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Enhanced photocatalytic properties of a chemically modified blue phosphorene†

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It is high time to placate the peak demand for an efficient, economic and green fuel in the form of H₂ through photocatalytic water splitting. Several low dimensional materials have been explored for their photocatalytic properties on account of their surface to volume ratio. The present study illustrates the excellent photocatalytic potential of a two-dimensional material, viz. a chemically tempered blue-phosphorene sheet, with single atom thickness and high carrier mobility. Metal-free element, sulphur, is explored as a dopant in a 32-atom blue-phosphorene sheet. The dopant is inserted at three locations viz. central, edge and central edge positions with varying concentrations from 3.125% to 18.75% (corresponding to $n = 1$ to 6 sulphur atoms within a 32-atom blue-phosphorene sheet, P_{32-n}S_n). The cohesive energy studies predict the higher stability of even number S doped sheets as compared to their odd counterparts. Photocatalytic activity is studied in terms of band gap and band alignment for different concentrations of the former. Studies reveal that edge doping demonstrates better water molecule activation independent of S atom concentration. The edge doped systems not only provide the chemical activity to activate water, but also show feasible HER overpotentials of 1.24–1.29 eV at neutral medium. Finally, this work opens up a driving lead of non-corrosive catalysts for water molecule splitting.

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Introduction

Low dimensional materials have been a buzzing topic of research in the last two decades due to their characteristic physico-chemical properties. Various quantum phenomena such as phonon–electron interactions, spin interaction, quantum tunnelling effect and quantum confinement within these materials are responsible for their enhanced and yet tuneable electronic, thermal, magnetic and optical properties with far reaching applications in novel devices such as spintronics, photonics, quantum computing and catalytic materials. However, the most widely applied area of these materials is catalysis on account of their higher surface to volume ratio and higher electron transferability making it cost effective.^{1–6} Among the various low dimensional nano-structures such as quantum dots/nanoparticles (zero dimensional), 1D nanotubes/nanorods and 2D nanofilms/nanosheets, 2D based materials have the upper hand with respect to other dimensional substances as catalysts due to the aforesaid reasons. (a) Firstly, each site on

their surface is equally accessible for catalytic process.⁷ (b) Secondly, these materials offer higher thermal and mechanical stability and thus, operational ability at various temperatures.⁸ (c) Due to their promising thermal conductivity and large surface space, they also act as efficient heat sinks during exothermic reaction.⁹ (d) 2D sheet being atomic thick, reduces charge carrier distance from the core to top, thereby reducing electron–hole recombination and enhancing the redox process marginally.¹⁰ (e) They offer wide range of options for feasible chemical engagement, thereby enabling charge storage.¹¹ (f) Lastly, quantum confinement and unsaturated surfaces leads to facile movement of electrons along in-plane direction, thus increasing the lifetime of charge carrier.^{12,13}

The journey of 2D materials based catalysts started with graphene, a single atom thick carbon sheet held by sp² hybrid bonds arranged in hexagonal manner.¹⁴ Employment of graphene-based materials, to catalysis quickly has captured the interests of researchers in the last few decades.^{14,15} However, due to its intrinsic zero band gap and consequently its instantaneous electron–hole recombination, its application in photocatalysis remains challenging.^{16,17} Although, its oxygenated counterpart is seen to have finite band gap of 2.4 eV to 4.3 eV,^{18–20} it fails to be within the acceptable slot of 1.6–2.2 eV so as to be activated efficaciously under natural solar light irradiation.²¹ Several other 2D materials such as silicene,²² germanene,²³ borophene,^{24,25} stanene^{26,27} and mono-chalcogenides²⁸ have been explored for their potential

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application as photocatalysts.²⁸ However, absence of a tunable band gap within the acceptable slot has limited their practical applications.^{29,30}

In this context, phosphorene, a mono-layer/multi-layer sheet of black phosphorous,³¹ with its tunable band gap ranging between 0.3–2.5 eV has been found to be more attractive as a photocatalyst as compared to other 2D materials.^{32–35} Following the fact finding journey, many other allotropes/polytypes of phosphorene *viz.*, red, black, violet, green phosphorene have been realized^{36–39} and some of them have been explored for water splitting application.^{40–43} More recently, a new member of phosphorene polytype was realized through the synthesis of mono-layered honeycomb buckled blue-phosphorene (B-phosphorene) on an Au (111) sheet using a molecular beam epitaxial method with black-phosphorous as precursor.⁴⁴ It has been reported that a specific dislocation in black phosphorene induces a new structure with zigzag puckering, called B-phosphorene with a characteristic tunable bandgap upto 2 eV.⁴⁴ This B-phosphorene is found to be more stable thermodynamically as compared to black-phosphorene,^{45,46} which is of great importance for an efficient photocatalyst.^{47–49} These properties inspired the researchers to further examine the photocatalytic properties of nanotube B-phosphorene,⁵⁰ doped⁵¹ and several hetero-structures of B-phosphorene.^{52–54} Research works also focused on engineering the band gap of B-phosphorene by application of external electric field,⁵⁵ using stacking effect,^{56,57} functionalization^{58–60} and doping.^{61–63} Doping is the most attractive mode among all the methods for modulating physico-chemical properties^{64–75} of phosphorenes (or for any given material).^{76,77} From the experimental point of view, single atomic layered 2D structures can be doped by traditional substitutional doping methods such as implantation,⁷⁸ diffusion processes^{79–81} or new techniques such as surface charge transfer method,⁸² intercalation,^{83,84} and field effect modulation⁸⁵ methods.

