



# One-dimensional weak antilocalization in single-crystal $\text{Bi}_2\text{Te}_3$ nanowires

Wei Ning<sup>1,2</sup>, Haifeng Du<sup>1,2</sup>, Fengyu Kong<sup>1,2</sup>, Jiyong Yang<sup>1,2</sup>, Yuyan Han<sup>1,2</sup>, Mingliang Tian<sup>1,2,3</sup> & Yuheng Zhang<sup>1,2</sup>

<sup>1</sup>High Magnetic Field Laboratory, Chinese Academy of Science, Hefei 230031, Anhui, P. R. China, <sup>2</sup>Hefei National Laboratory for Physical Science at The Microscale, University of Science and Technology of China, Hefei 230026, People's Republic of China, <sup>3</sup>The Center for Nanoscale Science and Department of Physics, The Pennsylvania State University, University Park, Pennsylvania 16802-6300.

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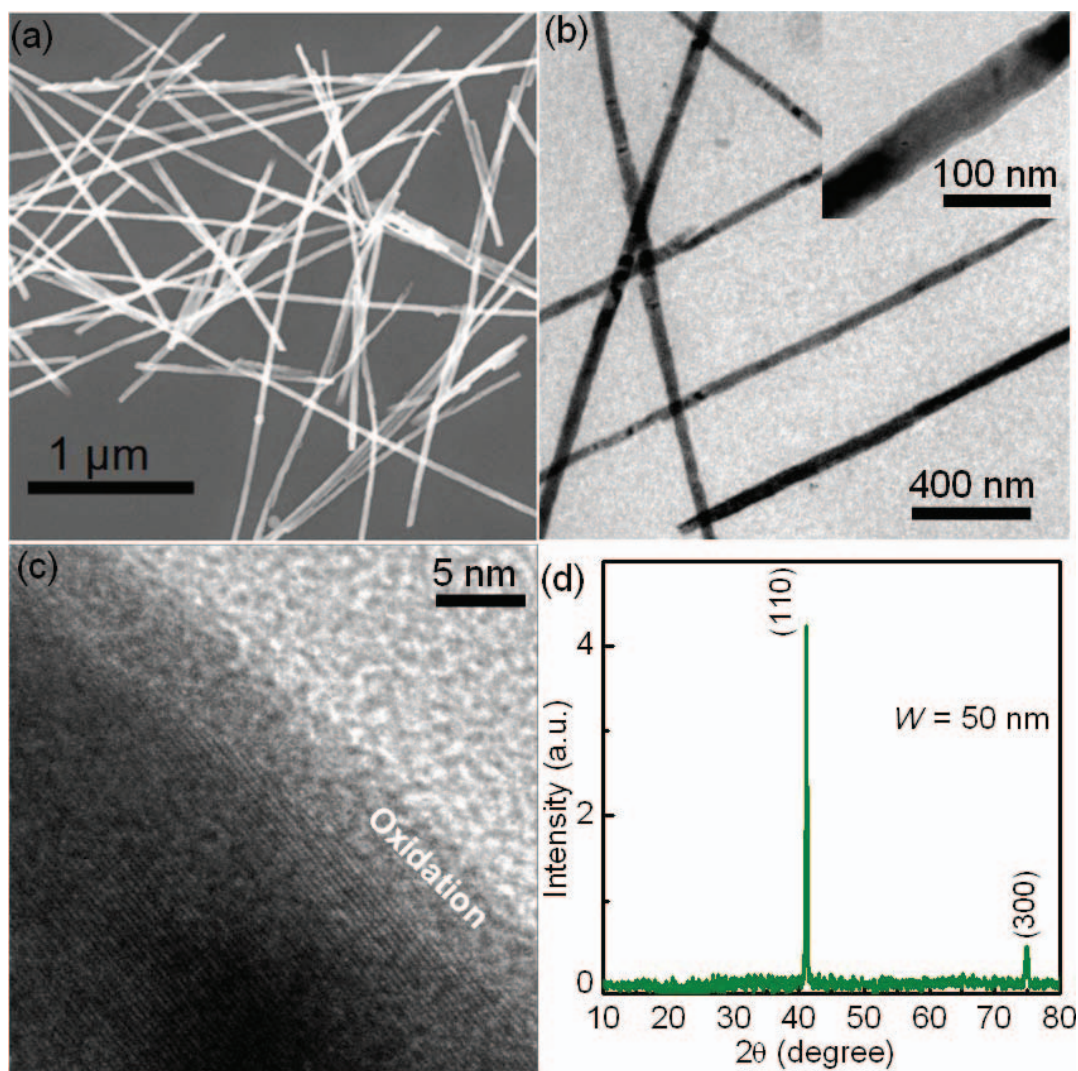
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Correspondence and  
requests for materials  
should be addressed to  
M.T. (tianml@hmf.ac.  
cn)

