Semiclassical theory of anomalous transport in type-II topological Weyl semimetals

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Weyl semimetals possess low-energy excitations which act as monopoles of Berry curvature in momentum space. These emergent monopoles are at the heart of the many novel transport properties that Weyl semimetals exhibit. The singular nature of the Berry curvature around the nodal points in Weyl semimetals allows for the possibility of large anomalous transport coefficients in zero applied magnetic field. Recently, a new class, termed type-II Weyl semimetals, has been demonstrated in a variety of materials, where the Weyl nodes are tilted. We present here a theoretical study of anomalous transport in this new class of Weyl semimetals. We find that the parameter governing the tilt of these type-II Weyl points is intimately related to the zero-field transverse transport properties. We also find that the temperature dependence of the chemical potential plays an important role in determining how the transport coefficients can effectively probe the Berry curvature of the type-II Weyl points. In particular, we find that the transverse thermoelectric transport coefficient L_{xy}^{ET} is strongly enhanced with the tilt of the type-II Weyl nodes and with increasing temperature. We also discuss the experimental implications of our work for time-reversal breaking type-II Weyl semimetals.

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I. INTRODUCTION

Topological Weyl semimetals have sparked tremendous recent interest in condensed matter physics [1–3]. These materials host low-energy excitations with massless, linear dispersions, known as Weyl fermions [4]. A Weyl node is a monopole of Berry curvature, which acts as a magnetic field in momentum space. Weyl nodes must come in pairs of opposite chirality [5] and the sign of their monopole charge corresponds to their chirality. A direct consequence of the topological nature of the bulk Berry curvature in Weyl semimetals is the presence of topologically protected Fermi arcs that reside in the surface Brillouin zone and form open contours of states [6]. The bulk Weyl fermions and the surface Fermi arcs provide the key signatures of Weyl semimetals and are responsible for their many novel features.

Weyl fermions have been predicted in a variety of condensed matter systems [6–11] and were first experimentally realized in the transition-metal monopnictides, where signatures of the bulk nodes and the surface Fermi arcs were detected by angle-resolved photoemission spectroscopy [12–16]. Shortly after their discovery, a new class of Weyl semimetals, called type-II Weyl semimetals, was predicted [17–20] and experimentally discovered [21–24] in the transition-metal dichalcogenides MoTe₂ and WTe₂. Subsequently, a number of other examples of this new class of Weyl semimetals have been discovered [25–29]. In type-II Weyl semimetals, the Weyl nodes are tilted and therefore have a finite density of electrons and holes at the Weyl energy.

The bulk Berry curvature of Weyl fermions is known to result in a plethora of unique transport phenomena in Weyl semimetals. In parallel magnetic and electric fields, Weyl semimetals exhibit negative longitudinal magnetoresistance as a result of the chiral anomaly [30–36]. The surface Fermi arcs also lead to a remarkable mixed real- and momentum-space

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channel of charge transport [37–40] and have also been predicted to lead to a novel mechanism for entropy transport [41]. In bulk thermoelectric transport, Weyl semimetals have been predicted to have a number of distinct signatures [42–45], most notably a Nernst thermopower at zero applied magnetic field in time-reversal breaking Weyl semimetals [43]. Recent experiments have shown extraordinary thermoelectric properties of NbP, including a large Nernst effect [46] and evidence for a mixed axial-gravitational anomaly [47].

Although there have been some preliminary predictions of transport in type-II Weyl semimetals [18,48–50], it remains comparatively less well understood. There have been signatures of the chiral anomaly in WTe_{1.98} [51] as well as evidence of viscous electronic and thermal transport in the type-II Weyl semimetal WP₂ [52]. There are also strong candidates for type-II time-reversal breaking Weyl semimetals [29,53] and two such candidates, Mn₃Sn and Mn₃Ge, have shown tantalizing signatures of a large anomalous Hall effect [54–56]. Furthermore, experimental signatures of the anomalous Nernst effect and anomalous thermal Hall effect have also been detected in Mn₃Sn [57].

Although ferromagnetic metals are known to possess anomalous transport coefficients in zero field [58–60], Mn₃(Ge,Sn) is instead a weakly canted *antiferromagnet*. It has been suggested that the real-space magnetic texture can account for a large anomalous Hall effect in Mn₃Ge if the spins are noncoplanar [61], however, experiments have shown that the Mn₃(Ge,Sn) system does possess a large anomalous Hall effect in the planar magnetic phase with a Hall coefficient that is much larger than its weakly canted moment would suggest [54]. Thus, many puzzles remain.

Motivated in part by some of these experimental puzzles, we study anomalous transport in a lattice model of a time-reversal breaking Weyl semimetal [62]. The model we use allows for tuning through the type-I to type-II transition as well as between different type-II phases with distinct Fermi surface connectivities. Although the Berry curvature of a Weyl node is independent of its type, the occupation of states immediately surrounding the Weyl nodes is strongly dependent

on the precise Fermiology of the material. Furthermore, the chemical potential in low-density systems, such as semimetals, is a strong function of temperature [46,63]. A detailed understanding of the interplay between the tilt of the nodes, the connectivity of the Fermi pockets, and the temperature dependence of the chemical potential is required for a complete theoretical picture of anomalous transport in type-II Weyl semimetals.

One of the central results of our work is the strong enhancement of the transverse thermoelectric transport coefficient L_{xy}^{ET} with temperature and with increasing nodal tilt. This term is zero in the strictly type-I limit, but we find that by increasing the tilt of the Weyl nodes in the type-II regime, L_{xy}^{ET} increases due to the change in the net Berry curvature of occupied states. We also find that L_{xy}^{ET} is sensitive to many of the Lifshitz transitions that occur throughout the type-II Weyl semimetal regime. This has broad implications for the possible utilization of Weyl semimetals in thermoelectric applications.

The remainder of this paper is organized as follows. In Sec. II we present a time-reversal breaking model for a type-II Weyl semimetal that allows for tuning between various Fermi pocket connectivities. We discuss how the chemical potential depends on temperature for this model. In Sec. III, we investigate how the topological features determine anomalous transverse transport coefficients in the type-I and type-II regimes, including the crossover between them. For the model described in Sec. II, we calculate the anomalous Hall coefficient, the anomalous transverse thermoelectric

coefficient, and the thermal Hall coefficient. In Sec. IV, we discuss our results and conclude with experimental implications.

