Inhomogeneous Weyl and Dirac Semimetals: Transport in Axial Magnetic Fields and Fermi Arc Surface States from Pseudo-Landau Levels

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Topological Dirac and Weyl semimetals have an energy spectrum that hosts Weyl nodes appearing in pairs of opposite chirality. Topological stability is ensured when the nodes are separated in momentum space and unique spectral and transport properties follow. In this work, we study the effect of a spacedependent Weyl node separation, which we interpret as an emergent background axial-vector potential, on the electromagnetic response and the energy spectrum of Weyl and Dirac semimetals. This situation can arise in the solid state either from inhomogeneous strain or nonuniform magnetization and can also be engineered in cold atomic systems. Using a semiclassical approach, we show that the resulting axial magnetic field \mathbf{B}_5 is observable through an enhancement of the conductivity as $\sigma \sim \mathbf{B}_2^2$ due to an underlying chiral pseudomagnetic effect. We then use two lattice models to analyze the effect of \mathbf{B}_5 on the spectral properties of topological semimetals. We describe the emergent pseudo-Landau-level structure for different spatial profiles of \mathbf{B}_5 , revealing that (i) the celebrated surface states of Weyl semimetals, the Fermi arcs, can be reinterpreted as n = 0 pseudo-Landau levels resulting from a **B**₅ confined to the surface, (ii) as a consequence of position-momentum locking, a bulk B_5 creates pseudo-Landau levels interpolating in real space between Fermi arcs at opposite surfaces, and (iii) there are equilibrium bound currents proportional to \mathbf{B}_{5} that average to zero over the sample, which are the analogs of bound currents in magnetic materials. We conclude by discussing how our findings can be probed experimentally.

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I. INTRODUCTION

Electronic and lattice degrees of freedom are inevitably intertwined in solid-state physics [1]. With the advent of graphene [2], a remarkable effect was soon acknowledged: Elastic deformations of the lattice originating from strain can couple to the low-energy Dirac quasiparticles of graphene as a pseudomagnetic or axial-magnetic-vector potential [3–5]. In order to preserve time-reversal symmetry, the axial-vector potential couples to the two valleys with an opposite sign. Spatially inhomogeneous strain generates an effective axial magnetic field \mathbf{B}_5 and gives rise to a pseudo-Landau-level spectrum at low energies [6]. Strain-induced pseudo-Landau levels have been observed with scanning-tunneling microscopy both in real [7] and artificial graphene [8,9]. The realization of Landau levels without real external magnetic fields results in effective fields as high as 300 T and is a direct consequence of

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the Dirac nature of the carriers, exemplifying the unique response of this class of systems to strain.

In this respect, Wevl and Dirac semimetals in three dimensions (3D) [10,11] are expected to behave as 3D cousins of graphene and host similar effects. A Weyl semimetal is a state with pairs of band-touching points with linear dispersion, also called Weyl nodes. The Weyl nodes are sources and sinks of Berry curvature, i.e., Berry curvature monopoles, in momentum space. The charge of a monopole defines the chirality of the corresponding node, and the two partners of a pair of nodes must have opposite chirality. Since monopoles can only annihilate in pairs, Weyl nodes are topologically protected as long as they are separated in momentum space by a vector **b**. Dirac semimetals can then be regarded as special cases of Weyl semimetals, where nodes of opposite chiralities are located at the same momentum but additional symmetries constrain the system to remain gapless.

The vector **b** can be alternatively interpreted as an axial gauge field since it couples with an opposite sign to Weyl nodes of opposite chirality [12–17]. This interpretation suggests that if **b** is varied in space, it will generate a nonzero axial magnetic field $\mathbf{B}_5 = \nabla \times \mathbf{b}$ that couples to

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fermions of opposite chirality with an opposite sign. This observation, together with the recent theoretical evidence that effective gauge fields can emerge in strained Weyl semimetals [18–21], is an important motivation of our study.

The main purpose of this work is therefore to address the effect of an axial magnetic field \mathbf{B}_5 arising in any type of Weyl and Dirac semimetal. We first discuss that the physical mechanism for the emergence of B_5 depends on the presence or absence of time-reversal symmetry. When a Weyl semimetal preserves time-reversal symmetry, each pair of nodes has a time-reversed partner (hence, the minimal number of nodes is four). In this case, \mathbf{B}_5 can be generated by an inhomogeneous strain profile and has an opposite sign for each time-reversed pair of nodes such that time-reversal symmetry is preserved. In contrast, timereversal breaking Weyl semimetals are not subjected to this constraint, and B₅ can emerge either from an inhomogeneous magnetization or strain [13,18]. Then, a richer set of phenomena can occur, with the example of a finite angular momentum for the electronic states [22].

Here, we naturally unify these mechanisms under a single framework by considering any Weyl semimetal in the presence of a spatially varying Weyl node separation. We discuss the effect of inhomogeneous nodal separation on (i) the changes in the spectrum due to the emergence of pseudo-Landau levels and (ii) transport and, in particular, the effect on the conductivity. In exploring (i), we will show that the Fermi arcs, the characteristic surface states of topological semimetals [23,24], can be reinterpreted as the chiral n = 0 pseudo-Landau level associated with a large \mathbf{B}_5 confined to the surface. On the other hand, the presence of a finite \mathbf{B}_5 in the bulk also creates pseudo-Landau levels. The n = 0 pseudo-Landau level of the bulk is then smoothly connected to the surface arcs. We find that this follows from a known phenomenon in the two-dimensional quantum Hall effect, sometimes referred to as positionmomentum locking; the average center position of a Landau-level wave function in one planar direction, say $\langle y \rangle$, is determined by its momentum in the perpendicular direction, k_x . Such a concept, considered recently in some detail for real magnetic fields in Weyl semimetals [25,26], will also be of use here. We find that these results significantly depart from the naive expectation gained from studies of strained graphene [3-5] from which the bulk zeroth pseudo-Landau level is expected to have opposite chirality with respect to the Fermi arcs [13,27]. Regarding (ii), we will show, based on both field theoretic arguments and a rigorous semiclassical calculation, that a finite bulk \mathbf{B}_5 enhances the conductivity of a Weyl semimetal as $\sigma \sim \mathbf{B}_{5}^{2}$. This response is related to a charge anomaly, much like the topological negative magnetoresistance $\sigma \sim \mathbf{B}^2$ is rooted in the chiral anomaly [28].

Moreover, we find that a time-reversal breaking Weyl semimetal may support bound currents originating in a chiral pseudomagnetic effect driven by \mathbf{B}_5 . This effect

exists in equilibrium, unlike the standard chiral magnetic effect [14–16,29–33] which is strictly zero in a nondy-namical situation [34–38].

The advances made on the experimental front in recent years allows us to explore the feasibility and detectability of these effects while considering a wide range of possible platforms. While Dirac and Weyl semimetal material growth thrives in condensed matter [39–68], they could also be engineered with cold atoms [69]. This provides access to inversion-breaking semimetals that are currently the standard in the solid state (with notable potential exceptions [67,68]), as well as to time-reversal breaking semimetals in cold atoms. We end this paper with an estimate of the magnitude of these effects based on real material parameters and discuss how the phenomena described here might be detected.

The structure of this paper is as follows. In Sec. II, we review how inhomogeneous strain (magnetization) in timereversal-invariant (broken) Weyl and Dirac semimetals leads to an effective axial magnetic field or, alternatively, a space-dependent Weyl node separation. In Sec. III, we predict the enhancement of the conductivity of Weyl and Dirac semimetals due to a uniform axial magnetic field and relate it to the underlying associated chiral pseudomagnetic effect. The latter leads to bound currents flowing within the material and along the boundary, even in equilibrium, as well as an enhanced bulk longitudinal conductivity. In Sec. IV, we introduce two lattice models that illustrate the spectral breakdown into pseudo-Landau levels and explicitly calculate the bound current distribution within a finite sample. We describe position-momentum locking and show that any equilibrium bulk current is compensated by surface currents. As an example of the relevance of our findings to different contexts, we discuss the implications of our results on the situation where two Weyl semimetals with different Weyl node separation are brought into proximity. Finally, in Sec. V, we provide a discussion and proposals to experimentally observe the discussed phenomena.

II. EMERGENCE OF SPACE-DEPENDENT NODE SEPARATION

In this section, we review how a nonuniform magnetization or strain can lead to a space-dependent node separation. This discussion of the physical origin of axial magnetic fields in solid-state systems will set the stage for our general study presented in the next sections, by introducing the main concepts and notation.

A. Time-reversal symmetric Weyl semimetal

The existence of Weyl nodes relies on breaking either time-reversal (\mathcal{T}) or inversion symmetry (\mathcal{I}) . If both are present, all energy bands are manifestly twofold degenerate, and, in particular, point nodes must have a degenerate

partner with opposite chirality. If \mathcal{T} is broken, the two nodes of a pair are allowed to be separated in momentum space, with the separation determined by **b**. If \mathcal{I} is broken but \mathcal{T} is preserved, each pair must have a time-reversed partner, such that the minimum number of Weyl nodes in \mathcal{I} -broken Weyl semimetals is four. In an \mathcal{I} -broken Weyl semimetal, the Hamiltonian of the low-energy Weyl fermions for each pair is given by

$$\mathcal{H}_{\text{Weyl}} = \sum_{i=x,y,z} \sum_{s=\pm} s v_i (k_i + s b^i) \sigma^i, \qquad (1)$$

where v_i are the Fermi velocities, σ^i are Pauli matrices spanning the orbital space of the Weyl fermions, and $s = \pm$ denotes the chirality of the nodes. It is clear from Eq. (1) that the vector $\mathbf{b} = (b^x, b^y, b^z)$ quantifies the separation of the two nodes of a pair in momentum space.

In general, in a crystalline material, a uniform strain of strength g alters the overlaps between wave functions in different orbitals [18,21]. In Ref. [21], for instance, which considered HgTe with spin-orbit-induced band inversion in the presence of strain, found that at low energies the spectrum consists of four pairs of Weyl nodes. Because of the presence of strain, the Hamiltonian of each pair then takes the form

$$\mathcal{H}_{\text{Weyl}} = \sum_{i,s=\pm} v_i(g) [sk_i \sigma^i + b^i(g)\sigma^i].$$
(2)

From Eq. (2), we observe that the effect of strain is twofold. First, it modifies the Fermi velocities through $v_i(g) \sim \sqrt{g}$. Second, it controls the Weyl node separation through the vector **b** with magnitude $|\mathbf{b}(g)| \sim \sqrt{g}$. Both of these effects are analogous to those predicted in graphene [3,5,70].

