Anomalous Hall effect and magnetoresistance in microribbons of the magnetic Weyl semimetal candidate PrRhC₂

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PrRhC₂ belongs to the rare-earth carbides family, whose properties are of special interest among topological semimetals due to the simultaneous breaking of both inversion and time-reversal symmetry. The concomitant absence of both symmetries grants the possibility to tune the Weyl nodes chirality and to enhance topological effects such as the chiral anomaly. In this paper, we report on the synthesis and compare the magnetotransport measurements of polycrystalline PrRhC₂ samples and a single-crystalline PrRhC₂ sample. Using a remarkable and sophisticated technique, the PrRhC₂ single crystal is prepared via focused-ion-beam cutting from the polycrystalline material. Our magnetometric and specific heat analyses reveal a noncollinear antiferromagnetic state below 20 K, as well as short-range magnetic correlations and/or magnetic fluctuations well above the onset of the magnetic transition. The transport measurements on the PrRhC₂ single crystal display an electrical resistivity peak at 3 K and an anomalous Hall effect below 6 K indicative of a net magnetization component in the ordered state. Furthermore, we study the angular variation of magnetoresistivities as a function of the angle between the in-plane magnetic field and the injected electrical current. We find that both the transverse and the longitudinal resistivities exhibit fourfold angular dependencies due to higher-order terms in the resistivity tensor, consistent with the orthorhombic crystal symmetry of PrRhC₂. Our experimental results may be interpreted as features of topological Weyl semimetallic behavior in the magnetotransport properties.

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I. INTRODUCTION

The realization of Weyl fermions in condensed matter systems is of great interest to prove novel topological theories while also having promising applications in the field of quantum computing, spintronics, and electro-optical devices [1-10]. Weyl nodes form a pair by breaking a symmetry such as inversion and/or time reversal. Although the majority of the so-far-discovered Weyl semimetals (WSMs) are of the parity-broken type, many experimental attempts have been performed to find magnetic WSMs that, unlike their nonmagnetic counterparts, combine magnetic order and a topologically nontrivial band character, giving rise to exotic

phenomena such as the intrinsic anomalous Hall effect and axion electrodynamics [11–18]. Among the proposed magnetic WSMs, rare-earth carbides [19,20] break both inversion and time-reversal symmetry due to their noncentrosymmetric point group and magnetic order, respectively [21–25].

The rare-earth carbides represent a promising material class to probe the "chiral anomaly" effect, considered a key fingerprint of topology. In the presence of parallel magnetic and electric fields, the electrons flow between Weyl nodes of opposite chirality thereby inducing a negative quadratic longitudinal magnetoresistance (MR) [26–29]. This chiral imbalance between the Weyl nodes is permitted in materials where both inversion and time-reversal symmetries are absent causing the net chirality picked up by experimental probes to vanish and, hence, making the chiral anomaly effect invisible. In turn, observation of a negative longitudinal MR, whose magnitude strongly depends on the angle between magnetic

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and electric fields, is considered a key observation indicative of Weyl physics via a topologically protected chiral charge [30]. As an alternative description, a negative MR can also be the result of extrinsic mechanisms, such as weak electron localization [31], ionic impurity scattering processes [32], or conductivity fluctuations [33], making the identification of the anomaly highly nontrivial. In semimetals with high mobility, the evidence of a chiral anomaly can also be strongly hindered by the "current jetting" effect that leads to a strong apparent negative longitudinal MR [34,35].

