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Berry curvature driven transverse thermoelectric generation in topological magnets

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ABSTRACT

Topological magnets such as magnetic Weyl and nodal-line semimetals possess topologically non-trivial band structures in magnetically ordered states. In this class of materials, the Berry curvature in momentum space can be significantly enhanced, resulting in thermoelectric responses that exceed empirical scaling laws based on magnetization. Such large transverse thermoelectric effects enable flexible thin-film-based lateral device structures, leading to novel energy-harvesting technologies and sensors beneficial for Internet of Things (IoT) sensors and wearable devices. In this review, we outline recent progress in the study of the large transverse thermoelectric effects in topological magnets.

KEYWORDS

topological magnets; Weyl semimetals; nodal line semimetals; anomalous Nernst effect

1. Introduction

Thermoelectric effects, such as the Seebeck effect, directly convert heat into electricity (and vice versa) in solids and have been extensively studied for applications in heat exchange, waste heat utilization, and as power sources for IoT sensors [1, 2, 3]. In particular, Peltier devices based on the Seebeck effect are now widely used as wearable coolers (Fig. 1). Recently, the transverse thermoelectric conversion in magnets, known as the anomalous Nernst effect (ANE), has attracted much attention because it allows a lateral configuration of the thermoelectric modules that can efficiently cover heat sources (Fig. 1) [4, 5, 6, 7]. Extensive studies on the transverse thermodynamic effects have identified topological magnets as a key material class exhibiting a large ANE [7], which we review in this article.

Recent developments in condensed matter physics have revealed that non-trivial topological band structures near the Fermi energy ε_F are important for understanding the unique properties of materials [8, 9, 10, 11]. Among them, topological magnets

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such as magnetic Weyl semimetals and nodal-line semimetals have been found to exhibit large transverse responses such as the ANE and anomalous Hall effect (AHE) [12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 5, 7, 28] owing to the non-vanishing momentum-space (k -space) Berry curvature. Various research groups have demonstrated ANE-based thermoelectric devices [4, 29, 30, 31, 32, 33, 34, 35], which may be useful for energy harvesting from waste heat, IoT sensors, and wearable devices.

In this article, we review the recent advances in research and development of transverse thermoelectric generation in topological magnets. We first introduce the ANE as the transverse thermoelectric generation for magnetic materials (Sec. 2). Then, we briefly illustrate the theoretical framework for the Berry curvature-driven intrinsic ANE in topological magnets (Secs. 3-5). We also review the experimental observations for ANE in various typical topological magnets (Sec. 6). Finally, we provide a summary and perspective (Sec. 7).

2. Anomalous Nernst effect –transverse thermoelectric effect in magnetic materials

Generally in solids, the electric current density \mathbf{J}_c , is proportional to the electric field \mathbf{E} , $\mathbf{J}_c = \sigma\mathbf{E}$, known as Ohm's law. Similarly, the heat current density \mathbf{J}_q is proportional to the temperature gradient ∇T , $\mathbf{J}_q = \kappa(-\nabla T)$, known as Fourier's law. Here, σ and κ are electric and thermal conductivity tensors. In metals and semiconductors, carriers (electrons or holes) transport both electric and heat current. This means that \mathbf{E} (∇T) also generates \mathbf{J}_q (\mathbf{J}_c). Thus, the above equations are generalized in the linear response regime as,

$$\mathbf{J}_c = \sigma\mathbf{E} + \alpha(-\nabla T), \quad (1)$$

$$\mathbf{J}_q = \Pi\mathbf{E} + \kappa(-\nabla T). \quad (2)$$

The newly added cross-correlated terms represent the thermoelectric effects with thermoelectric tensors α and Π . According to the Onsager reciprocal relations, α and Π are not independent and satisfy $\Pi = \alpha T$ [36, 37, 38]. Besides, the Onsager reciprocal relations also hold for the off-diagonal terms of the conductivity tensors,

$$\sigma_{ij}(B) = \sigma_{ji}(-B), \alpha_{ij}(B) = \alpha_{ji}(-B), \kappa_{ij}(B) = \kappa_{ji}(-B), \quad (3)$$

where σ_{ji} , α_{ji} and κ_{ji} are respectively called Hall conductivity, transverse thermoelectric conductivity and thermal Hall conductivity, and B is the magnetic field. The Eq. (3) implies $\sigma_{ji} = \alpha_{ji} = \kappa_{ji} = 0$ when both magnetic field B and magnetization M are zero ($B = M = 0$).

The Onsager reciprocal relations are fundamental for irreversible transport processes originating from microscopic time-reversal symmetry. Thus, they are never

violated as long as the system satisfies their precondition; the states at $+B$ and $-B$ are equivalent to each other after the time-reversal operation. In other words, the relations such as $\sigma_{ij}(B) \neq \sigma_{ji}(-B)$, $\alpha_{ij}(B) \neq \alpha_{ji}(-B)$, may occur in hysteretic regimes due to magnetic domains or exchange bias. Additionally, *non-Onsager* quantities such as the Nernst coefficient S_{ij} (defined as Eq. (8) below) do not obey this rule. Thus, $S_{ij}(B)$ is not necessarily equal to $S_{ji}(-B)$ as observed in YbMnBi₂ [25, 26]. (These two cases should not be called “violations” of the Onsager relations since they simply fall outside the scope of the principles.)

Anomalous Hall effect

The electrical resistivity ρ is the inverse of σ , $\rho = \sigma^{-1}$, and xx and xy components are given by

$$\sigma_{xx} = \frac{\rho_{yy}}{\rho_{xx}\rho_{yy} - \rho_{xy}\rho_{yx}}, \quad \sigma_{xy} = -\frac{\rho_{xy}}{\rho_{xx}\rho_{yy} - \rho_{xy}\rho_{yx}}, \quad (4)$$

where we consider the net flow of carriers along the xy plane, $\mathbf{J}_c \perp \mathbf{z} \parallel \mathbf{B}$. Thus, we get $\sigma_{xx} \sim 1/\rho_{xx}$, $\sigma_{yx} = -\sigma_{xy} \sim -\rho_{yx}/\rho_{xx}^2$ for conventional isotropic 3D metals where $\rho_{xx} \sim \rho_{yy} \gg |\rho_{yx}|$ and $\rho_{xy}(B) = -\rho_{yx}(B)$. Experimentally, we measure $\rho_{xx} = J_x/E_x$, and $\rho_{yx} = E_y/J_x$ and obtain σ_{xx} and σ_{xy} , where J_i and E_i are the applied current density and measured electric field along i -direction, respectively.

Historically, Edwin Hall first discovered the Hall effect in nonmagnetic metals and ferromagnets [39, 40]. Subsequent studies have established the empirical relation

$$\rho_{yx} = R_0 B + R_s M, \quad (5)$$

where R_0 and R_s are the ordinary Hall and anomalous Hall coefficients, respectively [41]. The first term is the ordinary Hall effect (OHE), related to the carrier density n and charge q as $R_0 = B/(nq)$. The second term represents the anomalous Hall effect (AHE), which typically dominates in ferromagnetic states. This has been intuitively interpreted as the consequence of the Lorentz force due to M , and thus it was long believed that AHE should not occur in antiferromagnets, where M is vanishingly small.

However, this empirical relation (Eq. (5)) and the corresponding intuitive interpretation have turned out to be inaccurate; they neglect contributions from Berry curvatures in both real and momentum space. The real-space Berry curvature originates, for example, from skyrmions in noncentrosymmetric magnets [42, 43, 44, 45, 46, 47, 48]. The k -space Berry curvature leads to the intrinsic AHE, which is distinct from the extrinsic AHE caused by disorder-related mechanisms such as skew scattering and side jump. Importantly, the intrinsic AHE is not necessarily proportional to M and can be significant in noncollinear and noncoplanar antiferromagnets (AFM) [49, 50, 51, 7, 28] and even finite in the collinear antiferromagnetic known as altermagnets [52, 53, 54, 55]. In Sec. 4, we discuss the intrinsic (k -space Berry curvature driven) AHE in detail and explain how large AHE signals can arise in both ferromagnets and antiferromagnets.

Anomalous Nernst effect

The longitudinal thermoelectric effect, the Seebeck coefficient S_{ii} , is obtained by setting $\mathbf{J}_c = 0, \nabla T \parallel \mathbf{x}$ and $\sigma_{ij} = \alpha_{ij} = 0$ in Eq. (1),

$$S_{xx} \equiv \frac{E_x}{-(\partial T/\partial x)} = \frac{V}{\Delta T} = \frac{\alpha_{xx}}{\sigma_{xx}}. \quad (6)$$

Note that the sample length cancels out and does not appear in Eq. (6) (see Fig. 2 (a)).

In contrast, the transverse thermoelectric effect (Nernst effect) can be finite by breaking the time-reversal symmetry i.e., $B \neq 0$ or $M \neq 0$. Similar to the Hall effect, B -linear and M -linear terms are called ordinary Nernst effect (ONE) and ANE, respectively.