Non-metal doped B-phosphorene are synthesized and reported in several experimental studies.^{86,87} Their physicochemical as well as electronic properties are also explored theoretically.^{88–91} Further, their corresponding application as catalytic materials is also studied by some groups.^{92–95} In the context of application as photocatalytic material, ample experimental studies of doped B-phosphorene has been reported where N is employed as dopant.^{96–98} The photocatalytic properties of N, Si, S, doped B-phosphorene have also been studied theoretically.^{99,100} Moreover, sulphur has also been earlier reported to be a successful catalyst in the context of sensors and opto-electronics.^{39,99} Further, S having the same atomic size to that of P atom, plays a vital role in improving its stability.¹⁰¹ The anionic substitutional doping of S results in puckered structure of phosphorene that facilitates anion atoms to get exposed to the outer surface as compared to planar 2D materials like graphene.¹⁰² From above studies and available literature reports it is clear that there are very few studies on the photocatalytic activity of sulphur doped B-phosphorene. More importantly, the influence of varying concentration of S on the stability and catalytic activity of B-phosphorene is not known in terms of water molecule activation.^{103,104} In the present work, using

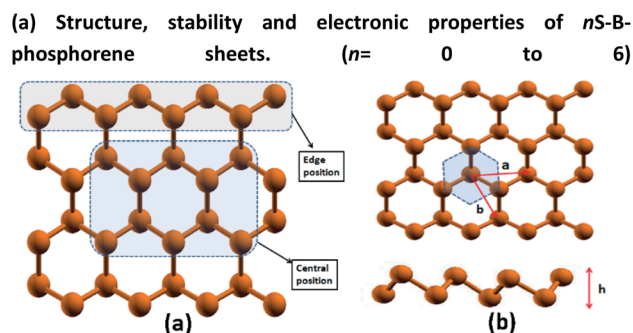


Fig. 1 (a) Central (C) and edge (E) atoms in B-phosphorene sheet. (b) Relaxed structure of B-phosphorene from top and side views (a and b are lattice parameters) Wigner–Seitz cells are shown by the shaded area.

density functional theory, we evaluate the role of S doping, its concentration and its position in modulating the catalytic activity of B-phosphorene in water molecule activation.

With reference to the above we vary the concentration of S from 3.125% to 18.75% in 32 B-phosphorene-sheet. The S atoms are doped at three different ways; (a) all S atoms are placed in the central position (see Fig. 1(a) for elucidation) (b) all S atoms are spread across both edge and central & edge position as shown in Fig. 1(a). Stability of thus doped systems is evaluated in terms of cohesive energies, band gap and band edges. pH is taken into account to ensure that the doped structures qualify to be photocatalyst practically. Finally, the models that qualify the photocatalytic properties are adsorbed with water molecule. Activation of water molecule is evaluated in terms of adsorption energy of water molecule, extent of O–H bond elongation from the optimal O–H bond length in water and the red shift in O–H stretching frequency. The thermostability of the sulphur-doped blue phosphorene systems has also been evaluated through *ab initio* molecular dynamics (AIMD) simulations at 500 K.

Computational details

The phosphorene sheets, pristine and doped, considered in this study are modelled using GaussView06 Software.¹⁰⁵ Density Functional Theory (DFT) calculations are carried out on these sheets using SIESTA¹⁰⁶ (Spanish Initiative for Electronics Simulations with Thousands of Atoms) which uses flexible linear combination of atomic orbital (LCAO) as a basis set. Norm-conserving pseudopotentials are taken up to describe the electron–ion interaction, which was yielded by implementing the Troullier–Martins scheme.¹⁰⁷ A Generalized Gradient Approximation–Perdew, Burke, Ernzerhof (GGA–PBE) functional is used for the calculation of correlation and exchange energies¹⁰⁸ along with a double-zeta plus (DZP) polarization basis set for all elements. Spin polarized optimization of all the structures are carried out using first-order minimization method, which is known as Conjugated Gradient (CG). All structures are relaxed until the force on each atom is less than $0.01 \text{ eV } \text{\AA}^{-1}$. In the Self-Consistent Field (SCF) process, the Brillouin Zone (BZ) of the reciprocal lattice is sampled through a $6 \times 6 \times 1$ grid



mesh, generating 3 special k -points in the Monkhorst-Pack scheme. For the computation of integrals and the representation of charge densities and potentials, mesh cut-off energy is kept at 300 Ry so as to depict the periodic plane waves in the grid without aliasing. To guarantee better contribution from dopant atoms and save computational time, we have considered a square-size supercell with 32 atoms. In order to prevent interaction between the neighbouring layers, a vacuum space of 20 Å thickness is maintained in the perpendicular direction of the monolayer. Upon optimization, water molecule is introduced and the adsorption energy of water molecule on these structures are computed. To measure the extent of water activation on the phosphorene sheets, the O–H stretching frequency of the adsorbed water molecule is measured using 'vibra', a post-processing tool of SIESTA. The AIMD simulations at 500 K, a Nosé–Hoover thermostat has been implemented and the n S-B-phosphorene systems are described with NVT (number, volume, temperature) ensemble.

Results and discussion

Structure, stability and electronic properties of n S-B-phosphorene sheets ($n = 0$ to 6)

B-phosphorene has a hexagonal primitive unit cell with two atoms per cell¹⁰⁹ as shown Fig. 1(b). We obtain an optimized lattice constant ($a = b$) of 3.27 Å and P–P bond length of 2.28 Å and P–P–P bond angle of 91.6°, which are in satisfactory agreement with previously reported calculations.¹¹⁰ The buckling length (h) of monolayer is found to be 1.24 Å, which is in good accordance with previous report.¹¹¹ The atoms in the sheet have been further classified as the central (C) and edge (E) atoms as discussed earlier are shown in Fig. 1(a). Accordingly, there are 10 central atoms and 22 edge atoms. In order to initially screen the influence of sulphur concentration on the stability and electronic effects of B-phosphorene, we have replaced phosphorous atoms with sulphur atoms only at the central (C) positions. Concentration of sulphur was varied in this sheet from 3.125% which translates to 1S atom in the sheet. As the concentration of S atom was increased beyond 18.75% the structure was seen to collapse. So, further structural studies are restricted to 18.75% which translates to doping of 1–6 P atoms with S in above sheet. Thus, there are 10 distinct conformations generated for single S atom doped B-phosphorene and are referred to as 1S-B-phosphorene conformations. Each of this conformation was optimized. The six lowest energy 1S-B-phosphorene conformations with their relative energies are shown in ESI Fig. S.F.1.[†] Next, the number of dopant atoms (S atoms) is increased to 2. The first S atom was placed at the 10 distinct C positions, followed by 9 distinct C positions for the second atom. Thus, totally 10×9 distinct conformations were generated. The six lowest energy 2S-B-phosphorene conformations are given in ESI Fig. S.F.2.[†] Similarly, the number of conformations generated for 3S doped B-phosphorene sheet are $10 \times 9 \times 8$ and so on. The six lowest energy conformation for 3S, 4S, 5S and 6S doped B-phosphorene are shown in ESI Fig. S.F.3–S.F.6,[†] respectively. In case of doped structures with more than 1S atom, it was

