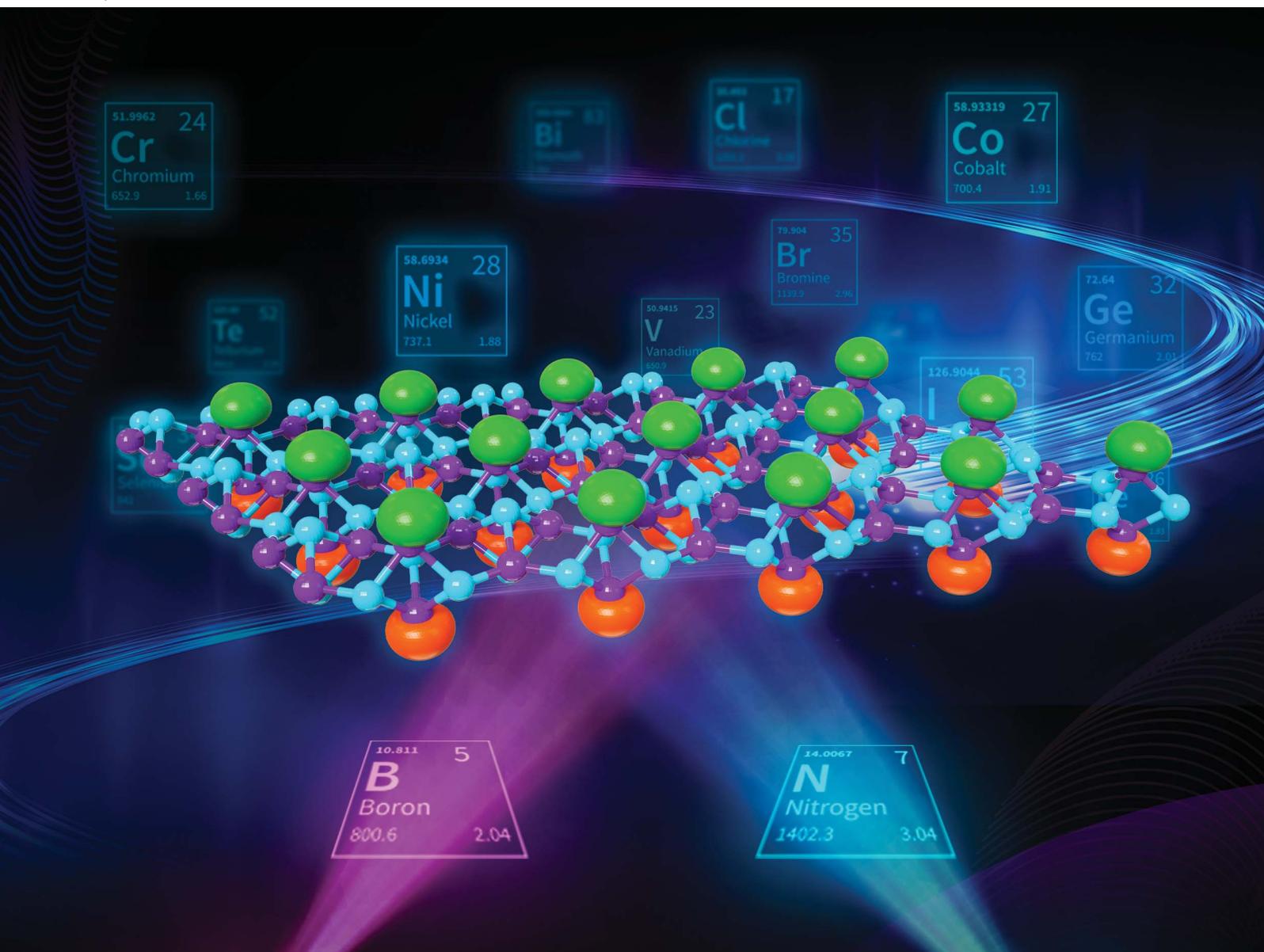


Nanoscale Advances

rsc.li/nanoscale-advances



ISSN 2516-0230

PAPER

Dong Zhang, Kai Chang *et al.*
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Cite this: *Nanoscale Adv.*, 2020, 2, 4421

Received 5th April 2020

Accepted 19th May 2020

DOI: 10.1039/d0na00270d

rsc.li/nanoscale-advances

B₅N₅ monolayer: a room-temperature light element antiferromagnetic insulator†

Dong Zhang,¹ Qihua Xiong^{2,3} and Kai Chang^{4,5}

We demonstrate theoretically that an intrinsic antiferromagnetic phase exists in monolayer materials consisting of non-magnetic light atoms, and propose that B₅N₅ with a decorated bounce lattice is a thermodynamically stable two-dimensional antiferromagnetic insulator by performing state-of-the-art density functional theory calculations. The antiferromagnetic phase originates from spontaneous symmetry breaking at the nearly flat bands in the vicinity of the Fermi energy. The flat bands are formed by purely s-p_z orbitals and are spin degenerate. A perpendicular electric field can remove the spin degeneracy and a prototype controllable dual spin filter with 100% spin polarization is proposed. Our proposal offers a possible two-dimensional atomically thick antiferromagnetic insulator.

