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1 Title: Quantum metric-induced nonlinear transport in a topological

2 antiferromagnet

Authors: Naizhou Wang¹, Daniel Kaplan², Zhaowei Zhang¹, Tobias Holder², Ning
Cao³, Aifeng Wang³, Xiaoyuan Zhou³, Feifei Zhou¹, Zhengzhi Jiang¹, Chusheng
Zhang¹, Shihao Ru¹, Hongbing Cai¹, Kenji Watanabe⁴, Takashi Taniguchi⁵, Binghai
Yan^{2*}, Weibo Gao^{1*}

¹ Division of Physics and Applied Physics, School of Physical and Mathematical
 Sciences, Nanyang Technological University, Singapore, Singapore

⁹ ² Department of Condensed Matter Physics, Weizmann Institute of Science, Rehovot,
 ¹⁰ Israel

¹¹ ³ Low Temperature Physics Laboratory, College of Physics and Center for Quantum

12 Materials and Devices, Chongqing University, Chongqing, China

¹³ ⁴Research Center for Functional Materials, National Institute for Materials Science, 1-

14 1 Namiki, Tsukuba, Japan

¹⁵ International Center for Materials Nanoarchitectonics, National Institute for Materials

- 16 Science, 1-1 Namiki, Tsukuba, Japan
- 17 *Corresponding authors' email: <u>binghai.yan@weizmann.ac.il</u>; <u>wbgao@ntu.edu.sg</u>
- 18

Abstract: The Berry curvature and quantum metric are the imaginary 19 part and real part, respectively, of the quantum geometric tensor 20 which characterizes the topology of quantum states¹. The former is 21 22 known to generate a zoo of important discoveries such as quantum Hall effect and anomalous Hall effect (AHE)^{2,3}, while the consequences 23 of the quantum metric have rarely been probed by transport. Here we 24 25 report the observation of quantum metric-induced nonlinear transport, including both nonlinear AHE and diode-like nonreciprocal 26 longitudinal response, in thin films of a topological antiferromagnet, 27 MnBi₂Te₄. Our observation reveals that the transverse and 28 longitudinal nonlinear conductivities reverse signs when reversing the 29 30 antiferromagnetic order, diminish above the Néel temperature, and are insensitive to disorder scattering, thus verifying their origin in the 31 32 band structure topology. They also flip signs between electron and hole-doped regions, in agreement with theoretical calculations. Our 33 work provides a pathway to probe the quantum metric through 34 nonlinear transport and to design magnetic nonlinear devices. 35

37 Main text:

Nonlinear transport provides a powerful tool for probing topological 38 physics in solids⁴⁻²¹. A prime example of this is the nonlinear anomalous 39 Hall effect (NLAHE) which reveals the Berry curvature dipole on the 40 Fermi surface and has recently been observed in nonmagnetic topological 41 semimetals (as shown in Fig. 1a)^{6,9,10,12,14,15}. This provides a powerful 42 means of investigating quantum geometry in flat space. Another NLAHE 43 caused by the quantum metric, the counterpart of the Berry curvature in the 44 quantum geometry, was also predicted for magnetic topological materials, 45 but is yet to be realized in experiments^{1,22-30}. The quantum metric measures 46 the amplitude distance between two neighboring Bloch states and 47 determines the electronic properties of a crystal. A very recent theory^{26,30} 48 finds that the quantum metric actually gives rise to both NLAHE and the 49 nonreciprocal longitudinal response, the latter of which is characterized by 50 a diode-like longitudinal resistance, in inversion-breaking magnetic 51 materials (as shown in Fig. 1b)⁷. The asymmetric quantum metric 52 (quantum metric dipole) will induce an anomalous motion of the wave 53 packet and generate nonlinear responses in both the longitudinal and 54 transverse directions. Since the quantum metric is antisymmetric under 55 momentum inversion $(k \rightarrow -k)$, both time reversal symmetry (T) and 56 inversion symmetry (P) must be broken in order to generate a net quantum 57 metric dipole. Additionally, the combined symmetry PT can exclude the 58 contribution from the Berry curvature dipole⁴. As a result, a system which 59 breaks both P and T but preserves PT is expected to be an ideal platform 60 for studying the quantum metric effect (see supplementary information S1 61 for detailed symmetry analysis). 62

