Quantum transport properties of the topological Dirac semimetal α-Sn

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We report on measurements of the electrical resistivity (ρ) and thermoelectric power (*S*) in a thin film of strained single-crystalline α -Sn grown by molecular beam epitaxy on an insulating substrate. The temperature (*T*) dependence of the resistivity of α -Sn can be divided into two regions: below $T^* \approx 135$ K $\rho(T)$ shows metalliclike behavior, while above this temperature, an increasing contribution from thermally excited holes to electrical transport is observed. However, it is still dominated by highly mobile electrons, resulting in a negative sign of the Seebeck coefficient above T = 47 K. In the presence of the magnetic field (*B*) applied along an electric field or thermal gradient, we note negative magnetoresistance or a negative slope of *S*(*B*), respectively. The theoretical prediction for the former (calculated using density functional theory) agrees well with the experiment. However, these characteristics quickly disappear when the magnetic field is deviated from an orientation parallel to the electrical field or the thermal gradient. We indicate that the behavior of the electrical resistivity and thermoelectric power can be explained in terms of the chiral current arising from the topologically nontrivial electronic structure of α -Sn. Its decay at high temperature is a consequence of the decreasing ratio between the intervalley Weyl relaxation time to the Drude scattering time.

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I. INTRODUCTION

Over the past two decades, topological quantum materials have gained significant attention due to their nontrivial momentum-space topology [1–4]. For example, the Dirac semimetals, which could be considered three-dimensional (3D) analogs to graphene, host fourfold-degenerate Dirac points protected by topological constraints [5–7]. Dirac nodes split into two chirally distinct Weyl nodes when at least one of the symmetries protecting the Dirac cone gets broken [8–10]. Dirac semimetals exhibit unique and exciting features such as ultrahigh mobility [11], large magnetoresistance [11,12], and chiral magnetic effect [13,14]. These exotic phenomena have been experimentally reported in a number of topological Dirac semimetals, namely, Na₃Bi [15,16], Cd₂As₃ [11,17,18], ZrTe₅ [19,20], Bi_{1-x}Sb_x [21], YbMnBi₂ [22], and TIBiSSe [12].

Recently, gray tin, or α -Sn, a zero-gap semiconductor, has emerged as an exciting material due to its nontrivial band topology [23–25]. It is the elemental candidate showing many topological phases that can be tailored by various conditions, such as changing the thickness, imposing the strain, and applying electric and magnetic fields [26,27]. The application of the in-plane tensile strain transforms α -Sn into a robust 3D topological insulator with a large topological gap [24] and a high Fermi velocity [28], whereas in-plane compressive strain makes it a topological Dirac semimetal protected by

fourfold rotational symmetry [26,27]. The in-plane compressive strain, which modifies the electronic structure of α -Sn, can be obtained experimentally by epitaxially growing a thin film on a substrate with lattice constants that do not exactly match those of gray tin. To obtain the Dirac semimetal phase, the desired mismatch is achieved by selecting the appropriate substrates, such as InSb(111) [29], InSb(001) [30], CdTe(111) [31], and GaAs(001) [32]. The degeneracy of the Dirac cones is lifted in the presence of an external magnetic field, turning α -Sn into a Weyl semimetal (WSM). The presence of a pair of Weyl points (WPs) in momentum space can lead to the appearance of a peculiar phenomenon when the electric and magnetic fields are applied parallel to each other. It is known as a chiral anomaly and can be observed as a positive magnetoconductivity in real crystalline materials, as suggested by Nielsen and Ninomiya [33]. Its origin is the charge pumping between WPs of opposite chirality that occurs in a WSM subjected to electric and magnetic fields applied in parallel along the direction of the WP separation [13]. Such negative magnetoresistance has been reported by various groups in topological Dirac and WSMs [14,15,20,34,35], but there exist other mechanisms that could underlie this effect without involving charge pumping between the Weyl nodes [36–38]. In this regard, measurements of the thermoelectric effects offer an opportunity to complement electrical measurements and gain valuable insight into electronic transport properties [39-42].