II. MODEL

In the continuum limit, a type-II Weyl node can be described by the following Hamiltonian:

$$\hat{H}_W = \gamma k_z \hat{\sigma}_0 + \chi \hbar v_F (\mathbf{k} - \mathbf{k}_W) \cdot \sigma, \tag{1}$$

where σ_0 is the 2×2 identity matrix, σ is the vector of Pauli matrices, γ describes the tilt of the node, \mathbf{k}_W is the momentum of the node, v_F is the Fermi velocity, taken to be a constant, and χ is the chirality of the node. The type-II to type-II transition for this continuum model occurs at $\gamma = \hbar v_F$. For $\gamma = 0$, the continuum model above has proven to be profoundly useful in understanding the properties of type-I Weyl semimetals. However, in the type-II case, this continuum model is manifestly unphysical. For $\gamma > \hbar v_F$, Eq. (1) describes an electron and a hole pocket that are unbounded and never close at large \mathbf{k} . Since various aspects of type-II Weyl semimetals are strongly dependent on the nature of the extended Fermi pockets surrounding the nodes [62], it is necessary to instead consider a lattice model.

In order to avoid the difficulties noted above caused by the unbounded Fermi pockets of continuum models for type-II Weyl semimetals, we consider the following model for a time-reversal symmetry-breaking system with two Weyl nodes:

$$\hat{H} = \gamma [\cos(k_z a) - \cos(k_0 a)] \hat{\sigma}_0 - 2t \sin(k_x a) \hat{\sigma}_1 - 2t \sin(k_y a) \hat{\sigma}_2 - [2t_z [\cos(k_z a) - \cos(k_0 a)] + m[2 - \cos(k_x a) - \cos(k_y a)] + \gamma_z [\cos(3k_z a) - \cos(3k_0 a)]] \hat{\sigma}_3,$$
(2)

where σ_0 is the 2 × 2 identity matrix, $\hat{\sigma}_j$ is the *j*th Pauli matrix, a is the lattice spacing, t, m, γ_z , and t_z are hopping amplitudes, k_0 sets the node separation, and γ sets the tilt of the Weyl nodes. This model supports two sets of electron and hole pockets, with each Weyl point being comprised of a separate pair of electron and hole pockets. The type-II to type-II Lifshitz transition happens at $\gamma = 2t_z - 3\gamma_z$.

A. Lifshitz transitions

In Fig. 1, we plot the energy band structure given by Eq. (2) for the parameters m=3t, $t_z=t$, $k_0a=\pi/2$, and $\gamma_z=0.5t$ for several values of γ . For these model parameters, the type-I to type-II transition occurs at $\gamma=0.5t$. The Fermi surfaces of the initial type-II band structure are shown in Fig. 1(f), where we see a pair of electron pockets (red) and hole pockets (blue) meeting at a pair of Weyl points. As γ increases, each of these pockets grow in size.

There is also a second set of topological Lifshitz transitions where the separate electron and hole pockets each merge into a single pocket. In Fig. 1(g), we see that the electron pockets have merged across the $k_z = \pi$ plane at $\gamma \approx 2t$ and a hole pocket within the electron pocket also forms for a small range of parameters within the electron pocket. For the model parameters above, the hole pockets similarly merge into a

single Fermi pocket across the $k_z=0$ plane at $\gamma\approx 2.5t$. The resulting band structure deeper in the type-II limit is shown in Fig. 1(h). The tunability of this model between these four different regimes makes it the minimal model to understand type-II Weyl semimetals. We will show how each regime has its own signature in anomalous transport.

B. Berry curvature

We can rewrite the Hamiltonian in Eq. (2) as $\hat{H} = d_0(\mathbf{k})\hat{\sigma}_0 + \mathbf{d}(\mathbf{k}) \cdot \sigma$ and upon doing so we can calculate the Berry curvature of the *n*th band through [64]

$$\Omega_{n,i}(\mathbf{k}) = \epsilon_{ijl} (-1)^n \frac{\mathbf{d} \cdot (\partial_{k_j} \mathbf{d} \times \partial_{k_l} \mathbf{d})}{2|\mathbf{d}|^3}, \tag{3}$$

where ϵ_{ijl} is the perfectly antisymmetric tensor. The Berry curvature plays a central role in the theory of anomalous transport. The parameter γ which determines the tilt, and therefore the type of the Weyl nodes, is embedded in $d_0(\mathbf{k})$, which does not enter Eq. (3). We note that due to the extended nature of the pockets in type-II Weyl semimetals, the anomalous transport coefficients are strongly dependent on the details of the Fermiology.

We can better understand the interplay between band Fermiology and the Berry curvature by introducing what we

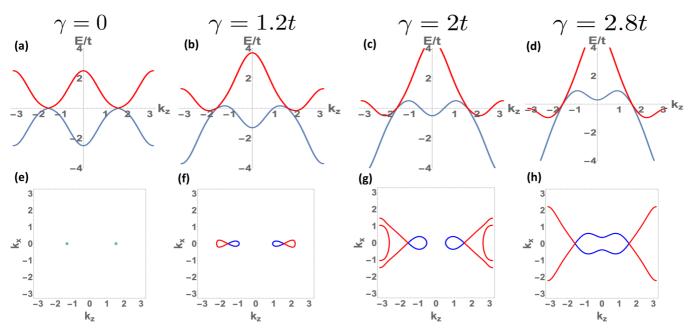


FIG. 1. Cuts through the band structure given by the Hamiltonian in Eq. (2). In (a)–(d), we show energy versus k_z cuts for $k_x = k_y = 0$. Here, we have chosen m = 3t; $t_z = t$; $k_0 a = \pi/2$; $\gamma_z = 0.5t$ for $\gamma = 0$ (a), $\gamma = 1.2t$ (b), $\gamma = 2t$ (c), and $\gamma = 2.8t$ (d). In (e)–(h), we show constant energy cuts for the band structure defined by Eq. (2). Panels (a) and (e) are in the type-I limit; (b) and (f) are in the type-II limit with distinct pockets making up each nodes; (c) and (g) are in the type-II limit after the electron pockets have merged; and (d) and (h) are in the type-II regime where the Weyl nodes share only a single electron and single hole pocket. Thus, as γ is increased, we can see the successive Lifshitz transitions described in the text.

call the net Berry curvature defined by

$$\Omega_z^{\text{net}}(\mathcal{E}) = \sum_n \int \frac{d\mathbf{S}}{(2\pi)^3} \frac{\Omega_{n,z}(\mathbf{k})}{|\nabla_{\mathbf{k}}\mathcal{E}|},\tag{4}$$

where \mathcal{E} is the energy and the integral in Eq. (4) is over surfaces of constant energy in the Brillouin zone. We plot $\Omega_z^{\rm net}(\mathcal{E})$ as a function of energy for various values of γ in Figs. 2(a)–2(d) for the model given by Eq. (2). We see that, for $\gamma=0$, $\Omega_z^{\rm net}(\mathcal{E})$ is a strictly odd function of energy. However, as γ increases, $\Omega_z^{\rm net}(\mathcal{E})$ has the same sign for positive and negative energies close to $\mathcal{E}=0$. As the electron pockets merge in Figs. 2(c), 2(g), and 2(k), we see that the net Berry curvature peaks. This interplay of Berry curvature and node tilt has a strong effect on zero-field transverse transport.

