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# **Quantum metric-induced nonlinear transport in a topological antiferromagnet**

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# **Title: Quantum metric-induced nonlinear transport in a topological**

# **antiferromagnet**

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**Abstract: The Berry curvature and quantum metric are the imaginary part and real part, respectively, of the quantum geometric tensor which characterizes the topology of quantum states<sup>1</sup> . The former is known to generate a zoo of important discoveries such as quantum Hall effect and anomalous Hall effect**  $(AHE)^{2,3}$ **, while the consequences of the quantum metric have rarely been probed by transport. Here we report the observation of quantum metric-induced nonlinear transport, including both nonlinear AHE and diode-like nonreciprocal longitudinal response, in thin films of a topological antiferromagnet, MnBi2Te4. Our observation reveals that the transverse and longitudinal nonlinear conductivities reverse signs when reversing the antiferromagnetic order, diminish above the Néel temperature, and are insensitive to disorder scattering, thus verifying their origin in the band structure topology. They also flip signs between electron and hole-doped regions, in agreement with theoretical calculations. Our work provides a pathway to probe the quantum metric through nonlinear transport and to design magnetic nonlinear devices.** Fractional Fraction Compares the topology<br>
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#### **Main text:**

Nonlinear transport provides a powerful tool for probing topological 39 . physics in solids<sup>4-21</sup>. A prime example of this is the nonlinear anomalous Hall effect (NLAHE) which reveals the Berry curvature dipole on the Fermi surface and has recently been observed in nonmagnetic topological 42 semimetals (as shown in Fig. 1a) $6,9,10,12,14,15$ . This provides a powerful means of investigating quantum geometry in flat space. Another NLAHE caused by the quantum metric, the counterpart of the Berry curvature in the quantum geometry, was also predicted for magnetic topological materials, 46 but is yet to be realized in experiments<sup>1,22-30</sup>. The quantum metric measures the amplitude distance between two neighboring Bloch states and 48 determines the electronic properties of a crystal. A very recent theory<sup>26,30</sup> finds that the quantum metric actually gives rise to both NLAHE and the nonreciprocal longitudinal response, the latter of which is characterized by a diode-like longitudinal resistance, in inversion-breaking magnetic 52 materials (as shown in Fig. 1b)<sup>7</sup>. The asymmetric quantum metric (quantum metric dipole) will induce an anomalous motion of the wave packet and generate nonlinear responses in both the longitudinal and transverse directions. Since the quantum metric is antisymmetric under 56 momentum inversion  $(k \rightarrow -k)$ , both time reversal symmetry (*T*) and inversion symmetry (*P*) must be broken in order to generate a net quantum metric dipole. Additionally, the combined symmetry *PT* can exclude the contribution from the Berry curvature dipole<sup>4</sup>. As a result, a system which breaks both *P* and *T* but preserves *PT* is expected to be an ideal platform for studying the quantum metric effect (see supplementary information S1 for detailed symmetry analysis). a powerful tool for probing topological<br>example of this is the nonlinear anomalous<br>reveals the Berry curvature dipole on the<br>y been observed in nonmagnetic topological<br>2.  $1a)^{69,10,12,14,15}$ . This provides a powerful<br>m

63 After careful consideration, we chose  $MnBi<sub>2</sub>Te<sub>4</sub>$  as the candidate platform to investigate the quantum metric. Compared to the magnetic doped topological insulator, the intrinsic magnetic topological insulator, MnBi<sub>2</sub>Te<sub>4</sub>, gained a lot of interest recently due to its unique properties<sup>17,31-</sup> . As shown in Fig. 1c and e, the crystal structure of MnBi<sub>2</sub>Te<sub>4</sub> consists of alternating layers of Te-Bi-Te-Mn-Te-Bi-Te, known as septuple layers  $(SLs)^{33,34}$ . MnBi<sub>2</sub>Te<sub>4</sub> has an A-type antiferromagnetic (AFM) ground state, in which the Mn spins in each SL are aligned ferromagnetically with an out-of-plane easy axis but are coupled antiparallel to adjacent SLs. For <sup>17</sup> inversion symmetry (*P*) must be browntrive inversion symmetry (*P*) must be browntrive inversion symmetry (*P*) must be browntrive contribution from the Berry curvature of the summetric of the summetry analysis).<br>
6 72 even-layer MnBi<sub>2</sub>Te<sub>4</sub>, both *P* and *T* symmetry are broken but the combined

*PT* symmetry is preserved below the Néel temperature<sup>37</sup>. Fig. 1d shows the spatial reflectance magnetic circular dichroism (RMCD) mapping of 3SL and 4SL MnBi2Te4 measured at zero magnetic field, indicating fully 76 compensated AFM states in even layer  $MnBi_2Te_4^{40}$ . In addition,  $MnBi_2Te_4$ possesses three-fold rotational symmetry (from the optical second harmonic generation shown in Fig. 1f), which suppresses the Berry curvature dipole even in the absence of *PT*-symmetry, making the 80 contribution from the quantum metric much easier to observe  $30,41$ .