Reference [21] analyzed the effect of strain as a homogeneous perturbation characterized by q. However, in practice, strain can have a nonuniform profile $q(\mathbf{r})$. This can occur, for instance, when there is a lattice mismatch between a substrate and the semimetal sample or, alternatively, via chemically induced strain. In the former scenario, for thin enough samples, it can be expected that the strain profile relaxes smoothly as a function of the distance from the substrate (quantitative estimates will be provided in Sec. V). Let us then choose the direction of the strain inhomogeneity to be y such that $q(\mathbf{r}) = q(y)$, and assume that q(y) changes slowly with y, on a length scale that is much smaller than any other microscopic length scale. In such a perturbative regime, $g(y) \simeq$ $g_0 + \delta g(y)$ with $\delta g(y)/g_0 \ll 1$, it is reasonable to expand Eq. (2) to lowest order in $\delta g(y)/g_0$. Using $v[g(y)] \simeq$ $v(g_0)(1+\delta g(y)/2g_0)$ and $b^i[g(y)] \simeq b_i(g_0)(1+\delta g(y)/2g_0)$, one obtains

$$\mathcal{H}_{\text{Weyl}} \simeq \sum_{i,s=\pm} v(g_0) \sigma^i \left[si\partial_i + b^i(g_0) \left(1 + \frac{\delta g(y)}{g_0} \right) \right].$$
(3)

We first note that the effect of the space-dependent Fermi velocity due to strain [70] is of order $[\delta g(y)/g_0]\partial_i$, so it can be neglected in the low-energy linear approximation. An inhomogeneous velocity has been predicted recently to result in an anomalous Hall signal transverse to the direction in which the strain is applied [71,72]. However, these works neglected the effect of emergent inhomogeneous Weyl node separation that, according to the above analysis, is of lower order in $\delta g(x)$ and therefore dominant.

Motivated by the above effective model, we consider the following real-space Hamiltonian:

$$\mathcal{H}_{\text{Weyl}} \simeq \sum_{i,s=\pm} v[si\partial_i - b_i^{(0)} + \delta b_i(\mathbf{r})]\sigma^i, \qquad (4)$$

where $\mathbf{b}(\mathbf{r}) \equiv \mathbf{b}^{(0)} - \delta \mathbf{b}(\mathbf{r})$ encodes the strain profile. From such a model with a single pair of nodes, the Dirac semimetal and the inversion-breaking Weyl semimetal can be constructed by adding time-reversal partners of Eq. (4) in the appropriate location in momentum space in order to restore time-reversal symmetry.

We note that in the literature, the vector \mathbf{b} has been referred to, depending on the context, as chiral, axial, or pseudovector potentials. In this work, we will refer to \mathbf{b} as the axial-vector potential, as is customary in high-energy physics where this field was first discussed.

B. Time-reversal-breaking Weyl semimetal

In \mathcal{T} -breaking Weyl semimetals, the axial-vector potential **b** that separates the Weyl nodes in momentum space can have its physical origin in a finite magnetization. A hallmark example of a Weyl semimetal of this kind is the proposal by Burkov and Balents [73], which is based on topological-insulator-trivial-magnetic-insulator heterostructures. Another example is the Weyl semimetal state induced by magnetic order at a quadratic band crossing point on pyrochlore iridates [74]. In addition, a timereversal-breaking Weyl semimetal has been argued to be consistent with ARPES measurements on YbMnBi₂ [67]. Other promising recent proposals [68] suggest, relying on ab initio calculations, that magnetic Heusler compounds can host Weyl quasiparticles. Remarkably, alloys of the latter kind could realize the minimal model of a \mathcal{T} -breaking Weyl semimetal with only two nodes and a node separation that spans a significant fraction of the Brillouin zone.

Similar to other magnetic materials, the magnetization that induces the Weyl semimetal state by separating two nodes of opposite chirality in momentum space may itself be nonuniform and become a function of position [75]. As a result, the Weyl node separation becomes a function of position, generating an axial-magnetic-field strength. To lowest order in the spatial variation of the Weyl node separation (i.e., magnetization), the effective model (4) will describe this state. We note as well that interfaces between magnetic domains, either abrupt or continuous, could be modeled by the spatial profiles of **b** that we consider in this work. In addition, it is plausible to expect that coating one of the surfaces with a ferromagnetic layer or enough surface magnetic dopants will also result in a spatially varying magnetization. The successful realization of magnetically doped topological insulators [76] suggests that similar techniques could be applied in Weyl samples as well.

In addition to an inhomogeneous magnetization, a nonuniform strain can generate an axial magnetic field in \mathcal{T} -breaking Weyl semimetals. Once \mathcal{T} is broken and a finite **b** exists, strain can allow us to effectively change **b** from its bare value, thus changing the separation of Weyl nodes in momentum space. Note, however, that strain on its own cannot break time-reversal symmetry and thus cannot induce a finite **b** from **b** = 0.

C. Effective realization in cold atomic systems

While the above sections discuss realizations of inhomogeneous Dirac and Weyl semimetals in the solid state, our discussion is also relevant for cold atomic systems [69]. Two appealing aspects of the latter are that different topological properties can be extracted easily from experimental data [77–79] and models that respect or break time reversal could be engineered.

There are two further advantages of considering cold atomic systems that are relevant to probe the effects we discuss. First, engineering a space-dependent Weyl node separation is experimentally feasible. A natural way to achieve this is to smoothen the confining trapping potential that contains the lattice [80]. Although plausible, it does not offer the degree of control that will expose all of the salient features we are after, but only a subset of them. A more controlled and realistic approach is to create domain walls between topological states [81] as already described in Ref. [82] for Chern insulators. We elaborate on more particulars of this proposal in the concluding section.

The second advantage of cold atomic systems is the controlled experimental access to transport properties that was recently discussed in the context of Chern insulators [83,84]. The key observable is the center-of-mass velocity $\mathbf{v}_{c.m.}$ since it is simple to measure experimentally and is connected to the current density \mathbf{j} and particle density *n* through [83]

$$\mathbf{v}_{\rm c.m.} = \frac{\mathbf{j}}{n}.$$
 (5)

Although, in general, n also depends on external fields, in our particular case, it acts as an unimportant constant factor as we describe below. Thus, via Eq. (5), it is possible to directly probe the longitudinal conductivity.

III. ENHANCED TOPOLOGICAL LONGITUDINAL CONDUCTANCE

Many peculiar transport properties have been attributed to topological semimetals [10,11], while not many of them have proven easy to measure. One particularly striking feature that has been confirmed in several experiments [39–66,85] is the appearance of a nonsaturating negative magnetoresistance [86,87]. Conductance is one of the most accessible experimental observables to characterize the properties of a material. Therefore, we aim here to explore the effect of an inhomogeneous Weyl node separation on this quantity. We show that a space-dependent axial-vector potential **b** enhances the longitudinal conductance of a Weyl semimetal through its corresponding axial magnetic field $\mathbf{B}_5 = \nabla \times \mathbf{b}$. We begin by introducing a coupling to an external electromagnetic field and discussing the role of gauge invariance. We follow this by extending a quantum field theory argument previously used in Ref. [40] to include pseudo-gauge fields. We then support our claims with a more rigorous semiclassical calculation by solving the semiclassical kinetic equation.

A. Coupling to an external electromagnetic field and the role of gauge invariance

In order to calculate the conductivity, we couple the Hamiltonian (4) to an external electromagnetic field $A_{\mu} = (A_0, \mathbf{A})$ using the minimal substitution

$$\mathcal{H}_{\text{Weyl}} = \sum_{s=\pm} v\{s[i\partial_i - eA_i(x)] - b_i(x)\}\sigma_i + A_0. \quad (6)$$

The first important property of this Hamiltonian is that left-handed (s = 1) and right-handed (s = -1) Weyl fermions are decoupled, experiencing a chirality-dependent gauge field, $a_{\mu}^{s} = A_{\mu} + sb_{\mu}$, where $b_{\mu} = [0, \mathbf{b}(\mathbf{r})]$. For what follows, it is useful to define the left- and right-handed field strengths $f_{s}^{\mu\nu} = \partial_{\mu}a_{\nu}^{s} - \partial_{\nu}a_{\mu}^{s}$.

The continuity equation for the current for each species $j_s^{\mu} = (n_s, \mathbf{j}_s)$ is determined by [13,88]

$$\partial_{\mu}j_{s}^{\mu} = s \frac{e^{3}}{4\pi^{2}\hbar^{2}} \mathbf{E}_{s} \cdot \mathbf{B}_{s} - \frac{e}{\tau}(n-n_{s}), \qquad (7)$$

which we now discuss in some depth. The first term on the right-hand side of Eq. (7) is composed of the left and right effective electromagnetic fields $\mathbf{E}_s = \mathbf{E} + s\mathbf{E}_5$ and $\mathbf{B}_s = \mathbf{B} + s\mathbf{B}_5$, where (\mathbf{E}, \mathbf{B}) , $(\mathbf{E}_5, \mathbf{B}_5)$, and $(\mathbf{E}_s, \mathbf{B}_s)$ are built out of the gauge field strength $F^{\mu\nu} = \partial_{\mu} A_{\nu} - \partial_{\nu} A_{\mu}$, the axial-gauge-field strength $f_b^{\mu\nu} = \partial_{\mu} b_{\nu} - \partial_{\nu} b_{\mu}$, and $f_s^{\mu\nu}$, respectively. The second term on the right-hand side is a scattering term that acts to equilibrate the number density n_s of the left and right chiralities [89,90] with a typical scattering time τ [91].

The conservation laws for the current $j^{\mu} = \sum_{s} j_{s}^{\mu}$ and axial current $j_{5}^{\mu} = \sum_{s} s j_{s}^{\mu}$ can be obtained from the addition or subtraction of the two equations composing Eq. (7). A first inspection reveals a seemingly striking feature; neither of the two currents is conserved. The nonconservation of j_{5}^{μ} is not forbidden, and it is referred to as the chiral anomaly [11,88,92]. If both $\mathbf{E}_{5} = 0$ and $\mathbf{B}_{5} = 0$, then one recovers

the celebrated chiral anomaly that schematically reads $\partial_{\mu}j_{5}^{\mu} \sim \mathbf{E} \cdot \mathbf{B}$ [28,88]. In this case, the vector current satisfies $\partial_{\mu}j^{\mu} = 0$, and thus charge is conserved. However, it seems that charge is not conserved when either $\mathbf{E}_{5} \neq 0$ or $\mathbf{B}_{5} \neq 0$. Indeed, from Eq. (7), we find that the vector current satisfies $\partial_{\mu}j^{\mu} \sim \mathbf{E}_{5} \cdot \mathbf{B} + \mathbf{E} \cdot \mathbf{B}_{5}$ [13,17].

In fact, there is no contradiction with current conservation, which can be understood in two different ways. First, the axial-vector potential **b** is an observable and thus must be single valued, with a zero vacuum expectation value [11]. Thus, the total flux of the resulting **B**₅ must be zero over a surface enclosing the entire sample; regions with **B**₅ and $-\mathbf{B}_5$ compensate each other by generating an equal number of left- and right-handed fermions separated in real space. Then, upon applying an electric field, charge flows from one region to the other, respecting global charge conservation.

Second, at the quantum-field-theory level, the nonconservation of charge is fixed by defining a consistent current that preserves gauge invariance. The procedure has been described both in the context of high-energy physics [93] and Weyl semimetals [33]: The current calculated above (the covariant current) is complemented by Chern-Simons currents (the Bardeen-Zumino polynomials) of the form $-[1/(4\pi^2)]\epsilon_{\mu\nu\rho\sigma}b^{\nu}F^{\mu\nu}$, which exactly cancel the anomaly and define the consistent current. This procedure effectively imposes a boundary condition for the spectral flow at the energy of the cutoff that bounds the field theory. In a lattice system, such a cutoff is a natural quantity, and the spectral flow is bounded by construction. Note also that this procedure applies wherever there is a finite axial magnetic field \mathbf{B}_5 and therefore includes the boundaries where **b** jumps from zero to a finite value or vice versa.

