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Conduction band-edge valley splitting in two-dimensional ferroelectric AgBiP_2S_6 by magnetic doping: towards electron valley-polarized transport†

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Two-dimensional valleytronic systems, using the valley index of carriers to perform logic operations, serves as the basis of the next-generation information technologies. For efficient use of the valley degree of freedom, the major challenge currently is to lift the valley degeneracy to achieve valley splitting. In this work, using first-principles calculations, we propose that valley splitting can be readily achieved in a ferroelectric AgBiP_2S_6 monolayer by TM doping (TM = V, Cr, Mn, Fe, Co, and Ni), which is highly feasible in experiments. In sharp contrast to most previous reports of valley-related features in the valence band-edge, the pristine AgBiP_2S_6 monolayer has a direct band-gap located at K/K' points of the Brillouin zone and harbors strong coupled spin and valley physics around the conduction band-edge, due to inversion symmetry breaking combined with strong spin-orbit coupling. By TM-doping, the local magnetic moment can be introduced into the system, which can destroy the valley degeneracy of the conduction band-edge and induce valley splitting. Especially in a V-doped system, accompanied with a large valley splitting (26.8 meV), there is a serious n-type doping in AgBiP_2S_6 . The efficient electron-doping moves the Fermi level just located between the conduction band minimum of the K/K' valleys, which is suitable for valley-polarized transport. Moreover, the valley-polarized index can be flipped by applying a small magnetic field to rotate the magnetocrystalline direction. The magnitude of valley splitting relies on the strength of orbital hybridization between the TM-d and Bi-p states and can be tuned continually by applying biaxial strain. Under an in-plane electric field, such valley degeneracy breaking would give rise to the long-sought anomalous valley Hall effect, which is crucial to design a valleytronic device.

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Introduction

Manipulating different degrees of freedom (DOF) of electrons is at the heart of modern information processing technology. In addition to charge and spin, electrons are endowed with an additional DOF, namely, the valley DOF.^{1–5} The utilization of valley DOF is analogous to charge for electronics and spin for spintronics, leading to the vision of valley-based information processing and storage, which has now expanded into a well-known field named valleytronics.^{3,6–10} In crystalline solids, valley refers to the degenerate energy extreme of a conduction

band or valence band at special k points, have a large separation in the momentum space that enables valley pseudospin to be very robust against phonon and impurity scatterings.^{3,11,12} As the first stable two-dimensional (2D) crystal produced by mechanical exfoliation,¹³ graphene has been demonstrated to have many interesting valleytronic properties. However, in such a nonmagnetic centrosymmetric system, the combination of inversion with time-reversal symmetries ensures Berry curvature and magnetic moments to be zero, thus limiting the constitution of a binary valley index. The valleys can be distinguishable by breaking the inversion symmetry *via* stacking of graphene on hexagonal boron nitride or an application of out-of-plane electric field to bilayer graphene.^{14,15} Different from graphene, monolayer transition-metal dichalcogenides (TMDs; MX_2 , where M = Mo and W, and X = S and Se) with the hexagonal nature inherit broken inversion symmetry because the M and X atoms occupy one of the two inequivalent triangular sublattice. These materials have two valleys at the K and K' , which are related to each other by time-reversal symmetry and no longer being equivalent.¹⁶ The electrons and holes

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† Electronic supplementary information (ESI) available: Total DOS and orbital-resolved PDOS for each atom of AgBiP_2S_6 monolayer calculated by the HSE06 method (Fig. S1); calculated binding energies of TM-doped AgBiP_2S_6 (Fig. S2); spin charge density of TM-doped AgBiP_2S_6 monolayers (Fig. S3), lattice vectors and fractional coordinates of each atom in TM-doped AgBiP_2S_6 (Tables S1–S6). See <https://doi.org/10.1039/d2ra01697d>



residing at the K and K' valleys possess opposite spin, orbital magnetic moment, and Berry curvature.^{11,17} The broken inversion symmetry with strong spin–orbit coupling (SOC) introduce valley-contrasting physical properties in TMDs.¹⁸

In order to distinguish and manipulate the carriers at defined valleys, the K and K' valley degeneracy should be lifted to introduce the valley splitting, and previous studies have provided various strategies. One is to use an ultrafast circularly polarized laser pump to break the valley degeneracy through the optical Stark effect,^{6,19} another is to apply a vertical magnetic field by taking advantage of the Zeeman effect.^{20–22} However, the optical pumping technology may only be used for experimental measurements but not for practical applications because of the restricted lifetime of carriers and the efficiency of external magnetic field is usually too low that a 1 T magnetic field can only give rise to a splitting of 0.1–0.2 meV. Thus, magnetic proximity coupling is proposed and studied intensively, where magnetic substrates, such as EuO ,^{23,24} CoO ,²⁵ MnO ,^{12,26} EuS ,^{27,28} Fe_3O_4 ,²⁹ and YMnO_3 ,³⁰ contribute to the interfacial magnetic exchange. A giant valley splitting in WS_2/EuS heterostructures have been experimentally realized.³¹ But the bulk substrates intrinsically limit the device applications in nanoscale. A 2D magnetic van der Waals (vdW) heterostructure $\text{WSe}_2/\text{CrI}_3$, which is favored for forming a relatively clean interface to eliminate impurity scattering and can minimize the effect of lattice mismatch, has been both theoretically predicted and experimentally realized, showing sizable valley splitting and optical-controllable valley pseudospin.^{32,33} Compared with the construction of 2D magnetic heterostructure, internal magnetic moments induced by magnetic dopants should also be able to engineer the valley splitting and is more feasible.^{34–36} With the innovation of laboratory equipment and methods, transition-metal (TM) doped 2D TMDs have been reported in many experiments, and can avoid the formation of metal clusters.^{37–41} However, for most 2D TMDs, the valence band undergoes larger spin splitting with SOC which varies from 150 meV in MoS_2 to 460 meV in WSe_2 .^{42,43} We expect that the holes in a given valley may retain their spin information even at room temperature. Thus, p-type TMDs are promising candidates for valleytronic applications. Although magnetic doping can break valley degeneracy and induce valley splitting, the Fermi level must be tuned to the valence band maximum (VBM) by subsequent gating or hole doping, and then the longitudinal transport will be both spin and valley-polarized.

