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Magnetic and electronic structure of Dirac semimetal candidate EuMnSb_2

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We report an experimental study of the magnetic order and electronic structure and transport of the layered pnictide EuMnSb_2 , performed using neutron diffraction, angle-resolved photoemission spectroscopy (ARPES), and magnetotransport measurements. We find that the Eu and Mn sublattices display antiferromagnetic (AFM) order below $T_N^{\text{Eu}} = 21(1)$ K and $T_N^{\text{Mn}} = 350(2)$ K, respectively. The former can be described by an A-type AFM structure with the Eu spins aligned along the c axis (an in-plane direction), whereas the latter has a C-type AFM structure with Mn moments along the a -axis (perpendicular to the layers). The ARPES spectra reveal Dirac-like linearly dispersing bands near the Fermi energy. Furthermore, our magnetotransport measurements show strongly anisotropic magnetoresistance and indicate that the Eu sublattice is intimately coupled to conduction electron states near the Dirac point.

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I. INTRODUCTION

Topological semimetals can host quasiparticle excitations which masquerade as massless fermions due to the linearly dispersing electronic bands created by interactions with the crystal lattice. The Dirac or Weyl nodes, where the conduction and valence bands meet in momentum space, are robust against small perturbations due to the protection afforded by crystalline symmetries or the topology of the electronic bands [1–5]. Topological semimetals exhibit exceptional electronic transport properties (e.g., extremely high carrier mobility and large linear magnetoresistance) and the control of these exotic charge carriers could help realize a new generation of spintronic devices with low power consumption [6–8].

Such control can potentially be realized in materials in which magnetic order coexists with nontrivial electronic band topology. Recent ARPES, quantum oscillation, neutron diffraction, and *ab initio* band-structure studies suggest that materials in the AMnSb_2 ($A = \text{Ca, Sr, Ba, Eu, Yb}$) family display many of the required properties [9–18]. The two-dimensional zigzag layer of Sb atoms [Fig. 1] in these 112-pnictides play host to fermions which can be described by the relativistic Dirac or Weyl equations. Furthermore, the electronic transport in this family of materials also displays large magnetoresistive effects, suggesting a coupling between the magnetism and charge carriers [9–17]. These effects could be driven by changes in the electronic band-structure topology due to changes in the symmetry of the spin structures induced by the applied field [19].

Within the AMnSb_2 family, EuMnSb_2 is of particular interest because the conducting zigzag layer of Sb atoms is sandwiched between two interpenetrating magnetic sub-

lattices (Eu and Mn), as shown in Fig. 1(a). Such a structure may lead to an enhancement of the coupling between the topological quasiparticles and magnetism, compared to that in compounds with a nonmagnetic atom on the A site. The dramatic magnetoresistive behavior observed in a recent paper [20] is evidence for the importance of this coupling. Up to now, however, the nature of the magnetic order in EuMnSb_2 , which determines the character of any topological band crossings, has yet to be determined. Moreover, although the magnetic susceptibility of EuMnSb_2 has an anomaly at 20 K, which is attributed to the AFM order of Eu, no evidence for magnetic order of the Mn spins has been detected up to 400 K [20]. This is surprising given that all other AMnX_2 ($X = \text{Sb, Bi}$) systems studied thus far exhibit AFM order of Mn near room temperature. [9–17, 21–35].

In light of this, we set out in this study to (i) determine the spin configuration of the two magnetic sublattices by powder neutron diffraction, (ii) investigate the nature of electronic states by mapping the band structure with ARPES, and (iii) shed light on the coupling between magnetism and the charge carriers through magnetotransport measurements. We find that the Mn sublattice displays long-range order below $T_N^{\text{Mn}} = 350(2)$ K with the magnetic moments aligned along the a axis in a C-type AFM structure. Upon further cooling, we find that the Eu spins exhibit an A-type AFM ordering at $T_N^{\text{Eu}} = 21(1)$ K with the spins aligned along the c axis. Our ARPES results show that EuMnSb_2 displays Dirac-like linearly dispersing electronic bands near the Fermi energy. Furthermore, we demonstrate that these electronic states are strongly coupled to the Eu magnetic sublattice, evidenced by the magnetotransport data presented in this paper.

