

# Angle-resolved magnetoresistance in strongly anisotropic quantum magnet $\text{TmB}_4$

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Precise angle-resolved magneto-resistance (ARMR) measurements in various magnetic fields enabled to create illustrative distributions of  $\Delta\rho/\rho(\varphi, H)$  in  $\text{TmB}_4$ , where  $\varphi$  is the angle between the sample  $c$  axis and applied magnetic field  $H$ . These distributions reveal the charge transport anisotropy in this strongly Ising anisotropic quantum antiferromagnet with a geometrically frustrated Shastry-Sutherland lattice exhibiting fractional magnetization plateaus. While in the paramagnetic region  $\Delta\rho/\rho(\varphi, H)$  reaches its maxima for  $H \perp c$ , below Néel temperature  $T_N = 11.7$  K is the situation different. Here the main MR features appear for  $H \parallel c$ , i.e. along the easy axis of magnetic anisotropy, and correspond to magnetic phases and phase transitions between them. Expressive are above all the features (maxima) related with the scattering of conduction electrons on spin magnetic structure related with fractional magnetization plateaus. With increasing  $\varphi$  MR anomalies shift to higher fields. Above the field of magnetic saturation, moreover, significant MR maxima have been observed at certain angles which correspond to specific directions in the crystal lattice, what is pointing to field directions in which the scattering of conduction electrons on the magnetic structure is the highest. Thus, ARMR appears to be a sensitive experimental tool reflecting the angular dependence of the interplay between charge carriers and magnetic structure as a function of temperature and applied magnetic field.

## I. INTRODUCTION

Properties of quantum spins with antiferromagnetic (AF) coupling on frustrated lattices have attracted widespread interest in recent years due to the discovery of a variety of new quantum ground states as e.g. spin ice [1], quantum spin liquid-like states [2], and fractional magnetization plateaus on the Shastry-Sutherland lattice (SSL) [3], among them the insulating  $\text{SrCu}_2(\text{BO}_3)_2$  [4] as well as the family of metallic rare earth (RE) tetraborides,  $\text{REB}_4$  [5-17]. In case of RE tetraborides, even if they crystallize in the same frustrated magnetic lattice as  $\text{SrCu}_2(\text{BO}_3)_2$ , the phase diagrams of  $\text{REB}_4$  magnetic compounds show different properties. In insulating  $\text{SrCu}_2(\text{BO}_3)_2$  the exchange interaction is of the Heisenberg type, while on the other hand in metallic  $\text{REB}_4$  magnets the AF exchange interaction between their magnetic moments is of long-range Ruderman-Kittel-Kasuya-Yosida (RKKY) type mediated by conduction electrons. In such systems also the transport properties can be strongly influenced by the magnetic structure. This interplay between charge carriers and magnetism therefore allows the use of transport experiments as an indirect probe of the magnetic anisotropy that is present both in the paramagnetic and magnetically ordered phases in these model systems.

Thulium tetraboride  $\text{TmB}_4$ , probably is the most investigated compound of this series. It orders antiferromagnetically at  $T_N = 11.7$  K and has attracted attention for its rich magnetic phase diagram which is strongly biased by crystal field effects at  $\text{Tm}^{3+}$  ion sites that lift the degeneracy of the  $J = 6$  multiplet and lead to a  $M_J = \pm 6$  ground state doublet [5, 18-22].  $\text{TmB}_4$  thus exhibits strong Ising anisotropy where the magnetic field of saturation along the  $c$  axis is at least 10 times smaller than in the perpendicular  $a$ - $b$  plane. The most distinctive features of magnetization  $M$  along the  $c$  axis are various fractional magnetization plateaus which depend in applied field  $H$ . A wide main plateau with  $M/M_{\text{SAT}} = 1/2$  arises in fields between about 17.5 kOe and 36 kOe, and narrow fractional plateaus with  $M/M_{\text{SAT}} = 1/11, 1/9, 1/8$  and  $1/7$  in fields between about 14 kOe and 17.5 kOe (see e.g. [5, 23]). At low fields, below about 14 kOe, the antiferromagnetic ground state phase consists of an AF arrangement of thulium ion dimers, where the dimer spins are ferromagnetic (see Fig. 1). Very recent results concerning the properties of fractional plateaus based mainly on magnetization and heat capacity measurements can be found in [24-26].

Nevertheless, in electrically conducting  $\text{TmB}_4$ , as mentioned above, also conducting electrons can provide additional information about various magnetic states. Such

information, based on resistivity, magnetoresistance (MR) and Hall effect measurements, were recently obtained in [22, 27, 28]. They show that electronic transport as a function of temperature and magnetic field is a very sensitive probe of scattering processes on the magnetic order / disorder in these frustrated systems.

Here, we present a detailed study of angle-resolved magneto-resistance (ARMR) measurements that allow the construction of a complex  $\Delta\rho/\rho(\varphi, H)$  mapping which provides information on scattering processes of charge carriers in various magnetic phases of  $\text{TmB}_4$  when the applied magnetic field changes its magnitude  $H$  and orientation  $\varphi$  (see Fig. 1), and identify field directions where the scattering due to the magnetic structure is highest.

## II. EXPERIMENTAL

The single crystalline  $\text{TmB}_4$  samples were grown by an inductive, crucible-free zone melting method, with residual resistivity ratio larger than  $\approx 14$ , documenting their high quality. All samples were cut from one large, oriented single crystal. More information about sample preparation can be found e.g. in [29].

Magnetoresistance  $\Delta\rho/\rho(\varphi, H) = [\rho(\varphi, H) - \rho(\varphi, H=0)]/\rho(\varphi, H=0)$  measurements were performed using a standard low-frequency ac technique in a commercial PPMS unit equipped with a sample rotation option. The current  $I = 5$  mA was applied along the [110] direction of the Shastry-Sutherland ( $a$ - $b$ ) plane. The sample orientation to the external magnetic field  $H$  direction was changed with a stepsize of  $\Delta\varphi = 1^\circ$  from [001] via  $[\bar{1}11]$  and  $[\bar{1}10]$  to  $[00\bar{1}]$ . For this orientation there is always  $H \perp I$ , but the field can change its orientation from perpendicular to parallel alignment to the Shastry-Sutherland plane (for details see Fig. 1) with the rotation.