B. Quantum-field-theory approach

Our aim is to use the chiral anomaly Eq. (7) to find the longitudinal conductance in the presence of a finite axial magnetic field **B**₅. To promote the derivation in Ref. [40], we first use that, for three-dimensional Weyl fermions, $n_s = \mu_s^3/(6\pi^2\hbar^3v^3)$, which we employ to rewrite the steady-state form of Eq. (7) as

$$\mu_s = \left[\mu^3 - \frac{3}{2}s\hbar v^3 e^2 \tau \mathbf{E}_s \cdot \mathbf{B}_s\right]^{1/3}.$$
 (8)

We now recall that the component of the current parallel to **B** and \mathbf{B}_5 is given by

$$\mathbf{j} = \frac{e^2}{2\pi^2 \hbar^2} [\mu_5 \mathbf{B} + \mu \mathbf{B}_5]. \tag{9}$$

The first term is the chiral magnetic effect [29,30] and has been thoroughly studied in the context of Weyl semimetals [14–16,31–33]. This term must be zero in equilibrium [36,37,94–96] since it is proportional to a chemical

potential imbalance $\mu_5 = \frac{1}{2} \sum_s s\mu_s$. The second term on the left-hand side is the key to our results: It represents the analog of the chiral anomaly in the presence of an axial magnetic field **B**₅ and can be finite in equilibrium. In the context of dense relativistic matter, it was shown that it is possible to generate an axial current **j**₅ from a vector field **B** with a conductivity proportional to the chemical potential $\mu = \frac{1}{2} \sum_s \mu_s$ [97]. It follows that there is a contribution to the vector current **j** generated by an axial field with the *same* coefficient [98]. We refer to this term as the chiral pseudomagnetic effect.

In order to obtain the longitudinal conductivity, we combine Eqs. (8) and (9),

$$\mathbf{j} = \frac{e^2}{h^2} \sum_{s} s \mu_s \mathbf{B}_s \tag{10}$$

$$=\frac{e^2}{h^2}\sum_{s}s\mu\left[1-s\frac{3}{2\mu^3}\hbar v^3e^2\tau\mathbf{E}_s\cdot\mathbf{B}_s\right]^{1/3}\mathbf{B}_s.$$
 (11)

Assuming that

$$\frac{3}{2\mu^3}\hbar v^3 e^2 \tau \mathbf{E}_s \cdot \mathbf{B}_s \ll 1, \tag{12}$$

and for the case where $\mathbf{B} = B\hat{z}$ and $\mathbf{B}_5 = B_5\hat{z}$, $\mathbf{B}_s = (B + sB_5)\hat{z}$, and $E_5 = 0$, we obtain

$$j_z = \frac{v^3 e^4 \tau}{4\pi^2 \mu^2 \hbar} E(B^2 + B_5^2) + \frac{e^2}{2\pi^2 \hbar^2} \mu B_5, \qquad (13)$$

to linear order in **E**. The first term is a transport (or free) current. For the case when $B_5 = 0$, it reproduces the chiralanomaly-enhanced magnetoconductivity [86]. The central result here is that for $\mathbf{B}_5 \neq 0$, there is a contribution to the bulk longitudinal conductivity coming entirely from the spatial dependence of $\mathbf{b}(\mathbf{r})$: Inhomogeneous strain or magnetization enhances conductance in a Weyl or a Dirac semimetal.

We now provide a physical interpretation of the second term as a magnetization (or bound) current. Recall that $\mathbf{b}(\mathbf{r})$ is analogous to a finite magnetization $\mathbf{M}(\mathbf{r})$. Thus, local bound currents given by $\mathbf{j}_b \propto \nabla \times \mathbf{M}(\mathbf{r}) \propto \mathbf{B}_5$ are allowed, consistent with the form of the second term in Eq. (13). Note that in the related context of the quantum Hall effect, a finite bound current proportional to μ and the curl of the magnetization is expected on general grounds [99]. In that case, the bound current has its origin on the edge states, while here they can be associated with the bulk.

We remark that the second term in Eq. (13) highlights an important difference between semimetals that respect time-reversal symmetry and those that do not. In the former, the sum over time-reversed pairs of nodes will cancel out the μ **B**₅ term. In Sec. III C, we provide further arguments that support the interpretation of this term as a bound current,

while in Sec. IV, we corroborate these findings by numerically studying the bound current profile within specific lattice models.

C. Boltzmann equation approach

In this section, we outline a different and more rigorous derivation of Eq. (13) that relies on solving the Boltzmann equation. Within this approach, the semiclassical equations of motion are extended to include an anomalous velocity term that arises because of the existence of a nonzero Berry curvature [100]. Typically, for a Weyl semimetal, the equations of motion are written for a single flavor of chiral fermions accounting for the physics in momentum space centered around a particular Weyl node [86]. Both chiralities feel the same electric and magnetic fields. The key difference in the present analysis is that the effective external fields are now chirality dependent because of the axial vector potential; both chiralities are still decoupled but feel different effective fields.

More precisely, the effect of the effective magnetic fields $\mathbf{B}_s = (\mathbf{B} + s\mathbf{B}_5)$ on the left (s = +1) and right (s = -1) chiralities can be incorporated by promoting the semiclassical equations of motion [100] to

$$\dot{\mathbf{r}}_s = \partial_{\mathbf{p}} \mathcal{E}^s_{\mathbf{p}} + \dot{\mathbf{p}} \times \Omega^s_{\mathbf{p}}, \qquad (14)$$

$$\dot{\mathbf{p}}_s = e\mathbf{E} + \frac{e}{c}\dot{\mathbf{r}} \times \mathbf{B}_s. \tag{15}$$

Here, $\mathcal{E}_{\mathbf{p}}^{s} = \boldsymbol{\varepsilon}_{\mathbf{p}}^{s} - \mathbf{m}_{\mathbf{p}}^{s} \cdot \mathbf{B}_{s}$ and $\Omega_{\mathbf{p}}^{s}$ are, respectively, the dispersion relation, which includes a correction due to the magnetic orbital moment $\mathbf{m}_{\mathbf{p}}^{s}$, and the corresponding Berry curvature for each chirality *s*. For each chirality, the unperturbed dispersion relation of the upper band is $\boldsymbol{\varepsilon}_{\mathbf{p}}^{s} = sv|\mathbf{p}|$. The Berry curvature and magnetic orbital moment, in this case, take the simple form $\Omega_{\mathbf{p}}^{s} = s[1/(2|\mathbf{p}|^{2})]\hat{\mathbf{p}}$ and $\mathbf{m}_{\mathbf{p}}^{s} = -ev|\mathbf{p}|\Omega_{\mathbf{p}}^{s}$. We emphasize that, consistent with the discussion in Sec. II, we neglect effects of the higher-order corrections due to the inhomogeneous Fermi velocity, which can enter through a space-dependent Berry curvature [71,72,101].

The distribution function for each chirality $f_{\mathbf{p}}^{s}$ satisfies a semiclassical kinetic equation

$$\frac{\partial f_{\mathbf{p}}^{s}}{\partial t} + \dot{\mathbf{r}} \cdot \frac{\partial f_{\mathbf{p}}^{s}}{\partial \mathbf{r}} + \dot{\mathbf{p}} \cdot \frac{\partial f_{\mathbf{p}}^{s}}{\partial \mathbf{p}} = I_{\text{coll}}\{f\}, \quad (16)$$

where $I_{coll}{f}$ is the collision integral. Using Eqs. (14) and (15), we can write

$$\dot{\mathbf{r}}_{s} = D_{\mathbf{p},s}^{-1} \bigg(\mathbf{v}_{\mathbf{p}}^{s} + e\mathbf{E} \times \Omega_{\mathbf{p}}^{s} + \frac{e}{c} (\Omega_{\mathbf{p}}^{s} \cdot \mathbf{v}_{\mathbf{p}}^{s}) \mathbf{B}^{s} \bigg), \quad (17)$$

where $D_{\mathbf{p},s} = [1 + (e/c)\mathbf{B}^s \cdot \Omega_{\mathbf{p}}^s]$ and $\mathbf{v}_{\mathbf{p}}^s = \partial \mathcal{E}_{\mathbf{p}}^s / \partial \mathbf{p}$ is the perturbed velocity. We find that the contribution of a single chirality to the current density is given by

$$\mathbf{j}^{s} = e \int \frac{d^{3}p}{(2\pi)^{3}} D_{\mathbf{p},s} \dot{\mathbf{r}}_{s} f_{\mathbf{p}}^{s}(\mathcal{E}_{\mathbf{p}}^{s}),$$

$$= e \int \frac{d^{3}p}{(2\pi)^{3}} \left(\mathbf{v}_{\mathbf{p}}^{s} + e\mathbf{E} \times \Omega_{\mathbf{p}}^{s} + \frac{e}{c} (\Omega_{\mathbf{p}}^{s} \cdot \mathbf{v}_{\mathbf{p}}^{s}) \mathbf{B}^{s} \right) f_{\mathbf{p}}^{s}(\mathcal{E}_{\mathbf{p}}^{s}).$$
(18)

Within the relaxation time approximation, $I_{\text{coll}}{f} = -(f_{\mathbf{p}}^s - f_{\mathbf{p},s}^0)/\tau$, where τ is a scattering time [102] and $f_{\mathbf{p},s}^0(\mathcal{E}_{\mathbf{k},s})$ is the equilibrium distribution function to be evaluated at the modified dispersion relation $\mathcal{E}_{\mathbf{k},s}$. We are interested in a stationary and homogeneous solution to the kinetic equation, which takes the form

$$\dot{\mathbf{p}} \cdot \frac{\partial f_{\mathbf{p}}^s}{\partial \mathbf{p}} = -\frac{(f_{\mathbf{p}}^s - f_{\mathbf{p},s}^0)}{\tau}.$$
(19)

Expanding the left-hand side of the previous equation to lowest order in the fields and rearranging terms, we obtain

$$f_{\mathbf{p}}^{s} = f_{\mathbf{p},s}^{0} - \tau D_{\mathbf{p},s}^{-1} \left(e\mathbf{E} + \frac{e}{c} \mathbf{v}_{\mathbf{p}}^{s} \times \mathbf{B}^{s} + \frac{e^{2}}{c} (\mathbf{E} \cdot \mathbf{B}^{s}) \Omega_{\mathbf{p}}^{s} \right) \\ \times \frac{\partial f_{\mathbf{p},s}^{0}}{\partial \mathbf{p}}.$$
 (20)

Inserting the first term in Eq. (20) into Eq. (18) and ignoring the contribution transverse to the electric field leads to

$$\mathbf{j}_0^s = e \int \frac{d^3 p}{(2\pi)^3} \frac{e}{c} \left(\Omega_{\mathbf{p}}^s \cdot \mathbf{v}_{\mathbf{p}}^s \right) \mathbf{B}^s f_{\mathbf{p},s}^0(\mathcal{E}_{\mathbf{p}}^s).$$
(21)

To leading order in the magnetic field, we can make the replacement $\mathcal{E}_{\mathbf{p}}^s \to \mathcal{E}_{\mathbf{p}}^s$ in both the equilibrium distribution function and the definition of the velocity. The integral results in

$$\mathbf{j}_0^s = s \frac{e^2}{h} \mu \mathbf{B}^s, \tag{22}$$

with μ the chemical potential, consistent with earlier findings [19,32]. Summing Eq. (22) over chiralities results in the second term in Eq. (13).