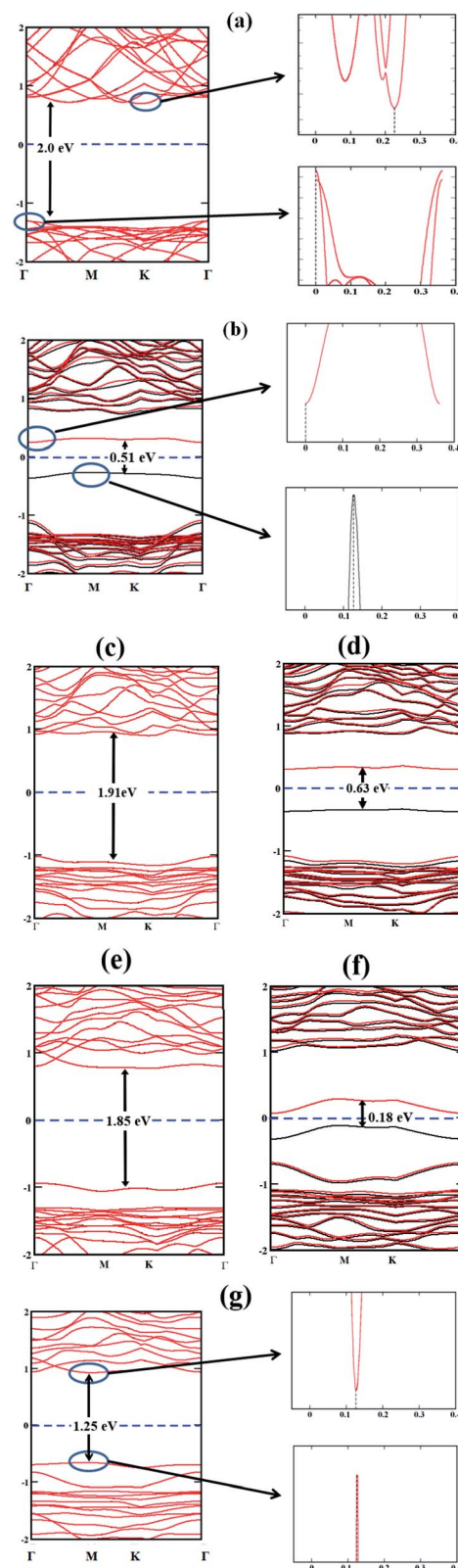


Fig. 2 Band structures of (a) B-phosphorene; (b) 1S-B-phosphorene (c) 2S-B-phosphorene (d) 3S-B-phosphorene (e) 4S-B-phosphorene (f) 5S-B-phosphorene (g) 6S-B-phosphorene with their respective band gaps indicated with vertical lines. The position of the band edges is indicated by the sharp band lines as shown for pristine B-phosphorene, 1S-B-phosphorene and 6S-B-phosphorene.



noted that defects are formed around the S atoms in the sheet, on account of its lower valency, higher electronegativity and smaller atomic radius as compared to that of P (see ESI Fig. S.F.1–S.F.6,† respectively). This defect formation was also noted in the previous work where blue phosphorene nano-ribbons were doped with S atoms.¹¹² These defects are anticipated to play a vital role in enhancing photocatalysis by averting the recombination of photogenerated charges.¹¹³

In order to evaluate the potential of a material as a photocatalyst, it is instructive to probe its band structure. Therefore, we analyse the electronic properties of the lowest energy conformation for each *n*S-B-phosphorene (*n* = 0–6). Fig. 2 shows the spin resolved band structure for B-phosphorene and *n*S-B-phosphorene sheets (*n* = 0, 1, 2, 3, 4, 5, 6). Spin-up (black-lines) and spin-down (red-lines) energy levels are highlighted and follow the representation adopted by Zhu *et al.*¹¹⁴ It is noted that the first Brillouin zone of hexagonal B-phosphorene is projected between the conduction band minimum and valence band maximum through the high symmetric path Γ –*M*–*K*– Γ (see Fig. 2(a)). Our calculations further reveal an indirect band gap of 2 eV for the B-phosphorene monolayer, which is also substantiated by other reported data.^{115,116} Coming to the discussion on the type of band gap pertinent in the context of visible light driven catalysis, the effective participation of the photogenerated electrons and holes is determined by recombination rate of these pairs. On recombination there is liberation of energy in the form of photon with the excited electrons needing to fulfil the following transition selection rules of momentum conservation¹¹⁷

$$E_g = \hbar\omega_{\text{photon}} \quad (1)$$

$$\hbar k_{\text{ev}} - \hbar k_{\text{ec}} = \pm \hbar q_{\text{phonon}} \quad (2)$$

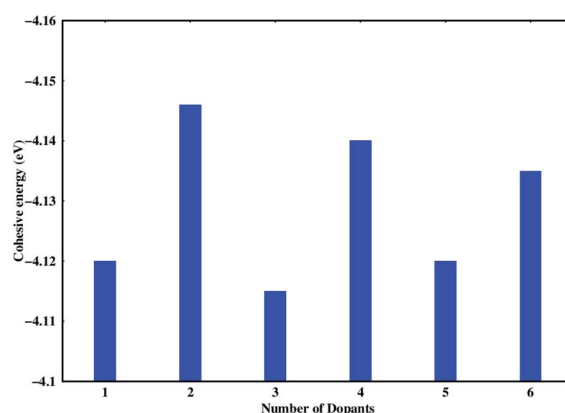
where \hbar is the reduced Planck constant, k_{ev} and k_{ec} are the electron wave vectors at the VBM and CBM, respectively, q_{phonon} is the wave vector of the assisted phonon, E_g is the band gap of semiconductor, and ω_{photon} is angular frequency of the emitted photon. It is seen in case of pristine B-phosphorene, that $k_{\text{ev}} \neq k_{\text{ec}}$ with an indirect band gap characteristic of semiconductors. Thus, the excited electrons cannot recombine directly and easily with holes, resulting in an increase of the photoexcited electron–hole lifetime, increase in diffusion length and time. Thus, B-phosphorene has better photocatalytic activity than direct band gap semiconductors as reported in a previous study.¹¹⁸ Analysis of the band structures of the lowest energy *n*S-B-phosphorene (*n* = 1–6) reveals a dissimilarity in valence orbital occupancy of dopant S with respect to phosphorous ($3s^23p^4$ as compared $3s^23p^3$), bringing in distinct changes in the electronic structure of doped sheets with respect to their pristine counterpart. *n*S-B-phosphorene (*n* = 1, 2, 3, 4, 5) sheets demonstrate n-type semiconducting characteristics with Conduction Band Minimum (CBM) lying nearer to Fermi level as compared to Valence Band Maximum (VBM). On the other hand, 6S-B-phosphorene is found to have a direct band gap as seen from Fig. 2(g). The band gap results for pristine and doped B-phosphorene are quantified in Table 1, from which clearly, we