II. EXPERIMENTAL DETAILS

Bulk EuMnSb_2 single crystals were grown via the flux method, as described in Ref. [20]. The unit cell

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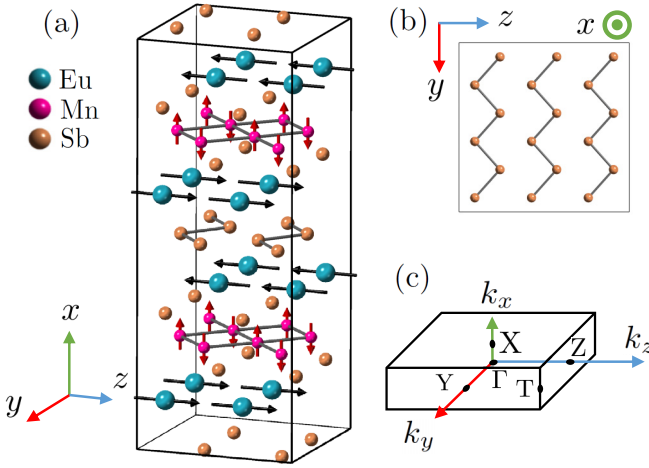


FIG. 1. (a) The crystal and magnetic structure of EuMnSb₂ (1 × 2 × 2 unit cells). The Eu and Mn sublattices exhibit A-type and C-type AFM order, respectively. (b) View down the x axis of the orthorhombic unit cell [space group *Pnma*, lattice parameters $a = 22.567(5)$ Å, $b = 4.371(1)$ Å, $c = 4.409(1)$ Å at $T = 300$ K] showing the zigzag network of Sb atoms in the y - z plane. (c) High symmetry points of the orthorhombic Brillouin zone.

can be described by the *Pnma* space group with the Eu and Mn atoms at two symmetry-inequivalent 4c Wyck-off positions [0.38637(2), 0.25, 0.76977(5)] and [0.24970(3), 0.25, 0.27005(14)], respectively ($T = 300$ K). Magnetotransport measurements were performed on a Quantum Design Physical Properties Measurement System for fields up to 13 T in a Hall configuration with the standard five-contact method. The longitudinal component, ρ_{zz} , was obtained by symmetrizing the data measured in positive and negative fields, respectively. Here we define $\rho_{zz} = E_z/j_z$, where E_z and j_z are the electric field and current density along z , respectively.

Powder neutron diffraction measurements of EuMnSb₂ were performed on the high-flux diffractometer WISH (ISIS Facility, UK) [36]. The EuMnSb₂ single crystals were crushed in argon and loaded in a vanadium can. The can had a relatively small diameter (3 mm) to reduce the attenuation due to the high neutron absorption of Eu ($\sigma_a^{\text{Eu}} = 4530$ barns). ARPES spectra of EuMnSb₂ were recorded with soft x rays on the SX-ARPES end station of the ADDRESS beamline (Swiss Light Source, Switzerland) [37]. Measurements were performed with a SPECS analyzer at a photon energy of 790 eV with right-circularly polarized light ($C+$). The sample was cleaved normal to the [100] direction and measured at $T \sim 20$ K at a vacuum better than 1×10^{-10} mbar.

III. RESULTS AND ANALYSIS

Figure 2 shows the temperature dependence of three powder neutron diffraction peaks, with d -spacings of 4.09 Å, 4.38 Å, and 11.25 Å, respectively. The intensity of the first two peaks increases sharply below $T_N^{\text{Mn}} = 350(2)$ K, signaling the onset of magnetic order on the Mn sublattice. The intensity at 4.09 Å continues to grow and saturates as the temperature

