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First-principles investigation of possible room-temperature topological insulators in monolayers

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A Quantum Spin Hall (QSH) insulator with a large bulk band gap and tunable topological properties is crucial for both fundamental research and practical application. Chemical functionalization has been proposed as an effective route to realize the QSH effect. Using the ABINIT package, we have investigated the properties of (1) TIP, the functionalized monolayers TIPX_2 ($X = \text{F}, \text{Cl}, \text{Br}, \text{I}$); (2) TIAs, the functionalized monolayers TIAsX_2 ($X = \text{F}, \text{Cl}, \text{Br}, \text{I}$), and (3) GaGeTe, InGeTe, and InSnTe systems. The topological nature is verified by the calculation of the Z_2 topological invariant. We discovered TIPF_2 , TIPCl_2 , TIPBr_2 , TIPI_2 , TIAs, TIAsF_2 , TIAsCl_2 , TIAsBr_2 , and TIAsI_2 were promising 2D TIs with bulk band gaps as large as 0.21 eV. Each monolayer was suitable for room-temperature application, and show great potential for their future applications in quantum computers, nanoelectronics, and spintronics.

1. Introduction

Topological insulators (TIs) are a class of materials that have created a surge of research activities in the past decade.^{1–4} Due to their intriguing properties and promising applications in spintronics and quantum computations, they are attracting worldwide interest.^{5–8} The term “topological insulator” was coined by Moore and Balents in their paper to propose the existence of TIs in 3-Dimensional (3D) systems,^{1,9} but their story started long before the discovery of 3D TIs. The Quantum Hall System, discovered in 1980, is considered to be the first TI that became known to physicists.¹ The Spin Hall Effect was experimentally confirmed by Kato *et al.* in 2004,^{1,10} which led Murakami, Nagaosa, and Zhang to propose the idea of a Spin Hall Insulator,¹¹ a gapped insulator with zero charge conductivity but with a finite spin Hall conductivity due to a finite Berry phase of the occupied states.¹² Although this proposal could not generate spin currents in the absence of any electrons at the Fermi level, it triggered Kane and Mele’s proposal of its quantized version, the Quantum Spin Hall (QSH) insulator.^{1,13}

Many materials have been predicted to be TIs, but only some have been addressed experimentally. The first material that was experimentally identified as a time-reversal (TR) invariant TI was the CdTe/HgTe/CdTe quantum well,^{1,14} or a thin layer of HgTe sandwiched by CdTe. Afterwards, the AlSb/InAs/GaSb/AlSb quantum well was theoretically predicted¹⁵ and experimentally confirmed^{16,17} to be a 2D TI system. The first 3D TI material that was experimentally identified was $\text{Bi}_{1-x}\text{Sb}_x$,¹⁸ an alloy of Bi and Sb. Discoveries of both 2D and 3D TI materials are ongoing and strongly called for. However, 2D TIs have both

an insulating bulk and conducting edge states that display unique advantages over 3D TIs with regard to flexibility, a higher charge carrier mobility, and controllability owing to their atomic thickness.⁵

The last decade has seen an immense development of interest in monolayer materials.¹⁹ A monolayer forms when the thickness of material is reduced down to a single atom.¹ A common feature of materials that show band structures with room temperature 2D TI properties are they most likely have 2D hexagonal honeycomb-like crystal structures, indicating that a 2D hexagonal lattice could be an excellent cradle to breed QSH insulators with the influence of spin-orbit coupling (SOC).⁵ TLX ($X = \text{N}, \text{P}, \text{As}, \text{Sb}$) monolayers were recently reported to possess structural stability.^{5,19} First-principles calculations confirmed that a two-dimensional TIP monolayer could convert into a topological insulator with the effect of bromination accompanied by a large bulk band gap of 76.5 meV, which meets the requirement for room-temperature application.²⁰

GaGeTe, InSnTe, and InGeTe were promising monolayers due to GaGeTe’s high carrier mobility and tunable band structure.²¹ Each monolayer has a layered crystal structure stacked from six-atom thick building blocks. First-principles calculations have identified the GaGeTe-type periodic structures as a potential host for topological phases.²² The layered In-SnTe bulk material is predicted to be a 3D strong topological insulator with $Z_2 = 1$; (111).²²

Additionally, the tetrahedral atomic coordination in the GaGeTe-type structures closely resembles the topological materials with the diamond-like cubic lattices.^{14,23,24} Possible ways to induce topological order could be doping GaGeTe with larger isovalent p-elements, such as In and Sn.

The crucial bottleneck of reported 2D TIs is their small bulk band gaps, which are too weak to be reflected in modern

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experimental conditions. Graphene, for example, was the first material predicted to realize a TI, but the gap was unobservably small due to carbon's weak spin-orbital coupling.²⁵ A large bulk band gap is crucial for protecting the edge current against the interference of thermally activated carriers.⁵ Fortunately, chemical functionalization of 2D materials is a powerful tool for creating new materials with desirable features.²⁶ Chemical functionalization of topological insulator monolayers is an effective method of tuning the band gap, while preserving the nontrivial topological order.^{27–29} For example, pristine stanene has a band gap of 0.1 eV, but with functional groups, the band gap reaches 0.3 eV.²⁷ This research studies possible 2D TIs by performing first-principles calculations on both monolayers and chemically functionalized monolayers. We performed calculations on the monolayers TlP and TlAs, which were chemically functionalized with halogens, and on the monolayers GaGeTe, InSnTe, and InGeTe. These findings may endow the monolayers with the potential to fabricate new quantum devices operating at room temperature in nanoelectronics and spintronics.

2. Method

We performed first-principle calculations based on Density Functional Theory (DFT) using the Generalized Gradient Approximation (GGA) exchange–correlation in the Perdew–Burke–Ernzerhof (PBE)³⁰ format implemented in the ABINIT^{31,32} code. We use the Projected Augmented Wave (PAW) method³³ with projectors generated using the AtomPAW code.^{34,35} The electron configurations and radial cutoffs used to generate the PAW pseudopotentials are shown in Table 1.

In total energy calculations, self-consistent cycles were recognized when the total energy difference was less than 1.0×10^{-5} Hartree twice consecutively. The kinetic energy cutoff, the Monkhorst–Pack grid, and the vacuum height of the unit cell were converged for each monolayer. The converged values corresponding to each monolayer were used for calculations with and without SOC.

The Broyden–Fletcher–Goldfarb–Shanno (BFGS) minimization algorithm was used to perform structural optimization.