In the class of rare-earth carbide materials, it has been predicted that an appropriate tilt of the magnetization can stabilize an odd number of Weyl nodes near the Fermi surface, giving a nonzero net chirality that can be easily experimentally probed through the chiral anomaly effect [19,20]. The class of rare-earth-transition-metal carbides, which, according to the theory, combine topology and magnetism, includes GdCoC₂, GdNiC₂, NdRhC₂, and PrRhC₂ [19]. Magnetization measurements of single crystals reveal that GdCoC₂ is antiferromagnetic (AFM) below Néel temperature $T_N = 15.6$ K [36] and shows an order-order transition at T = 14.0 K. Reference [37] describes the same material as a ferromagnet with Curie temperature $T_C \simeq 15$ K. On the other hand, GdNiC₂ single crystals, like most of the Ni-based members in this family [38], were proven to reach an antiferromagnetic state below $T_N = 20$ K [36]. Experimental reports on polycrystals of NdRhC2 and PrRhC2 have been limited to their high-temperature susceptibility [39], and single crystals of NdRhC₂ and PrRhC₂ have not been grown, so far. The hightemperature susceptibility measurements suggest that these two compounds possess an AFM ground state with critical temperatures of ~ 13 K and close to 0 K, respectively. Density functional calculations based on the generalized gradient approximation have predicted, however, a ferromagnetic (FM) ground state for NdRhC₂, whereas PrRhC₂ is theoretically found to exhibit either an AFM or FM ground state, depending on the simulation approximations for the two unpaired 4felectrons [19]. Therefore the growth of a PrRhC₂ single crystal is highly desirable to experimentally verify its magnetic structure. Here, we circumvent this experimental challenge by cutting single-crystal microribbons from a multigrain sample using a focused ion beam. In addition to analyzing the magnetic order of the polycrystalline PrRhC2 material through magnetometric measurements, we are therefore able to investigate the magnetotransport in the single crystal. Since for the PrRhC₂ crystal structure only ferromagnetic order and noncollinear antiferromagnetic order [40,41] produce the anomalous Hall effect, this lets us draw conclusions about the type of magnetic order.

II. PREPARATION AND MAGNETIC CHARACTERIZATION OF THE POLYCRYSTAL

Polycrystalline samples are prepared by arc-melting slugs or pieces of >99.9% by mass metals-based pure elements on a water-cooled copper hearth under purified argon. The metal blends are melted three times for homogenization keeping the evaporation losses below 1 wt %. Samples are then individually wrapped in protective Ta foil, sealed in evacuated silica ampoules, and annealed at 1000 °C for

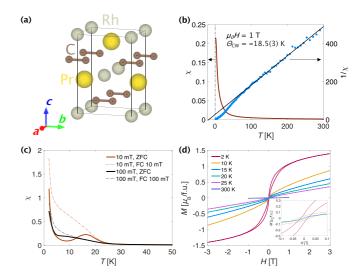


FIG. 1. (a) Crystal structure of PrRhC₂ (orthorhombic symmetry, space group *Amm2*) including the expected carbon dimers. The unit cell is indicated by black solid lines. (b) Magnetic susceptibility (SI units) of PrRhC₂ polycrystal measured while increasing the temperature in the presence of an external magnetic field of 1 T (brown curve) and the reciprocal of the susceptibility vs temperature (blue dots) and a Curie-Weiss (CW) fit to the data (solid black line). This linear fit yields a CW temperatures recorded in different magnetic fields following zero-field-cooled (ZFC) and field-cooled (FC) protocols. (d) Magnetization curves as a function of magnetic field at different temperatures. Inset: Magnetization at low field and low temperatures (2, 10, and 15 K).

1 week followed by quenching the ampoules in cold water. The crystal structure of polycrystalline samples is determined by x-ray powder diffraction analysis (STOE STADI in transmission geometry, CoK α 1 radiation, equipped with a germanium monochromator and a DECTRIS MYTHEN 1K detector). The lattice parameters are obtained from a structural model derived by Rietveld analysis yielding a = 0.36876(1), b = 0.47103(1), c = 0.66124(1) nm, $R_p = 0.0281$, $R_{wp} = 0.0371$, $R_I = 0.0563$, and $\chi^2 = 1.28$. More details can be found in Table S1 in the Supplemental Material [42].

The magnetic structure of the polycrystal is investigated by magnetometry measurements in a superconducting quantum interference device (SQUID) down to 2 K. The magnetic susceptibility χ of PrRhC₂ [Fig. 1(b)], measured at a 1 T field from 2 to 300 K, displays a steep increase below 10 K, indicating a net magnetization at low temperature. The inverse susceptibility above 50 K is well fitted by the Curie-Weiss (CW) law $1/\chi = (T - \Theta_{CW})/C$, where C is related to the effective magnetic moment per unit cell and Θ_{CW} is the CW temperature, connected to the sum of all magnetic interactions in the system. The obtained effective magnetic moment $\mu_{\text{eff}} = 3.7(2) \,\mu_B$ is in agreement with the value expected for Pr^{3+} , 3.58 μ_B [43]. The CW temperature is negative [$\Theta_{CW} = -18.5(3)$ K], indicating AFM interactions. On the other hand, the magnetic susceptibility [Fig. 1(c)], measured from low to high temperature in 10 and 100 mT fields, displays a pronounced zero-field-cooled-field-cooled (ZFC-FC) splitting starting around 24 K. In the FC curve, a

