

ARTICLE



<https://doi.org/10.1038/s41467-021-26915-3>

OPEN

Spin-neutral currents for spintronics

Ding-Fu Shao¹ [✉], Shu-Hui Zhang² ², Ming Li¹, Chang-Beom Eom³ & Evgeny Y. Tsymbal¹ [✉]

Electric currents carrying a net spin polarization are widely used in spintronics, whereas globally spin-neutral currents are expected to play no role in spin-dependent phenomena. Here we show that, in contrast to this common expectation, spin-independent conductance in compensated antiferromagnets and normal metals can be efficiently exploited in spintronics, provided their magnetic space group symmetry supports a non-spin-degenerate Fermi surface. Due to their momentum-dependent spin polarization, such antiferromagnets can be used as active elements in antiferromagnetic tunnel junctions (AFMTJs) and produce a giant tunneling magnetoresistance (TMR) effect. Using RuO₂ as a representative compensated antiferromagnet exhibiting spin-independent conductance along the [001] direction but a non-spin-degenerate Fermi surface, we design a RuO₂/TiO₂/RuO₂ (001) AFMTJ, where a globally spin-neutral charge current is controlled by the relative orientation of the Néel vectors of the two RuO₂ electrodes, resulting in the TMR effect as large as -500%. These results are expanded to normal metals which can be used as a counter electrode in AFMTJs with a single antiferromagnetic layer or other elements in spintronic devices. Our work uncovers an unexplored potential of the materials with no global spin polarization for utilizing them in spintronics.

¹Department of Physics and Astronomy & Nebraska Center for Materials and Nanoscience, University of Nebraska, Lincoln, NE 68588-0299, USA. ²College of Mathematics and Physics, Beijing University of Chemical Technology, Beijing 100029, People's Republic of China. ³Department of Materials Science and Engineering, University of Wisconsin-Madison, Madison, WI 3706, US. ✉email: dfshao@unl.edu; tsymbal@unl.edu

The field of spintronics utilizes the spin degree of freedom in condensed matter for information processing and storage¹. Most spintronic applications rely on electric currents with sizable spin polarization for detection or manipulation of the magnetic order parameter in spintronic devices. A typical and widely used spintronic device is the magnetic tunnel junction (MTJ), where a longitudinal charge current spin polarized by one ferromagnetic metal quantum-mechanically tunnels into another ferromagnetic metal through an insulating barrier layer^{2,3}. Conductance of the MTJ is controlled by the relative magnetization orientation of the two ferromagnetic electrodes, resulting in a tunneling magnetoresistance (TMR) effect⁴.

Contrary to the spin-polarized currents, spin-neutral currents are usually considered impractical for spintronics due to being unable to directly interact with the magnetic order parameter. This fact challenges spintronics based on non-ferromagnetic materials, such as compensated antiferromagnets, which normally do not support spin-polarized currents. Due to being robust against magnetic perturbations, the absence of stray fields, and ultrafast spin dynamics, antiferromagnets are considered as outstanding candidates to replace the widely used ferromagnets in the next generation spintronics^{5–9}. This promising route has been recently stimulated by the demonstrated control of the antiferromagnetic Néel vector by spin-orbit torques^{10,11}. However, the absence of a net magnetization and hence spin-independent conductance makes the electrical detection of the Néel vector using conventional methods, such as TMR measurements, unfeasible⁵. So far, the electrical detection of the Néel vector has been performed using anisotropic^{10,11} or spin-Hall^{12–15} magnetoresistance. Unfortunately, both methods suffer from relatively small signals easily influenced by perturbations¹⁶ and require multiple in-plane terminals resulting in large device dimensions⁷. Antiferromagnetic spin valves^{17–20} and antiferromagnetic tunnel junctions (AFMTJs)^{21,22} have been theoretically proposed, promising, in some cases, sizable magnetoresistance effects. However, these magnetoresistance effects rely on perfect interfaces and switching the interfacial magnetic moment alignment between parallel and antiparallel. This mechanism is not robust against disorder and interface roughness inevitable in experimental conditions. Recent efforts have been aimed at exploring unconventional methods for the Néel vector detection based on topological properties^{9,23–25} but require an experimental confirmation.

One promising direction is to create spin-polarized currents in antiferromagnets. Recently, it has been predicted that certain types of compensated antiferromagnets exhibit a momentum-dependent spin splitting of the Fermi surface^{26–28}, resulting in spin-polarized currents along certain crystallographic orientations^{29–31}. These predictions indicate that these antiferromagnets can work as ferromagnets in spintronic devices, which broadens the range of materials useful for spintronics.

Here, we embark on a different path and argue that *globally spin-independent* conductance in compensated antiferromagnets can be efficiently used in spintronics, provided their crystal symmetry supports a non-spin-degenerate Fermi surface and thus momentum-dependent spin polarization. While such a spin polarization is cancelled out in the net conductance due to being antisymmetric with respect to certain symmetry operations, its presence in the momentum space can be functionalized if such an antiferromagnet is combined with another similar antiferromagnet in a spintronic device such as an AFMTJ. In this case, the resistance change of the AFMTJ occurs in response to the orientation of the antiferromagnetic Néel vector due to changing matching conditions between the spin-polarized conduction channels in the two metal electrodes. These considerations can be expanded to normal (nonmagnetic) metals with spin-orbit

coupling where the combined space inversion-time reversal symmetry is broken, indicating that they can also be utilized in spintronics despite globally spin-neutral currents.

Results

Spin polarized conduction channels. To explore the possible use of spin-neutral currents in a spintronic device, we first consider ballistic conductance of a material under investigation. Since the ballistic conductance is determined by the number of conduction channels, i.e. propagating Bloch states at the Fermi energy, it can provide an important characteristic of a spintronic device where this material is used as a metal electrode^{32,33}. In the absence of spin-orbit coupling, the ballistic conductance g per unit area along the z direction can be obtained in terms of two spin components as follows³⁴.

$$g = g^\uparrow + g^\downarrow = \frac{e^2}{h} \sum_{k_\parallel} (N_\parallel^\uparrow + N_\parallel^\downarrow), \quad (1)$$

$$N_\parallel(\vec{k}_\parallel) = \frac{\hbar}{2} \sum_n \int |\nu_{nz}^\sigma| \frac{df}{\partial E_n^\sigma(\vec{k})} dk_z. \quad (2)$$

Here σ denotes the spin component \uparrow or \downarrow , \vec{k} is the wave vector in the three-dimensional Brillouin zone, $N_\parallel^\sigma(\vec{k}_\parallel)$ is the number of conduction channels (integer) at the transverse wave vector $\vec{k}_\parallel = (k_x, k_y)$ for spin σ , E_n^σ is energy for the n -th band, $\nu_{nz}^\sigma = \frac{\partial E_n^\sigma(\vec{k})}{\hbar \partial k_z}$ is the band velocity along the transport z direction, and f is the Fermi distribution function.