Table 1 Electron configurations and radial cutoffs used to generate the PAW pseudopotentials for the elements used in this study

Element	Atomic number	Electron configuration	Radius cutoff (Bohr)
Tl	81	[Xe 4f ¹⁴] 6s ² 6p ¹ 5d ¹⁰	2.42
P	15	[Ne] 3s ² 3p ³	1.91
As	33	[Ar] 4s ² 4p ³ 3d ¹⁰	2.10
F	9	[He] 2s ² 2p ⁵	1.40
Cl	17	[Ne] 3s ² 3p ⁵	1.80
Br	35	[Ar 3d ¹⁰] 4s ² 4p ⁵	2.20
I	53	[Kr 4d ¹⁰] 5s ² 5p ⁵	2.30
Ga	31	[Ar] 4s ² 4p ¹ 3d ¹⁰	2.10
Ge	32	[Ar] 4s ² 4p ² 3d ¹⁰	2.30
Te	52	[Kr 4d ¹⁰] 5s ² 5p ⁴	2.31
In	49	[Kr] 5s ² 5p ¹ 4d ¹⁰	2.51
Sn	50	[Kr] 5s ² 5p ² 4d ¹⁰	2.51

The atomic structure was relaxed until the maximum atomic forces were less than 5.0×10^{-5} Hartree Bohr^{−1}. The relaxed structure corresponding to each monolayer was used for band structure calculations with and without SOC.

The Z_2 topological invariant was computed by tracking the evolution of the hybrid Wannier charge centers by implementing the Wannier 90 package^{36,37} and the Z2Pack software package.^{38–40} The Wannier charge centers are based on the notion of Wannier orbitals.

These are given by Fourier transforming the Bloch States

$$|Rn\rangle = \frac{V^r}{(2\pi)^d} BZ^e \cdot \text{ik} \cdot R |\psi_{n,k}\rangle dk \quad (1)$$

where d is the dimensionality of the system and V is the unit cell volume. These orbitals can be changed by a Gauge transformation which affects their localization and position in real space. To compute topological invariants, hybrid Wannier orbitals are introduced: they are Fourier transforms performed only in one spatial direction, for example

$$|R_x, k_y, k_z; n\rangle = \frac{a_x^r}{(2\pi)^{-\pi/a_x}} e^{-ik_y R_x} |\psi_{nk}\rangle \quad (2)$$

3. Results and discussion

The atomic structure, band structures with and without SOC, and Z_2 topologies of each monolayer will be described for TlP, TlAs, GaGeTe, InGeTe, InSnTe, and the chemically functionalized monolayers TlPX₂ and TlAsX₂, where X = F, Cl, Br, and I. The band structures were calculated using the high-symmetry k -points M (1/2, 1/2, 0.0), K (2/3, 1/3, 0.0), 2 2 3 3 and Γ (0.0, 0.0, 0.0).

3.1. TlPX₂ (X = F, Cl, Br, I)

The side and top views of the atomic structures are presented in Fig. 1. The monolayers have a honeycomb lattice. Similar to silicene, the Tl and P atoms occupy two sublattices with a buckled height of 1.14 Bohr and a lattice constant of 8.14 Bohr. It is characterized by the sp² hybridization. The calculated lattice constants and bond lengths of each monolayer can be found in Table 2. When TlP is chemically functionalized, each Tl (P) atom is bonded to a X atom and three P (Tl) atoms, which is analogous to a typical III–V bulk counterpart. Thus, the sp³ hybridization is naturally formed, leading to an increase in the buckled height and bond lengths. Our results confirm that chemically functionalizing TlP leads to a consistent increase in each monolayer's buckled height, in agreement with previous results.²⁰ The buckled height increased by 0.1 Bohr, 0.36 Bohr, 0.39 Bohr, and 0.39 Bohr respectively for the monolayers TlPF₂, TlPCL₂, TlPBr₂, and TlPI₂. A similar phenomenon has been observed in previous research.²⁰ The bond length Tl–P increased from 4.84 Bohr for the monolayer TlP to 5.12 Bohr, 5.11 Bohr, 5.12 Bohr, and 5.16 Bohr for the chemically functionalized monolayers TlPF₂, TlPCL₂, TlPBr₂, and TlPI₂, respectively.



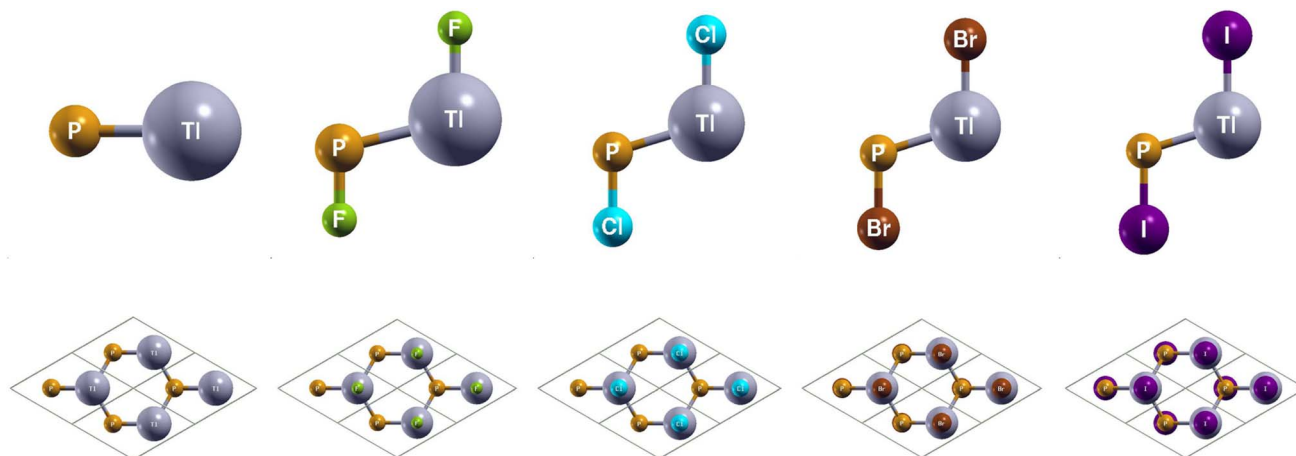


Fig. 1 Optimized atomic structures of TIP and TIPX_2 ($X = \text{F}, \text{Cl}, \text{Br}, \text{I}$) are shown from left to right. The atomic structures can be identified by the labels on each atom. Here, gray, orange, green, cyan, brown, and purple represent Tl, P, F, Cl, Br, and I atoms, respectively.