#### 81 **Electron nonlinearity from spin order**

82 To investigate the electron nonlinearity, we fabricated several high-quality, 83 dual gated even-layer MnBi2Te4 devices. Here, we focus on a 4SL-84 MnBi2Te4 device, the schematic structure of which is shown in Fig. 2a. The 85 dual-gate structure allows us to independently control the carrier density 86 and vertical displacement field. As per theoretical predictions, the quantum 87 metric gives rise to both NLAHE and the nonreciprocal longitudinal 88 response<sup>30</sup>. The former is directly measurable via an alternating current 89 (AC) method, while the latter exhibits diode-like nonreciprocal resistance 90 behavior, expressed as:  $V_x = R_0(I + \gamma I^2)$  where *I* represents the 91 applied current,  $R_0$  is the linear resistance, and  $\gamma$  denotes the coefficient 92 characterizing the nonreciprocity<sup>7</sup>. Considering that the nonreciprocal 93 longitudinal response stems from the quadratic term of the current, we 94 adopted the AC method to measure the second-harmonic voltage  $V_x^{2\omega}$  to 95 reflect the non-reciprocal behavior in longitudinal direction, which offers 96 a better signal-to-noise ratio (see supplementary information section S4). 97 To measure the linear and non-linear responses of the device, we employed 98 a standard lock-in technique. As depicted in Fig. 2a, an AC current with frequency of  $\omega$  ( $I^{\omega}$ ) was injected, and the linear voltage  $V^{\omega}$  and secondharmonic voltage  $V^{2\omega}$  are simultaneously probed. In our experiment, the *x*-101 axis is defined as the current direction and the *y*-axis as the transverse 102 direction to the current. All the measurements are conducted at  $T = 1.8$  K 103 in the absence of a magnetic field unless otherwise specified. Fig. 2b shows the linear longitudinal  $(V_x^{\omega})$  and transverse  $(V_y^{\omega})$  voltages, indicating good 105 ohmic contact and a negligible misalignment in the Hall bar geometry from 106 the vanishing of  $V_y^{\omega}$ . As and 4SL MnBi-Tse ineusured at zero magnetic field, indicating fully<br>
79 compensated APM states in even in Mar Minki-Te<sup>4</sup>. In addition. Minki-Te<sub>1</sub>.<br>
77 passesses there-Gold rotational symmetry (from the optical second

107 Next, we focus on the nonlinear responses of  $4SL-MnBi<sub>2</sub>Te<sub>4</sub>$  in opposite 108 AFM states. Utilizing a method akin to those reported previously for

109 controlling AFM states in CrI<sub>3</sub> or even-layer MnBi<sub>2</sub>Te<sub>4</sub>, we prepared the 110 AFM-I and AFM-II states  $37,42$  with opposite Néel orders. As shown in Fig. 111 2c and 2e, the AFM-I state is prepared by sweeping the magnetic field from  $112 - 7$  T to zero, while the AFM-II state is achieved by sweeping the magnetic 113 field from +7 T to zero. For the linear responses, both AFM states exhibited 114 identical longitudinal  $(V_x^{\omega})$  voltages, and the transverse  $(V_y^{\omega})$  voltages 115 remained at zero (Extended Data Fig. 1). This observation aligns with the 116 fully compensated AFM orders in  $4SL-MnBi<sub>2</sub>Te<sub>4</sub>$  (Extended Data Fig. 2). 117 However, the nonlinear responses differ significantly from their linear 118 counterparts. Focusing first on the AFM-I state, Fig. 2c shows that while 119 the linear transverse  $(V_y^{\omega})$  voltage remains at zero, there is a significant 120 negative nonlinear transverse voltage (notated as  $V_y^{2\omega}$ ) that scales 121 quadratically with the injected current  $I^{\omega}$ . More importantly, we also 122 detected a prominent negative nonlinear longitudinal voltage (denoted as 123  $V_x^{2\omega}$ ), with the same order of magnitude as  $V_y^{2\omega}$ . This observation aligns 124 with the prediction that the quantum metric dipole can induce both 125 nonlinear Hall and nonreciprocal longitudinal response, distinctly different 126 from the response observed in the nonlinear Hall effect caused by the Berry 127 curvature dipole, where only a Hall response is allowed and the 128 longitudinal response is absent<sup>9,10</sup>. The nonreciprocal longitudinal response 129 in MnBi2Te4 can also be observed through DC measurements (see 130 supplementary information section S4). Subsequently, we prepared the 131 AFM-II state. As shown in Fig. 2d, in sharp contrast, although the 132 amplitudes of  $V_y^{2\omega}$  and  $V_x^{2\omega}$  remain nearly the same with the AFM-I 133 state, the sign of both flips to positive. The sign reversal of the nonlinear 134 response suggests that the nonlinear response observed in  $MnBi<sub>2</sub>Te<sub>4</sub>$  is 135 associated with the AFM order. We carefully examined and ruled out 136 potential alternative origins, such as thermal effect or accidental diode 137 junction that could lead to a nonlinear effect. We also exclude the 138 possibility that the nonlinear response observed in even-layer  $MnBi<sub>2</sub>Te<sub>4</sub>$  is 139 originated from the residual magnetization (See supplementary 140 information section S2). To further confirm that the nonlinear signal is an 141 intrinsic response originating from the *PT*-symmetric AFM order, we 142 investigate whether the nonlinear response complies with the rotational 143 symmetry of AFM even-layer MnBi<sub>2</sub>Te<sub>4</sub>, where  $\sigma_{yxx} = -\sigma_{yyy}$  and <sup>129</sup> in MnBi<sub>2</sub>Te<sub>4</sub> can also be obsert<br><sup>130</sup> supplementary information section<br><sup>131</sup> AFM-II state. As shown in Fig.<br><sup>132</sup> amplitudes of  $V_y^{2\omega}$  and  $V_x^{2\omega}$  rem<br><sup>133</sup> state, the sign of both flips to posit<br><sup>134</sup> resp 144  $\sigma_{xyy} = -\sigma_{xxx}$ . When y aligns with the in-plane crystal axis, it should repared by sweeping the magnetic field from<br>I state is achieved by sweeping the magnetic<br>linear responses, both AFM states exhibited<br>voltages, and the transverse  $(V_p^{\omega})$  voltages<br>tata Fig. 1). This observation aligns wit

follow that  $\sigma_{xyy} = -\sigma_{xxx} = 0$ . The coexistence of  $\sigma_{yxx}^{2\omega}$  and  $\sigma_{xxxx}^{2\omega}$  in 146 this 4SL-MnBi2Te4 device suggests some misalignment between *x/y* and 147 the crystal axes, consistent with our optical SHG measurement in Fig. 1f. 148 To substantiate this further, we fabricated another  $4SL-MnBi<sub>2</sub>Te<sub>4</sub>$  device 149 with radially distributed electrodes aligned to the crystalline axis (dashed 150 line in Fig. 2e). Fig. 2f demonstrates the nonlinear transverse and 151 longitudinal response upon rotation of the current injection direction. Both 152 transverse  $V_y^{2\omega}$  and longitudinal  $V_x^{2\omega}$  exhibit three-fold symmetry. 153 These observed features align consistently with the symmetry of the AFM 154 even-layer  $MnBi<sub>2</sub>Te<sub>4</sub>$ .