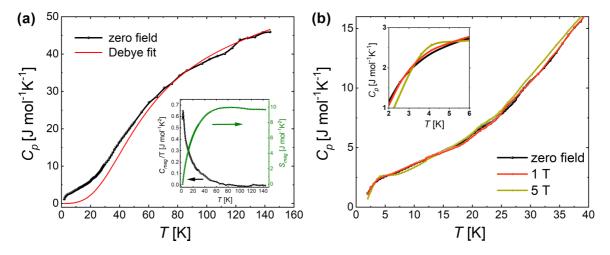


FIG. 2. (a) Zero-field specific heat of $PrRhC_2$. The red curve represents the lattice specific heat obtained using a Debye model; for details, see text. Inset: zero-field magnetic specific heat plotted as C_{mag}/T vs T (left scale) together with the magnetic entropy (right scale). (b) Temperature dependence of the specific heat for applied magnetic fields of 1 and 5 T. The inset shows a zoom of the low-temperature region.

ferromagneticlike transition is observed at $T \approx 22$ K (determined using the second-derivative method, i.e., the transition temperature is defined as the inflection point of the curve). For the ZFC magnetization, a kink between 10 and 20 K, possibly related to a rearrangement of the magnetic structure, is rapidly suppressed by applying an external magnetic field. Figure 1(d) depicts the magnetization as a function of external magnetic fields, at different base temperatures, which indicates the ferromagneticlike nature of the compound, exhibiting clear hysteresis loops with coercive fields smaller than 30 mT below 15 K. Additionally, by increasing the applied magnetic field, the magnetization at the lowest temperature measured, 2 K, approaches a value of 1.4 μ_B , which is below the expected $2 \mu_B$ for the $4f^2$ (J = 4) state, assuming that Rh atoms have no magnetic moment. Altogether, our observations point to a noncollinear, canted AFM ground state with finite net magnetization which, nonetheless, is not fully saturated even up to 7 T (Fig. S2 in the Supplemental Material [42]). This finding makes the presence of magnetic domains unlikely. In ferromagnetic materials with well-organized domains, the magnetization reaches a saturation point where it no longer increases with an increasing magnetic field, corresponding to a single domain phase. Further Kerr microscopy and neutron diffraction experiments are necessary to investigate the saturation field and to understand the microscopic nature of the magnetic ordering. This is, however, out of the scope of this work. It is worth noticing that we observe the same magnetic properties in two additional polycrystalline samples, whose surfaces are treated with CH₃ and HCl, respectively (not shown).

The magnetic behavior of $PrRhC_2$ is further analyzed by specific heat measurements as a function of temperature and applied magnetic field. The measurements are performed on four single crystals using a heat-pulse relaxation method in a Physical Property Measurement System (PPMS) from Quantum Design. In order to obtain the intrinsic specific heat, the temperature- and field-dependent addenda are thoroughly subtracted from the measured data. The temperature-dependent specific heat is shown in Fig. 2. As can be observed in Fig. 2(a), $C_p(T)$ at zero field does not show a clear anomaly associated with the magnetic transition observed in magnetic susceptibility around 22 K: Instead, a subtle bump appears, as well as a broad feature detected below 5 K. In order to better understand this behavior and disentangle the magnetic contribution to the heat capacity, the lattice contribution is estimated using the following Debye model:

$$C_{\rm ph} = 9R \sum_{i=1}^{2} n_i \left(\frac{T}{\Theta_{\rm Di}}\right)^3 \int_0^{\Theta_{\rm Di}/T} \frac{x^4 e^x}{(e^x - 1)^2} dx, \qquad (1)$$

where R is the universal gas constant and the index i sums over two different Debye temperatures Θ_{Di} , each one representing a part of the total unit cell formed by n = 4 atoms ($n_1 = 2$) and $n_2 = 2$). The values $\Theta_{D1} = 240$ K and $\Theta_{D2} = 1540$ K are determined as the best parameters. The total estimated lattice contribution is shown by the red curve in Fig. 2(a). By subtracting the lattice contribution thus calculated from the experimental data, the magnetic contribution to the specific heat is obtained, which is plotted as C_{mag}/T as a function of temperature in the inset of Fig. 2(a) (left axis). The magnetic entropy S_{mag} is evaluated by integrating C_{mag}/T [Fig. 2(a), right axis]. An entropy release is observed up to $T \sim 70$ K, followed by a plateau with $S_{\text{mag}} \approx 9.6$ J mol⁻¹ K⁻¹. Despite the limits of our phononic background, this value is close to the expected value for an S = 1 ground state, $(R \ln (2S + 1)) =$ 9.1 J mol⁻¹ K⁻¹, probably overestimated by the nonperfect estimation of the lattice contribution. Most significant is the fact that magnetic fluctuations and/or short-range correlations seem to be present well above $T \approx 22$ K, and that the entropy related to the magnetic order is gradually released over a wide temperature range. Last but not least, the temperature dependence of the specific heat under applied magnetic field is shown in Fig. 2(b). While the subtle bump around 22 K does not change noticeably, the broad feature below 5 K slightly shifts to higher temperature, consistent with a robust ferromagneticlike character of the canted AFM moments ordering at low temperature.