Table 1 Band gap energy of pristine and various *n*S-B-phosphorene

System	Band gap energy (eV)
Pristine phosphorene	2.00
1S-B-phosphorene	0.51
2S-B-phosphorene	1.91
3S-B-phosphorene	0.63
4S-B-phosphorene	1.85
5S-B-phosphorene	0.18
6S-B-phosphorene	1.25

can note that while the pristine and even number S doped (*n* = 2, 4, 6) systems have a gap which is higher than the electrolytic potential of water *viz.*, 1.23 eV, the odd number S doped systems (*n* = 1, 3, 5) fails to meet this essential criterion. The even *n*S-doped systems with band gaps above 1.23 eV are potential photocatalyst for water splitting and will be explored further. The odd *n*S-doped B-phosphorene systems possess a smaller band gap and would inherently show higher carrier mobility and electrical conductivity. However, the adsorption efficiency of H^+ (dissociated from water) decreases eventually leading to lower solar energy to hydrogen evolution efficiency, thereby reducing the photocatalytic efficiency for HER.

Stability of the generated catalytic systems are studied in terms of their cohesive energies and it is calculated from the total electronic energies.^{119–121} The cohesive energies of different systems are shown in Fig. 3. Evaluation of cohesive energies suggest that even doped *n*S-B-phosphorene systems have higher cohesive energies with nearly difference of 0.02 eV than their odd counterparts. This results in better thermal stability of *n*S-B-phosphorene (*n* = 2, 4, 6) sheets indicating that a particular concentration of dopants is more favourable (or number of dopant) *viz.* 6.25%, 12.5% and 18.75%. Based on the cohesive energy results and band gap energy results, we have further restricted our studies to the even doped *n*S-B-phosphorene (*n* = 2, 4, 6) sheets. In these dopant concentrations, the most favourable dopant substitution sites are more rigorously

Fig. 3 Cohesive energies of various studied *n*S-B-phosphorene systems.

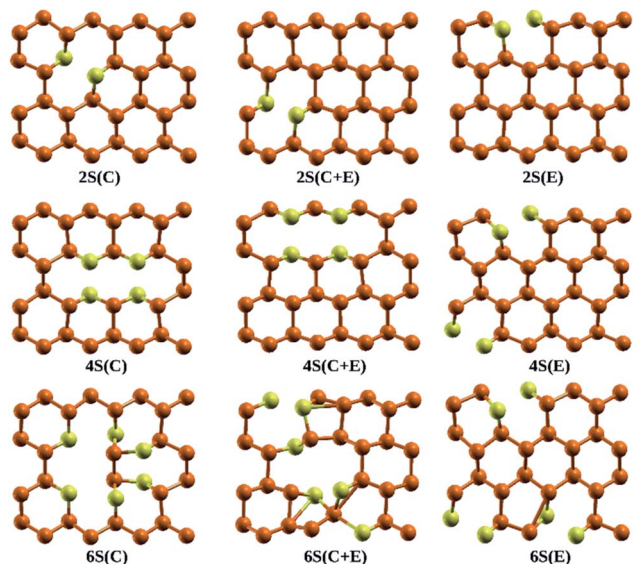


Fig. 4 Lowest energy conformations of S atom occupation in 2S-B-phosphorene, 4S-B-phosphorene, 6S-B-phosphorene, respectively.

explored taking central (C), edge (E) and central & edge (C + E) sites into account. The three sites of S-atom occupation are described as follows; (i) all S atoms occupy central sites of Fig. 1(a) as elaborated in Fig. 4 and this is referred to as *n*S-B-phosphorene (C). (ii) The S atoms are distributed among various edge and central sites of Fig. 1(a), these conformations are referred to as *n*S-B-phosphorene (C + E) and representative structure of this is shown in Fig. 4. (iii) All S atoms occupy edge sites of Fig. 1(a) as shown in Fig. 4, and is referred to as *n*S-B-phosphorene (E).

As already discussed, for a material to be an efficient photocatalyst for water molecule activation, band gap must be at least 1.23 eV. Besides, the Conduction Band Minimum (CBM) should be more negative than the redox potential of H^+/H_2 (0 V vs. NHE) and the Valence Band Maximum (VBM) should be more positive than the redox potential of $\text{O}_2/\text{H}_2\text{O}$.¹²⁰ A material will act as a catalyst for OER when its VBM is more positive than the redox potential of $\text{O}_2/\text{H}_2\text{O}$ whereas, it will act as a catalyst for HER when its CBM is more negative than the redox potential of H^+/H_2 . The water redox potential energies are also reported to be dependent on the pH of the medium.¹²⁰ To analyse the potential of the studied materials we first look at the redox potential of water molecule at 0 and 7 pH values using the below standard equation.^{122–124}

$$\text{EO}_2/\text{H}_2\text{O} = -5.67 \text{ eV} + \text{pH} \times 0.059 \text{ eV} \quad (3)$$

$$\text{EH}^+/\text{H}_2 = -4.44 \text{ eV} + \text{pH} \times 0.059 \text{ eV} \quad (4)$$

It is known that the reduction potential of H^+/H_2 is 0.00 V at pH 0 vs. NHE, however -0.41 V over-potential is required for the evolution of hydrogen in aqueous solutions at pH 7 vs. NHE. Similarly, potential for water oxidation $\text{O}_2/\text{H}_2\text{O}$ is at $+1.23$ V vs. NHE at pH 0 (or $+0.82$ V vs. NHE at pH 7).¹²⁵ This, reflects as to why a band gap of 1.23 eV is required for a material to drive overall water splitting reaction. To calculate the band edge position at pH = 7 of the medium, the values of CBM and VBM are computed from the data obtained. The Conduction Band (CB) potential of a semiconducting material in aqueous solution usually exhibits a pH dependence explained by $E_{\text{CB}} = E_{\text{CB}}^0 (\text{pH } 0) - 0.059 \times \text{pH}$; where E_{CB}^0 is the standard reduction potential.^{126,127} Further, the location of the VBM should be more