Angle-dependent magnetoconductance was measured on an individual surface-curved  $\text{Bi}_2\text{Te}_3$  single-crystal nanowire fabricated by electrochemical deposition, where the evolution of surface conduction with wire diameters was investigated. It was found that the magnetoconductance of these nanowires in low field regime can be well described by one-dimensional (1D) weak antilocalization (WAL) model, where the dephasing length of the electrons follows  $T^{-1/3}$  dependence but insensitive to the wire diameter. Meanwhile, such a 1D surface WAL was found to be enhanced significantly with the decrease of the wire diameter.

Topological insulator (TI) is a new state of matter. It is characterized by a full insulating gap in the bulk and gapless spin-resolved surface state (SS) in which the spin of the electrons is locked perpendicular to its momentum by the strong spin-orbit interaction. These spin helical properties of Dirac fermions in SS have made them relevant for exotic new physics<sup>1–3</sup> and applications on improved spintronic devices and potentially useful for quantum computing<sup>4–7</sup>. In three-dimensional (3D) TIs, the topological surface states have been proved by angle-resolved photoemission spectroscopy (ARPES)<sup>8,9</sup>, scanning tunneling microscopy<sup>10,11</sup> and electrical transport measurements including two-dimensional (2D) weak antilocalization (WAL)<sup>12–17</sup>, Shubnikov-de Haas (SdH) or quantum Hall (QH) effect<sup>18</sup>, where a 2D character was determined due to a fact that all of these effects were found only sensitive to the perpendicular component of the magnetic field. However, direct probing the surface state of an individual nanowire by ARPES is almost impossible because of the limitation of sample size. Instead, electrical transport measurement become the most efficient way to obtain the surface information due to its unique geometry and extreme large surface to volume ratio<sup>19,20</sup>. Several quasi-1D nanoribbons have been synthesized, including  $\text{Bi}_2\text{Se}_3$ ,  $\text{Bi}_2\text{Te}_3$  and  $\text{Bi}_2(\text{Se}_x\text{Te}_{1-x})_3$ <sup>21–25</sup>, where the Aharonov-Bohm and Atshuler-Aronov-Spivak<sup>20,21</sup>, Aharonov-Casher effect<sup>22</sup> and 2D WAL of the SS<sup>23,25</sup> were found. Compared with the TI nanoribbons with two well-defined parallel surfaces, the nature of the surface state in a surface-curved 1D cylindrical nanowire has not been well explored experimentally although a number of unique and fascinating properties<sup>26–28</sup> have been predicted theoretically, such as the possible helical Luttinger liquid<sup>28</sup> that was ever described as the edge mode of the 2D quantum spin Hall (QSH) topological insulator.

In this work, we report angle-dependent magnetoresistance (MR) studies on an individual surface-curved  $\text{Bi}_2\text{Te}_3$  nanowire fabricated by electrochemical deposition. The evolution of the surface conduction with wire diameters was carefully investigated. It was found that the nanowires of different diameters showed similar positive MR behavior, where the variations of their magnetoconductance at different angles,  $\Delta G(\theta, B)$ , by subtraction of the magnetoconductance at  $\theta = 0^\circ$  can be surprisingly normalized to one curve with the perpendicular component,  $B_\perp = B \sin \theta$ , of the magnetic field, where  $\theta$  is the angle between  $B$  and the wire axis as shown schematically in the inset of Fig. 3(a). Such a behavior is a direct indication of the WAL of the surface states of nanowire. Theoretical analysis based on the WAL model showed that the dephasing length of electrons in these thin curved nanowires follows  $T^{-1/3}$  temperature dependence but insensitive to the wire diameter, indicating the 1D WAL character from the surface channels of the nanowire instead of the expected 2D WAL reported in nanoribbons<sup>23,25</sup> and bulk<sup>12–15</sup>. These results are also consistent with our observation that the upper limit for the range of the magnetic fields which is appropriate for fitting the observed 1D WAL increases significantly with the decrease of the wire diameter, *i.e.*, an enhancement of quasi-one dimensionality in electrical transport of surface states in small diameter wire.