By setting $\mathbf{J}_c = 0, \nabla T \parallel \mathbf{x}$ in Eq. (1), we get

$$\alpha_{yx} = \sigma_{yx} S_{xx} + \sigma_{yy} S_{yx}, \quad (7)$$

where the Nernst coefficient S_{yx} is defined as

$$S_{yx} \equiv \frac{E_y}{-(\partial T/\partial x)} = \frac{\ell_T}{\ell_V} \frac{V_y}{\Delta T}. \quad (8)$$

Interestingly, the Nernst voltage V_y is proportional not only to the temperature difference ΔT but also to the sample geometry ℓ_V/ℓ_T , where ℓ_V and ℓ_T respectively represent distances between voltage and temperature terminals (Fig. 2 (b)). Thus, one can enhance V_y in thin-film thermopile structure by reducing ℓ_T and increasing ℓ_V (Fig. 1 (b)), as demonstrated by various experiments [5, 4, 30, 31, 32, 56, 34]. Eq. (7) immediately provides

$$-S_{yx} = -\alpha_{yx} \rho_{yy} + \sigma_{yx} \rho_{yy} S_{xx} \equiv S_1 + S_2. \quad (9)$$

Here, S_1 and S_2 respectively indicate the contributions directly from α_{yx} and longitudinal thermoelectric effect S_{xx} converted to the transverse direction through σ_{yx} . Thus, in addition to large $|\alpha_{yx}|$, these two contributions should be constructive (namely $\text{sgn}(-\alpha_{yx}) = \text{sgn}(\sigma_{yx} S_{xx})$) for enhancing $|S_{yx}|$.

3. Berry phase, Berry connection and Berry curvature

Topology in condensed matter physics appears from the Hilbert space geometry [57]. For simplicity, let us assume a nondegenerate complete orthonormal system $\{|u_n\rangle\}$

$$H(\mathbf{R})|u_n(\mathbf{R})\rangle = E_n(\mathbf{R})|u_n(\mathbf{R})\rangle, \quad (10)$$

where E_n is the eigenenergy for the n -th eigen state $|u_n\rangle$, and $\mathbf{R}(\mathbf{t})$ is a parameter for the Hamiltonian $H(\mathbf{R})$ such as the wave vector \mathbf{k} and real space position \mathbf{r} . Then, the time-

evolution for the initial state $|\psi(t)\rangle = \sum_n c_n |u_n(\mathbf{R})\rangle$ obtained from the time-dependent Schrödinger equation $i\hbar \frac{d}{dt} |\psi(t)\rangle = \mathcal{H}(\mathbf{R}) |\psi(t)\rangle$ is

$$i\hbar \frac{d}{dt} c_n + i\hbar c_n \langle u_n | \frac{d}{dt} |u_n\rangle + i\hbar \sum_{m \neq n} \frac{\langle u_n | \frac{d}{dt} \mathcal{H} |u_m\rangle}{E_m - E_n} c_m = c_n E_n \quad (11)$$

On the other hand, the time derivative of Eq. (10) gives $\langle u_n | \frac{d}{dt} |u_m\rangle = \frac{\langle u_n | \frac{d}{dt} \mathcal{H} |u_m\rangle}{E_m - E_n}$ for $n \neq m$. This indicates the last term on the left-hand side of Eq. (11) is negligible compared to the diagonal term $\langle u_n | \frac{d}{dt} |u_n\rangle$ in the adiabatic approximation, where the time-evolved state stays at the initial eigenstate with a certain phase difference ϕ ; $|u_n(\mathbf{R}(t))\rangle \sim |u_n(\mathbf{R}(0))\rangle \exp(i\phi)$. Solving Eq. (11), we obtain

$$c_n(t) = c_n(0) \exp(i\theta_n(\mathbf{R})) \exp(i\gamma_n(\mathbf{R})), \quad (12)$$

where $\theta_n(\mathbf{R}) \equiv -\frac{1}{\hbar} \int_{\mathbf{R}_0}^{\mathbf{R}} E_n(\mathbf{R}') d_{\mathbf{R}}$ and $\gamma_n(\mathbf{R}) \equiv i \int_{\mathbf{R}_0}^{\mathbf{R}} \langle u_n(\mathbf{R}') | \frac{\partial}{\partial \mathbf{R}'} |u_n(\mathbf{R}')\rangle d_{\mathbf{R}'}$ are known as the dynamical and geometrical phases, respectively. The former is the conventional term originating from the time evolution of the eigenstate. The latter is the topological term depending on the path C from \mathbf{R}_0 to \mathbf{R} . While the above $\gamma_n(\mathbf{R})$ is gauge-dependent and not a physical observable, the closed-loop integral so-called the Berry phase

$$\gamma_n \equiv \int_C \mathbf{A}_n(\mathbf{R}') d_{\mathbf{R}'} = \int_S \mathbf{\Omega}_n(\mathbf{R}') d\mathbf{S}' \gamma(C) \quad (13)$$

becomes gauge-independent. Here,

$$\mathbf{A}_n(\mathbf{R}) \equiv i \langle u_n(\mathbf{R}) | \nabla_{\mathbf{R}} |u_n(\mathbf{R})\rangle \quad (14)$$

and

$$\mathbf{\Omega}_n(\mathbf{R}') \equiv \nabla_{\mathbf{R}'} \times \mathbf{A}_n(\mathbf{R}') \quad (15)$$

are the Berry connection and Berry curvature, respectively. In real space, $\gamma_n(C), \mathbf{A}_n(\mathbf{r}), \mathbf{\Omega}_n(\mathbf{r})$ correspond to the Aharonov-Bohm phase, vector potential and magnetic field, respectively. In this review, we focus on the k -space Berry curvature $\mathbf{\Omega}_n(\mathbf{k}) = \nabla_{\mathbf{k}} \times \mathbf{A}_n(\mathbf{k}) = i \nabla_{\mathbf{k}} \times \langle u_n(\mathbf{k}) | \nabla_{\mathbf{k}} |u_n(\mathbf{k})\rangle$ as an origin of the large transverse responses in topological magnets, where $|u_n(\mathbf{k})\rangle$ is the periodic part of the Bloch wave function $|\psi_{n,\mathbf{k}}\rangle = e^{i\mathbf{k}\mathbf{r}} |u_n(\mathbf{k})\rangle$.

4. Berry curvature driven intrinsic transverse thermoelectric effect

Similar to σ_{ij} , α_{ij} can originate from both intrinsic and extrinsic mechanisms. Here, the term ‘‘intrinsic’’ refers to the contribution from the k -space Berry curvature as discussed in Sec. 2. In MnBi, the experimentally observed α_{ij} is one order of magnitude larger than the theoretical value assuming the intrinsic mechanism, indicating large

extrinsic contributions [58]. However, this is so far an exception; the intrinsic mechanism dominates in most of magnetic materials with large ANE. Thus, in this review, we focus on the intrinsic ANE driven by k -space Berry curvature. In this section, we first introduce the mechanism for the intrinsic AHE as it is related to the origin of the intrinsic ANE.

Intrinsic Anomalous Hall conductivity

Using Kubo-formula we obtain [59, 60]

$$\sigma_{xy}(T, \mu) = -\frac{e^2}{\hbar} \int \frac{d^D k}{(2\pi)^D} \sum_n f_{nk} \Omega_n^z(\mathbf{k}), \quad (16)$$

where $D(= 2,3)$, e , \hbar , μ , ε_{nk} , and $f_{nk} = 1/(e^{\varepsilon_{nk}-\mu}/k_B T + 1)$ are the spatial dimension of the system, elementary charge, reduced Planck constant, chemical potential, energy for the band n , and the Fermi-Dirac distribution function, respectively. The equation (16) yields the quantum Hall effect for 2D-band insulator at $T = 0$

$$\sigma_{xy}^{\text{QHE}} = -\frac{e^2}{\hbar} \int_{BZ} \frac{d^2 k}{(2\pi)^2} \sum_n^{\text{occupied}} \Omega_n^z(\mathbf{k}) = -\frac{e^2}{\hbar} C, \quad (17)$$

where $C = 1, 2, 3 \dots$ is the Chern number and matches with the number of occupied states [61]. Thus, the intrinsic AHE in 3D metals can be viewed as an "unquantized" version of the quantum Hall effect. The Berry curvature $\Omega_n^z(\mathbf{k})$ in Eq. (16) is numerically computed by using

$$\Omega_n^z(\mathbf{k}) = -2\text{Im} \sum_{m \neq n} \frac{\langle u_n(\mathbf{k}) | \frac{\partial H(\mathbf{k})}{\partial k_x} | u_m(\mathbf{k}) \rangle \langle u_m(\mathbf{k}) | \frac{\partial H(\mathbf{k})}{\partial k_y} | u_n(\mathbf{k}) \rangle}{(\varepsilon_{m\mathbf{k}} - \varepsilon_{n\mathbf{k}})^2}, \quad (18)$$

which means that the magnitude of the Berry curvature $|\Omega_n^z(\mathbf{k})|$ becomes singularly large when two bands touch. This explains why topological magnets with band crossings (Fig. 4 and Sec. 5) show a large AHE.

In ferromagnets, the spontaneous time-reversal symmetry breaking ensures a finite σ_{xy} . By contrast, antiferromagnets require additional constraints for the non-vanishing AHE, i.e. $\sigma_{xy} = 0$ for the simplest collinear antiferromagnets, where the spin structure is invariant time-reversal + half-unit cell translation [62, 63, 28]. Examples include Mn_3X ($X = \text{Sn, Ge}$) as shown in Sec 6.

