Unification of Nonlinear Anomalous Hall Effect and Nonreciprocal Magnetoresistance in Metals by the Quantum Geometry

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The quantum geometry has significant consequences in determining transport and optical properties in quantum materials. Here, we use a semiclassical formalism coupled with perturbative corrections unifying the nonlinear anomalous Hall effect and nonreciprocal magnetoresistance (longitudinal resistance) from the quantum geometry. In the dc limit, both transverse and longitudinal nonlinear conductivities include a term due to the normalized quantum metric dipole. The quantum metric contribution is intrinsic and does not scale with the quasiparticle lifetime. We demonstrate the coexistence of a nonlinear anomalous Hall effect and nonreciprocal magnetoresistance in films of the doped antiferromagnetic topological insulator $MnBi_2Te_4$. Our work indicates that both longitudinal and transverse nonlinear transport provide a sensitive probe of the quantum geometry in solids.

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Introduction.-The quantum geometry of wave functions significantly impacts transport properties in quantum materials [1-3]. It is encoded in the quantum geometric tensor [4], which includes the Berry curvature [5] and a quantum metric [4,6,7]. As is well-known, the Berry curvature causes the intrinsic anomalous Hall effect in magnetic materials [8,9]. In similar vein, it has been suggested that the Berry curvature dipole [10] and the quantum metric [11] generate a nonlinear anomalous Hall effect (NLAHE). The former was predicted [12,13] and shortly after realized experimentally in Weyl semimetals (e.g., WTe₂ and MoTe₂) [14–16]. More recently, the latter has been predicted to appear in antiferromagnetic metals (e.g., CuMnAs) [17-19]. The NLAHE might be useful in optoelectronic applications such as terahertz detection and radio frequency rectification [20-22].

Another nonlinear transport phenomenon that has recently gained attention is the nonreciprocal magnetoresistance (NMR) (also called electric magnetochiral anisotropy) [23]. Here, the longitudinal resistance exhibits a secondorder correction that can be reversed by the magnetism or a magnetic field, acting as a magnetic diode. It is extensively studied in noncentrosymmetric or chiral materials [24–29]. The NMR is believed to originate either from the inelastic scattering by magnons [30] and spin clusters [31] or alternatively from the second-order Drude conductivity caused by an asymmetric band structure [32,33]. We note that previous theories based on the Berry curvature or quantum metric [11,17,18,34] lead to a vanishing second-order conductivity in the longitudinal direction.

In this Letter, we propose a unification of the NLAHE and NMR in the same theory framework of second-order perturbation theory, based on the quantum geometry. For the NLAHE, we find a significant interband correction related to the quantum metric. This quantum metric correction leads to a compact expression [i.e., Eq. (12)] for the nonlinear transport in metals and concomitantly predicts a NMR (i.e., a nonvanishing longitudinal nonlinear conductivity). Here, this NMR is created partially by the normalized quantum metric dipole, besides by the known Drude term. We further provide a scaling relation with the linear conductivity (σ_{\parallel}) to separate three contributions at second order: namely, the (i) normalized quantum metric dipole (independent of σ_{\parallel}), (ii) Berry curvature dipole (linear to σ_{\parallel}), and (iii) Drude weight (quadratic to σ_{\parallel}). These three contributions can be distinguished by symmetry restrictions, as summarized in Table I. We demonstrate the coexistence of a large NLAHE and the NMR in thin films of the well-studied, doped antiferromagnetic (AFM) topological insulator, MnBi₂Te₄ [35], and call for experimental verification. Our findings indicate that



FIG. 1. Schematics of the nonlinear anomalous Hall effect (NLAHE) and nonreciprocal magnetoresistance (NMR) due to the anomalous motion of wave packets induced by the quantum geometry. The dependence of the current components J_x , J_y with respect to the electric field E_x are illustrated on the right.

nonlinear longitudinal and transverse transports provide a sensitive probe for the quantum geometry in solids, as illustrated in Fig. 1.