Remarkably, we have also performed a rate of entropy production calculation in the spirit of Ref. [86] and find this term to be absent. The reason for this absence is that the rate of entropy production is determined exclusively by the *free* current density rather than the total current density. This result implies, as anticipated above, that Eq. (22) should be physically interpreted as a bound current proportional to the curl of the magnetization in the system and as such does not contribute to transport.

Next, we insert the second term in Eq. (20) into Eq. (18) to calculate the correction to the current corresponding to

the deviation $\delta f_{\mathbf{p}}^{s} = f_{\mathbf{p}}^{s} - f_{\mathbf{p},s}^{0}$ up to first order in **E** and second order in **B**_s, which reads

$$\delta \mathbf{j}^{s} = -e\tau \int \frac{d^{3}p}{(2\pi)^{3}} D_{\mathbf{p},s}^{-1} \left[e\mathbf{v}_{\mathbf{p}}^{s} \left(\mathbf{E} \cdot \frac{\partial f_{\mathbf{p}}^{s}}{\partial \mathbf{p}} \right) + \frac{e^{2}}{c} \mathbf{v}_{\mathbf{p}}^{s} (\mathbf{E} \cdot \mathbf{B}^{s}) \left(\Omega_{\mathbf{p}}^{s} \cdot \frac{\partial f_{\mathbf{p}}^{s}}{\partial \mathbf{p}} \right) + \frac{e^{2}}{c} \mathbf{B}^{s} (\Omega_{\mathbf{p}}^{s} \cdot \mathbf{v}_{\mathbf{p}}^{s}) \left(\mathbf{E} \cdot \frac{\partial f_{\mathbf{p}}^{s}}{\partial \mathbf{p}} \right) + \frac{e^{3}}{c^{2}} \mathbf{B}^{s} (\Omega_{\mathbf{p}}^{s} \cdot \mathbf{v}_{\mathbf{p}}^{s}) (\mathbf{E} \cdot \mathbf{B}^{s}) \left(\Omega_{\mathbf{p}}^{s} \cdot \frac{\partial f_{\mathbf{p}}^{s}}{\partial \mathbf{p}} \right) \right].$$
(23)

Using that for the lower band $\Omega_{\mathbf{p}}^{s} = -s[1/(2|\mathbf{p}|^{2})]\hat{\mathbf{p}}$ and $\mathbf{m}_{\mathbf{p}}^{s} = -ev|\mathbf{p}|\Omega_{\mathbf{p}}^{s}$, and after tedious but straightforward manipulations, we obtain

$$\delta j_z^s = -\tau \left(\frac{1}{3} e^2 v^2 \nu(\mu) + \frac{v^3 e^4}{30 \pi^2 \hbar \mu^2} B_s^2 \right) E, \qquad (24)$$

where $\nu(\varepsilon) = \varepsilon^2 / 2\pi^2 \hbar^3 v^3$ is the unperturbed density of states of a Weyl node at energy ε . The first term encodes the conductivity due to a finite Fermi surface [103,104]. The second term is novel to this work and, upon summing chiralities, leads to a longitudinal contribution to the conductivity that reads

$$\sigma_{zz} = \tau \frac{v^3 e^4}{15\pi^2 \hbar \mu^2} B_5^2, \tag{25}$$

for B = 0, and represents the main result of this section. Up to the numerical coefficient, this contribution has exactly the same form as the first term of Eq. (13), confirming that an inhomogeneous strain or magnetization contributes to increase the conductivity. The numerical differences between Eqs. (13) and (25) can be traced back to the expansion of the modified density of states $D_{\mathbf{p},s}$ appearing in the denominator of Eq. (23) and the inclusion of the magnetic moment in the semiclassical calculation. We note as well that, as with conventional magnetoresistance, we expect that the term Eq. (24) is also supplemented by a Fermi surface contribution due to the Lorentz force when the Fermi surface is anisotropic [105].

We conclude this section by computing the center-ofmass velocity in this approach, an observable quantity in cold atomic experiments. From Eq. (5), to obtain the centerof-mass velocity and relate it to the conductivity, we must compute the electron density n that to leading order reads

$$n = \sum_{s} \int \frac{d\mathbf{p}}{(2\pi)^3} D_{\mathbf{p},s} f_{\mathbf{p}}^s \sim \frac{\mu^3}{3\pi^2 v^3 \hbar^3}.$$
 (26)

The relevant center-of-mass velocity component is obtained by inserting Eqs. (24) and (26) into Eq. (5) to obtain

$$\delta v_{\rm c.m.}^z = \frac{\delta j_z}{n} = -\tau \left(\frac{e^2 v^2}{\mu} + \frac{2}{5} \frac{\hbar^2 v^6 e^4}{\mu^5} B_5^2 \right) E, \quad (27)$$

which is valid in the absence of magnetic field ($\mathbf{B} = 0$). Therefore, through Eq. (27), a cold atomic experiment can probe the anomalous longitudinal conductivity $\sigma \sim \mathbf{B}_5^2$ that we predict by monitoring the center-of-mass motion of an atomic cloud.

IV. SPECTRAL PROPERTIES OF INHOMOGENEOUS SEMIMETALS

Based on the field-theoretic and semiclassical axialgauge-field coupling in Weyl semimetals, we now study the effects of magnetization or strain inhomogeneities within a microscopic lattice model realization. This allows us to numerically corroborate the arguments presented in the previous section. We consider two canonical lattice models whose low-energy long-wavelength limit is given by Eq. (4). For simplicity, we choose lattice realizations of Weyl semimetals with the minimum number of low-energy Weyl fermions, i.e., one Weyl fermion of each chirality, which implies that time-reversal symmetry is broken in both models. Our results, however, generalize to timereversal symmetric (but inversion symmetry-breaking) models, and occasionally, we explicitly comment on such generalizations.

The first minimal lattice model we use to describe Weyl fermions coupled to axial gauge fields is motivated by the solid-state realizations of Weyl and Dirac semimetals. It can be obtained starting from a lattice-regularized version of a topological insulator $\mathbf{k} \cdot \mathbf{p}$ Hamiltonian [106], such as Bi₂Se₃ or related materials. A simple generalization of this model has been used to describe the Dirac semimetals Cd₂As₃ and Na₃Bi [34,81]. The model has four bands, originating from an orbital degree of freedom *A*, *B* and spin \uparrow, \downarrow , and the corresponding electron operators are defined as $c_{\mathbf{r}} = (c_{\mathbf{r}A\uparrow}, c_{\mathbf{r}A\downarrow}, c_{\mathbf{r}B\uparrow}, c_{\mathbf{r}B\downarrow})^T$. The Hamiltonian H_{4b} of this four-band model is the sum of two terms and is given by

$$H_{4b} = \sum_{\mathbf{k},j} D^j(\mathbf{k}) c_{\mathbf{k}}^{\dagger} \Gamma^j c_{\mathbf{k}} + \sum_{\mathbf{r},j} b^j(\mathbf{r}) c_{\mathbf{r}}^{\dagger} \Gamma_b \Gamma^j c_{\mathbf{r}}.$$
 (28)

The first term is the topological-insulator Hamiltonian with Γ matrices $\Gamma^{j} = (\sigma^{z}s^{y}, \sigma^{z}s^{x}, \sigma^{y}s^{0}, \sigma^{x}s^{0}), \ \Gamma_{b} = \sigma^{y}s^{z}$, and with the components of the **D** vector given by

$$D^{j}(\mathbf{k}) = -\left(\sin k_{x}a, \sin k_{y}a, \sin k_{z}a, \sum_{i}\cos k_{i}a - M\right).$$
(29)

Here, we have set the kinetic energy scale t = 1. The matrices $\sigma^{x,y,z}$ and $s^{x,y,z}$ are Pauli matrices acting on orbital and spin degrees of freedom, respectively (σ^0 and s^0 are

identity matrices). The Dirac mass M, which describes a hybridization of the orbitals into bonding and antibonding states, controls whether the material is on a trivial- or topological-insulator phase (which may be strong or weak). Here, we set $M \equiv 3 + m$, such that m = 0 corresponds to a Dirac semimetal state with a 3D Dirac point at $\mathbf{k} = 0$, and m > 1 (m < 0) corresponds to a trivial (topological) insulator.

Whereas the first term in Eq. (28) respects both timereversal symmetry (\mathcal{T}) and inversion symmetry (\mathcal{I}), a nonzero $\mathbf{b} = (b^x, b^y, b^z)$ breaks \mathcal{T} . To see how it is responsible for generating the Weyl semimetal phase, one may expand Eq. (28) to linear order in $\delta \mathbf{k}$ around $\mathbf{k} = 0$ and obtain

$$H(\mathbf{k}) = \sum_{j} (\delta k_j + b^j \Gamma_b) \Gamma^j + m \Gamma^4.$$
(30)

This result shows that **b** is responsible for the separation of the two nodes in momentum space and thus couples as an axial gauge field. Indeed, Eq. (30) should be compared to the Weyl Hamiltonian of Eq. (4), where the node separation was identified with the axial gauge field. In Eq. (30), the matrix Γ_b takes the role of giving the axial gauge field **b** a different sign at the two nodes. It should be noted, however, that in Eq. (30) the node separation depends on both **b** and m. In particular, the node separation is proportional to $\sqrt{|\mathbf{b}|^2 - m^2}$ and vanishes (i.e., the system is insulating) for $m > |\mathbf{b}|$ [14,81].

In the following, we will study Hamiltonian (28) numerically in the presence of a spatially nonuniform $\mathbf{b}(\mathbf{r})$, which gives rise to axial gauge fields $\mathbf{B}_5 \simeq \nabla \times \mathbf{b}(\mathbf{r}) \neq 0$. It is important to note that $\nabla \times \mathbf{b}(\mathbf{r})$ can only have the meaning of an axial magnetic field coupled to Weyl fermions in regions where $m < |\mathbf{b}(\mathbf{r})|$. We always take $m \ge 0$ since we are not interested in the regime where **D** describes a topological insulator. A full phase diagram of the model defined by Eq. (28) was described in Ref. [81].

The second lattice model we use to verify our results consists of effectively spinless electrons, has only two bands, and can be regarded as one of the two time-reversal partners that compose the model in Ref. [69]. It is therefore motivated by two practical considerations: (i) It is plausible to implement it in the cold atomic context, and (ii) it falls into the class of models tailored for the methods presented in Ref. [82] to generate domain walls in topological systems. In this case, the electron operators are given by $c_{\mathbf{r}} = (c_{\mathbf{r}A}, c_{\mathbf{r}B})$, where *A*, *B* represent some generalized orbital degree of freedom, and the Hamiltonian takes the form

$$H_{2b} = \sum_{\mathbf{k},j} d^j(\mathbf{k}) c_{\mathbf{k}}^{\dagger} \sigma^j c_{\mathbf{k}}.$$
 (31)

Here, $\sigma^z = \pm 1$ again represents the orbital degree of freedom, and the components of **d** are given by

$$d^{j}(\mathbf{k}) = -\left(\sin k_{x}a, \sin k_{y}a, \sum_{i}\cos k_{i}a - M\right). \quad (32)$$

This model breaks time-reversal symmetry; it has a pair of linearly dispersing Weyl cones at $\mathbf{k} = \{0, 0, \pm \cos^{-1}(M/t-2)\}$ for 1 < |M/t| < 3, two pairs of Weyl cones for |M/t| < 1, and is a gapped insulator when |M/t| > 3. The parameter M/t controls the distance between Weyl nodes; interpolating between M/J = 2 and M/t > 3 simulates the boundary between a Weyl semimetal with two nodes and an insulator. Therefore, we promote $M \to M(y)$, which sets the Weyl node separation as $\mathbf{b}(y)$ by

$$\mathbf{b}(y) = (0, 0, 2 \arccos(-2 + M(y)/t)).$$
(33)

We note that, although this model separates the Weyl nodes in k_z , any other separation direction can be chosen by redefining **d** appropriately.