We further investigated the temperature dependence of the nonlinear response in MnBi2Te4. Fig 3a and 3b shows the temperature dependent 157 nonlinear voltages  $V_y^{2\omega}$  and  $V_x^{2\omega}$  which are prepared at AFM-I and AFM-II states, respectively. As the temperature increases, we found that the nonlinear voltage gradually decreases and vanishes above the Néel temperature of MnBi2Te4, indicating that the nonlinear response is absent in the non-magnetic states. In addition, the nonlinear response in even-layer MnBi2Te4 is only presented at the AFM states and vanishes when all spins are aligned in one direction (see details in supplementary information section S3). In summary, we can conclude that the nonlinear response observed is associated with the nonlinearity of electron motion originating 166 from the AFM order in 4SL-MnBi<sub>2</sub>Te<sub>4</sub>. th our optical SHG measurement in Fig. 1f.<br>
we fabricated another 4SL-MnBi<sub>2</sub>Te<sub>4</sub> device<br>
rodes aligned to the crystalline axis (dashed<br>
emonstrates the nonlinear transverse and<br>
lation of the current injection direction

### 167 **Quantum metric induced nonlinear transport**

168 We now systematically investigate the microscopic origin of the nonlinear 169 response observed in 4SL-MnBi2Te4. To quantify the strength of the 170 nonlinear response, we use the nonlinear conductivity, as it is independent 171 of the sample size. In our experiment, we can extract the longitudinal and transverse nonlinear conductivity from our data as  $\sigma_{xxxx}^{2\omega} = \frac{J_x^{2\omega}}{(\epsilon \omega)}$ The transverse nonlinear conductivity from our data as  $\sigma_{xxxx}^{2\omega} = \frac{f_x}{(E_{xx}^{\omega})^2}$  $V_x^{2\omega}$  L  $\frac{V_x^{2\omega} L}{(I_x^{\omega})^2 R_{xx}^3}$  and  $\sigma_{yxx}^{2\omega} = \frac{J_y^{2\omega}}{(E_{xx}^{\omega})^2}$  $\frac{J_y^{2\omega}}{(E_{xx}^{\omega})^2} = \frac{V_y^{2\omega}}{(I_x^{\omega})^2}$  $(I_x^{\omega})^2 R_{xx}^3$ 173  $\frac{V_x^{2\omega} L}{(I_x^{\omega})^2 R_{xx}^3}$  and  $\sigma_{yxx}^{2\omega} = \frac{I_y^{2\omega}}{(E_x^{\omega})^2} = \frac{V_y^{2\omega}}{(I_x^{\omega})^2 R_{xx}^3} \frac{L^3}{W^2}$ , respectively, where *L* and *W* are 174 the longitudinal and transverse length of the Hall bar device. In theory, the 175 nonlinear conductivity due to an applied electric field  $E_x$  in a metal includes 176 three contributions<sup>30</sup>, 164 section S5). In summary, we can<br>
165 observed is associated with the non<br>
166 from the AFM order in 4SL-MnBi<sub>2</sub><br>
167 **Quantum metric induced nonline**<br>
168 We now systematically investigate<br>
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177 
$$
J_c = \sigma_{cxx} E_x^2 = (\sigma_{cxx}^2 + \sigma_{cxx}^1 + \sigma_{cxx}^0) E_x^2
$$
 (Eq. 1)

178 where  $\sigma_{cxx}^i$  has the *i*-th order of  $\tau$  dependence  $(c = x \text{ or } y)$ . Here  $\sigma_{cxx}^2 =$ 179  $-\frac{e^{3}\tau^{2}}{\hbar^{3}}\int d^{2}k f_{n} \partial_{k_{c}} \partial_{k_{x}} \partial_{k_{x}} \varepsilon_{n}(\mathbf{k})$  is the nonlinear Drude weight, and  $\varepsilon_{n}(\mathbf{k})$ 180 is the energy of Bloch state  $n$  at momentum  $k$ , and  $f_n$  is the Fermi Dirac 181 distribution for band *n*.  $\sigma_{cxx}^1 = -\frac{e^{3\tau}}{\hbar^2} \int d^2k f_n(2\partial_{k_x}\Omega_n^{xc})$ , is the Berry 182 curvature dipole contribution<sup>4</sup>. The conductivity due to the normalized 183 dipole of the quantum metric is  $30$ ,

184 
$$
\sigma_{baa}^0 = -\frac{e^3}{\hbar^2} \int d^2k f_n (2 \partial_{k_b} G_n^{aa} - \partial_{k_a} G_n^{ab})
$$
 (Eq. 2)