1 Introduction

Magnetism, the ordered arrangement of neighbouring spins over macroscopic length scales, tends to be destroyed by thermal fluctuations above certain critical temperature, and the Curie temperatures depend critically on the dimensionality of the systems. In one dimensional (1D) systems, the long-range magnetic order only exists at $T = 0$,¹ while in three dimensional (3D) systems, the magnetic phase transition occurs at a finite temperature. In two dimensional (2D) systems, the situation becomes more complicated. In continuous and isotropic 2D systems, according to the Mermin-Wagner theorem,² a macroscopic magnetic order can be destroyed by long-range fluctuations at a finite temperature, because the gapless spin waves have a finite density of states and can be easily excited at any non-zero temperature. However, the exact Onsager solution to the 2D Ising model³ indicates that a magnetic phase transition occurs at a critical temperature $T_c > 0$. The controversial theoretical predictions make the search for 2D magnets extremely attractive in terms of both basic physics and potential applications.

The family of atomically thick magnetic 2D materials has expanded rapidly in recent years, and pioneering efforts have been taken through exfoliation of monolayer and few-layer

magnetic layered bulk materials, such as NiPS₃,⁴ FePS₃,⁵⁻⁷ and CrSiTe₃.⁸ However, macroscopic magnetic orders are only observed in thin films. The atomically-thick CrI₃ (ref. 9) and Cr₂Ge₂Te₆ (ref. 10) down to the monolayer and bilayer limit have been successfully fabricated and the corresponding experiments finally demonstrated the existence of intrinsic ferromagnetism (FM). Afterwards, a number of 2D ferromagnets such as Fe₃GeTe₂,¹¹⁻¹³ 1T-VSe₂,¹⁴ MnSe_x,¹⁵ etc., mushroomed. All the monolayer 2D FM materials display significant magnetic anisotropy which opens a magnon excitation gap and lifts the Wagner-Mermin criteria to a finite T_c by suppressing thermal fluctuations. Discoveries of monolayer ferromagnets not only enriched the 2D family, but also provided us new platforms to investigate emergent interlayer exchange interactions in van der Waals (vdW) heterostructures.^{16,17} Compared to the rapid progress of 2D ferromagnets, low-dimensional antiferromagnets only appear as interlayer couplings in vdW ferromagnetic multilayers,^{9,18} and are rarely reported in the monolayer limit. This scenario requires precise control of the number of multilayers, since an even number of layers leads to ferromagnetism and only an odd number of layers causes antiferromagnetism.⁹

Apart from the magneto-crystalline anisotropy preserved magnetic order in 2D materials, s-p electron magnetism provides an alternative way to realize low-dimensional magnets and shows particular advantages of long spin relaxation times and lengths due to weak intrinsic spin orbit coupling. It has been initially demonstrated that structural defects and topological defects in 2D materials can host unpaired spins and even observable magnetic orders,¹⁹⁻²² but the robustness of the macroscopic magnetic order is still debatable. In addition to the randomly distributed defects, the localized edge states of zigzag graphene nanoribbons have been theoretically proposed to be a robust antiferromagnet,²³⁻²⁵ and such an antiferromagnetic feature has been experimentally confirmed at room

¹SKLSM, Institute of Semiconductors, Chinese Academy of Sciences, P.O. Box 912, Beijing 100083, China. E-mail: zhangdong@semi.ac.cn; kchang@semi.ac.cn

²Center for Excellent in Topological Quantum Computation, University of Chinese Academy of Sciences, Beijing 100190, China

³Division of Physics and Applied Physics, School of Physical and Mathematical Sciences, Nanyang Technological University, Singapore, Singapore

⁴State Key Laboratory of Low-Dimensional Quantum Physics, Department of Physics, Tsinghua University, Beijing 100084, China

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0na00270d



temperatures.^{26,27} Although the magnetic orderings sensitively depend on the widths and edge configurations of graphene nanoribbons, these novel 2D magnets originate from the spontaneous symmetry breaking, which occurs at the flat bands with high density of states (DOS) in the vicinity of the Fermi levels, making 2D materials consisting of light elements and hosting flat bands promising candidates to realize 2D magnets.