Since the magnetization of fractional magnetization plateaus depends on field history [22, 26], measuring protocols were developed that reproduced the same starting points for subsequent magnetoresistance or magnetization measurements below  $T_N$  when measurements were repeated.

## III. RESULTS AND DISCUSSION

Before presenting our experimental results, it is useful to elucidate the effectiveness of ARMAR as a tool for anisotropy study in correlated systems. In uncorrelated compounds, magnetoresistance usually originates from the bending of the carrier trajectory in applied magnetic field. For systems with strong anisotropic behavior in magnetic field, the scattering of conduction electrons can reveal a rich intrinsic structure. The transport properties can be strongly influenced by the magnetic structure and this is reflected in very sharp changes of the MR at specific angles between applied magnetic field and crystallographic

orientation of the system. However, for a successful application of ARMAR high-quality single crystalline samples are inevitable, too.

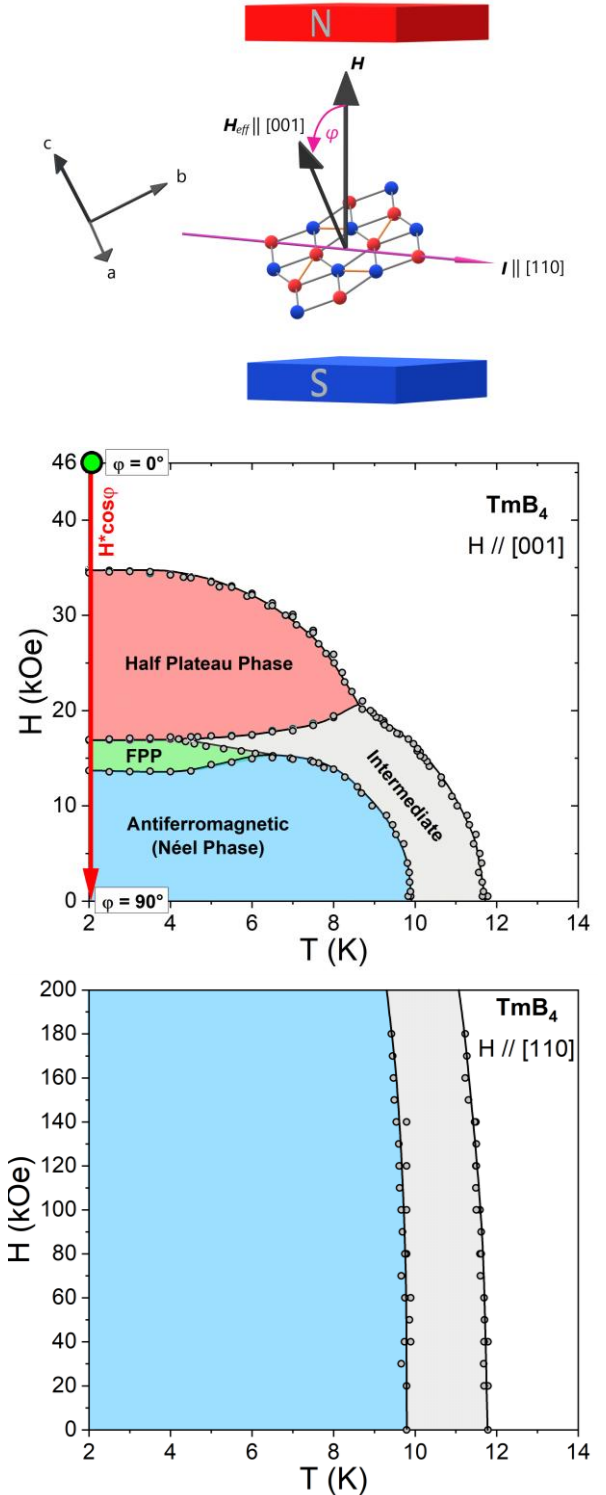


Fig. 1. Top – schematic view of angle-resolved magneto-resistance (ARMAR) measurements in  $\text{TmB}_4$ . During measurements the electrical current  $I$  flows along crystallographic direction [110]. The orientation of applied

magnetic field  $H$ , which is always perpendicular to  $I$ , changes from direction  $[001]$  via  $[\bar{1}11]$  and  $[\bar{1}10]$  to direction  $[00\bar{1}]$ , when  $\varphi$  is changed from  $0^\circ$  to  $180^\circ$ . Spin configuration of the antiferromagnetic ground state of the Shastry-Sutherland lattice in the  $a$ - $b$  plane is displayed by red (spin “up”) and blue (spin “down”) spheres. Middle – phase diagram of  $\text{TmB}_4$  for  $H \parallel c$ . The red arrow indicate the change of effective magnetic field,  $H_{\text{eff}} = H \cos \varphi$ , along  $c$ -axes during sample rotation in field  $H$  from  $\varphi = 0^\circ$  to  $90^\circ$ , which corresponds to the transition of sample from the saturation phase (the big green point at 2 K and  $H_{\text{eff}} = 46$  kOe) to antiferromagnetic Néel phase (the end of red arrow at 2 K and  $H_{\text{eff}} = 0$  kOe). Bottom – phase diagram of  $\text{TmB}_4$  for  $H \parallel [110]$ .

A very good example is  $\text{TmB}_4$ , where the main MR features appear for  $H \parallel c$ , i.e. along the easy axis of magnetic anisotropy, and correspond to magnetic phases and phase transitions between them. Results of the angular dependence of magnetoresistance  $\Delta\rho/\rho(\varphi, H)$  in magnetic field up to 46 kOe and at temperatures of 13 K and 2 K are shown in Fig. 2. During these measurements, as mentioned and shown above, the current  $I$  flowed in direction  $[110]$  and the magnetic field (always  $H \perp I$ ) varied between principal crystallographic directions  $[001]$ ,  $[\bar{1}11]$  and  $[\bar{1}10]$  (see Fig. 1). Thus, the MR measurements covered the full anisotropy of the tetragonal  $\text{TmB}_4$  lattice. From Fig. 2a it can be seen that at 13 K (above  $T_N$ ) the angular dependence of  $\Delta\rho/\rho(\varphi, H)$  exhibits only basic features of the  $\text{TmB}_4$  magnetic anisotropy: The MR reaches minima for  $H \parallel c$ , i.e. when  $H$  is parallel to the easy axis of magnetization (in this case  $\varphi = 0^\circ$  and/or  $\varphi = 180^\circ$ ). Maxima are reached, as expected, when  $H \perp c$ , i.e. when the field is perpendicular to the easy axis. In this case the Lorentz force acting on current  $I$  forces it to flow in the direction of the  $c$  axis where the resistivity is higher than in the  $a$ - $b$  plane ( $\rho_c > \rho_{a,b}$ , see e.g. [22]).