C. Temperature dependence of the chemical potential

In metals with high densities of electrons at degenerate temperatures, the chemical potential is nearly constant with respect to temperature. However, in low-density semimetals, it has been shown that the chemical potential has a strong temperature dependence at experimentally relevant temperatures [46,63]. For the lattice model in Eq. (2), we calculate the temperature dependence of the chemical potential by self-consistently solving for $\mu(T)$ for fixed density:

$$n = \int_{-\infty}^{\infty} d\mathcal{E} \frac{g(\mathcal{E})}{1 + e^{\frac{\mathcal{E} - \mu(T)}{k_B T}}},$$
 (5)

where n is the density, T is the temperature, and $g(\mathcal{E})$ is the density of states found through

$$g(\mathcal{E}) = -\frac{1}{\pi} \sum_{n} \operatorname{Im} \left[\int \frac{d^{3}k}{(2\pi)^{3}} G_{n}(\mathbf{k}, \mathcal{E}) \right], \tag{6}$$

where $G_n(\mathbf{k}, \mathcal{E})$ is the Green function of the *n*th band.

For isolated type-I Weyl points, the minimum of $g(\mathcal{E})$ will generically occur at the Weyl points [46] due to the symmetry of the particle and hole bands. This results in the chemical potential shifting to the Weyl points with increasing temperature in type-I Weyl semimetals. However, in general, the hole and electron pockets are not symmetric about the Weyl energy in type-II Weyl semimetals, and the minimum of $g(\mathcal{E})$ will occur above or below the Weyl nodes. In Figs. 2(e)–2(h) we plot the density of states $g(\mathcal{E})$ of the model in Eq. (2) for several values of the tilt parameter γ at energies around the node energy. For $\gamma = 0$, we see the density of states is minimum at the Weyl energy. However, for larger values of γ , in the type-II regime, the tilt of the Weyl cones breaks particle-hole symmetry and shifts the minimum of $g(\mathcal{E})$ away from the Weyl energy. In the large- γ limit where both the electron and hole pockets merge [Fig. 2(h)], the minimum of the density of states shifts far from the Weyl energy. In Figs. 2(i)-2(1) we show how the chemical potential evolves with temperature for each value of γ . For low values of γ in Figs. 2(i) and 2(j), we see that the chemical potential shifts to the Weyl energy roughly on a temperature scale of the distance E_F is away from the energy for which $g(\mathcal{E})$ is minimized. However, as the nodes become increasingly tilted, the temperature scale over which $\mu(T)$ shifts becomes much larger than the relevant scales in transport that we will consider.

D. Definition of transport coefficients

From Onsager's generalized transport equations [65], we have

$$\begin{pmatrix} \mathbf{J}^{e} \\ \mathbf{J}^{q} \end{pmatrix} = \begin{pmatrix} \mathbf{L}^{EE} & \mathbf{L}^{ET} \\ \mathbf{L}^{TE} & \mathbf{L}^{TT} \end{pmatrix} \cdot \begin{pmatrix} \mathbf{E} \\ -\nabla T \end{pmatrix}, \tag{7}$$

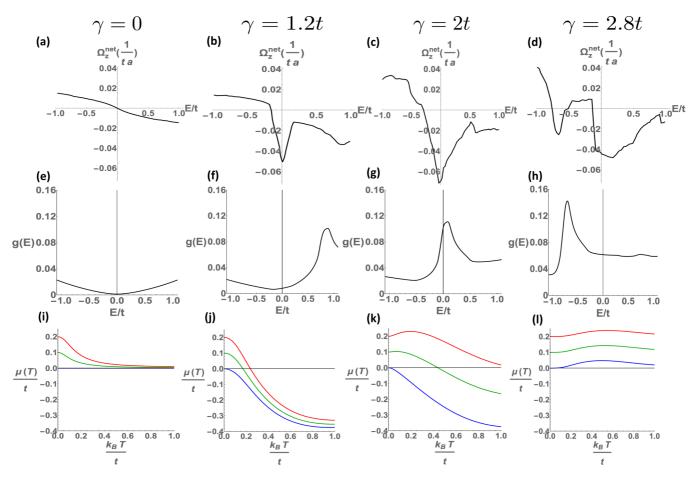


FIG. 2. Each column corresponds to a particular γ with $\gamma=0$ [(a), (e), (i)], $\gamma=1.2t$ [(b), (f), (j)], $\gamma=2t$ [(c), (g), (k)], and $\gamma=2.8t$ [(d), (h), (l)]. (a)–(d) Show the net Berry curvature in the z direction $\Omega_z^{\rm net}(\mathcal{E})$ defined by Eq. (4). We see that for nonzero γ , the net Berry curvature around the nodes is of the same sign. (e)–(h) The density of states for m=3t, $t_z=t$, $k_0a=\pi/2$, and $\gamma_z=0.5t$, for different values of the tilt parameter. (i)–(l) Illustrate the temperature dependence of the chemical potential $\mu(T)$. Each plot shows three separate values of E_F : $E_F=0$ (blue), $E_F=0.1$ (green), and $E_F=0.2$ (red). We see that for smaller values of γ , γ , γ , γ , γ , this minimum shifts far from the nodes. This has a strong effect on the shift of the chemical potential with temperature.

where \mathbf{J}^e is the charge current density, \mathbf{J}^q is the heat current density, \mathbf{E} is the electric field, and ∇T is the applied temperature gradient. In general, the transport coefficients in Eq. (7) must be obtained by solving for the nonequilibrium distribution function using the Boltzmann formalism. However, in the absence of an applied magnetic field, the situation simplifies dramatically. In this case, the modified velocity of the nth band is found to be

$$\dot{\mathbf{r}}_n = \mathbf{v}_n(\mathbf{k}) + \frac{e}{\hbar} (\mathbf{E} \times \Omega_n(\mathbf{k})), \tag{8}$$

where $\mathbf{v}_n(\mathbf{k}) \equiv \frac{1}{\hbar} \nabla_{\mathbf{k}} \mathcal{E}_n(\mathbf{k})$ is the usual group velocity and where $\frac{e}{\hbar}(\mathbf{E} \times \Omega_n(\mathbf{k}))$ is the anomalous velocity due to the Berry curvature [66]. The Berry curvature is sufficient to generate nonzero transverse transport coefficients L_{xy}^{EE} , L_{xy}^{ET} , and L_{xy}^{TT} in the presence of zero external magnetic field. We consider the effects in type-II Weyl semimetals on each of these coefficients below. For the remainder of this paper, we will only consider the case of zero magnetic field.