Table 2 Calculated optimized structural parameters of TIP and TIPX_2 ($X = \text{F}, \text{Cl}, \text{Br}, \text{I}$): the lattice constant (a) in Bohr, bond length P – X ($d_{\text{P-X}}$) in Bohr, bond length Tl – P ($d_{\text{Tl-P}}$) in Bohr, bond length Tl – X ($d_{\text{Tl-X}}$) in Bohr, and buckled height (d_b) in Bohr. A dash denotes the information is not applicable. Band gaps and topologies of TIP and TIPX_2 , including the band gap without SOC E_g in electron-volts and the band gap with SOC $E_{g-\text{SOC}}$ in electron-volts. d and id stand for direct and indirect band gap, respectively. Z_2 is the topological index

System	a (Bohr)	$d_{\text{P-X}}$ (Bohr)	$d_{\text{Tl-P}}$ (Bohr)	$d_{\text{Tl-X}}$ (Bohr)	d_b (Bohr)	E_g (eV)	$E_{g-\text{SOC}}$ (eV)	Z_2
TIP	8.14	—	4.84	—	1.14	0.23 (d)	0.18 (d)	0
TIPF_2	8.60	3.07	5.12	3.90	1.24	0.00 (d)	0.04 (id)	1
TIPCl_2	8.46	3.89	5.11	4.60	1.50	0.00 (d)	0.03 (id)	1
TIPBr_2	8.46	4.22	5.12	4.87	1.53	0.00 (d)	0.07 (id)	1
TIPI_2	8.54	4.65	5.16	5.27	1.53	0.00 (d)	0.15 (id)	1

Next, we turn towards the electronic properties of TIP and the chemically functionalized TIP monolayers. The band structures with and without an SOC effect are displayed in Fig. 2. Without applying SOC, TIP is a semiconductor with a small direct band gap of 0.23 eV at the Γ point. TIPF_2 , TIPCl_2 , TIPBr_2 , and TIPI_2 have zero energy band gaps with the valence band maximum (VBM) and the conduction band minimum (CBM) degenerate at the Γ point, forming a single Dirac point. The band gaps are displayed in Table 2.

Typically, the effect of SOC on the electronic structures of semiconductors is negligible.⁴¹ However, it is crucial for the case of semimetals or metallic systems. When SOC is taken into account, TIPF_2 , TIPCl_2 , TIPBr_2 , and TIPI_2 have indirect energy gaps of 0.04 eV, 0.03 eV, 0.07 eV, and 0.15 eV, respectively. Unlike the chemically functionalized monolayers, TIP's valence band and conduction band appear relatively closer together with SOC and the band gap decreases from 0.23 eV to 0.18 eV. The band gaps of TIP and the chemically functionalized monolayers, TIPF_2 , TIPCl_2 , TIPBr_2 , and TIPI_2 , most likely change dramatically due to the strong SOC of the Tl and P atoms.⁵ The SOC-induced energy gaps are larger than the thermal energy at room temperature (0.026 eV), implying that the band gaps for each monolayer would be feasibly measured at room temperature.

The band structures of TIP and the TIPX_2 monolayers are projected on the p orbitals of the elements. These band structures can be found in the third column of Fig. 2. Band inversion is absent in TIP, and present in some of the other chemically functionalized TIP monolayers. Band inversion has long been cited as a strong indicator of a topological insulator.^{20,41} However, our results confirm the prediction that band inversion is conceptually not a necessary outcome of topological phase transitions.⁴²

To verify our curiosities about the QSH state in the chemically functionalized TIP monolayers, the Z_2 topological invariant is an important decision criteria for a material's classification as a TI.²⁰ We directly calculate the Z_2 topological invariants and list them in Table 2. Fig. 3 shows the trajectories of the Wannier charge centers at the surfaces $k_z = 0$ and $k_z = 0.5$. The hybrid Wannier charge centers (HWCC) form bands, similar to the band structure of a dispersion relation. The Z_2 index can be calculated using the Wilson loop using the evolution of the Wannier charge centers. We calculate the Z_2 invariants on a coarse k-mesh, considering the points marking the middle of the largest gap at each k_x . These points are marked by blue diamonds on Fig. 3. Whenever the location of the middle of the gap changes between two adjacent k_x values, we count the number of HWCC that exist between the two gap centers and



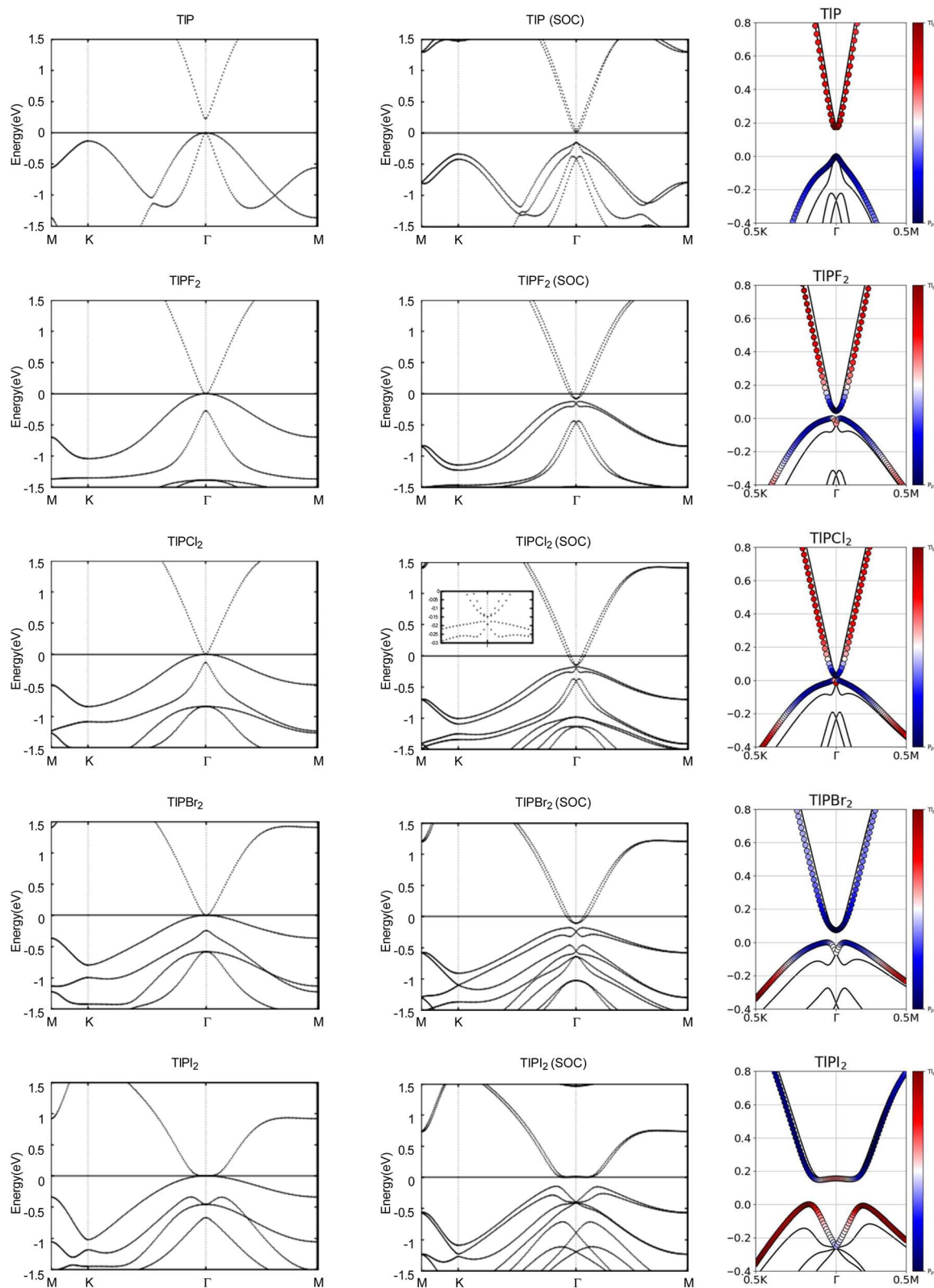


Fig. 2 Calculated band structures with and without SOC of TIP and TIPX₂ (X = F, Cl, Br, I), with the orbital-resolved band structures with SOC. The Fermi energy is set to 0 eV. The red and blue denote contributions from the 6p and 3p orbitals, respectively. The band structures can be identified by their titles.