On the other hand, at 2 K (far below  $T_N$ ) the  $\Delta\rho/\rho(\varphi, H)$  dependence shows distinct features of the magnetic phase transitions in  $\text{TmB}_4$  as a function of increasing angle  $\varphi$  between magnetic field and the sample  $c$  axis, and as a function of applied field intensity. Moreover, the obtained  $\Delta\rho/\rho(\varphi, H)$  dependence in fields higher than approximately 15 kOe are not symmetric, but exhibit clear features of hysteresis. This hysteresis (asymmetry with respect to  $\varphi = 180^\circ$ ) might be surprising, however, it only reflects the fact that the rotation of strongly anisotropic  $\text{TmB}_4$  in external magnetic field,  $H$ , manifests itself as a  $H_{\text{eff}} \sim H \cos \varphi$  change of the field along the sample  $c$  direction (see Fig. 1 and [30]). Thus, e.g. sample or field rotation between  $0^\circ$  and  $90^\circ$  can be considered as a field lowering and the further angular change from  $90^\circ$  to  $180^\circ$  as a field increase (see Fig. 3). And, also direct  $\rho(H)$  measurements between 46 kOe  $\rightarrow$  0 kOe  $\rightarrow$  - 46 kOe at 2 K show (Fig. 3b) that

during such a field change hysteresis appears in fields where magnetic phase transitions occur.

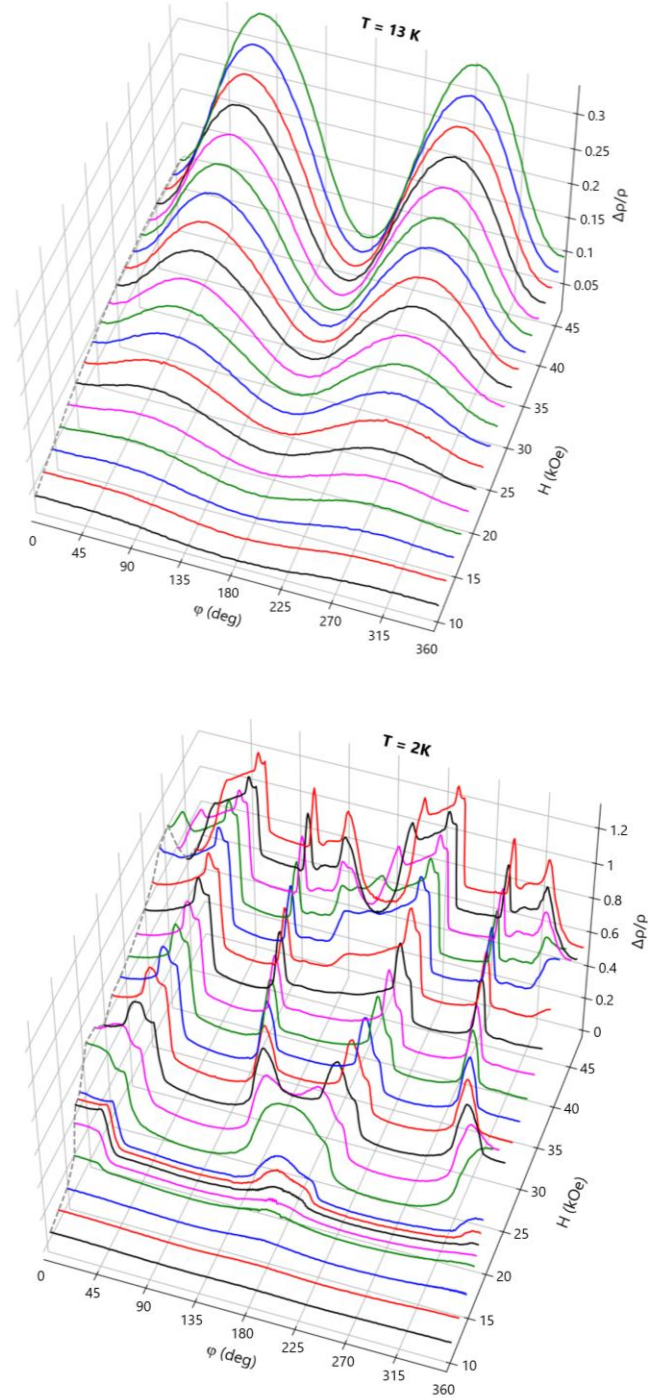


Fig. 2. Angular dependence of magnetoresistance  $\Delta\rho/\rho(\varphi, H)$  of  $\text{TmB}_4$  in various magnetic fields at 13 K (above  $T_N$ ) and 2 K (below  $T_N$ ).  $\varphi$  denotes the angle between the applied field and the  $c$ -axis of sample.

