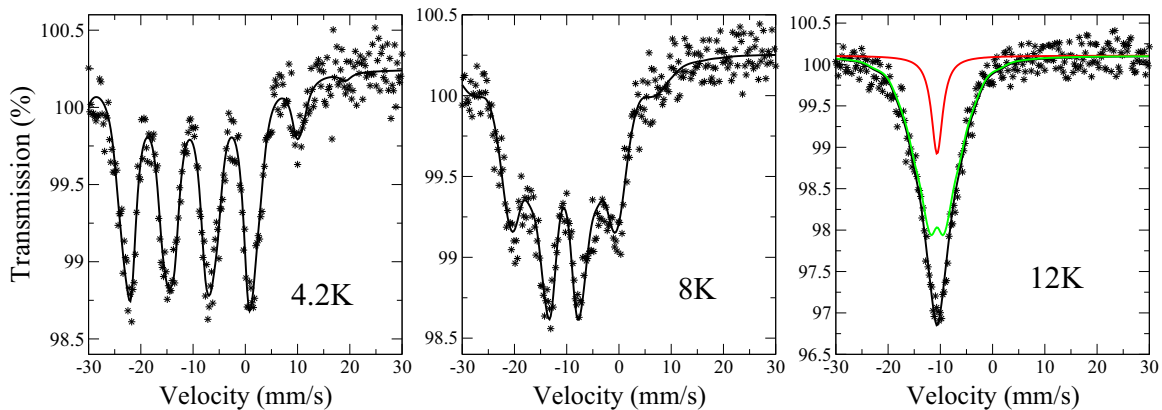


Fig. 13. ¹⁵¹Eu Mössbauer spectra at selected temperatures in EuIrGe₃, in the equal moment phase (4.2 K), close to the equal moment incommensurate transition (8 K) and close to the incommensurate-paramagnetic transition (12 K). At 12 K, the green subspectrum represents the incommensurate pattern and the red subspectrum the paramagnetic pattern. (For interpretation of the references to color in this figure caption, the reader is referred to the web version of this paper.)

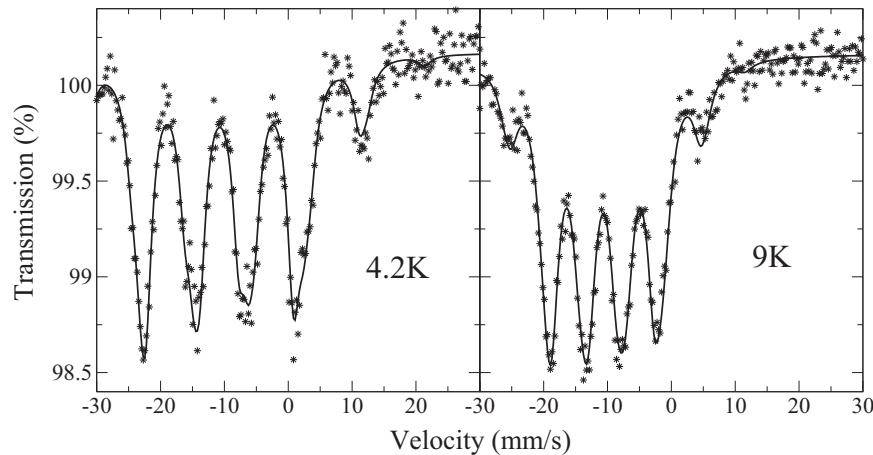


Fig. 14. ^{151}Eu Mössbauer spectra in EuRhGe_3 at 4.2 K and 9 K fitted to a single hyperfine field pattern.

points to a probable first order character of the paramagnetic to incommensurate magnetic phase transition. For EuRhGe_3 , the Mössbauer spectra recorded at 4.2 and 9 K, shown in Fig. 14, are single hyperfine field patterns, with values respectively 29.8 T and 20.7 T. So, in agreement with the previous thermodynamic measurements, EuRhGe_3 presents a single magnetic transition.

Of the five compounds EuPtSi_3 , EuNiGe_3 , EuIrGe_3 , EuPtGe_3 and EuRhGe_3 in which the ^{151}Eu Mössbauer data have been taken, the first three show a cascade of magnetic transitions, the intermediate phase being amplitude modulated. In EuPtGe_3 , where the magnetisation is rather isotropic, no intermediate phase is present and it was conjectured that multiple transitions may be linked to anisotropy [5]. The isothermal magnetisation of EuIrGe_3 is rather similar to that of EuPtGe_3 , yet it shows a cascade of transitions. This shows that other factors are important.

Neutron diffraction studies, now relatively feasible in Eu materials, are clearly required to determine the moment configurations in these compounds. We note that, in addition to the two present germanides, the Pt based EuPtSi_3 and EuPtGe_3 materials should show a spiral magnetic structure in their equal moment phase, as witnessed by the dips in their magnetisation curves, which had not been interpreted in our previous publications [1,2].

References

- [1] N. Kumar, S.K. Dhar, A. Thamizhavel, P. Bonville, P. Manfrinetti, *Phys. Rev. B* 81 (2010) 144414.
- [2] N. Kumar, P.K. Das, R. Kulkarni, A. Thamizhavel, S.K. Dhar, P. Bonville, *J. Phys.: Condens. Matter* 24 (2012) 036005.
- [3] D. Kaczorowski, B. Belan, R. Gladyshevskii, *Solid State Commun.* 152 (2012) 839.
- [4] R.J. Goetsch, V.K. Anand, D.C. Johnston, *Phys. Rev. B* 87 (2012) 064406.
- [5] A. Maurya, P. Bonville, A. Thamizhavel, S.K. Dhar, *J. Phys.: Condens. Matter* 26 (2014) 216001.
- [6] Oleksandr Bednarchuk, Anna Gagor, Dariusz Kaczorowski, *J. Alloys Compd.* 622 (2015) 432.
- [7] Juan Rodriguez-Carvajal, *Physica B* 192 (1993) 55.
- [8] G. Venturini, M. Méot-Meyer, B. Malaman, B. Roques, *J. Less-Common Met.* 113 (1985) 197.
- [9] A. Herpin, in: *Théorie du magnétisme*, Presses Universitaires de France, Paris, 1968.
- [10] J.A. Blanco, D. Gignoux, D. Schmitt, *Phys. Rev. B* 43 (1991) 13145.
- [11] N. Kumar, P.K. Das, N. Kumar, R. Kulkarni, S.K. Dhar, A. Thamizhavel, *J. Phys.: Condens. Matter* 24 (2012) 146003.
- [12] V.K. Anand, D.C. Johnston, *J. Phys.: Condens. Matter* 26 (2012) 286002.
- [13] Z. Hossain, C. Geibel, N. Senthilkumaran, M. Deppe, M. Baenitz, F. Schiller, S. L. Molodtsov, *Phys. Rev. B* 69 (2004) 014422.