In this paper, we investigate the electrical and thermoelectrical properties of a α -Sn thin film, which is a topological Dirac semimetal when deposited on a CdTe/GaAs (001) substrate. To ensure the best quality, the sample was grown using

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molecular beam epitaxy (MBE). We report on the quadratic variation in the field dependencies of the electrical conductivity and thermopower, which can be explained because of the chiral anomaly. We also calculated the ratio of intervalley scattering time to mean free time, which indicates the role of chiral Weyl fermions in the emergence of the negative longitudinal magnetoresistance (NLMR) and the negative slope of thermopower at the high magnetic field.

II. MATERIALS AND METHODS

Epitaxial film of 200-nm-thick α -Sn was grown by MBE on (001) GaAs substrate with a 4 µm CdTe buffer layer. CdTe buffer provides the necessary ~0.1% compressive in-plane strain to induce the Dirac semimetal phase in gray tin. Extensive structural characterization confirmed the high quality of the obtained film without any inclusions of the metallic β -Sn phase as well as the value of the strains. More details about the growth procedure as well as the structural characterization of the studied film can be found in Ref. [32].

To perform the electric transport measurements, we cut the sample with a length (*a* axis) of 2.2 mm and a width (*b* axis) of 0.6 mm. This reasonable aspect ratio of the sample was chosen to minimize the geometric effects on our experimental data. We measured the resistivity using the standard four-probe method. The electrical contacts were made with 25- μ m-thick gold wires and DuPont 4929 silver paint. The current contacts were made along the entire width of a sample to minimize the possibility of the current jetting effect occurrence [43]. The dc electrical current was applied using a Keithley 6221 current source, and voltages along the sample were measured with a Keithley 2182A nanovoltmeter.

To measure the thermoelectric properties, the sample was mounted between the two phosphor bronze clamps with two Cernox thermometers attached. These were used to determine the thermal gradient along the sample generated by a Micro-Measurements strain gage heater (10 k Ω resistance), which was connected to a Keithley 6221 current source. Thermoelectric voltage data were collected by an EM Electronics A20a DC subnanovolt amplifier working in conjunction with a Keithley 2182A nanovoltmeter. The temperature dependence of thermoelectric power was measured with the heater off and on method, while for the magnetic field (±14.5 T) sweeps, the heater was continuously turned on.

The electronic structure calculations were carried out with the projector augmented-wave approach within the density functional theory (DFT) framework, utilizing VASP [44]. A plane-wave energy cutoff of 650 eV was employed in this paper. We have performed the calculation using a metageneralized gradient approximation approach, which is based on the modified Becke-Johnson (MBJ) exchange potential together with local density approximation for the correlation potential scheme with the parameter CMBJ = 1.215to get the experimental band ordering [45]. The calculations of electronic structures were performed with a 12×12 $\times 10$ Monkhorst-Pack k mesh [46], incorporating the spinorbit coupling (SOC) self-consistently. The VASPWANNIER90 interface was employed in this paper, and we utilized s and p orbitals of Sn atoms to generate an ab initio tightbinding Hamiltonian without performing the procedure for



FIG. 1. Temperature dependences of the resistivity (ρ) and the thermoelectric power (*S*) of 200-nm-thick α -Sn thin film where the current (*J*) or thermal gradient (∇T) is applied parallel to the *a* axis. Inset shows low-temperature thermoelectric power data.

maximizing localization [47,48]. The calculation of the surface state was performed using the semi-infinite Green's function approach incorporated in WANNIERTOOLS [49,50].