The net transport spin polarization is defined by

$$p = \frac{g^\uparrow - g^\downarrow}{g} \quad (3)$$

and represents an important quantity useful in spintronics. For example, in a crude approximation of \vec{k}_\parallel -independent transmission between two ferromagnetic electrodes with spin polarizations p_1 and p_2 in an MTJ, the TMR effect is given by the well-known Julliere's formula² $TMR = \frac{2p_1 p_2}{1 - p_1 p_2}$. Clearly, a larger spin polarization of the electrodes favors a larger TMR.

A large spin polarization p is generally expected for ferromagnets where the finite net magnetization breaks time reversal symmetry \hat{T} . The latter flips the spin σ and changes sign of \vec{k}_\parallel , resulting in $\hat{T} N_\parallel^\uparrow(\vec{k}_\parallel) = N_\parallel^\downarrow(-\vec{k}_\parallel)$. Compensated antiferromagnets do not have net magnetization and hence (with some exceptions^{29–31}) do not support the macroscopic spin-polarized current. However, even though macroscopically the net transport spin polarization p is absent, microscopically the conductance could be spin polarized as reflected in the spin polarization of conduction channels at \vec{k}_\parallel :

$$p_\parallel(\vec{k}_\parallel) = \frac{N_\parallel^\uparrow - N_\parallel^\downarrow}{N_\parallel^\uparrow + N_\parallel^\downarrow}. \quad (4)$$

If both electrodes in a two-terminal spintronic device are made of materials with zero net spin polarization but have spin-polarized conduction channels, this momentum-dependent spin polarization will be reflected in the device conductance and can be functionalized through the antiferromagnetic Néel vector. Indeed, in the transport regime conserving spin (no spin-orbit coupling) and wave vector \vec{k}_\parallel (no diffuse scattering), the device conductance is largely affected by the spin matching of the conduction channels \vec{k}_\parallel of the electrodes. If their spin polarization changes in response to the Néel vector rotation in an antiferromagnetic electrode, this alters the net conductance of

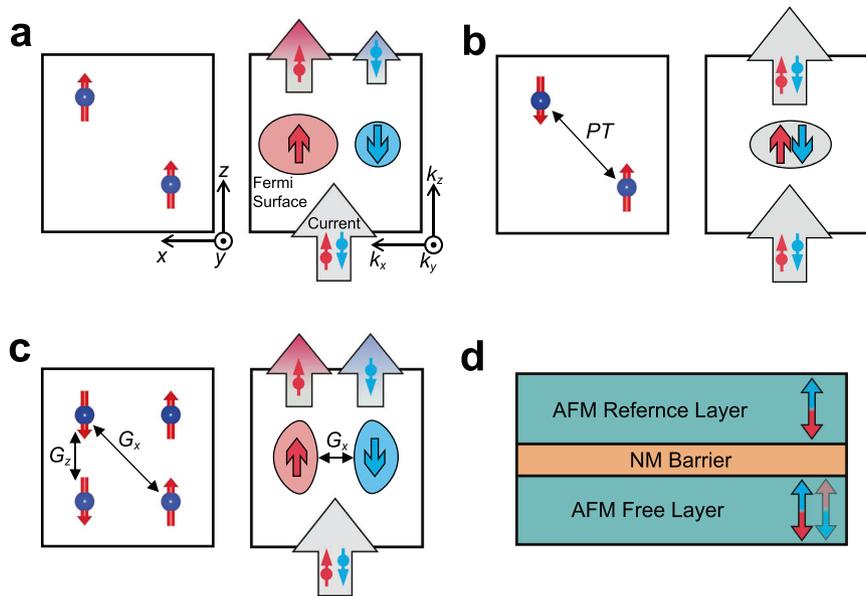


Fig. 1 Spin-polarized conduction channels in different types of magnetic materials. **a** Schematics of the atomic structure (left) and spin-polarized Fermi surface (right) for a ferromagnet. A spin-neutral current passing through the ferromagnet becomes spin polarized. **b** Schematics of the atomic structure (left) and spin-degenerate Fermi surface (right) for a compensated antiferromagnet where the net magnetization is forbidden by $\hat{P}\hat{T}$ symmetry. A spin-neutral current passing through this antiferromagnet remains spin neutral. **c** Schematics of the atomic structure (left) and non-spin-degenerate Fermi surface congruent for opposite spins (right) for a compensated antiferromagnet where the net magnetization is forbidden by glide symmetries \hat{G}_x and \hat{G}_z . A spin-neutral current passing through this antiferromagnet remains globally spin neutral but has a momentum-dependent spin polarization. **d** Schematics of an AFMTJ where two antiferromagnetic (AFM) layers are separated by a nonmagnetic (NM) barrier layer. The Néel vector (indicated by arrows) of the bottom free layer can be switched resulting in the TMR effect.

the device. Thus, although the spin polarization of the charge current remains zero, the device conductance reflects the \vec{k}_{\parallel} -dependent spin polarization of the conduction channels.

Next, we identify magnetic space group symmetry requirements for crystals to exhibit the \vec{k}_{\parallel} -dependent spin polarization. Obviously, in ferromagnets, the conduction channels are spin polarized due to the spin-dependent Fermi surface. Thus, passing a spin-neutral current through a ferromagnetic material makes it spin polarized (Fig. 1a). On the contrary, most compensated antiferromagnets contain symmetries that not only prevent the net magnetization but also lead to a spin-degenerate Fermi surface and thus spin-independent conduction channels. For example, if a compensated antiferromagnet exhibits $\hat{P}\hat{T}$ symmetry, where \hat{P} and \hat{T} are space inversion and time reversal symmetries, respectively, $p_{\parallel} = 0$ due to $\hat{P}\hat{T}N_{\parallel}^{\uparrow}(\vec{k}_{\parallel}) = N_{\parallel}^{\downarrow}(\vec{k}_{\parallel})$. This property follows from the spin-degenerate Fermi surface due to $\hat{P}\hat{T}E_{n}^{\uparrow}(\vec{k}) = E_{n}^{\downarrow}(\vec{k})$ (Fig. 1b). The spin degeneracy also appears in compensated antiferromagnets with $\hat{T}\hat{t}$ symmetry (\hat{t} is half a unit cell translation) in the absence of spin-orbit coupling.