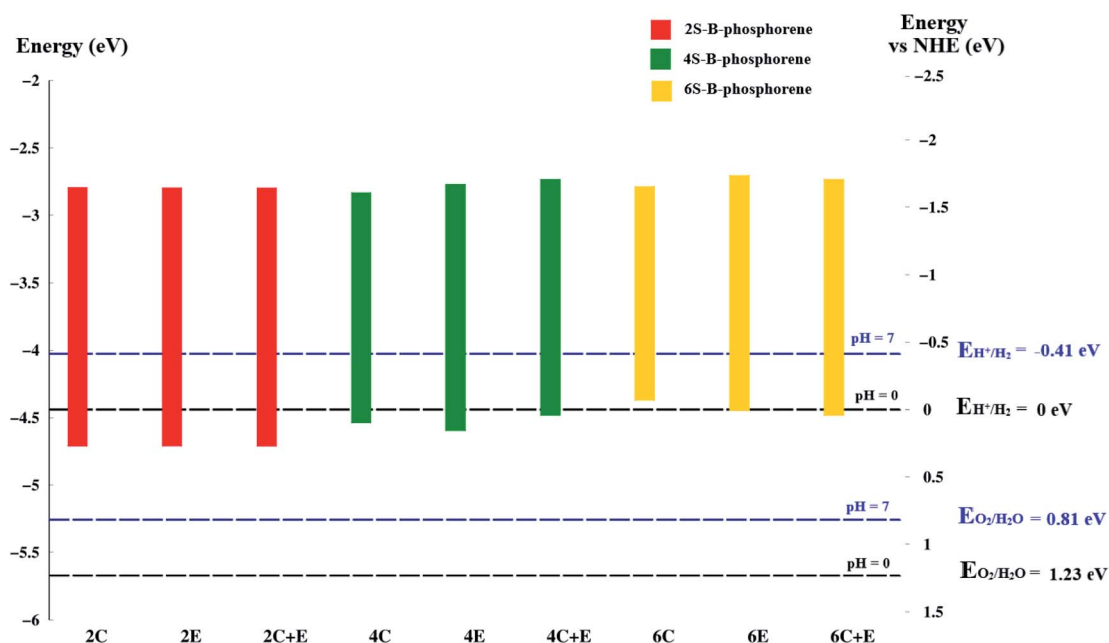


Fig. 5 Band edge positions of *n*S-B-phosphorene (*n* = 2, 4, 6), at (C), (E), (C + E) locations in vacuum energy and NHE energy scale. The horizontal black dashed line is the redox potential of water splitting at pH = 0 and the blue dashed line is the redox potential of water splitting at pH = 7.



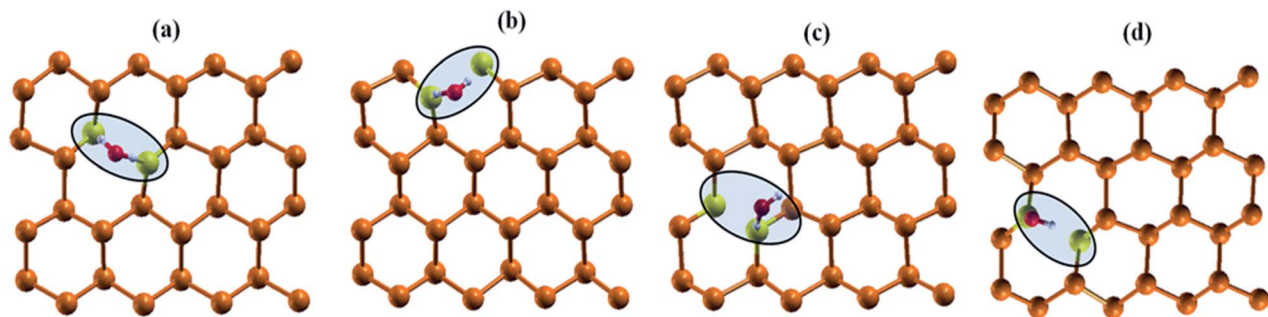


Fig. 6 Interaction of water molecule at various dopant sites (a) *n*S-B-phosphorene (C) (b) *n*S-B-phosphorene (E) (c) *n*S-B-phosphorene (C+E) (d) *n*S-B-phosphorene (C+E). The highlighted and red coloured letter in C + E corresponds to the site on which water molecule is adsorbed.

positive than +0.82 V while the position of the CBM should be more negative than −0.41 V vs. NHE at pH 7, respectively.

Coming to the present work, we plot the band edge locations for the lowest energy *n*S-B-phosphorene (*n* = 2, 4, 6) where dopants positions are *n*S-B-phosphorene (C), *n*S-B-phosphorene (E) and *n*S-B-phosphorene (C + E), as shown in Fig. 5 with the redox potentials of hydrogen evolution (H^+/H_2) and oxygen evolution (O_2/H_2O) at pH 0 and 7. The band width of all the systems surpasses the required band gap of 1.23 eV and are favourable for photocatalytic water splitting at acidic (pH = 0) as well as neutral medium (pH = 7) without any application of an external potential. The redox potential of oxygen evolution (O_2/H_2O) is found to be more positive than the VBM of all *n*S-B-phosphorene systems, see Fig. 5. Thus, these systems will fail to

catalyse the oxygen evolution reaction (OER). However, the CBM is seen to be more negative than the redox potential of hydrogen evolution (H^+/H_2) at both pH values; thereby all *n*S-B-phosphorene systems can behave as efficient catalysts for HER. An acidic medium can lead to etching or degradation of the material; therefore, a neutral medium is always preferred for carrying out experiments. At neutral medium, *i.e.*, pH = 7, all 2S-B-phosphorene structures show an overpotential of 1.24 eV for HER in all three sites (*i.e.* C, E and C + E). 4S-B-phosphorene shows varying overpotentials ranging from 1.2 eV in C-site, 1.26 eV in E-site and 1.3 eV in C + E site; while the overpotentials in 6S-B-phosphorene sites range from 1.24 eV in C-site, 1.29 eV in E-site to 1.32 eV in C + E site. The *n*S-B-phosphorene (C) systems show a lower overpotential