In view of the above analysis, materials with larger SOC-induced spin splitting at the conduction bands may have more advantages for magnetic doping and practical application. Our previous reports have demonstrated that the broken inversion symmetry with strong SOC in 2D ferroelectric AgBiP_2S_6 will introduce large spin splitting at the conduction band-edge.⁴⁴ Here, we performed density functional theory (DFT) and Berry curvature calculations to explore the valley splitting related properties of AgBiP_2S_6 induced by the TM doping (TM = V, Cr, Mn, Fe, Co, and Ni). In previous report, B-site (In) substitution effect in ferroelectric CuInP_2S_6 monolayer have been investigated.⁴⁵ Ferromagnetism and ferroelectricity can coexist in both CuCrP_2S_6 and CuVP_2S_6 . However, complete

substitution may induce large structural distortion or even phase transformation. Thus, A-site (Ag) substitution doping in ferroelectric AgBiP_2S_6 can destroy the time-reversal symmetry of the system by maintaining the valley behavior related to the heavy element Bi at the conduction bands, which is beneficial to the subsequent investigation of valley splitting. We find that pristine AgBiP_2S_6 monolayer has a direct band-gap located at K/K' points of the Brillouin zone and harbors coupled spin and valley physics around the conduction band-edge. By TM-doping, the local magnetic moment can be introduced into the system, which can destroy the valley degeneracy of the conduction band-edge and induce valley splitting. Especially in V-doped system, accompany with a large valley splitting (26.8 meV), there is a serious n-type doping in AgBiP_2S_6 . The efficient electron-doping moves the Fermi level just located between the conduction band minimum (CBM) of the K/K' valleys, which is suitable for valley-polarized transport. Moreover, the valley-polarized index can be flipped by applying a small magnetic field to rotate the magnetocrystalline direction. The magnitude of valley splitting relies on the strength of orbital hybridization between the TM-d and Bi-p states and can be tuned continually by applying biaxial strain. Under an in-plane electric field, such valley degeneracy breaking would give rise to the long-sought anomalous valley Hall effect, which is crucial to design a valleytronic device.

Computational details and models

Density functional theory (DFT) calculations for structural relaxation and electronic structures are performed by using the Perdew–Burke–Ernzerhof generalized gradient approximation (PBE-GGA) for the exchange and correlation function as implemented in the Vienna *ab initio* simulation package (VASP).^{46–48} The projector augmented wave (PAW) is used to describe the electron–ion interaction.^{49,50} The cutoff energy is set to 500 eV for plane-wave expansion after convergence tests. A $2 \times 2 \times 2$ AgBiP_2S_6 supercell with one substituted dopant atom is used. The T -centered k -grids of $2 \times 2 \times 1$ and $4 \times 4 \times 1$ are adopted to sample the Brillouin zone for geometry optimization and static electronic calculation, respectively. During the structural relaxation, the energy convergence criterion is 10^{-5} eV per unit cell, and the forces on all relaxed atoms are less than $0.01 \text{ eV } \text{Å}^{-1}$. The vacuum space larger than 20 Å is adopted to eliminate the interaction between adjacent layers. The PBE + U method with $U_{\text{eff}} = 3.0 \text{ eV}$ for TM atoms is used to take strong correlation effects into account,⁵¹ which has been widely used in previous works.^{35,52} The SOC effect explicitly included in the band structure calculations. For the calculations of Berry curvature and anomalous Hall conductivity, the maximally localized Wannier functions (MLWFs) implemented in the WANNIER90 package are employed.⁵³

Results and discussion

TM thio/selenophosphates is a broad class of vdW layered structures with two sulfur or selenium layers sandwiching a layer of metal ions and P_2 pairs and general compositions of

$M_4[P_2X_6]^{4-}$, $[M^{2+}]_2[P_2X_6]^{4-}$, and $M^{1+}M^{3+}[P_2X_6]^{4-}$, where $M^{1+} = \text{Cu, Ag}$; $M^{3+} = \text{Cr, V, Al, Ga, Bi}$; and $X = \text{S, Se}$.^{54,55} Especially, those quaternary compounds exhibit mixed electron-ionic conductivity, promising optical and catalytic properties.⁵⁶ As shown in Fig. 1(a), AgBiP_2S_6 is a typical quaternary compounds with a rhombohedral structure and contains a sulfur framework with the octahedral voids filled by Ag, Bi, and P-P triangular patterns. The $[P_2S_6]^{4-}$ anions are bridged by alternating and ordered arrangement of Ag and Bi metal centers. The inset of Fig. 1(b) shows the side view of the crystal structure. Both Ag^+ and Bi^{3+} ions are distorted from the octahedral environment, constituting a collinear two-sublattice system. They shift to opposite directions and the Ag^+ ion off-centering is much greater than that of the Bi^{3+} ions. Consequently, spontaneous polarization emerges with the polar axis normal to the layer plane.⁵⁷ The non-centrosymmetric unit cell illustrated by the dashed line belongs to the space group $P3$. The optimized lattice constant of AgBiP_2S_6 monolayer is 6.45 Å, while the thickness d is 3.44 Å. The Ag^+ and Bi^{3+} ions shifting opposite away from the center of the layer along the z axis by 0.49 Å and 0.17 Å, respectively. These structure parameters are in good agreement with the previous work.⁵⁵ The average electric potential energy of AgBiP_2S_6 is shown in Fig. 1(b), suggesting a potential gradient arising from the broken mirror symmetry in the vertical direction, and the net electric field pointing from Bi to Ag. According to the Helmholtz equation, the energy difference between the

two vacuum levels is actually the change of work function ($\Delta\phi = 0.11$ eV), which is proportional to the dipole moment.⁵⁸ The phonon dispersion relations of AgBiP_2S_6 monolayer display no imaginary frequency (see Fig. 1(c)), suggesting that it is dynamically stable.^{59,60} The band structure of AgBiP_2S_6 without considering SOC is shown in Fig. 1(d). It can be seen that AgBiP_2S_6 monolayer is a nonmagnetic semiconductor with a direct band-gap of 2.03 eV, wherein the band edges located at the same high symmetry K point. In a hexagonal system with broken inversion symmetry, the valley DOF will be present when a local minimum in the conduction band or local maximum in the valence band appears at the K/K' points.^{16,43} Apart from the VBM and CBM, there are also some valleys appearing in the other conduction and valence bands, and we do not consider them because they lie too far away from the Fermi level to be useful. Fig. 1(e) shows the total density of states (DOS) and the orbital-resolved partial density of states (PDOS) for each atom of AgBiP_2S_6 . The bands near the VBM are mainly contributed by Ag- d and S- p orbitals, while the bands near the CBM arise from Bi- p and S- p orbitals. These phenomena can also be obtained intuitively from the related band decomposed charge density of the VBM and CBM in Fig. 1(f). Since PBE is known to underestimate the band-gap, the hybrid Heyd-Scuseria-Ernzerhof (HSE06) method is used to predict a more reasonable energy band-gap.⁶¹ Fig. S1† shows the total DOS and orbital-resolved PDOS calculated by the HSE06 method. It is worth