From our calculations, we find that both models qualitatively exhibit the same behavior, so we will focus mainly on results obtained for the four-band model of Eq. (28).

A. Lattice pseudo-Landau-level structure of B₅

We begin by specifying the spatial profiles of $\mathbf{b}(\mathbf{r})$ that generate the axial magnetic configurations we study. In what follows, we always take $\mathbf{b}(\mathbf{r}) = b^x(y)\hat{x}$, such that it only depends on y and corresponds to an axial magnetic field along z. Since nonzero b^x implies a separation of Weyl nodes along the x axis in momentum space, $b^x(y)$ describes Weyl nodes whose separation Δk_x depends on the y coordinate. Note that this effectively corresponds to a Landau gauge, and consequently, (k_x, k_z) remain good quantum numbers. Furthermore, it follows that the axial magnetic field $\nabla \times \mathbf{b}$ is orthogonal to the direction of node separation. The extension of the system in the y direction is L_y (similarly for x and z), which we measure in units of the lattice constant a, and we assume $b^x(1) = b^x(L_y)$.

We first consider the series of profiles $b^{x}(y)$ shown schematically in the bottom row of Fig. 1. The form of $b^{x}(y)$ is such that it increases linearly from b_{1}^{x} to b_{2}^{x} between $y = \ell_1$ and $y = \ell'_1$, then stays flat at b_2^x , and subsequently decreases linearly again from b_2^x back to b_1^x between $y = \ell_2$ and $y = \ell'_2$. As a result, between ℓ_1 and ℓ'_1 , we have a constant and negative $\nabla \times \mathbf{b} \simeq \mathbf{B}_5$, whereas between ℓ_2 and ℓ'_2 , we have a positive constant $\nabla \times \mathbf{b} \simeq \mathbf{B}_5$, with a magnitude that depends on the ratio $\Delta b^{x} / \Delta \ell_{1,2}$, with $\Delta b^{x} = b_{2}^{x} - b_{1}^{x}$ and $\Delta \ell_{1,2} = \ell_{1,2}' - \ell_{1,2}$. For fixed Δb^x , the strength of the effective axial magnetic field increases for decreasing $\Delta \ell_{1,2}$. By taking $\Delta \ell_{1,2} \rightarrow 0$, as shown in the rightmost configuration of Fig. 1, we create two sharp interfaces between regions of constant $\mathbf{b} = b_1^x \hat{x}$ and $\mathbf{b} = b_2^x \hat{x}$. We choose *m* [see Eqs. (28) and (30)] such that the latter corresponds to a boundary between a Weyl

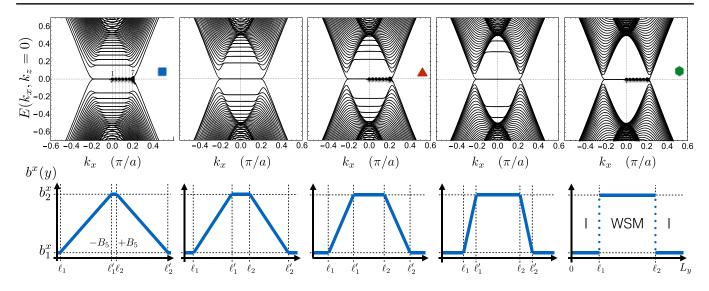


FIG. 1. The top row shows the energy spectra obtained by solving Eq. (28) in the presence of a spatially varying $b^x = b^x(y)$, as a function of k_x and for $k_z = 0$. The schematic profile of $b^x(y)$ corresponding to each spectrum is presented below the energy panel. From left to right, the region of linear increase (decrease) of b^x , set by $\ell'_1 - \ell_1$ ($\ell'_2 - \ell_2$), gradually shrinks and becomes an abrupt stepwise increase (decrease) at ℓ_1 (ℓ_2). The latter case describes an interface between an insulator (I) and a Weyl semimetal (WSM) (all spectra are calculated for m = 0.5), and the energy spectrum exhibits the zero-energy Fermi arc surface states connecting the bulk nodes. The panel on the left shows the pseudo-Landau-level structure of the low-energy states, arising as a result of a bulk axial magnetic field $B_5^z = \pm B_5$. As $\ell_1' - \ell_1 = \ell_2' - \ell_2$ is decreased from left to right, the strength of the axial magnetic field is increased and the $n \ge 1$ pseudo-Landau levels (energy scales as $\sim \sqrt{nB_5}$) are pushed out of the spectrum. The n = 0, however, remains at E = 0 and morphs into the Fermi arc associated with the boundary surface shared by I and WSM. In these calculations ($b_1^x = 0.0, b_2^x = 1.0$), $\ell_1' - \ell_1 = \ell_2' - \ell_2 = (80, 60, 40, 20), (\ell_1, \ell_2) = (45, 135)$ (rightmost panel), with system size $L_x \times L_y \times L_z = 120 \times 180 \times 120$ (all lengths measured in units of lattice constant a). The solid black dots in the upper panels indicate the (zero-energy) states for which the wave-function support is shown in Fig. 3.

semimetal and an insulator, i.e., $m > b_1^x$ [81]. Thus, the configurations of Fig. 1 interpolate between axial magnetic fields in the 3D bulk of the system and a sharp 2D boundary between a Weyl semimetal and an insulator. The axial magnetic field is gradually confined to a 2D surface while increasing its strength. The energy spectra corresponding to these axial-magnetic-field profiles are shown in the top panels of Fig. 1, where we have set $b_1^x = 0.0$, $b_2^x = 1.0$, and m = 0.5.

Before turning to an analysis of the energy spectra obtained from the lattice model, it is useful to recall the Landau-level spectrum associated with axial magnetic fields in the continuum, cf. Eq. (4). We refer to these Landau levels as pseudo-Landau levels. For Weyl fermions coupled to a uniform axial magnetic field \mathbf{B}_5 along the *z* direction, B_5^z , the $n \ge 1$ pseudo-Landau levels have energies $E_{n\pm}(k_z) = \pm \hbar v \sqrt{k_z^2 + 2|B_5^z|n}$ and thus disperse in k_z . In addition, there is an n = 0 or zeroth pseudo-Landau level with energy $E_0(k_z) = \mathrm{sgn}(B_5^z)\hbar v k_z$. Importantly, all energy levels are doubly degenerate: one for each of the two Weyl nodes. In particular, the two Weyl nodes, which have opposite chiralities, have *the same* $E_0(k_z)$ dispersion of the zeroth Landau level. In contrast, in the presence of a (vector) magnetic field B^z , the chirality of the n = 0

branch, i.e., the upward or downward slope as a function of k_z , depends on the chirality of the Weyl nodes [28].

The pseudo-Landau-level structure of the continuum is reflected in the low-energy part of the lattice energy spectra shown in Fig. 1. In particular, in the leftmost panel, we observe both a flat branch of zero energy states and flat branches of states at higher energies. These can be identified with pseudo-Landau levels since these do not disperse in k_x . Moreover, labeling the flat branches of states at nonzero energies by an index n, we find that the energies indeed scale as $\sim \sqrt{n}$. In the rightmost panel, where the axial magnetic field is confined to the sharp boundary between the Weyl semimetal and the insulator, we still find a branch of zero-energy states. The flat branches at higher energies are absent. The zero-energy states are simply the Fermi arc surface states, which must exist at such a surface boundary because of the topology of the Weyl nodes, and they connect the projections of the bulk Weyl nodes. Figure 1 thus suggests that as $\ell'_{1,2} - \ell'_{1,2}$ is gradually taken to zero, confining the axial magnetic field to a narrower region along y while increasing its magnitude, the n = 0pseudo-Landau level becomes the Fermi arc. The energy of the higher $n \ge 0$ pseudo-Landau levels scales as $\sim \sqrt{B_5}$ and is therefore "pushed" out of the spectrum as B_5 increases, as can be observed from left to right in Fig. 1.

The key implication of Fig. 1 is that the Fermi arc surface states of a Weyl semimetal can be thought of as an n = 0 pseudo-Landau level corresponding to an axial magnetic field spatially confined to the surface boundary, i.e., $\sim B_5^z \delta(y - \ell_{1,2})$. The chirality of the Fermi arc states, i.e., their dispersion in k_z (discussed in more detail below), corresponds to the chirality of the n = 0 pseudo-Landau-level modes and depends on $\text{sgn}(B_5^z)$.

B. Fermi arcs as n = 0 pseudo-Landau level

To study this correspondence in more detail, we now consider a different set of $b^x(y)$ profiles, which are shown in the middle row of Fig. 2. As is schematically demonstrated, in this set of profiles, $b^x(y)$ increases and decreases linearly from $-b^x$ to b^x , and we take $b^x = 0.5$. Furthermore, *m* is set to zero, such that the profiles interpolate between bulk axial magnetic fields on the left and an interface between two Weyl semimetals with inverted Weyl node separation b^x on the right. The corresponding energy spectra are shown in the top panels of Fig. 2.

Focusing first on the right panel, i.e., the sharp interface between two Weyl semimetals with inverted b^x , we find that the branch of zero-energy states, which connect the projections of the bulk nodes, is fourfold degenerate. These are the Fermi arcs localized at the boundaries between the Weyl semimetals, two for each boundary. This is consistent with the number of arcs mandated by topology. Similar to Fig. 1, the spectrum in the left panel exhibits the pseudo-Landau-level structure at low energies. The flat branch of zero-energy states corresponding to the n = 0 pseudo-Landau level is fourfold degenerate, in agreement with the degeneracy of the Fermi arcs $(2 \times 2 = 4)$. The pseudo-Landau-level degeneracy can be understood by recalling that each Weyl node contributes one Landau level, which is doubled because of the two spatially separated regions of $\pm B_5$. (The counting in Fig. 1 is more subtle, which we explain below.) From Fig. 2, we again observe that the Fermi arcs are adiabatically connected to n = 0 pseudo-Landau levels as the profile of $b^{x}(y)$ is varied. The n > 1pseudo-Landau levels are pushed to higher energies (and are eventually absent from the spectrum) because of the increasing axial-magnetic-field strength.