The spin degeneracy is however broken in compensated antiferromagnets belonging to magnetic space groups with violated $\hat{P}\hat{T}$ and $\hat{T}\hat{t}$ symmetries²⁶. The vanishing net magnetization in such antiferromagnets originates from the combination of some other magnetic space group symmetries of the crystal. For example, Fig. 1c shows a collinear antiferromagnet with the Néel vector pointing along the z direction. The zero net magnetization in this antiferromagnet is guaranteed by two glide symmetries \hat{G}_x and \hat{G}_z , where $\hat{G}_i = \{\hat{M}_i|\hat{t}\}$ represents mirror symmetry \hat{M}_i with a mirror plane normal to vector \vec{t} combined with translation \hat{t} . The symmetry transformation $\hat{G}_x N_{\parallel}^{\uparrow}(k_x, k_y) = N_{\parallel}^{\downarrow}(-k_x, k_y)$ flips the spin and thus according to Eqs. (1) and (2) results in a

vanishing net spin polarization p for the current along the z direction. On the other hand, due to the Néel vector pointing along the z axis, the symmetry transformation $\hat{G}_z E_{n}^{\sigma}(\vec{k}_{\parallel}, k_z) = E_{n}^{\sigma}(\vec{k}_{\parallel}, -k_z)$ conserves the spin $\sigma = \uparrow, \downarrow$ of the conduction modes at \vec{k}_{\parallel} and there is no symmetry operation which would enforce $p_{\parallel} = 0$. The presence of spin-polarized conduction channels in this type of antiferromagnets can be understood in terms of two congruent (but not identical) up- and down-spin Fermi surfaces, which are transformed to each other by the symmetry transformation \hat{G}_x (Fig. 1c). In this case, each conduction channel is spin polarized (except high-symmetry \vec{k}_{\parallel} points invariant to \hat{G}_x), whereas the net conductance is spin neutral.

Due to the non-spin-degenerate Fermi surface and spin-polarized conduction channels, the globally spin-neutral conduction of the compensated antiferromagnets can be exploited in spintronic devices, such as AFMTJs. Fig 1d shows an AFMTJ which contains two identical antiferromagnetic electrodes separated by a nonmagnetic insulating spacer. The antiferromagnets are assumed to have spin-polarized conduction channels along the out-of-plane transport direction. The functionality of the AFMTJ is controlled by the relative orientation of the Néel vector of the two antiferromagnetic electrodes. In the parallel state, the spin-polarized conduction channels of the electrodes perfectly match, resulting in a low resistance state. In the antiparallel state, the spin polarized conduction channels are mismatched, resulting in a high resistance state.

Electronic structure of RuO₂. To demonstrate this spintronic functionality, we consider the recently discovered room-temperature antiferromagnetic metal RuO₂³⁵ suitable for realizing the proposed AFMTJ. RuO₂ exhibits interesting properties

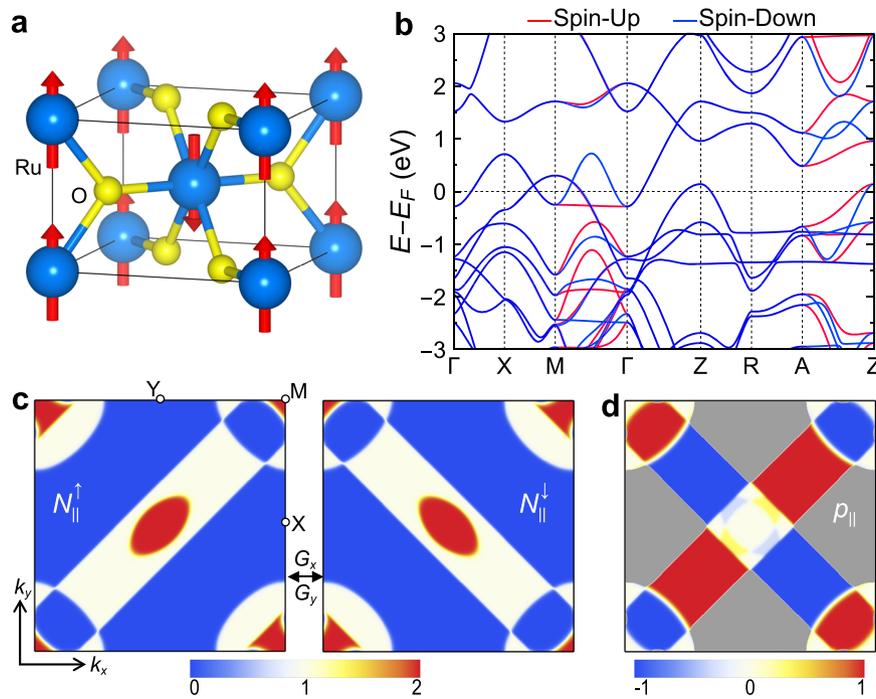


Fig. 2 Electronic properties of RuO₂. **a** The atomic and magnetic structures of RuO₂. **b** The calculated band structure of RuO₂. **c** The number of \vec{k}_{\parallel} -resolved conduction channels in the 2D Brillouin zone of RuO₂ for spin up N_{\parallel}^{\uparrow} (left) and spin down $N_{\parallel}^{\downarrow}$ (right). High-symmetry \vec{k}_{\parallel} -points are indicated. N_{\parallel}^{\uparrow} and $N_{\parallel}^{\downarrow}$ can be transformed to each other by the glide transformations \hat{G}_x or \hat{G}_y . **d** Spin polarization of conduction channels $p_{\parallel}(\vec{k}_{\parallel})$. Gray contrast indicates regions where $N_{\parallel}^{\uparrow} = N_{\parallel}^{\downarrow} = 0$ and thus p_{\parallel} undefined.