Semiclassical kinetic equation.—We derive the nonlinear conductivity for a generic metal from a modification to the Boltzmann equation. Similar calculations have been performed very recently using the Kubo formalism [36–38], where the manipulations can be done order by order in the lifetime. In the following, we show that by combining perturbation theory and the Boltzmann approach, such an analysis reveals that the renormalization of quasiparticle properties enables longitudinal nonlinear current response, stemming from the quantum geometry.

Within the semiclassical approach, the nonlinear conductivity arises from a renormalization of the distribution function and quasiparticle operators due to the applied field, consistent with other approaches. In the semiclassical approach, corrections that are independent of scattering time arise *solely* due to correction to ground state quantities, while the scattering time enters through the time evolution of the distribution function. For the electron distribution function $f(\mathbf{r}, \mathbf{k}, t)$, the Boltzmann equation is,

$$\partial_t f + \frac{\mathbf{F}}{\hbar} \nabla_{\mathbf{k}} f + \mathbf{v} \nabla_{\mathbf{r}} f = \mathcal{I}(f). \tag{1}$$

Here, **F** is the semiclassical force in the presence of electric field, $\mathbf{F} = e\mathbf{E}$, **v** is the band diagonal velocity given by $v_n^c(\mathbf{k}) = \partial_{k_c} \varepsilon_n(\mathbf{k})$, with $\varepsilon_n(\mathbf{k})$ being the energy of Bloch state *n* with momentum **k**. For a uniform perturbation, we drop the spatial gradient acting on the distribution function. \mathcal{I} refers to the collision integral, and throughout this work we adopt the relaxation time approximation and set $\mathcal{I}(f) = -[(f - f_0)/\tau]$, where τ is the scattering time and f_0 is the Fermi Dirac distribution. The solution to the equation follows by an order-by-order (in electric field) expansion of the density, $f = f_0 + f_1 + f_2 + \cdots$. The zeroth order evolves trivially in time, as expected. Working in the frequency domain, the two relevant deviations from equilibrium are

$$i\omega f_1 + \frac{e\mathbf{E}}{\hbar}\partial_{\mathbf{k}}f_0 = -\frac{f_1}{\tau}, \quad i\omega f_2 + \frac{e\mathbf{E}}{\hbar}\partial_{\mathbf{k}}f_1 = -\frac{f_2}{\tau}.$$
 (2)

As the current is given by $j = -e \int_k vf$, it is imperative to consider correction to second order in the applied field not only in *f* but also in *v*.

We study these corrections using a modified form of the Luttinger-Kohn method [39]. In the presence of an electric field, the band Hamiltonian H_0 is modified by the coupling term $H_1 = -e\mathbf{E} \cdot \mathbf{r}$, where \mathbf{r} is the position operator. The effective low energy degrees of freedom are therefore given by a renormalized band Hamiltonian H', which can be obtained through the unitary transformation $H' = e^S H e^{-S}$. To fix *S*, we expand $e^S H e^{-S}$ to first order in \mathbf{E} , with the condition

$$H_1 + [S, H_0] = 0. (3)$$

This yields the matrix elements

$$S_{nm} = -e \frac{E_a \mathcal{A}_{nm}^a}{\varepsilon_{nm}}, \qquad n \neq m, \tag{4}$$

where $S_{nn} = 0$ and $\varepsilon_{nm} = \varepsilon_n - \varepsilon_m$. The interband Berry connection is defined as $\mathcal{A}^a_{nm} = \langle n\mathbf{k} | \hat{r} | m\mathbf{k} \rangle$. The immediate effect of the transformation *S* is in the renormalization of operators. The diagonal part of the Berry connection is analogously transformed, $\mathcal{A} \to e^S \mathcal{A}^{-S}$,

$$A_n^{\prime a} = \mathcal{A}_n^a + [S, \mathcal{A}^a]_n = \mathcal{A}_n^a - eE^bG_n^{ba}.$$
 (5)

