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Direct comparison of current-induced spin polarization in topological insulator Bi₂Se₃ and InAs Rashba states

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Three-dimensional topological insulators (TIs) exhibit time-reversal symmetry protected, linearly dispersing Dirac surface states with spin-momentum locking. Band bending at the TI surface may also lead to coexisting trivial two-dimensional electron gas (2DEG) states with parabolic energy dispersion. A bias current is expected to generate spin polarization in both systems, although with different magnitude and sign. Here we compare spin potentiometric measurements of bias current-generated spin polarization in $Bi_2Se_3(111)$ where Dirac surface states coexist with trivial 2DEG states, and in InAs(001) where only trivial 2DEG states are present. We observe spin polarization arising from spin-momentum locking in both cases, with opposite signs of the measured spin voltage. We present a model based on spin dependent electrochemical potentials to directly derive the sign expected for the Dirac surface states, and show that the dominant contribution to the current-generated spin polarization in the TI is from the Dirac surface states.

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he quest for efficient generation and electrical control of spin has motivated the search for materials and structures that exhibit strong spin splitting of their electronic states^{1,2}. A successful platform has been the two-dimensional electron gas (2DEG) in semiconductor heterostructures, where structural inversion asymmetry along surface normal lifts spin degeneracy via spin-orbit coupling³. In 2DEGs, with a parabolic energy dispersion, the Rashba form of spin-orbit coupling leads to a pair of Fermi surfaces that exhibit counter-rotating chiral spin texture, locking spin to the linear momentum^{2,4–8}. The further demonstration of electrical gate control of the strength of such spin splitting has led to prospects for prototypical semiconductor field effect spintronic devices^{8,9}.

Helical spin-momentum locking is also exhibited in the recently discovered quantum phase of matter, three-dimensional topological insulators (TIs), where linearly dispersing metallic surface states populated by massless Dirac fermions coexist with a semiconducting bulk^{10–14}. The 2D Dirac states are occupied by a single spin, and topologically protected by time-reversal symmetry, making them robust against scattering. This helical spin texture has been observed by spin- and angle-resolved photoemission spectroscopy (ARPES)^{15–17}. These measurements have also shown that trivial Rashba spin-split 2DEG states may coexist with the Dirac surface states in TI materials that exhibit surface carrier accumulation arising from band bending^{18,19}.

An unpolarized bias current is predicted to create a net spin polarization due to spin-momentum locking for both the topologically protected TI surface states^{20–22} and the Rashba 2DEG states^{3,4}. The spin helicities of the two have been shown in momentum-resolved measurements such as spin-ARPES to be opposite¹⁸. In transport measurements, however, the measured spin polarization is momentum integrated, and both Dirac and Rashba 2DEG states can contribute to the spin voltage measured at the detector contact. Calculations treating a model Bi₂Se₃ surface in which these states coexist found that the spin polarization and sign of the corresponding spin voltage measured at the detector contact is indeed opposite for the TI Dirac state and Rashba 2DEG contributions^{23,24}. These calculations were performed for a three-terminal potentiometric geometry for both ballistic and diffusive regimes, using a spin-orbit coupling coefficient of $\alpha = 0.79 \text{ Å}$ obtained from ARPES measurements on Bi₂Se₃ (ref. 19). Furthermore, because the Rashba 2DEG states exists as spin-split pairs with Fermi level momenta k_1 and k_2 , where $\mathbf{k}_1 < \mathbf{k}_2$, their spin contribution given by $(\mathbf{k}_2 - \mathbf{k}_1)/(\mathbf{k}_2 + \mathbf{k}_1)$ mostly cancels, and the net spin polarization was shown to be dominated by the TI surface states 13,22,23.

We recently demonstrated the first direct electrical detection of spin polarization resulting from spin-momentum locking in TI surface states in Bi₂Se₃ using spin potentiometric measurements, where the projection of the current-generated spin onto the magnetization of a ferromagnetic/tunnel barrier detector contact

was measured as a voltage²⁵. Similar investigations on various TI materials using similar FM/tunnel contacts^{26–31} have subsequently been reported. However, inconsistent results were reported regarding the sign of the spin signal, perhaps due to variations in material quality, device structure and measurement geometry, or the potential coexistence of the two spin systems. Lee *et al.*²⁹ measure a spin–voltage consistent with ours^{25,31} in electrically gated (BiSb)₂Te₃ samples where the Fermi energy is systematically moved from the conduction band edge through the Dirac point to the valence band edge, while others report a spin voltage of opposite sign with a markedly different temperature dependence²⁸. This underscores the need to independently probe and compare the characteristics of the Dirac and Rashba systems.