2 Results

Inspired by the flat band induced s-p magnetism in graphene nanoribbons and artificially designed two-dimensional materials,^{28–30} we searched for 2D materials that possess flat bands which are protected by the crystal symmetries. To further enhance the magnetocrystalline anisotropy, in this work, we propose a new type of allotrope for h-BN with a decorated bounce lattice as shown in Fig. 1, since the 2-nd row elements boron and nitride behave like glue atoms and form versatile allotropes.³¹ The crystalline structure of the 2D B₅N₅ hosts the D_{3h} symmetry with the space group *P6̄m2* (no. 187), and possesses a trilayer configuration. In the middle of the trilayer, there lies a bounce lattice connecting adjacent boron and nitrogen equilateral triangles with a pair of B–N bonds, and the upper and lower layers possess a honeycomb lattice with boron and nitrogen atoms taking positions above and beneath the centers of the nitrogen and boron triangles alternatively, as illustrated in Fig. 1(a).

The unitcell of this type of allotrope is indicated in Fig. 1(b) with dashed lines, and in each unitcell, there are two dual triangular bipyramids consisting of B₃N₂ and B₂N₃ clusters, respectively. Considering the unbalanced chemical compositions of the two bipyramids, structural distortions are expected and verified by structure relaxation. From Fig. 1(b), one can find out that the B₂N₃ and the B₃N₂ clusters are not symmetric, but shrink along horizontal and perpendicular directions, respectively. The fully relaxed lattice constant of the unitcell is $a = b =$

4.875 Å, and the lengths of the B–N bonds within the B₂N₃ and B₃N₂ clusters, and those bridging the two clusters are 1.612 Å, 1.579 Å, and 1.533 Å, respectively. To investigate the stability of the optimized crystal, we calculated the phonon dispersions by employing the frozen phonon method,³² and the phonon dispersions are shown in Fig. 1(c). No imaginary frequencies can be found throughout the whole Brillouin Zone (BZ) in phonon dispersions, indicating thermodynamic structural stability of the 2D structure. Therefore, we adopted the relaxed lattice parameters for all the subsequent calculations.

To calculate the electronic structures of this new type of monolayer allotrope, we set up a vacuum slab as thick as 20 Å, to ensure separation between adjacent monolayers to eliminate interlayer interactions, and perform the electronic structure calculations. The band structures of monolayer B₅N₅ in the vicinity of the Fermi level in the first BZ are shown in Fig. 2. From Fig. 2, one can see that the band gap is about 1.68 eV, and the highest valence band is nearly absolutely flat, with a bandwidth of about 20 meV. We analyse the orbital components of the flat band along a typical reciprocal path K–Γ–M, and denote each orbital with different coloured curves as displayed in Fig. 2(b), and the weight of each orbital is represented by the size of the corresponding circle. From Fig. 2(b), one can see that the s and p_z orbitals are dominant in the flat band (the highest valence band), the lowest two conduction bands are almost pure p_z and the second and third valence bands are comprised of almost equal p_x and p_y orbitals. The corresponding densities of states of orbitals are shown in Fig. 2(c), and are denoted by solid lines with the same colours as in Fig. 2(b).

To clarify the origin of the flat band, we calculated its spatial charge distributions as illustrated in Fig. 3(b). From Fig. 3(b), it can be found that the wave functions of the flat band locate at the top and bottom boron atoms, and possess typical hybridized s-p_z orbital characteristics. In such a nearly absolute flat band, the kinetic potentials of the carriers are almost quenched, and the electron correlations become more important. Spin polarized band structure calculations, as shown in Fig. 3(a), reveal

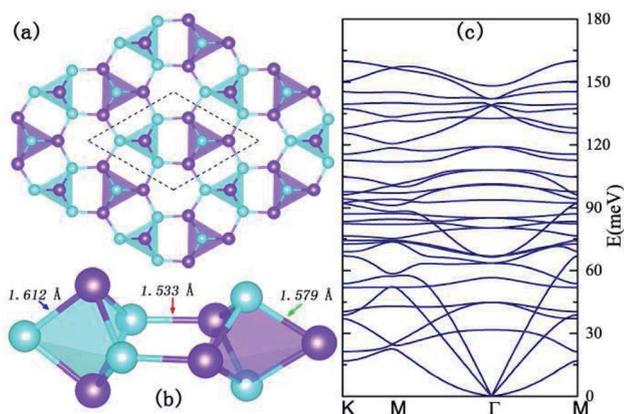


Fig. 1 Crystal structure and stability. (a) Top-view of the B₅N₅ allotrope, the unitcell is indicate in the dashed area and the boron atoms and nitride atoms are indicated with purple and azure spheres, respectively. (b) The side view of the relaxed structure within a unitcell. (c) The phonon dispersions of the B₅N₅ monolayer in the Brillouin zone.