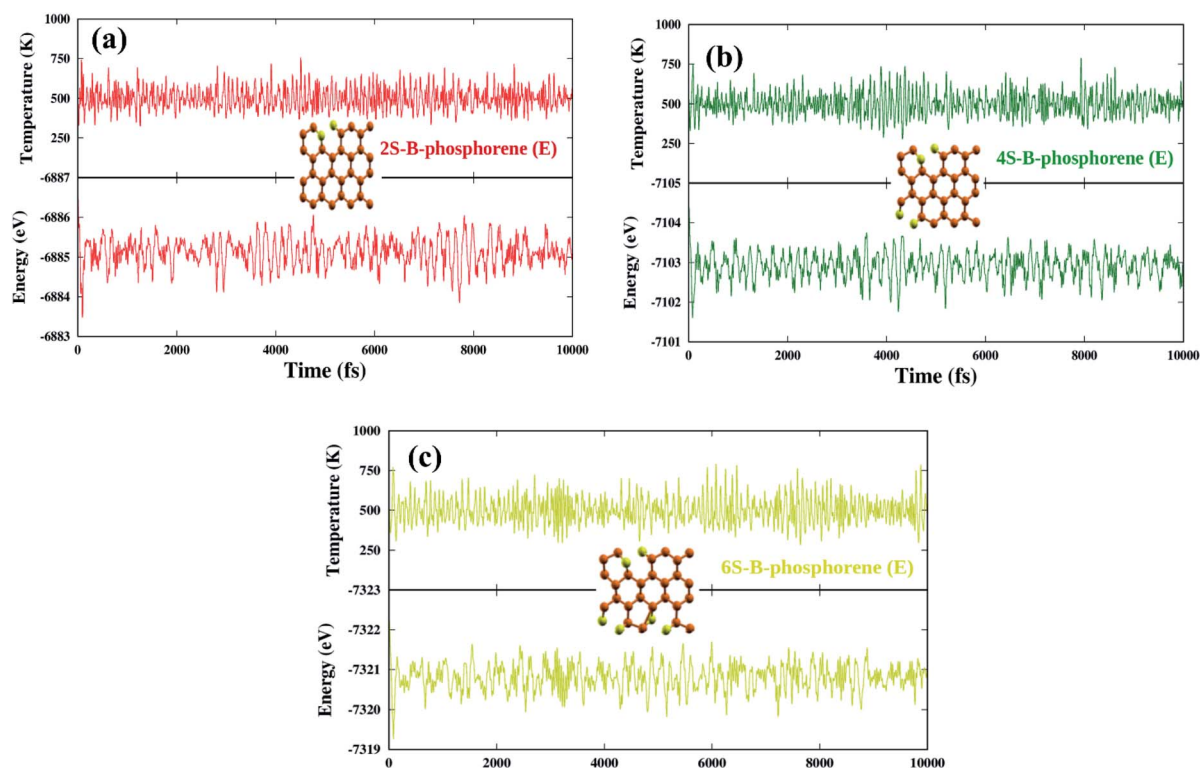


Fig. 7 AIMD simulations of (a) 2S-B-phosphorene (E) (b) 4S-B-phosphorene (E) and (c) 6S-B-phosphorene systems for 10 ps at 500 K. Relaxed geometries of *n*S-B-phosphorene (E) systems are as shown inset.



compared to systems with dopants at other sites. Also, these *n*S-B-phosphorene (C) systems with lower overpotentials also possess a lower band gap compared to *n*S-B-phosphorene (E) or *n*S-B-phosphorene (C + E) systems. Therefore, all *n*S-B-phosphorene (C) systems will show higher photocatalytic activity towards HER; in particular, the 4S-B-phosphorene (C) with overpotential of 1.2 eV will show highest activity at pH = 7. Interestingly, the even *n*S-B-phosphorene systems are also found to show high thermostability without any distortion in their geometry at temperature as high as 500 K. AIMD simulations of the *n*S-B-phosphorene (E) systems carried out for 10 ps and the geometries generated for every 1 fs is as shown in Fig. 7. Considering the band edge alignment at ambient pH condition and the thermostability at 500 K, the *n*S-B-phosphorene (E) systems potential photocatalysts that can be pragmatically implemented at room temperature and neutral pH.

Activation of water molecule on *n*S-B-phosphorene sheet (*n* = 2, 4, 6)

The electronic properties of *n*S-B-phosphorene (*m*), *n* = 2, 4, 6 and *m* = C, E, C + E is further corroborated by explicit adsorption and activation of water molecule on the sheet. The water

molecule in all cases is adsorbed on the dopant site, *viz.*; S atom. The activation is analysed in terms of adsorption energy of water molecule and activation of O–H bond through analysis of its red shift and O–H bond elongation. Adsorption energy is indicative of chemical interaction the *n*S-B-phosphorene and the water molecules and is calculated by using the following equation¹²⁸

$$E_{\text{ads}} = E_{\text{water+system}} - (E_{\text{water}} + E_{\text{system}}) \quad (5)$$

In the above equation, E_{ads} is the adsorption energy, $E_{\text{water+system}}$ is the electronic energy of water molecule adsorbed B-phosphorene, E_{water} is the electronic energy of a water molecule and E_{system} is the electronic energy of B-phosphorene sheet.

As mentioned earlier, water molecule is adsorbed on the dopant sites (S atom) of lowest energy conformation of *n*S-B-phosphorene (C), *n*S-B-phosphorene (E) and *n*S-B-phosphorene (C + E), *n* = 2, 4, 6. In case of *n*S-B-phosphorene (C + E), water molecule can be adsorbed on centrally placed S atom or edge S atom giving rise to *n*S-B-phosphorene (C+E) and *n*S-B-phosphorene (C+E), respectively (see Fig. 6(c) and (d)). The adsorption energies, OH stretching frequencies of the activated OH bond and activated OH bond length is given in Fig. 8. The