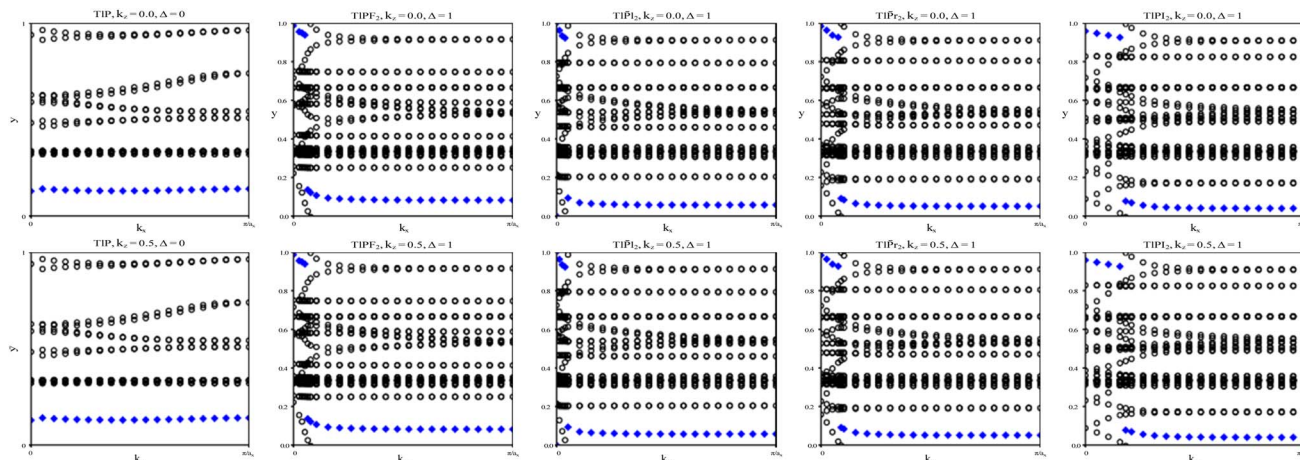


Fig. 3 The trajectories of the Wannier charge centers for TIP and TIPX_2 ($X = \text{F}, \text{Cl}, \text{Br}, \text{I}$). The open circles represent hybrid Wannier charge centers. The first row is at the surface $k_z = 0$. The second row is at the surface $k_z = 0.5$. The plots can be identified by their titles.

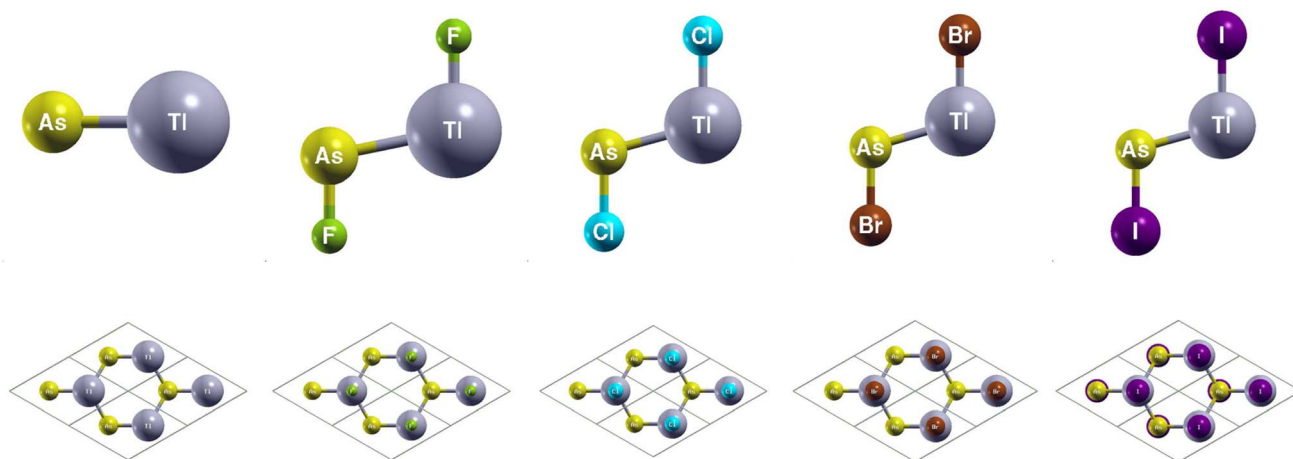


Fig. 4 Optimized atomic structures of TIAs and TIAsX_2 ($X = \text{F}, \text{Cl}, \text{Br}, \text{I}$) monolayers are shown from left to right. The atomic structures can be identified by the labels on each atom. Here, gray, yellow, green, cyan, brown, and purple represent TI, P, F, Cl, Br, and I atoms, respectively.

sum this number for all the crossings as k_x goes from 0 to π . If this value is even, $Z_2 = 0$, and if its odd, $Z_2 = 1$. The Z_2 calculations confirm that TlPF_2 , TlPcl_2 , TlPBr_2 , and TlPI_2 are promising 2D TIs, with $Z_2 = 1$, while TIP is a normal insulator, with $Z_2 = 0$. Our results for TIP corroborate Li *et al.*'s calculations with the plane-wave basis Vienna *ab initio* simulation package (VASP).⁵

3.2. TIAsX_2 ($X = \text{F}, \text{Cl}, \text{Br}, \text{I}$)

You can find the side and top views of the atomic structures in Fig. 4. The monolayers have a honeycomb lattice. Similar to TIP, the TI and As atoms occupy two sublattices with a buckled height of 1.32 Bohr and a lattice constant of 8.14 Bohr. The calculated lattice constants and bond lengths of each monolayer can be found in Table 3. When TIAs is chemically

Table 3 Calculated optimized structural parameters of TIAs and TIAsX_2 ($X = \text{F}, \text{Cl}, \text{Br}, \text{I}$): the lattice constant (a) in Bohr, bond length As – X ($d_{\text{As-X}}$) in Bohr, bond length TI – As ($d_{\text{TI-As}}$) in Bohr, bond length TI – X ($d_{\text{TI-X}}$) in Bohr, and buckled height (d_b) in Bohr. A dash denotes the information is not applicable. Band gaps and topologies of TIAs and TIAsX_2 , including the band gap without SOC E_g in electron-volts and the band gap with SOC $E_{g-\text{SOC}}$ in electron-volts. d and id stand for direct and indirect band gap, respectively. Z_2 is the topological index