The energy is accordingly renormalized,

$$\varepsilon'_n = \varepsilon_n(\mathbf{k}) - e\mathbf{E} \cdot \mathbf{r}'_n = \varepsilon_n(\mathbf{k}) + e^2 G_n^{ab} E_a E_b, \quad (6)$$

where $G_n^{ab} = \sum_{m \neq n} [(\mathcal{A}_{nm}^a \mathcal{A}_{mn}^b + \mathcal{A}_{nm}^b \mathcal{A}_{mn}^a)/\varepsilon_{nm}]$ is the bandnormalized quantum metric and \mathbf{r}'_n is the linear correction to \mathcal{A}_n , Eq. (5) (the full derivation of the correction to the Hamiltonian is found in the Supplemental Material [40]. The velocity operator incurs both a first- and second-order correction via dressing with *S*, giving

$$v_n^{\prime a} = \frac{\partial \varepsilon_n^{\prime}}{\partial k_a} + [S, v^a]_{nn} = v_n^{\prime a} - e\mathbf{E} \times \Omega_n^{\prime}.$$
(7)

The term linear in E is the familiar anomalous velocity, which is itself corrected by the electric field. The corrected Berry curvature is then obtained via

$$\Omega^{\prime\alpha\beta} = \partial_{\alpha}\mathcal{A}_{n}^{\prime\beta} - \partial_{\beta}\mathcal{A}_{n}^{\prime\alpha}.$$
 (8)

Scattering time analysis.—The solutions to Eq. (2) give terms that depend explicitly on the scattering time τ . Working iteratively, it is

$$f_1 = \frac{-e\mathbf{E}\nabla_{\mathbf{k}}f_0}{\hbar(i\omega+1/\tau)}, \qquad f_2 = \frac{e^2E_aE_b}{\hbar^2(i\omega+1/\tau)^2}\frac{\partial f_0}{\partial k_a\partial k_b}.$$
 (9)

We arrive at two charge current pieces along the c direction at second order in the electric field, which depend explicitly on the scattering time

$$j_{1}^{c} = -\frac{eE_{a}E_{b}}{\hbar} \sum_{n} \int_{\mathbf{k}} \frac{e^{2}}{(i\omega+1/\tau)^{2}} \frac{\partial^{2}f_{n}}{\partial k_{a}\partial k_{b}} \frac{\partial \varepsilon_{n}}{\partial k_{c}},$$

$$j_{2}^{c} = \frac{eE_{a}E_{b}}{\hbar} \sum_{n} \int_{\mathbf{k}} \frac{e^{2}}{(i\omega+1/\tau)} \frac{\partial f_{n}}{\partial k_{a}} \Omega_{n}^{bc} + (a \leftrightarrow b). \quad (10)$$

Here, j_1^c results from combining the second-order correction to the density f_2 [right-hand term in Eq. (9)] with the unperturbed band velocity $v_n^c = (\partial \varepsilon_n / \partial k_c)$. j_2^c , similarly

draws from the perturbed velocity in Eq. (7) with the firstorder correction in f_1 [left-hand term in Eq. (9)]. Note that the sum is over all ground state single-particle bands *n*. Besides the terms generated by the perturbation of the semiclassical density $(j_{1,2}^c)$, there are extra corrections to the operators. The associated currents are due to corrections to the dispersion (j_{disp}^c) and to the anomalous velocity (j_{anom}^c) , respectively. We stress that although these currents couple to the equilibrium density, they exist only for finite electric field. They are given by

$$j_{\rm disp}^c = -\frac{eE_a E_b}{\hbar} \int_{\mathbf{k}} f_0 v_n^{\prime c} = -\frac{e^3 E_a E_b}{\hbar} \sum_n \int_{\mathbf{k}} f_n \partial_{k_c} G_n^{ab},$$

$$j_{\rm anom}^c = \frac{e^2}{\hbar} \int_{\mathbf{k}} \frac{f_0}{2} (E_a \Delta \Omega_n^{ac} + E_b \Delta \Omega_n^{bc}), \tag{11}$$

where in the last line the expressions are written manifestly symmetric with respect to $(a \leftrightarrow b)$. We also include the linear-in-field contribution to the Berry curvature with the notation $\Delta\Omega^{ac} = -(eE_a\partial_{k_a}G_n^{ba} - eE_b\partial_{k_c}G_n^{cb})$. Assembling all terms $(j_1^c, j_2^c, j_{disp}^c, j_{anom}^c)$, one arrives at the conductivity, which is given by j^c/E^aE^b ,