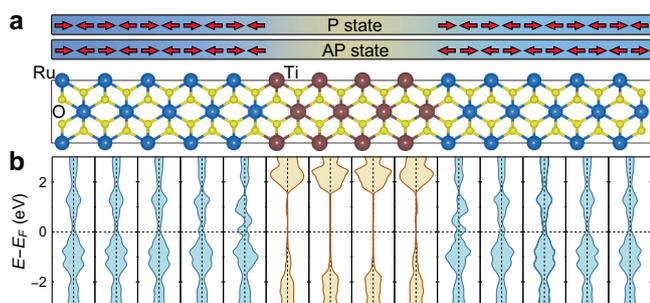


Fig. 3 Atomic and electronic structure of a RuO₂/TiO₂/RuO₂ supercell.

a, b The atomic structure (**a**) and layer resolved density of states (DOS) (**b**) of the RuO₂/TiO₂/RuO₂ supercell. Each panel in **b** contains two atomic layers of MO₂ (M = Ru, Ti) and has left and right subpanels corresponding to up- and down-spin states, respectively. The arrows in (**a**) indicate the magnetic moments of Ru atoms.

such as spin splitting³⁶, a crystal Hall effect³⁷, and a magnetic spin Hall effect^{31,38}. RuO₂ has a rutile structure with an out-of-plane Néel vector (Fig. 2a) and magnetic space group $P4_2'/mmm'$, which contains glide $\hat{G}_x = \{\hat{M}_x | (\frac{1}{2}, \frac{1}{2}, \frac{1}{2})\}$, $\hat{G}_y = \{\hat{M}_y | (\frac{1}{2}, \frac{1}{2}, \frac{1}{2})\}$ and mirror \hat{M}_z symmetries. In the absence of spin-orbit coupling, the energy bands are spin degenerate at the k -planes invariant to \hat{G}_x and \hat{G}_y , such as $k_x = 0, \frac{\pi}{2}$ or $k_y = 0, \frac{\pi}{2}$. This is evident from our first-principles density functional theory (DFT) calculations. As seen from Fig. 2b, the energy bands of RuO₂ are spin degenerate along the Γ -X, Γ -Z, X-M, Z-R, and R-A directions lying in these glide-invariant planes. On the other hand, a large spin splitting appears along the directions away from these planes, such as Γ -M and Z-A.

The spin-dependent band structure leads to the momentum-dependent spin polarization of the conduction channels along the z direction. We explicitly demonstrate this by calculating the

number of conduction channels $N_{\parallel}^{\sigma}(\vec{k}_{\parallel})$ in the two-dimensional (2D) Brillouin zone of RuO₂. As seen from Fig. 2c, the distributions of N_{\parallel}^{\uparrow} and $N_{\parallel}^{\downarrow}$ in the (k_x, k_y) plane have congruent shapes and are symmetric with respect to the \hat{G}_x and \hat{G}_y symmetry transformations, which enforce a zero global spin polarization in the conductance along the z direction. On the contrary, as seen from Fig. 2d, the \vec{k}_{\parallel} -dependent spin polarization p_{\parallel} remains finite across a sizable portion of the 2D Brillouin zone. This demonstrates that RuO₂ (001) exhibits a globally spin-neutral conductance through spin-polarized conduction channels.

TMR in a RuO₂/TiO₂/RuO₂ AFMTJ. Next, we design an AFMTJ using RuO₂ (001) as electrodes and TiO₂ (001) as an insulating barrier layer. Due to both having a rutile structure and a similar lattice constant, this AFMTJ is feasible in practice. Fig. 3a shows the atomic structure of the RuO₂/TiO₂/RuO₂ (001) supercell, which is used in our DFT and quantum transport calculations and includes 8 TiO₂ layers in the center and 10 RuO₂ layers on each side. We find that a wide band gap of TiO₂ is well maintained in this heterostructure, and the Fermi energy E_F is located deeply inside the band gap (Fig. 3b).

The RuO₂/TiO₂/RuO₂ (001) structure in Fig. 3a is then used as the scattering region of the AFMTJ connected to two semi-infinite RuO₂ (001) electrodes for calculating transmission. The transmission is obtained for parallel (Fig. 4a) and antiparallel (Fig. 4b) alignments of the Néel vectors of the electrodes. For the parallel-aligned AFMTJ, the \vec{k}_{\parallel} -resolved transmission $T_p^{\sigma}(\vec{k}_{\parallel})$ is shown in Fig. 4c for spin up (left panel) and spin down (right panel). The clearly seen spin asymmetry between $T_p^{\uparrow}(\vec{k}_{\parallel})$ and $T_p^{\downarrow}(\vec{k}_{\parallel})$ reflects the related asymmetry in the distribution of the spin-polarized conduction channels in RuO₂ (Fig. 2c). The suppressed transmission near the Brillouin zone corners is due to