In this work, we report a direct comparison of the current-induced spin polarization measured using identical Fe/Al₂O₃ tunneling spin-potentiometric contacts measurement geometries in two prototype systems: the TI Bi₂Se₃, where both Dirac surface states and Rashba 2DEG states are known to coexist; and InAs that exhibits only the Rashba 2DEG states. We show that the sign of the spin signal measured in the Bi₂Se₃ and InAs samples is indeed opposite, and the temperature dependence is markedly different. We further develop a model based on spin-dependent electrochemical potentials to explicitly illustrate the measurement and derive the sign of the spin voltage expected for the TI surface states, which corroborates our experimental observation here and in previous work²⁵. These results show that the dominant contribution to the bias current-generated spin voltage in the TI is from the Dirac surface states in spin potentiometric measurements. This direct electrical access of the helical spin texture of Dirac and Rashba 2DEG states, along with recent demonstration of electrical-fieldcontrolled magnetization switching in a magnetically doped TI³², highlight the exciting opportunities for future TI-based novel devices, such as the recently proposed TI/ferromagnet junction where transport can be altered from a spin-valve to an amplifier³³.

Results

Potentiometric measurement of spin-momentum locking. A schematic diagram of the spin-momentum locking textures of the Rashba 2DEG is shown in Fig. 1a, where the Rashba form of spin-orbit coupling leads to a pair of Fermi surfaces with counter-rotating spins. For the Dirac surface states of TI (Fig. 1b), only one such Fermi surface exists and the 2D surface states are occupied by a single spin. For TI materials that exhibit surface carrier accumulation due to band bending, trivial Rashba spin-split 2DEG states can coexist with the TI Dirac surface states (Fig. 1c). The trivial 2DEG states are nested within the linear dispersing Dirac states^{18,19} such that the spin orientation of the higher **k** 2DEG state is opposite that of the nearest Dirac state.

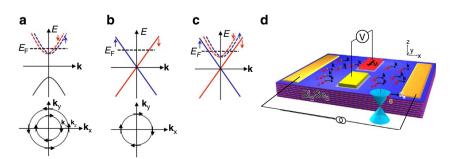


Figure 1 | Spin-momentum locking and experimental concept. Schematic diagram of the spin-momentum locking textures of the Rashba 2DEG (a), Dirac surface states of TI (b), and coexistence of both (c). (d) Experimental concept of the potentiometric measurement.

These spin textures arising from spin-momentum locking can be probed electrically using potentiometric measurements as shown in Fig. 1d, where the current-generated spin orientation is detected by its projection onto the ferromagnetic detector magnetization.

Current-generated spin in TI Dirac states. Bi₂Se₃ samples were grown by molecular beam epitaxy (MBE) on Al₂O₃(0001) substrates using a two-step process (see Methods). Structural and electronic properties of these films are provided in the Supplementary Figs 1–4. Fe/Al₂O₃ spin detector contacts were deposited on both Bi₂Se₃ and InAs samples in the same separate MBE system. The samples were then processed into identical device structures illustrated in Fig. 2a,b. Note that the geometry of the ferromagnetic detector contacts are specifically designed to cancel out the effect of fringe fields at the edges of the contacts (Supplementary Note 1). For the spin potentiometric measurements, a bias current is applied between the two Ti/Au current leads on either end of the device mesa, and a voltage is measured between the pairs of ferromagnetic (red) detector and corresponding non-magnetic Au/Ti (yellow) reference contacts.

When an unpolarized current flows between the two outer Ti/Au contacts, a spontaneous spin polarization is produced in the ${\rm Bi_2Se_3}$ surface states throughout the channel due to spin–momentum locking. The projection of this spin onto the magnetization of the ferromagnetic detector contact is recorded as a voltage with a high-impedance voltmeter (>1 Giga-ohm). An in-plane magnetic field is applied to rotate the magnetization of the Fe detector contact, so that the projection of the current-generated spins onto the detector magnetization changes, resulting in a change in sign and magnitude of the detector voltage. Here we define the positive current to be holes flowing from left to right along the +x axis, and the positive magnetic field to be pointing in the +y direction.