Intrinsic transverse thermoelectric conductivity

Similar to σ_{xy} , the transverse thermoelectric conductivity α_{xy} also arises from the intrinsic Berry curvature mechanism [64], i.e.,

$$\alpha_{xy}(T, \mu) = \frac{|e|}{T\hbar} \int \frac{d\mathbf{k}}{(2\pi)^3} \Omega_{n,z}(\mathbf{k}) \{(\varepsilon_{n\mathbf{k}} - \mu) f_{n\mathbf{k}} + k_B T \log[1 + e^{-\beta(\varepsilon_{n\mathbf{k}} - \mu)}]\} \quad (19)$$

$$= \frac{1}{|e|} \int d\varepsilon \sigma_{xy}(0, \varepsilon) \frac{\varepsilon - \mu}{T} \left(-\frac{\partial f}{\partial \varepsilon} \right) \quad (20)$$

$$= -\frac{k_B}{|e|} \int d\varepsilon \frac{\partial \sigma_{xy}(T, \varepsilon)}{\partial \varepsilon} s(\varepsilon, T), \quad (21)$$

where $s = -f \ln(f) - (1 - f) \ln(1 - f)$ is the entropy density and $\partial \sigma_{xy} / \partial \varepsilon =$

$\sum_{n\mathbf{k}} \Omega_{n,z}(\mathbf{k}) \delta(\varepsilon - \varepsilon_{n\mathbf{k}})$. Using the Sommerfeld expansion we obtain the low-temperature asymptotic form called the Mott relation

$$\alpha_{xy} = -\frac{\pi^2 k_B^2 T}{3|e|} \left. \frac{\partial \sigma_{xy}(0, \varepsilon)}{\partial \varepsilon} \right|_{\varepsilon = E_F}. \quad (22)$$

This implies that α_{xy}/T becomes large when σ_{xy} varies rapidly as a function of ε owing to topological band structures (Fig. 3) [65].

σ_{xy} and α_{xy} are often numerically computed using Eq. (16) and Eq. (19) by firstprinciples calculations, respectively. While σ_{xy} is given by the sum of the Berry curvature below $\varepsilon \sim \mu$ (Eq. (16)), α_{xy} depends on the Berry curvature and density of states (DOS) near $\varepsilon \sim \mu$ (Eq. (19)). Thus, topological flat band structures that produce both large Berry curvature and DOS near the Fermi energy are essential for enhancing α_{xy} . Examples include the titled Weyl cones (Co₂MnGa) [18], nodal web structure (Fe₃Ga) [21, 66], and nodal plane (Fe₃Sn) [24], as discussed in Sec. 5.

5. Topological band structures inducing large Berry curvatures

As discussed in Sec. 4, Berry curvature is highly enhanced near band crossings. Such band crossings occur at either (i) a k point (Weyl point, Fig. 4(a)), (ii) a line (nodal line, Fig. 4(b)), or (iii) a plane (nodal plane, Fig. 4(c)). The nodal line and plane are defined in the absence of the spin-orbit coupling (SOC). After introducing SOC, Weyl points may persist, otherwise a small gap opens all over the nodal line (plane).

The Weyl point denotes a non-degenerate linear band crossing $\varepsilon_{\mathbf{k}} \sim \mathbb{V}|\mathbf{k} \pm \mathbf{k}_0|$ and their model Hamiltonian is identical to the time-independent Weyl equation $H\psi_{\mathbf{k}} \sim \mathbb{V}(\mathbf{k} \pm \mathbf{k}_0) \sigma \psi_{\mathbf{k}} = \varepsilon_{\mathbf{k}} \psi_{\mathbf{k}}$ [12, 13, 67]. In particular, non-vanishing AHE and ANE appear in magnetic Weyl semimetals (i.e. Weyl magnets), possessing Weyl points near the Fermi energy ε_F in a magnetically ordered state. Each Weyl point carries the chirality of ± 1 and always pairs with the opposite signs [68]. These Weyl points act as either sources or sinks of the Berry curvature (Fig. 4(a)), inducing AHE and ANE.

6. Experimental realization -large ANE in topological magnets

Until now, various topological magnets have been discovered that exhibit large ANEs. In this section, we first describe the universal features shared across these materials, followed by details on specific compounds. The materials highlighted here are some of

the the best-studied topological magnets, i.e., Mn_3X ($X = Sn, Ge$) [15, 16, 17, 22], Co_2MnGa [18], Fe_3X ($X = Ga, Al$) [21], $YbMnBi_2$ [25, 26], Fe_3Sn [24], $Co_3Sn_2S_2$ [19, 20] and $UCo_{0.8}Ru_{0.2}Al$ [23].

(1) Dominant intrinsic contribution to σ_{xy} and α_{xy} .

The intrinsic Berry curvature contribution dominates both σ_{xy} and α_{xy} in topological magnets. For example, first-principles calculation that consider only the intrinsic contribution (Eq. (16) and Eq. (19)) successfully reproduce the experimental σ_{xy} and α_{xy} . The enhanced σ_{xy} and α_{xy} are associated with topological band structures, which are often confirmed by the ARPES (Angle-resolved Photoemission Spectroscopy) measurements. In most of these topological magnets, electron correlation effects are included only through the bandwidth renormalization, characterized by the ratio of Sommerfeld coefficient between experiment and theory, $\gamma_{exp}/\gamma_{theory}$.

Another piece of evidence supporting the intrinsic contribution is the scaling relation between the anomalous Hall conductivity σ_{xy} and longitudinal conductivity σ_{xx} [69]. According to the scaling relation, σ_{xy} is almost constant and reaches $\sim e^2/(ha) \sim 10^3 \Omega^{-1}cm^{-1}$ in the good metal regime $\sigma_{xx} \sim 3 \times 10^3 \Omega^{-1}cm^{-1} - 5 \times 10^5 \Omega^{-1}cm^{-1}$ [69]. As shown in Figs. 5 (a) and (b), the topological magnets exhibit $|\sigma_{ij}| \sim e^2/(ha)$ and satisfy these conditions.

(2) Relation between ANE and magnetization.

As mentioned above, the intrinsic σ_{xy} and α_{xy} originate from the k -space Berry curvature and thus they do not necessarily scale with the saturation magnetization. However, this is typically observed only at low T (where M is nearly saturated) and at high B (single-domain state). In other words, at high T and low B , ANE and AHE are usually related to the magnetization.

(i) Temperature dependence (Fig. 7 (a))

The AHE and ANE vanish above Curier temperature T_C (or N'eel temperature T_N) since they require spin-split band structure due to the time-reversal symmetry breaking. For instance, in Weyl magnets, Weyl points with opposite chirality annihilate into a four-fold degenerate Dirac point above $T_{C,N}$. On cooling below $T_{C,N}$, local magnetic moments $\langle m \rangle$ develop and induce spin splitting in the band structure. The size of $\langle m \rangle$ ($\langle m \rangle \propto M$ for ferromagnets) is usually a good indicator for the magnitude of spin-splitting (and distance between Weyl points for Weyl magnets). Thus T dependence of ANE and AHE resembles that for M near $T_{C,N}$. In particular, this is evident for low $T_{C,N}$ materials such as $YbMnBi_2$, $Co_3Sn_2S_2$, and $UCo_{0.8}Ru_{0.2}Al$ (Fig. 11(b)) [25, 26, 20, 23]. Since the first-principles calculations normally fix the moment size $\langle m \rangle$, a comparison to the experiment should be performed at $T \ll T_C$, where the magnetization becomes nearly T independent.

(ii) Magnetic field dependence (Fig. 7 (b))

At high B , the Berry curvature Ω_z dominates σ_{xy} and α_{xy} , leading to the violation of the empirical scaling relation with the saturation magnetization M_s . On the other hand, at low B , the curves of $S_{xy}(B)$ and $\rho_{xy}(B)$ resemble that of $M(B)$, as magnetic domains are not fully aligned. Namely, integrating the contributions from all the magnetic domains

that have either parallel or antiparallel M to the magnetic field direction, S_{xy} and ρ_{xy} are generally proportional to M since the sign of each domain's contribution depends on its magnetization direction (Fig. 7 (b)).

The zero-field spontaneous ANE is important for thermoelectric applications. This occurs in hard magnets with large magnetocrystalline anisotropy K_u [J/m³] and small saturation magnetization M_s [A/m], i.e., larger magnetic hardness parameter κ_h than unit; $\kappa_h = \sqrt{\frac{K_u}{\mu_0 M_s^2}} > 1$ [70]. For soft magnets such as Co₂MnGa, Fe₃X ($X =$ Ga, Al) and Fe₃Sn shown below, zero-field spontaneous ANE occurs only in thin-film form under perpendicular temperature gradient to the film plane since the shape anisotropy stabilizes the in-plane magnetization.

(3) Diverging behavior of α_{xy}/T (Fig. 8)

At low T , α_{xy}/T should be constant as expected from the Mott relation (Eq. (22)), and finally $\alpha_{xy} = 0$ at $T = 0$. On the other hand, various topological magnets show $\alpha_{xy}/T \sim -\ln T$ at high T (Fig. 8 (a)). This logarithmic enhancement arises from the anomalous enhancement of $\sigma_{xy}(\epsilon)$ and $\alpha_{xy}(\epsilon)$ due to the topological flat band structure (Sec 4 & Fig. 3), which is also reproduced by the first-principles calculations for Co₂MnGa and Fe₃Ga (Fig. 8 (b)) [18, 21, 66]. As the topological flat band approaches the Fermi energy E_F , α_{xy} shows $-\ln T$ divergence to lower temperatures (Fig. 8 (b)).