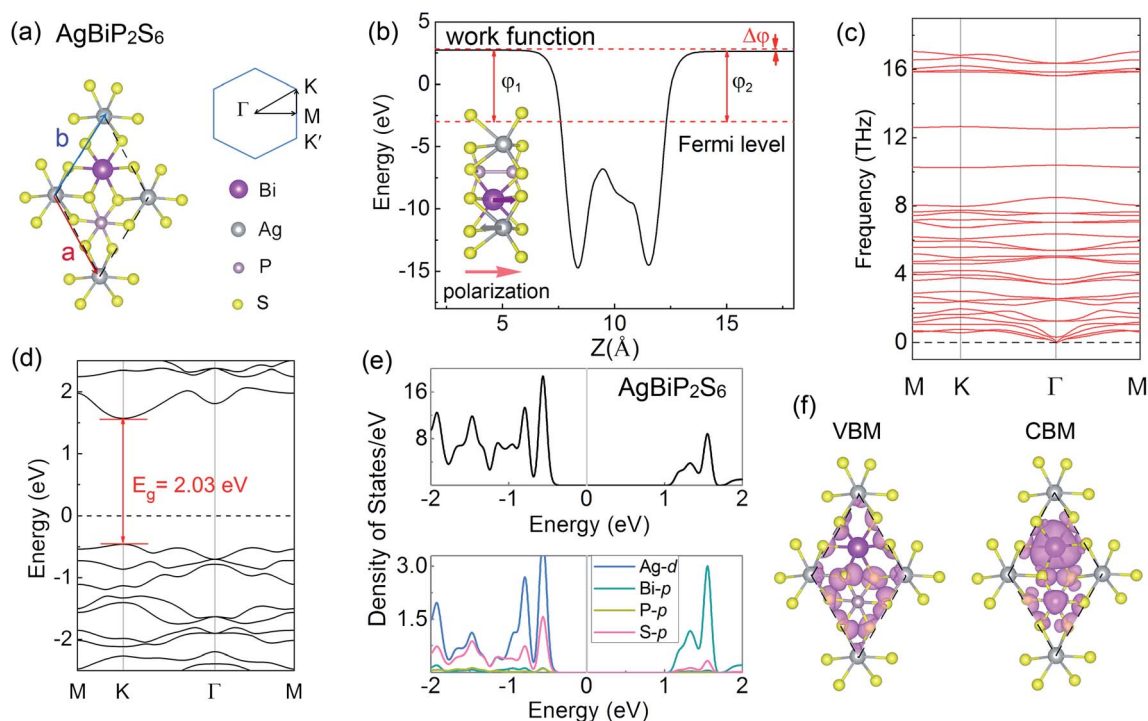


Fig. 1 (a) The top view of the crystal structure of the AgBiP_2S_6 monolayer and hexagonal 2D Brillouin zone labeled with high-symmetry points. The dashed line is the unit cell. (b) The average electronic potential energy in the vertical direction of the AgBiP_2S_6 monolayer. Inset is the side view of AgBiP_2S_6 . The purple and grey arrows represent the polarized contributions from the Bi^{3+} and Ag^+ ions, respectively. The pink arrow represents the total polarization of the AgBiP_2S_6 monolayer. (c) Calculated phonon band dispersion. (d) Band structure of the AgBiP_2S_6 monolayer without SOC and the Fermi level is set to zero. (e) The total DOS and orbital-resolved PDOS for each atom of AgBiP_2S_6 monolayer. (f) The related band decomposed charge density of the VBM and CBM.

mentioning that the HSE06 functional only enhances the band-gap to 2.45 eV, but have no significant changes in the orbital contributions of the VBM and CBM.

Upon considering SOC, as shown in Fig. 2(a), the spin degeneracy for the valleys of AgBiP₂S₆ monolayer is deformed because of the broken inversion symmetry. And the time-reversal symmetry leads to the opposite ordering of the spin-up and spin-down states at the *K* and *K'* valleys. The SOC-induced spin splitting is 423 meV in conduction band (Δ_c), whereas the splitting of spin-up and spin-down states near valence band (Δ_v) is found to be very small (~ 30 meV), which is exactly opposite to that of TMDs,⁴³ where large SOC-induced spin splitting is observed at valence band. This phenomenon is attributed to the fact that the valence and conduction band-edges are dominated by different orbitals. The conduction band-edges are mainly contributed by heavy Bi atoms as plotted in the orbital-resolved PDOS and band decomposed charge density, thus the strong SOC of Bi-p orbitals will lift the out-of-plane spin degeneracy of the band-edges at *K* and *K'* by a significant amount. Because of the time-reversal symmetry, the *K* and *K'* valleys of AgBiP₂S₆ exhibit opposite spin signs but are energetically degenerate. To make use of the valley index as an information carrier, the crucial step is to lift this degeneracy,

thereby producing valley splitting. Hence, 3d TM (V, Cr, Mn, Fe, Co, and Ni) with large atomic magnetic moments are selected as the magnetic dopants to break the time-reversal symmetry of AgBiP₂S₆. For retaining the Bi-p orbital-related large spin splitting at the conduction band-edge, substitution doping of Ag atom is performed in a 2×2 supercell. The TM atom replaces one of the four Ag atoms in the supercell with a doping concentration of 25%. It should be mentioned that the electronic structures of the 2×2 supercell at the *K/K'* points are the same as those of the unit cell. Thus, we do not need to consider the band unfolding technique here.^{35,52} The binding energy (E_b) is given by the expression

$$E_b = E_{\text{doped}} - (E_{\text{vacancy}} + E_{\text{TM}}) \quad (1)$$

where E_{doped} , E_{vacancy} , and E_{TM} represent the energies of TM-doped AgBiP₂S₆, AgBiP₂S₆ supercell with one Ag vacancy, and TM dopants, respectively. The calculated E_b for V-, Cr-, Mn-, Fe-, Co-, and Ni-doped systems are -2.64 , -3.12 , -2.74 , -0.97 , -0.92 , and -0.40 eV, respectively (see Fig. S2[†]), which are comparable to those of V/Cr-doped MoSSe.³⁵ These negative values are related to the exothermic formation process, indicating strong thermal stability and experimental feasibility of