The detector voltage as a function of magnetic field for $+2 \,\mathrm{mA}$ bias current at $T=10 \,\mathrm{K}$ is shown in Fig. 2c, after a simple linear background subtraction²⁵ and centring around the vertical axis. Electron flow in the -x direction generates a spin in the +y direction due to spin-momentum locking in the TI Dirac states. At positive magnetic field>60 Oe, when the detector magnetization is saturated and completely parallel to the spin direction, a constant low voltage is observed. As the magnetic field decreases to small negative values around $-50 \,\mathrm{Oe}$ (coercive

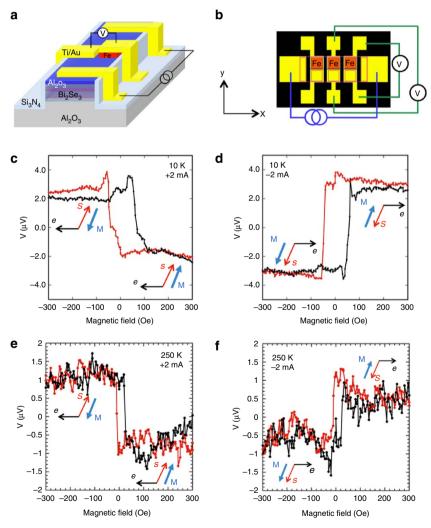


Figure 2 | Device schematic and electrical detection of current-generated spin in TI. Schematic (a) and top view (b) of contact layout with two parallel rows of collinear detector contacts, top row is ferromagnetic (Fe, red), bottom row is non-magnetic reference (Ti/Au). Magnetic field dependence of the voltage measured at the ferromagnetic detector contact with the magnetization collinear with the induced TI spin for bias currents of $+2 \,\mathrm{mA}$ (c) and $-2 \,\mathrm{mA}$ (d). Similar measurements at 250 K at $+2 \,\mathrm{mA}$ (e) and $-2 \,\mathrm{mA}$ (f).

field of the magnetic contact; red trace), an abrupt increase in detector voltage is seen as the detector magnetization reverses to be antiparallel with the TI surface-state spin, and the overall scan exhibits a single step-like behaviour. When the field sweep direction is reversed to increase from negative to positive values (black trace), the detector voltage is constant until reaching the positive coercive field of 50 Oe, where the voltage abruptly decreases as the detector magnetization again switches and becomes parallel with the spin orientation of the TI surface state.

Changing the current direction to $-2 \, \text{mA}$ (Fig. 2d), i.e., electrons flowing from left to right in the +x direction generate a spin in the -y direction. At positive magnetic field above saturation, the detector magnetization is antiparallel to the current-generated spin, and a constant high voltage is observed, while at negative magnetic field where the detector magnetization is parallel to the spin, a constant low voltage is seen. Comparing Fig. 2c,d, the hysteresis loop simply inverts around the x axis. This behaviour is very reproducible for temperatures up to 250 K, as is shown in Fig. 2e,f for both currents where clear hysteresis curves can still be seen.

Temperature and bias dependence. The temperature dependence of the magnitude of the spin signal, |V(+M) - V(-M)|, measured at $\pm 2 \,\text{mA}$ is shown in Fig. 3a. It decreases monotonically with increasing temperature to 175 K, exhibits a small increase between 175 and 250 K, and then disappears at 275 K. The small increase in the spin voltage in the range 175-250 K, and its abrupt suppression by 275 K are not well understood at present. This temperature dependence is similar to our previous observations of Bi₂Se₃ on graphene/n-SiC substrates²⁵, although in our previous work the spin-voltage was observed only up to 150 K. The higher temperature achieved in this case for Bi₂Se₃ films grown directly on an insulating Al₂O₃ substrate could be attributed to the fact that there is no current shunting through the epi-graphene/SiC conductive substrate used previously, so that a higher fraction of the bias current flows in the TI surface layer to produce the measured spin polarization.