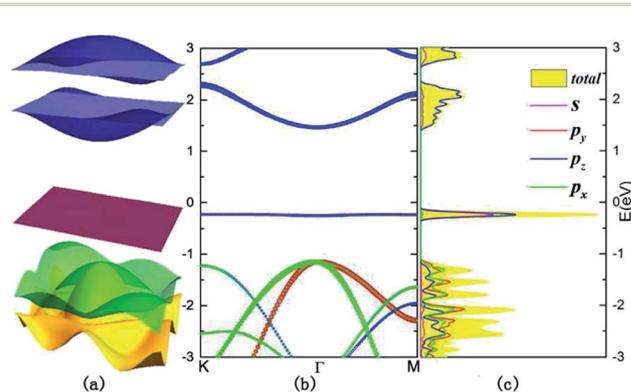


Fig. 2 Electronic structures. (a) 3-D band structures of B₅N₅ within the first BZ. (b) The band structures of B₅N₅, the Fermi surface is set to be 0 and the orbital components are indicated by separate color circles, magenta for s, red for p_y, blue for p_z, and green for p_x orbitals, respectively. (c) The densities of states (DOS) on the same energy scale as (b), the projected DOS are indicated by solid lines with the corresponding color codings.



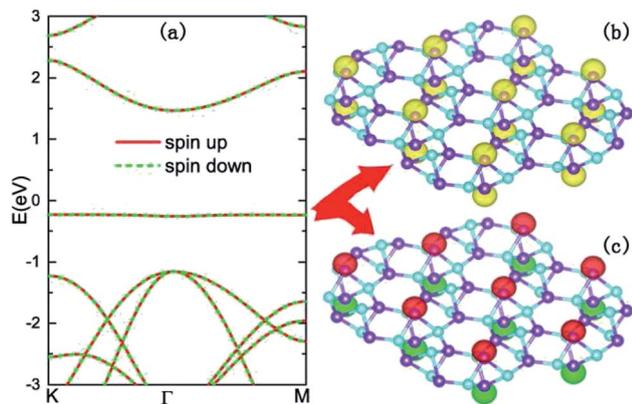


Fig. 3 Antiferromagnetic insulator. (a) Spin polarized band structures of B_5N_5 , the spin-up and spin-down dispersions are denoted with red solid lines and green dashed lines, respectively. (b) Spatial charge density distributions (yellow isosurfaces) of the flat-bands in the vicinity of the Fermi level. (c) Spatial spin density distributions of the spin-degenerate flat-bands, the spin-up and spin-down components are denoted with red and green isosurfaces, respectively.

that all the energy bands are doubly degenerate with anti-parallel spin states. However, the spatial distributions of opposite spins are separated. In Fig. 3(c), the upward and downward spins display intralayer ferromagnetic and interlayer antiferromagnetic orderings, making the monolayer B_5N_5 a 2D antiferromagnetic insulator.

To demonstrate that the antiferromagnetic insulating phase is the ground state of B_5N_5 , various calculations, such as the rotationally invariant DFT+U approach with a series of Hubbard U corrections up to 5 eV, and tunable exchange interactions within the HSE06 scheme are performed. The antiferromagnetic insulator feature remains unchanged (see ESI Part I†). The ground state possesses an A-type antiferromagnet and aligns perpendicular to the 2D material, and it can be conveniently described using an effective Hamiltonian as follows,

$$\hat{H}_{\text{eff}} = \sum_{i=1}^N S_{i\downarrow} S_{i\uparrow} J_{\perp} + \sum_{u=\uparrow,\downarrow} \sum_{j=1}^6 S_u S_{u,j} J_{\parallel} \quad (1)$$

where J_{\perp} and J_{\parallel} indicate the interlayer and intralayer magnetic exchange coupling parameters, u and d indicate upper and lower sub-layers, and i and j are the nearest in-plane lattice sites of decorated boron atoms. S is the net magnetic moment at the boron sites where spin accumulates.

In order to determine quantitatively the magnetic exchange coupling parameters J_{\perp} and J_{\parallel} , various magnetic orderings were studied based on constrained local spin density approximations (CLSDA),³³ and Fig. 4 exhibits the relative total energy differences between the antiferromagnetic structure, ferromagnetic structure and two intermediate magnetic structures. To compare the total energy differences, the collinear magnetic structures are calculated and displayed with a 3×3 supercell. The antiferromagnetic structure has the minimal total energy with a 0 net magnetic moment, because the doubly degenerate flat band is occupied by two electrons with opposite spins as indicated by the density of states in the lower panel of Fig. 4.