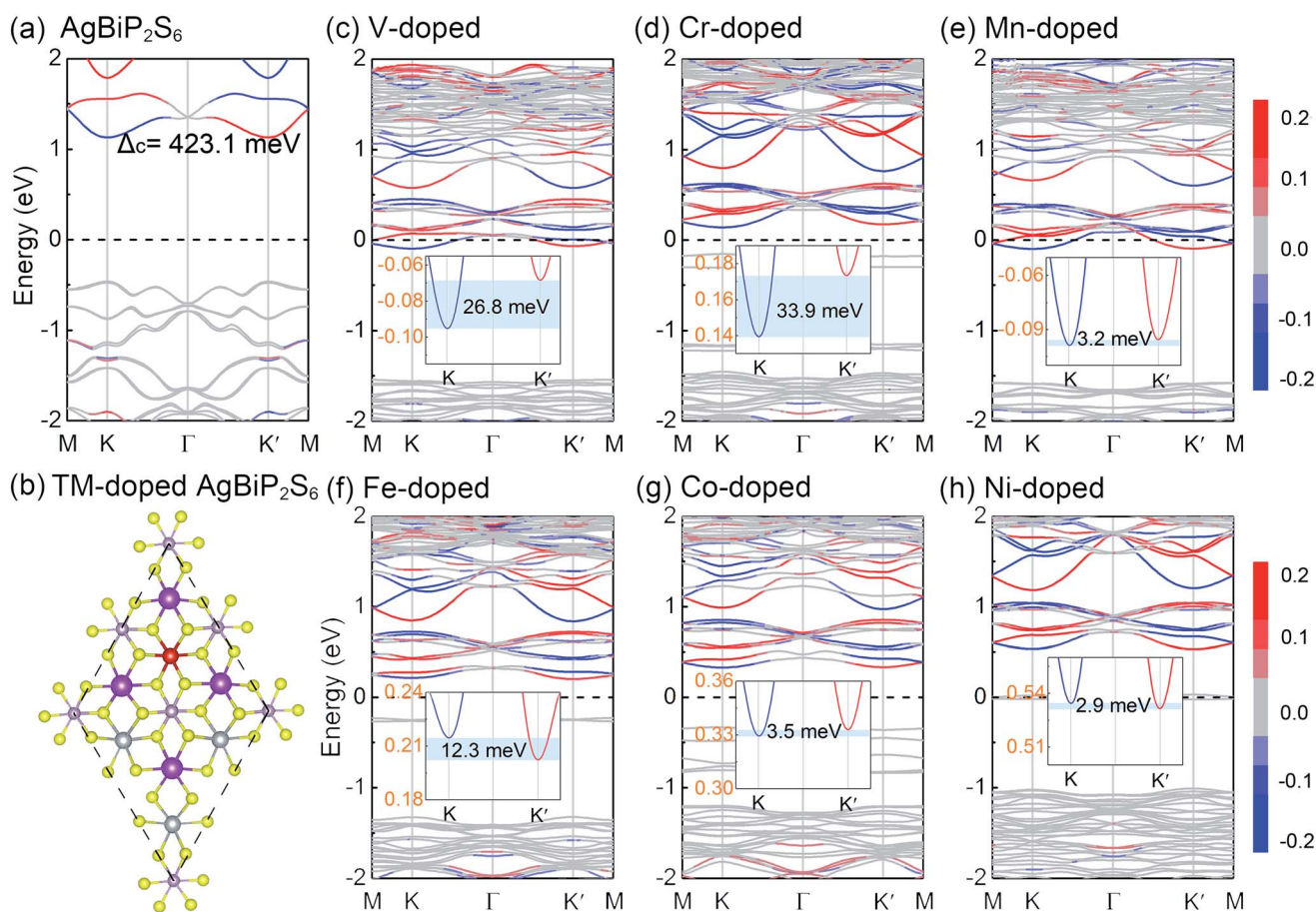


Fig. 2 (a) Band structure for pristine AgBiP₂S₆ monolayer with SOC. (b) The top view of the structure of TM-doped AgBiP₂S₆ monolayer. The red ball denotes the TM atoms. (c)–(h) Band structures of TM-doped AgBiP₂S₆ calculated by the PBE method with SOC. The spin projections along *z* direction are represented by red and blue lines, which represent opposite spin states, respectively.

the doped systems. The spin charge density of the TM-doped AgBiP₂S₆ monolayers is calculated and shown in Fig. S3.† In the pristine system, the spin density is distributed symmetrically, leading to no spin polarization. Our calculations show that all TM-doped systems are spin-polarized, and the obtained local magnetic moments are approximately 3.27, 5.00, 4.87, 3.00, 2.00, and 1.00 μ_B as Ag-d¹⁰s¹ substitutional doping by V-d³s², Cr-d³s¹, Mn-d⁵s², Fe-d⁶s², Co-d⁷s², and Ni-d⁸s², respectively. This can be attributed to the difference of valence electrons and the crystal field distortion induced by doping elements with small ionic radius. The spin density mainly distributes around the dopants, and a little with an opposite sign is distributed on the adjacent S atoms, reflecting weak antiferromagnetic coupling and strong interaction between dopant and AgBiP₂S₆.

The band structures of all TM-doped systems with considering SOC are shown in Fig. 2(c)–(h).

We can see that all of the TM-doped AgBiP₂S₆ monolayers maintain the semiconducting nature with different degree of n-type doping, which mainly depends on the valence states of dopants. Moreover, some impurity states appear in the band-gap, but fortunately, the *K* and *K'* valleys at the conduction band-edge are preserved well. Especially for the cases of V, Mn, and Ni-doped AgBiP₂S₆, the valleys in the conduction bands lie far away from the defect states, making it easy to realize further manipulations and practical applications. Upon considering SOC, the degeneracy of the CBM at *K* and *K'* valleys is lifted for all TM-doped systems, successfully achieving the valley splitting. Herein, valley splitting is quantified by the energy difference between the *K* and *K'* valleys of the CBM as $\Delta E_{KK'}$ = $|E_K - E_{K'}|$. In V and Cr-doped AgBiP₂S₆, the valley splitting in the conduction bands can reach 26.8 and 33.9 meV, respectively, which are comparable to those of the previously reported systems, like V/Cr-doped MoSSe (10–59 meV),³⁵ V/Mn-doped CrS₂ (40–65 meV),⁶² and N/C-doped Tl₂O (23–44 meV).⁶³ Considering that the external magnetic field can also induce valley splitting with a value of 0.1–0.2 meV/T,⁶⁴ the obtained $\Delta E_{KK'}$ of V and Cr-doped AgBiP₂S₆ even corresponds to an equivalent magnetic field of about 150–350 T. Such a large external magnetic field, of course, is difficult to be achieved in experiments, strongly suggesting that magnetic doping is a very promising method for realizing the valley splitting. From the PDOS of the doped systems shown in Fig. 3, we can see that V, Mn, and Ni-doped AgBiP₂S₆, have a clean band-gap, the TM-d orbitals only hybridize with Ag-d and S-p orbitals near the VBM, but in Cr, Fe, and Co-doped systems, the TM-d orbitals will also form impurity state in the band-gap. In any case, the Bi-p orbitals contribution at the CBM is not affected, indicating that the TM-doping destroys the time-reversal symmetry of the system by introducing local magnetic moments, providing an ideal platform for studying the conduction band-edge valley splitting. More surprisingly, with V-doping, the high-valence substitution also moves the Fermi level through the valleys at the CBM by n-type doping, which is conducive to the device application of electron valley-polarized transport. We further discuss the effect of doping concentration on valley splitting. In Fig. S4(a) and (b),† when Ag atoms in the AgBiP₂S₆ are

completely replaced with a doping concentration of 100%, VBIP₂S₆ monolayer is a semiconductor with an indirect band-gap of about 0.12 eV. The electronic states at the CBM and VBM are mainly contributed from the d orbitals of V and p orbitals of Bi, respectively. The carriers at the CBM are spin polarized and the degeneracy of the CBM at *K* and *K'* valleys is lifted with an energy difference of about 171 meV due to the SOC effect. This intrinsic valley splitting in such a magnetic semiconductor is similar to the previous reports, such as Nb₃I₈,⁶⁵ NbSe₂,⁶⁶ and LaBr₂.⁶⁷ In Fig. S4(c) and (d),† with a lower doping concentration of 11% in a 3 × 3 AgBiP₂S₆ supercell, the valley polarized states at *K* and *K'* points in the unit cell are folded to Γ point. It should be noted that, the coordinate overlap of the *K* and *K'* points in a 3 × 3 supercell does not affect the electronic properties at the two valleys. The two nondegenerate and inequivalent valleys still exist with an energy difference of about 7.2 meV. Also, the electronic bands at the two non-equivalent valleys are mainly contributed by V-d and Bi-p orbitals. These results indicate that the value of valley splitting in AgBiP₂S₆ is proportional to the doping concentration of magnetic elements.