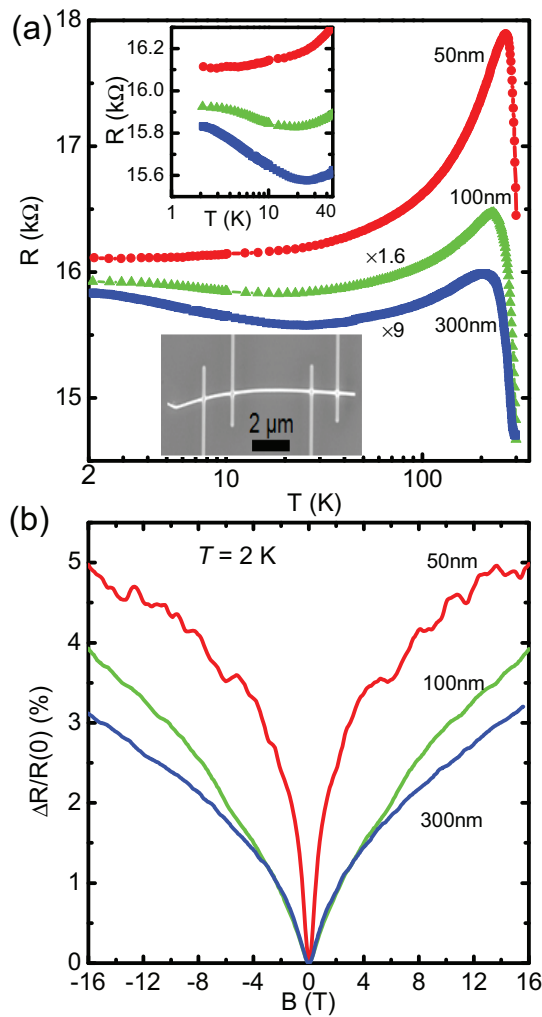
**Figure 1** | (a) and (b) are respectively the low magnification SEM and TEM images of nanowires with  $W = 50$  nm. The inset of (b) shows an enlargement of an individual nanowire. (c) High-resolution TEM image of 50 nm wire selected randomly. An oxidation layer of approximately  $\sim 5.0$  nm is visible on the surface. (d) x-ray diffraction spectra carried out on the nanowire arrays, where the (110) diffraction is the dominant peak in the spectra.

## Results

All of  $\text{Bi}_2\text{Te}_3$  nanowires used in this work were synthesized by electrochemical deposition via customer-made porous anodic aluminum oxide (AAO) membrane. The wire diameter was exactly controlled by the pore size of the template<sup>29</sup> so that we are able to carry out a systematic comparative study on wires of various diameters by selecting the templates with proper pore size. While the diameter of each wire along its length is uniform, it shows variations from wire to wire within an uncertainty of  $\sim 20\%$ , verified by both scanning electron microscopy (SEM) and transmission electron microscopy (TEM). It was shown that the majority of the nanowires have a length of  $8\sim 15\ \mu\text{m}$  with single-crystal morphology as shown in Figs. 1 (a), 1(b) and 1(c). An oxidation layer of approximately  $\sim 5.0$  nm, which was probably formed after the nanowires were released from the membrane, is visible on the surface. All samples of different diameters, such as 300 nm, 100 nm and 50 nm fabricated under the same controlled conditions showed very similar structure, verified by x-ray diffraction measurements on the nanowire arrays (see Fig. 1d), where the (110) diffraction is the dominant peak in the spectra.

Standard 4-probe transport measurements were carried out on an individual nanowire using a physical property measurement system (PPMS). Four platinum (Pt) strips of 100 nm (width)  $\times$  100 nm (thickness) as shown in the lower inset of Fig. 2 (a) were deposited