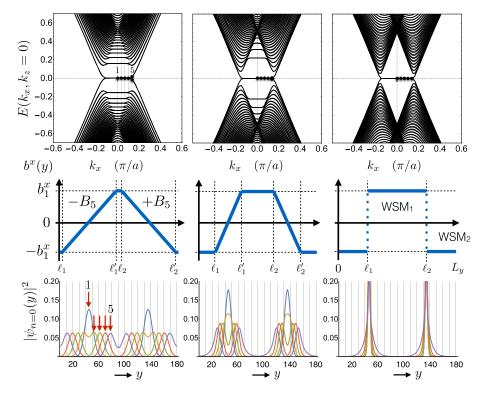


FIG. 2. Energy spectra (top row) obtained from Eq. (28) in the presence of a spatially varying $b^x = b^x(y)$, shown in the corresponding central panels. The spectra were calculated using m = 0.0 and $b_1^x = 0.5$, such that the rightmost panels describe an interface between two Weyl semimetals (WSM₁ and WSM₂) with an inverted **b** axial vector. As a result, at each of the two boundaries, ℓ_1 and ℓ_2 , there will be two Fermi arcs, and the degeneracy of the zero-energy states connecting the bulk nodes is doubled compared to Fig. 1. The energy-level structure of the leftmost panel can be understood from the perspective of bulk axial magnetic fields. In each of the two regions, $B_5^z = +B_5$ and $B_5^z = -B_5$, two n = 0 pseudo-Landau levels are present, one for each Weyl node, giving a total of four n = 0 pseudo-Landau levels. The bottom panels show the support of the wave function of some E = 0 states labeled by solid black circles as a function of y. The wave-function support demonstrates that the Fermi arc states (on the right) are the Landau orbitals of the n = 0 pseudo-Landau level (on the left). We used $\ell_1' - \ell_1 = \ell_2' - \ell_2 = (80, 40)$ for the first two columns, (ℓ_1, ℓ_2) = (45, 135) for the rightmost column, and a system size of $L_x \times L_y \times L_z = 120 \times 180 \times 120$.

More insight can be gained by studying the wave functions of the Fermi arc states and comparing them to Landau orbital wave functions. Recall that wave functions of the lowest n = 0 Landau-level orbitals are given by

$$\Psi_{k_{z}k_{z}}^{n=0}(\mathbf{r}) \propto e^{ik_{x}x} e^{ik_{z}z} e^{-(y-k_{x}l_{b}^{2})^{2}/2l_{b}^{2}},$$
(34)

where l_b is the magnetic length. The Landau orbitals are centered around $y = k_x l_b^2$, which locks the y coordinate to momentum k_x . The distance between the Landau orbitals along y, as well as the spread of the wave function, is determined by the magnetic length l_b . Since $l_b \sim 1/\sqrt{B}$, the magnetic length decreases as the field strength B increases.

In Fig. 2, we plot the support of the wave functions of the zero-energy states corresponding to different values of k_x . The states for which wave functions are shown are indicated by solid black dots in the upper panels. From the bottom left panel of Fig. 2, we see that at each momentum there are indeed four states, with a wavefunction support similar to that of Landau orbitals. At $k_x = 0$, these are localized at the center of each of the two regions of positive and negative axial magnetic fields, $B_5^z = \pm B_5$. As k_x increases, the support of the wave function is shifted along y, and in opposite directions for Landau orbitals associated with Weyl nodes of opposite chiralities. This result follows from the position-momentum locking expressed in Eq. (34).

As the profile of $b^x(y)$ is changed from left to right, the axial-magnetic-field strength is increased, and thus the effective magnetic length is decreased. This is clearly reflected in the wave-function support of the zero-energy states: The spread becomes narrower, and they move closer together. Eventually, when the axial magnetic field is confined to the interfacial boundary between the Weyl semimetals at ℓ_1 and ℓ_2 , the Landau orbitals are localized at the boundaries and should be viewed as the wave functions of the Fermi arc states.

An analogous picture arises when we plot the wavefunction support of zero-energy states of the spectra in Fig. 1, which is presented in Fig. 3. Figure 3(a) shows the wave functions of the n = 0 pseudo-Landau-level states of the leftmost panel in Fig. 1, demonstrating the positionmomentum locking $\langle y \rangle \propto k_x$. Note that there are only half the number of Landau orbitals as compared to Fig. 2, reflecting the different degeneracy of zero-energy states: twofold versus fourfold. As is shown in Fig. 3(a), the Landau orbitals in regions where $b^x(y) < m$ are absent. The region where $b^x(y) > m$ divides into a part where B_5^z is positive and a part where B_5^z is negative. In each region, both Weyl nodes contribute n = 0 pseudo-Landau-level orbitals, but of opposite momentum. Figure 3(a) shows the contribution of only one Weyl node (chirality).

Similar to Fig. 2, we observe that in Fig. 3, as the magnetic length decreases towards zero from (a) to (c), the pseudo-Landau orbitals become the wave functions of

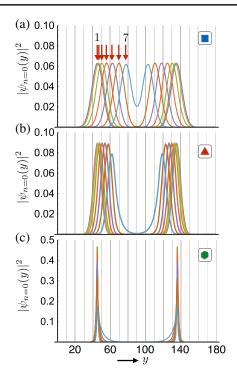


FIG. 3. Plot of the support of the wave function, as a function of y, of selected zero-energy states corresponding to the spectra shown in Fig. 1. Panels (a), (b), and (c) correspond to leftmost, middle, and rightmost panels of Fig. 1, respectively (graphically indicated by the square, triangle, and hexagon). Here, in each panel, the seven different curves correspond to different values of k_x , which are indicated by solid black dots in Fig. 1 and red arrows in (a). Two pseudo-Landau orbitals correspond to each value of k_x , as is most clearly seen in (a). This matches the degeneracy of n = 0 pseudo-Landau levels and is consistent with the number of Fermi arcs: one Fermi arc per boundary. Note that as compared to the bottom row of Fig. 2, the Landau orbitals centered in the region where $b^x(y) < m$ are absent, which explains the difference in degeneracy (twofold vs fourfold as discussed in the main text).

the Fermi arc states localized at the surface boundary between the insulator and Weyl semimetal. From the perspective of Fermi arc surface states, the degeneracy of states can be inferred from topology: The number of Fermi arcs at each boundary is equal to the Chern-number change across the boundary. The Chern number can be defined when Weyl nodes of opposite (or different) Berry monopole charge are separated in momentum space. This suggests that, per the adiabatic connection between Fermi arcs and n = 0 pseudo-Landau levels, the degeneracy of pseudo-Landau levels in solid-state systems has a topological origin.

The pseudo-Landau levels of the continuum are flat as a function of k_x but disperse in k_z . Therefore, it is useful to compare the spectra of the lattice model as a function of k_z . The energy spectra, obtained for the two $b^x(y)$ profiles of the leftmost and rightmost panels of Fig. 1, are shown in Fig. 4. The low-energy part of the spectrum in the left panel

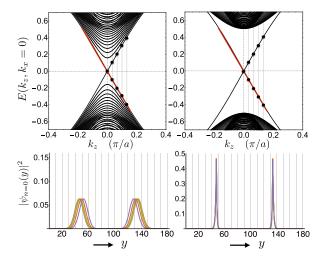


FIG. 4. Energy spectra, as a function of k_z (with $k_x = 0$), and wave-function support obtained from Eq. (28) in the presence of a bulk axial magnetic field B_5^z (left) and for an interface boundary between an insulator and a Weyl semimetal (right). The spatial profile of $b^x(y)$ corresponding to the left and right panels here are given in the leftmost and rightmost bottom panels of Fig. 1, respectively. On the left, the low-energy branch of the spectrum has the structure of pseudo-Landau levels. The linearly dispersing chiral (black) and antichiral (red) modes are the n = 0 pseudo-Landau levels. The wave-function support of n = 0 modes is shown for states with k_z values, indicated by solid black dots. On the right, the linearly dispersing modes correspond to the Fermi arc surface states and are sharply localized at the boundary interfaces ℓ_1 and ℓ_2 (see Fig. 1). The same systems sizes as in Figs. 1 and 2 are used.

of Fig. 4 clearly exhibits the pseudo-Landau-level structure, with $n \ge 1$ pseudo-Landau-level energies dispersing in k_z as $\sim \sqrt{k_z^2 + nB_5}$. In addition, there is an n = 0 chiral $(E \propto +k_z)$ and antichiral $(E \propto -k_z)$ mode, which are localized in different spatial regions corresponding to opposite B_5^z . This is confirmed by the wave-function support of states with different k_z in the bottom panel. Note that the momentum k_z is not locked to the y coordinate. The right panel of Fig. 4 shows the dispersion of the Fermi arc states in k_z . The wave functions (bottom panel) are localized at the surface boundaries. Once more, one may think of the Fermi arcs as the n = 0 chiral (and antichiral) modes of the n = 0 pseudo-Landau level. The difference in the spread of the wave functions between the bottom-left and bottom-right panels originates from the difference in effective magnetic length.

At this stage, two remarks regarding the generality of our results are in order. The first concerns the profiles of $b^x(y)$ shown in Figs. 1 and 2. In all these cases, the increase or decrease of $b^x(y)$ is chosen to be linear, giving rise to a constant B_5^z . In solid-state materials, axial-vector potentials are due to strain or magnetization inhomogeneities, and one may expect the axial-magnetic-field strength to have a more general dependence on position, i.e., $\mathbf{B}_5 = \mathbf{B}_5(\mathbf{r})$. To verify

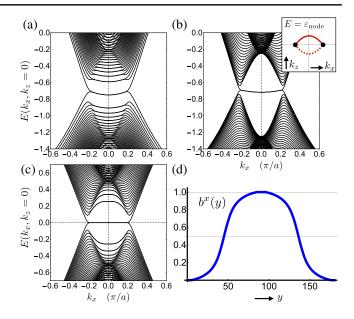


FIG. 5. Effect of the Fermi arc curvature and deviations from constant axial magnetic fields. (a,b) Plot of the energy spectra with $-\delta t \sum_i \cos k_i a \sigma^0 s^0$ added to Eq. (28) and δt set to $\delta t = 0.25$. The $b^x(y)$ profiles corresponding to (a) and (b) are given in the leftmost and rightmost bottom panels of Fig. 1, respectively, with m = 0.5. The effect of finite δt is to give k_x dispersion to the pseudo-Landau levels (a) and the Fermi arc states (b). The inset of (b) shows the Fermi arc states in the $k_x - k_z$ plane at the node energy ε_{node} . (c) Energy spectrum obtained from Eq. (28) with $b^x(y)$ as shown in (d), with m = 0.5. Profile (d) leads to a nonuniform $\mathbf{B}_5 = \mathbf{B}_5(y) \approx \nabla \times \mathbf{b}$, which is reflected in (c), where the $n \ge 1$ pseudo-Landau levels disperse in k_x because of position-momentum locking.

the application of our results to the more general case, we calculate the energy spectrum of Hamiltonian (28) with $b^{x}(y)$ as shown in Fig. 5(d). The corresponding spectrum is shown in Fig. 5(c), which demonstrates two important features. First, the n = 0 pseudo-Landau level remains nondispersive in k_x and at zero energy (for $k_z = 0$). Second, the higher pseudo-Landau levels appear to have acquired a dispersion and shifted in energy. This result is in agreement with the energies of the pseudo-Landau levels, $E_0 \propto \operatorname{sgn}(B_5^z)k_z$ and $E_{n\pm} \propto \sqrt{k_z^2 + 2|B_5^z|n}$, respectively. Replacing B_5^z by $B_5^z(y)$ and noting that $y \propto \text{sgn}(B_5^z)k_x$ because of Eq. (34), it follows that the $n \ge 1$ pseudo-Landau levels should disperse in k_x , whereas the energy of the n = 0 pseudo-Landau level should not change. This shows that our results hold for a general axialmagnetic-field configuration.