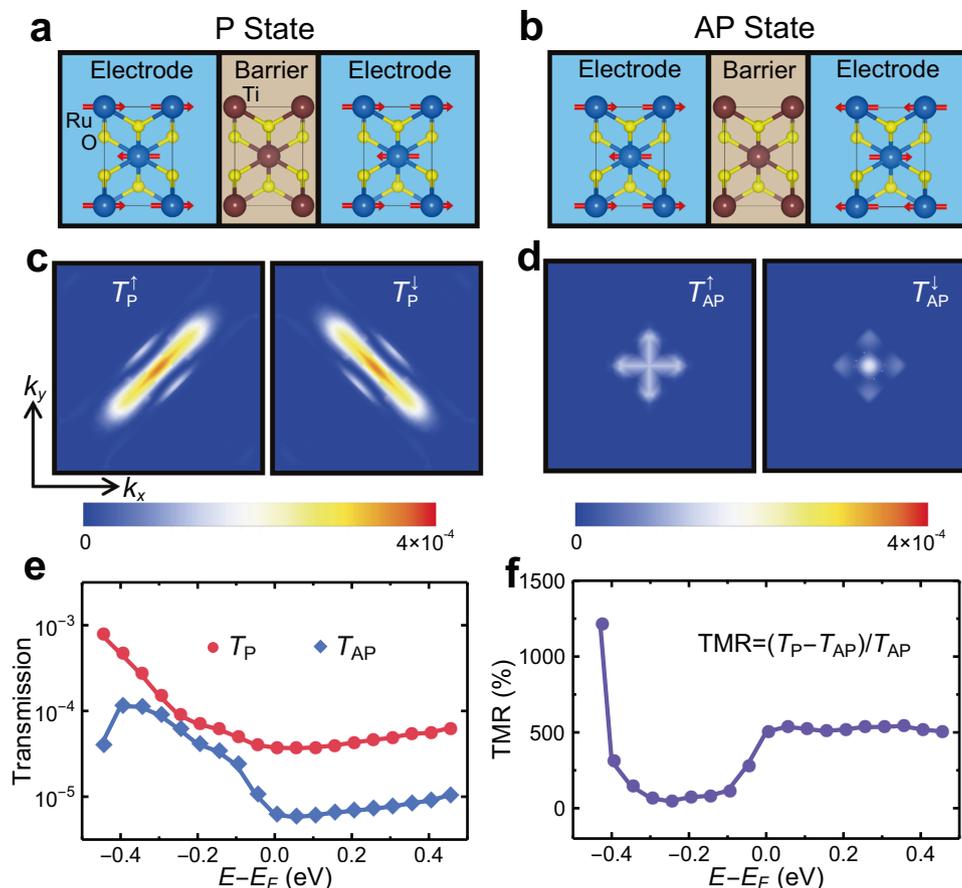


Fig. 4 Giant TMR in RuO₂/TiO₂/RuO₂ AFMTJ. **a, b** The atomic and magnetic structures of RuO₂/TiO₂/RuO₂ AFMTJ for parallel (P) (**a**) and antiparallel (AP) (**b**) alignment of the Néel vectors. **c, d** The calculated \vec{k}_{\parallel} -resolved transmission in the 2D Brillouin zone for the AFMTJ in P (**c**) and AP (**d**) states. **e** Total transmission as a function of energy for the AFMTJ in P (red dots) and AP (blue dots) states. **f** TMR as a function of energy.

a larger decay rate of the evanescent states for \vec{k}_{\parallel} away from the zone center.

For the antiparallel-aligned AFMTJ, the transmission $T_{AP}^{\sigma}(\vec{k}_{\parallel})$ is blocked for the wave vectors \vec{k}_{\parallel} with no conduction channels in one of the spin states, i.e. $N_{\parallel}^{\sigma}(\vec{k}_{\parallel}) = 0$ for $\sigma = \uparrow$ or $\sigma = \downarrow$. These are the regions where $p_{\parallel} = \pm 1$ or $N_{\parallel}^{\uparrow} = N_{\parallel}^{\downarrow} = 0$ in Fig. 2c and d. As a result, only the states near the zone center where the spin channels are degenerate (enforced by \hat{G}_x and \hat{G}_y) contribute to the transmission (Fig. 4d). This leads to the total transmission being much smaller for the antiparallel state (T_{AP}) than for the parallel state (T_P) (Fig. 4e). At the Fermi energy E_F , we find the TMR ratio $(T_P - T_{AP})/T_{AP}$ as large as $\sim 500\%$. This value is comparable to the values obtained for the well-known Fe/MgO/Fe (001) MTJs^{39,40} which are currently used in magnetic random-access memories. The giant TMR appears not only for $E = E_F$ but also for the energies around the Fermi level, with the smallest value of $\sim 50\%$ at $E = E_F - 0.25$ eV (Fig. 4f). This fact indicates that the large TMR will be sustained under an applied bias voltage.

The predicted TMR is largely independent of the interface terminations and the relative alignment of the interface magnetic moments, as follows from our explicit DFT calculations (Supplementary Figs. S1 and S2). This distinguishes our results from the previous findings^{21,22}, where the interface termination controls TMR and implies that the predicted TMR effect is likely less sensitive to the interface roughness than that in the previous studies. The bulk origin of TMR in the proposed AFMTJs makes

it also more robust against other types of disorder, as long as the crystallinity of the tunnel junction and the direct tunneling transport mechanism are maintained (see Section C of Supplemental Material).

Discussion

The above properties are sustained in the presence of spin-orbit coupling. This is due to the momentum-dependent spin polarization in rutile antiferromagnets being inherited from the antiferromagnetic order rather than spin-orbit coupling^{27,31,37}. In RuO₂, the \hat{M}_z symmetry transformation reverses the wave vector component k_z , conserves the spin component σ_z but flips σ_x and σ_y . As a result, the conduction modes at \vec{k}_{\parallel} are spin polarized purely along the z axis and the x - and y -spin components vanish (see Supplementary Section D). Supplementary Fig. S5e shows the results of the DFT calculation for RuO₂ in the presence of spin-orbit coupling. It is seen that the spin polarization of the most conduction channels is well preserved, indicating that the giant TMR is robust against spin-orbit coupling in the RuO₂ based AFMTJ.

The spin-polarized conduction channels are not limited to RuO₂, but typical for a wide group of materials with violated $\hat{P}\hat{T}$ symmetry, including those with a noncollinear antiferromagnetic order²⁹. For the antiferromagnets with magnetization being compensated by combined mirror and/or rotation symmetries, the spin-polarized conduction channels can exist purely due to the antiferromagnetic order, even in the absence of spin-orbit coupling^{26–31}. On the other hand, in compensated

antiferromagnets with $\hat{T}\hat{t}$ symmetry, the spin-polarized conduction channels appear due to the spin degeneracy lifted by spin-orbit coupling. Such antiferromagnets can also be used in AFMTJs if they have sizable spin-orbit splitting.

These considerations can be expanded to normal metals with broken space inversion symmetry, where the band spin degeneracy is lifted by spin-orbit interaction. In these materials, the two conduction channels with opposite spin polarizations are linked by the time reversal symmetry operation. For example, in topological metal TaN^{41,42}, the conduction channels along the [001] direction carry the spin polarization pointing along the same [001] direction (Supplementary Fig. S6). This property is enforced by the \hat{M}_z mirror symmetry. While the momentum-dependent spin polarization in non-centrosymmetric normal metals is fixed by their crystal symmetry and band structure, they can be used in spintronics in conjunction with antiferromagnets. For example, the antiferromagnetic reference layer in the AFMTJ in Fig. 1d can be replaced by a normal metal layer. Alternatively, one can create an antiferromagnet/normal metal interface. In such systems with a single antiferromagnetic layer, the spintronic functionality is controlled by the Néel vector orientation that regulates a matching of the conduction channels in the antiferromagnetic and normal metal layers.

Note that in junctions with a single antiferromagnetic layer, reversal of the Néel vector is equivalent to the time-reversal transformation which does not change the resistance. However, the resistance changes with rotation of the Néel vector, resulting in a tunnelling anisotropic magnetoresistance (TAMR) effect⁴³.

Another possibility is to utilize non-centrosymmetric insulators as a tunneling barrier layer in an AFMTJ. Due to the broken space inversion symmetry and spin-orbit coupling, the evanescent states in these insulators are spin-polarized^{44,45}. Therefore, the Néel vector of the free antiferromagnetic layer can be used to control the matching between the propagating Bloch states in the antiferromagnetic electrode and the evanescent gap states in the barrier resulting in a TAMR effect. An additional useful functionality of this kind of tunnel junctions may be provided by a switchable polarization of the non-centrosymmetric insulating barrier layer if it is ferroelectric⁴⁵.