The temperature dependence of the spin signal is similar to that of the conductivity shown in Fig. 3b (which necessarily includes both surface state and bulk contributions), suggesting that both are reduced by scattering events as the temperature increases. These scattering events may arise from impurity scattering or interactions between surface state and bulk carriers^{34–38}. For example, Dirac surface states can be scattered

by impurities, leading to Dirac point fluctuations³⁴, and phonons from the bulk states, which includes acoustic phonons that dominate at lower temperatures with a strong temperature dependence, as well as optical phonons that dominate at higher temperatures with a weaker temperature dependence³⁷. Hybridization with bulk states can also lead to reduction of spin polarization³⁸. In addition, the spin voltage may also be reduced by scattering between Rashba and Dirac states within the surface layer. Further experimental and theoretical work is required to understand such complex mechanisms and the resultant temperature dependence.

The dependence of the spin signal measured at the detector contact $(\Delta V = V(+M) - V(-M))$ as a function of the bias current at $T = 10 \,\mathrm{K}$ is shown in Fig. 3b, where a nearly linear dependence is observed. This linear behaviour of the spin signal with bias current is consistent with model calculations²³, where the voltages measured on the FM detector V(M) were directly related to the bias current and spin polarization by $[V(+M) - V(-M)] = I_b R_B P_{FM}$ (**p·M**_u), (bold case denotes a vector). Here I_h is the (hole) bias current in the +x direction, R_B is the ballistic resistance of the channel, and P_{FM} is the transport spin polarization of the FM detector metal. Mu is a unit vector along the detector magnetization M, and p is the degree of spin polarization induced per unit current by both spin-momentum locking in TI Dirac surface states and Rashba spin-orbit coupling in the 2DEG. From the spin signal we measure (for example, Fig. 3b), assuming that the bias current is shunted equally by each quintuple layer of the Bi_2Se_3 film²⁵, and taking P_{FM} (Fe) ~ 0.4, and $k_F \sim 0.15 \,\text{Å}^{-1}$, we estimate $\mathbf{p} \sim -0.15$, with a sign that's indicative of the TI Dirac states²³

Current-generated spin in InAs Rashba states. To further distinguish the sign of the spin signal measured for the TI Dirac surface states from that of potential trivial 2DEG states, we performed similar measurements on InAs(001) samples where only the surface 2DEG states are known to exist^{39–43}. It is well known that the downward band bending of the conduction band at the InAs(001) surface leads to an electron accumulation layer and the formation of a surface 2DEG^{39–43} (Fig. 4a) that extends ~20 nm into the sample⁴³. The InAs samples were processed to produce the same Fe/Al₂O₃ contact geometry used for the Bi₂Se₃ measurements (Materials and Methods). As noted earlier, the Rashba spin–orbit-induced polarization is predicted to exhibit the opposite sign to that of TI Dirac states²³ for a given bias current.

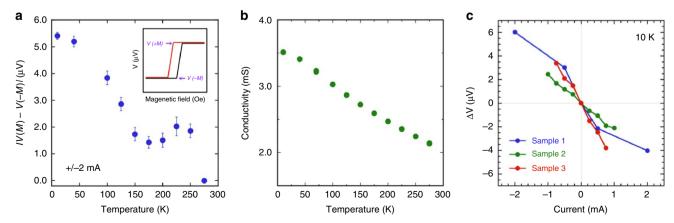


Figure 3 | Temperature and bias dependence of TI spin voltage. (a) Temperature dependence of the spin voltage at +/-2 mA bias current. (Inset: illustration of how $\Delta V = V(M) - V(-M)$ is determined. Error bars are determined based on variations from a 10-point moving average.) (b) Temperature dependence of the conductivity. (c) Bias current dependence of the ferromagnetic detector voltage for several samples and devices showing a general linear dependence (Sample 1: 10 nm Fe/Al₂O₃/20 nm Bi₂Se₃/Al₂O₃).

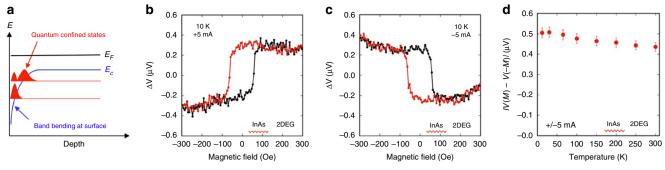


Figure 4 | Electrical detection of current-generated spin in InAs 2DEG. (a) Schematic of InAs surface 2DEG formation. Magnetic field dependence of the voltage measured at the ferromagnetic detector contact with the magnetization collinear with the induced 2DEG spin in InAs for bias currents of $+5 \,\text{mA}$ (b) and $-5 \,\text{mA}$ (c). (d) Temperature dependence of the spin voltage at $+/-5 \,\text{mA}$ bias current. (Error bars are determined based on variations from a 10 point moving average.)