System	a (Bohr)	$d_{\text{As-X}}$ (Bohr)	$d_{\text{TI-As}}$ (Bohr)	$d_{\text{TI-X}}$ (Bohr)	d_b (Bohr)	E_g (eV)	$E_{g-\text{SOC}}$ (eV)	Z_2
TIAs	8.14	—	5.06	—	1.32	0.01 (d)	0.14 (id)	1
TIAsF_2	8.60	3.35	5.32	3.93	1.07	0.00 (d)	0.18 (id)	1
TIAsCl_2	8.46	4.13	5.31	4.63	1.40	0.00 (d)	0.15 (id)	1
TIAsBr_2	8.46	4.43	5.31	4.90	1.49	0.00 (d)	0.18 (id)	1
TIAsI_2	8.54	4.86	5.34	5.35	1.44	0.00 (id)	0.21 (id)	1



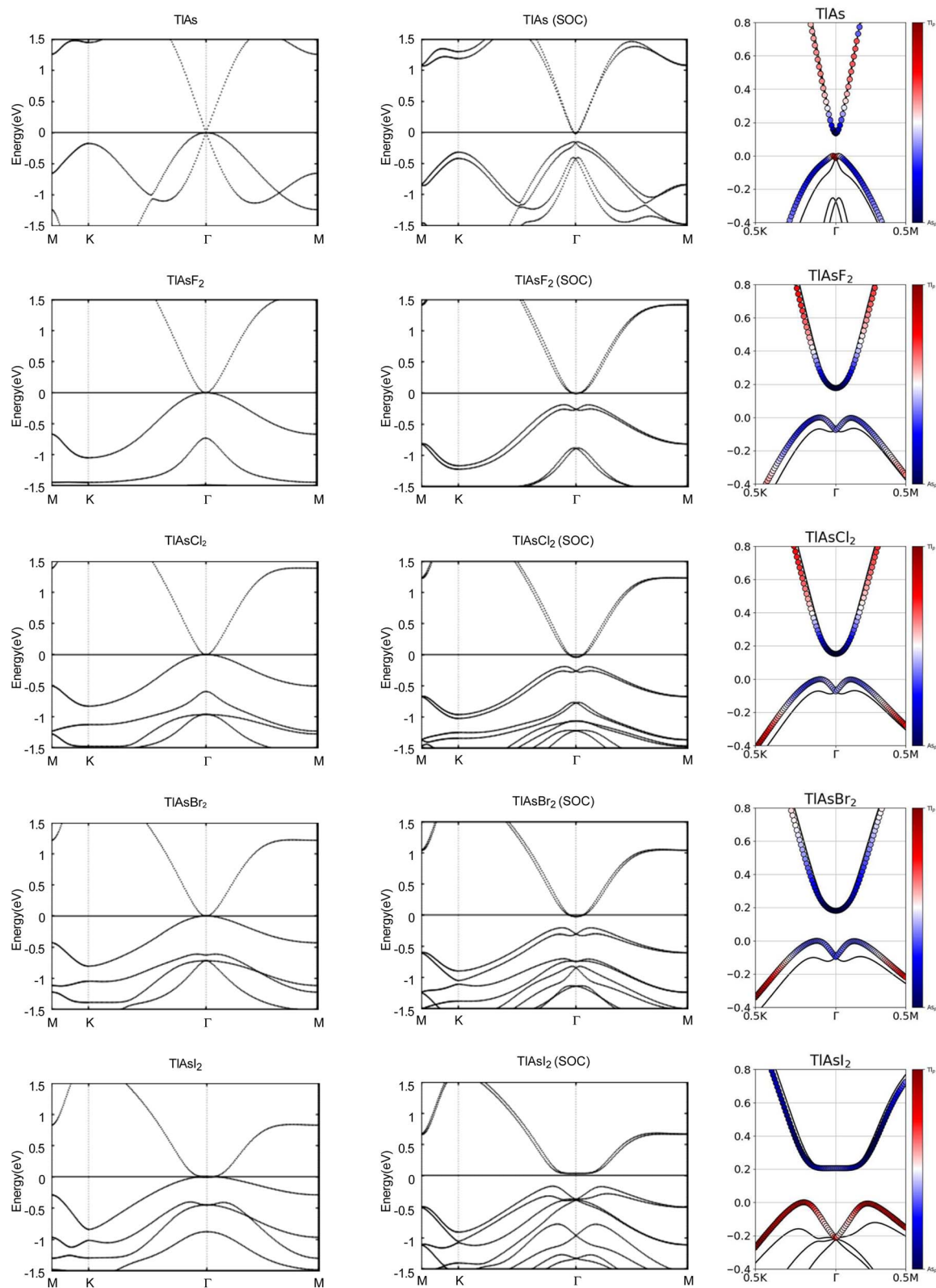


Fig. 5 Calculated band structures with and without SOC of TIAs and TIAsX₂ (X = F, Cl, Br, I), with the orbital-resolved band structures with SOC. The Fermi energy is set to 0 eV. The red and blue denote contributions from the 6p and 4p orbitals, respectively. The band structures can be identified by their titles.



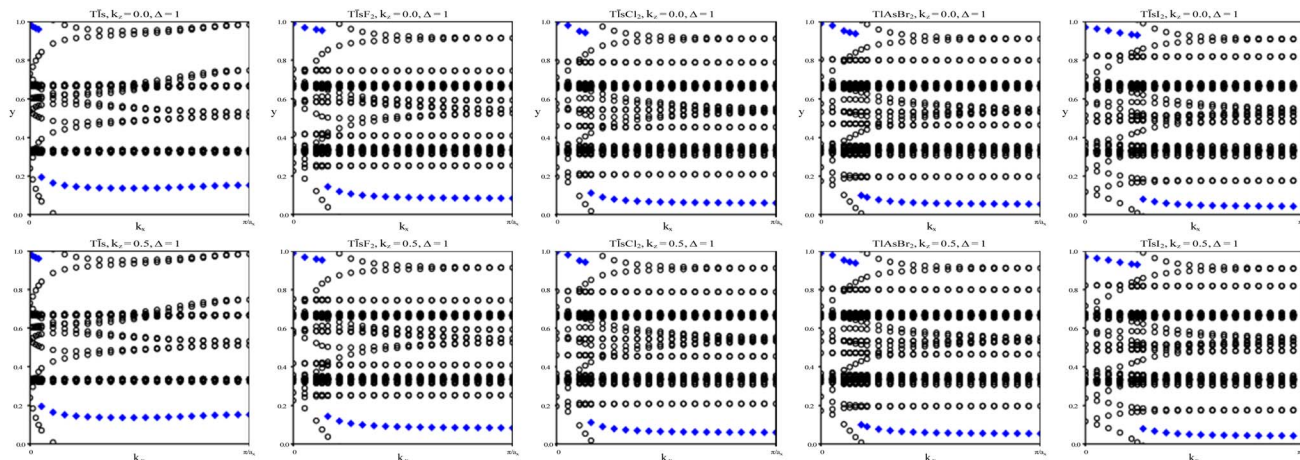


Fig. 6 The trajectories of the Wannier charge centers for TIAs and TIAsX₂ (X = F, Cl, Br, I). The open circles represent hybrid Wannier charge centers. The first row is at the surface $k_z = 0$. The second row is at the surface $k_z = 0.5$. The plots can be identified by their titles.