III. RESULTS AND DISCUSSIONS

The thermoelectric power (S) and electrical resistivity (ρ) were measured, respectively, with the thermal gradient (∇T) or electrical current (*j*) applied along the *a* axis of the α -Sn thin film. Figure 1 presents the temperature dependence of resistivity in zero magnetic field (B) measured in the temperature range 2–300 K. The entire $\rho(T)$ dependence can be divided into two temperature regions, namely, for $T \ge 135$ K, we observe semiconducting behavior that can be attributed to the increasing temperature contribution to electrical transport from thermally excited holes [51]. Those holes originate from the thermally driven transitions between valence and conduction parts of the Γ_{8vc}^+ band as well as from the indirect transition $L_{6c}^+ - \Gamma_{8vc}^+$ found in the calculated electronic structure of α -Sn [52]. In the latter, the magnitude of the band gap in the epitaxially stretched α -Sn thin film is temperature dependent and becomes larger as the temperature decreases [53]. In consequence, thermal hole excitation is prevented for $T \leq 135$ K, and $\rho(T)$ behaves in a metalliclike manner. This is due to the electrons in the vicinity of Γ bands dominating the transport features of α -Sn at low temperatures [51]. Interestingly, even at a very low temperature, some contribution to the electronic transport from holelike charge carriers is still present [32].

It appears that the temperature dependence of the thermopower is also affected by the presence of two types of charge carriers. In almost the entire temperature range, the thermopower is negative, reflecting a dominating role of highly mobile electrons. The absolute value of *S* reaches its maximum of 22.5 μ V/K at $T \approx 150$ K. The upturn in *S*(*T*) above this temperature likely reflects the above-discussed increasing contribution from the thermally excited holes. For a multiband conductor, the total thermoelectric power is a



FIG. 2. (a) Normalized magnetoresistance vs magnetic field of α -Sn for selected temperatures when both magnetic field and current are applied parallel to the *a* axis ($B \parallel j$). Magnetothermopower of α -Sn measured with the configuration of applied thermal gradient and magnetic field parallel to the *a* axis ($\nabla T \parallel B$), (b) at low temperatures, and (c) at high temperatures.

sum of individual band contributions weighted by the respective conductivities. Within the two-band model, this can be expressed as $S = (S_e \sigma_e + S_h \sigma_h)/\sigma$, where $\sigma = \sigma_e + \sigma_h$, where subscripts e and h denote contributions from electrons and holes, respectively. Therefore, at high temperatures, the increasing positive participation from thermally excited holes will result in a decreasing absolute value of negative S dominated by electrons. Below $T \approx 50$ K, the Seebeck coefficient becomes $<1 \mu V/K$ and attains a positive value in the low temperature limit (see inset of the Fig. 1), which can be also seen in the S(B) dependences presented in Fig. 2(b). This may be due to phonon drag but also contribution from the low-mobility holes, which were recently reported to be present at low temperatures in α -Sn thin films [32]. For the former, S(T) should exhibit a maximum at the temperature of about $\theta_D/5$ [54], where θ_D is the Debye temperature. In gray tin $\theta_{\rm D} \approx 260 \, \text{K}$ [55], but the maximum we observe is at the temperature somewhat lower than the expected $T \approx$ 50 K. However, this is only an estimated position, which may also be effectively shifted by the contribution from diffusive thermopower. Nevertheless, this positive contribution appears to be weakly dependent on the magnetic field, so does not affect our analysis.

The magnetoresistance of α -Sn was measured at different temperatures in the parallel configuration of the electrical current and magnetic field (j||B); the results are presented in Fig. 2(a). Below $T \approx 100$ K, a Landau quantization occurs, leading to the appearance of pronounced Shubnikov-de Haas (SdH) effect in high magnetic field. The resulting strong oscillations in $\rho(B)$ are shown in Fig. S1(a) in the Supplemental Material (SM) [56] after subtraction of slowly varying background (third-order polynomial). The fast Fourier transform (FFT) spectrum (see Fig. S1(b) in the SM [56]) reveals a single slow frequency $F \sim 12$ T, that can be assigned to a bulk Dirac point near the Fermi level of Alpha-Sn, in agreement with the previous study [32]. The SdH oscillations can be described by the Lifshitz-Kosevich theory [57]:

$$\Delta \rho \propto R_T R_D R_S \cos\left[2\pi \left(\frac{F}{B} - \frac{1}{2} + \beta \pm \delta\right)\right].$$
(1)