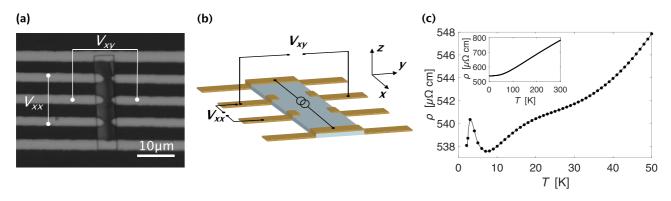


FIG. 3. (a) Close-up microscopy image of the device. The lamella is obtained with a FIB cut, and electrical contacts are defined by standard optical lithography. (b) Schematic of the device with the coordinate system. (c) Electrical resistivity measured in a four-point geometry at low temperatures (main panel) and in the full range of temperatures (inset).

III. PREPARATION AND MAGNETOTRANSPORT OF THE SINGLE CRYSTAL

For magnetotransport studies, a single crystal of 28 μ m \times $6 \,\mu\text{m} \times 300 \,\text{nm}$ size is mechanically extracted from the ascast alloy, and the relative crystal structure is determined using single-crystal x-ray diffraction, which confirms the orthorhombic space group Amm2 phase with lattice parameters consistent with the ones from the polycrystal. The microstructure and chemical composition of single-crystalline and polycrystalline samples are analyzed by scanning electron microscopy (SEM) on a ZEISS EVO MA 15 equipped with an energy-dispersive x-ray (EDX) detector operated at 30 kV. Carbon is a rather light element; so any quantification with EDX is not reliable. Therefore only the Pr and Rh compositions are determined at several positions of each sample [Pr 50.2(4), Rh 49.8(4) at. %] confirming their 1 : 1 ratio in agreement with the PrRhC2 composition. For cutting the microribbon from the bulk material, we utilize a focused ion beam (FIB) of Ga_{2+} ions. A typical position to yield a single crystal is indicated in Fig. S1 in the Supplemental Material [42]. After FIB preparation, the ribbons are then lifted out of the crystal and transferred to the substrate for further device preparation. Details on the FIB preparation procedure can be found in Ref. [44].

The electrical contacts are defined by the optical lithography and lift-off process with 300 nm of sputtered Au on a thin Cr adhesive layer [Fig. 3(a)]. Figure 3(b) shows the device layout with the electrical connections and the coordinate system. Figure 3(c) reports the electrical resistivity ρ as a function of temperature measured using a four-point geometry. This data set shows a metallic behavior with a hump around 20 K, followed by a peak at 3 K. These features go along with those observed in the magnetic susceptibility and specific heat measurements. The local increase of the electrical resistivity at low temperature could be attributed to the enhanced carrier scattering by spin fluctuation due to rearrangement of the magnetic structure [45].