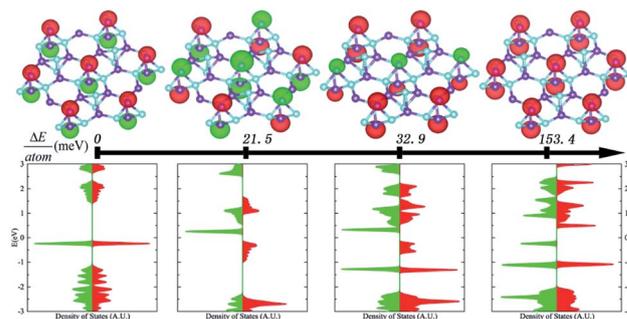


Fig. 4 Antiferromagnetic insulator as the ground state. Total energy differences per atom between various magnetic structures. In the upper panel and from the left to right, the magnetic structures of antiferromagnetic, intermediate and ferromagnetic structures are displayed. The middle arrow indicates the increase of the energy differences. And the corresponding spin densities of states are illustrated in the lower panel. The spin-up and spin-down states are denoted with red and green color coding, respectively.

The ferromagnetic counterpart has the maximal total energy with a net magnetic moment of $2 \mu_B$. Since the spin densities of the two configurations are identical in magnitude, the net magnetic moment at the boron sites S is $1 \mu_B$, and shows intrinsic s-p electron magnetism. By constructing magnetic orderings with total energy differences, the magnetic exchange coupling parameters can be numerically derived by applying the following formula with the output of constrained DFT. The total energy of the studied magnetic orderings can be expressed as the formula as follows:

$$E_{\text{total}} = \frac{\sum_{i=1}^N S_{i\downarrow} S_{i\uparrow} J_{\perp} + \sum_{u=\uparrow,\downarrow} \sum_{j=1}^6 S_u S_{u,j} J_{\parallel}}{N} + \Delta \quad (2)$$

where N is the multiple of the calculated supercell with respect to the original unitcell, as well as the number of magnetic dipoles. Specifically, $N = 1$ for AFM and FM orderings, while $N = 3$ for the intermediate model 2 and $N = 2$ for the intermediate model 3. $j = 1, \dots, 6$ is the index of the coordinate spin accumulated sites. And the total energies are divided by N to be compared with an equal number of atoms, and Δ is the non-magnetic total energy of the unitcell. (The details of the solved equations are shown in ESI Part II.†)

The magnetic exchange coupling parameters J_{\perp} and J_{\parallel} are mapped and $J_{\perp} = 76.7$ meV and $J_{\parallel} = -88.9$ meV. The positive J_{\perp} and negative J_{\parallel} indicate the interlayer antiferromagnetic and intralayer ferromagnetic orderings of monolayer B_5N_5 . With these parameters, the Néel temperature T_N can be approximately estimated as 629.22 K by fitting the modified random phase approximation (RPA) like relation,³⁴ $T_N = -2\pi\rho_s/[b - \ln(-2J_{\perp}/J_{\parallel})]$, where $\rho_s = 0.183J_{\parallel}$, and $b = 2.43$ is an empirical constant. Although the mean-field approximation usually overestimates T_N , the 2D antiferromagnet could be observed at room temperatures.

Antiferromagnetism surviving at room temperatures makes monolayer B_5N_5 a possible building block to construct flexible 2D spintronic devices. To mimic realistic interfacial conditions



such as charge transfer induced electric fields and lattice-mismatch induced stress in van der Waals heterostructures, the electronic structures of B_5N_5 under external stress and electric fields were investigated. The responses of B_5N_5 to external biaxial stress are relatively complicated, where tensile biaxial stress enhances the interlayer coupling strength but weakens the intralayer coupling strength. In the reciprocal space, the second valence band is pushed away from the isolated flat band, and the magnetic structure remains an A-type antiferromagnetic insulator (intralayer ferromagnetic and interlayer antiferromagnetic). In contrast, compressive biaxial stress enhances the intralayer coupling but weakens the interlayer coupling, and mixed the p_x - p_y dominant valence band with the s - p_z flat band. The band gap reduces significantly and the magnetic structure of the system changes to a G-type antiferromagnetic semiconductor (both intralayer and interlayer antiferromagnetic). The A-type to G-type phase transition at a critical compressive biaxial strain of about -3% and the evolution of band structures and spin orderings under compressive biaxial stress are exhibited (see ESI Part III†). Nevertheless the antiferromagnetic feature remains under both tensile and compressive stress.