functionalized, each Tl (As) atom is bonded to a X atom and three As (Tl) atoms. Our results confirm that chemically functionalizing TIAs leads to a consistent increase in each monolayer's buckled height for the monolayers TIAsCl₂, TIAsBr₂, and TIAsI₂. The buckled height increased by 0.08 Bohr, 0.17 Bohr, and 0.12 Bohr, respectively. However, the buckled height decreased by 0.25 Bohr for the monolayer TIAsF₂. This may be due to the difference in electronegativities, as the electronegativity of F is much higher than Cl, Br, or I. When As is functionalized with a more electronegative atom, there can be

a stronger pull on the electrons. Because As is one period below P, it is heavier and has more electron shielding, leading to the changes in geometric structure. The bond length Tl–As increased from 5.06 Bohr for the monolayer TIAs to 5.32 Bohr, 5.31 Bohr, 5.31 Bohr, and 5.34 Bohr for the chemically functionalized monolayers TIAsF₂, TIAsCl₂, TIAsBr₂, and TIAsI₂, respectively. Previous literature has concluded that TIAs has a buckled height of 1.40 Bohr and a lattice constant of 8.54 Bohr.^{5,20}

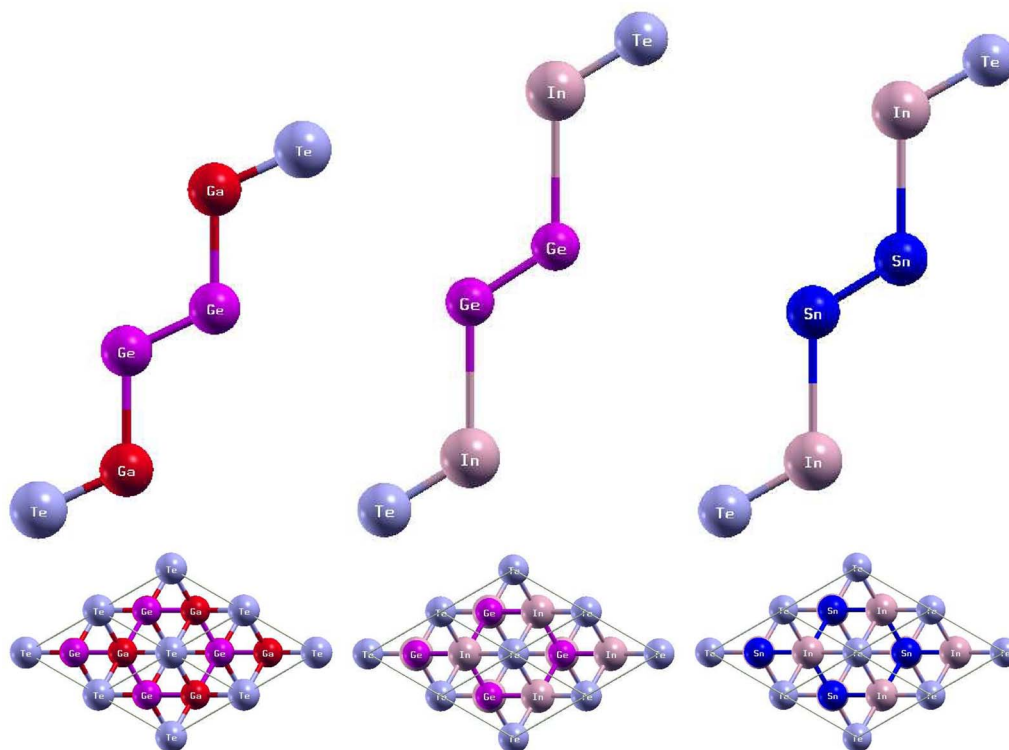


Fig. 7 Optimized atomic structures of the GaGeTe, InGeTe, and InSnTe monolayers are shown from left to right. The atomic structures can be identified by the labels on each atom. Here, magenta, red, violet, and pink atoms represent Ge, Ga, Te, and In atoms, respectively.



Table 4 Calculated optimized structural parameters of GaGeTe, InGeTe, and InSnTe: the lattice constant (a) in Bohr, bond length Te – M ($d_{\text{Te-M}}$) in Bohr where M = Ga, In, bond length

System	a (Bohr)	$d_{\text{Te-M}}$ (Bohr)	$d_{\text{M-N}}$ (Bohr)	$d_{\text{N-N}}$ (Bohr)	E_g (eV)	$E_{g-\text{SOC}}$ (eV)	Z_2
GaGeTe	8.14	4.80	4.72	4.83	0.87 (d)	0.74 (d)	0
InGeTe	8.60	5.39	5.03	4.88	0.08 (id)	0.02 (id)	0
InSnTe	8.47	5.52	5.37	5.31	0.24 (d)	0.07 (d)	1

Next, we investigated the electronic properties of TIAs and the chemically functionalized TIAs monolayers. The band structures with and without an SOC effect are displayed in Fig. 5. Without applying SOC, TIAs has a small direct band gap of 0.01 eV at the Γ point. TIAsF_2 , TIAsCl_2 , and TIAsBr_2 have zero energy band gaps. TIAsI_2 has a zero energy indirect band gap, with the VBM and CBM degenerate at the k -points (0.0402, 0.0402, 0.0000) and (0.0345, 0.0345, 0.0000), respectively. The band gaps are displayed in Table 3.

When SOC is taken into account, TIAs , TIAsF_2 , TIAsCl_2 , TIAsBr_2 , and TIAsI_2 have indirect energy gaps of 0.14 eV, 0.18 eV, 0.15 eV, 0.18 eV, and 0.21 eV, respectively. The band gaps of TIAs , TIAsF_2 , TIAsCl_2 , TIAsBr_2 , and TIAsI_2 most likely change dramatically due to the strong SOC of the TI and As atoms. The SOC-induced energy gaps are much larger than the thermal energy at room temperature (0.026 eV), implying that the band gaps for each monolayer would be feasibly measured at each room temperature.