It is known that in systems with broken inversion symmetry, the charge carriers in the *K* and *K'* valleys will acquire a nonzero Berry curvature Ω along the out-of-plane direction. The Berry curvature induced intrinsic transverse velocity shows the physical nature of the anomalous Hall effect. The Berry curvature can be written as⁶⁸

$$\Omega(k) = \pm \frac{2\tau a^2 t^2 (\Delta - \tau s_z \lambda)}{[(\Delta - \tau s_z \lambda)^2 + 4a^2 t^2 k^2]^{3/2}}, \quad (2)$$

where the valley order τ has opposite signs for the two valleys, and the + and – signs represent the valence and conduction bands, respectively. The integral Berry curvature over the occupied states gives the contribution to the anomalous Hall conductivity (AHC)

$$\sigma_H = -\frac{e^2}{h} \int \frac{d^2k}{(2\pi)^2} \Omega(k). \quad (3)$$

In the absence of a magnetic field, the net effect is a pure valley Hall current with a vanishing charge Hall current. However, an external magnetic field will cause a net charge current because the Hall currents from the two valleys do not completely cancel. Thus, the internal magnetic field induced by introducing magnetic dopants in AgBiP₂S₆ is expected to cause AHC owing to the broken time-reversal symmetry. In Fig. 4(a), the calculated Berry curvatures for the *K* and *K'* valleys have the same absolute value but opposite signs, which reveals that the valley-contrasting characteristic remains in AgBiP₂S₆ even strongly hybridized with magnetic dopants. Under an in-plane electric field *E*, Bloch electrons in this system will acquire an anomalous velocity $v \sim E \times \Omega(k)$. As signs of Berry curvature are opposite, the electrons in the *K* and *K'* valleys will move to opposite edges of V-doped AgBiP₂S₆. Since the Fermi level is just located between the *K* and *K'* valleys in the conduction band-edges, as shown from the inset of Fig. 4(c), the spin-down

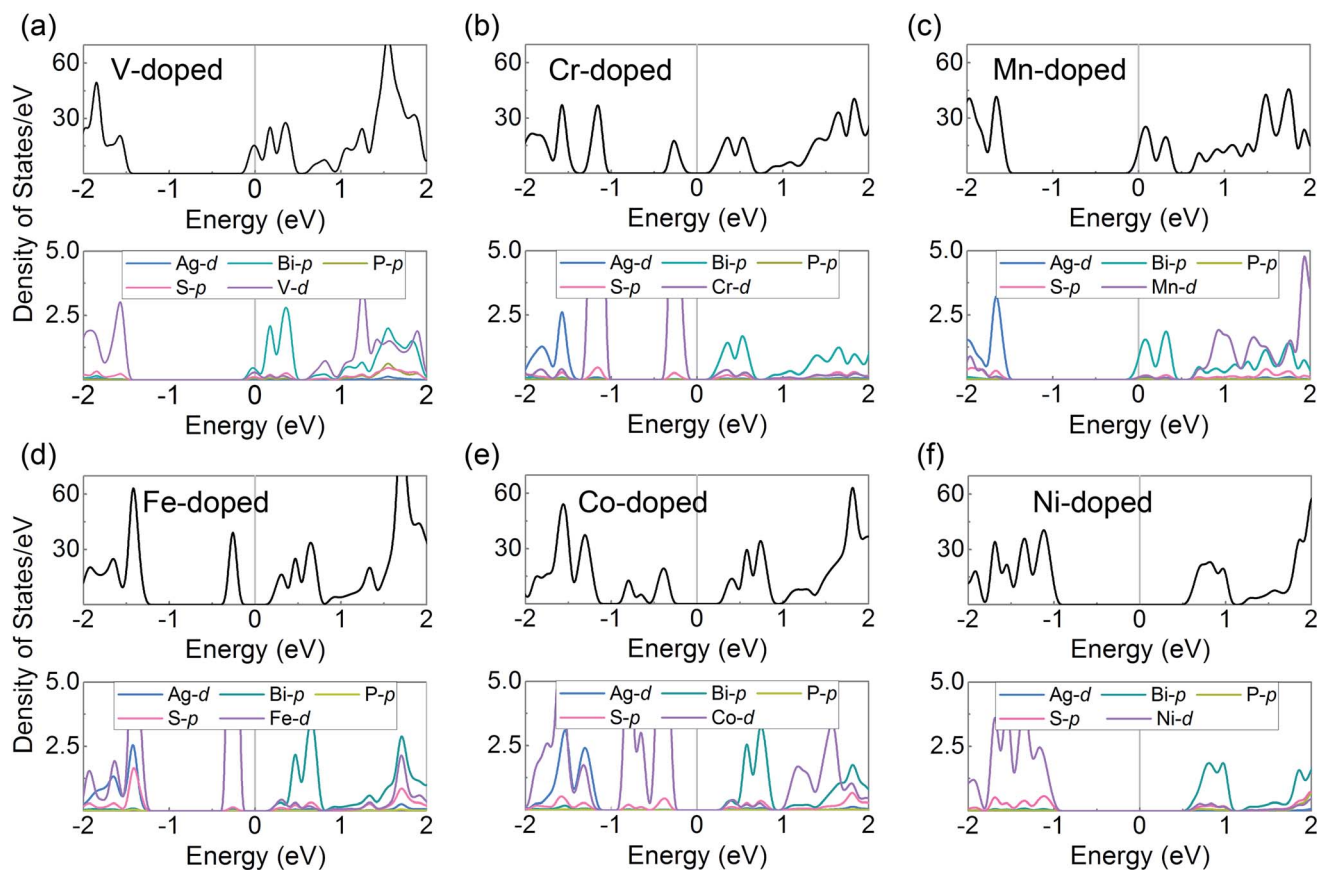


Fig. 3 (a)–(f) The total DOS and orbital-resolved PDOS for each atom of TM-doped AgBiP_2S_6 monolayers by the PBE method with SOC. The Fermi level is set to zero.

electrons from the K valley are accumulated at the downside of the sample under an in-plane electric field. When such a system is magnetized in the opposite direction, the spin-up electrons in the K' valley will move to the upside of the sample in the presence of an in-plane electric field, realizing the anomalous valley Hall effect. As electrons are only accumulated at one edge of the sample, a more easily measurable charge Hall current as a voltage can be detected, offering a possible avenue to realize data storage using valley splitting. The AHC is also calculated

and shown in Fig. 4(b). The valley splitting of the AHC is prominent because it changes from a small finite value to zero when the Fermi level shifts from the CBM_K to the $\text{CBM}_{K'}$. When the Fermi level lies between the CBM_K to the $\text{CBM}_{K'}$ as denoted by the shaded region, a fully spin- and valley-polarized Hall conductivity will be generated. This provides us with a method to detect the valley pseudospin by electrical measurements and forms the basis for the application of the valleytronic device.