After careful consideration, we chose MnBi₂Te₄ as the candidate platform 63 to investigate the quantum metric. Compared to the magnetic doped 64 topological insulator, the intrinsic magnetic topological insulator, 65 MnBi₂Te₄, gained a lot of interest recently due to its unique properties^{17,31-} 66 ⁴¹. As shown in Fig. 1c and e, the crystal structure of MnBi₂Te₄ consists of 67 alternating layers of Te-Bi-Te-Mn-Te-Bi-Te, known as septuple layers 68 (SLs)^{33,34}. MnBi₂Te₄ has an A-type antiferromagnetic (AFM) ground state, 69 in which the Mn spins in each SL are aligned ferromagnetically with an 70 out-of-plane easy axis but are coupled antiparallel to adjacent SLs. For 71 even-layer MnBi₂Te₄, both P and T symmetry are broken but the combined 72

PT symmetry is preserved below the Néel temperature³⁷. Fig. 1d shows the 73 74 spatial reflectance magnetic circular dichroism (RMCD) mapping of 3SL and 4SL MnBi₂Te₄ measured at zero magnetic field, indicating fully 75 compensated AFM states in even layer MnBi₂Te₄⁴⁰. In addition, MnBi₂Te₄ 76 possesses three-fold rotational symmetry (from the optical second 77 harmonic generation shown in Fig. 1f), which suppresses the Berry 78 curvature dipole even in the absence of PT-symmetry, making the 79 contribution from the quantum metric much easier to observe^{30,41}. 80

81 Electron nonlinearity from spin order

To investigate the electron nonlinearity, we fabricated several high-quality, 82 dual gated even-layer MnBi₂Te₄ devices. Here, we focus on a 4SL-83 MnBi₂Te₄ device, the schematic structure of which is shown in Fig. 2a. The 84 dual-gate structure allows us to independently control the carrier density 85 and vertical displacement field. As per theoretical predictions, the quantum 86 metric gives rise to both NLAHE and the nonreciprocal longitudinal 87 response³⁰. The former is directly measurable via an alternating current 88 (AC) method, while the latter exhibits diode-like nonreciprocal resistance 89 behavior, expressed as: $V_x = R_0(I + \gamma I^2)$ where I represents the 90 applied current, R_0 is the linear resistance, and γ denotes the coefficient 91 characterizing the nonreciprocity⁷. Considering that the nonreciprocal 92 longitudinal response stems from the quadratic term of the current, we 93 adopted the AC method to measure the second-harmonic voltage $V_x^{2\omega}$ to 94 reflect the non-reciprocal behavior in longitudinal direction, which offers 95 a better signal-to-noise ratio (see supplementary information section S4). 96 97 To measure the linear and non-linear responses of the device, we employed a standard lock-in technique. As depicted in Fig. 2a, an AC current with 98 frequency of ω (I^{ω}) was injected, and the linear voltage V^{ω} and second-99 harmonic voltage $V^{2\omega}$ are simultaneously probed. In our experiment, the x-100 axis is defined as the current direction and the y-axis as the transverse 101 direction to the current. All the measurements are conducted at T = 1.8 K 102 in the absence of a magnetic field unless otherwise specified. Fig. 2b shows 103 104 the linear longitudinal (V_x^{ω}) and transverse (V_y^{ω}) voltages, indicating good 105 ohmic contact and a negligible misalignment in the Hall bar geometry from the vanishing of V_v^{ω} . 106

Next, we focus on the nonlinear responses of $4SL-MnBi_2Te_4$ in opposite AFM states. Utilizing a method akin to those reported previously for

controlling AFM states in CrI₃ or even-layer MnBi₂Te₄, we prepared the 109 AFM-I and AFM-II states^{37,42} with opposite Néel orders. As shown in Fig. 110 2c and 2e, the AFM-I state is prepared by sweeping the magnetic field from 111 112 -7 T to zero, while the AFM-II state is achieved by sweeping the magnetic field from +7 T to zero. For the linear responses, both AFM states exhibited 113 identical longitudinal (V_x^{ω}) voltages, and the transverse (V_y^{ω}) voltages 114 remained at zero (Extended Data Fig. 1). This observation aligns with the 115 116 fully compensated AFM orders in 4SL-MnBi₂Te₄ (Extended Data Fig. 2). However, the nonlinear responses differ significantly from their linear 117 counterparts. Focusing first on the AFM-I state, Fig. 2c shows that while 118 the linear transverse (V_v^{ω}) voltage remains at zero, there is a significant 119 negative nonlinear transverse voltage (notated as $V_y^{2\omega}$) that scales 120 quadratically with the injected current I^{ω} . More importantly, we also 121 122 detected a prominent negative nonlinear longitudinal voltage (denoted as $V_x^{2\omega}$), with the same order of magnitude as $V_y^{2\omega}$. This observation aligns 123 with the prediction that the quantum metric dipole can induce both 124 nonlinear Hall and nonreciprocal longitudinal response, distinctly different 125 from the response observed in the nonlinear Hall effect caused by the Berry 126 curvature dipole, where only a Hall response is allowed and the 127 longitudinal response is absent^{9,10}. The nonreciprocal longitudinal response 128 in MnBi₂Te₄ can also be observed through DC measurements (see 129 130 supplementary information section S4). Subsequently, we prepared the AFM-II state. As shown in Fig. 2d, in sharp contrast, although the 131 amplitudes of $V_y^{2\omega}$ and $V_x^{2\omega}$ remain nearly the same with the AFM-I 132 state, the sign of both flips to positive. The sign reversal of the nonlinear 133 134 response suggests that the nonlinear response observed in MnBi₂Te₄ is associated with the AFM order. We carefully examined and ruled out 135 potential alternative origins, such as thermal effect or accidental diode 136 137 junction that could lead to a nonlinear effect. We also exclude the 138 possibility that the nonlinear response observed in even-layer MnBi₂Te₄ is originated from the residual magnetization (See supplementary 139 140 information section S2). To further confirm that the nonlinear signal is an intrinsic response originating from the *PT*-symmetric AFM order, we 141 142 investigate whether the nonlinear response complies with the rotational symmetry of AFM even-layer MnBi₂Te₄, where $\sigma_{yxx} = -\sigma_{yyy}$ and 143 $\sigma_{xyy} = -\sigma_{xxx}$. When y aligns with the in-plane crystal axis, it should 144