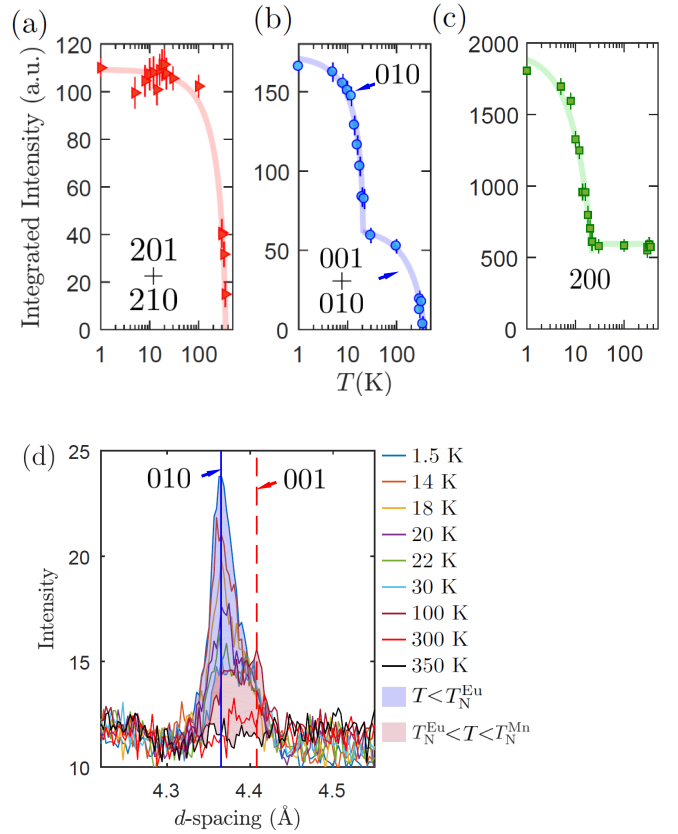


FIG. 2. (a)–(c) Temperature dependence (log scale) of the integrated intensity of the powder neutron diffraction peaks at d -spacings 4.09 Å, 4.38 Å, and 11.25 Å, respectively. (d) For $T_N^{\text{Eu}} < T < T_N^{\text{Mn}}$, the purely magnetic peak at 4.38 Å comprises the peaks 010 and 001. Below T_N^{Eu} , only the intensity of the 010 reflection grows.

approaches 1.5 K. On the other hand, the intensity of the reflection at 4.38 Å displays another sharp increase below $T_N^{\text{Eu}} = 21$ K, which is also observed at 11.25 Å. This transition is associated with the magnetic ordering of the Eu moments, and coincides with the anomaly in the temperature-dependent magnetotransport data (see later).

For $T_N^{\text{Eu}} < T < T_N^{\text{Mn}}$, the peak at 4.38 Å comprises two reflections at slightly different d -spacing, namely 4.37 Å and 4.41 Å, which can be indexed by the Miller indices 010 and 001. These peaks are not well separated as the ratio $c/b = 4.41/4.37$ is close to unity [see Fig. 2(d)]. The 010 and 001 peaks are purely magnetic as they are forbidden nuclear reflections in the *Pnma* space group. The peak at 4.09 Å is similarly made up of two reflections, 201 and 210, at d -spacings 4.10 Å and 4.08 Å, respectively. These are weak nuclear reflections in the paramagnetic phase which display additional magnetic intensity below T_N^{Mn} .

The peak at 11.25 Å, which is the nuclear-allowed 200 reflection, displays additional intensity due to magnetic scattering only for $T < T_N^{\text{Eu}}$, Fig. 2(c). The two magnetic orderings are also represented in Fig. 2(d), which shows the temperature dependence of the composite magnetic peak at 4.38 Å. As the intensity of the 001 component at 4.41 Å is already saturated by 100 K, the increase in the integrated intensity for $T < T_N^{\text{Eu}}$

TABLE I. Representational analysis for the 4c Wyckoff site of the $Pnma$ space group ($\mathbf{k} = \mathbf{0}$). The first column shows the relative spin arrangement at the atomic positions $(x, 0.25, z)$, $(-x + 0.5, 0.75, z + 0.5)$, $(-x, 0.75, -z)$, and $(x + 0.5, 0.25, -z + 0.5)$, respectively. The last column denotes whether the irrep can be described by one of the A_i , C_i , or G_i types of AFM, or by a ferromagnet, F_i , where $i = x, y, z$ indicates the spin orientation.

Basis			Structure	
vector	Orientation	Irrep	Eu	Mn
(++++)	x	Γ_3^+	F_x	F_x
	y	Γ_4^+	F_y	F_y
	z	Γ_2^+	F_z	F_z
(++--)	x	Γ_4^-		A_x
	y	Γ_3^-		A_y
	z	Γ_1^-		A_z
(+--+)	x	Γ_2^+		G_x
	y	Γ_1^+		G_y
	z	Γ_3^+		G_z
(+--++)	x	Γ_1^-	A_x	C_x
	y	Γ_2^-	A_y	C_y
	z	Γ_4^-	A_z	C_z

[Fig. 2(b)] only arises from the 010 peak at 4.37 Å. Additional peaks, which can be indexed by 211, 401, 410, 600, and 400, were also observed below T_N^{Eu} .

All of the observed magnetic peaks are consistent with the magnetic propagation vector of $\mathbf{k} = \mathbf{0}$ for both Mn and Eu. As both magnetic species have the same magnetic propagation vector and are on the same Wyckoff site (but different positions in the lattice), the magnetic order on both sublattices can be described by the same set of irreps. The magnetic Γ -point representation for the 4c Wyckoff position $(x, 0.25, z)$ of the $Pnma$ space group decomposes into eight distinct one-dimensional irreducible representations (irreps), $\Gamma = \Gamma_1^+ + 2\Gamma_1^- + 2\Gamma_2^+ + \Gamma_2^- + 2\Gamma_3^+ + \Gamma_3^- + \Gamma_4^+ + 2\Gamma_4^-$, with the associated basis vectors listed in Table I (the Miller and Love notation is adopted [38]). In both cases, the irreps which describe the ferromagnetic sublattices with moments along x , y , and z , namely Γ_3^+ , Γ_4^+ , and Γ_2^+ , can be excluded by the bulk magnetization [20]. We consider all of the remaining irreps for the Eu and Mn sublattices systematically and discuss the magnetism on each in turn.