Mn_3X ($X = Sn, Ge$)

The realization of the spontaneous AHE without net magnetization has been a great challenge in condensed matter physics. It was first discovered in the chiral spin liquid state of Pr₂Ir₂O₇ at low temperatures [51] and later found in the inverse-triangular spin state of Mn₃Sn at room temperature [15]. Mn₃X ($X = Sn, Ge$) has a hexagonal Ni₃Sn-type ($D0_{19}$) crystal structure (space group $P6_3/mmc$), where Mn atoms form breathing-type kagome layers in the ab plane [71]. Mn₃X exhibits inverse-triangular non-collinear antiferromagnetic ordering at $T_N \sim 430$ K ($X = Sn$) and 372 K ($X = Ge$) (Table 1) with a vanishingly small net spontaneous magnetization of several $m\mu_B$ in ab plane [72, 73, 74, 75]. Nevertheless, Mn₃X shows a large spontaneous Hall effect [15, 76, 77], Nernst effect [16, 17, 22] and magneto-optical Kerr effect [78, 22, 79] comparable to those for ferromagnets. Besides, Mn₃X also exhibits spintronic phenomena such as the magnetic spin Hall effect [80], electric switching of AFM domains [81, 82, 83] and tunneling magnetoresistance [84]. These interesting properties are intimately linked to the magnetic structure and k -space topology: (1) The magnetic point group symmetry $m'm'm$ for Mn₃Sn is compatible with ferromagnetism, which allows the finite spontaneous AHE and ANE. This situation can be naturally interpreted by considering the magnetic structure as a cluster multipole [63, 85]. (2) The ARPES measurements on Mn₃Sn reveal Weyl points around K point near E_F , consistent with the first-principles calculations [86]. These Weyl points act as sources or sinks of the Berry curvature, enhancing the AHE and ANE.

Mn₃X has been intensively studied for developing antiferromagnetic spintronics devices such as low-power and ultrafast nonvolatile memory [87, 88, 89]. For thermoelectric applications, the absence of stray fields and shape anisotropy in randomly-oriented polycrystalline films has proven better for designing a highly dense

array of the spintronic thermopiles with high heat flux sensitivity, compared to the conventional sensors based on ferromagnets [31].

Co₂MnGa, Co₂MnAl_{1-x}Si_x

The above-mentioned ANE for Mn₃Sn is surprisingly large for AFM and comparable to that in ferromagnets. However, its magnitude is still small $|S_{xy}| \sim 0.5 \mu\text{V/K}$ at room T . A much large ANE $|S_{xy}| \sim 6\text{-}8 \mu\text{V/K}$, over an order of magnitude larger than in conventional magnets, was first reported in the Weyl ferromagnet Co₂MnGa [18, 90, 91, 92]. This remains the largest room- T value reported to date (Fig. 6(a)). The crystal

structure of Co₂MnGa is the $L2_1$ ordered cubic full Heusler (space group $Fm\bar{3}m$) and the Curie temperature is $T_C \sim 694$ K (Table 1) [93]. The Hall conductivity σ_{xy} gradually increases on cooling and reaches $\sigma_{xy} \sim 2000 \Omega^{-1}\text{cm}^{-1}$ (Fig. 6(b)), consistent with the first-principles calculations. The transverse thermoelectric conductivity α_{xy} for Co₂MnGa first increases on cooling, peaks at ~ 150 K, and finally vanishes at low T (Fig. 6(c)), again consistent with the first-principles calculations after considering the renormalization factor due to the correlation effect. The first-principles calculations attribute the large transverse responses to the enhanced Berry curvature near the tilted Weyl cones (Fig. 9), which ARPES measurements confirm later [94, 95, 96, 97]. The tilted Weyl cone in Co₂MnGa is interpreted as a Lifshitz transition between type-I and type-II Weyl semi-metallic states, where one of the bands becomes nearly flat at $E \sim E_F$. This topological flat band is the key for the large α_{xy} in Co₂MnGa. The large transverse responses for Co₂MnGa are also promising for thermoelectric and spintronics device applications [98, 99, 32].

To date, most thermoelectric applications of the ANE focus on heat flux sensors utilizing the large voltage in the thin-film thermopile structure. Since ANE materials are typically metallic with relatively high thermal conductivity, they are suitable for efficient heat flux sensing. However, power generation from waste heat remains a significant challenge. A problem is the low conversion efficiency $\eta_{ANE} = \eta_C \sqrt{1 - Z_{yx} T_{av}}$ due to $1 + 1 - Z_{yx} T_{av}$

to the small figure of merit $Z_{yx} T = \sigma_{xx} S_{yx}^2 T / \kappa_{yy}$. For example, even for Co₂MnGa, the ANE material with the largest S_{yx} at room T , $Z_{yx} T \sim 0.0001$ [18] partly due to the high thermal conductivity. Here, $T_{av} = (T_h + T_c)/2$ and $\eta_C = (T_h - T_c)/T_h$ are the average T between high T_h and low T_c temperatures, and Carnot efficiency, respectively [100]. We note that $\eta_{ANE} \rightarrow \eta_C$ at $Z_{yx} T \rightarrow 1$, which is different from the

Seebeck effect case, where $\eta_{SE} = \eta_C \frac{1 + Z T_{av} - 1}{\sqrt{1 + Z T_{av} + T_c/T_h}} \rightarrow \eta_C$ at $Z T = \sigma_{xx} S_{xx}^2 T / \kappa_{xx} \rightarrow \infty$.

When considering applications, material cost is also an important factor. In the case of Co₂MnGa, Ga is rather expensive compared to Co and Mn. Although the ANE $S_{xy} \sim 1 \mu\text{V/K}$ for pure Co₂MnAl is not so large as $6 \sim 8 \mu\text{V/K}$ in Co₂MnGa, the Si-doping enhances the ANE upto $S_{xy} \sim 5.7 \mu\text{V/K}$ in Co₂MnAl_{0.63}Si_{0.37} [101]. Since both Al and Si are low-cost and naturally abundant elements, the system is promising for industrial applications.

Fe₃X (X=Ga, Al)

The discovery of the large ANE in Co₂MnGa has stimulated the study of thermoelectric applications based on ANE. In addition to the large ANE, low material cost is an important factor for such applications. Iron is abundant and inexpensive, but the ANE for body-centered cubic (bcc) iron (α -Fe) is very small ($|S_{xy}| \sim 0.3 \mu\text{V/K}$). With the help of high-throughput first-principles calculations, a large ANE has been identified in Fe₃X ($X = \text{Ga, Al}$) [21]. Fe₃X ($X = \text{Ga, Al}$) crystallizes in a bcc $D0_3$ structure and shows ferromagnetic order below $T_c \sim 720 \text{ K}$ (Fe₃Ga) and $\sim 600 \text{ K}$ (Fe₃Al) [102, 103]. As shown in Fig. 6(a), Fe₃X exhibits a large ANE $-S_{yx} \sim 5.7 \mu\text{V/K}$ (Fe₃Ga) and $\sim 4.0 \mu\text{V/K}$ (Fe₃Al) at room temperature [21], which is more than an order of magnitude larger than that for bcc iron [104]. The large spontaneous ANE is also confirmed in epitaxial thin films (Fig. 10 (a)). Besides, the room- T α_{xy} is also very large $\alpha_{xy} \sim 5 \text{ (Am}^{-1}\text{K}^{-1}\text{)}$. The first-principles calculations reveal that a nodal web structure – multiple interconnected nodal lines – near the L point gives rise to large Berry curvatures (Fig. 10(b) and (c)) [21, 66]. The large ANE in Fe₃X is robust against chemical doping as found in Fe_xGa_{4-x} ($3.07 < x < 3.12$) and Fe₃Ga_{1-x}Al_x ($0 \leq x \leq 0.6$) polycrystals [105, 106], as well as in epitaxial Fe-Ga thin films without post-annealing [107], which are all found beneficial for thermoelectric applications. For example, flexible heat flux sensors have been fabricated with mass-producible roll-to-roll sputtering method (Fig. 10(d) and (e)) [34]. Ribbon-shaped Fe₃Al alloys have also been produced, which can be directly wrapped around heat pipes (Fig. 10(f)) [35]. Fe₃X is also promising for spintronics device applications due to the low Gilbert damping constant [108, 109].

YbMnBi₂

Topological antiferromagnets are attractive even for thermoelectric applications in electronic devices owing to their large responses and small stray field. The noncollinear antiferromagnet YbMnBi₂ provides a large ANE $-S_{zy} \sim 10 \mu\text{V/K}$ at $T \sim 200 \text{ K}$ (Fig. 6(a)) [25, 26]. YbMnBi₂ has a primitive tetragonal lattice (nonsymmorphic space group $P4/nmm$) and shows nearly collinear (C-type) AFM along c axis with a small cant along ab -plane below $T_N \sim 280 \text{ K}$ [110, 111, 112, 25, 26]. The electronic band structure for YbMnBi₂ is highly anisotropic, resulting in highly anisotropic transport properties such as $\rho_{zz} > \rho_{yy}$, $S_{yy} < 0 < S_{zz}$ and $S_{yz} = -S_{zx}$. On the other hand, samples are highly sensitive to oxygen and easily change their properties in the air [26]. For instance, the violation of the Onsager relation for σ_{ji} reported in [25] may be caused by the sample degradation [26]. Thus, industrial applications using this material are currently not straightforward.