III. ANOMALOUS TRANSPORT

In this section, we consider anomalous transport in a two-band lattice model for the type-II time-reversal breaking Weyl semimetal described in Sec. II. We calculate the anomalous Hall coefficient L_{xy}^{EE} , the anomalous transverse thermoelectric coefficient L_{xy}^{ET} , and the anomalous thermal Hall coefficient L_{xy}^{TT} .

A. Anomalous Hall effect

The anomalous Hall effect has long been studied in the context of Weyl semimetals. For a time-reversal breaking Weyl semimetal, it was shown that the anomalous Hall conductivity L_{xy}^{EE} was directly proportional to the net separation between Weyl nodes [67]. In the presence of Berry curvature, we have seen that the equations of motion are modified and, as a result [59,66,68,69], the anomalous Hall coefficient takes the form

$$L_{xy}^{EE} = \frac{e^2}{\hbar} \sum_{n} \int \frac{d^3k}{(2\pi)^3} \Omega_{n,z}(\mathbf{k}) f_0(\mathbf{k}), \tag{9}$$

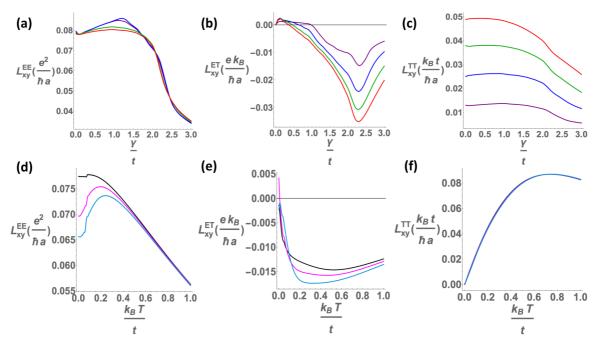


FIG. 3. In (a)–(c), we plot each anomalous transport coefficient L_{xy}^{EE} , L_{xy}^{ET} , and L_{xy}^{TT} for the Hamiltonian given by Eq. (2) with parameters m=3t, $t_z=t$, $k_0a=\pi/2$, and $\gamma_z=0.5t$, as a function of γ for the following temperatures: T=0.05t (purple), T=0.1t (blue), T=0.15t (green), and T=0.2t (red). In (d)–(f), we show L_{xy}^{EE} , L_{xy}^{ET} , and L_{xy}^{TT} for the same values as in (a)–(c), with $\gamma=1.2t$, plotted as functions of temperature for various Fermi energies: $E_F=0$ (black), $E_F=0.1t$ (magenta), and $E_F=0.2t$ (blue).

where $f_0(\mathbf{k}) = (1 + e^{\frac{\mathcal{E}(\mathbf{k}) - \mu}{k_B T}})^{-1}$ is the equilibrium Fermi-Dirac distribution. Since $f_0(\mathbf{k})$ only depends on \mathbf{k} through the energy \mathcal{E} , we can write Eq. (9) in terms of $\Omega_z^{\rm net}(\mathcal{E})$ in the following way:

$$L_{xy}^{EE} = \frac{e^2}{\hbar} \int d\mathcal{E} \,\Omega_z^{\text{net}}(\mathcal{E}) f_0(\mathcal{E}). \tag{10}$$

The anomalous Hall coefficient in Eq. (9) integrates the Berry curvature over all filled states. In the untilted regime $(\gamma=0)$, Eq. (2) will have particle-hole symmetry and since, from Eq. (3), the Berry curvature is opposite in sign but equal in magnitude at a particular point ${\bf k}$ for two bands. Therefore, equal energy Fermi surfaces at positive and negative energies will have opposite net integrated Berry curvature $\Omega_z^{\rm net}(\mathcal{E})$. Hence, at $\gamma=0$ and $E_F=0$, the filled states all contribute to one sign of L_{xy}^{EE} . As the tilt γ is increased, the Fermi surfaces become increasingly asymmetric, and the distributions of $\Omega_{n,z}({\bf k})$ over the occupied states undergo substantial change, as shown in Figs. 2(a)–2(d).

In Fig. 3(a), we plot the anomalous Hall coefficient L_{xy}^{EE} as a function of the tilt parameter γ for several fixed temperatures at $E_F = 0$. Other than a small increase in L_{xy}^{EE} for $\gamma < 1.5t$ for small temperatures, we note that the curves are substantially similar. We can understand the increase of L_{xy}^{EE} by noting that the asymmetry in the Berry curvature distribution enhances the Hall coefficient for small temperatures but that this effect is washed out as the temperature increases and the Fermi function broadens. When the separate electron pockets comprising the Weyl nodes merge into a single-electron pocket at $\gamma = 2t$, we see that the Hall coefficient uniformly decreases for all temperatures.

In Fig. 3(d), we show the temperature dependence of the anomalous Hall coefficient L_{xy}^{EE} as a function of temperature for fixed $\gamma = 1.2t$ for various Fermi energies E_F . We see that for $E_F = 0$, L_{xy}^{EE} decreases monotonically as a function of temperature. This is because as the temperature is raised, the Fermi function broadens and electronic states with the opposite sign of $\Omega_{n,z}(\mathbf{k})$ begin to become occupied. This is true for the type-I case as well. However, for $E_F \neq 0$, the Hall coefficient will in general be lower than its $E_F = 0$ value at $T \approx 0$. For $E_F > 0$ in Fig. 3(d), we see that L_{xy}^{EE} attains a maximum at T > 0. This is due to the movement of the chemical potential with temperature [see Figs. 2(i)-2(1)]. As the chemical potential crosses the Weyl energy, a maximum is attained in the anomalous Hall coefficient as a function of temperature. This effect is seen to a lesser extent in the type-I case, however, there are two distinct differences in the type-II regime:

- (i) In the type-I case, $\mu(T)$ always shifts to the node energy. However, in the type-II case, the minimum of the density of states $g(\mathcal{E})$ occurs generically at a different energy, causing $\mu(T)$ to cross the Weyl energy rather than approach it asymptotically. This leads to a sharper rise of L_{xy}^{EE} with temperature.
- (ii) Due to the higher density of states around the nodes, where the Berry curvature is stronger, the anomalous Hall coefficient is more sensitive to the shift of the chemical notential.

At higher temperatures, the chemical potential reaches the energy where the density of states is minimized and the temperature dependence of L_{xy}^{EE} is uniform across each value of E_F . Although here we have only shown the results for a single value of γ , the broad conclusions that we have drawn remain true. There is a peak in temperature where $\mu(T)$ crosses

the node energy. In the type-II regime, this increase over the T=0 value is, in general, greater than that in the type-I regime for other similar model parameters. However, we note that for values of γ such that the electron (or the hole pockets) have merged, the temperature dependence becomes much weaker.