follow that $\sigma_{xyy} = -\sigma_{xxx} = 0$. The coexistence of $\sigma_{yxx}^{2\omega}$ and $\sigma_{xxx}^{2\omega}$ 145 in this 4SL-MnBi₂Te₄ device suggests some misalignment between x/y and 146 the crystal axes, consistent with our optical SHG measurement in Fig. 1f. 147 To substantiate this further, we fabricated another 4SL-MnBi₂Te₄ device 148 149 with radially distributed electrodes aligned to the crystalline axis (dashed line in Fig. 2e). Fig. 2f demonstrates the nonlinear transverse and 150 longitudinal response upon rotation of the current injection direction. Both 151 transverse $V_y^{2\omega}$ and longitudinal $V_x^{2\omega}$ exhibit three-fold symmetry. 152These observed features align consistently with the symmetry of the AFM 153154 even-layer MnBi₂Te₄.

We further investigated the temperature dependence of the nonlinear 155response in MnBi₂Te₄. Fig 3a and 3b shows the temperature dependent 156 nonlinear voltages $V_y^{2\omega}$ and $V_x^{2\omega}$ which are prepared at AFM-I and 157 AFM-II states, respectively. As the temperature increases, we found that 158 the nonlinear voltage gradually decreases and vanishes above the Néel 159 temperature of MnBi₂Te₄, indicating that the nonlinear response is absent 160 in the non-magnetic states. In addition, the nonlinear response in even-161 layer MnBi₂Te₄ is only presented at the AFM states and vanishes when all 162 spins are aligned in one direction (see details in supplementary information 163 section S3). In summary, we can conclude that the nonlinear response 164 observed is associated with the nonlinearity of electron motion originating 165 from the AFM order in 4SL-MnBi₂Te₄. 166

167 Quantum metric induced nonlinear transport

168 We now systematically investigate the microscopic origin of the nonlinear response observed in 4SL-MnBi₂Te₄. To quantify the strength of the 169 nonlinear response, we use the nonlinear conductivity, as it is independent 170 of the sample size. In our experiment, we can extract the longitudinal and 171 transverse nonlinear conductivity from our data as $\sigma_{xxx}^{2\omega} = \frac{J_x^{2\omega}}{(E_{xxx}^{\omega})^2} =$ 172 $\frac{V_x^{2\omega} L}{(I_x^{\omega})^2 R_{xx}^3}$ and $\sigma_{yxx}^{2\omega} = \frac{J_y^{2\omega}}{(E_{xx}^{\omega})^2} = \frac{V_y^{2\omega}}{(I_x^{\omega})^2 R_{xx}^3} \frac{L^3}{W^2}$, respectively, where L and W are 173the longitudinal and transverse length of the Hall bar device. In theory, the 174 nonlinear conductivity due to an applied electric field E_x in a metal includes 175 three contributions³⁰, 176

$$J_c = \sigma_{cxx} E_x^2 = (\sigma_{cxx}^2 + \sigma_{cxx}^1 + \sigma_{cxx}^0) E_x^2$$
(Eq. 1)

where σ_{cxx}^{i} has the *i*-th order of τ dependence (c = x or y). Here $\sigma_{cxx}^{2} = -\frac{e^{3}\tau^{2}}{\hbar^{3}}\int d^{2}kf_{n}\partial_{k_{c}}\partial_{k_{x}}\varepsilon_{n}(\mathbf{k})$ is the nonlinear Drude weight, and $\varepsilon_{n}(\mathbf{k})$ is the energy of Bloch state n at momentum \mathbf{k} , and f_{n} is the Fermi Dirac distribution for band n. $\sigma_{cxx}^{1} = -\frac{e^{3}\tau}{\hbar^{2}}\int d^{2}kf_{n}(2\partial_{k_{x}}\Omega_{n}^{xc})$, is the Berry curvature dipole contribution⁴. The conductivity due to the normalized dipole of the quantum metric is³⁰,