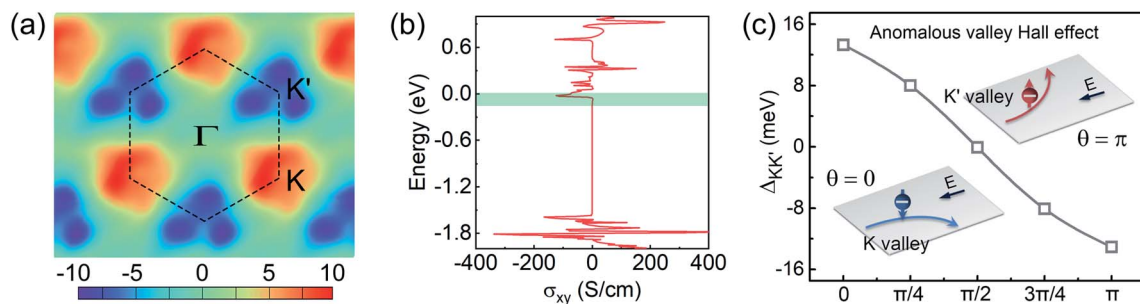


Fig. 4 (a) Contour map of Berry curvature of V-doped AgBiP_2S_6 over the entire 2D Brillouin zone. (b) AHC with the Fermi level in the same energy ranges as the band structure. The shaded region denotes the energy levels of the two valleys in the conduction band-edge. (c) Relation between valley splitting $\Delta E_{KK'}$ and magnetocrystalline angle θ . Insets are the illustrations of anomalous valley Hall effect in V-doped AgBiP_2S_6 with the magnetization direction $\theta = 0$ and $\theta = \pi$, respectively.

Finally, we focus on engineering the valley splitting in V-doped AgBiP_2S_6 . The external magnetic field, which can modulate the spin orientation, is expected to be an effective approach to control the valley splitting. As shown in Fig. 4(c), valley splitting is strongly correlated with the magnetization direction. When the spin orientation is along the in-plane direction ($\theta = \pi/2$), the valley splitting of the doped system vanishes. When the spin orientation is reversed ($\theta = \pi$), the valley splitting is also reversed. Furthermore, since the magnetic dopant induced valley splitting is mediated with the orbital hybridization, we expect that its magnitude can be modulated by changing the hybridization strength. In experiment, one can apply a specific substrate supporting the system to achieve the strain, and then the lattice constant, atomic distance, and orbital hybridization can be adjusted. The biaxial strain is defined as $\varepsilon = (a - a_0)/a_0$, where a_0 and a are the lattice constants of the unstrained and strained systems. The related band structures under various biaxial strain are shown in Fig. 5(a)–(g). Fig. 5(h) shows the variation of the valley splitting as a function of the biaxial strain ε (%). We can see that compressed strain will inhibit the spin splitting in both K and K' valleys, decreasing the valley splitting. When tensile strain is applied, the SOC-induced spin splitting in K valley is enhanced, but suppressed in K' valley. Particularly, when $\varepsilon = +2\%$ (see Fig. 5(b)), valley splitting increases to 63 meV, and its morphology is similar to the recent reported “single-valley state” in 2D antiferromagnetic BiFeO_3 .⁶⁹ With further

increasing the tensile strain, this phenomenon gradually disappears, suggesting that the V-doped AgBiP_2S_6 under appropriate tensile strain is an ideal valleytronic material.

Conclusions

In conclusion, using first-principles calculations, we demonstrate that valley splitting can be readily achieved in TM-doped 2D ferroelectric AgBiP_2S_6 . In sharp contrast to most previous reports of valley-related features in the valence band-edge, pristine AgBiP_2S_6 monolayer has a direct band-gap located at K/K' points of the Brillouin zone and harbors strong coupled spin and valley physics around the conduction band-edge, due to inversion symmetry breaking combined with strong spin-orbit coupling. By TM-doping, the local magnetic moment can be introduced into the system, which can destroy the valley degeneration of the conduction band-edge and induce valley splitting. Especially in V-doped system, accompany with a large valley splitting (26.8 meV), the high-valence substitution also moves the Fermi level through the valleys at the CBM by n-type doping, which is conducive to the device application of electron valley-polarized transport. Moreover, the valley-polarized index can be flipped by applying a small magnetic field to rotate the magnetocrystalline direction. The magnitude of valley splitting relies on the strength of orbital hybridization between the TM-d and Bi-p states and can be tuned continually by applying biaxial strain. Under an in-plane electric field, such valley

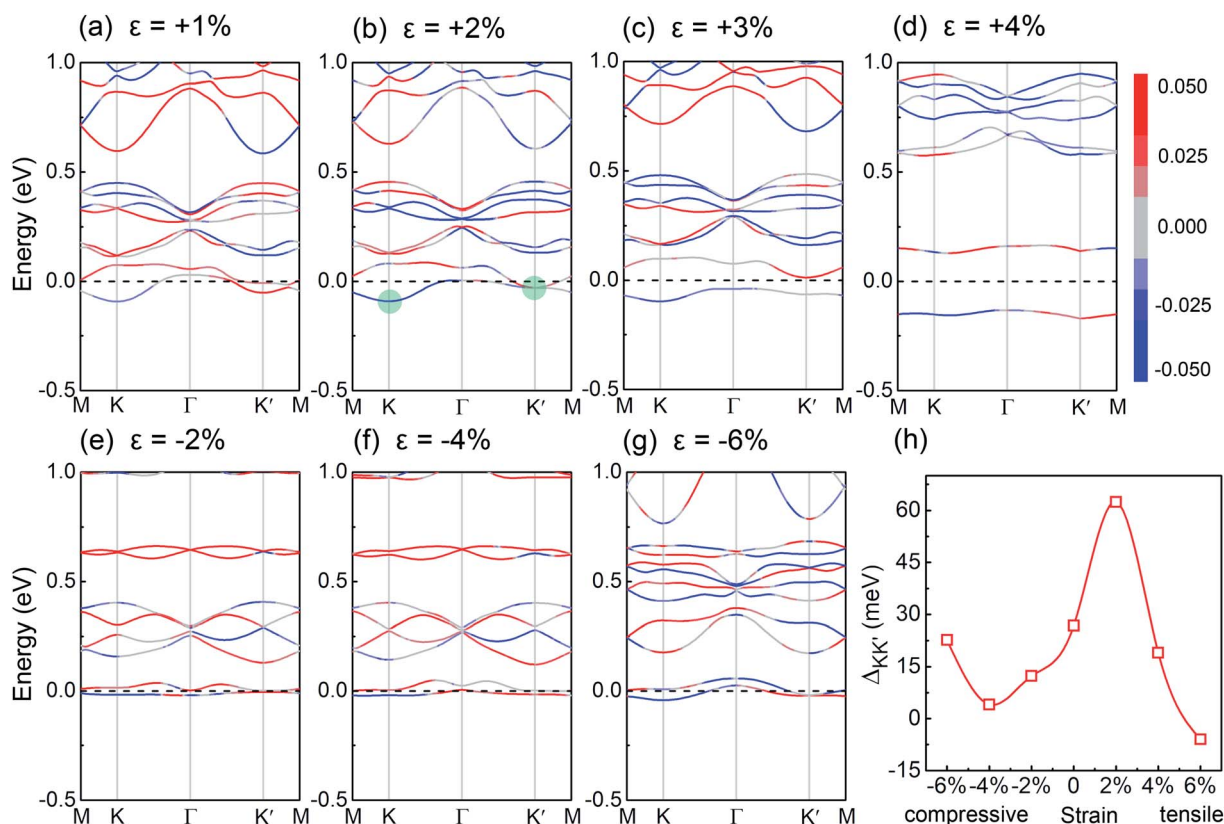


Fig. 5 (a)–(g) Calculated band structures with various biaxial strain. (h) Valley splitting as a function of the biaxial strain ε (%).

degeneracy breaking would give rise to the long-sought anomalous valley Hall effect for both spin/valley filter, which is crucial to design a valleytronic device.