For the Mn sublattice, the basis vector $(++--)$ [see Table I] which describes A-type AFMs (Γ_4^- , Γ_3^- , and Γ_1^-) does not produce the observed 201, 210, 001, and 010 magnetic reflections. On the other hand, the G-type AFMs $(+-+-)$ basis vector] with Mn moments along x , y , and z do not give rise to the 210 and 010 magnetic reflections. The basis vector $(+--++)$ with Mn moments along y and z is not predicted to have 010 and 001 magnetic reflections, respectively. Hence, the only irrep that is consistent with the observed peaks is the Γ_1^- irrep, which can be described by a C-type AFM with Mn moments along x .

For the Eu sublattice, the basis vector $(++--)$ with moments along x , y , and z , corresponding to the irreps Γ_4^- ,

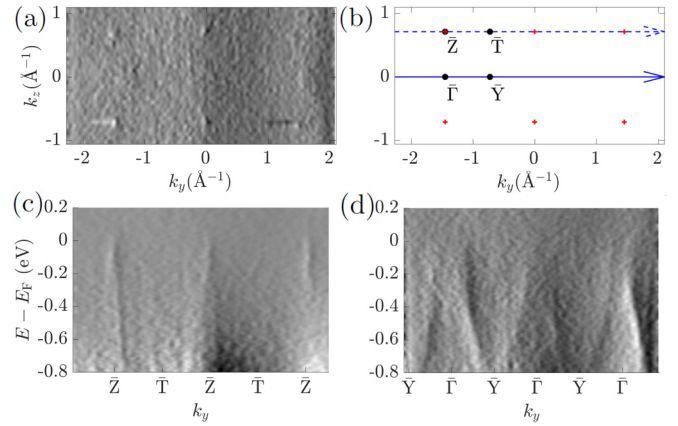


FIG. 3. First derivative of the ARPES data. (a) Constant energy surface measured at 790 eV with right-circularly polarized light at $E = E_F$. (b) High symmetry points in the k_y - k_z plane, as defined for the BZ shown in Fig. 1(c). (c), (d) Electronic dispersion along the high-symmetry line cuts indicated in (b) (dashed and solid blue arrows, respectively). Note that the first derivative is taken to emphasize the weak features in the data. While the features in the plot may not be at exactly the same \mathbf{k} positions as the bands, the raw data which do accurately show the bands can be found in the Supplemental Material (Ref. [39]).

Γ_3^- , and Γ_1^- respectively, do not produce the observed 010 and 200 magnetic reflections. The same is true for the basis vector $(+-+-)$ with irreps Γ_2^+ , Γ_1^+ , and Γ_3^+ . For the basis vector $(+--++)$, the irreps with Eu moments along x and z predict additional intensity at 001 which is not observed below T_N^{Eu} . This leaves us with an A-type AFM with Eu moments along y which can be described by the Γ_2^- irrep.

For $T_N^{\text{Eu}} < T < T_N^{\text{Mn}}$, the symmetry of the EuMnSb₂ crystal can be described by the $Pn'm'a'$ Shubnikov group due to the Mn order. This symmetry is lowered to $P2_1/a'$ (unique axis c , $\gamma \neq 90^\circ$), below the Eu ordering temperature, due to the combined effect of the two irreps. The magnetic structures for the Mn and Eu sublattices are presented in Fig. 1(a).

Figures 3(a), 3(c) and 3(d) illustrate the first derivative of our ARPES measurements of the electronic structure of EuMnSb₂, which were made at $T \simeq 20$ K. The constant energy map in the k_y - k_z plane at 0 eV binding energy (where $E = E_F$), as shown in Fig. 3(a), reveals small Fermi pockets at the \bar{Z} high-symmetry points across the six Brillouin zones [marked by the red crosses in Fig. 3(b)]. Furthermore, the \mathbf{k} - E dispersion slice in Fig. 3(c) for \mathbf{k} along the \bar{Z} - \bar{T} - \bar{Z} high symmetry line [depicted by the dashed blue arrow in Fig. 3(b)] shows steeply dispersing linear bands converging at the \bar{Z} high-symmetry points at E_F . This provides direct evidence for Dirac fermions which could be important for electronic transport.