184
$$\sigma_{baa}^{0} = -\frac{e^{3}}{\hbar^{2}} \int d^{2}k f_{n} (2\partial_{k_{b}}G_{n}^{aa} - \partial_{k_{a}}G_{n}^{ab}) \qquad (\text{Eq. } 2$$

Where a/b = x/y, $G_n^{ab} = \sum_{m \neq n} \frac{A_{nm}^a A_{mn}^b + A_{nm}^b A_{mn}^a}{(\varepsilon_n - \varepsilon_m)}$ is the band-energy 185 normalized quantum metric and $\partial_{k_a} G_n^{ab}$ represents the normalized 186 quantum metric dipole. In the specific case of 4SL-MnBi₂Te₄, the PT-187 symmetry excludes any contribution from the Berry curvature dipole⁴. 188 Additionally, the Berry curvature dipole contribution, which is even under 189 time-reversal, contradicts the sign reversal of the nonlinear signal between 190 the AFM-I and II phases. It is also worth noting that the skew scattering 191 mechanism can also contribute to the nonlinear response^{21,43,44}, which is 192 significantly suppressed by PT-symmetry^{43,45}. Furthermore, we can 193 exclude its contribution based on the scaling analysis discussed 194 subsequently. Therefore, the intrinsic contributions to the nonlinear 195 conductivity in MnBi₂Te₄ are determined by the nonlinear Drude weight 196 and the normalized quantum metric dipole. 197

To verify the dominant contribution from the quantum metric, we 198 investigate the scaling relationship between nonlinear conductivities 199 $(\sigma_{yxx}^{2\omega}$ and $\sigma_{xxx}^{2\omega}$) and the linear conductivity σ_{xx}^{ω} (i.e. scattering time τ) 200 as a function of temperature. Fig. 3c shows the temperature dependence of 201 longitudinal conductivity σ_{xx}^{ω} and charge carrier density n_e . The charge 202 carrier density n_e remains almost unchanged at the temperature of 1.6 to 10 203 204 K, suggesting that the τ is the main component to determine the conductivity σ_{xx}^{ω} in this range. By adopting this range, we plot the 205 $\sigma_{yxx}^{2\omega}$ and $\sigma_{xxx}^{2\omega}$ as a function of conductivity σ_{xx}^{ω} , 206

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

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

dipole (η_0 , τ -independent term). The nonlinear Drude contribution (η_2 , 210 τ^2 -dependent term) is nearly negligible compared to the intrinsic one, 211 212 which is reasonable considering the small conductivity and low mobility in MnBi₂Te₄. The dominant τ -independent nonlinear conductivity further 213 214 exclude contribution from the impurity scattering, because its lowest order contribution to the nonlinear conductivity begins at τ^1 (see detailed 215 discussion in supplementary information S7.2)^{21,44}. In conclusion, we 216 217 attribute the nonlinear response observed in 4SL-MnBi₂Te₄ to the nonlinear Drude weight and the normalized quantum metric dipole, with 218 the latter being the dominant contributor. 219