The proposed use of spin-neutral currents in spintronics is feasible from the experimental perspective. For example, the proposed RuO₂/TiO₂/RuO₂ (001) AFMTJ has all rutile structure with a good match of the RuO₂ and TiO₂ lattice constants and thus can be grown epitaxially preserving crystallinity of the overall heterostructure. The Néel vector of the antiferromagnetic free layer can be switched by a spin-orbit torque via the spin current from an adjacent heavy metal layer generated by an in-plane charge current^{8,46}. With the in-plane writing path and out-of-plane reading path, only two in-plane terminals and one out-of-plane terminal are required for such an AFMTJ, which is desirable for nanoscale spintronic applications. In addition, the large magnitude of TMR indicates a possibility of a strong spin transfer torque in the AFMTJs, which may be robust against disorder⁴⁷ and may offer an alternative way to switch the Néel vector.

In conclusion, we have proposed that globally spin-neutral currents flowing through oppositely spin-polarized conduction channels can be efficiently used in spintronics. Such currents exist in compensated antiferromagnets and normal metals with the magnetic space group symmetries which lift the spin-degeneracy of the Fermi surface. In the heterostructures, such as antiferromagnetic tunnel junctions or antiferromagnet/normal metal interfaces, these currents can be controlled by the Néel vector orientation providing a useful functionality for spintronics. Based on first-principles density functional theory combined with

quantum transport calculations, we have demonstrated such functionality using a room-temperature antiferromagnetic metal RuO₂ as electrodes in a RuO₂/TiO₂/RuO₂ AFMTJ and predicted a giant TMR effect of ~500%. Our work uncovers an unexplored potential of the materials with no global spin polarization for utilizing them in spintronics. We hope therefore that our predictions will stimulate experimental investigations of these materials and the associated phenomena.

Note added: During the review of this manuscript, we became aware of the relevant work by Šmejkal *et al.* posted recently⁴⁸.

Methods

The atomic and electronic structures shown in Figs. 2a, b, 3, S5a, and S6a, b of the systems are calculated using the projector augmented wave (PAW) method⁴⁹ implemented in the VASP code⁵⁰. A plane-wave cut-off energy of 500 eV and a $16 \times 16 \times 16$ \vec{k} -point mesh in the irreducible Brillouin zone are used in the calculations. The exchange and correlation effects are treated within the generalized gradient approximation (GGA) developed by Perdew-Burke-Ernzerhof (PBE)⁵¹. The GGA+U functional^{52,53} with $U_{\text{eff}} = 2$ eV on Ru 4d orbitals and $U_{\text{eff}} = 5$ eV on Ti 3d orbitals is included in the calculations.

The transport properties shown in Fig. 4 and S1–S3 are calculated using the non-equilibrium Green's function formalism (DFT+NEGF approach)^{54,55}, as implemented the Atomistic Simulation Toolkit (ATK) distributed in the QuantumWise package (Version 2015.1) (ATOMISTIX TOOLKIT version 2015.1 Synopsys QuantumWise (www.quantumwise.com)). QuantumWise A/S is now part of Synopsys, and from the upcoming version ATK will be part of the QuantumATK suite⁵⁶. The atomic structures are relaxed by VASP and the nonrelativistic Fritz-Haber-Institute (FHI) pseudopotentials using a single-zeta-polarized basis. The spin polarized GGA+U functional^{51,52} with $U_{\text{eff}} = 2.3$ eV on Ru 4d orbitals and $U_{\text{eff}} = 5$ eV on Ti 3d orbitals is included in the calculations. A cut-off energy of 75 Ry and a $11 \times 11 \times 101$ \vec{k} -point mesh are used for the self-consistent calculations to eliminate the mismatch of the Fermi level between the electrodes and the central region. Unless mentioned in the text, the transmission is calculated using an adaptive \vec{k} -point mesh. These parameters are confirmed to yield a good balance between the computational time and accuracy.

The tight-binding Hamiltonians of RuO₂ and TaN are obtained using Wannier90 code⁵⁷ utilizing the maximally localized Wannier functions⁵⁸. A $500 \times 500 \times 500$ \vec{k} -point mesh and the adaptive smearing method⁵⁹ are used to calculate the \vec{k}_{\parallel} -resolved ballistic conductance shown in Fig. 2c, d, S5e and S6c, d. The spin-projected Fermi surfaces of RuO₂ with spin-orbit coupling shown in Figs. S5b–d are calculated using WannierBerri code^{60,61}.

Figures are plotted using VESTA⁶², FermiSurfer⁶³, gnuplot⁶⁴, and the SciDraw scientific figure preparation system⁶⁵.

Data availability

The data that support the findings of this study are available from the corresponding author upon reasonable request.

Received: 29 March 2021; Accepted: 25 October 2021;

Published online: 03 December 2021

References

1. Tsymbal, E. Y. & Žutić I. *Spintronics Handbook: Spin Transport and Magnetism*, 2-nd edition. (CRC press, 2019).
2. Julliere, M. Tunneling between ferromagnetic films. *Phys. Lett.* **54A**, 225–226 (1975).
3. Moodera, J. S., Kinder, L. R., Wong, T. M. & Meservey, R. Large magnetoresistance at room temperature in ferromagnetic thin film tunnel junctions. *Phys. Rev. Lett.* **74**, 3273–3276 (1995).
4. Tsymbal, E. Y., Mryasov, O. N. & LeClair, P. R. Spin-dependent tunneling in magnetic tunnel junctions. *J. Phys.: Condens. Matter* **15**, R109–R142 (2003).
5. Baltz, V. et al. Antiferromagnetic spintronics. *Rev. Mod. Phys.* **90**, 015005 (2018).
6. Jungwirth, T., Marti, X., Wadley, P. & Wunderlich, J. Antiferromagnetic spintronics. *Nat. Nanotech.* **11**, 231–241 (2016).
7. Jungwirth, T. et al. The multiple directions of antiferromagnetic spintronics. *Nat. Phys.* **14**, 200–203 (2018).
8. Železný, J., Wadley, P., Olejník, K., Hoffmann, A. & Ohno, H. Spin transport and spin torque in antiferromagnetic devices. *Nat. Phys.* **14**, 220–228 (2018).
9. Šmejkal, L., Mokrousov, Y., Yan, B. & MacDonald, A. H. Topological antiferromagnetic spintronics. *Nat. Phys.* **14**, 242–251 (2018).