Meanwhile perpendicular electric fields are effective to remove the degeneracy of the flat band (the dependency of band splitting on perpendicular electric fields is shown in ESI Part IV†), and separate the spin-up and spin-down channels without overlapping, and therefore a dual spin filter with high spin selectivity can be expected. The underlying physics comes from the feature of the flat bands, which are degenerate in the energy space while separated in the real space. In the presence of the vertical distance between the upper and lower boron sub-layers, a perpendicular electric field will induce an electrostatic potential between the states of opposite spins. As a consequence, perpendicular electric fields are effective to lift the two-

fold degenerate flat bands, and drive the system into a half metallic phase.

A prototype dual spin filter device is shown in Fig. 5. In Fig. 5(a), two top gates are applied perpendicularly on the monolayer B_5N_5 ; the first gate V_1 is responsible for separating spin-up and spin-down carriers, and the second gate V_2 works as a spin valve to switch on or off the spin channels. When the directions of V_1 and V_2 are parallel/anti-parallel, the spin channels are switched on/off, and the dual spin filter works in the ON/OFF state, as illustrated in Fig. 5(b) and (c). In Fig. 5(b) and (c), red and green solid lines are energy dispersions all over the BZ of spin-up and spin-down carriers in the vicinity of the Fermi surface, and both the gate voltages are $|V_1| = |V_2| = 3$ V, and the energetic relative locations of spin-up and spin-down states can be switched by the direction of V_1 , which makes spin states tunable. Since the upper states possess a narrow band width, a perpendicular gate voltage exceeding 0.58 V can eliminate energy overlapping between opposite spin states, and the spin polarization of the current can be as high as nearly 100%.

3 Discussion

In recent years, antiferromagnets have been recognized as promising future spintronic materials rather than merely passive components in exchange bias applications. The renaissance of antiferromagnets is due to their advantageous properties such as insensitivity to external magnetic fields and high intrinsic frequency dynamics.^{35–37} Owing to the rapid progress of 2D spintronics, electrically tunable and robust monolayer antiferromagnets are of particular interest.

In this work, we demonstrate that the intrinsic magnetic ordering survives even at room temperature in monolayer B_5N_5 with a decorated bounce lattice, which is a thermodynamically stable A-type 2D antiferromagnetic insulator. The antiferromagnetism in such a 2D material arises from the nearly flat bands in the vicinity of the Fermi energy, and is robust under external biaxial stress. A perpendicular electric field can remove the flat band degeneracy and a prototype dual spin filter is proposed. Our proposal not only offers a promising antiferromagnet candidate in the atomically-thin limit, but also provides an alternative way to explore antiferromagnetic insulators consisting of super-light and non-magnetic atoms.

4 Methods

Phonon dispersions are calculated with the PHONOPY code.³⁸ Electronic structure calculations of B_5N_5 are performed by using the Vienna *ab initio* simulation package (VASP)³⁹ within the screened-nonlocal exchange Heyd–Scuseria–Ernzerhof (HSE06)⁴⁰ hybrid functional of the generalized Kohn–Sham scheme. The projector augmented-wave (PAW) pseudopotentials⁴¹ are adopted, and the kinetic energy cut-off is set to be 560 eV for wave-function expansions, and a Γ -centred k -point grid is densely sampled by summing over $13 \times 13 \times 1$.⁴² Since both boron and nitride are super-light species with literally

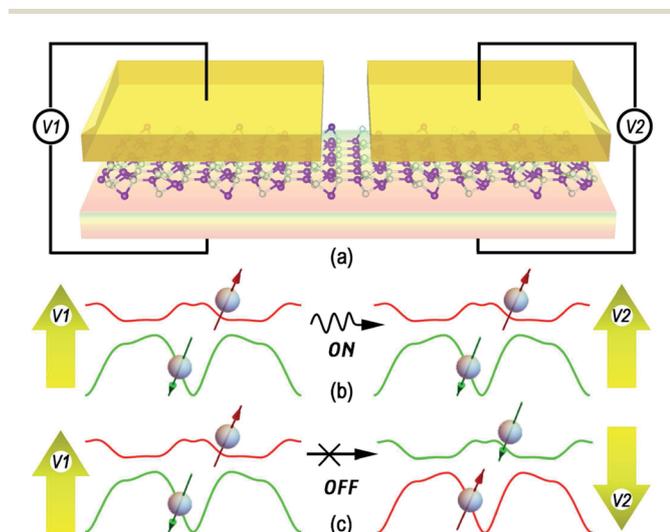


Fig. 5 Dual spin filter. (a) Schematic of the dual spin filter based on monolayer B_5N_5 , (b) on state and (c) off state of pure spin current obtained by applying parallel and anti-parallel voltages. Spin-up and spin-down states are denoted with red and green colour codings, respectively.