The three damping factors are the thermal reduction factor $R_T = \frac{\chi}{\sinh(\chi)}$, the Dingle damping factor $R_D = \exp(-\chi \frac{T_D}{T})$ (T_D is the Dingle temperature), and the spin-splitting term $R_S = \cos(\frac{p\pi}{2}\frac{gm^*}{m_e})$. The parameter $\chi = \frac{2\pi^2 k_B T m^*/m_e}{e\hbar B}$, where $k_B = 1.381 \times 10^{-23}$ J K⁻¹ is the Boltzman constant, $e = 1.602 \times 10^{-19}$ C and $m_e = 9.108 \times 10^{-31}$ kg are the electron charge and mass, $\hbar = 1.054 \times 10^{-34}$ J s is the reduced Planck's constant, and m^* is the cyclotron mass of the charge carrier. Within the R_S term, g is the Landé factor, and p is the harmonic order. Under the cosine function in Eq. (1), F is the frequency of quantum oscillations (QOs), $2\pi\beta$ is the Berry phase, and $\delta = \pm \frac{1}{8}$ is the phase shift related to the dimensionality of the electronic structure of a 3D system.

The effective mass can be calculated from the temperature dependence of the amplitude of the oscillations described by the thermal reduction factor R_T (see Fig. S1(c) in the SM [56]), and the resulting small effective mass $m^* \approx 0.01 m_e$ is close to that previously reported [32]. The frequency of the QOs is directly related to the Fermi surface cross-section area via the Onsager relation: $F = (\frac{\hbar}{2\pi e})A_F$, where A_F is the Fermi surface cross-section area. Since the Fermi momentum $k_F =$ $\sqrt{\frac{A_F}{\pi}}$, we have deduced the Fermi velocity $v_F = \frac{\hbar k_F}{m^*}$ and the Fermi energy $E_F = m^* v_F^2$ under the assumption that charge carriers have linear energy dispersion [58]. The estimated v_F and E_F are $\sim 2.2 \times 10^6$ m/s and 277 meV, respectively. Figure S1(d) in the SM [56] shows that the experimental data can be well modeled with Eq. (1) from which another parameter that can be obtained from the analysis of QO is the Dingle temperature, which was estimated to be $T_D = 11$ K at 9.9 K, again in good agreement with results reported in Ref. [32]. From the Dingle temperature, we can calculate the quantum lifetime $\tau_q = \frac{\hbar}{2\pi k_B T_D}$ and quantum mobility $\mu_q = \frac{e\tau_q}{m^*}$ [59], which are $\tau_q \approx 1 \times 10^{-13}$ s and $\mu_q \approx 5900 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$, respectively. Such a high value of μ_q confirms the excellent transport properties of α -Sn.

Next, we will focus on a nonoscillatory component in $\rho(B)$ of α -Sn, which for j||B exhibits a large NLMR at high field and low temperature. With increasing temperature, NLMR decreases and vanishes above $T \approx 200$ K. Intriguingly, the presence of NLMR for j || B was also reported in early studies of QOs in bulk single crystals of α -Sn [60]. Furthermore, the authors observed an additional $\frac{1}{8}$ phase shift of the SdH oscillation, which is the expected result for the extra Berry phase of carriers from a 3D band with linear dispersion [61]. These experimental facts initially remained uninterpreted but have recently been linked to the presence of Dirac cones [26]. This is interesting because α -Sn in its pristine form should be, as mentioned above, a semiconductor with conduction and valence bands touching each other at the vertices. Perhaps even a small perturbation, such as the presence of an internal strain or a strain applied unwillingly during the experiment, can slightly shift the electronic bands, creating a nontrivial electron structure in the bulk sample [62].

The appearance of NLMR was reported for many Dirac and WSMs [15,20,34,35,63], and it has been ascribed to pumping of the chiral charge between the Weyl nodes. On the other hand, questions were also raised as to whether the effect could be due to extrinsic mechanisms [36,38]. The current jetting contribution in highly mobile WSMs caused concerns about the detection of the chiral current in electrical measurements [43]. Therefore, the complementary measurements of other phenomena, such as the thermoelectric power, can provide a unique opportunity to investigate the effects of the chiral anomaly without a current jetting artifact [41,43].