Conflicts of interest

The authors declare no competing financial interest.

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References

- 1 S. A. Wolf, D. D. Awschalom, R. A. Buhrman, J. M. Daughton, S. von Molnar, M. L. Roukes, A. Y. Chtchelkanova and D. M. Treger, *Science*, 2001, **294**, 1488–1495.
- 2 J. R. Schaibley, H. Yu, G. Clark, P. Rivera, J. S. Ross, K. L. Seyler, W. Yao and X. Xu, *Nat. Rev. Mater.*, 2016, **1**, 16055.
- 3 A. Rycerz, J. Tworzydło and C. W. J. Beenakker, *Nat. Phys.*, 2007, **3**, 172–175.
- 4 M. Ezawa, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2013, **88**, 161406.
- 5 Z. Zhu, A. Collaudin, B. Fauqué, W. Kang and K. Behnia, *Nat. Phys.*, 2011, **8**, 89–94.
- 6 T. Cao, G. Wang, W. Han, H. Ye, C. Zhu, J. Shi, Q. Niu, P. Tan, E. Wang, B. Liu and J. Feng, *Nat. Commun.*, 2012, **3**, 887.
- 7 H. Yu and W. Yao, *Nat. Mater.*, 2017, **16**, 876–877.
- 8 D. Xiao, G.-B. Liu, W. Feng, X. Xu and W. Yao, *Phys. Rev. Lett.*, 2012, **108**, 196802.
- 9 A. Castellanos-Gomez, M. Poot, G. A. Steele, H. S. van der Zant, N. Agrait and G. Rubio-Bollinger, *Adv. Mater.*, 2012, **24**, 772–775.
- 10 H. Fang, S. Chuang, T. C. Chang, K. Takei, T. Takahashi and A. Javey, *Nano Lett.*, 2012, **12**, 3788–3792.
- 11 X. Xu, W. Yao, D. Xiao and T. F. Heinz, *Nat. Phys.*, 2014, **10**, 343–350.
- 12 L. Xu, M. Yang, L. Shen, J. Zhou, T. Zhu and Y. P. Feng, *Phys. Rev. B*, 2018, **97**, 041405.
- 13 K. S. Novoselov, A. K. Geim, S. V. Morozov, D. Jiang, Y. Zhang, S. V. Dubonos, I. V. Grigorieva and A. A. Firsov, *Science*, 2004, **306**, 666–669.
- 14 R. V. Gorbachev, J. C. Song, G. L. Yu, A. V. Kretinin, F. Withers, Y. Cao, A. Mishchenko, I. V. Grigorieva, K. S. Novoselov, L. S. Levitov and A. K. Geim, *Science*, 2014, **346**, 448–451.
- 15 M. Sui, G. Chen, L. Ma, W.-Y. Shan, D. Tian, K. Watanabe, T. Taniguchi, X. Jin, W. Yao, D. Xiao and Y. Zhang, *Nat. Phys.*, 2015, **11**, 1027–1031.
- 16 D. Xiao, W. Yao and Q. Niu, *Phys. Rev. Lett.*, 2007, **99**, 236809.
- 17 K. F. Mak, K. L. McGill, J. Park and P. L. McEuen, *Science*, 2014, **344**, 1489–1492.
- 18 Y. Ye, J. Xiao, H. Wang, Z. Ye, H. Zhu, M. Zhao, Y. Wang, J. Zhao, X. Yin and X. Zhang, *Nat. Nanotechnol.*, 2016, **11**, 598–602.
- 19 H. Zeng, J. Dai, W. Yao, D. Xiao and X. Cui, *Nat. Nanotechnol.*, 2012, **7**, 490–493.
- 20 G. Aivazian, Z. Gong, A. M. Jones, R.-L. Chu, J. Yan, D. G. Mandrus, C. Zhang, D. Cobden, W. Yao and X. Xu, *Nat. Phys.*, 2015, **11**, 148–152.
- 21 D. MacNeill, C. Heikes, K. F. Mak, Z. Anderson, A. Kormanyos, V. Zolyomi, J. Park and D. C. Ralph, *Phys. Rev. Lett.*, 2015, **114**, 037401.
- 22 Y. Li, J. Ludwig, T. Low, A. Chernikov, X. Cui, G. Arefe, Y. D. Kim, A. M. van der Zande, A. Rigosi, H. M. Hill, S. H. Kim, J. Hone, Z. Li, D. Smirnov and T. F. Heinz, *Phys. Rev. Lett.*, 2014, **113**, 266804.
- 23 J. Qi, X. Li, Q. Niu and J. Feng, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2015, **92**, 121403.
- 24 Q. Zhang, S. A. Yang, W. Mi, Y. Cheng and U. Schwingenschlogl, *Adv. Mater.*, 2016, **28**, 959–966.
- 25 G. Yang, J. Li, H. Ma, Y. Yang, C. Li, X. Mao and F. Yin, *Phys. Rev. B*, 2018, **98**, 235419.
- 26 W. Zhou, Z. Yang, A. Li, M. Long and F. Ouyang, *Phys. Rev. B*, 2020, **101**, 045113.
- 27 C. Zhao, T. Norden, P. Zhang, P. Zhao, Y. Cheng, F. Sun, J. P. Parry, P. Taheri, J. Wang, Y. Yang, T. Scrace, K. Kang, S. Yang, G. X. Miao, R. Sabirianov, G. Kioseoglou, W. Huang, A. Petrou and H. Zeng, *Nat. Nanotechnol.*, 2017, **12**, 757–762.
- 28 X. Liang, L. Deng, F. Huang, T. Tang, C. Wang, Y. Zhu, J. Qin, Y. Zhang, B. Peng and L. Bi, *Nanoscale*, 2017, **9**, 9502–9509.
- 29 Y. Song, Q. Zhang, W. Mi and X. Wang, *Phys. Chem. Chem. Phys.*, 2016, **18**, 15039–15045.
- 30 Y. Song, X. Wang and W. Mi, *Adv. Electron. Mater.*, 2017, **3**, 1700245.
- 31 T. Norden, C. Zhao, P. Zhang, R. Sabirianov, A. Petrou and H. Zeng, *Nat. Commun.*, 2019, **10**, 4163.
- 32 K. Zollner, P. E. Faria and J. Fabian, *Phys. Rev. B*, 2019, **100**, 085128.
- 33 K. L. Seyler, D. Zhong, B. Huang, X. Linpeng, N. P. Wilson, T. Taniguchi, K. Watanabe, W. Yao, D. Xiao, M. A. McGuire, K. C. Fu and X. Xu, *Nano Lett.*, 2018, **18**, 3823–3828.
- 34 Y. C. Cheng, Q. Y. Zhang and U. Schwingenschlögl, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2014, **89**, 155429.
- 35 R. Peng, Y. Ma, S. Zhang, B. Huang and Y. Dai, *J. Phys. Chem. Lett.*, 2018, **9**, 3612–3617.
- 36 X. W. Zhao, Y. Li, R. D. Liang, G. C. Hu, X. B. Yuan and J. F. Ren, *Appl. Surf. Sci.*, 2020, **504**, 144367.
- 37 J. Zhou, J. Lin, H. Sims, C. Jiang, C. Cong, J. A. Brehm, Z. Zhang, L. Niu, Y. Chen, Y. Zhou, Y. Wang, F. Liu, C. Zhu, T. Yu, K. Suenaga, R. Mishra, S. T. Pantelides, Z. G. Zhu, W. Gao, Z. Liu and W. Zhou, *Adv. Mater.*, 2020, **32**, 1906536.
- 38 S. J. Yun, D. L. Duong, D. M. Ha, K. Singh, T. L. Phan, W. Choi, Y. M. Kim and Y. H. Lee, *Adv. Sci.*, 2020, **7**, 1903076.