To quantify the MR ratio, defined as (R(H) - R(0))/R(0), and the Hall effect, a small dc current of $I = 50 \mu A$ along the x direction is injected, and the longitudinal (V_{xx}) and transversal (V_{yx}) voltages are simultaneously measured in an out-of-plane magnetic field sweep. The longitudinal resistivity corresponds to the diagonal components of the resistivity tensor, and their variation in a magnetic field (magnetoresistivity) is typically an even function of the magnetic field. The transversal signal corresponds to the off-diagonal components of the resistivity tensor, which are typically odd under magnetic field reversal. In our measured signals we have contributions from both longitudinal and transversal voltage due to the small misalignment between the electrical contacts on the PrRhC₂ lamella [Fig. 3(a)]. When an electrical current is applied along the high-symmetry direction, we can isolate the longitudinal and transversal signals, separating the measured signal into symmetric and antisymmetric components, respectively. The MR traces measured in an out-of-plane magnetic field at different base temperatures are displayed in Fig. 4(a). The negative MR at low temperature suggests a weak localization effect, a phenomenon that originates from the constructive interference of backscattered electronic wave functions, which increases the probability of localizing the electron [46]. The coherent superposition of the wave functions is destroyed by applying a magnetic field, resulting in a decrease of the resistivity and thus a negative MR. This effect disappears in our sample above 30 K, due to the dominant incoherent scattering at higher temperatures. The positive MR with the characteristic quadratic dependency on the external field above 30 K can be explained by the Lorentz force acting on the carrier motion. The absolute transversal resistivity (ρ_{vx}) curves versus the external magnetic field, taken at very low temperatures, are shown in Fig. 4(b), after subtracting the linear ordinary Hall background with a positive slope of 1.06 $\mu\Omega$ cm T⁻¹. Below 6 K, an anomalous contribution to the Hall signal is visible around zero magnetic field, consistent with the temperature dependence of electrical resistivity data [Fig. 3(c)]. Assuming a single-band model, from the linear region of the ρ_{yx} at high magnetic fields of Fig. S3 in the Supplemental Material [42], we estimate the carrier concentration $n_0 = 1.32 \times 10^{21} \text{ cm}^{-3}$ (electron type) and the carrier mobility $\mu = 5.7 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ at 300 K.

In order to study the in-plane anisotropy, we perform azimuthal-angle-dependent magnetoresistance measurements at 2 K under an in-plane magnetic field with constant amplitude of 5 T. The azimuthal angular variation of longitudinal resistivity shown in Fig. 5(a) can be well approximated with $\rho_{xx} = C_0 + C_2 \cos^2(\phi - \phi_0) + C_4 \cos^4(\phi - \phi_0)$

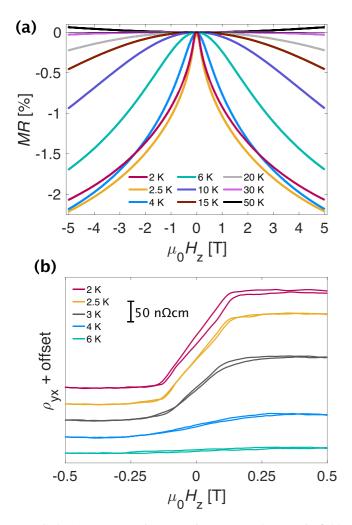


FIG. 4. (a) Magnetoresistance ratio vs external magnetic field at several temperatures. (b) Transversal resistivity as a function of magnetic field at some low temperatures. The linear background associated with the ordinary Hall effect is subtracted, and an offset is added for better visualization.

[47]. This equation is a modified version of a well-known phenomenological model for anisotropic magnetoresistance in magnetic systems. The modified equation includes a misalignment angle ϕ_0 and a fourfold term, both of which have been previously introduced in studying anisotropic magnetoresistance (AMR) in other systems arising from the crystal symmetry [48] or heterointerfaces [47,49,50]. The misalignment angle ϕ_0 can be attributed to the misalignment between magnetization and the external field, as well as the orientation of the crystallographic axis with respect to the transport measurement geometry. Typically, in the absence of crystalline anisotropy, rotating the magnetic field within the two-dimensional (2D) plane leads to a conventional twofold-symmetric form for the in-plane anisotropic magnetoresistance as a function of the angle between the magnetic field and the current directions. Our magnetotransport measurement on a single crystal reveals fourfold-symmetric contributions, which are consistent and also reflect the underlying crystalline symmetry of the material [47,51,52]. Especially at the intermediate-field limit, a very intricate

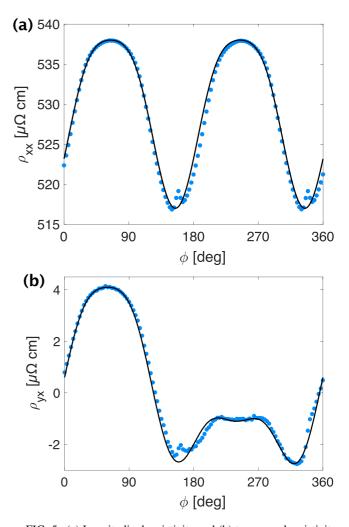


FIG. 5. (a) Longitudinal resistivity and (b) transversal resistivity in the presence of an in-plane 5 T field as a function of the angle between the current line and the magnetic field direction at 2 K. The solid black curves are fits based on the phenomenological relations given in the text for ρ_{xx} and ρ_{yx} . The fitted values for the parameters of longitudinal signal are $C_0 = 521$, $C_2 = 21$, $C_4 = -4$, and $\phi_0 = 65^\circ$. For the transverse resistivity the fitting coefficients are $C'_1 = 2.6$, $C'_2 = 2.0$, $C'_4 = 0.5$, $\phi'_0 = 14.5^\circ$, and $\theta_{tilt} = 30^\circ$. The unit of all coefficients C_i , and C'_i is $\mu\Omega$ cm, the same as the resistivities.