onto the wire as the contact electrodes using FEI Helios NanoLab 600i FESEM/FIB dual beam system. Resistance as a function of temperature for three individual  $\text{Bi}_2\text{Te}_3$  nanowires of 300 nm, 100 nm and 50 nm showed very similar transport behavior in high temperature range (see Fig. 2a), such as a resistance hump near 220 K and a subsequent metallic behavior down to  $\sim 25$  K. We noticed that the hump behavior of the resistance usually appears in TI-samples with the reduction of bulk carrier density less than  $10^{17}/\text{cm}^3$  by element-doping (e.g.,  $\text{Na}^{30}$ ,  $\text{Pb}^{16}$  or  $\text{Sb}^{18}$  doping and so on), where the surface contribution to the total conduction become significant. Therefore, the similar hump behavior observed in our nanowires might have the same origin due to the competition of conduction between the surface and bulk. When the temperature decreases, the contribution of the bulk conductivity decreases sharply and thus the surface conductivity will be dominant in low- $T$  range. Below about 25 K, both larger diameter wires ( $W = 100$  nm and 300 nm) show slightly an upturn in resistance while the small diameter wire ( $W \sim 50$  nm) displays a plateau down to 2 K. It was also noted that the upturn behavior becomes more significant with the increase of the wire diameter, as shown in the upper inset of Fig. 2 (a). The tendency of the weakened insulating feature with the decrease of the wire diameter at low temperatures means that the contribution of the metallic surface states to the total conduction might be enhanced in such a thinner nanowire.



**Figure 2** | (a) Resistance versus temperature for three nanowires of  $W = 300$  nm,  $100$  nm and  $50$  nm. The upper inset shows the resistance in low temperatures. The lower inset is the SEM image of the four-probe configuration of wire for transport measurement. (b) Normalized magnetoresistance in perpendicular magnetic field for  $W = 50$  nm,  $100$  nm and  $300$  nm wires. The MR oscillations superimposed on the positive MR curves were observed. The amplitude of the MR oscillations was enhanced when the diameter decreases from  $300$  nm to  $50$  nm.

Fig. 2 (b) shows the normalized resistance as a function of the magnetic field aligned perpendicular to the axis of the nanowires at  $T \sim 2$  K. The MR quantum oscillations or fluctuations were seen in all three diameter wires but the amplitude of the oscillations in  $50$  nm wire was much larger than those of  $100$  nm or  $300$  nm wires, which is consistent with our expectation that the metallic surface contribution is probably enhanced in thinner nanowires due to the high ratio of the surface to bulk volume. The details related to the nature of these quantum oscillations in high field regime has been discussed elsewhere<sup>31</sup>. Here we focused our attention on the MR behavior in low-H regime.

The overall MR behavior as shown in Fig. 2 (b) is very similar to those reported in TI-bulk, films and nanoribbons, namely all three wires display a positive MR over entire magnetic field regime we measured and a valley in low-H range. Such a positive MR in TIs was usually attributed to the weak antilocalization (WAL) of the surface states because of the strong spin-orbit coupling and the helicity of the surface states<sup>32,33</sup>. The WAL is a negative quantum correction to the classical weak localization due to the wave nature of the electrons in a disordered metallic system.

## Discussion

To quantitatively understand the size dependent properties of the MR in these thin surface-curved TI nanowires, Figure 3 (a), 3(c) and 3(e), respectively, show the angle-dependent magnetoconductance,  $G(\theta, B) = 1/R(\theta, B)$ , of three samples with diameters of  $W = 300$  nm,  $100$  nm and  $50$  nm. It was known that the WAL induced by 2D surface states is characterized by a sole dependence on the perpendicular component,  $B_{\perp} = B \sin(\theta)$ , of the applied field. The fact is that, as usually expected the magnetoconductance of surface state at  $\theta = 90^\circ$ , the magnetoconductance at  $\theta = 0^\circ$  was also observed, indicating that bulk electrons also contribute to the total conductance. Therefore, to extract the pure surface state contribution, we subtract the 3D WAL contribution at  $\theta = 0^\circ$  from the total magnetoconductance at various angles, i.e.,  $\Delta G(\theta, B) = G(\theta, B) - G(0, B)$ , with  $G(\theta, B) = 1/R(\theta, B)$  and  $G(0, B) = 1/R(0, B)$ . Figs. 3(b), 3(d) and 3(f) show  $\Delta G(\theta, B)$  versus  $B_{\perp} = B \sin(\theta)$  for nanowires with  $W = 300$  nm,  $100$  nm and  $50$  nm, respectively. The surprising feature is that all of the traces at various angles can be approximately normalized to a single curve in low field regime although the angle  $\theta$  defined here has an uncertainty of at least  $\pm 5$  degree. These results clearly indicate that the magnetoconductance shown in Figs. 3 (b), 3(d) and 3(f) is mainly a result of the surface conductivity.