The spin-voltage transport data for these InAs samples are shown in Fig. 4b,c. The measurement procedures were identical to those used for the Bi₂Se₃ samples. Similar hysteresis loops are observed where a constant high and low voltage is measured when the detector magnetization is fully aligned with the applied field. However, for a given current/electron flow direction, the hysteresis loop is clearly inverted about the horizontal axis relative to that observed for the Bi₂Se₃ samples (Fig. 2c-f): for a positive bias current, a high-voltage signal is seen at positive fields above the coercive field of the Fe contact, and a low voltage is observed at negative fields. Given that the spin-detecting contacts (Fe/Al₂O₃) and contact geometry are the same, and are sensitive only to the orientation of the induced spin polarization regardless of the source, this observation indicates that the bias current-induced spin polarization due to spin-momentum locking in the InAs 2DEG is opposite to that of the Bi₂Se₃ TI Dirac states, consistent with the theory²³. In addition, the spin voltage exhibits a weak temperature dependence, decreasing by only $\sim 10\%$ from 10 to 300 K, and persists to at least 300 K, as shown in Fig. 4d, consistent with the metallic nature of the 2DEG. This is markedly different than the strong temperature dependence observed for the TI Dirac surface state (Fig. 3a).

Discussion

While both the topologically protected Dirac surface states and the Rashba spin-split 2DEG states of the Bi₂Se₃ are expected to produce a bias current induced spin polarization due to spin-momentum locking, the current-induced spin density is expected to be substantially larger for the Dirac surface states for several reasons. First, as discussed above, for a given momentum direction, the Rashba 2DEG states exist as spin-split pairs of opposite spin orientation, and the net induced spin polarization is proportional to $(\mathbf{k}_2 - \mathbf{k}_1)/(\mathbf{k}_2 + \mathbf{k}_1)$ (ref. 23), where $\mathbf{k}_2 > \mathbf{k}_1$ (Fig. 1a). Consequently, the contributions from these spin-split states tend to cancel. In contrast, the Dirac state has only one spin orientation, and no such cancellation occurs. Second, the induced spin polarization is enhanced by a factor $v_F/\alpha >> 1$ for the TI Dirac surface states, where v_F is the Fermi velocity of the TI (on the order of $10^5\,\mathrm{m\,s^{-1}}$) and α the strength of the Rashba spin-orbit coefficient in the 2DEG in units of velocity (on the order of 10^3 m s⁻¹)^{13,20,22}. The fact that in degenerate Bi₂Se₃ samples where both the Dirac and Rashba states coexist, the sign of the spin signal we observe corresponds to that of the TI Dirac states, corroborates the prediction and expectation that the signal should be dominated by contributions from the Dirac states.

As noted earlier, there are inconsistencies in the sign of the spin signal [V(+M) - V(-M)] reported for nominally identical measurements of the bias current-induced spin polarization attributed to

the TI Dirac states, even for the same $\mathrm{Bi}_2\mathrm{Se}_3$ material (and stated conventions for current and magnetic field directions). This is reflected in whether a low- or high-voltage signal is observed when the detector magnetization is parallel or antiparallel to the induced spin. In our measurements of the TI Dirac surface states, we observe [V(+M)-V(-M)]<0, that is, a low-voltage signal when the detector magnetization and TI spin are parallel, and a high signal when antiparallel (Fig. 2; ref. 25), as have other groups²⁹. In contrast, others report the opposite behaviour^{28,30}, as well as markedly different temperature dependence for the spin voltage in $\mathrm{Bi}_2\mathrm{Se}_3$ films³⁰.

Hence, to directly derive the sign of the spin voltage that should be expected, we develop a model based on the spin-dependent electrochemical potentials generated and their detection by a ferromagnetic detector contact. We note that similar models have been reported in refs 28,30, although the gradients and/or the reference of the electrochemical potentials are inconsistent with conventional usage in the spintronics community. To better illustrate and contrast the discrepancies, we construct our diagram using notation similar to that of ref. 28.