Figure 3(d), which plots the \mathbf{k} - E dispersion for \mathbf{k} along the $\bar{\Gamma}$ - \bar{Y} - $\bar{\Gamma}$ line cut, as depicted by the solid blue arrow in Fig. 3(b), shows linearly dispersing bands that terminate at E_F . These bands, unlike those in Fig. 3(c), do not converge at E_F , but instead form hole pockets centered around the $\bar{\Gamma}$ point.

Furthermore, we also observe a smearing out of the bands beyond binding energies of about 0.5 eV and 0.7 eV in

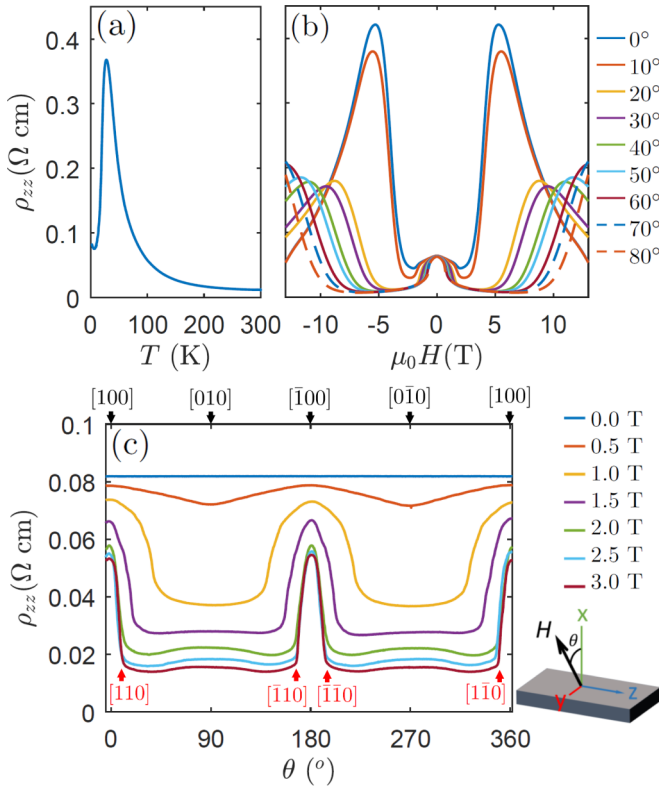


FIG. 4. (a) The in-plane resistivity shows a metal-to-insulator transition which peaks at $T_N^{\text{Eu}} \sim 20$ K before falling at lower temperatures. (b) Field-dependent ρ_{zz} at various field directions measured at $T = 2$ K. Here the angles [see insert of Fig. 4(c)] are defined with respect to the x axis where the field direction is fixed to be within the x - y plane. (c) Angular dependence of ρ_{zz} at various field strengths demonstrate the strong anisotropy in the magnetotransport of EuMnSb_2 .

Figs. 3(c) and 3(d), respectively. This effect can arise from the Eu $4f$ bands which are strongly correlated due to the highly localized nature of the orbitals. Hence, ARPES shows that the $4f$ states, which give rise to the Eu magnetism, are not expected to play a direct role in electronic transport. Our magnetotransport data, which we will now show, demonstrate that the $4f$ electrons can nonetheless influence charge transport indirectly through exchange interactions.

The longitudinal magnetotransport behavior of EuMnSb_2 with the current along z is summarized in Fig. 4. Figure 4(a) plots ρ_{xx} as a function of temperature in zero field. The resistivity increases strongly on cooling, reaching a maximum at T_N^{Eu} with $\rho_{xx}^{20\text{K}}/\rho_{xx}^{300\text{K}} \sim 31$, then decreases at lower temperatures. This behavior is consistent with paramagnetic scattering of conduction electrons by the Eu spins, and is evidence that the charge carriers in EuMnSb_2 are strongly coupled to spin fluctuations associated with the magnetic ordering on the Eu sublattice.

To help understand the magnetoresistance behavior further, we also measured ρ_{zz} at $T = 2$ K as a function of applied field in several directions within the x - y plane, where the angle θ is defined with respect to the crystallographic a axis [see insert of Fig. 4(c)]. The data are plotted in Fig. 4(b). At

$\theta = 0^\circ$, three distinct features can be identified in ρ_{zz} : (i) In the low field regime ($\mu_0 H \lesssim 3$ T), there is a small drop in ρ_{zz} . (ii) In the intermediate regime ($3 \text{ T} \lesssim \mu_0 H \lesssim 5.3$ T), there is a sharp increase in ρ_{zz} , with the fractional change in $\rho_{zz}(H)$ reaching 9.4 at the maximum. (iii) In the high field regime ($\mu_0 H > 5.3$ T), the anomalous resistivity decreases again with increasing field.