Fe₃Sn

As discussed in Sec. 4, topological flat bands are key for enhancing the transverse thermoelectric coefficient. Kagome metals are good candidates since s -electrons on the kagome lattice are known to provide a flat band owing to the destructive interference [113, 114]. The kagome ferromagnet Fe₃Sn shares the same crystal structure as Mn₃X ($X = \text{Sn, Ge}$), hexagonal Ni₃Sn type ($D0_{19}$, space group $P6_3/mmc$), where Fe atoms form

a kagome lattice in ab plane (Table 1) [115]. The Curie temperature for Fe_3Sn is $T_C \sim 760$ K, which is high among topological magnets (Table 1). The large ANE $S_{ij} \sim 3 \mu\text{V}/\text{K}$ at room T in Fe_3Sn was first reported in bulk polycrystals [24] and later in amorphous and epitaxial thin films [116, 117]. As shown in Fig. 6(a), S_{ij} for polycrystalline Fe_3Sn increases linearly at $100 \text{ K} \leq T \leq 400 \text{ K}$. The first-principles calculations reveal that the large ANE for Fe_3Sn originates from a sharp peak in σ_{yz} at $E - E_F \sim 50 \text{ meV}$, which comes from a nodal plane –nearly degenerate, partially flat bands– near the Brillouin zone boundary [24]. These partially flat bands might originate from the destructive interference in the kagome lattice. The topological ANE found in the thin film based on the amorphous form of Fe_3Sn makes this material attractive for the thermoelectric applications.

Co₃Sn₂S₂

A Co-based shandite $\text{Co}_3\text{Sn}_2\text{S}_2$ is another kagome ferromagnet with $T_C \sim 177$ K. It crystallizes in a rhombohedral structure (space group $R\bar{3}m$). The magnetic Weyl semimetal phase in $\text{Co}_3\text{Sn}_2\text{S}_2$ is confirmed by various techniques, including ARPES (Fermi-arcs and linear bulk band crossings), STM (Scanning Tunneling Microscope) (quasi-particle interference), and magnetoresistance (chiral anomaly) [118, 19]. Owing to the strong magnetocrystalline anisotropy and relatively small saturation magnetization $M_s \sim 0.8 \mu_B/\text{f.u.}$, the ANE shows clear hysteresis with the sizeable spontaneous value $S_{xy} \sim 3 \mu\text{V}/\text{K}$ even in bulk form when B is applied along the easy axis $B \parallel c$ (Fig. 11) [20]. While the Curie temperature is lower than room T , these properties are attractive for thermoelectric applications at low T .

UCo_{0.8}Ru_{0.2}Al

Another approach for enhancing the DOS at the Fermi energy is via strong electronic correlations, especially in f -electron systems. The U-based ferromagnet $\text{UCo}_{0.8}\text{Ru}_{0.2}\text{Al}$ shows the largest ANE $S_{xy} \sim 23 \mu\text{V}/\text{K}$ (Figs. 6(a) and 11(b)) and the transverse thermoelectric conductivity $\alpha_{ij} \sim 14 \text{ Am}^{-1}\text{K}^{-1}$ at $T \sim 40 \text{ K}$ (Fig. 6(c)) [23]. The crystal structure is noncentrosymmetric hexagonal ZrNiAl-type (space group $P\bar{6}2m$) and U atoms form the distorted Kagome lattice in the ab plane (Table 1). The Sommerfeld coefficient for $\text{UCo}_{0.8}\text{Ru}_{0.2}\text{Al}$ is $\gamma \sim 50 \text{ mJ}/(\text{mol K}^2)$, indicating moderately heavy charge carriers probably due to electronic correlations. The large σ_{xy} and α_{xy} for $\text{UCo}_{0.8}\text{Ru}_{0.2}\text{Al}$ are reproduced by the first-principles calculations, indicating the large Berry curvature due to topological band structures. Indeed, the first-principles calculations find a large number of Weyl points (148 within $E - E_F \sim \pm 60 \text{ meV}$), which may contribute to enhancing the Berry curvature and α_{xy} . Although this material is not suitable for applications due to the uranium and low Curie temperature $T_C \sim 56 \text{ K}$ [119], this work provides valuable insight into further enhancement of the ANE.

7. Summary and Perspective

Recent advances in topological condensed matter physics have led to the discovery of large ANEs in a variety of topological magnets, opening the way for their application at room temperature. Since the direction of the ANE-induced electromotive force is controlled by magnetization, even polycrystalline samples can generate strong signals [24, 105, 106], which is beneficial for industrial applications. ANE is better suited for flexible thin-wire circuit architectures, compared to the conventional Seebeck effect. Various ANE-based thermoelectric devices including high-sensitivity thin-film heat flow sensors have been fabricated, and some are fabricated by the mass-producible roll-to-roll sputtering method [34] and ready to be available commercially.

On the other hand, there is still room for improving their functionalities, in particular, for power generation. As discussed in Sec. 6, the magnitude of ANE is still much smaller than the Seebeck effect, resulting in a small figure of merit $Z_{xy}T \sim 0.001$ [18]. In addition to the band engineering relying on the intrinsic AHE, the extrinsic contributions such as magnon-driven ANE (e.g., MnBi [58]) would provide a new route for enhancing ANE. Another important aspect is the zero-field spontaneous ANE in bulk form. As already known in the field of permanent magnets [70], rare-earth magnets with large magnetocrystalline anisotropy would be suitable, though they are costlier. In addition to these functionalities, high Curie temperature, cost-effectiveness, and figure of merit in polycrystalline samples are important factors, while satisfying all the conditions would be rather challenging. Nevertheless, we believe that the innovative work reviewed in this article will inspire future research and uncover novel physical properties and functionalities in magnetic materials to improve the ANE.

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Figures

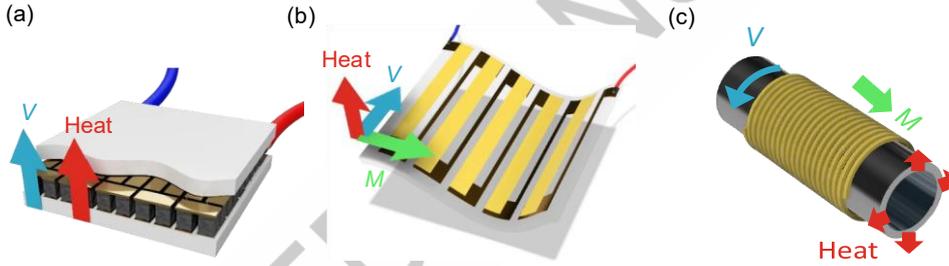
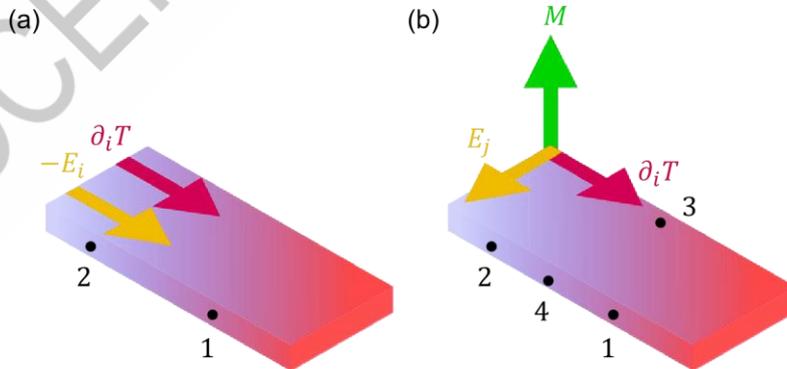


Figure 1. Schematic figures of thermoelectric devices based on the longitudinal (a) and transverse (b, c) thermoelectric effects (Adapted from [21]). In the Seebeck-effect-based (Peltier) device, a large number of p and n-type semiconductor pillars are necessary to enhance the output voltage. In contrast, ANE-based devices can achieve increased output voltage in a thin-film form, which is suitable for covering large areas and curved heat sources.



$$S_{ii} = \frac{E_i}{-\partial_i T} = \frac{(V_1 - V_2)/\ell}{(T_1 - T_2)/\ell} = \frac{\Delta V_{1-2}}{\Delta T_{1-2}}$$

$$S_{ji} = \frac{E_j}{-\partial_i T} = \frac{(V_3 - V_4)/\ell_V}{(T_1 - T_2)/\ell_T} = \frac{\ell_T \Delta V_{3-4}}{\ell_V \Delta T_{1-2}}$$

Figure 2. Schematic figures illustrating the Seebeck effect (a) and ANE (b). For the Seebeck coefficient S_{ii} , the sample length ℓ appears in both numerator and denominator and cancels out. As a result, S_{ii} is determined solely by the ratio of voltage to temperature difference $\Delta V_{1-2}/\Delta T_{1-2}$. In contrast, the sample dimension remains for the anomalous Nernst coefficient S_{ij} as $S_{ij} = (\ell_T/\ell_V)(\Delta V_{3-4}/\Delta T_{1-2})$, where $\ell_T = \ell_{1-2}$ and $\ell_V = \ell_{3-4}$ respectively represent length between thermometers along the heat current direction and length between transverse voltage terminals. This means that the ANE voltage ΔV_{3-4} increases by reducing ℓ_T and increasing ℓ_V , as realized in thin-film ANE-based thermoelectric modules (see Fig. 1(b) and (c)).