10. Wadley, P. et al. Electrical switching of an antiferromagnet. *Science* **351**, 587–590 (2016).
11. Bodnar, S. Y. et al. Writing and reading antiferromagnetic Mn₂Au by Néel spin-orbit torques and large anisotropic magnetoresistance. *Nat. Commun.* **9**, 348 (2018).
12. Hoogeboom, G. R., Aqeel, A., Kuschel, T., Palstra, T. T. M. & van Wees, B. J. Negative spin Hall magnetoresistance of Pt on the bulk easy-plane antiferromagnet NiO. *Appl. Phys. Lett.* **111**, 052409 (2017).
13. Chen, X. Z. et al. Antidamping-torque-induced switching in biaxial antiferromagnetic insulators. *Phys. Rev. Lett.* **120**, 207204 (2018).
14. Fischer, J. et al. Spin Hall magnetoresistance in antiferromagnet/heavy-metal heterostructures. *Phys. Rev. B* **97**, 014417 (2018).
15. Baldrati, L. et al. Full angular dependence of the spin Hall and ordinary magnetoresistance in epitaxial antiferromagnetic NiO(001)/Pt thin films. *Phys. Rev. B* **98**, 024422 (2018).
16. Chiang, C. C., Huang, S. Y., Qu, D., Wu, P. H. & Chien, C. L. Absence of evidence of electrical switching of the antiferromagnetic Néel vector. *Phys. Rev. Lett.* **123**, 227203 (2019).
17. Núñez, A. S., Duine, R. A., Haney, P. & MacDonald, A. H. Theory of spin torques and giant magnetoresistance in antiferromagnetic metals. *Phys. Rev. B* **73**, 214426 (2006).
18. Saidaoui, H. B. M., Manchon, A. & Waintal, X. Spin transfer torque in antiferromagnetic spin valves: From clean to disordered regimes. *Phys. Rev. B* **89**, 174430 (2014).
19. Xu, Y., Wang, S. & Xia, K. Spin-transfer torques in antiferromagnetic metals from first principles. *Phys. Rev. Lett.* **100**, 226602 (2008).
20. Manchon, A. Spin diffusion and torques in disordered antiferromagnets. *J. Phys. Condens. Matter* **29**, 104002 (2017).
21. Merodio, P., Kalitsov, A., Ba, H., Baltz, V. & Chshiev, M. Spin-dependent transport in antiferromagnetic tunnel junctions. *Appl. Phys. Lett.* **105**, 122403 (2014).
22. Stamenova, M., Mohebbi, R., Seyed-Yazdi, J., Rungger, I. & Sanvito, S. First-principles spin-transfer torque in CuMnAl/GaP/CuMnAs junctions. *Phys. Rev. B* **95**, 060403 (2017).
23. Šmejkal, L., Železný, J., Sinova, J. & Jungwirth, T. Electric control of Dirac quasiparticles by spin-orbit torque in an antiferromagnet. *Phys. Rev. Lett.* **118**, 106402 (2017).
24. Shao, D.-F., Gurung, G., Zhang, S.-H. & Tsymbal, E. Y. Dirac nodal line metal for topological antiferromagnetic spintronics. *Phys. Rev. Lett.* **122**, 077203 (2019).
25. Shao, D.-F., Zhang, S.-H., Gurung, G., Yang, W. & Tsymbal, E. Y. Nonlinear anomalous Hall effect for Néel vector detection. *Phys. Rev. Lett.* **124**, 067203 (2020).
26. Hayami, S., Yanagi, Y. & Kusunose, H. Momentum-dependent spin splitting by collinear antiferromagnetic ordering. *J. Phys. Soc. Jpn.* **88**, 123702 (2019).
27. Yuan, L.-D., Wang, Z., Luo, J.-W., Rashba, E. I. & Zunger, A. Giant momentum-dependent spin splitting in centrosymmetric low-Z antiferromagnets. *Phys. Rev. B* **102**, 014422 (2020).
28. Yuan, L.-D., Wang, Z., Luo, J.-W. & Zunger, A. Prediction of low-Z collinear and noncollinear antiferromagnetic compounds having momentum-dependent spin splitting even without spin-orbit coupling. *Phys. Rev. Mater.* **5**, 014409 (2021).
29. Železný, J., Zhang, Y., Felser, C. & Yan, B. Spin-polarized current in noncollinear antiferromagnets. *Phys. Rev. Lett.* **119**, 187204 (2017).
30. Naka, M. et al. Spin current generation in organic antiferromagnets. *Nat. Commun.* **10**, 4305 (2019).
31. González-Hernández, R. et al. Efficient electrical spin-splitter based on non-relativistic collinear antiferromagnetism. *Phys. Rev. Lett.* **126**, 127701 (2021).
32. Landauer, R. Spatial variation of currents and fields due to localized scatterers in metallic conduction. *IBM J. Res. Dev.* **32**, 306–316 (1988).
33. Datta, S. *Electronic Transport in Mesoscopic Systems* (Cambridge University Press, 1995).
34. Schep, K. M., Kelly, P. J. & Bauer, G. E. W. Ballistic transport and electronic structure. *Phys. Rev. B* **57**, 8907–8926 (1998).
35. Berlijn, T. et al. Itinerant antiferromagnetism in RuO₂. *Phys. Rev. Lett.* **118**, 077201 (2017).
36. Ahn, K.-H., Hariki, A., Lee, K.-W. & Kuneš, J. RuO₂ Antiferromagnetism in as *d*-wave Pomeranchuk instability. *Phys. Rev. B* **99**, 184432 (2019).
37. Šmejkal, L., González-Hernández, R., Jungwirth, T. & Sinova, J. Crystal Hall effect in collinear antiferromagnets. *Sci. Adv.* **6**, eaaz8809 (2020).
38. Bose, A. et al. Tilted spin current generated by the collinear antiferromagnet RuO₂. *arXiv* **2108**, 09150 (2021).
39. Yuasa, S., Nagahama, T., Fukushima, A., Suzuki, Y. & Ando, K. Giant room-temperature magnetoresistance in single-crystal Fe/MgO/Fe magnetic tunnel junctions. *Nat. Mater.* **3**, 868–871 (2004).
40. Parkin, S. S. P. et al. Giant tunnelling magnetoresistance at room temperature with MgO (100) tunnel barriers. *Nat. Mater.* **3**, 862–867 (2004).
41. Zhu, Z., Winkler, G. W., Wu, Q., Li, J. & Soluyanov, A. A. Triple point topological metals. *Phys. Rev. X* **6**, 031003 (2016).
42. Weng, H., Fang, C., Fang, Z. & Dai, X. Topological semimetals with triply degenerate nodal points in θ -phase tantalum nitride. *Phys. Rev. B* **93**, 241202 (2016).
43. Park, B. G. et al. & Jungwirth, T. A spin-valve-like magnetoresistance of an antiferromagnet-based tunnel junction. *Nat. Mater.* **10**, 347–351 (2011).
44. Mishra, S., Thulasi, S. & Satpathy, S. Spin polarization via electron tunneling through an indirect-gap semiconductor barrier. *Phys. Rev. B* **72**, 195347 (2005).
45. Tao, L. L. & Tsymbal, E. Y. Perspectives of spin-textured ferroelectrics. *J. Phys. D: Appl. Phys.* **54**, 113001 (2020).
46. Manchon, A. et al. Current-induced spin-orbit torques in ferromagnetic and antiferromagnetic systems. *Rev. Mod. Phys.* **91**, 035004 (2019).
47. Saidaoui, H., Manchon, A. & Waintal, X. Robust spin transfer torque in antiferromagnetic tunnel junctions. *Phys. Rev. B* **95**, 134424 (2017).
48. Šmejkal, L., Hellenes, A. B., González-Hernández, R., Sinova, J. & Jungwirth, T. Giant and tunneling magnetoresistance effects from anisotropic and valley-dependent spin-momentum interactions in antiferromagnets. *arXiv* **2103**, 12664 (2021).
49. Kresse, G. & Joubert, D. From ultrasoft pseudopotentials to the projector augmented-wave method. *Phys. Rev. B* **59**, 1758–1775 (1999).
50. Kresse, G. & Furthmüller, J. Efficient iterative schemes for ab initio total-energy calculations using a plane-wave basis set. *Phys. Rev. B* **54**, 11169–11186 (1996).
51. Perdew, J. P., Burke, K. & Ernzerhof, M. Generalized gradient approximation made simple. *Phys. Rev. Lett.* **77**, 3865–3868 (1996).
52. Dudarev, S. L., Botton, G. A., Savrasov, S. Y., Humphreys, C. J. & Sutton, A. P. Electron-energy-loss spectra and the structural stability of nickel oxide: An LSDA+U study. *Phys. Rev. B* **57**, 1505–1509 (1998).
53. Anisimov, V. I., Zaanen, J. & Andersen, O. K. Band theory and Mott insulators: Hubbard U instead of Stoner. *I. Phys. Rev. B* **44**, 943–954 (1991).
54. Taylor, J., Guo, H. & Wang, J. Ab initio modeling of quantum transport properties of molecular electronic devices. *Phys. Rev. B* **63**, 245407 (2001).
55. Brandbyge, M., Mozos, J. L., Ordejón, P., Taylor, J. & Stokbro, K. Density-functional method for nonequilibrium electron transport. *Phys. Rev. B* **65**, 165401 (2002).
56. Smidstrup, S. et al. QuantumATK: An integrated platform of electronic and atomic-scale modelling tools. *J. Phys. Condens. Matter* **32**, 015901 (2019).
57. Pizzi, G. et al. Wannier90 as a community code: new features and applications. *J. Phys. Cond. Mat.* **32**, 165902 (2020).
58. Marzari, N., Mostofi, A. A., Yates, J. R., Souza, I. & Vanderbilt, D. Maximally localized Wannier functions: Theory and applications. *Rev. Mod. Phys.* **84**, 1419 (2012).
59. Yates, J. R., Wang, X., Vanderbilt, D. & Souza, I. Spectral and Fermi surface properties from Wannier interpolation. *Phys. Rev. B* **75**, 195121 (2007).
60. Tsirkin, S. S. High performance Wannier interpolation of Berry curvature and related quantities with WannierBerri code. *npj Comput. Mater.* **7**, 33 (2021).
61. Destraz, D. et al. Magnetism and anomalous transport in the Weyl semimetal PrAlGe: Possible route to axial gauge fields. *npj Quantum Mater.* **5**, 5 (2020).
62. Momma, K. & Izumi, F. VESTA 3 for three-dimensional visualization of crystal, volumetric and morphology data. *J. Appl. Crystallogr.* **44**, 1272–1276 (2011).
63. Kawamura, M. FermiSurfer: Fermi-surface viewer providing multiple representation schemes. *Comp. Phys. Commun.* **239**, 197–203 (2019).
64. Williams T. & Kelley, C. Gnuplot 5.2: An interactive plotting program, <http://www.gnuplot.info> (2017).
65. Caprio, M. A. LevelScheme: a level scheme drawing and scientific figure preparation system for Mathematica. *Comp. Phys. Commun.* **171**, 107–118 (2005).