B. Anomalous thermoelectric effect

In the presence of Berry curvature, the electronic wave function acquires an orbital magnetization that is responsible for nontrivial anomalous thermoelectric properties [70]. It is found that in this case, the anomalous transverse thermoelectric coefficient is given by

$$L_{xy}^{ET} = \frac{k_B e}{\hbar} \sum_{n} \int \frac{d^3 k}{(2\pi)^3} \Omega_{n,z}(\mathbf{k}) s(\mathbf{k}), \tag{11}$$

where $s(\mathbf{k})$ is the electronic entropy given by

$$s(\mathbf{k}) = -f_0(\mathbf{k}) \ln[f_0(\mathbf{k})] - [1 - f_0(\mathbf{k})] \ln[1 - f_0(\mathbf{k})], \quad (12)$$

and where $f_0(\mathbf{k})$ is the equilibrium Fermi-Dirac distribution defined above. The entropy, as defined in Eq. (12), is sharply peaked around $\mu(T)$. Therefore, unlike the anomalous Hall coefficient, the anomalous thermoelectric coefficient L_{xy}^{ET} is only sensitive to the Berry curvature around the chemical potential. As in the case of L_{xy}^{EE} above, we can also write Eq. (11) in terms of $\Omega_z^{\text{net}}(\mathcal{E})$ since $s(\mathbf{k})$ is also only a function of momentum through the energy. Doing so, we obtain

$$L_{xy}^{ET} = \frac{k_B e}{\hbar} \int d\mathcal{E} \,\Omega_z^{\text{net}}(\mathcal{E}) s(\mathcal{E}). \tag{13}$$

The tilt γ of the model for a type-II Weyl semimetal can lead to a change in the distribution of net Berry curvature that is large enough to flip the sign of $\Omega_z^{\text{net}}(\mathcal{E})$ above or below the Weyl node in energy, for energies small compared with the bandwidth $\mathcal{E} \ll t$, as shown in Figs. 2(b)–2(d). Hence, at energies just above and below the Weyl energy, $\Omega_z^{\text{net}}(\mathcal{E})$ has the same sign for a tilted Weyl cone. This leads to a large enhancement of the anomalous thermoelectric coefficient for energies around the node. We see this in Fig. 3(b), where L_{rv}^{ET} is plotted as a function of γ for various temperatures. In the type-I case, L_{xy}^{ET} is small in magnitude and positive in sign. After the Lifshitz transition to the type-II regime at $\gamma = 0.5t$, we see a change in sign of L_{xy}^{ET} and substantial increase in its magnitude. This is precisely because $\Omega_z^{\text{net}}(\mathcal{E})$ has the same sign and a large value. The anomalous thermoelectric coefficient L_{xy}^{ET} attains a maximum at $\gamma \approx 2.3t$ after the electron pockets have merged and just as the hole pockets are about to merge (with increasing γ). Hence, we see that measurable quantities, such as the Nernst effect, which depend on L_{xy}^{ET} may be quite sensitive to Lifshitz transitions between various regimes of Fermi pocket connectivity in type-II Weyl semimetals.

In Fig. 3(e), we plot the anomalous thermoelectric coefficient L_{xy}^{ET} as a function of temperature for various Fermi energies E_F at $\gamma=1.2t$. Here, we see that there is an increase with temperature in the magnitude of L_{xy}^{ET} for small T. This occurs because, as the entropy $s(\mathbf{k})$ broadens, L_{xy}^{ET} is enhanced from the entropy including a larger range of energies in the integral in Eq. (11). At a temperature on the order of the energy for which the density of states $g(\mathcal{E})$ attains its minimum,

 L_{xy}^{ET} reaches its maximum absolute value. For Fermi energies E_F farther from the node energy, we see that the maximum absolute value of L_{xy}^{ET} increases. This occurs for the same reason as in the anomalous Hall coefficient above; for a type-II Weyl semimetal, the chemical potential $\mu(T)$ will pass over the Weyl energy where $\Omega_{\tau}^{\rm net}(\mathcal{E})$ is enhanced.

Finally, we also note that since the Berry curvature is symmetric in magnitude but opposite in sign for an untilted type-I Weyl node, as shown in Fig. 2(a), at exactly $E_F = 0$, L_{xy}^{ET} must vanish. Hence, at precisely $E_F = 0$, L_{xy}^{ET} can only take a nonzero value for a type-II Weyl semimetal where the shifted occupation of states can lead to a nonzero net Berry curvature when Eq. (11) is evaluated at the node energy.

C. Anomalous thermal Hall effect

As in the anomalous transverse thermoelectric effect above, similar effects of orbital magnetization in the presence of Berry curvature also lead to an anomalous thermal Hall effect in the absence of magnetic field [71–73]. In the presence of Berry curvature, the anomalous thermal Hall coefficient is given by

$$L_{xy}^{TT} = \frac{k_B^2 T}{\hbar} \sum_{n} \int \frac{d^3 k}{(2\pi)^3} \Omega_{n,z}(\mathbf{k}) \left(\frac{\pi^2}{3} + \frac{(\mathcal{E} - \mu)^2}{(k_B T)^2} f_0(\mathbf{k}) - \ln\left(1 + e^{-\frac{\mathcal{E} - \mu(T)}{k_B T}}\right) + 2\text{Li}_2(1 - f_0(\mathbf{k})) \right),$$
(14)

where $f_0(\mathbf{k})$ is again the equilibrium Fermi-Dirac distribution and where $\text{Li}_m(z)$ is the polylogarithm function of order m defined by

$$\operatorname{Li}_{m}(z) = \sum_{k=1}^{\infty} \frac{z^{k}}{k^{m}}.$$
 (15)

Recognizing that, aside from $\Omega_{n,z}(\mathbf{k})$, the integrand of Eq. (14) only depends on the energy, we can write

$$L_{xy}^{TT} = \frac{k_B^2 T}{\hbar} \int d\mathcal{E} \,\Omega_z^{\text{net}}(\mathcal{E}) \left(\frac{\pi^2}{3} + \frac{(\mathcal{E} - \mu)^2}{(k_B T)^2} f_0(\mathcal{E})\right)$$
$$-\ln\left(1 + e^{-\frac{\mathcal{E} - \mu(T)}{k_B T}}\right) + 2\text{Li}_2(1 - f_0(\mathcal{E})). \tag{16}$$

Like the anomalous Hall coefficient, the anomalous thermal Hall coefficient defined by Eq. (14) integrates over many states below the chemical potential. However, unlike the simple Fermi distribution $f_0(\mathbf{k})$, the kernel of the integrand multiplying $\Omega_{n,z}(\mathbf{k})$ in Eq. (14) has a broader inflection point at $\mathcal{E} = \mu(T)$ than $f_0(\mathbf{k})$ and, hence, L_{xy}^{TT} is sensitive to Berry curvature over a wider range than L_{xy}^{EE} . In Fig. 3(c), we plot L_{xy}^{TT} as a function of γ and we see that it is quite similar with the anomalous Hall coefficient plotted in Fig. 3(a). However, we note that the anomalous thermal Hall coefficient increases with increasing γ for all temperatures up to the Lifshitz transition where the electron pockets merge. We also see that after the Fermi pockets do merge, the decrease of L_{xy}^{TT} is less rapid than L_{xy}^{EE} with increasing γ . We also note that the temperature dependence of L_{xy}^{TT} is increasing rather than decreasing as in L_{xy}^{EE} at large T. Finally, the dependence of L_{xy}^{TT} on E_F