negligible intrinsic spin-orbit couplings (SOC), only the spin polarized band structure calculations are performed.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the National Key R&D Programmes of China, Grant No. 2017YFA0303400, 2016YFE0110000, National Natural Science Foundation of China, Grant No. 11574303, 11504366, Youth Innovation Promotion Association of Chinese Academy of Sciences, Grant No. 2018148, and the Strategic Priority Research Program of Chinese Academy of Sciences, Grant No. XDB28000000. D. Zhang is thankful for the discussion with Prof. Xiong and inspiration from his beloved daughter, Miss 4D. Zhang.

Notes and references

- 1 R. Peierls, *Math. Proc. Cambridge Philos. Soc.*, 1936, 477–481.
- 2 N. D. Mermin and H. Wagner, *Phys. Rev. Lett.*, 1966, 17, 1133.
- 3 L. Onsager, *Phys. Rev.*, 1944, 65, 117.
- 4 C. T. Kuo, M. Neumann, K. Balamurugan, H. J. Park, S. Kang, H. W. Shiu, J. H. Kang, B. H. Hong, M. Han, T. W. Noh and J. G. Park, *Sci. Rep.*, 2016, 6, 20904.
- 5 K. Z. Du, X. Z. Wang, Y. Liu, P. Hu, M. I. B. Utama, C. K. Gan, Q. Xiong and C. Kloc, *ACS Nano*, 2016, 10, 1738–1743.
- 6 J.-U. Lee, S. Lee, J. H. Ryoo, S. Kang, T. Y. Kim, P. Kim, C.-H. Park, J.-G. Park and H. Cheong, *Nano Lett.*, 2016, 16, 7433–7438.
- 7 X. Wang, K. Du, Y. Y. F. Liu, P. Hu, J. Zhang, Q. Zhang, M. H. S. Owen, X. Lu, C. K. Gan, P. Sengupta, C. Kloc and Q. Xiong, *2D Mater.*, 2016, 3, 1–9.
- 8 M. W. Lin, H. L. Zhuang, J. Yan, T. Z. Ward, A. A. Puretzky, C. M. Rouleau, Z. Gai, L. Liang, V. Meunier, B. G. Sumpter, P. Ganesh, P. R. Kent, D. B. Geohegan, D. G. Mandrus and K. Xiao, *J. Mater. Chem. C*, 2016, 4, 315–322.
- 9 B. Huang, G. Clark, E. Navarro-Moratalla, D. R. Klein, R. Cheng, K. L. Seyler, D. Zhong, E. Schmidgall, M. A. McGuire, D. H. Cobden, W. Yao, D. Xiao, P. Jarillo-Herrero and X. Xu, *Nature*, 2017, 546, 270–273.
- 10 C. Gong, L. Li, Z. Li, H. Ji, A. Stern, Y. Xia, T. Cao, W. Bao, C. Wang, Y. Wang, Z. Q. Qiu, R. J. Cava, S. G. Louie, J. Xia and X. Zhang, *Nature*, 2017, 546, 265–269.
- 11 B. Chen, J. H. Yang, H. D. Wang, M. Imai, H. Ohta, C. Michioka, K. Yoshimura and M. H. Fang, *J. Phys. Soc. Jpn.*, 2013, 82, 124711.
- 12 Z. Fei, B. Huang, P. Malinowski, W. Wang, T. Song, J. Sanchez, W. Yao, D. Xiao, X. Zhu, A. F. May, W. Wu, D. H. Cobden, J. H. Chu and X. Xu, *Nat. Mater.*, 2018, 17, 778–782.
- 13 Y. Deng, Y. Yu, Y. Song, J. Zhang, N. Z. Wang, Z. Sun, Y. Yi, Y. Z. Wu, S. Wu, J. Zhu, J. Wang, X. H. Chen and Y. Zhang, *Nature*, 2018, 563, 94–99.
- 14 M. Bonilla, S. Kolekar, Y. Ma, H. C. Diaz, V. Kalappattil, R. Das, T. Eggers, H. R. Gutierrez, M. H. Phan and M. Batzill, *Nat. Nanotechnol.*, 2018, 13, 289–293.
- 15 D. J. O'Hara, T. Zhu, A. H. Trout, A. S. Ahmed, Y. K. Luo, C. H. Lee, M. R. Brenner, S. Rajan, J. A. Gupta, D. W. McComb and R. K. Kawakami, *Nano Lett.*, 2018, 18, 3125–3131.