Acknowledgements

The authors thank Bo Li for helpful discussions. This work was supported by the Vannevar Bush Faculty Fellowship (ONR grant N00014-20-1-2844) (C.B.E) and by the National Science Foundation (NSF) through the MRSEC (NSF Award DMR-1420645) and EPSCoR RII Track-1 (NSF Award OIA-2044049) programs (E.Y.T.). S.-H.Z. thanks the support of National Science Foundation of China (NSFC Grant No. 12174019). Computations were performed at the University of Nebraska Holland Computing Center.

Author contributions

D.-F.S. and E.Y.T. conceived this project. D.-F.S. performed symmetry analysis and calculated electronic properties for RuO₂ and TaN. S.H.Z. calculated the ballistic conductance of RuO₂ and TaN. D.-F.S. and M.L. calculated the transport properties of RuO₂/TiO₂/RuO₂ AFMTJ. D.F.S., S.H.Z., M.L. and E.Y.T. analyzed the results and discussed with C.B.E. D.-F.S. and E.Y.T. wrote the manuscript. All authors contributed to the final version of the manuscript.

Competing interests

The authors declare no competing interests.

Additional information

Supplementary information The online version contains supplementary material available at <https://doi.org/10.1038/s41467-021-26915-3>.

Correspondence and requests for materials should be addressed to Ding-Fu Shao or Evgeny Y. Tsymbal.

Peer review information *Nature Communications* thanks the anonymous reviewer(s) for their contribution to the peer review of this work.

Reprints and permission information is available at <http://www.nature.com/reprints>

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>.

© The Author(s) 2021