$$\sigma^{ab;c} = -\frac{e^3\tau^2}{\hbar^3} \sum_n \int_k f_n \partial_{k^a} \partial_{k^b} \partial_{k^c} \varepsilon_n \tag{12a}$$

$$-\frac{e^{3}\tau}{\hbar^{2}}\sum_{n}\int_{k}f_{n}(\partial_{k^{a}}\Omega_{n}^{bc}+\partial_{k^{b}}\Omega_{n}^{ac})$$
(12b)

$$-\frac{e^3}{\hbar}\sum_n \int_k f_n \left(2\partial_{k^c} G_n^{ab} - \frac{1}{2}(\partial_{k^a} G_n^{bc} + \partial_{k^b} G_n^{ac})\right). \quad (12c)$$

We observe that $\partial_{k^a} G_n^{bc}$ is the band-normalized quantum metric dipole. Equation (12a) refers directly to the nonlinear Drude weight [36] and is obtained by integrating twice by parts the first line in Eq. (10). Similarly, the Berry curvature dipole Eq. (12b) is obtained by integrating by parts the second line of Eq. (10), in agreement with Ref. [10]. The intrinsic contribution, which is τ -independent in Eq. (12c), is caused by the band-normalized quantum metric dipole. The Fermi surface contribution to the current of this intrinsic term is obtained again by integrating by parts. It should be noted that Eq. (12c) violates the (a, b, c) cyclic permutation symmetry, the source of which is a gravitational anomaly [19].

Discussion.—As demonstrated, Eq. (12c) naturally decomposes into j_{disp}^c , which propagates with the current direction $\partial_{k^c} G_n^{ab}$, and the purely transverse current j_{anom}^c , related to $\partial_{k^c} G_n^{ab} - \frac{1}{2}(\partial_{k^a} G_n^{bc} + \partial_{k^b} G_n^{ac})$, which vanishes whenever a = b = c [11,46]. Therefore, Eq. (12) naturally unifies both longitudinal and transverse effects at nonlinear order. Specifically, we find that the longitudinal component (i.e., the NMR) has contributions from both the Drude

conductivity ($\sim \tau^2$) and the band-normalized quantum metric dipole, independent of τ . On the other hand, the NLAHE is constituted by the Berry curvature dipole at order τ , and again the band-normalized quantum metric dipole. The band-normalized quantum metric G_n^{ab} thus enters in both the longitudinal and the transverse components.

The second-order conductivity has received much attention lately, with some variation in the precise form of the expressions due to the large number of terms involved [11,38,47,48]. While there is some disagreement between different approaches regarding coefficients, in all works there exists a broad consensus that the nonlinear conductivity is nonzero at all orders of τ . A simple criterion to validate our result is that Eq. (12) respects the intrinsic permutation symmetry when exchanging E_a and E_b , as required by the dc response. The common geometric origin of longitudinal and transverse components shows that unlike for the linear conductivity, at second order, valuable information about the band structure geometry is accessible in either spatial component.

We point out two possible limitations of our calculation. First, second-order perturbation theory captures the instantaneous response of the system, but is insensitive to nonperturbative effects like a steady-state equilibration at long times. Second, we focused on effects of the band structure in a system with finite quantum lifetime τ . Further extrinsic contributions to the conductivity beyond the relaxation time approximation were neglected. In principle, the nonlinear Hall effect is composed of extrinsic contributions [49,50] that induce corrections to the nonlinear Hall conductivity to all three contributions, at order τ^2 , τ , and τ^0 . However, for a collinear antiferromagnet (such as MnBi₂T₄ discussed in the following section) skew scattering contributions are expected to be significantly suppressed [47]. More exotic mechanisms such as anomalous skew scattering proposed by Ref. [51] require breaking of C_{3z} symmetry to appear, and are therefore not relevant in MnBi₂Te₄.