The second remark concerns the shape of the Fermi arcs. In Figs. 1, 2, and 4, Fermi arc states at zero energy connect the bulk Weyl nodes in a straight line located on the k_x axis because of the nonfundamental symmetry of the model (28). In general, the shape of the Fermi arcs is not restricted, and the condition $E(k_x, k_z) = \varepsilon_{node}$, where ε_{node} is the energy of the bulk nodes (assuming they are both at the same energy), may define a curve connecting the bulk nodes with arbitrary shape. To study the more general case, we add $-\delta t \sum_i \cos k_i a \sigma^0 s^0$ to Hamiltonian (28) and calculate the energy spectra for two different $b^x(y)$ profiles. As shown in Figs. 5(a) and 5(b), both the pseudo-Landau levels (a) and the Fermi arcs (b) disperse in k_x , implying that the Fermi arcs trace out a curve of the form shown in the inset of (b). We conclude from Figs. 5(a) and 5(b) that

C. Calculation of the bound current density

the correspondence between Fermi arcs and n = 0 pseudo-

Landau levels holds for more general Fermi arc shapes.

As pointed out in Sec. III, the axial-vector potential **b** in time-reversal-broken Weyl semimetals physically corresponds to a magnetization vector and is therefore expected to give rise to bound currents \mathbf{j}_b . Generally, the total current in an electronic system with a finite magnetization can be expressed as a sum of bound \mathbf{j}_b and free current \mathbf{j}_f densities [107]. The former has the property that it must average to zero over the system's volume, but it is allowed to be nonzero locally. It therefore can be expressed as the curl of a local vector $\mathbf{M}(\mathbf{r})$, the magnetization, such that $\mathbf{j}_b = \mathbf{\nabla} \times \mathbf{M}$. In the case where **M** is constant in the bulk, the bound currents only exist as surface currents $\mathbf{j}_b^{\text{surf}}$ that are perpendicular to the surface normal $\hat{\mathbf{n}}$ such that $\mathbf{j}_{b}^{\text{surf}} = \hat{\mathbf{n}} \times \mathbf{M}$.

We now provide numerical evidence that justifies the interpretation of the second term in Eq. (13), i.e., $\mu \mathbf{B}_5$, as a bound current density, by calculating the current density from our microscopic lattice models. To this end, we consider five different linear profiles of the axial-vector potential of the form $\mathbf{b} = b_a^x(y)\hat{x}$, parametrized by a = 1, 2, 3, 4, 5. These trace out a finite Weyl semimetal with a Weyl node separation that increases linearly in the *y* direction [see Fig. 6(b)]. The current density $j_z(y)$ is computed through the expression

$$j_{z}(y) = \langle \hat{J}_{z}(y) \rangle$$

= $\frac{1}{L_{y}L_{z}} \sum_{n,k_{x},k_{z}} \langle u_{k_{x}k_{z}}^{n} | J_{z}(y) | u_{k_{x}k_{z}}^{n} \rangle f(\varepsilon_{k_{x}k_{z}}^{n}), \quad (35)$

where $J_z(y) = \partial H_{k_x k_z}(y) / \partial k_z$ is the current operator, $|u_{k_x k_z}^n\rangle$ are the single-particle eigenstates, and $f(\varepsilon_{k_x k_z}^n)$ is the Fermi-Dirac distribution function evaluated at the *n*th eigenvalue $\varepsilon_{k_x k_z}^n$. Its real-space distribution is shown in Fig. 6(a) for different values of b_a^x , with an offset for clarity. From Fig. 6(a), we observe two main features. First, for the flat profile b_1^x , the current is localized at the boundaries with equal weight but opposite sign. This is entirely consistent with what is expected of a bound current; for a constant magnetization, the bound currents are localized at the interface and are normal to it $(\mathbf{j}_{b}^{\text{surf}} \exp | \mathbf{a} + \mathbf{j})$, we observe

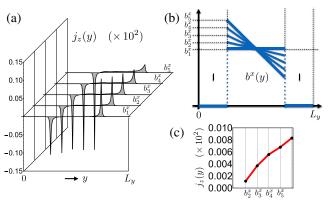


FIG. 6. Bound current density in an inhomogeneous Weyl semimetal. Panel (a) shows the five different current densities $j_z(y)$ corresponding to the five different profiles of the axial-vector potential shown in (b), labeled b_a^x with a = 1, 2, 3, 4, 5. Larger values of a support larger axial magnetic fields $\mathbf{B}_5 = \mathbf{\nabla} \times \mathbf{b}$, which create bulk pseudo-Landau levels [see Fig. 7(c)] that compensate for the missing current density at the boundary. Plot (c) shows the bound current density at a point $y = L_y/2$ belonging to the Weyl semimetal bulk for different values of b_a^x .

that the weight associated with one boundary is transferred to the bulk, but the total current density remains zero overall.

In order to understand both of these features in more detail, we express each of these linear profiles mathematically as

$$b_a^x(y) = \left(\frac{b_f^x - b_i^x}{(\ell_f - \ell_i)}(y - \ell_i) + b_i^x\right) \\ \times [\Theta(y - \ell_i) - \Theta(y - \ell_f))], \tag{36}$$

which is schematically shown in Fig. 7(b) by the light blue curve. The corresponding $\mathbf{B}_5 = \nabla \times \mathbf{b}$ profile is set to a constant in the bulk (which is zero for the flat profile outlined by b_1^x and nonzero for $b_{a\neq 1}^x$) and to two Dirac delta functions of opposite sign corresponding to each boundary.

Setting $b_i^x = b_f^x = b_1^x$ results in the spectral profile already discussed in Fig. 1 (lower row, rightmost panel): a Weyl semimetal with two Fermi arc surface states. As chiral surface states, the two Fermi arcs carry two compensating current densities, consistent with what is observed for the b_1^x case shown in Fig. 6(a). As the difference $b_f^x - b_i^x$ increases, the profile traces higher values of *a* in $b_{a\neq1}^x$, and an increasing and constant $\mathbf{B}_5 = \nabla \times \mathbf{b}$ emerges in the bulk. The relevant energy spectrum for this situation is shown in Fig 7(c); pseudo-Landau levels emerge, and the zeroth chiral pseudo-Landau level seems indistinguishable from a Fermi arc.

In fact, it is possible to distinguish the surface and bulk contribution due to position-momentum locking. As was discussed following Eq. (34), the average guiding center position $\langle y \rangle$ is tied to the momentum k_x such that $\langle y \rangle \sim k_x$.

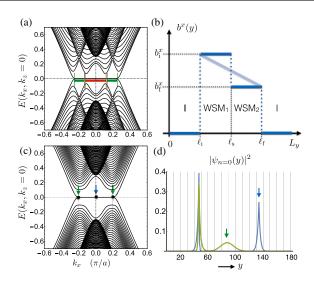


FIG. 7. Plot of the energy spectra (left panels) as a function of k_x obtained for the $b^x(y)$ profiles shown in (b). The first case (a) corresponds to a heterostructure of insulator (I) and a Weyl semimetal in which the separation of Weyl nodes changes abruptly from b_i^x (WSM₁) to b_f^x (WSM₂) at ℓ_s ; see Eq. (36). In (a), the zero energy states in the green region are associated with the "mini-arcs" localized at l_s , whereas the states marked by red are associated with the interface between the Weyl semimetals and the insulator. In the second case (c), the separation of Weyl nodes is changed linearly between ℓ_i and ℓ_f [see Eq. (37)], as shown by the dark blue line. The wave-function support of states indicated by solid black dots in (c) is presented in (d).

Thus, by tracking the dependence of the wave function of the zeroth Landau Level $\Psi_{n=0}(y)$ as a function of k_x , we can extract real-space information.

In Fig. 7(d), we show $|\Psi_{n=0}(y)|^2$ for two representative values of k_x marked in Fig. 7(c). At $k_x = 0$, the wave function only has a finite weight at the edges, outlining a purely surface state. However, when $b_{\rm f}^x < |k_x| < b_{\rm i}^x$, the wave function acquires weight in the bulk associated with the zeroth pseudo-Landau level emerging in this momentum-space region. Importantly, both regions are continuously connected as a function of k_x , and thus, from positionmomentum locking, they are also connected in real space. The bulk \mathbf{B}_5 creates bulk pseudo-Landau levels that connect to the surface arcs as a consequence of positionmomentum locking. This is a central result of this work, and it allows us to understand the different instances of Fig. 6(a): As **B**₅ increases, a zeroth bulk pseudo-Landau level forms that connects the unbalanced surface states. The current density is sensitive to this fact; the current density lost by one Fermi arc is compensated and transferred to a bulk zeroth pseudo-Landau level. In other words, the bound current at one surface is compensated by the sum of bound currents in the bulk and the remaining surface.

Taking one step back, our results can be summarized by the identification of the second term in Eq. (13) with a bound current due to the curl of a magnetization, $\mathbf{j}_b = \mu \mathbf{B}_5 = \mathbf{\nabla} \times \mathbf{M}$. We have further corroborated this result by explicitly checking that for small fields the magnitude of the bound current in the bulk of Fig. 6(a) increases linearly with \mathbf{B}_5 [cf. Fig. 6(c)], and it is only finite if $\mu \neq 0$.

All these arguments combined establish the three main points collected in the abstract: (i) the Fermi arcs can be reinterpreted as n = 0 pseudo-Landau levels resulting from a **B**₅ confined to the surface, (ii) a bulk **B**₅ creates bulk pseudo-Landau levels that connect to the surface arcs as a consequence of position-momentum locking, and (iii) there are bound currents proportional to **B**₅ and the chemical potential that average to zero over the sample, as occurs in magnetic materials.

It is important to stress that the effects we discuss are sensitive to the pseudomagnetic-field direction, in particular, whether it is parallel or perpendicular to the Weyl node separation. In the parallel case, the bulk and surface behave as expected: The two bulk zeroth pseudo-Landau levels have the same chirality, which is opposite to the chirality of the Fermi arcs at the boundaries. The surface spectral weight is evenly distributed between each arc and compensates the bulk [27]. In contrast, in the perpendicular case considered in this work, the surface states are allowed to have a chirality that matches that of the bulk zeroth pseudo-Landau levels, a fact that is enforced by positionmomentum locking. The Fermi arcs at different surfaces do not have the same spectral weight, rendering the rich spectral structure reported here.

D. Interface between two Weyl semimetals as a finite bulk B₅

Given the above discussion, it is interesting to consider the case of two Weyl semimetals of different Weyl node separation that share a boundary. At this point, we can intuitively predict the outcome: If the system abruptly changes the Weyl node separation, this is equivalent to the appearance of a B_5 that is confined to the interface. This case occurs when the change from b_i^x to b_f^x is localized at a single point in space, as shown in Fig. 7(b) (dark curve). It represents two Weyl semimetals with two constant axial-vector potentials b_i^x and $b_{\rm f}^{\rm x}$ brought into contact. From the previous section, we expect that currents bound to the surfaces of the two semimetals have a different magnitude, and their difference is proportional to $\nabla \times (\mathbf{b}_{i} - \mathbf{b}_{f})$. Alternatively, the extensions of their Fermi arcs in momentum space are different. Therefore, we expect that there is a finite amount of current bound to the interface that compensates for that difference.