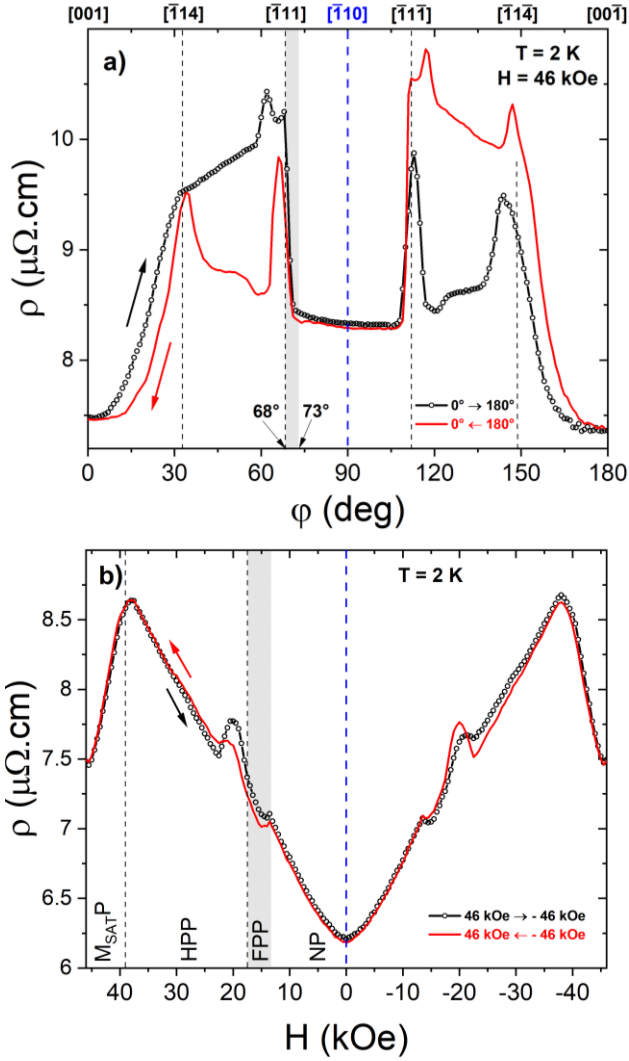


Fig. 3. a) – angular dependence of the resistivity of  $\text{TmB}_4$  at 2 K and in a field of 46 kOe; b) – field dependence of the resistivity  $\rho(H)$  at 2 K. The starting point at 2 K and 46 kOe was reached by cooling the sample from the paramagnetic state at 20 K in field of 46 kOe. Subsequently the field was lowered to -46 kOe and then increased back to 46 kOe. Note that the hysteresis features of  $\rho(H)$  appear on all magnetic phase transitions ( $M_{\text{SAT}}^{\text{P}}$  – Magnetization Saturation Phase, HPP – Half Plateau Phase, FPP – Fractional Plateau Phase, NP – Néel Phase) that involve fractional magnetization plateaus. The shadow area on both graphs represents the fractional plateau phase.

A complete view of the angular dependence at 2 K from Fig. 2 is shown in Fig. 4, where the magnitude of  $\Delta\rho/\rho(\varphi, H)$  is shown in different colors. At 13 K (Fig. 4b), above  $T_N$ , the highest MR is observed, as shown and explained above, for  $H \perp c$  (in this case  $\varphi = 90^\circ$  or  $270^\circ$ ) and the lowest for  $H \parallel c$  ( $\varphi = 0^\circ$  or  $180^\circ$ ).

Below  $T_N$  the situation changes (Fig. 4a). One can see that the angular dependence of the MR is completely different, and that the largest  $\Delta\rho/\rho(\varphi, H)$  anomalies appear for  $H \parallel c$  and correspond to magnetic phases and phase transitions between them (similarly to the  $\rho(H)$  dependence, see Fig. 3b). This seemingly is related to the strong molecular field in the ordered state the origin of which is associated with quantum mechanical exchange interaction. Thus, in the ordered state the magnetic phases dominate the scattering processes of charge carriers. On the other hand, for  $H \perp c$  ( $\varphi = 90^\circ$  and/or  $270^\circ$ ) no such features can be seen. This difference is related to the fact that for  $H \perp c$  the field of magnetic saturation is at least 10 times higher than this for  $H \parallel c$ , and thus in a field of 46 kOe the Néel state still persists.

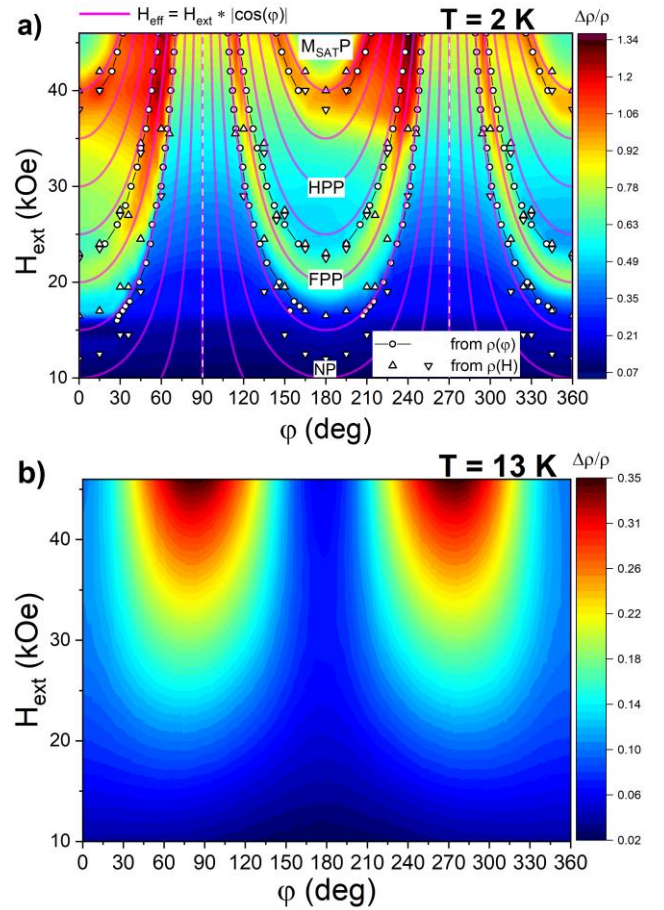


Fig. 4. Magnitude of magnetoresistance as function of angle  $\varphi$  and applied external magnetic field  $H_{\text{ext}}$  for current direction  $I \parallel [110]$  at 2 K and 13 K. Squares and circles represent phase transitions obtained from angular and field dependencies of resistivity ( $\rho(\varphi)$  and  $\rho(H)$ ), respectively. See text for a detailed description.