- 39 Y. T. H. Pham, M. Liu, V. O. Jimenez, Z. Yu, V. Kalappattil, F. Zhang, K. Wang, T. Williams, M. Terrones and M. H. Phan, *Adv. Mater.*, 2020, **32**, 2003607.
- 40 F. Zhang, B. Zheng, A. Sebastian, D. H. Olson, M. Liu, K. Fujisawa, Y. T. H. Pham, V. O. Jimenez, V. Kalappattil, L. Miao, T. Zhang, R. Pendurthi, Y. Lei, A. L. Elias, Y. Wang, N. Alem, P. E. Hopkins, S. Das, V. H. Crespi, M. H. Phan and M. Terrones, *Adv. Sci.*, 2020, **7**, 2001174.
- 41 S. Fu, K. Kang, K. Shayan, A. Yoshimura, S. Dadras, X. Wang, L. Zhang, S. Chen, N. Liu, A. Jindal, X. Li, A. N. Pasupathy, A. N. Vamivakas, V. Meunier, S. Strauf and E. H. Yang, *Nat. Commun.*, 2020, **11**, 2034.
- 42 H. Yuan, M. S. Bahramy, K. Morimoto, S. Wu, K. Nomura, B.-J. Yang, H. Shimotani, R. Suzuki, M. Toh, C. Kloc, X. Xu, R. Arita, N. Nagaosa and Y. Iwasa, *Nat. Phys.*, 2013, **9**, 563–569.
- 43 D. Xiao, G. B. Liu, W. Feng, X. Xu and W. Yao, *Phys. Rev. Lett.*, 2012, **108**, 196802.
- 44 B. Zhou, *Nanoscale*, 2020, **12**, 5533–5542.
- 45 J. Qi, H. Wang, X. Chen and X. Qian, *Appl. Phys. Lett.*, 2018, **113**, 043102.
- 46 G. Kresse and J. Furthmüller, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1996, **54**, 11169–11186.
- 47 J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **77**, 3865–3868.
- 48 M. Fuchs and M. Scheffler, *Comput. Phys. Commun.*, 1999, **119**, 67–98.
- 49 G. Kresse and D. Joubert, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1999, **59**, 1758–1775.
- 50 P. E. Blochl, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **50**, 17953–17979.
- 51 S. L. Dudarev, G. A. Botton, S. Y. Savrasov, C. J. Humphreys and A. P. Sutton, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1998, **57**, 1505–1509.
- 52 T. Zhang, Y. Ma, X. Xu, C. Lei, B. Huang and Y. Dai, *J. Phys. Chem. C*, 2020, **124**, 20598–20604.
- 53 A. A. Mostofi, J. R. Yates, G. Pizzi, Y.-S. Lee, I. Souza, D. Vanderbilt and N. Marzari, *Comput. Phys. Commun.*, 2014, **185**, 2309–2310.
- 54 W. Bai, Z. Hu, C. Xiao, J. Guo, Z. Li, Y. Zou, X. Liu, J. Zhao, W. Tong, W. Yan, Z. Qu, B. Ye and Y. Xie, *J. Am. Chem. Soc.*, 2020, **142**, 10849–10855.
- 55 M. A. Gave, D. Bilc, S. D. Mahanti, J. D. Breshears and M. G. Kanatzidis, *Inorg. Chem.*, 2005, **44**, 5293–5303.
- 56 L. Ju, J. Shang, X. Tang and L. Kou, *J. Am. Chem. Soc.*, 2020, **142**, 1492–1500.
- 57 B. Xu, H. Xiang, Y. Xia, K. Jiang, X. Wan, J. He, J. Yin and Z. Liu, *Nanoscale*, 2017, **9**, 8427–8434.
- 58 Q.-F. Yao, J. Cai, W.-Y. Tong, S.-J. Gong, J.-Q. Wang, X. Wan, C.-G. Duan and J. H. Chu, *Phys. Rev. B*, 2017, **95**, 165401.
- 59 A. Togo, F. Oba and I. Tanaka, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2008, **78**, 134106.
- 60 A. Togo and I. Tanaka, *Scr. Mater.*, 2015, **108**, 1–5.
- 61 J. Heyd, G. E. Scuseria and M. Ernzerhof, *J. Chem. Phys.*, 2003, **118**, 8207–8215.
- 62 C. Lei, Y. Ma, T. Zhang, X. Xu, B. Huang and Y. Dai, *New J. Phys.*, 2020, **22**, 033002.
- 63 X. Xu, Y. Ma, T. Zhang, C. Lei, B. Huang and Y. Dai, *J. Phys. Chem. Lett.*, 2019, **10**, 4535–4541.
- 64 N. Singh and U. Schwingenschlogl, *Adv. Mater.*, 2017, **29**, 1600970.
- 65 R. Peng, Y. Ma, X. Xu, Z. He, B. Huang and Y. Dai, *Phys. Rev. B*, 2020, **102**, 035412.
- 66 Y. Zang, Y. Ma, R. Peng, H. Wang, B. Huang and Y. Dai, *Nano Res.*, 2020, **14**, 834–839.
- 67 P. Zhao, Y. Ma, C. Lei, H. Wang, B. Huang and Y. Dai, *Appl. Phys. Lett.*, 2019, **115**, 261605.
- 68 T. Cai, S. A. Yang, X. Li, F. Zhang, J. Shi, W. Yao and Q. Niu, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2013, **88**, 115140.
- 69 X. Xu, Z. He, Y. Dai, B. Huang and Y. Ma, *Phys. Rev. B*, 2021, **104**, 205430.