	Sensitive only to states near μ	Enhanced by γ	Enhanced with T	E_F dependence
L_{xy}^{EE}	No	No	For $T \lesssim E_F$	Yes
L_{xy}^{ET}	Yes	Yes	For $T \lesssim E_F$	Yes
L_{xy}^{EE} L_{xy}^{ET} L_{xy}^{TT}	No	No	Yes	No

TABLE I. Summary of properties of anomalous transport coefficients in type-II Weyl semimetals.

is essentially negligible due to the broadened kernel in the integrand in Eq. (14).

IV. RELATION TO MEASURABLE QUANTITIES

In the previous section, we studied transverse transport in zero magnetic field and calculated the transport coefficients L_{xy}^{EE} , L_{xy}^{ET} , and L_{xy}^{TT} . From Eq. (7), we immediately recognize L_{xx}^{EE} as simply the electrical conductivity. Under isothermal conditions, the Nernst effect is defined by

$$\alpha_{xyz} = \frac{E_y}{-\nabla_x T},\tag{17}$$

under the conditions

$$J_x = J_y = 0 \tag{18}$$

and

$$\nabla_{\mathbf{v}}T = 0. \tag{19}$$

From Eq. (7), we can find α_{xyz} to be given by

$$\alpha_{xyz} = \frac{E_y}{-\nabla_x T} = \frac{L_{xx}^{EE} L_{xy}^{ET} - L_{xy}^{EE} L_{xx}^{ET}}{\left(L_{xx}^{EE}\right)^2 + \left(L_{xy}^{EE}\right)^2}.$$
 (20)

We note that Eq. (20) depends not only on the transverse coefficients L_{xy}^{EE} and L_{xy}^{ET} , but also on the longitudinal coefficients L_{xx}^{EE} and L_{xx}^{ET} . We can find the transverse thermal conductivity to be

$$\kappa_{xy} = L_{xy}^{TT} - T \frac{\left[\left(L_{xy}^{ET} \right)^2 - \left(L_{xx}^{ET} \right)^2 \right] L_{xy}^{EE} + 2 L_{xx}^{EE} L_{xy}^{ET} L_{xx}^{ET}}{\left(L_{xx}^{EE} \right)^2 + \left(L_{xy}^{EE} \right)^2}.$$
(21)

Similar to the Nernst thermopower given above, we note that the thermal conductivity κ_{xy} depends on L_{xy}^{TT} as well as the second, explicitly temperature-dependent term which we recognize as the ambipolar thermal conductivity that originates from a Seebeck effect generated by a Peltier current. Thus, we expect the effects of the nodal tilt to be strongly reflected in each of anomalous Hall transport, the anomalous Nernst thermopower, and in the anomalous thermal Hall effect.

So far, we have only considered the transverse transport coefficients in zero applied external magnetic field. However, in general, each $L_{ij}^{\alpha\beta}$ will be nonzero. Furthermore, all transverse coefficients will have additional contributions at nonzero applied magnetic field. Full expressions for the transport coefficients of lattice models for Weyl semimetals can be found in Ref. [43]. All contributions other than the anomalous terms discussed in the previous section will depend explicitly on the transport scattering time. From Matthiessen's rule, the scattering rate from independent processes can be written as

a sum of the scattering rates of each process. We expect that impurity scattering, electron-electron scattering, and phonon-induced scattering will all play a role in general. The various contributions to the scattering time in Weyl semimetals have been investigated previously [74–77], however, the nature of scattering in type-II Weyl semimetals remains an open problem. We have limited the scope of our work to the Berry curvature induced anomalous transport and so we leave the effects of scattering for future investigations.

V. DISCUSSION AND CONCLUSION

We have calculated how the anomalous transport coefficients reveal signatures of type-II Weyl semimetals. Our results are summarized in Table I. In particular, the anomalous transverse thermoelectric coefficient L_{xy}^{ET} is greatly enhanced by tilting the Weyl nodes. However, we have also seen that even deeper in the type-II phase, the various anomalous coefficients are sensitive to changes in Fermi surface topology even quite far from the Weyl nodes. Each of L_{xy}^{EE} and L_{xy}^{TT} show a marked decrease in magnitude as the electron pockets merge, while L_{xy}^{ET} peaks just after they merge and L_{xy}^{ET} decreases for γ large enough that the hole pockets similarly merge.

Previous calculations have shown [62] that tuning through the various Lifshitz transitions within the type-II Weyl semimetal phase is associated with changing connectivities of the topological Fermi arcs. The bulk Fermi surface Berry curvature is deeply linked to the topology of the bulk band structure, and it is enlightening to see the anomalous transport coefficients reflect this bulk-boundary correspondence. In some inversion-breaking Weyl semimetals [28], it has been shown that strain may be possible to tune between a type-I to type-II phase transition. Although this has not yet been demonstrated in a time-reversal breaking Weyl semimetal, it should be possible in principle. This would allow for an experimental verification of our results as well as allowing for deeper explorations of connections between anomalous transport and changes in Fermi arc topology. We note that the coefficients that we have calculated in this work are manifestly independent of scattering time. However, the experimentally measurable quantities, such as the Nernst thermopower, will necessarily involve the longitudinal transport coefficients, such as L_{xx}^{EE} and L_{xx}^{ET} , which depend on scattering

We have also seen that the temperature dependence of the anomalous transport coefficients is quite distinct for each of L_{xy}^{EE} , L_{xy}^{ET} , and L_{xy}^{TT} . The temperature dependence of the chemical potential has already been demonstrated to lie at the heart of the ordinary Nernst thermopower in the Weyl semimetal NbP [46]. However, similar effects in zero-field anomalous transport have remained, until this point,

unexplored. We have demonstrated that, through the location of the Fermi energy E_F , the temperature at which L_{xy}^{EE} attains its maximum can be tuned, as can the strength of the anomalous thermoelectric coefficient L_{xy}^{ET} .

The generation of a transverse response to an applied electric field or thermal gradient unifies the zoo of the various Hall effects. The ability to generate such a response in the absence of an externally applied magnetic field opens the door to a wide variety of technological applications. Weyl semimetals have been predicted to generate various anomalous transport phenomena in zero field due to their Berry curvature. However, the lack of experimental realizations of time-reversal breaking Weyl semimetals has stymied their application. Recently, it has been proposed that time-reversal breaking Weyl semimetal candidates are type-II in nature. Our calcu-

lations serve not only to guide future experiments, but also to demonstrate that through changing various experimentally accessible properties, it may be possible to tune the anomalous transport coefficients in type-II Weyl semimetal to an extent not possible in the type-I case.