Where  $a/b = x/y$ ,  $G_n^{ab} = \sum_{m \neq n} \frac{A_{nm}^a A_{mn}^b + A_{nm}^b A_{mn}^a}{(s - s)}$ 185 Where  $a/b = x/y$ ,  $G_n^{ab} = \sum_{m \neq n} \frac{A_n m A_{mn} + A_n m A_{mn}}{(\varepsilon_n - \varepsilon_m)}$  is the band-energy 186 normalized quantum metric and  $\partial_{k_a} G_n^{ab}$  represents the normalized 187 quantum metric dipole. In the specific case of 4SL-MnBi2Te4, the *PT*-188 symmetry excludes any contribution from the Berry curvature dipole<sup>4</sup>. 189 Additionally, the Berry curvature dipole contribution, which is even under 190 time-reversal, contradicts the sign reversal of the nonlinear signal between 191 the AFM-I and II phases. It is also worth noting that the skew scattering 192 mechanism can also contribute to the nonlinear response<sup>21,43,44</sup>, which is 193 significantly suppressed by  $PT$ -symmetry<sup>43,45</sup>. Furthermore, we can 194 exclude its contribution based on the scaling analysis discussed 195 subsequently. Therefore, the intrinsic contributions to the nonlinear 196 conductivity in MnBi2Te4 are determined by the nonlinear Drude weight 197 and the normalized quantum metric dipole. 176 = T<sub>2</sub>(1)  $V_{1/20}V_{1$ 

198 To verify the dominant contribution from the quantum metric, we 199 investigate the scaling relationship between nonlinear conductivities 200  $(\sigma_{yxx}^{2\omega}$  and  $\sigma_{xxxx}^{2\omega}$  ) and the linear conductivity  $\sigma_{xx}^{\omega}$  (i.e. scattering time  $\tau$ ) 201 as a function of temperature. Fig. 3c shows the temperature dependence of 202 longitudinal conductivity  $\sigma_{xx}^{\omega}$  and charge carrier density  $n_e$ . The charge 203 carrier density  $n_e$  remains almost unchanged at the temperature of 1.6 to 10 204 K, suggesting that the  $\tau$  is the main component to determine the conductivity  $\sigma_{xx}^{\omega}$  in this range. By adopting this range, we plot the 206  $\sigma_{yxx}^{2\omega}$  and  $\sigma_{xxx}^{2\omega}$  as a function of conductivity  $\sigma_{xx}^{\omega}$ ,

$$
\sigma^{2\omega} = \eta_2 (\sigma_{xx}^{\omega})^2 + \eta_0 \quad \text{(Eq. 3)}
$$

208 we find that the predominant contribution to both  $\sigma_{xxxx}^{2\omega}$  and  $\sigma_{yxx}^{2\omega}$  is the 209 intrinsic nonlinear conductivity from the normalized quantum metric

210 dipole ( $\eta_0$ ,  $\tau$ -independent term). The nonlinear Drude contribution ( $\eta_2$ , 211  $\tau^2$ -dependent term) is nearly negligible compared to the intrinsic one, 212 which is reasonable considering the small conductivity and low mobility 213 in MnBi<sub>2</sub>Te<sub>4</sub>. The dominant  $\tau$ -independent nonlinear conductivity further 214 exclude contribution from the impurity scattering, because its lowest order 215 contribution to the nonlinear conductivity begins at  $\tau^1$  (see detailed 216 discussion in supplementary information  $S7.2$ ) <sup>21,44</sup>. In conclusion, we 217 attribute the nonlinear response observed in 4SL-MnBi2Te4 to the 218 nonlinear Drude weight and the normalized quantum metric dipole, with 219 the latter being the dominant contributor.

#### 220 **Fermi energy dependent nonlinear response**

221 We next study how the nonlinear conductivity in 4SL-MnBi2Te4, 222 originating from normalized quantum metric dipole, is influenced by the 223 vertical displacement electric field (denoted as *D*) and the electron charge 224 carrier density (denoted as  $n_e$ ). We first examine the effect of *D* on the 225 nonlinear response. Although *PT* symmetry is instrumental in isolating the 226 quantum metric dipole contribution, the induced nonlinear response 227 remains even when *PT* is broken by *D*. Furthermore, we note that the 228 Berry curvature dipole induced NLAHE still vanishes because of three-229 fold rotational symmetry. To demonstrate this, we investigate the 230 dependence of  $\sigma_{yxx}^{2\omega}$  on the electric field *D* with fixed carrier density at 231  $n_e \approx -3 \times 10^{12}$  cm<sup>-2</sup>. As shown in Fig. 4b,  $\sigma_{yxx}^{2\omega}$  changes only 232 slightly when tuning the electric field away from *D*=0. Next, we investigate 233 the effect of charge carrier density  $n$  on the nonlinear response. Fig. 4c 234 shows the nonlinear conductivity  $\sigma_{yxx}^{2\omega}$  and  $\sigma_{xxxx}^{2\omega}$  as a function of carrier 235 density *n*, respectively. When the Fermi level is tuned into the charge 236 neutral gap, both  $\sigma_{yxx}^{2\omega}$  and  $\sigma_{xxxx}^{2\omega}$  vanish. This is in line with the fact 237 that the quantum metric dipole is a Fermi surface property<sup>30</sup>. Furthermore, 238 both  $\sigma_{yxx}^{2\omega}$  and  $\sigma_{xxxx}^{2\omega}$  exhibit a sign reversal when tuning the carrier 239 density between hole and electron regimes. Lastly, given the evident 240 nonreciprocal longitudinal response, we can evaluate the nonreciprocity 241 coefficient  $\gamma$  in MnBi<sub>2</sub>Te<sub>4</sub>. We find the coefficient  $\gamma$  can reach  $\gamma \approx 7 \times$  $242 \quad 10^{-11} m^2 A^{-1}$ , which is two or three orders larger than traditional heavy 243 metal (HM)/ferromagnetic metal (FM) heterostructures (See details in 244 supplementary information section S4.3)<sup>46,47</sup>. 229 fold rotational symmetry. To d<br>
230 dependence of  $\sigma_{yxx}^{2\omega}$  on the electri<br>
231  $n_e \approx -3 \times 10^{12} \text{ cm}^{-2}$ . As show<br>
232 slightly when tuning the electric fiel<br>
233 the effect of charge carrier density<br>
234 shows t mg the small conductivity and low mobility<br>
i-independent nonlinear conductivity further<br>
impurity scattering, because its lowest order<br>
r conductivity begins at  $\tau^1$  (see detailed<br>
information S7.2)<sup>21,44</sup>. In conclusi