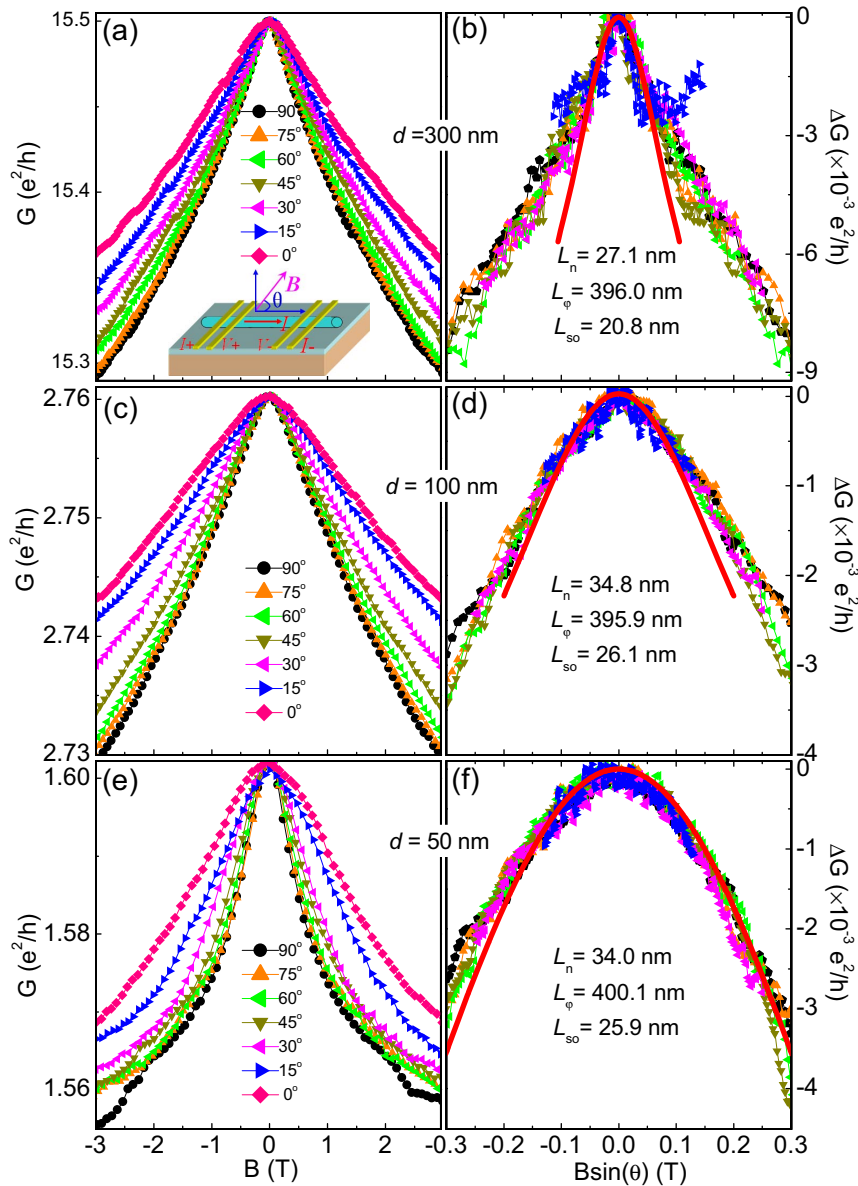
We now turn to a more quantitative analysis of the WAL behavior. As we noticed from Fig. 3 that the magnetoconductance of the surface states in  $300$  nm large diameter wire display a cusp-like shape while a parabolic-like shape was obtained with the decrease of the wire diameter. The cusp-like shape of the surface magnetoconductance was always seen in bulk and thin film TIs, where a 2D WAL dominates in the surface state. Because the typical dephasing length in TIs is usually on the order of  $L_{\phi} \sim 330$  nm<sup>14</sup>, which is larger than the diameter of our  $\text{Bi}_2\text{Te}_3$  nanowires ( $d = 300 \sim 50$  nm), a quasi-1D localization theory was used to fit the observed WAL. In a 1D system, the weak localization correction at a magnetic field  $B$  that includes the electron-electron (e-e) interactions and spin-orbit (s-o) interactions to the conductance is given by<sup>34,35</sup>:

$$\Delta G = \frac{\sqrt{2}e^2}{\pi h} \frac{L_n}{L} \left[ \frac{3}{2} \frac{A_i(\frac{2L_n^2}{L_1})}{A'_i(\frac{2L_n^2}{L_1})} - \frac{1}{2} \frac{A_i(\frac{2L_n^2}{L_2})}{A'_i(\frac{2L_n^2}{L_2})} \right] \quad (1)$$

where  $\Delta G$  is the conductance quantum correction in 1D,  $L$  and  $L_n$  are the wire length and the Nyquist length, respectively,  $A_i$  represents the Airy function,  $L_1 = (4/3L_{so}^2 + L_{\phi}^2)^{-1/2}$ , and  $L_2 = (1/L_{\phi}^2 + W^2/3L_{so}^2)^{-1/2}$ . Here,  $L_{so} = \sqrt{D\tau_{so}}$  is the spin-orbit length that characterizes the strength of spin-orbit coupling, where  $D$  is the diffusion constant.  $L_{\phi} = \sqrt{D\tau_{\phi}}$ ,  $L_H = (\hbar/eB)^{1/2}$  are the dephasing length and the magnetic length,  $W$  is the diameter of the wire.

Considering the oxide layer of nanowires, the actual widths,  $W$ , of the three wires quoted by  $300$  nm,  $100$  nm, and  $50$  nm were selected as  $290$  nm,  $90$  nm and  $40$  nm in the theoretical fitting, respectability. The solid red curves in Fig. 3(b), 3(d) and 3(f) are the fitting results with 1D localization theory, yielding  $L_n = 27.1$  nm,  $L_{\phi} = 396.0$  nm,  $L_{so} = 20.8$  nm for nanowire of  $300$  nm,  $L_n = 34.8$  nm,  $L_{\phi} = 395.9$  nm,  $L_{so} = 26.1$  nm for nanowire of  $100$  nm, and  $L_n = 34.0$  nm,  $L_{\phi} = 400.1$  nm,  $L_{so} = 25.9$  nm for nanowire of  $50$  nm. It was noted that  $L_{so} < L_{\phi}$ , which means the spin-orbit scattering time  $\tau_{so}$  is significantly shorter than  $\tau_{\phi}$ , being consistent with the results obtained from other similar topological insulators<sup>17,36</sup>. Obviously, the fitting is more successful for small diameter wire, and the upper limit for the range of the magnetic field which is appropriate for the 1D WAL increases significantly with the decrease of the wire diameter. Since the dephasing length  $L_{\phi}$  shows no clear diameter dependence, indicating the surface nature of the 1D WAL (If the bulk contribution to the WAL effect were significant,  $L_{\phi}$  should change with the diameter). These observations are consistent