We begin with a simple 3-terminal measurement geometry similar to that of Hong $et~al.^{23}$ shown in Fig. 5a. We define the left contact as the positive terminal, and the right contact as the negative terminal, or the reference contact, as used in our measurements. The positive magnetic field direction (and detector magnetization) is again defined to be in the +y direction, with positive (hole) current flowing in the +x direction. We present a diagram of the electric field and voltage (V), where V is directly related to electrochemical potential (μ) by $\mu = -eV$ (refs 44,45), where e is the electron charge (taken to be a positive quantity). With these conventions, the voltage reference point and gradient of the electric field are unambiguously defined. In the following, we discuss the measurement in terms of both the voltage and electrochemical potential.

For a positive current (I>0; Fig. 5b,c), electron momentum \mathbf{k}_{e} is from right to left in the -x direction, and the voltage of the left electrode (V_{L}) is high relative to that of the right electrode (V_{R}). The right (reference) contact need not be grounded, so we indicate a zero reference $V=\mu=0$ by the yellow line common to Fig. 5b,c. Thus, the gradient of the electric field has a negative slope (Fig. 5b). The profile of the electrochemical potential (Fig. 5c), $\mu=-eV$, is merely the mirror image of the electric field/voltage profile across the $V=\mu=0$ axis. Here the left electrode (L) has a more negative (larger magnitude) electrochemical potential than the right electrode (R), and the profile has a positive slope.

When a bias current flows through the TI Dirac surface states, a net spin polarization is created due to spin-momentum locking with a direction determined by the electron momentum: for $\mathbf{k}_{\rm e}$ along the -x direction (positive bias current), the induced spin is

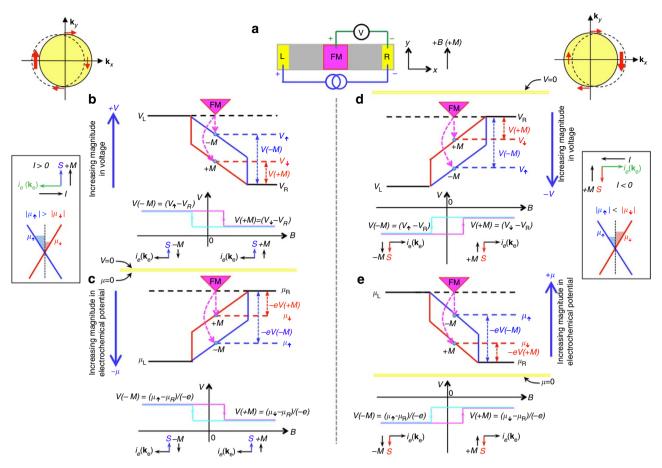


Figure 5 | Model to derive sign of spin signal expected in TI. (a) Schematic of a simplified 3-terminal device with a left (L) and right (R) electrode and central magnetic detector contact (FM), as well as definitions of voltage (V) terminals, magnetic field (B) and magnetization (M) directions. Model of the spin potentiometric measurement probing the current-induced spin polarization due to TI surface states, based on the spin-dependent electrochemical potentials: for positive bias currents using voltage profiles (\mathbf{b}) and electrochemical potential (μ) profiles (\mathbf{c}), and for negative bias currents in the form of voltage profiles (\mathbf{d}) and electrochemical potential profiles (\mathbf{e}). The definitions of current, spin (S), and magnetization directions are indicated for positive currents next to \mathbf{b} , and for negative currents next to \mathbf{d} , and red horizontal dashed lines indicate the levels been probed by corresponding magnetization directions. For example, for \mathbf{b} , \mathbf{c} , the spin-up (blue) voltage and electrochemical potentials are probed by the FM detector with -M magnetization (oriented along -y), which has its majority spin oriented along +y (due to the opposite alignment of majority spin to its magnetization in magnetic metals⁴⁷). The yellow horizontal lines indicate the zero axis for the voltage and electrochemical potential ($V = \mu = 0$), which is shared for \mathbf{b} , and individually marked for \mathbf{d} , \mathbf{e} .

oriented along the +y axis and referred to as spin-up. Consequently, the electrochemical potential splits for spin-up and spin-down electrons, as represented by the blue μ_{\uparrow} and red μ_{\downarrow} lines in Fig. 5c, where μ_{\uparrow} for the spin-up electrons is larger (in magnitude, that is, more negative) than that of the spin-down μ_{\downarrow} (i.e., $|\mu_{\uparrow}| > |\mu_{\downarrow}|$). The corresponding levels in the voltage diagram of Fig. 5b are again the mirror image across the horizontal axis.