245 To explain our experimental data, we evaluate  $\sigma_{yxx}^{2\omega}$  quantitatively by 246 constructing a slab model for  $4SL MnBi<sub>2</sub>Te<sub>4</sub>$  with density-functional theory (DFT) calculations. Fig. 4d shows the total conductivity as a function of carrier density, in which we assume  $\sigma_{yxx}^{2\omega}$  for an ideally aligned device (*y* aligned with the in-plane crystalline axis). Importantly, around the charge 250 neutrality gap ( $\mu = 0$ ), we observe a sign change which reproduces the experimental findings. (see supplementary information S7.1 for the band-resolved contributions to the quantum metric) The deviation of theoretical  $\sigma_{yxx}^{2\omega}$  compared to experimental values may be attributed to the known discrepancy in surface band structure between calculations and experiments on MnBi2Te4. Additionally, in our calculation, we find that the Drude contribution is small within a large energy range around the Fermi level, indicating that the nonlinear conductivity is primarily driven by the quantum metric dipole. We conducted similar measurements on another 6SL-MnBi2Te4 device, as detailed in supplementary information section S6. hows the total conductivity as a function of<br>ssume  $\sigma_{yxx}^{2\omega}$  for an ideally aligned device (y<br>talline axis). Importantly, around the charge<br>sperve a sign change which reproduces the<br>pplementary information S7.1 for th

As a probe, the quantum metric dipole-driven electron nonlinearity in MnBi2Te4 has several advantages and holds great promise for future applications. First, compared to nonlinearity originating from the Berry curvature dipole or skew scattering, the quantum metric dipole-driven electron nonlinearity is independent of the scattering time, suggesting its robustness against disorder scattering. Second, the quantum metric dipole-driven electron nonlinearity is distinct for opposite AFM spin orders and even robust against small perturbations of external magnetic fields (see details in supplementary information S3), making it a promising candidate for use as a nonlinear magnetic memory device. Finally, the quantum metric dipole-driven nonlinear response is more prominent than the "traditional" nonlinear response from the Berry curvature dipole. We 273 compared the nonlinear response observed in  $MnBi<sub>2</sub>Te<sub>4</sub>$  with other 2D material systems, as presented in Extended Table I. We find that our MnBi<sub>2</sub>Te<sub>4</sub> devices have a much larger nonlinear conductivity than WTe<sub>2</sub> 276 and strained WSe<sub>2</sub>, only smaller than twisted WSe<sub>2</sub> and graphene Moiré 277 superlattices, making  $MnBi_2Te_4$  a promising candidate for highly efficient 264 curvature dipole or skew scatter<br>
265 electron nonlinearity is independer<br>
266 robustness against disorder scatterir<br>
267 driven electron nonlinearity is disti<br>
268 even robust against small perturba<br>
269 details in s

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#### **Figure captions:**

**Figure 1 Quantum metric induced nonlinearity in a** *PT***-symmetric AFM. a.** The nonlinear Hall effect originating from Berry curvature dipole, which has been observed in systems with broken inversion symmetry. **b.** The nonlinear longitudinal and transverse response observed in a *PT*-symmetric system due to quantum metric dipole. **c.** The crystal structure of 410 even-layer MnBi<sub>2</sub>Te<sub>4</sub> from the side view. The arrows represent the spin momentum direction of the Mn atoms. **d.** The spatial reflectance magnetic circular dichroism (RMCD) mapping of MnBi2Te4 measured at zero magnetic field. The dashed line marks the boundary between the 4SL-MnBi2Te4 and 3SL- MnBi2Te4. **e.** The crystal structure of MnBi2Te4 from the top view. The dashed line marked with *M* represents the mirror line. **f.** The angle dependent optical second-harmonic generation (SHG) of the 4SL-MnBi2Te4 sample. The zero degree corresponds to the applied current direction in the device.

**Figure 2 Observation of nonlinear transport in AFM MnBi2Te4. a.** The 420 schematic view of the dual-gated  $4SL$ -layer MnBi<sub>2</sub>Te<sub>4</sub> device. The linear and nonlinear signals are measured simultaneously with the denoted 422 direction. **b.** The linear longitudinal  $V_x^{\omega}$  and transverse  $V_y^{\omega}$  voltage as a 423 function of current  $I_x^{\omega}$ . The solid line is the linear fitting of the data. **c and** 424 **d.** The nonlinear longitudinal  $V_x^{2\omega}$  and transverse  $V_y^{2\omega}$  voltage as a 425 function of current  $V_x^{\omega}$  for AFM-I and AFM-II states, respectively. At a fixed electric field, the different AFM states are prepared by: AFM-I, sweeping the magnetic field from -7 T to 0 T; AFM-II, sweeping the 428 magnetic field from  $+7$  T to 0 T. The solid lines are quadratic fits to the data. **e.** The optical image of another 4SL-MnBi2Te4 device with radially distributed electrodes. The dashed line indicates the in-plane crystalline 431 axis, determined by SHG measurement. scale bar:  $5 \mu m$ . **f.** The nonlinear 432 transverse  $V_y^{2\omega}$  and longitudinal  $V_x^{2\omega}$  response as a function of current 433 injection angle  $\theta$ . AT **A SET and the matrix in the control in the set of the matrix particle in the set of the matrix in the matrix in the set of the matrix system o** 

**Figure 3 Spin order related electron nonlinearity from bandnormalized quantum metric dipole. a.** Nonlinear transverse voltage  $V_y^{2\omega}$ of 4SL-MnBi2Te4 as a function of temperature for opposite AFM-I and AFM-II states, respectively. The amplitudes of  $V_y^{2\omega}$  are similar for the 438 two AFM states but their sign is reversed.  $V_y^{2\omega}$  vanishes when the temperature is above the Néel temperature of MnBi2Te4. **b.** The nonlinear 440 longitudinal voltage  $V_x^{2\omega}$  as a function of temperature for opposite AFM-141 I and AFM-II states, respectively, exhibiting similar trend as  $V_y^{2\omega}$ . **c.** The conductivity and fitted carrier density of 4SL-MnBi2Te4 at different temperatures. **d.** The scaling relationship between the nonlinear