dependence of the MR as a function of the angle between magnetization and current with respect to a reference crystallographic axis has been reported even for simple cubic lattices [53,54]. In the present case, the minimal model given above sufficiently mimics the observed AMR, as illustrated by the experimental data. Turning to the transversal signal, we observe transversal AMR, sometimes called the planar Hall effect (PHE), together with a cosinelike function as shown in Fig. 5(b). The PHE was originally proposed and measured in ferromagnetic thin films, where the Hall voltage scales quadratically with the in-plane magnetization as $\rho_{\rm vx} \propto M_{\parallel}^2 \sin 2\phi_{M,I}$, with $\phi_{M,I}$ representing the angle between the in-plane magnetization and the applied current [49,55,56]. Recently, a very strong PHE has been observed in Weyl semimetals due to the chiral anomaly in these systems [57,58]. In this case, the Hall conductivity depends quadratically on the in-plane magnetic field, and its angular dependence is determined by the angle between the current and the applied in-plane magnetic field. Hence a strong PHE may be observed (i) in FM films with in-plane magnetization and (ii) in Weyl semimetals caused by the chiral anomaly. PrRhC2 is a promising magnetic Weyl semimetal and, due to its canted AFM order, has a FM component with in-plane magnetization. Another complication arises from the role of magnetization in tailoring the Weyl cones [19], which may cause linear terms in the PHE proportional to $B\sin(\phi - \theta_{\text{tilt}})$ as predicted in Ref. [59]. This quantity could also be given by the ordinary Hall effect resulting from a finite out-of-plane component of the magnetic field and/or due to the unknown crystalline orientation, which might produce any term of the angledependent transverse resistivity. Taking all the effects into account, we can consider a phenomenological form for the transverse resistivity, $\rho_{yx} = C'_1 \sin(\phi + \theta_{\text{tilt}}) + C'_2 \sin[2(\phi - \theta_{\text{tilt}})]$ $\phi'_0)] + C'_4 \cos [4(\phi - \phi'_0)]$, that provides a good agreement with the measured resistivity, as illustrated in Fig. 5(b). The angles θ_{tilt} and ϕ'_0 account for asymmetries resulting from misalignments between the current direction and the crystallographic orientation of the material. More specifically, θ_{tilt} can be interpreted as the direction of Weyl nodes tilting relative to the device orientation, while ϕ'_0 represents the angle between the current direction and the principal in-plane crystal axis. For the quantified $C'_1 = 2.6 \ \Omega \text{ cm}$, an out-of-plane misalignment of 30° would be necessary for the ordinary Hall effect to be the origin of the linear term, and therefore this scenario can safely be discarded. Excluding the relative reduced AMR $(C'_2 = 2.0 \ \Omega \ \text{cm} < C'_1)$ and a misalignment of the sample, we assume the contribution to the transverse resistivity to be caused by the tilted Weyl cones. We emphasize that we are here dealing with a phenomenological model. Indeed, a more elaborate theoretical analysis is needed to understand the origin of different terms and their connection to the

topological Weyl phase of the system along with its magnetic properties.

IV. CONCLUSIONS

This study investigates the magnetic order of a polycrystal along with thermal and magnetotransport properties in PrRhC₂ single crystals. Our results strongly suggest a noncollinear antiferromagnetic state with a robust net magnetization at low temperature. This ferromagneticlike behavior is also confirmed by our magnetotransport study, which shows a resistivity peak around 3 K and a significant anomalous Hall signal. We also observe azimuthal angular variations in magnetoresistance, indicating the presence of higher-order terms in the resistance tensor and potentially Weyl phase character of the material. Our findings highlight the potential of PrRhC₂, as a unique material to study the interplay between magnetic and topological semimetallic properties, particularly due to the breaking of both time-reversal and inversion symmetries.

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