To detect the signatures of the quantum geometry in experiments, we suggest examining the scaling relation between the transverse second order $(\sigma_{\perp}^{(2)})$ and longitudinal linear conductivity (σ_{\parallel}) , which reads

$$\sigma_{\perp}^{(2)} = \eta_2(\sigma_{\parallel})^2 + \eta_1 \sigma_{\parallel} + \eta_0.$$
(13)

Here, the coefficients η_i denote the respective part of the nonlinear conductivity that contributes at order $\mathcal{O}(\tau^i)$. Namely, η_2 contains the nonlinear Drude term, and if present contributions from skew scattering [50]. η_1 contains the Berry curvature dipole term, as well as extrinsic contributions related to side jump scattering. η^0 contains the effect of the normalized quantum metric. Similarly, for the longitudinal components it holds that

$$\sigma_{\parallel}^{(2)} = \eta_2'(\sigma_{\parallel})^2 + \eta_1'\sigma_{\parallel} + \eta_0', \qquad (14)$$

TABLE I. Symmetry restrictions for three contributions in Eq. (12) for a two-dimensional system regarding the inversion symmetry (\mathcal{P}), time-reversal symmetry (\mathcal{T}), and combined \mathcal{PT} symmetry. With the rotational symmetries C_{3z} related to MnBi₂Te₄ film, both longitudinal (\mathbf{j}_{\parallel}) and transverse currents (\mathbf{j}_{\perp}) exists. We note that \mathbf{j}_{\parallel} is dissipative because $\mathbf{j} \cdot \mathbf{E} \neq 0$, despite that the quantum metric-induced \mathbf{j}_{\parallel} is τ -independent.

Mechanism	C_{3z}	\mathcal{P}	Τ	$\mathcal{P}\mathcal{T}$	j∥	j⊥
Nonlinear Drude	1	X	X	1	1	1
Berry curvature dipole	X	X	\checkmark	X	X	1
Quantum metric	\checkmark	X	×	\checkmark	\checkmark	1

where η'_1 will be nonzero only if there are extrinsic contributions. Using these scaling relations, Eqs. (13) and (14), it is therefore possible to isolate each of these terms and to quantify both the NLAHE and the NMR.

Realization in a magnetic metal.—In Eq. (12), the Drude term and the normalized quantum metric dipole are antisymmetric under momentum inversion $(k \rightarrow -k)$, which is related to the inversion symmetry (\mathcal{P}) or time-reversal symmetry (\mathcal{T}). Thus, breaking both \mathcal{P} and \mathcal{T} is required to obtain a nonzero contribution over the full Brillouin zone. In contrast, the Berry curvature dipole integral requires breaking both \mathcal{P} and the combined symmetry \mathcal{PT} (see Table I).

While the NLAHE induced by the Berry curvature dipole was already observed in many materials, it is much harder to realize the quantum metric dipole-driven NLAHE or likewise the NMR. We will now demonstrate the coexistence of both NLAHE and NMR in a doped AFM topological insulator, $MnBi_2Te_4$. $MnBi_2Te_4$ is a layered Van der Waals material with the A-type AFM structure. Thin film of this material with an even number of layers break \mathcal{P} and \mathcal{T} , but preserve \mathcal{PT} . The \mathcal{PT} symmetry specifically excludes the Berry curvature dipole

contribution in the NLAHE, so that we can focus on the effect of the quantum metric. Experimentally, \mathcal{PT} seems to be weakly broken in some MnBi₂Te₄ samples. For example, \mathcal{PT} breaking was witnessed by the finite anomalous Hall signal for a six-layer-thick film [52]. However, even in this case the Berry dipole contribution to the NLAHE is still strongly suppressed by the threefold rotational symmetry (C_{37}).

The crystal symmetry helps us understand the shape of nonlinear conductivity tensor. In a 2D film of $MnBi_2Te_4$, we set *x* along the lattice vector direction in the basal plane for convenience. Here, C_{3z} constrains that the NLAHE and NMR share the same amplitude but opposite sign, i.e., it holds that

$$\sigma^{yy;x} = -\sigma^{xx;x}, \qquad \sigma^{xx;y} = -\sigma^{yy;y}$$

The combined symmetry [53] by mirror reflection $(\mathcal{M}_x, x \to -x)$ and \mathcal{T} then enforces $\sigma^{xx;y} = -\sigma^{yy;y} = 0$. Therefore, we only have one independent nonlinear conductivity, the NLAHE conductivity $\sigma^{yy;x}$, or equivalently the NMR conductivity $\sigma^{xx;x}$. When rotating the sample, we obtain the angle (θ) dependence in the new coordinates (x', y') by $\sigma^{y'y';x'} = -\sigma^{x'x';x'} = \cos(3\theta)\sigma^{yy;x}$.