220 Fermi energy dependent nonlinear response

We next study how the nonlinear conductivity in 4SL-MnBi₂Te₄, 221 originating from normalized quantum metric dipole, is influenced by the 222 223 vertical displacement electric field (denoted as D) and the electron charge carrier density (denoted as n_e). We first examine the effect of D on the 224 nonlinear response. Although PT symmetry is instrumental in isolating the 225226 quantum metric dipole contribution, the induced nonlinear response remains even when PT is broken by D. Furthermore, we note that the 227 Berry curvature dipole induced NLAHE still vanishes because of three-228 fold rotational symmetry. To demonstrate this, we investigate the 229 dependence of $\sigma_{yxx}^{2\omega}$ on the electric field D with fixed carrier density at 230 $n_e \approx -3 \times 10^{12} \ cm^{-2}$. As shown in Fig. 4b, $\sigma_{yxx}^{2\omega}$ changes only 231 slightly when tuning the electric field away from D=0. Next, we investigate 232 the effect of charge carrier density n on the nonlinear response. Fig. 4c 233 shows the nonlinear conductivity $\sigma_{yxx}^{2\omega}$ and $\sigma_{xxx}^{2\omega}$ as a function of carrier 234 density n, respectively. When the Fermi level is tuned into the charge 235 neutral gap, both $\sigma_{yxx}^{2\omega}$ and $\sigma_{xxx}^{2\omega}$ vanish. This is in line with the fact 236 that the quantum metric dipole is a Fermi surface property³⁰. Furthermore, 237 both $\sigma_{yxx}^{2\omega}$ and $\sigma_{xxx}^{2\omega}$ exhibit a sign reversal when tuning the carrier 238 density between hole and electron regimes. Lastly, given the evident 239 240 nonreciprocal longitudinal response, we can evaluate the nonreciprocity coefficient γ in MnBi₂Te₄. We find the coefficient γ can reach $\gamma \approx 7 \times$ 241 $10^{-11}m^2A^{-1}$, which is two or three orders larger than traditional heavy 242 243 metal (HM)/ferromagnetic metal (FM) heterostructures (See details in supplementary information section S4.3)^{46,47}. 244

To explain our experimental data, we evaluate $\sigma_{vxx}^{2\omega}$ 245 quantitatively by constructing a slab model for 4SL MnBi₂Te₄ with density-functional theory 246 (DFT) calculations. Fig. 4d shows the total conductivity as a function of 247 carrier density, in which we assume $\sigma_{yxx}^{2\omega}$ for an ideally aligned device (y 248 aligned with the in-plane crystalline axis). Importantly, around the charge 249 neutrality gap ($\mu = 0$), we observe a sign change which reproduces the 250 experimental findings. (see supplementary information S7.1 for the band-251252 resolved contributions to the quantum metric) The deviation of theoretical $\sigma_{vxx}^{2\omega}$ compared to experimental values may be attributed to the known 253 discrepancy in surface band structure between calculations 254 and experiments on MnBi₂Te₄. Additionally, in our calculation, we find that 255256 the Drude contribution is small within a large energy range around the Fermi level, indicating that the nonlinear conductivity is primarily driven 257 by the quantum metric dipole. We conducted similar measurements on 258 259 another 6SL-MnBi₂Te₄ device, as detailed in supplementary information 260 section S6.

As a probe, the quantum metric dipole-driven electron nonlinearity in 261 MnBi₂Te₄ has several advantages and holds great promise for future 262 applications. First, compared to nonlinearity originating from the Berry 263 curvature dipole or skew scattering, the quantum metric dipole-driven 264 265 electron nonlinearity is independent of the scattering time, suggesting its 266 robustness against disorder scattering. Second, the quantum metric dipoledriven electron nonlinearity is distinct for opposite AFM spin orders and 267 even robust against small perturbations of external magnetic fields (see 268 details in supplementary information S3), making it a promising candidate 269 270 for use as a nonlinear magnetic memory device. Finally, the quantum 271 metric dipole-driven nonlinear response is more prominent than the 272 "traditional" nonlinear response from the Berry curvature dipole. We 273 compared the nonlinear response observed in MnBi₂Te₄ with other 2D material systems, as presented in Extended Table I. We find that our 274 MnBi₂Te₄ devices have a much larger nonlinear conductivity than WTe₂ 275 276 and strained WSe₂, only smaller than twisted WSe₂ and graphene Moiré 277 superlattices, making MnBi₂Te₄ a promising candidate for highly efficient rectifying devices^{9,10,13,16,19}. 278

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404 **Figure captions:**

405 Figure 1 Quantum metric induced nonlinearity in a *PT*-symmetric 406 **AFM. a.** The nonlinear Hall effect originating from Berry curvature dipole, 407 which has been observed in systems with broken inversion symmetry. **b**. The nonlinear longitudinal and transverse response observed in a PT-408 symmetric system due to quantum metric dipole. c. The crystal structure of 409 even-layer MnBi₂Te₄ from the side view. The arrows represent the spin 410 momentum direction of the Mn atoms. d. The spatial reflectance magnetic 411 circular dichroism (RMCD) mapping of MnBi₂Te₄ measured at zero 412 magnetic field. The dashed line marks the boundary between the 4SL-413 MnBi₂Te₄ and 3SL- MnBi₂Te₄. e. The crystal structure of MnBi₂Te₄ from 414 the top view. The dashed line marked with M represents the mirror line. f. 415 The angle dependent optical second-harmonic generation (SHG) of the 416 4SL-MnBi₂Te₄ sample. The zero degree corresponds to the applied current 417 direction in the device. 418