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- [1] B. Yan and C. Felser, Annu. Rev. Condens. Matter Phys. 8, 337 (2017).
- [2] M. Z. Hasan, S.-Y. Xu, I. Belopolski, and C.-M. Huang, Annu. Rev. Condens. Matter Phys. 8, 289 (2017).
- [3] N. P. Armitage, E. J. Mele, and A. Vishwanath, arXiv:1705.01111.
- [4] H. Weyl, Proc. Natl. Acad. Sci. USA 15, 323 (1929).
- [5] H. Nielsen and M. Ninomiya, Phys. Lett. B 105, 219 (1981).
- [6] X. Wan, A. M. Turner, A. Vishwanath, and S. Y. Savrasov, Phys. Rev. B 83, 205101 (2011).
- [7] A. A. Burkov and L. Balents, Phys. Rev. Lett. 107, 127205 (2011).
- [8] G. Xu, H. Weng, Z. Wang, X. Dai, and Z. Fang, Phys. Rev. Lett. **107**, 186806 (2011).
- [9] G. Volovik and M. Zubkov, Nucl. Phys. B **881**, 514 (2014).
- [10] H. Weng, C. Fang, Z. Fang, B. A. Bernevig, and X. Dai, Phys. Rev. X 5, 011029 (2015).
- [11] S.-M. Huang, S.-Y. Xu, I. Belopolski, C.-C. Lee, G. Chang, B. Wang, N. Alidoust, G. Bian, M. Neupane, C. Zhang *et al.*, Nat. Commun. 6, 7373 (2015).
- [12] S.-Y. Xu, I. Belopolski, N. Alidoust, M. Neupane, G. Bian, C. Zhang, R. Sankar, G. Chang, Z. Yuan, C.-C. Lee *et al.*, Science 349, 613 (2015).
- [13] B. Q. Lv, N. Xu, H. M. Weng, J. Z. Ma, P. Richard, X. C. Huang, L. X. Zhao, G. F. Chen, C. E. Matt, F. Bisti *et al.*, Nat. Phys. 11, 724 (2015).
- [14] S.-Y. Xu, N. Alidoust, I. Belopolski, Z. Yuan, G. Bian, T.-R. Chang, H. Zheng, V. N. Strocov, D. S. Sanchez, G. Chang *et al.*, Nat. Phys. 11, 748 (2015).
- [15] S.-Y. Xu, I. Belopolski, D. S. Sanchez, C. Zhang, G. Chang, C. Guo, G. Bian, Z. Yuan, H. Lu, T.-R. Chang *et al.*, Sci. Adv. 1, e1501092 (2015).
- [16] Z. K. Liu, L. X. Yang, Y. Sun, T. Zhang, H. Peng, H. F. Yang, C. Chen, Y. Zhang, Y. F. Guo, D. Prabhakaran *et al.*, Nat. Mater. 15, 27 (2016).
- [17] M. O. Goerbig, J.-N. Fuchs, G. Montambaux, and F. Piéchon, Phys. Rev. B 78, 045415 (2008).
- [18] A. A. Soluyanov, D. Gresch, Z. Wang, Q. Q.-S. Wu, M. Troyer, X. Dai, and B. A. Bernevig, Nature (London) 527, 495 (2015).

- [19] Z. Wang, D. Gresch, A. A. Soluyanov, W. Xie, S. Kushwaha, X. Dai, M. Troyer, R. J. Cava, and B. A. Bernevig, Phys. Rev. Lett. 117, 056805 (2016).
- [20] Y. Sun, S. C. Wu, M. N. Ali, C. Felser, and B. Yan, Phys. Rev. B 92, 161107 (2015).
- [21] F. Y. Bruno, A. Tamai, Q. S. Wu, I. Cucchi, C. Barreteau, A. de La Torre, S. McKeown Walker, S. Ricco, Z. Wang, T. K. Kim et al., Phys. Rev. B 94, 121112 (2016).
- [22] L. Huang, T. M. McCormick, M. Ochi, Z. Zhao, M.-T. Suzuki, R. Arita, Y. Wu, D. Mou, H. Cao, J. Yan *et al.*, Nat. Mater. 15, 1155 (2016).
- [23] Y. Wu, D. Mou, N. H. Jo, K. Sun, L. Huang, S. L. Bud'Ko, P. C. Canfield, and A. Kaminski, Phys. Rev. B 94, 121113 (2016).
- [24] C. Wang, Y. Zhang, J. Huang, S. Nie, G. Liu, A. Liang, Y. Zhang, B. Shen, J. Liu, C. Hu et al., Phys. Rev. B 94, 241119(R) (2016).
- [25] G. Autès, D. Gresch, M. Troyer, A. A. Soluyanov, and O. V. Yazyev, Phys. Rev. Lett. 117, 066402 (2016).
- [26] S.-Y. Xu, N. Alidoust, G. Chang, H. Lu, B. Singh, I. Belopolski, D. S. Sanchez, X. Zhang, G. Bian, H. Zheng *et al.* (Science Advances, 2017), Vol. 3, p. 1603266.
- [27] K. Koepernik, D. Kasinathan, D. V. Efremov, S. Khim, S. Borisenko, B. Büchner, and J. van den Brink, Phys. Rev. B 93, 201101 (2016).
- [28] G. Chang, S.-Y. Xu, D. S. Sanchez, S.-M. Huang, C.-C. Lee, T.-R. Chang, G. Bian, H. Zheng, I. Belopolski, N. Alidoust et al., Sci. Adv. 2, e1600295 (2016).
- [29] S. Borisenko, D. Evtushinsky, Q. Gibson, A. Yaresko, T. Kim, M. N. Ali, B. Buechner, M. Hoesch, and R. J. Cava, arXiv:1703.04700.
- [30] K.-S. Kim, H.-J. Kim, and M. Sasaki, Phys. Rev. B 89, 195137 (2014)
- [31] D. T. Son and B. Z. Spivak, Phys. Rev. B 88, 104412 (2013).
- [32] P. Goswami, J. H. Pixley, and S. Das Sarma, Phys. Rev. B **92**, 075205 (2015).
- [33] E. V. Gorbar, V. A. Miransky, and I. A. Shovkovy, Phys. Rev. B 89, 085126 (2014).
- [34] C.-L. Zhang, S.-Y. Xu, I. Belopolski, Z. Yuan, Z. Lin, B. Tong, G. Bian, N. Alidoust, C.-C. Lee, S.-M. Huang *et al.*, Nat. Commun. **7**, 10735 (2016).