- 16 M. Gibertini, M. Koperski, A. F. Morpurgo and K. S. Novoselov, *Nat. Nanotechnol.*, 2019, 14, 408–419.
- 17 C. Gong and X. Zhang, *Science*, 2019, 363, eaav4450.
- 18 S.-J. Gong, C. Gong, Y.-Y. Sun, W.-Y. Tong, C.-G. Duan, J.-H. Chu and X. Zhang, *Proc. Natl. Acad. Sci. U. S. A.*, 2018, 115, 8511–8516.
- 19 M. M. Ugeda, I. Brihuega, F. Guinea and J. M. Gómez-Rodríguez, *Phys. Rev. Lett.*, 2010, 104, 096804.
- 20 J. Červenka, M. I. Katsnelson and C. F. Flipse, *Nat. Phys.*, 2009, 5, 840–844.
- 21 B. Uchoa, V. N. Kotov, N. M. R. Peres and A. H. C. Neto, *Phys. Rev. Lett.*, 2008, 101, 26805.
- 22 R. R. Nair, M. Sepioni, I. L. Tsai, O. Lehtinen, J. Keinonen, A. V. Krasheninnikov, T. Thomson, A. K. Geim and I. V. Grigorieva, *Nat. Phys.*, 2012, 8, 199–202.
- 23 Y. W. Son, M. L. Cohen and S. G. Louie, *Nature*, 2006, 444, 347–349.
- 24 J. Jung, T. Pereg-Barnea and A. H. MacDonald, *Phys. Rev. Lett.*, 2009, 102, 227205.
- 25 D. Zhang, D.-B. Zhang, F. Yang, H.-Q. Lin, H. Xu and K. Chang, *2D Mater.*, 2015, 2, 041001.
- 26 G. Z. Magda, X. Jin, I. Hagymási, P. Vancsó, Z. Osváth, P. Nemes-Incze, C. Hwang, L. P. Biró and L. Tapasztó, *Nature*, 2014, 514, 608–611.
- 27 M. Slota, A. Keerthi, W. K. Myers, E. Tret'yakov, M. Baumgarten, A. Ardavan, H. Sadeghi, C. J. Lambert, A. Narita, K. Müllen and L. Bogani, *Nature*, 2018, 557, 691–695.
- 28 Z. Liu, Z.-F. Wang, J.-W. Mei, Y.-S. Wu and F. Liu, *Phys. Rev. Lett.*, 2013, 110, 106804.
- 29 M. Zhou, Z. Liu, W. Ming, Z. Wang and F. Liu, *Phys. Rev. Lett.*, 2014, 113, 236802.
- 30 P. Le, B. Hoi and M. Yarmohammadi, *AIP Adv.*, 2018, 8, 125317.
- 31 M. Shahrokhi, B. Mortazavi and G. R. Berdiyrov, *Solid State Commun.*, 2017, 253, 51–56.
- 32 K. Parlinski, Z. Q. Li and Y. Kawazoe, *Phys. Rev. Lett.*, 1997, 78, 4063–4066.
- 33 B. Kaduk, T. Kowalczyk and T. Van Voorhis, *Chem. Rev.*, 2012, 112, 321–370.
- 34 C. Yasuda, K. Hukushima, F. Alet, M. Keller, M. Troyer and H. Takayama, *Phys. Rev. Lett.*, 2005, 94, 217201.
- 35 V. Baltz, A. Manchon, M. Tsoi, T. Moriyama, T. Ono and Y. Tserkovnyak, *Rev. Mod. Phys.*, 2018, 90, 015005.
- 36 M. B. Jungfleisch, W. Zhang and A. Hoffmann, *Phys. Lett. A*, 2018, 382, 865–871.
- 37 T. Jungwirth, J. Sinova, A. Manchon, X. Marti, J. Wunderlich and C. Felser, *Nat. Phys.*, 2018, 14, 200–203.
- 38 A. Togo, F. Oba and I. Tanaka, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2008, 78, 134106.



- 39 G. Kresse and J. Furthmüller, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1996, **54**, 11169–11186.
- 40 J. Heyd, G. E. Scuseria and M. Ernzerhof, *J. Chem. Phys.*, 2006, **124**, 219906.
- 41 P. E. Blöchl, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **50**, 17953–17979.
- 42 H. J. Monkhorst and J. D. Pack, *Phys. Rev. B: Solid State*, 1976, **13**, 5188–5192.