Figure 2 Observation of nonlinear transport in AFM MnBi₂Te₄. a. The 419 schematic view of the dual-gated 4SL-layer MnBi₂Te₄ device. The linear 420 and nonlinear signals are measured simultaneously with the denoted 421 direction. **b.** The linear longitudinal V_x^{ω} and transverse V_y^{ω} voltage as a 422 function of current I_x^{ω} . The solid line is the linear fitting of the data. **c and d.** The nonlinear longitudinal $V_x^{2\omega}$ and transverse $V_y^{2\omega}$ voltage as a 423 424 function of current V_x^{ω} for AFM-I and AFM-II states, respectively. At a 425 fixed electric field, the different AFM states are prepared by: AFM-I, 426 sweeping the magnetic field from -7 T to 0 T; AFM-II, sweeping the 427 428 magnetic field from +7 T to 0 T. The solid lines are quadratic fits to the 429 data. e. The optical image of another 4SL-MnBi₂Te₄ device with radially distributed electrodes. The dashed line indicates the in-plane crystalline 430 axis, determined by SHG measurement. scale bar: 5 μ m. f. The nonlinear 431 transverse $V_y^{2\omega}$ and longitudinal $V_x^{2\omega}$ response as a function of current 432 injection angle θ . 433

Figure 3 Spin order related electron nonlinearity from band-434 **normalized quantum metric dipole. a.** Nonlinear transverse voltage $V_v^{2\omega}$ 435 of 4SL-MnBi₂Te₄ as a function of temperature for opposite AFM-I and 436 AFM-II states, respectively. The amplitudes of $V_y^{2\omega}$ are similar for the 437 two AFM states but their sign is reversed. $V_y^{2\omega}$ vanishes when the 438 temperature is above the Néel temperature of MnBi₂Te₄. **b.** The nonlinear 439 longitudinal voltage $V_x^{2\omega}$ as a function of temperature for opposite AFM-440 I and AFM-II states, respectively, exhibiting similar trend as $V_v^{2\omega}$. c. The 441 conductivity and fitted carrier density of 4SL-MnBi₂Te₄ at different 442 temperatures. d. The scaling relationship between the nonlinear 443

conductivity $\sigma_{yxx}^{2\omega}$ ($\sigma_{xxx}^{2\omega}$) and the square of the linear longitudinal conductivity σ_{xx}^2 . The scaling is carried out at the temperature range of 2 444 445 -10 K (marked with shadow area in c), in which the carrier density nearly 446 remains constant. The nonlinear transverse and longitudinal conductivities 447 are extracted by $\sigma_{yxx}^{2\omega} = \frac{V_y^{2\omega}}{(I_x^{\omega})^2 R_{xx}^3} \frac{L^3}{W^2}$ and $\sigma_{xxx}^{2\omega} = \frac{V_x^{2\omega} L}{(I_x^{\omega})^2 R_{xx}^3}$, respectively. 448 The dashed line is a fit of the data with the scaling relation $\sigma^{2\omega} =$ 449 $\eta_2(\sigma_{xx}^{\omega})^2 + \eta_0$. From the fitting, we conclude that the predominant 450 contribution is from the η_0 term, namely from the intrinsic nonlinear 451 conductivity due to the band-normalized quantum metric dipole. 452

453 Figure 4 The electric field and carrier density dependence of the nonlinear response. a. The resistance map of the 4SL-MnBi₂Te₄ device. 454 455 The vertical displacement field D and charge carrier density n_e can be independently controlled, as denoted in the figure. b. The nonlinear 456 transverse conductivity $\sigma_{yxx}^{2\omega}$ and longitudinal resistance R_{xx} as a function 457 of vertical displacement electric field. The carrier density is fixed at $n_e \approx$ 458 -3×10^{12} cm⁻² as D varies along the dashed line in **a**. **c**. The 459 measured nonlinear transverse conductivity $\sigma_{yxx}^{2\omega}$ and longitudinal 460 conductivity $\sigma_{xxx}^{2\omega}$ as a function of charge carrier density $n_{e.}$ **d.** The 461 calculated total nonlinear transverse conductivity $\sigma_{yxx}^{2\omega}$ (red line) as a 462 function of carrier density. The grey line denotes the Drude contribution. 463

464 Methods

465 MnBi₂Te₄ single Crystal growth

Single crystals of MnBi₂Te₄ were grown using a self-flux method^{48,49}. The 466 starting materials used in the single crystal processes are Mn slices, Bi 467 lumps, and Te chunks. MnTe and Bi₂Te₃ precursors were prepared by 468 reacting the mixed stoichiometric starting materials at 1100 °C and 900 °C 469 470 for 24 h, respectively. Then MnTe and Bi₂Te₃ were mixed according to the ratio MnTe: $Bi_2Te_3 = 15$: 85, loaded into an alumina crucible, and sealed 471 in a quartz tube. The quartz tube was heated to 650 °C in 10 h, dwelled for 472 473 12 h, and slowly cooled to 595 °C at a rate of 1 °C/h to grow the single crystals. Shiny single crystals with a typical size of $3 \times 2 \times 0.5$ mm³ can be 474 obtained after centrifuging. 475