From a spectral point of view, it is interesting to observe how the Fermi arcs behave in this case. To this end, we consider the profile of $b^x(y)$ shown by the dark curve in Fig. 7(b). Mathematically, it is described by

$$b^{x}(y) = b^{x}_{i}\Theta(y - \ell_{i}) + (b^{x}_{f} - b^{x}_{i})\Theta(y - \ell_{s})$$
$$-b^{x}_{f}\Theta(y - \ell_{f}).$$
(37)

The corresponding spectrum is shown in Fig. 7(a) and is characterized by the appearance of "mini-arcs", fragments of longer arcs that are surface states belonging to the uniform system with the larger Weyl node separation b_i^x . This situation is quite generic for an interface between two topological phases with the same topological invariant the surface states hybridize and gap out along the region of contact. For Weyl semimetals, we can consider this from the perspective of assigning a Chern number to twodimensional slices of momentum space that lie between two Weyl nodes [23]. The corresponding chiral edge states hybridize and gap out in the region of overlap in momentum space, which is equal to the length of the shorter axialvector potential b_f^x .

It is now straightforward to understand how the spectrum evolves when the interface between two Weyl semimetals is smeared across the entire sample, which corresponds to Fig. 7(b) (light blue curve). The spectrum in this case, presented in Fig. 7(c), shows that the smeared interface transforms the interfacial arc states at l_s into zeroth pseudo-Landau Levels. Moreover, this conclusion based on the spectral information of Fig. 7(c) is corroborated by the wave-function spread in real space shown in Fig. 7(d). As discussed in the last section, the latter portrays how, as k_x is changed, the real-space probability density shifts from edge to bulk. We stress that this is another complementary instance of the discussion around Figs. 1 and 2. The surface Fermi arcs blend with the zeroth Landau levels within a single $b^{x}(y)$ profile, Fig. 7(b) (light blue curve), rather than a sequence of them as in Figs. 1 and 2.

Moreover, the comparison between the light blue and dark blue profiles in Fig. 7(b) inspires the following physically appealing picture of how the current density profile is distributed for the linear profile studied in Fig. 6(b). The region between ℓ_i and ℓ_f in Fig. 7(b) (light blue curve) can be approximated by a collection of infinitesimal discontinuities in the spirit of the trapezoidal method for curve integration. At each discontinuity, there are Fermi arcs that meet and annihilate with part of the arcs of the neighboring layer. These bulk arcs carry current density and thus act as a collection of bulk sheets that "leak" current density into the bulk from the surface.

V. DISCUSSION AND CONCLUSION

In this work, we studied inhomogeneous Weyl and Dirac semimetals with a space-dependent Weyl node separation. We have discussed how such a scenario can arise either from inhomogeneous strain or magnetization in existing solid-state systems as well as cold atomic setups. Underlying our results is an axial magnetic field B_5 that couples to the electronic degrees of freedom with two main experimental consequences for both Dirac and Weyl semimetals, which we have addressed in depth.

The first experimental consequence is a drastic change of the spectral properties of topological semimetals. Using two different lattice models, we have established two novel spectral features attributed to the emergence of B_5 : (i) Fermi arcs are secretly zeroth pseudo-Landau levels due to a finite and large \mathbf{B}_5 at the boundary, and (ii) bulk pseudo-Landau levels form because of B_5 and compensate the difference in density of states of inequivalent Fermi arcs at opposite boundaries via position-momentum locking. Point (i) is an inevitable consequence of the boundary supporting a finite \mathbf{B}_5 since the Weyl node separation must vanish in vacuum. This allows us to identify the existence of Fermi arcs with the emergence of a boundary zeroth pseudo-Landau level due to \mathbf{B}_5 , even in the absence of bulk inhomogeneities. Such a correspondence provides a novel perspective on surface physics of Weyl and Dirac semimetals. Point (ii) is particularly relevant for spectral probes like ARPES or STM that are sensitive to a modification of the electronic spectrum [7–9]. Remarkably, for the particular case of time-reversal-broken realizations of inhomogeneous Weyl semimetals, the axial magnetic field results in an inhomogeneous distribution of bound currents throughout the sample, which exist in equilibrium and average to zero over the entire volume. The appearance and field dependence of these bound currents in the lattice realization that we study is also consistent with our semiclassical treatment of the response.

We envision two plausible routes to probe the effects resulting from the bound currents discussed. Magnetic sensors such as scanning superconducting quantum interference devices (SQUIDs) are a natural way to probe local distributions of bound currents. Such probes were proven very successful at detecting magnetization modulations and inhomogeneous current distributions at interfaces such as LAO/STO heterostructures, and in two- and threedimensional topological insulators (cf., Refs. [108–110]). It is important to emphasize that nonlocal currents due to an inhomogeneous magnetization and the bound currents due to \mathbf{B}_5 are physically analogous phenomena, and thus it is very likely for the latter to be detectable. The difference between bound currents in an ordinary magnetized material and bound currents within a Weyl semimetal is that those in the latter emerge from a unique coupling between the Weyl fermions and the background magnetization as described throughout this paper.

Another promising alternative are torque experiments, which have in fact already been conducted with Weyl semimetals [111]. The magnetic torque $\tau = \mathbf{M} \times \mathbf{B}$ is a direct measure of the magnetic anisotropy of a crystal and thus can reveal noncompensation of surface magnetic domains.

Within our semiclassical treatment, we have predicted a second experimentally relevant consequence, which is related to transport: Inhomogeneities enhance the longitudinal conductivity in Dirac or Weyl semimetals as $\sigma \sim \mathbf{B}_5^2$. We find that this transport property is routed in a chiral pseudomagnetic effect similar to the enhancement

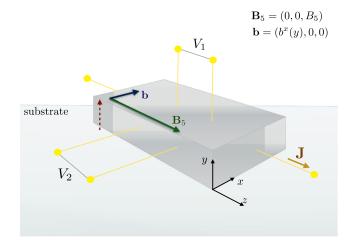


FIG. 8. A schematic point-contact transport setup to measure the current **J** (orange arrow) to determine the strain-dependent conductivity $\sigma \sim \mathbf{B}_5^2$ predicted in this work. Strain is induced by the substrate with a small lattice mismatch that relaxes along the y direction (red dashed arrow). If the Weyl node separation is $\mathbf{b} \| \hat{x}$ (blue arrow), then $\mathbf{B}_5 \| \hat{z}$ (green arrow). Contacts $V_1 (V_2)$ probe a surface with (without) Fermi arcs, resulting in an anisotropic contribution to the conductivity.

of the magnetoconductance due to the existence of the chiral magnetic effect. We stress that this prediction is applicable to all kinds of topological semimetals. Since this enhancement is insensitive to the sign of \mathbf{B}_5 , it will be generated by each pair of Weyl nodes, a number that is of order 10 in realistic materials.

A typical transport setup with point-contact probes, schematically shown in Fig. 8, suffices. It represents a thin film, or nanowire, of a topological semimetal that is strained by a substrate with a small lattice mismatch. This setup can be particularly relevant for strained HgTe/CdTe heterostructures discussed in Ref. [21]. As illustrated in Fig. 8, point-contact measurements have the additional freedom to choose the voltage probes to be on a surface with or without Fermi arcs ($V_{1,2}$, respectively), depending on wether the momentum-space projection of the nodes on that particular surface coincides or not. Although for the TaAs class of materials all accessible surfaces host Fermi arcs, Dirac materials like Cd₃As₂ do present surfaces free of arcs. Such anisotropy highlights the difference between topological metals and more conventional strained semiconductors. While in the latter conductance can also be enhanced via strain by modifying the band structure, changing the position of the voltage probes is not expected to result in anisotropic measurements if current jetting is negligible [45].

The above considerations suggests a way to probe the surface bound currents. As we have discussed extensively in Sec. IV, surface currents on opposite surfaces in the presence of bulk strain will not compensate each other completely since their difference will be carried by the bulk pseudo-Landau levels. It could be expected that two distinct

surfaces that are perpendicular to \mathbf{B}_5 will carry different currents. Put differently, the Fermi arcs of two opposite surfaces will not be of equal length in the presence of bulk strain. Thus, measuring an anisotropy in the surface currents could reveal the size of the bulk strain gradients. However, typical surface-current signals are small, and thus this detection mechanism is less feasible in practice. Nonetheless, we note that the special nature of the Fermi arcs can conspire to make the surface contribution sizable, as in other related situations [112].

In order to assess the significance of the effects we predict, we now estimate the magnitude of a bulk axial field \mathbf{B}_{5} . For a thin sample, it is expected that strain relaxes linearly with height over several unit cells if the lattice constant mismatch is small. For a typical sample, such as a Cd_3As_2 thin film, we can consider a height of the order of $\Delta L \sim 10$ nm, and a conservative value of strain is translated into a change in lattice constant of about 1% but can be as large as about 10%. If we assume that the Weyl node separation typically spans a tenth of the Brillouin zone $|\mathbf{b}| \sim 1/10 \text{ Å}^{-1}$, then the effective magnetic field is $|\mathbf{B}_5| \sim$ $[(\hbar\Delta b)/e\Delta L] \sim (\hbar/e)10^{15} \text{ m}^{-2} \sim 4 \text{ T}.$ Remarkably, this conservative estimate results in sizable magnetic fields, certainly above the detectable threshold of magnetic loops and SQUIDS. These fields will also induce detectable changes in conductance that can be probed by growing samples with different ΔL that result in different intrinsic **B**₅.

Cold atomic systems offer a controlled alternative. For these systems, we have proposed that the conductivity enhancement can be measured by monitoring the center-ofmass velocity $\mathbf{v}_{c.m.}$. This quantity is the quotient between the current density **j** and the particle density *n*, which will depend on the external fields in general. For small fields, the density reduces to a trivial constant, and the center-ofmass position is determined solely by the current density times a constant factor. This experiment could be performed in a cold atomic realization of the two-band model used in this work. Moreover, this model, along with the family of two-band models that host Weyl fermions [69], is ideal to apply the method proposed in Ref. [82] to study inhomogeneities. It relies on implementing a spacedependent offset between neighboring sites. Applying such an offset profile, which is feasible experimentally, results in a space-dependent on-site potential [M(y)] in our two-band model] leading to a finite axial magnetic field, as described in this work. These two considerations render cold atomic systems as a natural platform to engineer and observe the effects presented.

To conclude, we have shown that inhomogeneous strain and magnetization have profound and observable implications on the electronic spectrum and transport properties of Dirac and Weyl semimetals. Moreover, our general analysis provides an alternative angle to explore distinct topological-to-trivial and topologicalto-topological interfaces, both smooth and abrupt [81]. The fundamental correspondence between Fermi arcs and pseudo-Landau levels and their connection to bound currents departs from naive expectations and can inspire future theoretical and experimental studies of surface effects in Weyl and Dirac semimetals.

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Note added.—While preparing this manuscript, we became aware of the recent related work by Pikulin *et al.* [27] discussing the effect of strain generated by torsion in inversion-breaking Weyl semimetal nanowires. Their findings are consistent with and complementary to our results. In particular, torsion in wires generates an axial magnetic field \mathbf{B}_5 that is parallel to the Weyl node separation rather than the perpendicular component described here.

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