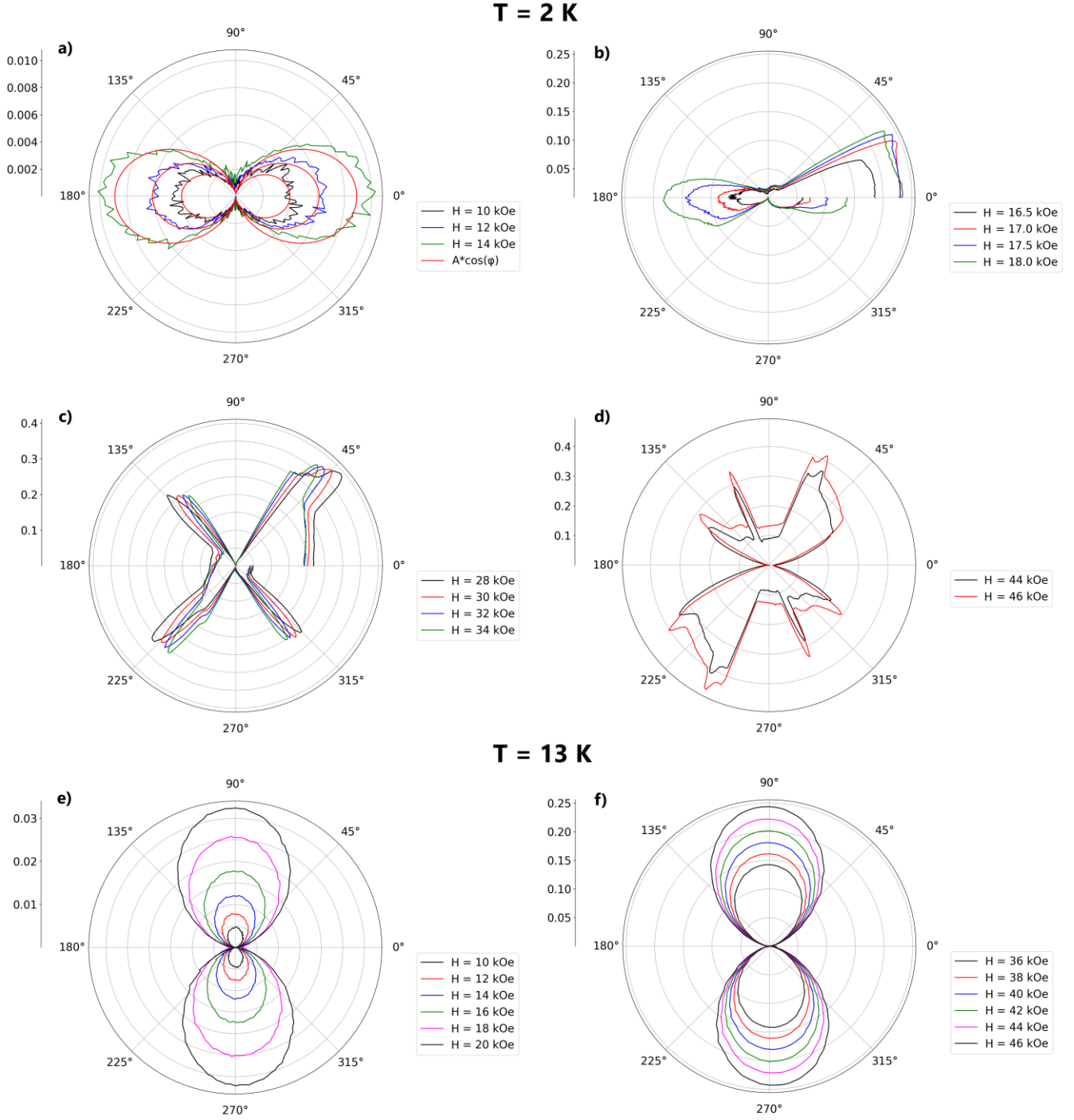


Fig. 5. Polar plots of the anisotropic magnetoresistance parameter  $R_{AMR}$  measured at  $T = 2$  K (a, b, c, d) and  $T = 13$  K (e, f) for various fields  $H$ . The  $R_{AMR}$  shows different shapes of lobes over the full range of  $\varphi$  depending on value of  $H$  i.e. on the position in magnetic phase diagram of  $TmB_4$ : a) - in the antiferromagnetic Néel phase, b) - in the fractional plateau phase, c) - in the half plateau phase and d) - in the saturated magnetization phase (for the location of individual phases in phase diagram see e.g. [26]). One can see that the shape of  $R_{AMR}$  depends on the magnetic phase and exhibits various symmetries (twofold, fourfold, ...). Note moreover, that in both plateau phases also strong features of hysteresis (seen e.g. in [26]) can be observed.

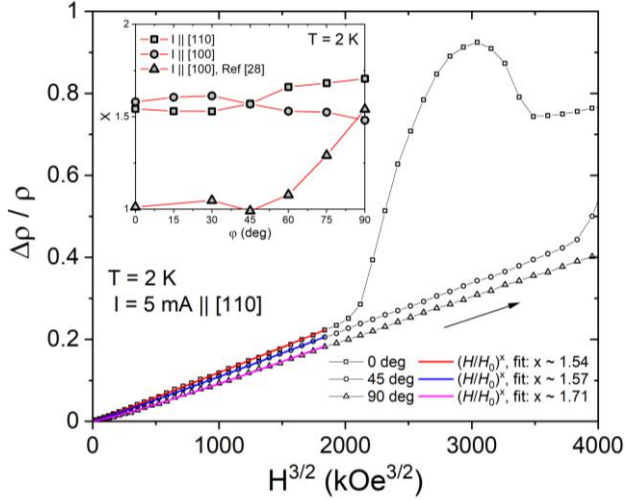


Fig. 6. The evolution of low field dependencies of magnetoresistance  $\Delta\rho/\rho(\varphi, H)$  ( $I \parallel [110]$ ) with increasing angle  $\varphi$  between the applied field  $H$  and the  $c$ -axis, shown in  $\Delta\rho/\rho$  vs.  $H^{3/2}$  representation. Inset shows the variation of exponent  $x$  with angle  $\varphi$  for two orientations of current,  $I \parallel [110]$  and  $I \parallel [100]$  (see [31] for the details).