Figure 2 shows the band structure and $\sigma^{yy;x}$ calculated on a eight-layer-thick (8L) film by first-principles methods. We carried out *ab initio* density-functional calculations [54] on slab models of 2,4,6,8 layered MnBi₂Te₄ with AFM order. We then projected the converged wave functions of each slab onto local Wannier functions [55] of Bi-*p* and Te*p* orbitals, which accurately span the energy window around the Fermi level. The energy dispersion is asymmetric between $\Gamma - \bar{K}$ and $\Gamma - K$ because both \mathcal{P} and \mathcal{T} are broken. Each energy state is furthermore doubly degenerate due to \mathcal{PT} symmetry. The lowest conduction bands and highest valence bands contribute opposite signs in the nonlinear conductivity. Between **k** and $-\mathbf{k}$, bands



FIG. 2. Band structure and nonlinear conductivity of antiferromagnetic MnBi₂Te₄ thin films ($\sigma^{yy;x} \equiv -\sigma^{xx;x}$ in this case). (a) The band structure of an eight-layer (8L) thick film. The band-decomposed contribution to $\sigma^{yy;x}$ is indicated by the color. (b)–(e) The energy dependence of $\sigma^{yy;x}$ for different films with the contribution from quantum metric (QM).

contribute opposite signs to $\sigma^{yy;x}$, but at different amplitude. Indeed, the asymmetry between **k** and $-\mathbf{k}$ leads to nonzero $\sigma^{yy;x}$. As varying the Fermi energy, $\sigma^{yy;x}$ shows sign changes when a group of new bands appear at the Fermi surface. It vanishes in the energy gap. Upon increasing the number of layers from two to eight layers, the region near to the lowest conduction band exhibits comparatively small changes in $\sigma^{yy;x}$, while the valence band region changes dramatically [see Fig. 2(c)]. This is related to the fact that the lowest conduction bands are composed of gapped Dirac surface states while the top valence states have a bulk origin [42]. We therefore conclude that films thicker than two layers have similar surface states in the lowest conduction bands.

Numerically, $\sigma^{yy;x}$ is in the order of magnitude of several mA nm V⁻², when using a relaxation time τ = 0.04 ps to evaluate the Drude weight. As MnBi₂Te₄ samples commonly suffer from defects and exhibit a low mobility [56], the Drude contribution may be smaller in reality. The presence of defects may also change the magnitude of the conductivity due to the decreased band gap and renormalization of surface state dispersion. A calculation for a defective slab is presented in the Supplemental Material. Given that similar transport devices have recently become readily available [29,57], MnBi₂Te₄ films are ideal candidates to explore the quantum metric-driven NLAHE and NMR.

Summary.—We have shown that signatures of the quantum geometry of the band structure are imprinted in the second-order conductivity not only in the transverse components but also in the longitudinal ones. To this end, we derived a NLAHE and NMR, which both appear due to the quantum metric, and explored their effect in thin films of the \mathcal{PT} -symmetric antiferromagnet MnBi₂Te₄. We found an intrinsic contribution to both NLAHE and NMR in antiferromagnetic thick films, providing an ideal platform to detect the quantum metric.

Our results further strengthen the observation that nonlinear responses carry more intricate and at the same time much more interesting information about the quantum geometry than linear response functions [58,59]. It is imperative to further explore these aspects systematically in theory, for example for finite frequency response functions, and also for the magnetotransport. At the same time, present sample quality and device technology have the capabilities to detect these phenomena experimentally.

Note added.—We note that the quantum metric-induced NLAHE and NMR were observed in $MnBi_2Te_4$ thin films by a very recent experiment [60].

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