- [35] X. Huang, L. Zhao, Y. Long, P. Wang, D. Chen, Z. Yang, H. Liang, M. Xue, H. Weng, Z. Fang, X. Dai, and G. Chen, Phys. Rev. X 5, 031023 (2015).
- [36] M. Hirschberger, S. Kushwaha, Z. Wang, Q. Gibson, S. Liang, C. A. Belvin, B. A. Bernevig, R. J. Cava, and N. P. Ong, Nat. Mater. 15, 1161 (2016).
- [37] A. C. Potter, I. Kimchi, and A. Vishwanath, Nat. Commun. 5, 5161 (2014).
- [38] J. Borchmann and T. Pereg-Barnea, Phys. Rev. B 96, 125153 (2017).
- [39] Y. Zhang, D. Bulmash, P. Hosur, A. C. Potter, and A. Vishwanath, Sci. Rep. 6, 23741 (2016).
- [40] P. J. W. Moll, N. L. Nair, T. Helm, A. C. Potter, I. Kimchi, A. Vishwanath, and J. G. Analytis, Nature (London) 535, 266 (2016).
- [41] T. M. McCormick, S. J. Watzman, J. P. Heremans, and N. Trivedi, arXiv:1703.04606.
- [42] R. Lundgren, P. Laurell, and G. A. Fiete, Phys. Rev. B 90, 165115 (2014).
- [43] G. Sharma, P. Goswami, and S. Tewari, Phys. Rev. B 93, 035116 (2016).
- [44] Q. Chen and G. A. Fiete, Phys. Rev. B 93, 155125 (2016).
- [45] G. Sharma, C. Moore, and S. Tewari, Phys. Rev. B 96, 195119 (2017).
- [46] S. Watzman, T. M. McCormick, C. Sekhar, S.-C. Wu, Y. Sun, A. Prakash, C. Felser, N. Trivedi, and J. P. Heremans, arXiv:1703.04700.
- [47] J. Gooth, A. C. Niemann, T. Meng, A. G. Grushin, K. Landsteiner, B. Gotsmann, F. Menges, M. Schmidt, C. Shekhar, V. Sueß *et al.*, Nature (London) **547**, 324 (2017).
- [48] T. E. O'Brien, M. Diez, and C. W. J. Beenakker, Phys. Rev. Lett. **116**, 236401 (2016).
- [49] G. Sharma, P. Goswami, and S. Tewari, Phys. Rev. B 96, 045112 (2017).
- [50] A. A. Zyuzin and R. P. Tiwari, JETP Lett. 103, 717 (2016).
- [51] Y.-Y. Lv, X. Li, B.-B. Zhang, W. Y. Deng, S.-H. Yao, Y. B. Chen, J. Zhou, S.-T. Zhang, M.-H. Lu, L. Zhang *et al.*, Phys. Rev. Lett. 118, 096603 (2017).
- [52] J. Gooth, F. Menges, C. Shekhar, V. Süß, N. Kumar, Y. Sun, U. Drechsler, R. Zierold, C. Felser, and B. Gotsmann, arXiv:1706.05925.
- [53] H. Yang, Y. Sun, Y. Zhang, W.-J. Shi, S. S. P. Parkin, and B. Yan, New J. Phys. 19, 015008 (2017).
- [54] S. Nakatsuji, N. Kiyohara, and T. Higo, Nature 527, 212 (2015).
- [55] N. Kiyohara, T. Tomita, and S. Nakatsuji, Phys. Rev. Applied 5, 064009 (2016).

- [56] A. K. Nayak, J. E. Fischer, Y. Sun, B. Yan, J. Karel, A. C. Komarek, C. Shekhar, N. Kumar, W. Schnelle, J. Kübler, C. Felser, and S. S. P. Parkin, Sci. Adv. 2, e1501870 (2016).
- [57] X. Li, L. Xu, L. Ding, J. Wang, M. Shen, X. Lu, Z. Zhu, and K. Behnia, Phys. Rev. Lett. 119, 056601 (2017).
- [58] M. Onoda and N. Nagaosa, J. Phys. Soc. Jpn. 71, 19 (2002).
- [59] N. Nagaosa, J. Sinova, S. Onoda, A. H. MacDonald, and N. P. Ong, Rev. Mod. Phys. 82, 1539 (2010).
- [60] S. J. Watzman, R. A. Duine, Y. Tserkovnyak, S. R. Boona, H. Jin, A. Prakash, Y. Zheng, and J. P. Heremans, Phys. Rev. B 94, 144407 (2016).
- [61] J. Kübler and C. Felser, Europhys. Lett. 108, 67001 (2014).
- [62] T. M. McCormick, I. Kimchi, and N. Trivedi, Phys. Rev. B 95, 075133 (2017).
- [63] Y. Wu, N. H. Jo, M. Ochi, L. Huang, D. Mou, S. L. Bud'ko, P. C. Canfield, N. Trivedi, R. Arita, and A. Kaminski, Phys. Rev. Lett. 115, 166602 (2015).
- [64] A. Bernevig and T. L. Hughes, Topological Insulators and Topological Superconductors (Princeton University Press, Princeton, NJ, 2013).
- [65] T. C. Harman and J. M. Honig, Thermoelectric and Thermomagnetic Effects and Applications (McGraw-Hill, New York, 1967)
- [66] D. Xiao, M.-C. Chang, and Q. Niu, Rev. Mod. Phys. 82, 1959 (2010).
- [67] K.-Y. Yang, Y.-M. Lu, and Y. Ran, Phys. Rev. B 84, 075129 (2011).
- [68] R. Karplus and J. M. Luttinger, Phys. Rev. 95, 1154 (1954).
- [69] J. M. Luttinger, Phys. Rev. 112, 739 (1958).
- [70] D. Xiao, Y. Yao, Z. Fang, and Q. Niu, Phys. Rev. Lett. 97, 026603 (2006)
- [71] D. L. Bergman and V. Oganesyan, Phys. Rev. Lett. 104, 066601 (2010).
- [72] T. Yokoyama and S. Murakami, Phys. Rev. B 83, 161407 (2011).
- [73] T. Qin, Q. Niu, and J. Shi, Phys. Rev. Lett. 107, 236601 (2011).
- [74] S. Das Sarma, E. H. Hwang, and H. Min, Phys. Rev. B 91, 035201 (2015).
- [75] J. H. Pixley, Y.-Z. Chou, P. Goswami, D. A. Huse, R. Nand-kishore, L. Radzihovsky, and S. Das Sarma, Phys. Rev. B 95, 235101 (2017).
- [76] D. Culcer, A. Sekine, and A. H. MacDonald, Phys. Rev. B 96, 035106 (2017).
- [77] A. Sekine, D. Culcer, and A. H. MacDonald, arXiv:1706.01200v2.