With increasing angle  $\varphi$  from  $0^\circ$  to  $90^\circ$  the MR maxima shift to higher magnetic fields. The dependence of this shift can be estimated by taking into account the above consideration that upon sample rotation the effective field along the sample  $c$  direction changes as  $H_{\text{eff}} \sim H \cos \varphi$ . Thus, as  $\varphi$  increases (e.g. from  $0^\circ$  to  $90^\circ$ ) higher fields are needed to induce the phase transitions at which MR maxima arise (see also figure S3 in [31]). It can be also seen that the strongest MR features are related to scattering of conduction electrons on the magnetic structure of the fractional plateau phase. The asymmetry of the MR, which can be seen e.g. with respect to  $\varphi = 180^\circ$  (MR to the right of  $\varphi = 180^\circ$  is higher / browner than the MR on the left), can be explained by considering the fact that the rotation of this strongly anisotropic system in magnetic field  $H$  manifests itself as a cosine change of field  $H$  along the sample  $c$  direction (mentioned already above in connection with the field dependence of resistivity in Fig. 3). The obtained precise angle-resolved magneto-resistance  $\Delta\rho/\rho(\varphi, H)$  data set for current direction  $I \parallel [110]$  can be displayed also in the polar presentation  $\Delta\rho/\rho = f(H, \varphi)$  which is shown in figure S1 of [31]. One can see that at 13 K (Fig. 4b) the ARMR distribution is symmetric and reflects in higher fields only the strong Ising magnetic anisotropy in  $\text{TmB}_4$ , which is present also in the paramagnetic phase above  $T_N$ . Below  $T_N$  (see Fig. 4a) the ARMR distribution is symmetric only in small magnetic fields, where magnetic hysteresis is not present. This suggests that ARMR is a rather sensitive experimental tool that provides additional information about the interplay between charge carriers, the various

magnetic phases as a function of temperature, and the applied magnetic field and crystallographic orientations.

A similar ARMR presentation for current direction  $I \parallel [001]$  is shown in figure S2 of [31]. In this case, however, the magnetic field ( $H \perp I$ ) can vary only between crystallographic directions  $[110]$ ,  $[100]$  and  $[1\bar{1}0]$  when  $\varphi$  is changed from  $0^\circ$  to  $90^\circ$ . It can be seen that for this current orientation the MR anomalies are much less pronounced and appear only in field directions, which lay between the directions mentioned above.

The obtained ARMR results are in accordance with our recent investigation of the detailed angular dependence of the rotating magneto-caloric effect (R-MCE), i.e. measurements of the adiabatic change of temperature with changing angle,  $\Delta T(\varphi, T, H)$  [30]. As it was shown [30], with increasing  $\varphi$  is the heating process (due to magnetic reversal) not monotonous, and except the expected peak at  $\varphi = 90^\circ$  (when the sample was rotated from  $H \parallel [001]$  to  $H \perp [001]$  below  $T_N$  in  $H$  above 16 kOe) it exhibits an anomaly also around  $\varphi \approx 60^\circ$ . In the same study [30], angular dependent magnetization measurements at temperatures below  $T_N$  and in various magnetic fields were performed,  $M(\varphi, T, H)$ . It was shown that  $M$  does not anymore exhibit a sinusoidal dependence (as above  $T_N$ ) when the sample is rotated from  $\varphi = 0^\circ$  to  $\varphi = 90^\circ$ , but a rather complicated course, with an anomaly at  $\varphi \approx 65^\circ$  [30]. If we assume that the rotation of an Ising system in magnetic field manifests itself as a  $H_{\text{eff}} \sim H \cos \varphi$  field change along the  $c$ -direction (see Fig. 1) then the observed anomalies in  $\text{TmB}_4$  below  $T_N$  and fields above 16 kOe in  $\Delta T(\varphi)$  at  $\varphi \approx 60^\circ$ , in  $M(\varphi)$  at  $\varphi \approx 65^\circ$  as well as in  $\rho(\varphi)$  at  $\varphi \approx 68^\circ$  (see Fig. 3a) represent the transition to fractional plateau phase. The irreversible behaviour of  $\rho(\varphi)$  as well as of  $\rho(H)$  (see Fig. 3) correspond very well to the fractional magnetization plateaus and to the  $M(H)$  hysteresis, the most distinctive feature of quantum magnet  $\text{TmB}_4$ .

Furthermore, we investigated the strong anisotropy in  $\text{TmB}_4$  by the calculation of anisotropic magnetoresistance parameter  $R_{\text{AMR}}$ , defined as  $R_{\text{AMR}} = (R(\varphi) - R_{\text{min}}) / R_{\text{min}}$ , where  $R(\varphi)$  is the resistance at any  $\varphi$ , measured at a constant  $H$  and  $T$ , and  $R_{\text{min}}$  is the minimum resistance obtained as  $\varphi$  is varied. In Fig. 5, we show the variation of  $R_{\text{AMR}}(\varphi)$  at  $T = 2$  and 13 K, respectively, for various fields. The data can be satisfactorily fitted by a  $\cos(\varphi)$  dependence in low fields (for  $H \leq 14$  kOe in ordered state at 2 K, see Fig. 5a). This dependence, according to [28], suggests a (quasi-)2D Fermi surface, where the MR responds to the perpendicular component of the applied field,  $H \cos(\varphi)$ , and  $R_{\text{AMR}}$  shows a twofold symmetry. In the half plateau phase (Fig. 5c), the twofold symmetry is changed to a fourfold one. The anisotropic MR also suggests anisotropy of the electronic effective mass [28].

To investigate the field dependence of MR in more detail, Fig. 6 shows the  $\Delta\rho/\rho(\varphi, H)$  dependencies for various angles between  $\varphi = 0^\circ$  and  $\varphi = 90^\circ$ . The analysis of the MR angle dependence shows that  $\Delta\rho/\rho(\varphi, H)$  is for all angles positive and that in low magnetic fields, i.e. below  $\approx 15$  kOe (Néel phase), it varies as  $\Delta\rho/\rho = (H/H_0)^x$ , with  $x$  almost independent on  $\varphi$  (see inset of Fig. 6). At  $\varphi = 0^\circ$  is  $x \approx 1.5$ , and at  $\varphi = 90^\circ$  is  $x \approx 1.7$ . In the case of current along  $a$ -axes ( $I \parallel [100]$ ), an approximately  $\Delta\rho/\rho \sim H^{3/2}$  dependence over the entire angle range was observed, too (see figures S5 and S6 in [31]). Obtained results are not in accordance with those in [28] (see inset of Fig. 6 as well as figure S7 in [31]), where MR changes from a linear dependence at  $\varphi = 0^\circ$  to a quadratic one at  $\varphi = 90^\circ$  were observed, and where it was explained by the anisotropy of the Fermi surface topology of TmB<sub>4</sub> [32]. However, it is interesting that the same exponent,  $x = 3/2$ , was obtained for the MR dependence at  $T = 2$  K and  $\varphi = 90^\circ$  (see inset of Fig. 6) in the mentioned study [28].