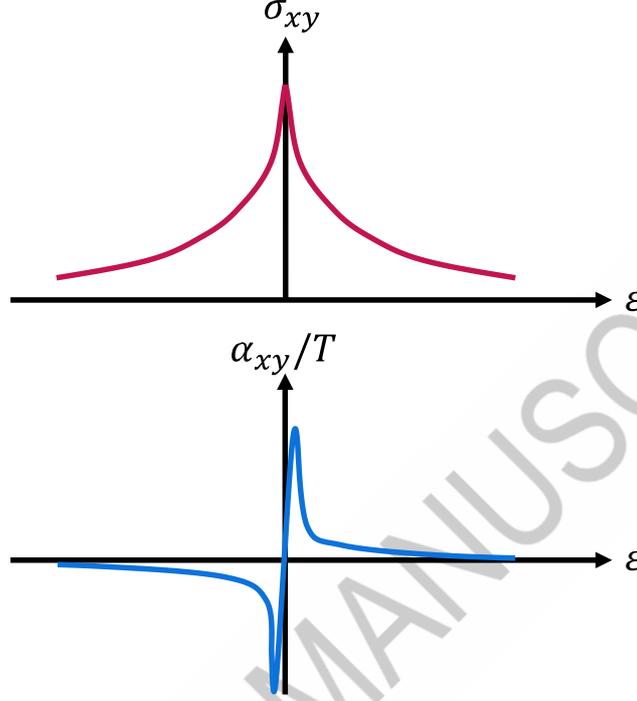


Figure 3. Schematic figures for the energy dependence of σ_{xy} (top) and α_{xy} (bottom) at low temperatures, where the Mott relation (Eq. (22)) holds [65]. Here, we assume topological band structure (e.g., Weyl points and nodal lines) inducing a large Berry curvature Ω_z at Fermi energy $\epsilon \sim 0$.

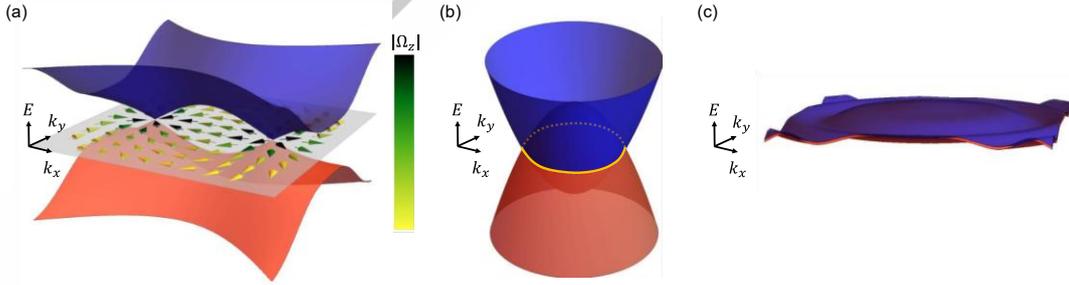


Figure 4. Schematic figures for topological band structures; (a) nodal points (Weyl points), (b) a nodal line and (c) a nodal plane. The arrows in (a) show the directions and magnitudes of the Berry curvature.

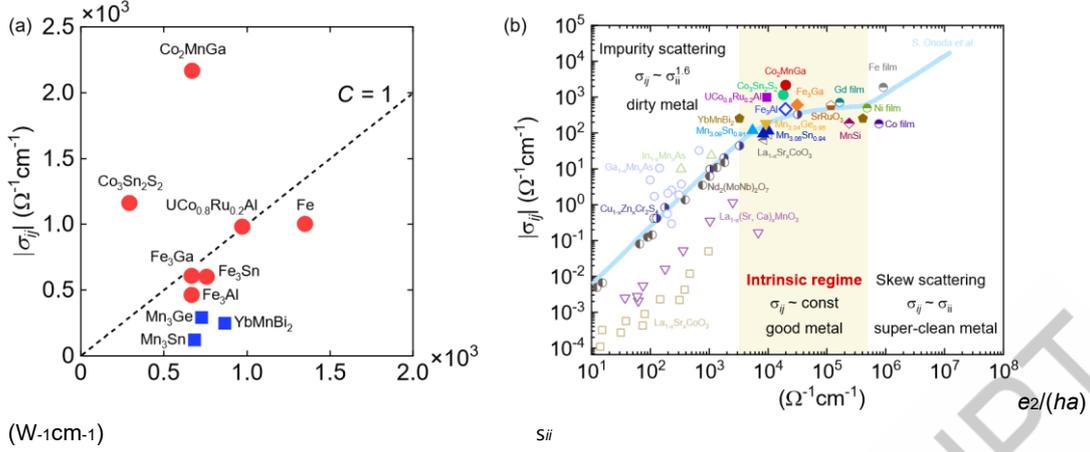


Figure 5. (a) The magnitude of the anomalous Hall conductivity $|\sigma_{ij}|$ compared to $e^2/(ha)$, where a is the lattice constant along the applied field direction; $a = b = c = 5.77$ Å for Co_2MnGa , 5.80 Å for Fe_3Ga , 5.79 Å for Fe_3Al , 2.89 Å for Fe , $a = b = 5.66$ Å for Mn_3Sn , $a = b = 5.33$ Å for Mn_3Ge , $a = b = 4.47$ Å for YbMnBi_2 , $c = 13.18$ Å for $\text{Co}_3\text{Sn}_2\text{S}_2$, $c = 3.97$ Å for $\text{UCo}_{0.8}\text{Ru}_{0.2}\text{Al}$. For polycrystalline Fe_3Sn , we used the averaged lattice constant $(a + b + c)/3 \sim 5.10$ Å. (b) The scaling relation between $|\sigma_{ij}|$ and the longitudinal conductivity σ_{ii} . Here, data taken at the lowest temperature ($T \sim 2$ K) are used except for Mn_3Sn ($T \sim 100$ K). The references include Refs. [15, 16, 22] for Mn_3Sn and Mn_3Ge , Ref. [18] for Co_2MnGa , Ref. [21] for Fe_3Ga and Fe_3Al , Ref. [26] for YbMnBi_2 , Ref. [24] for Fe_3Sn , Refs. [19, 20] for $\text{Co}_3\text{Sn}_2\text{S}_2$, Ref. [23] for $\text{UCo}_{0.8}\text{Ru}_{0.2}\text{Al}$, Ref. [120] for Fe , Co , Ni and Gd thin films, Ref. [121] for MnSi , Ref. [69] for perovskite oxides (SrRuO_3 , $\text{La}_{1-x}\text{Sr}_x\text{CoO}_3$, $\text{La}_{1-x}(\text{Sr,Ca})_x\text{MnO}_3$), spinels ($\text{Cu}_{1-x}\text{Zn}_x\text{Cr}_2\text{Se}_4$), pyrochlore ($\text{Nd}_2(\text{MoNb})_2\text{O}_7$) and magnetic semiconductor ($\text{Ga}_{1-x}\text{Mn}_x\text{As}$, $\text{In}_{1-x}\text{Mn}_x\text{As}$).

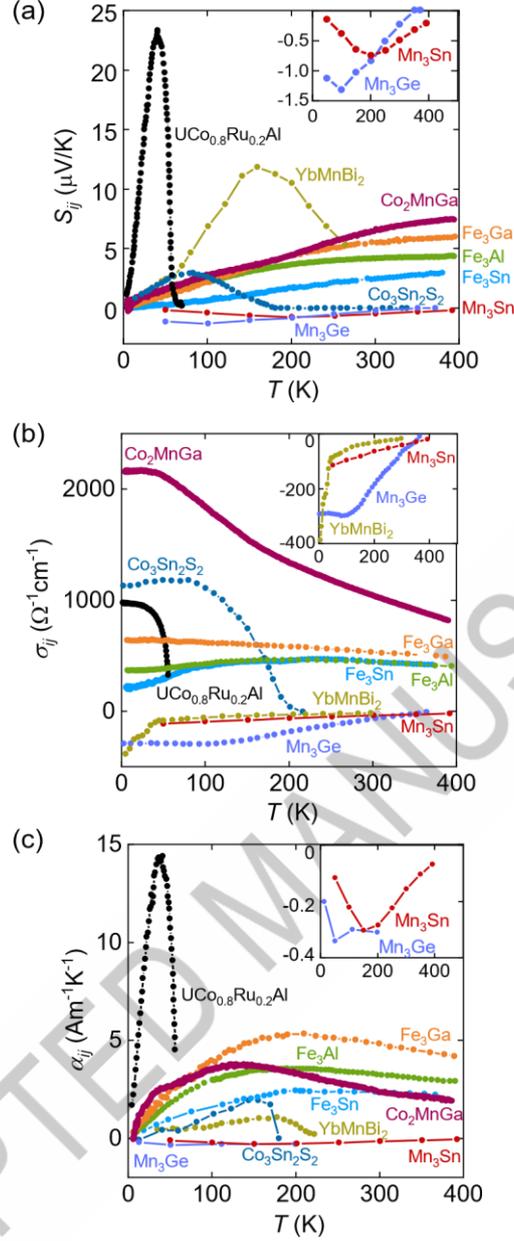


Figure 6. Temperature dependence of the anomalous Nernst coefficient S_{ij} (a), anomalous Hall conductivity σ_{ij} (b) and transverse thermoelectric conductivity α_{ij} (c) for various topological magnets. The measurement conditions, including the directions of electric current (\mathbf{I}) and heat current (\mathbf{Q}), are as follows: Mn_3Sn