Figure 2(b) presents the magnetic field dependences of the thermopower (with $B \parallel \nabla T$) for selected temperatures. As mentioned above, in addition to the negative thermopower attributable to electrons, we also observe a roughly field-independent positive contribution to the Seebeck coefficient. This appears in the S(B) data in Fig. 2(b) as a temperature-dependent vertical shift that does not affect the field-dependent part. At low magnetic field, S initially increases with B-such a dependence was also observed in other topological Dirac semimetals, and it was attributed to the process of Weyl node creation [43,64]. Below $T \approx 50$ K, we see a small peak in S(B) at $B \approx 5.2$ T, which can be attributed to the QOs. For the frequency obtained from the SdH effect (F = 12 T), the next peak in S(B) is expected at $B \approx 8.8 \text{ T}$, which coincides with the maximum that occurs at $B \approx 9$ T. The contribution from QO to the total value of S at the maximum can be calculated using the Lifshitz-Kosevich formula and is presented in Fig. S2 in the SM [56]. At $T \approx 30$ K, the peak value of the QO from the baseline is $\approx 0.026 \,\mu V \, K^{-2}$, which is $\approx 33\%$ of the total weight of *S* at $B \approx 9$ T. However, the calculated contribution from QO significantly decreases with increasing temperature. For example, this becomes $\sim 8\%$ at $T \approx 50$ K and vanishes completely above $T \gtrsim 55$ K, while the peak of S(B) linked to formations of WPs is still present

in the data. The maximum in S(B) shifts to higher magnetic fields and, at $B \approx 12$ T, is clearly visible even at $T \approx 150$ K [see Fig. 2(b)]. Thus, it is unlikely that the negative slope of S(B) observed at high magnetic field originates from QOs, and alternatively, this type of anomalous field dependence was indicated as a manifestation of the chiral anomaly in several Dirac and WSMs [43].

To support our experimental observation, we have performed first-principles calculations using VASP [44] and WANNIER90 [47]. Figure 3(a) shows the electronic band structure in the presence of SOC, where most of the electronic bands are contributed by the *s* and *p* orbitals of Sn atoms. We observe a fourfold-degenerate band crossing along the $\Gamma \rightarrow$ *Z* direction with camelback features [65,66] [see Fig. 3(b)]. The Dirac points are located 9 meV above the Fermi level $E_{\text{Dirac}} = E_f - 0.009 \text{ eV}$ at $(0, 0, \pm k_z)$, where $k_z = 0.398 \text{ Å}^{-1}$, although it should be mentioned that DFT calculations can overestimate or underestimate the Fermi energy.

The pair of Dirac points can be shown in the electronic band structure in the two-dimensional (2D) plane shown in Fig. 3(d). The application of magnetic field gives rise to a negative magnetoresistance demonstrated in the calculations when $\theta = 0^{\circ}$, as shown in Fig. 3(e), which is consistent with experimental observation. Furthermore, we demonstrate the Dirac semimetal phase by investigating the band structure projected in Fig. S3 in the SM [56], (a) along the (100) direction and (b) along the (001) direction. The presence of the topological surface state is evident with bulk band crossing points. In addition, we calculate the Fermi surface projected on (100) and (001) surfaces in the bottom panel of Fig. S3 in the SM [56]. The Fermi surface along (100) shows the presence of a close topological Fermi arc $k_y - k_z$ plane. These are C_{4z} rotational symmetry-protected Dirac nodes that are against gap formation. In the presence of a finite Zeeman field that breaks time-reversal symmetry, each Dirac node separates into two Weyl nodes with the opposite chirality. Notably, these paired Weyl nodes are still aligned with the high-symmetry direction protected by the crystal symmetry C_{4z} . The development of the chiral anomaly is facilitated by the combination of this property and the chiral nature of their lowest Landau level [27,43].