In addition, a very similar  $H^{3/2}$  behaviour was observed in field dependences of several other borides, like in simple metallic hexaboride LaB<sub>6</sub> [33] or in metallic high-pressure SmB<sub>6</sub> [34] as well as in nonmagnetic metallic dodecaboride LuB<sub>12</sub> [35]. In case of mentioned metallic borides, the authors hypothesize that  $\Delta\rho/\rho \sim H^x$  behaviour with  $x = 1.4 - 1.7$  and without saturation in external magnetic field  $H \parallel [001]$  is due to the magnetic breakdown resulting from topological changes of the orbits, while de Haas-van Alphen measurements confirm the presence of an open orbit/trajectories on the Fermi surface [34-36]. Recent studies of the crystal structure and anisotropy of the MR in LuB<sub>12</sub> [35] revealed the formation of dynamic charge stripes along the [110] directions, which develop due to the dynamic cooperative Jahn-Teller effect on B<sub>24</sub> clusters and are associated with the modulation of the degree of hybridization of  $5d-2p$  band states. It is necessary to emphasize that the filamentary electron density distribution is common for all rare-earth dodecaborides but can probably develop also in other borides containing boron clusters.

The local boron environment of Tm<sup>3+</sup> ions in TmB<sub>4</sub> can be described by B<sub>18</sub> clusters, as it was shown in [32]. Then the loosely bound states of Tm<sup>3+</sup> ions in TmB<sub>4</sub> are very similar to those in TmB<sub>12</sub> with B<sub>24</sub> clusters. It means that the static Jahn-Teller lattice distortion in combination with cooperative dynamic Jahn-Teller instability as well as the  $5d-2p$  hybridization in TmB<sub>4</sub> can play important role in modifying / changing the conduction band of this good metal. In addition,  $5d-2p$  hybridization around the  $Z$  point of Brillouin zone was predicted from the DFT calculation leading to four-hole pockets along  $\Gamma$ - $Z$  and two electron pockets around the  $Z$  point [32]. Calculated nesting of the Fermi surface around  $M$  and  $Z$  points should lead to the appearance of a charge density wave (CDW). We think that the scattering of charge carriers by

the CDW may be the reason for the  $3/2$  field dependence in TmB<sub>4</sub>, as well as in SmB<sub>6</sub> under pressure or LaB<sub>6</sub> in high fields, etc. Nesting of the Fermi surface in metallic SmB<sub>6</sub> just near the metal-insulator transition seems to be natural [33, 34]. Such CDW scattering in rare earth dodecaborides in the temperature range of 2-35 K can be the cause of the field dependence  $\sim H^{1.7}$  [35].

It should be noted that similar ARMR investigations were performed for the first time very recently on dodecaborides HoLuB<sub>12</sub>, HoB<sub>12</sub> and TmB<sub>12</sub> [37-39]. However, those compounds with a face-centred-cubic (*fcc*) crystal structure have a different crystal symmetry and exhibit completely different magnetic structures. It is therefore difficult to compare results in detail.

## IV. CONCLUSIONS

Precise ARMR measurements in various magnetic fields were used to obtain complete mappings of magneto-transport in the strong Ising antiferromagnet TmB<sub>4</sub>. The ARMR maps reveal field and crystalline directions where anomalies (maxima) of conduction electron scattering on the magnetic structure appear. It turns out that in the ordered state large scattering comes from the fractional plateau states. In fields above magnetic saturation significant MR maxima can be observed also at certain angles which correspond to specific directions in the crystal lattice. Thus, ARMR shows to be a sensitive tool that provides additional information about the interplay between charge carriers and various magnetic phases of the system.

## ACKNOWLEDGEMENTS

This work was supported by the Slovak Research and Development Agency under contract no. APVV-17-0020, by the Slovak Scientific Grant Agency under contract VEGA 2/0032/20, as well as projects DAAD-57561069, EU ERDF (European regional development fund) grant no. VA SR ITMS2014+ 313011W856 and European Union's Horizon 2020 Research and Innovation Programme under Grant Agreement No. 824109 (European Microkelvin Platform). Liquid nitrogen for experiments was sponsored by U.S. Steel Košice, s.r.o.

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- [1] A. P. Ramirez, A. Hayashi, R. J. Cava, R. Siddharthan, and B. S. Shastry, *Nature* (London) **399**, 333 (1999).
  - [2] J. S. Helton, K. Matan, M. P. Shores, E. A. Nytko, B. M. Bartlett, Y. Yoshida, Y. Takano, A. Suslov, Y. Qiu, J. H. Chung, D. G. Nocera, and Y. S. Lee, *Phys. Rev. Lett.* **98**, 107204 (2007).
  - [3] B. S. Shastry and B. Sutherland, *Physica* **108 B+C**, 1069 (1981).