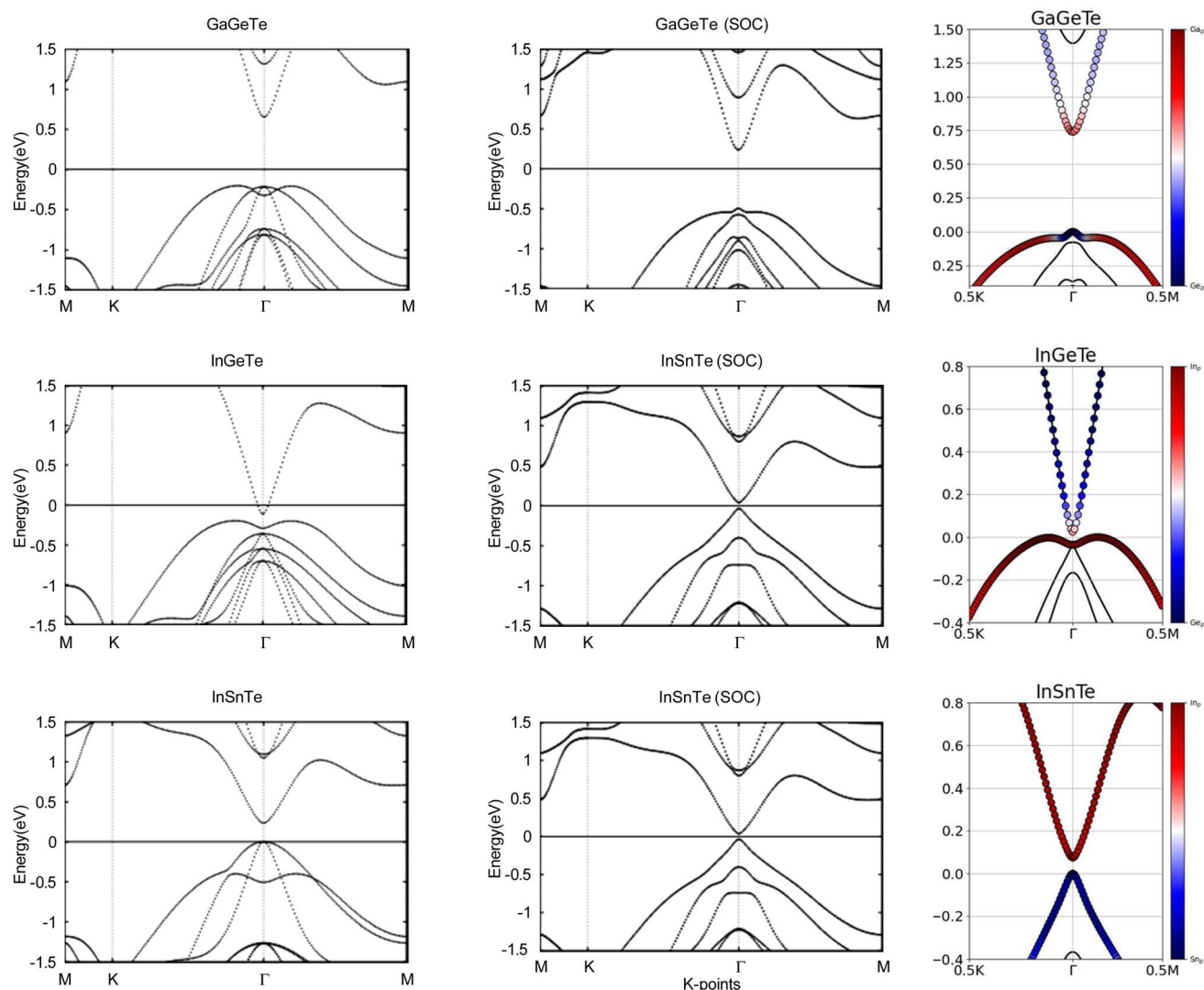


Fig. 8 Calculated band structures with and without SOC of GaGeTe, InGeTe, and InSnTe, with the orbital-resolved band structures with SOC. The Fermi energy is set to 0 eV. The red and blue denote contributions from the 4p and 4p orbitals, 5p and 4p orbitals, and 5p and 5p orbitals, respectively. The band structures can be identified by their titles.



The band structures of TIAs and the TIAsX₂ monolayers are projected on the p orbitals of the elements, similar to the TIP and the TIPX₂ monolayers. These band structures can be found in the third column of Fig. 5. Band inversion is present in some of the chemically functionalized TIAs monolayers.

We calculate the Z_2 topological invariants and list them in Table 3. Fig. 6 shows the trajectories of the Wannier charge centers at the surfaces $k_z = 0$ and $k_z = 0.5$. The Z_2 invariants are calculated identical to how they were calculated for TIP and the chemically functionalized TIP monolayers. The Z_2 calculations confirm that TIAsF₂, TIAsCl₂, TIAsBr₂, and TIAsI₂ are promising 2D TIs with $Z_2 = 1$. Although TIAs has a nonzero band gap of 0.01 eV without SOC, it is less than the thermal energy at room temperature (0.026 eV), and is therefore insignificant and does not affect our results. Because its Z_2 topological invariant is equal to 1, we can strongly conclude that TIAs is a promising 2D TI. Our results for TIAs corroborate Li *et al.*'s calculations with the plane-wave basis Vienna *ab initio* simulation package (VASP).

3.3. GaGeTe, InGeTe, InSnTe

The side and top views of the atomic structures are presented in Fig. 7. The monolayers have buckled honeycomb atomic arrangements.⁴³ When thinned from bulk to monolayer, GaGeTe undergoes a transition from semimetal to semiconductor.²¹ The calculated lattice constants and bond lengths of each monolayer can be found in Table 4. GaGeTe, InGeTe, and InSnTe have calculated lattice constants of 8.14 Bohr, 8.60

Bohr, and 8.47 Bohr, respectively. After substituting In atoms with the Ga atoms of GaGeTe, the bond length Te – In was 0.59 Bohr larger than the bond length Te – Ga. The bond length In – Ge was 0.31 Bohr larger than the bond length Ga – Ge. The bond length Ge – Ge increased by 0.05 Bohr. After substituting Sn atoms with the Ge atoms of InGeTe, the bond length Te – In increased by 0.13 Bohr. The bond length In – Sn was 0.34 Bohr larger than the bond length In – Ge. The bond length Sn – Sn was 0.43 Bohr larger than the bond length Ge – Ge. Previous literature has concluded that GaGeTe has a lattice constant of 7.41 Bohr, in disagreement with the calculated lattice constant of 8.14 Bohr.^{21,44,45}

Next, we turn towards the electronic properties of GaGeTe, InGeTe, and InSnTe. The band structures with and without an SOC effect are displayed in Fig. 8. Without applying SOC, GaGeTe and InSnTe are semiconductors with direct band gaps of 0.87 eV and 0.24 eV at the Γ point, respectively. InGeTe is a semiconductor with a small indirect band gap of 0.08 eV, with the VBM at (0.1200, 0.0600, 0.0000) and the CBM at (0.0000, 0.0000, 0.0000).