With increasing field applied along the $[100]$ direction ($\theta = 0^\circ$), the AFM-ordered Mn spins are expected to flop into the y - z plane. Subsequently, both the Mn and Eu spins will rotate toward the x axis and eventually become fully spin-polarized at their respective saturation fields. Strong spin fluctuations are expected during this evolution in the magnetic structures, which can lead to enhanced scattering of the conduction electrons. An exchange-induced modification of the electronic bands is also possible, arising out of the changes in the magnetic symmetry due to the field-induced alteration of spin ordering which can lift or add symmetry protection to the band crossing at high-symmetry points in the Brillouin zone [40,41]. These are two possible mechanisms which could account for the large magnetoresistive effects seen in Fig. 4(b).

The ρ_{zz} magnetoresistance displays strong angular dependence, with the peak moving from 5.3 T for $\theta = 0^\circ$, to about 8 T for $\theta = 20^\circ$, and eventually beyond the measurement window range of 13 T, for $\theta = 80^\circ$. For $\mu_0 H = \pm 5.3$ T, we find that the ratio of ρ_{zz} measured at $\theta = 0^\circ$ to that at $\theta = 80^\circ$, or $\rho_{zz}^{0^\circ}/\rho_{zz}^{80^\circ}$, is 54. Another view of this anisotropy is given in Fig. 4(c), which plots ρ_{zz} as a function of θ , measured at $T = 2$ K at various field strengths up to 3 T. For this measurement, the in-plane resistivity was continuously measured as the applied field was rotated about the current j_z . The field thus passes through the principal crystal axes $[100]$, $[010]$, $[\bar{1}00]$, and $[0\bar{1}0]$ at $\theta = 0^\circ, 90^\circ, 180^\circ$, and 270° , respectively [see insert of Fig. 4(c)]. The 180° periodicity in ρ_{zz} is in agreement with charge transport in a two-dimensional layer. Moreover, consistent with the $|\mu_0 H| \leq 3$ T data presented in Fig. 4(b), ρ_{zz} decreases in all field directions, forming sharp resistivity peaks at angles $\theta = 0^\circ, 180^\circ$, and 360° . At $\mu_0 H = 3$ T, the peaks have an angular width of $\delta\theta \sim 20^\circ$, and are suppressed when $\theta = 10^\circ, 170^\circ, 190^\circ$, and 350° , as denoted by the red arrows in Fig. 4(c). Intriguingly, these angles correspond to the $[110]$, $[\bar{1}10]$, $[\bar{1}\bar{1}0]$, and $[1\bar{1}0]$ directions of the orthorhombic crystal, respectively.

The results presented in Figs. 4(b)–4(c) demonstrate that the charge transport in EuMnSb_2 is extremely sensitive to the applied field direction. The large magnetoresistance which arises when the field is aligned with the $[100]$ direction (perpendicular to the in-plane Eu magnetic moments) is suppressed as soon as there is a small component of field along the $[010]$ direction.

The extent to which the electron bands are altered remains to be seen since ARPES cannot be performed in an applied magnetic field. Hence, a natural extension of the present paper would be an *ab initio* study of the electronic band structure with different spin configurations, the latter being determined experimentally by resonant elastic x-ray scattering (REXS) in an applied field. As REXS is element specific, it allows for the study of the spin configuration on the Eu and Mn magnetic sublattices separately.

IV. CONCLUSION

In conclusion, we have used powder neutron diffraction to establish that EuMnSb₂ displays magnetic ordering at two distinct temperatures, and we have determined the spin arrangement for both magnetic sublattices in spite of the large neutron absorption of Eu. Moreover, in the small energy window available to map the electronic spectrum due to the presence of strongly correlated Eu 4*f* bands, we have successfully recorded ARPES spectra which provide evidence for a Dirac-like linear dispersion at E_F . Finally, our magnetotransport results show that in EuMnSb₂ there is a strong coupling between magnetic order and charge transport. These three strands of evidence lend support to the prediction that

EuMnSb₂ is a promising material to realize magnetic control of topological quasiparticles.