($\text{Mn}_{3.06}\text{Sn}_{0.94}$, $ij = zx$, $\mathbf{B} \parallel [01\bar{1}0]$, $\mathbf{I}, \mathbf{Q} \parallel [2\bar{1}\bar{1}0]$) [16], Mn_3Ge ($\text{Mn}_{3.03}\text{Ge}_{0.97}$, $ij = zx$, $\mathbf{B} \parallel [01\bar{1}0]$, $\mathbf{I}, \mathbf{Q} \parallel [2\bar{1}\bar{1}0]$) [22], Co_2MnGa ($ij = xy$, $\mathbf{B} \parallel [100]$, $\mathbf{I}, \mathbf{Q} \parallel [001]$) [18], Fe_3X ($X = \text{Ga}, \text{Al}$) ($ij = xy$, $\mathbf{B} \parallel [100]$, $\mathbf{I}, \mathbf{Q} \parallel [001]$) [21], polycrystalline Fe_3Sn [24], $\text{Co}_3\text{Sn}_2\text{S}_2$ ($\mathbf{B} \parallel [0001]$, $\mathbf{I}, \mathbf{Q} \parallel ab$ plane) [19, 20], $\text{UCo}_{0.8}\text{Ru}_{0.2}\text{Al}$ ($\mathbf{B} \parallel c$, $\mathbf{I}, \mathbf{Q} \parallel a$) [23] and YbMnBi_2 ($-S_{xy}$, $\mathbf{B} \parallel x, \mathbf{I}, \mathbf{Q} \parallel z$) [26]. Insets show zoomed-in region where the y axis is negative.

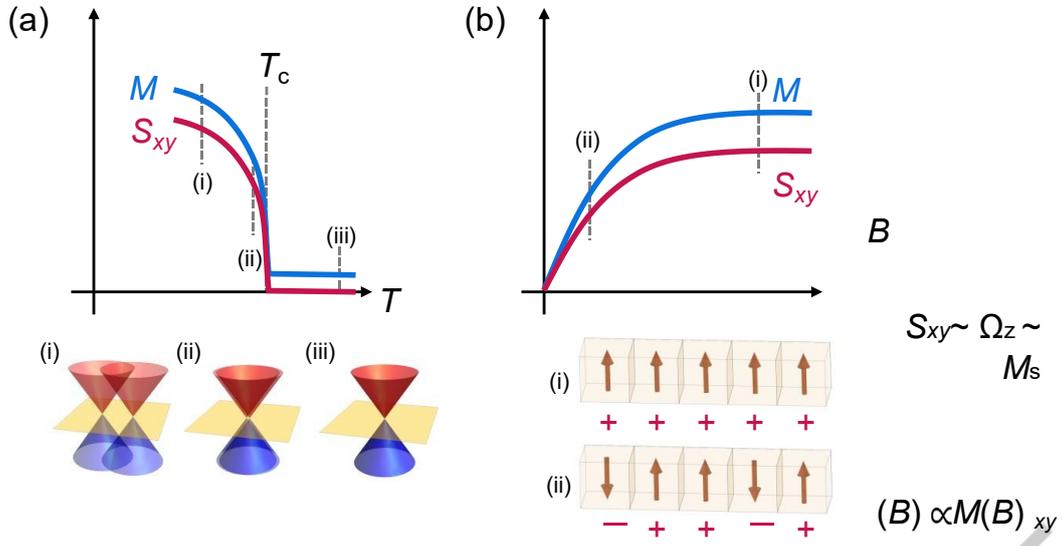


Figure 7. (a) Schematic figures for the T dependences of M and S_{ij} near the Curie temperature T_c (top), and Weyl (or Dirac) points for (i)-(iii). (b) Schematic figures for the B dependences of M and S_{xy} (top), magnetic domains (boxes & red vectors) and their contributions to S_{xy} (signs below boxes). The total M and S_{xy} are obtained by summing over each domain contribution (red vectors and signs, respectively). At high field (e.g., state at (i)), all the magnetic moments are parallel to the applied external field, and thus total S_{xy} reflects the Berry curvature Ω_z , leading to the violation of the empirical scaling relation between the *magnitude* of S_{xy} and saturation magnetization M_s . On the other hand, the *shape* of the B sweep curves are similar to each other; $S_{xy}(B) \propto M(B)$. This is because both the total $S_{xy}(B)$ and $M(B)$ are determined by integrating all the magnetic-domain contributions at low fields (e.g., the state at (ii)). Here, we assume the Ising anisotropy and B applied to the easy axis.

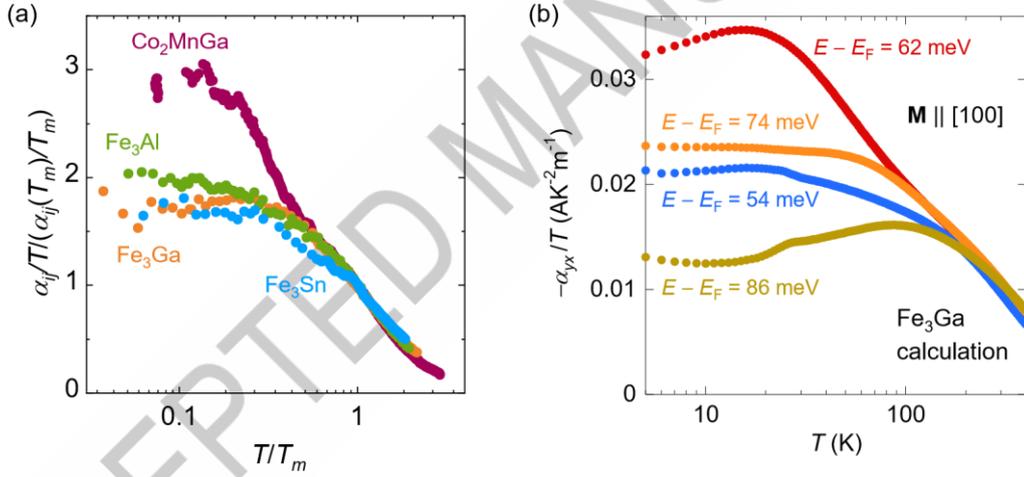


Figure 8. (a) The scaling relation of the transverse thermoelectric coefficient, $\alpha_{ij}/T / (\alpha_{ij}(T_m)/T_m)$ vs T/T_m , for Co_2MnGa [18], Fe_3X ($X = \text{Ga}, \text{Al}$) [21] and Fe_3Sn [24]. Here, T_m is the temperature where α_{ij} shows maximum. (b) T dependence of α_{xy}/T at various energy $E - E_F$ obtained from the first-principles calculation for Fe_3Ga . As the energy associated with the topological flat band approaches the Fermi energy, α_{xy} shows $-\ln T$ divergence down to much lower temperatures.

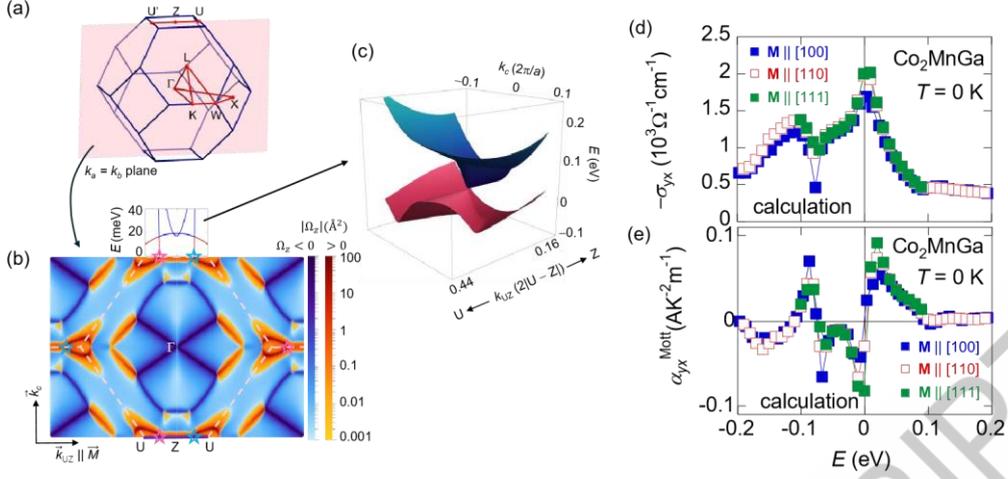


Figure 9. Topological band structure and large transverse responses obtained from the first-principles calculations for Co_2MnGa (Adapted from [18]). (a) First Brillouin zone with $k_a = k_b$ plane highlighted (pink). (b) Berry curvature on $k_a = k_b$ plane. (c) Tilted Weyl cone near the Z-U line. (d, e) Calculated anomalous Hall conductivity $-\sigma_{yx}$ (d) and transverse thermoelectric conductivity α_{yx} (e). Here α_{yx} is obtained from the Mott relation (Eq. 22).