The band gaps are displayed in Table 4.

When SOC is taken into account, GaGeTe and InSnTe have direct energy gaps of 0.74 eV and 0.07 eV. With SOC, InGeTe has an indirect energy gap of 0.02 eV, with the VBM at (0.0690, 0.0690, 0.0000) and the CBM at (0.0000, 0.0000, 0.0000). All three of these energy gaps decreased by 0.13 eV, 0.06 eV and 0.17 eV, respectively, from the energy gaps without SOC. This behavior opposes TIPX₂ and TIAsX₂.

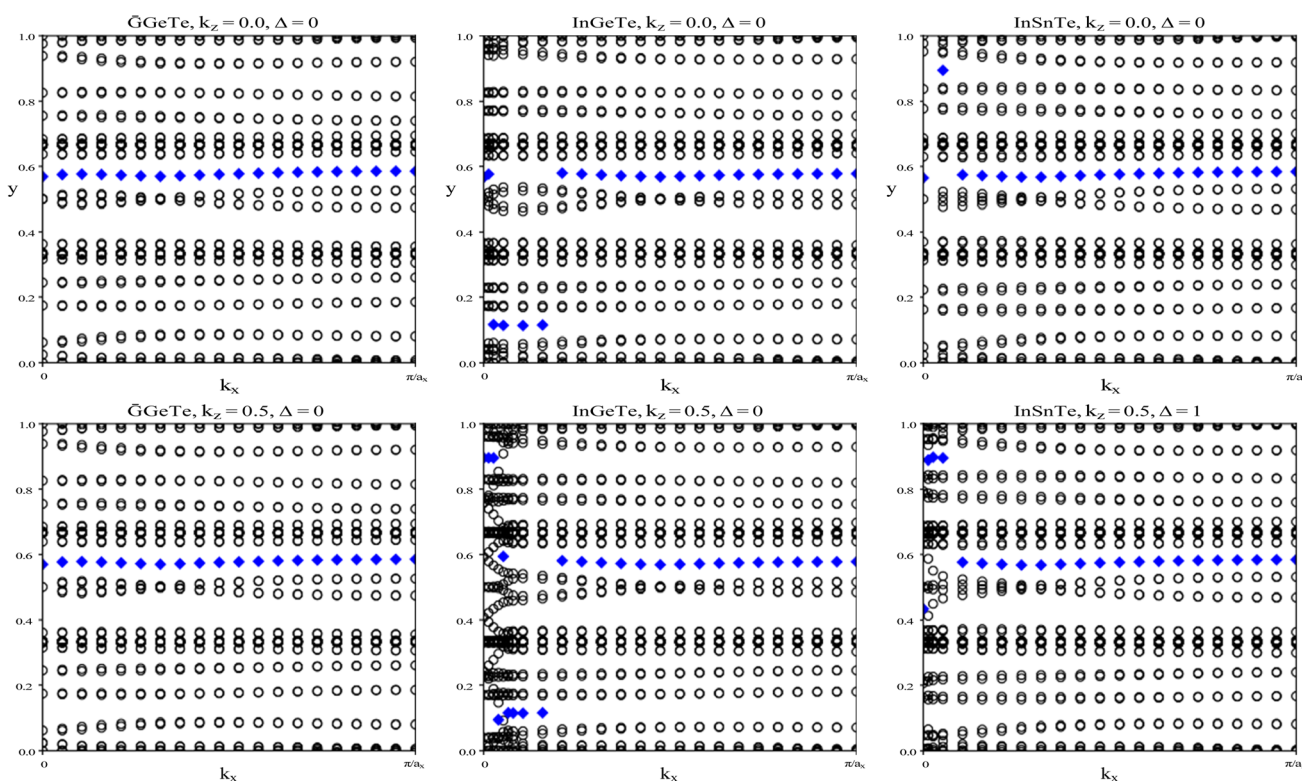


Fig. 9 The trajectories of the Wannier charge centers for GaGeTe, InGeTe, and InSnTe. The open circles represent hybrid Wannier charge centers. The first row is at the surface $k_z = 0$. The second row is at the surface $k_z = 0.5$. The plots can be identified by their titles.



The band structures of the GaGeTe, InGeTe, and InSnTe monolayers are projected on the p orbitals of the elements. These band structures can be found in the third column of Fig. 8. Band inversion is present in some of the monolayers.

Although we are inclined to believe that a QSH state does not exist in each GaGeTe, $M - N$ (d_{M-N}) in Bohr where $N = \text{Ge, Sn}$, and bond length $N - N$ (d_{N-N}) Bohr. Band gaps and topologies of GaGeTe, InGeTe, and InSnTe, including the band gap without SOC E_g in electron-volts and the band gap with SOC $E_{g,\text{SOC}}$ in electron-volts. d and id stand for direct and indirect band gap, respectively. Z_2 is the topological index.

InGeTe, and InSnTe monolayer, we directly calculate the Z_2 topological invariants and list them in Table 4. The trajectories of the Wannier charge centers can be found in Fig. 9 at the surfaces $k_z = 0$ and $k_z = 0.5$. The Z_2 calculations confirm that GaGeTe and InGeTe are normal insulators, with $Z_2 = 0$. Even though bulk GaGeTe has been investigated as a topological semimetal,⁴³ the GaGeTe monolayer does not have topological properties. Although it is true that band inversion can be seen in the GaGeTe and InGeTe monolayers, it does not alone guarantee a non-trivial topological phase.⁴² Because its Z_2 invariant is equal to 0, we can conclude that it is not a topological insulator. The conflicting results may be due to the strength of spin-orbit coupling. It is entirely likely despite band inversion, the parities of the occupied electronic states do not lead to a non-trivial topological insulator. The Z_2 calculation for InSnTe seems promising at first, with $Z_2 = 1$. However, since there are no edge states, we conclude that it is a normal insulator.

4. Conclusion

In conclusion, we have predicted a new family of large-gap 2D TIs using first-principles calculations by investigating the geometric, electric and topological properties of each monolayer. In TlP and TlPX₂ ($X = \text{F, Cl, Br, I}$), TlPF₂, TlPCl₂, TlPBr₂, and TlPI₂ are promising 2D TIs. In TlAs and TlAsX₂ ($X = \text{F, Cl, Br, I}$), TlAs, TlAsF₂, TlAsCl₂, TlAsBr₂, and TlAsI₂ are promising 2D TIs. In GaGeTe, InGeTe, InSnTe, our results show that none of the monolayers were predicted to be promising 2D TIs. Out of these predicted 2D TIs, the band gaps range from 0.03 eV to 0.21 eV. The QSH effect can be detected at room temperature (0.026 eV) for each predicted 2D TI. Our research provides an impressive advance in promising 2D TIs. We firmly believe that these 2D TIs are promising platforms for device application in quantum computers, nanoelectronics, and spintronics.

Conflicts of interest

There are no conflicts to declare.

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