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- [1] N. P. Armitage, E. J. Mele, and A. Vishwanath, *Rev. Mod. Phys.* **90**, 015001 (2018).
 - [2] A. A. Burkov, *Nat. Mater.* **15**, 1145 (2016).
 - [3] J. G. Rau, E. K.-H. Lee, and H.-Y. Kee, *Annu. Rev. Condens. Matter Phys.* **7**, 195 (2016).
 - [4] M. Z. Hasan and C. L. Kane, *Rev. Mod. Phys.* **82**, 3045 (2010).
 - [5] D. Pesin and L. Balents, *Nat. Phys.* **6**, 376 (2010).
 - [6] L. Šmejkal, T. Jungwirth, and J. Sinova, *Phys. Status Solidi B* **RRL 11**, 1700044 (2017).
 - [7] L. Šmejkal, J. Železný, J. Sinova, and T. Jungwirth, *Phys. Rev. Lett.* **118**, 106402 (2017).
 - [8] Z. Shi, M. Wang, and J. Wu, *Appl. Phys. Lett.* **107**, 102403 (2015).
 - [9] J. Y. Liu, J. Hu, H. Cao, Y. Zhu, A. Chuang, D. Graf, D. J. Adams, S. M. A. Radmanesh, L. Spinu, I. Chiorescu, and Z. Q. Mao, *Sci. Rep.* **6**, 30525 (2016).
 - [10] M. A. Farhan, G. Lee, and J. H. Shim, *J. Phys.: Condens. Matter* **26**, 042201 (2014).
 - [11] R. Kealhofer, S. Jang, S. M. Griffin, C. John, K. A. Benavides, S. Doyle, T. Helm, P. J. W. Moll, J. B. Neaton, J. Y. Chan, J. D. Denlinger, and J. G. Analytis, *Phys. Rev. B* **97**, 045109 (2018).
 - [12] Y.-Y. Wang, S. Xu, L.-L. Sun, and T.-L. Xia, *Phys. Rev. Mater.* **2**, 021201 (2018).
 - [13] S. V. Ramankutty, J. Henke, A. Schiphorst, R. Nutakki, S. Bron, G. Araizi-Kanoutas, S. K. Mishra, L. Li, Y. K. Huang, T. K. Kim, M. Hoesch, C. Schlueter, T. L. Lee, A. de Visser, Z. Zhong, J. van Wezel, E. van Heumen, and M. S. Golden, *SciPost Phys.* **4**, 010 (2018).
 - [14] J. Y. Liu, J. Hu, Q. Zhang, D. Graf, H. B. Cao, S. M. A. Radmanesh, D. J. Adams, Y. L. Zhu, G. Cheng, X. Liu, W. A. Phelan, J. Wei, M. Jaime, F. Balakirev, D. A. Tennant, J. F. DiTusa, I. Chiorescu, L. Spinu, and Z. Q. Mao, *Nat. Mater.* **16**, 905 (2017).
 - [15] J. B. He, Y. Fu, L. X. Zhao, H. Liang, D. Chen, Y. M. Leng, X. M. Wang, J. Li, S. Zhang, M. Q. Xue, C. H. Li, P. Zhang, Z. A. Ren, and G. F. Chen, *Phys. Rev. B* **95**, 045128 (2017).
 - [16] J. S. You, I. Lee, E. Choi, Y. Jo, J. Shim, and J. S. Kim, *Curr. Appl. Phys.* **19**, 230 (2019).
 - [17] X. Yan, C. Zhang, S.-S. Liu, Y.-W. Liu, D. W. Zhang, F.-X. Xiu, and P. Zhou, *Front. Phys.* **12**, 127209 (2017).
 - [18] S. J. Ray and L. Alff, *Phys. Status Solidi B* **254**, 1600163 (2017).
 - [19] S. Klemenz, S. Lei, and L. M. Schoop, *Annu. Rev. Mater. Res.* **49**, 185 (2019).
 - [20] C. Yi, S. Yang, M. Yang, L. Wang, Y. Matsushita, S. Miao, Y. Jiao, J. Cheng, Y. Li, K. Yamaura, Y. Shi, and J. Luo, *Phys. Rev. B* **96**, 205103 (2017).
 - [21] K. Wang, D. Graf, L. Wang, H. Lei, S. W. Tozer, and C. Petrovic, *Phys. Rev. B* **85**, 041101(R) (2012).
 - [22] A. Wang, I. Zaliznyak, W. Ren, L. Wu, D. Graf, V. O. Garlea, J. B. Warren, E. Bozin, Y. Zhu, and C. Petrovic, *Phys. Rev. B* **94**, 165161 (2016).
 - [23] E. Brechtel, G. Cordier, and H. Schäfer, *J. Less-Common Met.* **79**, 131 (1981).
 - [24] J. Park, G. Lee, F. Wolff-Fabris, Y. Y. Koh, M. J. Eom, Y. K. Kim, M. A. Farhan, Y. J. Jo, C. Kim, J. H. Shim, and J. S. Kim, *Phys. Rev. Lett.* **107**, 126402 (2011).
 - [25] Y. Feng, Z. Wang, C. Chen, Y. Shi, Z. Xie, H. Yi, A. Liang, S. He, J. He, Y. Peng, X. Liu, Y. Liu, L. Zhao, G. Liu, X. Dong, J. Zhang, C. Chen, Z. Xu, X. Dai, Z. Fang, and X. J. Zhou, *Sci. Rep.* **4**, 5385 (2014).
 - [26] H. Masuda, H. Sakai, M. Tokunaga, Y. Yamasaki, A. Miyake, J. Shiogai, S. Nakamura, S. Awaji, A. Tsukazaki, H. Nakao, Y. Murakami, T.-h. Arima, Y. Tokura, and S. Ishiwata, *Sci. Adv.* **2**, e1501117 (2016).
 - [27] Y. J. Jo, J. Park, G. Lee, M. J. Eom, E. S. Choi, J. H. Shim, W. Kang, and J. S. Kim, *Phys. Rev. Lett.* **113**, 156602 (2014).
 - [28] S. Borisenko, D. Evtushinsky, Q. Gibson, A. Yaresko, T. Kim, M. N. Ali, B. Buechner, M. Hoesch, and R. J. Cava, *arXiv:1507.04847*.
 - [29] J. Y. Liu, J. Hu, D. Graf, T. Zou, M. Zhu, Y. Shi, S. Che, S. M. A. Radmanesh, C. N. Lau, L. Spinu, H. B. Cao, X. Ke, and Z. Q. Mao, *Nat. Comms.* **8**, 646 (2017).
 - [30] A. F. May, M. A. McGuire, and B. C. Sales, *Phys. Rev. B* **90**, 075109 (2014).
 - [31] G. Lee, M. A. Farhan, J. S. Kim, and J. H. Shim, *Phys. Rev. B* **87**, 245104 (2013).
 - [32] Y. F. Guo, A. J. Princep, X. Zhang, P. Manuel, D. Khalyavin, I. I. Mazin, Y. G. Shi, and A. T. Boothroyd, *Phys. Rev. B* **90**, 075120 (2014).
 - [33] M. Chinotti, A. Pal, W. J. Ren, C. Petrovic, and L. Degiorgi, *Phys. Rev. B* **94**, 245101 (2016).