This spin imbalance is probed by the ferromagnetic detector contact. The magnetization of the ferromagnet aligns with the applied external magnetic field above saturation. However, its magnetic moment is opposite to the orientation of its majority spin⁴⁶. Hence, the FM detector with +M magnetization (oriented along +y) has its majority spin oriented along -y, and will probe the spin-down electrochemical potential $(\mu_{\downarrow}, V_{\downarrow})$ in the channel. Conversely, the detector with -M magnetization probes the spin-up levels $(\mu_{\uparrow}, V_{\uparrow})$.

Since the right electrode (R) is the reference, when the detector magnetization is saturated at positive magnetic field, its voltage V(+M) due to probing the spin-down electron band is $-eV(+M) = \mu_{\downarrow} - \mu_{R}$, or $V(+M) = (\mu_{\downarrow} - \mu_{R})/(-e)$. Similarly,

when the detector magnetization is saturated at negative magnetic field, its voltage due to probing the spin-up band is $V(-M) = (\mu_{\uparrow} - \mu_{R})/(-e)$. Since $|\mu_{\uparrow}| > |\mu_{\downarrow}|$, this yields a high voltage signal at the negative field, when the magnetization is antiparallel to the TI spin (spin-up), and a low voltage at positive field, when it is parallel to the TI spin, as depicted by the hysteresis loop below the potential diagram in Fig. 5c. Note that a simple linear background subtraction and centring around the vertical axis does not change the relative high and low signals.

A similar analysis can be made directly in terms of the voltage, as shown in the top panels of Fig. 5b. Electrons flowing from right to left in the -x direction create a net spin-up population oriented along +y, and hence the blue V_{\uparrow} level is higher (that is, larger in magnitude) than that of the spin-down V_{\downarrow} (that is, $V_{\uparrow} > V_{\downarrow}$), analogous to $|\mu_{\uparrow}| > |\mu_{\downarrow}|$. With the right electrode (R) as the reference, the detector voltage at positive magnetic field (+B, +M) probing the spin-down level is $V(+M) = (V_{\downarrow} - V_{R})$, and that at negative magnetic field (-B, -M) which probes spin-up is $V(-M) = (V_{\uparrow} - V_{R})$. And since $V_{\uparrow} > V_{\downarrow}$, at positive field (when magnetization is parallel to TI spin), a low voltage is expected, and at negative field (magnetization antiparallel to TI

spin), a high voltage is expected (or ΔV (= V(+M) - V(-M)) should be negative), exactly as we observe experimentally in Bi₂Se₃.

This exercise can be repeated when the current direction (and hence induced spin direction) is reversed, as shown in Fig. 5d,e. We note that discrepancies/mistakes can be made when discussing electrochemical potentials with negative values 28,30 . However, these can be avoided by establishing the voltage potential first, as derived above. Another potential complication is that the measured spin voltage from the trivial 2DEG states is also sensitive to the sign and value of the Rashba spin–orbit coupling parameter α (ref. 23), which can vary depending on the nature of the interface 47 . In the current case, a positive value of α has been reported for various types of TI and InAs in the literature, hence no sign reversal should be expected than that predicted in refs 23,24.

In summary, we directly compare electrical measurements of current-generated spin polarization due to spin-momentum locking in two complimentary systems: Bi₂Se₃ with the potential coexistence of both Dirac and trivial Rashba surface states, and InAs with only the Rashba states. We show that the spin voltages measured for the Dirac and Rashba systems are indeed opposite in sign, as predicted by theory²³. We further develop a model based on spin-splitting of the electrochemical potential to derive the sign of the spin voltage expected for the Dirac states from a potentiometric measurement using a ferromagnetic contact, further confirming that the spin signals we observe from the Bi₂Se₃ are consistent with the Dirac surface states. These results demonstrate that the currentgenerated spin polarization in non-trivial TI Dirac and trivial Rashba 2DEG states are indeed opposite, as expected from their different energy band dispersion, and that in a TI it is dominated by the Dirac surface states. These demonstrations of direct electrical detection of the helical spin texture of Dirac and Rashba states is an enabling step towards electrical manipulation of spins in future TI-based quantum devices.