If the negative slopes of $\rho(B)$ and S(B) in the high field are in fact signs of the chiral anomaly, they should disappear when the magnetic field is tilted away from being parallel to *j* or ∇T , respectively. The angular dependences of longitudinal transport coefficients at T = 60 K are presented in Fig. 4. In the measurement of the magnetoresistance [Fig. 4(a)], we observed the maximal NLMR when $\theta_{\rho} = 0^{\circ}$ (θ_{ρ} is the angle between B and j), whereas rotation of the magnetic field from in-plane to out-of-plane (from the a axis toward the c axis) quickly causes the dip in $\rho(B)$ to vanish. Similar behavior has been previously observed in topological semimetals, owing to the presence of a chiral current in the system [15,63,67]. The positive magnetoresistance for $\theta_{\rho} \gtrsim 2.5^{\circ}$ is due to the vanishing of the chiral anomaly influence and restoration of the orbital effect induced by the Lorentz force. Remarkably, the chiral anomaly appears to affect the magnetothermopower [Fig. 4(b)] in a similar manner. Namely, the negative slope of S(B) that occurs at a high magnetic field when $\theta_S = 0^\circ$ (θ_S



FIG. 3. (a) The electronic band structure of α -Sn in the presence of spin-orbit coupling using density functional theory (DFT) over the full Brillouin zone (BZ); the BZ is shown in (c); the closer look along $M \rightarrow \Gamma \rightarrow Z$ is shown in (b). (d) The two-dimensional (2D) band structure in the $(k_x - k_z)$ plane illustrates the position of two Dirac points at $(0,0, \pm k_z)$. (e) Magnetoresistance at $\theta = 0^\circ$ magnetic field orientations, influenced by Fermi surface topology.

is the angle between *B* and ∇T) becomes positive when *B* is away from ∇T . The threshold for θ_S , which is ~10°, is larger than that of θ_ρ , possibly because the relative contribution from out-of-plane positive magnetothermopower is not as large as that from orbital magnetoresistance.

The magnetic field breaks the time-reversal symmetry, leading to the formation of Weyl nodes in α -Sn through the degeneracy of the Dirac nodes. In the presence of parallel electric and magnetic fields, the imbalance in the number of Weyl fermions of different chirality leads to the generation of an additional current that contributes to the total electrical conductivity. In the semiclassical regime, this can be expressed as [68,69]

where $\sigma(0)$ is the Drude conductivity, $B_q = \frac{2E_F^2}{3e\hbar v_F^2}$ is the quantum magnetic field, and $\frac{\tau_i}{\tau}$ is the ratio of intervalley scattering time to the mean free time of charge carriers. The corresponding influence of the chiral anomaly on the thermoelectric power can be calculated using the Mott relation, which should obey if the chemical potential is larger than k_BT and the scattering is dominated by elastic processes [41]. If the mean free time is assumed to be independent of energy, the equation reads [68,69]

$$S(B) = S(0) - S(0)2\frac{\tau_i B^2}{3\tau B_a^2},$$
(3)

$$\sigma(B) = \sigma(0) + \sigma(0) \frac{1}{3} \frac{\tau_i}{\tau} \frac{B^2}{B_a^2},$$
(2)

where S(0) is the background thermopower unrelated to the chiral anomaly. Since the magnetic field can change the distance in k space between Weyl cones, e.g., Ref. [70], the



FIG. 4. (a) Resistivity (ρ_{xx}) in function of magnetic field (*B*) of Dirac semimetal α -Sn for selected angles (θ , where θ is the angle between *j* and *B*) at a constant temperature 60 K. (b) Magnetothermopower [$S_{xx}(B)$] of α -Sn for selected angles (θ , where θ is the angle between ∇T and *B*) at a constant temperature 60 K.



FIG. 5. (a) Conductivity (σ) in function of square of magnetic field (*B*) of α -Sn for several temperatures. For the sake of clarity, starting from $\sigma(B^2)$ for T = 38 K, the curves are successively shifted vertically by $10^2 \Omega^{-1} \text{ cm}^{-1}$ each for sake of clarity. (b) Normalized thermopower *S*(*B*) of α -Sn for selected temperatures. The dashed line in both panels shows the fit as calculated from Eqs. (2) and (3).