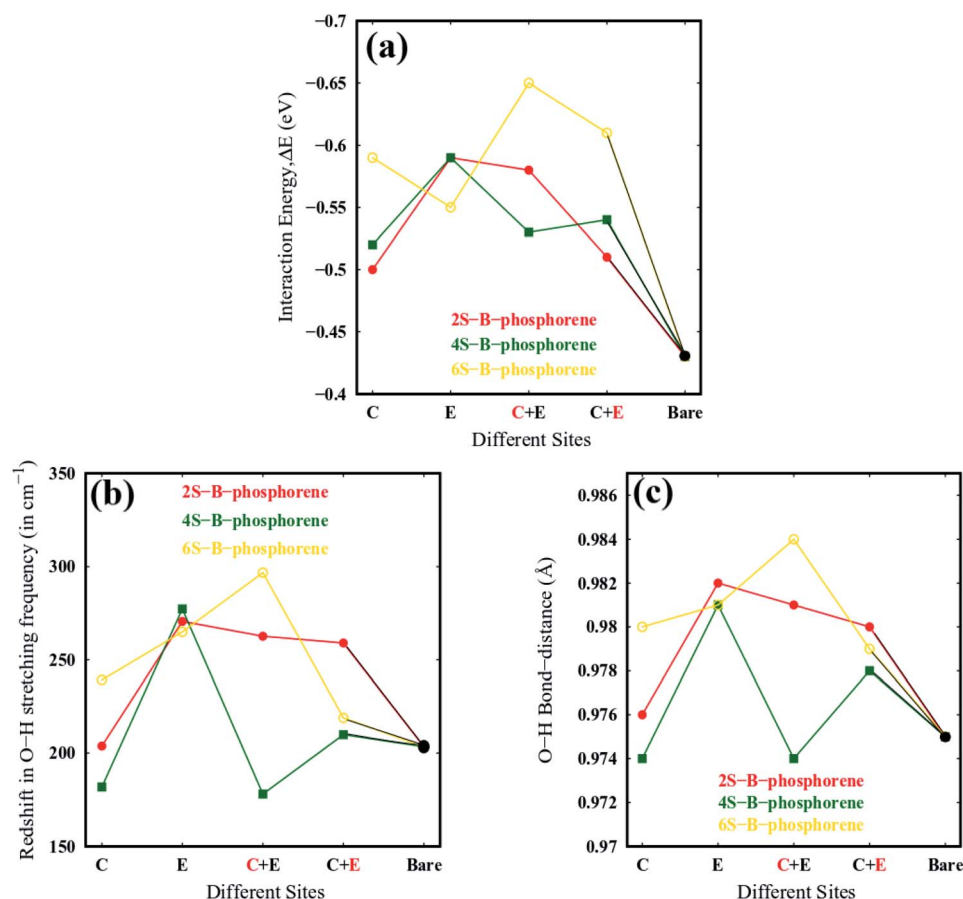


Fig. 8 (a) Water molecule adsorption energies, magnitude of the E_{ad} values are plotted (b) O–H stretching frequency of the activated bond in the water molecule, (c) activated O–H bond length of water molecule on various *n*S-B-phosphorene (*n* = 2, 4, 6). C, E, C + E on the x axis represents the doping sites of S atom *viz.*; namely, centre, edge, centre + edge. The highlighted and red coloured letter in C + E corresponds to the site on which water molecule is adsorbed.



absolute values for the same are given in ESI Table S.T 2.† The adsorption energy of water over B-phosphorene is noted to be 0.43 eV. All doped structures have higher interaction energies with the water. In particular, 6S-B-phosphorene (C+E) shows a higher adsorption energy (0.65 eV) in comparison to other *n*S-B-phosphorene structures. As we know experimentally, water splitting results in formation of adsorbed hydroxyl ions and hydrogen, respectively, on the catalyst interface. The obtained adsorption energy trends are further evaluated in terms of O–H bond elongation and red shift in O–H stretching frequencies. Red shifts in O–H stretching frequencies are obtained by taking the difference between O–H stretching frequency of free water molecule and that of water adsorbed on *n*S-B-phosphorene sheets; the values are as shown in Fig. 8(b). This shift in O–H stretching frequency is taken as a measure of O–H bond activation and the highest O–H bond activation is seen in the case of 6S-B-phosphorene (C+E) system with a red shift of 296.82 cm^{−1}. This is followed by *n*S-B-phosphorene (E) systems with their red shifts approximately at 270 cm^{−1}. The observed red shifts in O–H stretching frequencies are also corroborated through the analysis of the activated O–H bond lengths as shown in Fig. 8(c). O–H bond length in free water moiety is 0.96 Å however, the O–H bond length for 6S-B-phosphorene corresponds to 0.985 Å indicating that it gets activated upon adsorption on *n*S-B-phosphorene sheets resulting in higher interaction energy and maximum red shift in O–H stretching frequency.

While the 6S-B-phosphorene system shows a higher activation of water molecule when compared to rest of the systems; interestingly, all *n*S-B-phosphorene (E) systems show a relatively efficient activation of water molecule with the red shift in O–H red stretching frequency approximately at 270 cm^{−1}. Also, in several other 2D materials the edge sites have been reported to show higher catalytic activity when compared to the basal plane and our results are in correspondence to several such results. The overpotentials required for photocatalytic HER in these *n*S-B-phosphorene (E) systems range from 1.24 eV to 1.29 eV from our previous discussion on band edge alignment. When compared to the HER overpotential of 1.2 eV in 4S-B-phosphorene, the HER overpotentials in *n*S-B-phosphorene (E) systems are slightly higher but the chemically active edge sites available on these systems are an advantage when it comes to photocatalytic water splitting. These active sites bridge the energy required to activate the water molecule which the incident photons from solar light photons cannot provide sufficiently. Therefore, the *n*S-B-phosphorene (E) systems with catalytically active edge sites and HER overpotentials of 1.24–1.29 eV are efficient photocatalysts that can be experimentally explored.