- [4] H. Kageyama, K. Yoshimura, R. Stern, N. V. Mushnikov, K. Onizuka, M. Kato, K. Kosuge, C. P. Slichter, T. Goto, and Y. Ueda, *Phys. Rev. Lett.* **82**, 3168 (1999).
- [5] K. Siemensmeyer, E. Wulf, H. J. Mikeska, K. Flachbart, S. Gabani, S. Mat'as, P. Priputen, A. Efdokimova, and N. Shitsevalova, *Phys. Rev. Lett.* **101**, 177201 (2008).
- [6] G. Will, W. Schäfer, F. Pfeifer, F. Elf, and J. Etourneau, *J. Less Common. Met.* **82**, 349 (1981).
- [7] D. Okuyama, T. Matsumura, H. Nakao, and Y. Murakami, *J. Phys. Soc. Jpn.* **74**, 2434 (2005).
- [8] D. Okuyama, T. Matsumura, T. Mouri, N. Ishikawa, K. Ohoyama, and H. Hiraka, *J. Phys. Soc. Jpn.* **77**, 044709 (2008).
- [9] J. Y. Kim, B. K. Cho, and S. H. Han, *J. Appl. Phys.* **105**, 07E116 (2009).
- [10] R. Watanuki, G. Sato, K. Suzuki, M. Ishihara, T. Yanagisawa, Y. Nemoto, and T. Goto, *J. Phys. Soc. Jpn.* **74**, 2169 (2005).
- [11] J. A. Blanco, P. J. Brown, A. Stunault, K. Katsumata, F. Iga, and S. Michimura, *Phys. Rev. B* **73**, 212411 (2006).
- [12] R. Watanuki, T. Kobayashi, R. Noguchi, and K. Suzuki, *J. Phys. Conf. Ser.* **150**, 042229 (2009).
- [13] J. Y. Kim, N. H. Sung, B. Y. Kang, M. S. Kim, B. K. Cho, and J.-S. Rhyee, *J. Appl. Phys.* **107**, 09E111 (2010).
- [14] P. Farkašovský, H. Čenčariková, and S. Mataš, *Phys. Rev. B* **82**, 054409 (2010).
- [15] T. Verkholyak, J. Strečka, *SciPost Phys.* **12**, 056 (2022).
- [16] K. Wierschem, S. S. Sunku, T. Kong, T. Ito, P. C. Canfield, C. Panagopoulos, and P. Sengupta, *Phys. Rev. B* **92**, 214433 (2015).
- [17] S. Gabani, K. Flachbart, K. Siemensmeyer, and T. Mori, *J. Alloys Comp.* **821**, 153201 (2020).
- [18] F. Iga, A. Shigekawa, Y. Hasegawa, S. Michimura, T. Takabatake, S. Yoshii, T. Yamamoto, M. Hagiwara, and K. Kindo, *J. Magn. Magn. Mater.* **310**, e443 (2007).
- [19] S. Gabani, S. Matas, P. Priputen, K. Flachbart, K. Siemensmeyer, E. Wulf, A. Efdokimova, and N. Shitsevalova, *Acta Phys. Pol. A* **113**, 227 (2008).
- [20] S. Michimura, A. Shigekawa, F. Iga, T. Takabatake, and K. Ohoyama, *J. Phys. Soc. Jpn.* **78**, 024707 (2009).
- [21] M. C. Chang and M. F. Yang, *Phys. Rev. B* **79**, 104411 (2009).
- [22] S. S. Sunku, T. Kong, T. Ito, P. C. Canfield, B. S. Shastry, P. Sengupta, and C. Panagopoulos, *Phys. Rev. B* **93**, 174408 (2016).
- [23] S. Gabáni, I. Takáčová, M. Orendáč, G. Pristáš, E. Gažo, K. Siemensmeyer, A. Bogach, N. Sluchanko, N. Shitsevalova, J. Prokleška, V. Sechovský, K. Flachbart, *Solid State Sci.* **105**, 106210 (2020).
- [24] J. Trinh, S. Mitra, C. Panagopoulos, T. Kong, P. C. Canfield, and A. P. Ramirez, *Phys. Rev. Lett.* **121**, 167203 (2018).
- [25] D. Lançon, V. Scagnoli, U. Staub, O. A. Petrenko, M. Ciomaga Hatnean, E. Canevet, R. Sibille, S. Francoual, J. R. L. Mardegan, K. Beauvois, G. Balakrishnan, L. J. Heyderman, Ch. Rüegg, T. Fennell, *Phys. Rev. B* **102**, 060407 (2020).
- [26] Mat. Orendáč, S. Gabáni, P. Farkašovský, E. Gažo, J. Kačmarčík, M. Marcin, G. Pristáš, K. Siemensmeyer, N. Shitsevalova, K. Flachbart, *Sci. Reports* **11**, 6835 (2021).
- [27] L. Ye, T. Suzuki, J. G. Checkelsky, *Phys. Rev. B* **95**, 174405 (2017).
- [28] S. Mitra, J. G. S. Kang, J. Shin, J. Quan Ng, S. S. Sunku, T. Kong, P. C. Canfield, B. S. Shastry, P. Sengupta, and C. Panagopoulos, *Phys. Rev. B* **99**, 045119 (2019).
- [29] N. Shitsevalova, *Crystal Chemistry and Crystal Growth of Rare-Earth Borides*. In: D. S. Inosov, *Rare Earth Borides*, Jenny Stanford Publishing, Singapore 2021.
- [30] Mat. Orendáč, S. Gabáni, E. Gažo, G. Pristáš, N. Shitsevalova, K. Siemensmeyer, K. Flachbart, *Sci. Reports* **8**, 10933 (2018).
- [31] Supplementary Material.
- [32] J. Shin, Z. Schlesinger, B. S. Shastry, *Phys. Rev. B* **95**, 205140 (2017).
- [33] A. J. Arko, G. Crabtree, D. Karim, F. M. Mueller, L. R. Windmiller, J. B. Ketterson, Z. Fisk, *Phys. Rev. B* **13**, 5240 (1976).
- [34] J. C. Cooley, M. C. Aronson, A. Lacerda, Z. Fisk, P. C. Canfield, R. P. Guertin, *Phys. Rev. B* **52**, 7322 (1995).
- [35] K. M. Krasikov, A. N. Azarevich, V. V. Glushkov, S. V. Demishev, A. L. Khoroshilov, A. V. Bogach, N. Yu. Shitsevalova, V. B. Filippov, N. E. Sluchanko, *JETP Letters* **112**, 413 (2020).
- [36] S. Zhang, Q. Wu, Y. Liu, O. V. Yazyev, *Phys. Rev. B* **99** (3), 1 (2019).
- [37] A. L. Khoroshilov, V. N. Krasnorussky, K. M. Krasikov, A. V. Bogach, V. V. Glushkov, S. V. Demishev, N. A. Samarin, V. V. Voronov, N. Yu. Shitsevalova, V. B. Filipov, S. Gabani, K. Flachbart, K. Siemensmeyer, S. Yu. Gavrilkin, N. E. Sluchanko, *Phys. Rev. B* **99**, 174430 (2019).
- [38] K. Krasikov, V. Glushkov, S. Demishev, A. Khoroshilov, A. Bogach, V. Voronov, N. Shitsevalova, V. Filipov, S. Gabani, K. Flachbart, K. Siemensmeyer, N. Sluchanko, *Phys. Rev. B* **102**, 214435 (2020).
- [39] A. Azarevich, V. Glushkov, S. Demishev, A. Bogach, V. Voronov, S. Gavrilkin, N. Shitsevalova, V. Filipov, S. Gabáni, J. Kačmarčík, K. Flachbart, N. Sluchanko, *J. Phys.: Condens. Matter* **34**, 065602 (2022).