Methods

Sample growth. The growth of Bi_2Se_3 films was carried out on $Al_2O_3(0001)$ substrates in an ultrahigh vacuum (UHV) system (base pressure $\sim 1 \times 10^{-10}$ torr) that integrates two MBE chambers and a low temperature (500–300 K) scanning tunneling microscope. For the growth of Bi_2Se_3 , a two-step process is used⁴⁸: 2–3 quintuple layers (QL) of Bi_2Se_3 were first deposited at a reduced temperature of $100\,^{\circ}\text{C}$, and the substrate temperature was then slowly raised to $300\,^{\circ}\text{C}$ where the rest of the film was deposited. Bi and Se were supplied via separate Knudsen cells at 460 and $250\,^{\circ}\text{C}$, respectively⁴⁹. The samples were then removed to air and transferred to a separate MBE system where Fe/Al_2O_3 contacts were deposited as described previously⁵⁰ and below.

In the case of InAs, an undoped InAs(001) substrate was heated to $520\,^{\circ}\text{C}$ in an As flux to desorb the oxide. The sample was then cooled to room temperature and transferred under ultrahigh vacuum into an interconnected MBE system for the growth of Fe/Al₂O₃ (the same system used to deposit Fe/Al₂O₃ on Bi₂Se₃).

The Fe/Al₂O₃ contacts were formed on Bi₂Se₃ as follows. A 0.7 nm layer of polycrystalline Al was first deposited by MBE, and then oxidized in 200 torr O₂ for 20 min in the presence of ultraviolet light in the load-lock chamber. This step was then repeated for a total Al₂O₃ thickness of 2 nm. The sample was then transferred under UHV to an interconnected metals MBE chamber, where 20 nm of polycrystalline Fe was deposited at room temperature from a Knudsen cell.

Sample processing. The samples were processed into the device structures illustrated in Fig. 2a,b to enable transport measurements. Standard photolithography and chemical etching methods were used to define the Fe contacts, which ranged in size from $20\times20\,\mu\text{m}^2$ to $80\times80\,\mu\text{m}^2$, with adjacent contact separation ranging from 45 to $200\,\mu\text{m}$. Ion milling was used to pattern the Bi_2Se_3 mesa. Large Ti/Au contacts were deposited by lift-off in an electron beam evaporator as non-magnetic reference contacts and bias current leads. The Fe contacts were capped with 10 nm Ti/100 nm Au, and bond pads for wire bonded electrical connections are further electrically isolated using 100 nm of Si_3N_4 .

Transport measurements. Transport measurements were performed in a closed cycle cryostat equipped with an electromagnet (4–300 K, \pm 1,000 Oe). An unpolarized bias current was applied through the outer Ti/Au contacts on the opposite

ends of the ${\rm Bi_2Se_3}$ mesa, and the voltage on the detector contact was recorded as a function of the in-plane magnetic field applied orthogonal to the electron bias current direction in the TI.

Materials characterization. Topological insulator thin films are grown by molecular beam epitaxy and characterized by a variety of techniques. Synchrotron-based high-resolution X-ray Reflectivity was carried out at the Argonne National Lab in collaboration with Dr Jonathan Emery of Northwestern University. Raman spectroscopy was done at the University of Wisconsin, Milwaukee. High resolution transmission electron microscopy and associated high angle annular dark field imaging was done at the University of York, UK in collaboration with Dr Vlado Lazarov. Scanning tunneling microscopy and spectroscopy was carried out *in situ* after growth at 77 K at University of Wisconsin, Milwaukee.

Data availability. The data sets generated during and/or analysed during the current study are available from the corresponding author on reasonable request.

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Author contributions

C.H.L., L.L. and B.T.J. conceived and designed the experiments. S.R. and L.L. grew the Bi₂Se₃ films. C.H.L. fabricated the devices and performed the transport measurements with assistance from O.M.J.v.E. C.H.L., O.M.J.v.E. and B.T.J. Analysed the data. C.H.L., L.L. and B.T.J. wrote the paper. All authors discussed the results and commented on the manuscript.

Additional information

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