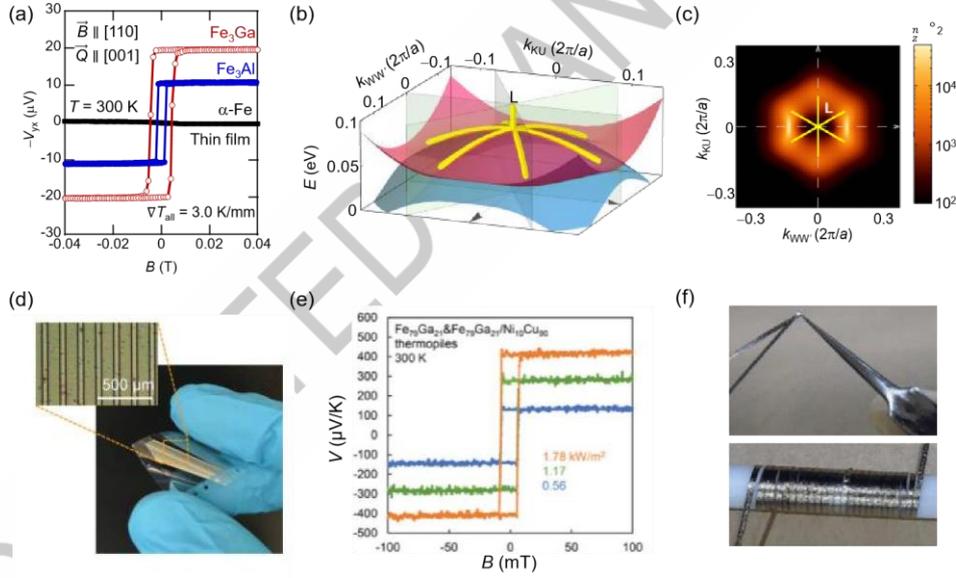


Figure 10. (a) Anomalous Nernst voltages for Fe_3Ga and Fe_3Al epitaxial thin films compared to that for $\alpha\text{-Fe}$. Heat current was applied perpendicular to the film $\mathbf{Q} \parallel [001]$ and with in-plane magnetic field, $\mathbf{B} \parallel [001]$. (b), (c) Topological band structure (nodal web) (b) and Berry curvature (c) near the L point obtained from the first-principles calculations for Fe_3X . ((a)-(c) Adapted from [21]) (d) Flexible Fe_3Ga -based heat flux sensor fabricated on PET (polyethylene terephthalate) substrate. (e) B dependence of the ANE voltage for the Fe_3Ga -based flexible heat flux sensor. Here, in-plane Seebeck effect is compensated by using $\text{Ni}_{10}\text{Cu}_{90}$ as electrodes since $S_{ii,\text{Co}_2\text{MnGa}} \sim -S_{ii,\text{Ni}_{10}\text{Cu}_{90}}$. ((d), (e) Adapted from [34]). (d) The ribbon-shaped Fe_3Al (Adapted from [35]).

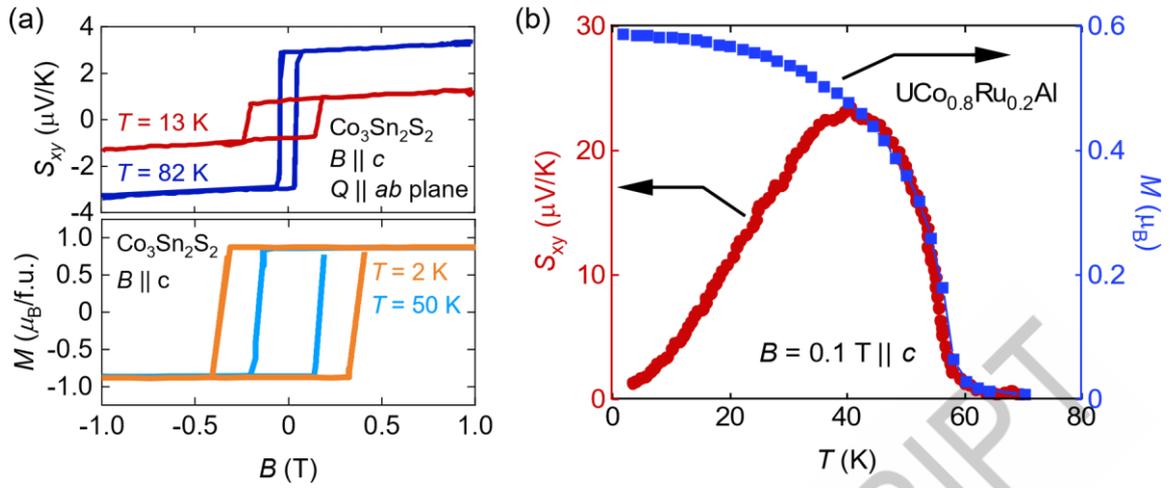
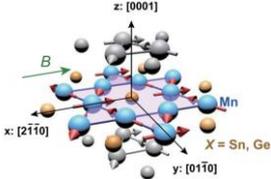
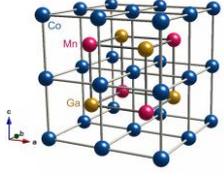
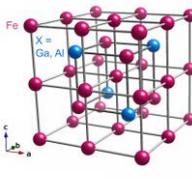
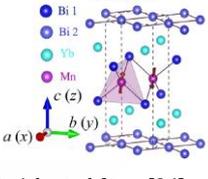
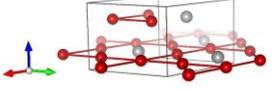
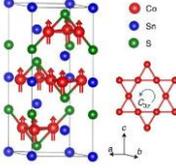
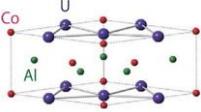


Figure 11. (a) B dependence of S_{xy} (top) and M (bottom) for $\text{Co}_3\text{Sn}_2\text{S}_2$. The data are taken from Ref. [20]. The difference in coercivity between S_{xy} and M can be attributed to the difference in T and sample shape. (b) T dependence of S_{xy} (left axis) and M (right axis) for $\text{UCo}_{0.8}\text{Ru}_{0.2}\text{Al}$. The data are taken from Ref. [23].

Table 1. Crystal structure and spin texture, type of magnetic order (FM or AFM), magnetic ordering temperature (T_C or T_N), bandstructure topology, S_{xy} at room temperature and related references for various topological magnets showing the large ANE.

Material	Crystal structure and spin texture	FM or AFM	T_C or T_N (K)	Band structure topology	S_{xy} at room T ($\mu\text{V}/\text{K}$)	Ref.
Mn_3X ($X = \text{Sn}, \text{Ge}$)	Hexagonal Ni_3Sn -type ($D0_{19}$)  Adapted from [22]	AFM	430 ($X = \text{Sn}$), 372 ($X = \text{Ge}$)	Weyl points	~ -0.5	ANE [16, 17, 22] AHE [15] Device [31] Review [27, 22, 28, 7]
Co_2MnGa , $\text{Co}_2\text{MnAl}_{1-x}\text{Si}_x$	Cubic full Heusler ($L2_1$) 	FM	694	Weyl points	~ 8	ANE & AHE [18, 90, 91, 101, 92], Device [32], Review [7]
Fe_3X ($X = \text{Ga}, \text{Al}$)	BCC derivative structure ($D0_3$) 	FM	720 ($X = \text{Al}$), 600 ($X = \text{Ga}$)	Nodal lines	~ 4 ($X = \text{Al}$), ~ 6 ($X = \text{Ga}$)	ANE [21, 107, 105, 106], AHE, Device [30, 34, 35], Review [7]
YbMnBi_2	tetragonal Canted AFM  Adapted from [26]	AFM	~ 290	Weyl points	0 (~ 10 $\mu\text{V}/\text{K}$ at $T \sim 200$ K)	AHE & ANE [25, 26]
Fe_3Sn	hexagonal Ni_3Sn -type ($D0_{19}$)  Adapted from [24]	FM	760	Nodal plane	~ 3	AHE & ANE [24, 116, 117]
$\text{Co}_3\text{Sn}_2\text{S}_2$	rhomboidal  Adapted from [19]	FM	177	Weyl points	0 (~ 3 $\mu\text{V}/\text{K}$ at $T \sim 75$ K)	ANE [20], AHE [19]

$UCo_{0.8}Ru_{0.2}Al$	<p>hexagonal ZrNiAl-type</p>  <p>Adapted from [23]</p>	<p>FM</p> <p>27</p>	<p>56</p>	<p>Weyl points</p>	<p>0 (~ 23 $\mu V/K$ at $T \sim 40$ K)</p>	<p>AHE [119] ANE [23]</p>
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ACCEPTED MANUSCRIPT



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We review recently developed transverse thermoelectric generation technology using topological magnets and cover its novel mechanism and applications.

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