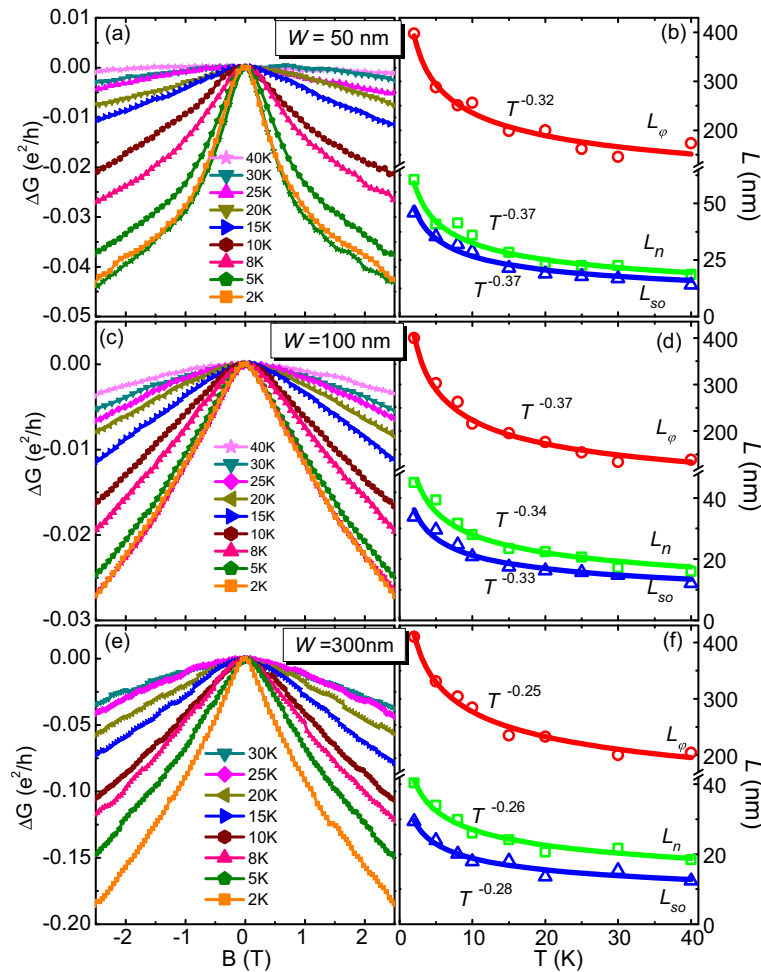
**Figure 3** (a), (c) and (e) are respectively the angle-dependent magnetoconductance of  $\text{Bi}_2\text{Te}_3$  nanowires ( $W = 300$  nm,  $100$  nm and  $50$  nm) measured at  $2$  K. The magnetoconductance persists at  $\theta = 0^\circ$ , indicating the contribution of bulk electrons to the WAL due to the bulk spin-orbit coupling. Inset: schematic of the measurement setup where  $\theta$  denotes the angle between the direction of the magnetic field and the current flow in the nanowires. (b), (d) and (f) are respectively the magnetoconductance change plotted in the perpendicular component,  $B\sin\theta$ , of the magnetic field for nanowires of  $W = 300$ ,  $100$  and  $50$  nm by subtracting the bulk contribution of the WAL at  $\theta = 0^\circ$ . The solid red lines are the fit with 1D localization theory. All the traces were normalized to a single curve in low field range, indicating the surface nature of the magnetoconductance. The upper limit for the range of the magnetic field which is appropriate for fitting the 1D WAL increases significantly with the decrease of the wire diameter.

with our expectation that quasi-one dimensionality in electrical transport of surface state is significantly enhanced in small diameter wire. A deviation of the 1D WAL fitting for  $300$  nm wire can be understood from a fact that the  $300$  nm is almost comparable to the phase coherence length,  $L_\phi$ . We have also tried the fitting with 2D WAL model for  $300$  nm wire, but unfortunately we obtained unphysical small fitting parameters compared with those reported for TI- bulk or film samples. A possible reason might be related to its cylindrical geometry with a curved surface or the diameter of  $300$  nm is just on the order of between 1D and 2D dimensionality.

To further investigate the observed WAL, the magnetoconductance as a function of temperature was also measured in several other  $\text{Bi}_2\text{Te}_3$  nanowires of different diameters ( $W = 50$  nm,  $100$  nm and  $300$  nm) in the field aligned perpendicular to the wire axis, as shown in Figs. 4 (a), 4(c) and 4(e), respectively. As  $T$  increases, the cusp of

the magnetoconductance in low field range is gradually broadened and finally disappears. Fitting the data at different  $T$  with eq.(1), the observed characteristic lengths of  $L_n$ ,  $L_\phi$  and  $L_{so}$  as a function of temperatures are shown in Fig. 4 (b), 4(d) and 4(f) for nanowires with  $W = 50$  nm,  $100$  nm and  $300$  nm, respectively. All characteristic lengths decrease with the increase of temperatures due to the increased thermal scattering.

It was well-known that the Nyquist length  $L_n$  and coherence length  $L_\phi$  are proportional to  $T^{-1/2}$  for the two dimensional system and  $T^{-1/3}$  for the one-dimensional one<sup>24,30,37</sup>. Fig. 4(b) shows that the Nyquist length  $L_n$ , coherence length  $L_\phi$  and spin-orbit length  $L_{so}$  for  $50$  nm wire are proportional to  $T^{-0.37}$ ,  $T^{-0.32}$  and  $T^{-0.37}$  which are very close to the exponent,  $-1/3$ , expected for the one-dimensional system, i.e., the electron conduction in  $\text{Bi}_2\text{Te}_3$  nanowires follows the quasi-one-dimensionality. Similar fitting were also done for nanowires with