476

477 **Device fabrication**

The MnBi₂Te₄ thin flakes were obtained using the Al₂O₃-assisted 478 exfoliation method^{35,50}. First, a 70 nm thick layer of Al₂O₃ was thermally 479 evaporated onto bulk MnBi₂Te₄ crystals. Second, the Al₂O₃ thin film was 480 lifted along with thin flakes of MnBi₂Te₄, which were cleaved from the 481 bulk crystal using thermal release tape. Then, the stacked MnBi₂Te₄/Al₂O₃ 482 was transferred onto a transparent Polydimethylsiloxane (PDMS) film. The 483 484 number of layers in the MnBi₂Te₄ thin flakes was determined using optical transmittance measurements. After confirming the layer number, the 485 MnBi₂Te₄ thin flake samples were transferred to a Si wafer coated with 486 487 285 nm of SiO₂. To ensure high-quality samples, we used a stencil mask method to deposit metal contacts (Cr/Au) on the samples. A 20-40 nm thick 488 h-BN flake was then transferred to the MnBi₂Te₄ devices as the top gate 489 490 dielectric layer, followed by the transfer of a graphite thin flake gate on top of the h-BN/MnBi₂Te₄ heterostructure. The entire device fabrication 491 process was carried out in a nitrogen-filled glove box with O₂ and H₂O 492 493 levels below 1 ppm.

494

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

low frequency of 17.777 Hz. During the transport measurements, the phase 501 of the first and second harmonic signals was confirmed to be 502 approximately 0 and 90 degrees, respectively (see supplementary 503 information S2.4). The gate voltages were applied using a Keithley 2636 504 SourceMeter. To independently control the charge carrier density n_e and 505 displacement electric field D, we used a dual gated device structure. The 506 charge carrier density *n* can be obtained by $n = \left(\frac{\varepsilon_0 \varepsilon^{BN} (V_{TG} - V_{TG0})}{d_{BN}} + \right)$ 507 $\varepsilon_0 (V_{BG} - V_{BG0}) / \left(\frac{d_{Al_2O_3}}{\varepsilon^{Al_2O_3}} + \frac{d_{SiO_2}}{\varepsilon^{SiO_2}} \right) / e$. Here we regarded the Al₂O₃ and SiO₂ 508 dielectric layer as two series connected capacitors. The electric 509 displacement field D can be obtained by $D = \left(\frac{V_{BG} - V_{BG0}}{\left(\frac{d_{Al_2O_3}}{\epsilon^{Al_2O_3}} + \frac{d_{SiO_2}}{\epsilon^{SiO_2}}\right)}\right)$ 510

 $\frac{\varepsilon^{BN}(V_{TG}-V_{TG0})}{d_{DN}})/2$. Here, ε_0 is the vacuum permittivity; V_{TG0} and V_{BG0} 511 correspond to the gate voltage of maximum resistance; $\varepsilon^{Al_2O_3}$, ε^{SiO_2} and 512 ε^{BN} are the relative dielectric constant of Al₂O₃, SiO₂ and h-BN; $d_{Al_2O_3}$, 513 d_{SiO_2} and d_{BN} are the thickness of Al₂O₃, SiO₂ and h-BN, respectively. 514 We noticed a minor asymmetry in $\sigma_{yxx}^{2\omega}$ and R_{xx} with respect to the 515 displacement electric field D, possibly attributed to charge trapping 516 induced by substrate effects or disorder⁵¹. The asymmetry is more 517 pronounced in $\sigma_{yxx}^{2\omega}$, as the quantum metric is related to energy differences 518 between bands and more sensitive to D. 519