444 conductivity  $\sigma_{yxx}^{2\omega}$  ( $\sigma_{xxxx}^{2\omega}$ ) and the square of the linear longitudinal 445 conductivity  $\sigma_{xx}^2$ . The scaling is carried out at the temperature range of 2 446 -10 K (marked with shadow area in **c**), in which the carrier density nearly 447 remains constant. The nonlinear transverse and longitudinal conductivities are extracted by  $\sigma_{yxx}^{2\omega} = \frac{V_y^{2\omega}}{(I_x^{0\omega})^2}$  $(I_x^{\omega})^2 R_{xx}^3$  $rac{L^3}{W^2}$  and  $\sigma_{xxx}^{2\omega} = \frac{V_x^{2\omega} L}{(I_x^{\omega})^2 R_x^3}$ 448 are extracted by  $\sigma_{yxx}^{2\omega} = \frac{y}{(I_x^{\omega})^2 R_{xx}^3} \frac{L}{W^2}$  and  $\sigma_{xxxx}^{2\omega} = \frac{v_x}{(I_x^{\omega})^2 R_{xx}^3}$ , respectively. 449 The dashed line is a fit of the data with the scaling relation  $\sigma^{2\omega}$  = 450  $\eta_2(\sigma_{xx}^{\omega})^2 + \eta_0$ . From the fitting, we conclude that the predominant 451 contribution is from the  $\eta_0$  term, namely from the intrinsic nonlinear 452 conductivity due to the band-normalized quantum metric dipole.

453 **Figure 4 The electric field and carrier density dependence of the**  454 **nonlinear response. a.** The resistance map of the 4SL-MnBi2Te4 device. 455 The vertical displacement field *D* and charge carrier density *ne* can be 456 independently controlled, as denoted in the figure. **b.** The nonlinear 457 transverse conductivity  $\sigma_{yxx}^{2\omega}$  and longitudinal resistance  $R_{xx}$  as a function 458 of vertical displacement electric field. The carrier density is fixed at  $n_e \approx$  $-3 \times 10^{12}$  cm<sup>-2</sup> as D varies along the dashed line in **a**. **c.** The 460 measured nonlinear transverse conductivity  $\sigma_{yxx}^{2\omega}$  and longitudinal 461 conductivity  $\sigma_{xxx}^{2\omega}$  as a function of charge carrier density  $n_e$ . **d.** The 462 calculated total nonlinear transverse conductivity  $\sigma_{yxx}^{2\omega}$  (red line) as a 463 function of carrier density. The grey line denotes the Drude contribution. 44 -10 K (marked with biadow are ain c), in which the carrier density many<br>44 -10 K (marked with shadow are aided by  $\sigma_{\mu\nu}^{2\mu} = \frac{\sqrt{\mu^2}}{(\mu^2)^2 R_{\pi\nu}^2}$  and  $\sigma_{\mu\nu}^{2\mu} = \frac{\sqrt{\mu^2}}{(\mu^2)^2 R_{\pi\nu}^2}$  and  $\sigma_{\mu\nu}^{$ 

#### 464 **Methods**

#### 465 **MnBi2Te4 single Crystal growth**

466 Single crystals of MnBi<sub>2</sub>Te<sub>4</sub> were grown using a self-flux method<sup>48,49</sup>. The 467 starting materials used in the single crystal processes are Mn slices, Bi 468 lumps, and Te chunks. MnTe and  $Bi<sub>2</sub>Te<sub>3</sub>$  precursors were prepared by 469 reacting the mixed stoichiometric starting materials at  $1100 \degree C$  and  $900 \degree C$ 470 for 24 h, respectively. Then MnTe and  $Bi<sub>2</sub>Te<sub>3</sub>$  were mixed according to the 471 ratio MnTe:  $Bi<sub>2</sub>Te<sub>3</sub> = 15: 85$ , loaded into an alumina crucible, and sealed 472 in a quartz tube. The quartz tube was heated to  $650 \degree C$  in 10 h, dwelled for 473 12 h, and slowly cooled to 595  $\degree$ C at a rate of 1  $\degree$ C/h to grow the single 474 crystals. Shiny single crystals with a typical size of  $3\times2\times0.5$  mm<sup>3</sup> can be 475 obtained after centrifuging.

476

#### 477 **Device fabrication**

478 The MnBi<sub>2</sub>Te<sub>4</sub> thin flakes were obtained using the  $Al_2O_3$ -assisted 479 exfoliation method<sup>35,50</sup>. First, a 70 nm thick layer of  $Al_2O_3$  was thermally 480 evaporated onto bulk  $MnBi<sub>2</sub>Te<sub>4</sub>$  crystals. Second, the  $Al<sub>2</sub>O<sub>3</sub>$  thin film was 481 lifted along with thin flakes of MnBi2Te4, which were cleaved from the 482 bulk crystal using thermal release tape. Then, the stacked  $MnBi_2Te_4/Al_2O_3$ 483 was transferred onto a transparent Polydimethylsiloxane (PDMS) film. The 484 number of layers in the MnBi2Te4 thin flakes was determined using optical 485 transmittance measurements. After confirming the layer number, the 486 MnBi2Te4 thin flake samples were transferred to a Si wafer coated with  $487$  285 nm of SiO<sub>2</sub>. To ensure high-quality samples, we used a stencil mask 488 method to deposit metal contacts (Cr/Au) on the samples. A 20-40 nm thick 489 h-BN flake was then transferred to the  $MnBi<sub>2</sub>Te<sub>4</sub>$  devices as the top gate 490 dielectric layer, followed by the transfer of a graphite thin flake gate on top 491 of the h-BN/MnBi<sub>2</sub>Te<sub>4</sub> heterostructure. The entire device fabrication 492 process was carried out in a nitrogen-filled glove box with  $O_2$  and  $H_2O$ 493 levels below 1 ppm. AN **NOTIFIELD SIDE CYSTRA (SPACE THE SECT ARTICLE THE SECT AND THE SECT AND THE SECT AND THE SECTION THE SECTION (SPACE THE SECTION THE SECTION AND THE SECTION ARTICLE THE SECTION (SPACE THE SECTION AND THE SECTION (SPACE** 