**Figure 4** | (a), (c) and (e) respectively show magnetoconductance at different temperatures with magnetic fields aligned perpendicular to the wire axis for nanowires with  $W = 50$  nm,  $100$  nm and  $300$  nm. (b), (d) and (f) respectively show the characteristic lengths of  $L_n$ ,  $L_\phi$  and  $L_{so}$  as a function of temperatures obtained from the fitting with the 1D localization theory. The solid lines in each plot show the power-law dependence on temperatures, yielding  $L_n$ ,  $L_\phi$  and  $L_{so}$  decay as  $\sim T^{-0.37}$ ,  $T^{-0.32}$ ,  $T^{-0.37}$  for  $50$  nm wire, and  $T^{-0.34}$ ,  $T^{-0.37}$ ,  $T^{-0.33}$  for  $100$  nm wire, and  $T^{-0.26}$ ,  $T^{-0.25}$ ,  $T^{-0.28}$  for  $300$  nm wire. The exponent of  $L_n$  and  $L_\phi$  closes to  $-1/3$  expected for the one-dimensionality in  $\text{Bi}_2\text{Te}_3$  nanowires.

$W = 100$  nm and  $300$  nm. The obtained results indicate that  $L_n$ ,  $L_\phi$  and  $L_{so}$  decay as  $\sim T^{-0.34}$ ,  $T^{-0.37}$ , and  $T^{-0.33}$  for  $W = 100$  nm, and  $T^{-0.26}$ ,  $T^{-0.25}$ ,  $T^{-0.28}$  for  $W = 300$  nm, as shown in Fig. 4 (d) and 4 (f), respectively. It is noted that, as shown in Fig. 4(f),  $L_\phi < W$  at  $T > 10$  K for nanowires with  $W = 300$  nm, which indicates that the transport behavior of electrons is beyond quasi-one-dimensional channel, but two or three dimensionally above this temperature.

In summary, single crystal  $\text{Bi}_2\text{Te}_3$  nanowires with diameters arranging from  $50$  nm to  $300$  nm were fabricated by electrodeposition. Both MR oscillations and weak antilocalization (WAL) behavior were observed for nanowires with different diameters, and both properties were enhanced with the decrease of wire diameters, indicating the enhancement of surface states in thinner nanowires. The obtained dephasing length  $L_\phi$  shows independence of diameter but follows  $T^{-1/3}$  dependence, suggesting the quasi-one-dimensional nature of magnetoconductance in the surface curved  $\text{Bi}_2\text{Te}_3$  nanowires.

## Methods

**Synthesis.**  $\text{Bi}_2\text{Te}_3$  nanowires were fabricated by direct-current electrodeposition into porous anodic aluminum oxide membrane (AAO). The plating solution was prepared by dissolving  $10$  mM Te (powder: 99.9998% purity) and  $7.5$  mM Bi (shots: 99.9999% purity) elements into concentrated nitric acid (69.1%), and then the solution was diluted to  $1$  M final concentration with distilled water<sup>38</sup>. The electrodeposition was carried out at a constant potential of  $-62$  mV, relative to an Ag/AgCl reference electrode at room temperature. A pure Pt-wire was used as the counter electrode and Au film evaporated on one side of the AAO membrane prior to the electrodeposition

was used as the working electrode. Freestanding nanowires were obtained by dissolving the membrane with  $2$  M aqueous NaOH solution, followed by precipitating the wires via centrifugation.

**Device fabrication and characterization.** To make standard four-probe devices on an individual  $\text{Bi}_2\text{Te}_3$  NW for transport measurement, we dispersed the NWs on a silicon substrate with a  $1 \mu\text{m}$  thick  $\text{Si}_3\text{N}_4$  insulating layer. Then, the sample was transferred into FEI NanoLab 600i SEM/FIB dual beam system for the deposition of Pt-electrodes. The size of the four Pt-stripe electrodes has a nominal width of  $100$  nm and a thickness of  $100$  nm deposited. Usually, the film by FIB-induced Pt deposition is amorphous consisting of Pt, C, Ga or O depending on the deposition condition. While the maximum spreading or diffusion distance of the actual Pt along the wire is on the order of  $300$  nm, the distance ( $L$ ) between the inner edges of the two voltage electrodes for all of the studied samples was kept larger than  $4 \mu\text{m}$ , which is much longer than the spreading distance.

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## Author contributions

M.T. supervised the research. W.N., H.D., Y.H., F.K. and J.Y. synthesized the  $\text{Bi}_2\text{Te}_3$  nanowires, fabricated the devices and carried out the transport measurement in high magnetic field and performed structural analysis. Y.Z. contributed to the analysis. W.N. and M.T. wrote the paper, together with the help of all other co-authors.

## Additional information

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

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