- [34] J. K. Wang, L. L. Zhao, Q. Yin, G. Kotliar, M. S. Kim, M. C. Aronson, and E. Morosan, *Phys. Rev. B* **84**, 064428 (2011).
- [35] J. B. He, D. M. Wang, and G. F. Chen, *Appl. Phys. Lett.* **100**, 112405 (2012).
- [36] L. Chapon, P. Manuel, P. Radaelli, C. Benson, L. Perrott, S. Ansell, N. Rhodes, D. Raspino, D. Duxbury, E. Spill, and J. Norris, *Neutron News* **22**, 22 (2011).
- [37] V. N. Strocov, T. Schmitt, U. Flechsig, T. Schmidt, A. Imhof, Q. Chen, J. Raabe, R. Betemps, D. Zimoch, J. Krempasky, X. Wang, M. Grioni, A. Piazzalunga, and L. Patthey, *J. Synchrotron Radiat.* **17**, 631 (2010).
- [38] S. C. Miller and W. F. Love, *Tables of Irreducible Representations of Space Groups and Co-Representations of Magnetic Space Groups* (Pruett Press, Boulder, Colorado, 1967).
- [39] See Supplemental Material at <http://link.aps.org/supplemental/10.1103/PhysRevB.100.174406> for raw ARPES data.
- [40] L. M. Schoop, A. Topp, J. Lippmann, F. Orlandi, L. Muehler, M. G. Vergniory, Y. Sun, A. W. Rost, V. Duppel, M. Krivenkov, S. Sheoran, P. Manuel, A. Varykhalov, B. Yan, R. K. Kremer, C. R. Ast, and B. V. Lotsch, *Sci. Adv.* **4**, eaar2317 (2018).
- [41] M. G. Vergniory, L. Elcoro, F. Orlandi, B. Balke, Y.-H. Chan, J. Nuss, A. P. Schnyder, and L. M. Schoop, *Eur. Phys. J. B* **91**, 213 (2018).