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#### 495 **Nonlinear Electrical transport measurements**

The electrical transport measurements were performed in a cryogenic-free cryostat equipped with a superconducting magnet (Cryomagnetic). We used a standard lock-in technique (Zurich MFLI) with excitation frequencies ranging from 10 to 200 Hz to measure both the first and second harmonic signals. The data presented in this manuscript was collected at a

501 low frequency of 17.777 Hz. During the transport measurements, the phase 502 of the first and second harmonic signals was confirmed to be 503 approximately 0 and 90 degrees, respectively (see supplementary 504 information S2.4). The gate voltages were applied using a Keithley 2636 505 SourceMeter. To independently control the charge carrier density *ne* and 506 displacement electric field *D*, we used a dual gated device structure. The 607 charge carrier density *n* can be obtained by  $n = \left(\frac{\varepsilon_0 \varepsilon^{BN} (V_{TG} - V_{TG_0})}{d_{BN}}\right) +$  $\varepsilon_0 (V_{BG} - V_{BG0}) / \left(\frac{d_{Al_2O_3}}{c_{Al_2O_3}}\right)$  $rac{d_{Al_2O_3}}{\varepsilon^{Al_2O_3}} + \frac{d_{SiO_2}}{\varepsilon^{SiO_2}}$  $\epsilon_0 (V_{BG} - V_{BG0}) / (\frac{\mu_{Al_2O_3}}{\varepsilon^{Al_2O_3}} + \frac{\mu_{SU_2}}{\varepsilon^{SiO_2}})$ )/e. Here we regarded the Al<sub>2</sub>O<sub>3</sub> and SiO<sub>2</sub> 509 dielectric layer as two series connected capacitors. The electric displacement field *D* can be obtained by  $D = \left(\frac{V_{BG} - V_{BG}}{d_{M,Q}}\right)^{d_{G,Q}}$  $\left(\frac{d_{Al_2O_3}}{d_{LO_2O_3}}\right)$  $rac{d_{Al_2O_3}}{d_{\epsilon}^{Al_2O_3}} + \frac{d_{SiO_2}}{d_{\epsilon}^{SiO_2}}$ 510 displacement field D can be obtained by  $D = \left(\frac{VBG - VBG}{dA} - \frac{dG}{dA} \right)$ degrees, respectively (see supplementary<br>oltages were applied using a Keithley 2636<br>dly control the charge carrier density  $n_e$  and<br>we used a dual gated device structure. The<br>interval and gated device structure. The<br> $\frac{a$ 

 $\varepsilon^{BN}(V_{TG}-V_{TG0})$  $\frac{\varepsilon^{(V)}(T_G - V_T G_0)}{d_{BN}}$  /2. Here,  $\varepsilon_0$  is the vacuum permittivity; *V<sub>TG0</sub>* and *V<sub>BG0</sub>* 512 correspond to the gate voltage of maximum resistance;  $\varepsilon^{Al_2O_3}$ ,  $\varepsilon^{SiO_2}$  and 513  $\varepsilon^{BN}$  are the relative dielectric constant of Al<sub>2</sub>O<sub>3</sub>, SiO<sub>2</sub> and h-BN;  $d_{Al_2O_3}$ , 514  $d_{SiO_2}$  and  $d_{BN}$  are the thickness of Al<sub>2</sub>O<sub>3</sub>, SiO<sub>2</sub> and h-BN, respectively. 515 We noticed a minor asymmetry in  $\sigma_{yxx}^{2\omega}$  and  $R_{xx}$  with respect to the 516 displacement electric field  $D$ , possibly attributed to charge trapping 517 induced by substrate effects or disorder<sup>51</sup>. The asymmetry is more 518 pronounced in  $\sigma_{yxx}^{2\omega}$ , as the quantum metric is related to energy differences  $519$  between bands and more sensitive to D.

## 520 **The nonlinear response in the MnBi2Te4 device with radially**  521 **distributed devices**

522 For the MnBi2Te4 device with radially distributed devices, when the 523 current aligns with the in-plane crystal axis ( $\theta = 0^{\circ}$ ), the transverse  $V_y^{2\omega}$ 524 is zero. In contrast, when the current is perpendicular to the crystalline axis 525 ( $\theta = 90^{\circ}$ ), the longitudinal  $V_x^{2\omega}$  is zero. In addition, the nonlinear 526 response should obey the symmetry of even-layer MnBi<sub>2</sub>Te<sub>4</sub>, with  $\sigma_{yxx}$  = 527  $-\sigma_{yyy}$  and  $\sigma_{xyy} = -\sigma_{xxx}$ . To substantiate this, we use the formula for nonlinear conductivity:  $\sigma_{xxxx}^{2\omega} = \frac{J_x^{2\omega}}{(\epsilon \omega)}$  $\frac{J_x^{2\omega}}{(E_{xx}^{\omega})^2} = \frac{V_x^{2\omega} L}{(I_x^{\omega})^2 R_x^3}$  $\frac{V_x^{2\omega} L}{(I_x^{\omega})^2 R_{xx}^3}$  and  $\sigma_{yxx}^{2\omega} = \frac{J_y^{2\omega}}{(E_{xx}^{\omega})^2}$ 528 nonlinear conductivity:  $\sigma_{xxxx}^{2\omega} = \frac{J_x}{(E_{xx}^{\omega})^2} = \frac{V_x}{(I_x^{\omega})^2 R_{xx}^3}$  and  $\sigma_{yxx}^{2\omega} = \frac{J_y}{(E_{xx}^{\omega})^2}$  $V_y^{2\omega}$  $(I_x^{\omega})^2 R_{xx}^3$ 529  $\frac{V_y^2 \omega}{(I_x^{\omega})^2 R_{xx}^3 W^2}$ . Here, *L* refers to the length and *W* to the width of the device. 530 The optical image (Fig. 2e) reveals that the length-to-width ratio,  $L/W$ , is 518 pronounced in  $\sigma_{yxx}^{2\omega}$ , as the quantum<br>
519 between bands and more sensitive t<br>
520 **The nonlinear response in th**<br>
521 **distributed devices**<br>
522 For the MnBi<sub>2</sub>Te<sub>4</sub> device with ra<br>
523 current aligns with the 531 approximately 0.85. Taking into consideration the geometric