The nonlinear response in the MnBi₂Te₄ device with radially distributed devices

For the MnBi₂Te₄ device with radially distributed devices, when the 522 current aligns with the in-plane crystal axis ($\theta = 0^{\circ}$), the transverse $V_{v}^{2\omega}$ 523 is zero. In contrast, when the current is perpendicular to the crystalline axis 524 $(\theta = 90^{\circ})$, the longitudinal $V_x^{2\omega}$ is zero. In addition, the nonlinear 525 response should obey the symmetry of even-layer MnBi₂Te₄, with $\sigma_{yxx} =$ 526 $-\sigma_{yyy}$ and $\sigma_{xyy} = -\sigma_{xxx}$. To substantiate this, we use the formula for 527 nonlinear conductivity: $\sigma_{xxx}^{2\omega} = \frac{J_x^{2\omega}}{(E_x^{\omega})^2} = \frac{V_x^{2\omega} L}{(I_x^{\omega})^2 R_{xx}^3}$ and $\sigma_{yxx}^{2\omega} = \frac{J_y^{2\omega}}{(E_x^{\omega})^2} =$ 528 $\frac{V_y^{2\omega}}{(I_w^{\omega})^2 R_{xx}^3} \frac{L^3}{W^2}$. Here, L refers to the length and W to the width of the device. 529 The optical image (Fig. 2e) reveals that the length-to-width ratio, L/W, is 530 consideration 0.85. Taking into 531 approximately the geometric 532 characteristics of the device, we conclude that the nonlinear conductivity

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

534 **Optical measurement**

For the RMCD measurement, a linearly polarized 633 nm HeNe laser was 535 modulated by a photoelastic modulators (PEM) at a frequency of 50 kHz. 536 537 The laser beam was focused on the sample at normal incidence using an Olympus MPLN50× objective with a 0.75 numerical aperture. The laser 538 spot size was approximately 2 µm, with a power of 1.8 µW. The reflected 539 light was collected by a photodetector and analyzed using a lock-in 540 amplifier set to the same frequency as the PEM. The RMCD measures the 541 differential reflection between left and right circularly polarized light, the 542 magnitude of which is proportional to the magnetic moments of the 543 544 MnBi₂Te₄ sample.

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 polarizationresolved measurement, a motorized polarizer was used to control the polarization of the incident laser beam as it was rotated next to the objective.

552 **DFT calculations**

We use VASP (Vienna ab-initio software package) with the PBE 553 functional in the generalized gradient approximation, to obtain the 554 electronic ground state⁵². We then project the ground state wavefunctions 555 on Wannier functions using Wannier 90⁵³ and create a tight-binding model 556 with 184 Wannier orbitals. For the Drude contribution we adopt a lifetime 557 of $\tau \approx 40$ fs estimated from our experimental conductivity and carrier 558 density in Fig. 2c and theoretical effective mass $(m^* = 0.14m_{e})$ for the 4 559 560 SL film.

561

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Author contributions: W.-b.G and B.Y conceived and supervised the 596 597 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 598 the theoretical calculations. N.W and F.Z performed the NV measurement 599 600 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. 601 K.W and T.T grew the hBN single crystals. N.W, B.Y and W.-b.G wrote 602 603 the manuscript with input from all authors.

604 **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
be addressed to Binghai Yan (binghai.yan@weizmann.ac.il) or Wei-bo
Gao (wbgao@ntu.edu.sg).

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612 Extended Figure 1 The linear conductivity of 4SL-MnBi₂Te₄ with 613 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 0 T, respectively. 615 c and d. The linear longitudinal V_x^{ω} and transverse V_y^{ω} voltage as a 616 function of current I_x^{ω} for AFM I and AFM II states, respectively. The solid 617 line is a linear fit to the experimental data. 618 Extended Figure 2 The fully compensated AFM order in 4SL-

- 619 MnBi₂Te₄ device. A. The magnetic field dependent longitudinal resistance
- 620 R_{xx} of the 4SL-MnBi₂Te₄ device. **B.** The magnetic field dependent Hall
- resistance R_{yx} of the 4SL-MnBi₂Te₄ device. In zero magnetic field, the
- 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
- transverse conductivity $\sigma_{yxx}^{2\omega}$ in even-layer MnBi₂Te₄ is summarized and compared with other reported material systems^{9,10,13,16,19}. The
- 627 corresponding mechanisms responsible for the nonlinear response are also
 628 listed for comparison.











Extended Data Fig. 1



Material	<i>T</i> (K)	$\sigma_{xx}^{1\omega}(10^{-3}\mathrm{S})$	$\sigma_{yxx}^{2\omega}$ (mA nm V ⁻²)	Origin
WTe ₂ (bilayer) ¹⁰	10	0.15	940	
WTe ₂ (few-layer) ⁹	2	2.1	3	BCD
Strained WSe ₂ ¹³	140	~ 0.41	~ 1200	
Twisted WSe ₂ ¹⁹	1.5	~ 0.3	~ 18000	
h-BN/graphene/h-BN ¹⁶	1.7	17.8	1.05×10^{7}	Skew scattering
MnBi ₂ Te ₄ (4-SL)	2	0.15	400	Quantum metric
MnBi ₂ Te ₄ (6-SL)	2	0.5	2400	dipole

PRENIEM

Extended Data Table 1

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