intervalley scattering time can in principle be expected to be field dependent. On the other hand, the model we used to describe the experimental data assumes τ_i to be field independent, which appears to be a sufficient approximation in the high field limit. Moreover, this agrees with optical studies of the Dirac semimetal Cd₃As₂, which concluded that chiral relaxation shows little field dependence [71]. Equations (2) and (3) describe the behavior of the nonoscillatory part of the respective signal, but this appears to be difficult to extract from field dependences of conductivity at low temperatures. This was done by simulating the SdH oscillations using the Lifshitz-Kosevich formula (see Fig. S4 in the SM [56]) up to B = 8 T and then extrapolating the oscillatory signal to higher fields. Finally, we subtract the oscillations to obtain the nonoscillatory part of $\sigma(B)$, However, the results of extrapolation were not perfect below T = 100 K; hence, we have chosen to restrict the upper limit of the fit to 10 T. As shown in Fig. 5(a), $\sigma(B)$ at this intermediate field follows a quadratic dependence on B expected for the chiral anomaly [41,68,69] and can be approximated with Eq. (2). Remarkably, the thermopower at a high magnetic field also appears to be well described by the model involving the chiral anomaly. Equation (3) predicts that the negative slope of the magnetothermopower is a B^2 -dependent component, which in fact appears in S(B) dependences, see Fig. 5(b). From the fitting of the magnetoresistance and magnetothermopower with Eqs. (2) and (3), we can estimate the ratio of the intervalley lifetime to the transport lifetime $\frac{\tau_i}{\tau}$, which is the parameter that defines whether an electronic system is in fact in the chiral limit [72]. The essential condition that must be met to observe the pumping of Weyl fermions from one node to another is $\tau_i > \tau$. Figure 6 presents the temperature dependences of $\frac{\tau_i}{\tau}$ calculated from $\sigma(B)$ and S(B), the latter multiplied by a prefactor $\frac{3}{2}$; the reason for that will be discussed later. The agreement between these two independent experimental results is good, i.e., the ratio $\frac{\tau_i}{\tau}$ decays with temperature, but both $\frac{\tau_i}{\tau}(T)$ do not coincide exactly. Similar behavior was also reported for the topological Dirac semimetal ZrTe₅ [68]. For α -Sn, the relaxation time ratio from thermoelectric data as calculated with Eq. (3) is smaller than one calculated from the electrical measurements. A likely reason for this discrepancy

may be the approximations made to derive Eq. (3). The first assumption is the energy (E) independent of carrier lifetime [68,69], which is usually not the case for real materials. The term in the Mott relations $\frac{d \ln \tau(E)}{dE}$ has significant contributions to the diffusion thermoelectric power [54]. For example, for metals at high temperature, the relation is expected to be $\tau(E) \propto E^{3/2}$ [73]. We include the $\frac{3}{2}$ factor in the relaxation time ratio calculated from thermoelectric data, but our experimental results suggest even stronger energy dependence. This in fact was suggested to be significantly enhanced in the Dirac material SnTe [74]. Another important assumption in Eq. (3) was the strict validity of the Mott relation [68]. Since the thermopower is measured under the condition of an imbalanced number but possibly also the energy of chiral Weyl fermions [75], some deviations from the Mott relation can be expected. An actual reason will be an interesting subject of further investigation.



FIG. 6. The ratio of intervalley Weyl scattering time to Drude relaxation time (τ_i/τ) of the α -Sn sample as a function of temperatures with the current (j) or thermal gradient (∇T) along with the magnetic field applied parallel to the *a* axis.

IV. CONCLUSIONS

We studied the transport properties of the Dirac semimetal α -Sn in the configuration where the magnetic field is nonorthogonal to the electric field or thermal gradient. At high field, we observed the NLMR and magnetothermopower, which also shows a negative slope. Both features, which vary with field like B^2 , can be attributed to the chiral anomaly and disappear at high temperatures. Further, a hint that we see manifestations of the chiral anomaly in α -Sn is a strong angular variation of the thermopower and resistivity. The calculated ratio of the intervalley scattering time to the mean free time satisfies the conditions for the chiral limit. We conclude that both the electrical and thermoelectric data indicate the presence of a Weyl system, which forms the chiral anomaly in the magnetic field.

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All of the relevant data that support the findings of this paper are available from the corresponding author upon reasonable request.

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