characteristics of the device, we conclude that the nonlinear conductivity

533 conforms to  $\sigma_{yxx} = -\sigma_{yyy}$  and  $\sigma_{xyy} = -\sigma_{xxx}$ .

#### **Optical measurement**

For the RMCD measurement, a linearly polarized 633 nm HeNe laser was modulated by a photoelastic modulators (PEM) at a frequency of 50 kHz. The laser beam was focused on the sample at normal incidence using an 538 Olympus MPLN50 $\times$  objective with a 0.75 numerical aperture. The laser 539 spot size was approximately 2  $\mu$ m, with a power of 1.8  $\mu$ W. The reflected light was collected by a photodetector and analyzed using a lock-in amplifier set to the same frequency as the PEM. The RMCD measures the differential reflection between left and right circularly polarized light, the magnitude of which is proportional to the magnetic moments of the MnBi2Te4 sample. **EVALUATION CONSULTERATE:** An interaction of the photological of Sin methods of the modulated by a photocheck of the set of the set of the last Fisure with a forest or the angular state of the set of the last Fisure with

The second harmonic generation (SHG) measurement was performed at room temperature using femtosecond pulse lasers with a central wavelength of 800 nm. The laser beam was focused at normal incidence on the sample using a Nikon ELWD 100x microscope objective. The SHG signals were collected using a spectrometer. To perform the polarization-resolved measurement, a motorized polarizer was used to control the polarization of the incident laser beam as it was rotated next to the objective.

**DFT calculations**

We use VASP (Vienna ab-initio software package) with the PBE functional in the generalized gradient approximation, to obtain the electronic ground state<sup>52</sup>. We then project the ground state wavefunctions on Wannier functions using Wannier  $90<sup>53</sup>$  and create a tight-binding model with 184 Wannier orbitals. For the Drude contribution we adopt a lifetime 558 of  $\tau \approx 40$  fs estimated from our experimental conductivity and carrier density in Fig. 2c and theoretical effective mass  $(m^* = 0.14m_e)$  for the 4 SL film.

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**Author contributions:** W.-b.G and B.Y conceived and supervised the project. N.W fabricated the devices, performed the transport and RMCD measurement with help from Z.Z and C.Z. D.K, T.H and B.Y performed 599 the theoretical calculations. N.W and F.Z performed the NV measurement with help from Z.J, S.R and H.C. N.W, W.-b.G, D.K, T.H and B.Y analysed the data. N.C, A.W and X.Z grew the MnBi2Te4 single crystals. K.W and T.T grew the hBN single crystals. N.W, B.Y and W.-b.G wrote the manuscript with input from all authors.

**Competing interests**: The authors declare no competing interests.

**Data availability:** Source data in the main text and extended figures are provided with this paper.

**Additional information:** Correspondence and requests for material should 608 be addressed to Binghai Yan (binghai.yan $(\partial w)$ weizmann.ac.il) or Wei-bo 609 Gao (wbgao@ntu.edu.sg).

**Extended Figure 1 The linear conductivity of 4SL-MnBi2Te4 with opposite AFM states. a and b.** The AFM-I and AFM-II states are prepared 614 by sweeping the magnetic field from -7 T to  $0T$  or +7 T to  $0T$ , respectively. **c and d.** The linear longitudinal  $V_x^{\omega}$  and transverse  $V_y^{\omega}$  voltage as a 616 function of current  $I_x^{\omega}$  for AFM I and AFM II states, respectively. The solid line is a linear fit to the experimental data. **Extended Figure 2 The fully compensated AFM order in 4SL-**Eta by sweeping the magnetic field from -7 T to OT or 17 T to OT, respectively,<br>
(ii) c and d. The linear longitudinal  $V_1^2$  and transverse  $V_1^*$  voltage as a<br>
first line is a linear til to the experimental data.<br>
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- 619 **MnBi2Te4 device. A.** The magnetic field dependent longitudinal resistance
- 620 *Rxx* of the 4SL-MnBi2Te4 device. **B.** The magnetic field dependent Hall
- 621 resistance  $R_{yx}$  of the 4SL-MnBi<sub>2</sub>Te<sub>4</sub> device. In zero magnetic field, the
- 622 AFM order is fully compensated and the Hall resistance  $R_{yx} = 0$ .
- 623 **Extended Table 1 Comparison of the nonlinear conductivity for**
- 624 **MnBi2Te4 and other two-dimensional material systems.** The nonlinear
- 625 transverse conductivity  $\sigma_{yxx}^{2\omega}$  in even-layer MnBi<sub>2</sub>Te<sub>4</sub> is summarized and 626 compared with other reported material systems<sup>9,10,13,16,19</sup>. The
- 627 corresponding mechanisms responsible for the nonlinear response are also
- 628 listed for comparison.











**Extended Data Fig. 1**





**Extended Data Table 1**