Conclusions

In this work, the photocatalytic reactivity of B-phosphorene and chemically modified B-phosphorene monolayer is analysed and explored in terms of band structure, S-dopant concentration at best probable atomistic positions, band alignment with respect to pH. Cohesive energy demonstrates a higher stability of even

*n*S-doped B-phosphorene monolayer at specific S-dopant concentration, *i.e.*; 6.25%, 12.5% and 18.75%. Further, band alignment studies suggest that even doped *n*S-B-phosphorene (*n* = 2, 4, 6) fulfils the band gap criteria and indirect band gap with an exception of 6S-B-phosphorene. Moreover, all *n*S-B-phosphorene (at all doping sites), respectively meet the redox potential level of H₂ evolution only at pH = 0 and 7, therefore they will behave as efficient photocatalysts for HER. Aside from an overpotential of ~1.24–1.29 eV for hydrogen evolution; these even *n*S-B-phosphorene systems show high feasibility to adsorb water molecule and activate the O–H bond and show high thermal stability at 500 K. The chemical activity of these systems are higher on the 6S-B-phosphorene (C + E) system followed by the *n*S-B-phosphorene (E) systems. Taking the photochemical activity *i.e.*; overpotentials required for HER at neutral medium, thermostability at 500 K as well as chemical activity towards water activation, the *n*S-B-phosphorene (E) systems are found to be effective photocatalysts. This computational investigation can be used to design and improvise phosphorene photocatalyst experimentally for water splitting.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 C. X. Lin, R. Li, M. Lu, C. Chen, D. Li, Y. Zhang and L. Jiang, *Fuel*, 2015, **162**, 271–280.
- 2 M. M. J. Li, C. Chin, T. Ayval, H. Suo, J. Zheng, I. F. Teixeira, L. Ye, H. Zou, D. Here and S. C. E. Tsang, *ACS Catal.*, 2018, **8**, 4390–4401.
- 3 S. Iguchi, K. Teramura, S. Hosokawa and T. Tanaka, *Phys. Chem. Chem. Phys.*, 2015, **17**, 17995–18003.
- 4 Z. Zhao, L. Zhang and Z. Xia, *J. Phys. Chem. C*, 2016, **120**(4), 2166–2175.
- 5 T. Oshima, T. Ichibha, K. S. Qin, K. Muraoka, J. J. M. Vequizo, K. Hibino, R. Kuriki, S. Yamashita, K. Hongo, T. Uchiyama, K. Fujii, D. Lu, R. Maezono, A. Yamakata, H. Kato, K. Kimoto, M. Yashima, Y. Uchimoto, M. Kakihana, O. Ishitani, H. Kageyama and K. Maeda, *Angew. Chem., Int. Ed.*, 2018, **57**, 8154–8158.
- 6 Q. Shao, P. Wang, T. Zhu and X. Huang, *Acc. Chem. Res.*, 2019, **52**, 3384–3396.
- 7 H. K. Chae, D. Y. Siberio-Pérez, J. Kim, Y. Go, M. Eddaoudi, A. J. Matzger, M. O'Keeffe and O. M. Yaghi, *Nature*, 2004, **427**, 523–527.
- 8 I. W. Frank, D. M. Tanenbaum, A. M. Van der Zande and P. L. McEuen, *J. Vac. Sci. Technol., B: Microelectron.*



- Nanometer Struct.–Process., Meas., Phenom.*, 2007, **25**, 2558–2561.
- 9 Y. Fu, J. Hansson, Y. Liu, S. Chen, A. Zehri, M. K. Samani, N. Wang, Y. Ni, Y. Zhang, Z. B. Zhang, Q. Wang, M. Li, H. Lu, M. Sledzinska, C. M. S. Torres, S. Volz, A. A. Balandin, X. Xu and J. Liu, *2D Mater*, 2020, 7(012001), 1–42.
 - 10 L. Wang, W. Chen, D. Zhang, Y. Du, R. Amal, S. Qiao, J. Wu and Z. Yin, *Chem. Soc. Rev.*, 2019, **48**, 5310–5349.
 - 11 R. Sahoo, A. Pal and T. Pal, *Chem. Commun.*, 2016, **52**, 13528–13542.
 - 12 B. Luo, G. Liu and L. Wang, *Nanoscale*, 2016, 6904–6920.
 - 13 D. Deng, K. S. Novoselov, Q. Fu, N. Zheng, Z. Tian and X. Bao, *Nat. Nanotechnol.*, 2016, **11**, 218–230.
 - 14 D. Dreyer, P. Jia and C. Bielawski, *Angew. Chem., Int. Ed. Engl.*, 2010, **49**, 6813–6816.
 - 15 H. Jia, D. Dreyer and C. Bielawski, *Tetrahedron*, 2011, **67**, 4431–4434.
 - 16 Z. H. Sheng, H. L. Gao, W. J. Bao, F. B. Wang and X. H. Xia, *J. Mater. Chem.*, 2012, **22**, 390–395.
 - 17 M. Burghard, H. Klauk and K. Kern, *Adv. Mater.*, 2009, **21**, 2586–2600.
 - 18 D. R. Dreyer, R. S. Ruoff and C. W. Bielawski, *Angew. Chem., Int. Ed.*, 2010, **49**, 9336–9344.
 - 19 J. Albero, D. Mateo and H. García, *Molecules*, 2019, **24**(906), 1–21.
 - 20 T. F. Yeh, J. M. Syu, C. Cheng, T. H. Chang and H. Teng, *Adv. Funct. Mater.*, 2010, **20**, 2255–2262.
 - 21 L. Clarizia, D. Russo, I. D. Somma, R. Andreozzi and R. Marotta, *Energies*, 2017, **10**(1624), 1–21.
 - 22 P. Vogt, P. D. Padova, C. Quaresima, J. Avila, E. Frantzeskakis, M. C. Asensio, A. Resta, B. Ealet and G. L. Lay, *Phys. Rev. Lett.*, 2012, **108**, 155501–155505.
 - 23 A. Acun, L. Zhang, P. Bampoulis, M. Farmanbar, A. Houselt, A. N. Rudenko, M. Lingensfelder, G. Brocks, B. Poelsema, M. I. Katsnelson and H. J. W. Zandvliet, *J. Phys.: Condens. Matter*, 2015, **27**, 443002–443012.
 - 24 L. Shi, C. Ling, Y. Ouyang and J. Wang, *Nanoscale*, 2017, **8**, 533–537.
 - 25 S. H. Mir, S. Chakraborty, P. C. Jha, J. Wärmå, H. Soni, P. K. Jha and R. Ahuja, *Appl. Phys. Lett.*, 2016, **109**, 53903–53907.
 - 26 P. Garg, I. Choudhuri, A. Mahata and B. Pathak, *Phys. Chem. Chem. Phys.*, 2017, **19**, 3660–3669.
 - 27 H. Tada, T. Mitsui, T. Kiyonaga, T. Akita and K. Tanaka, *Nat. Mater.*, 2006, **5**, 782–786.
 - 28 C. J. Rupp, S. Chakraborty, J. Anversa, R. J. Baierle and R. Ahuja, *ACS Appl. Mater. Interfaces*, 2016, **8**, 1536–1544.
 - 29 C. Chowdhury, S. Karmakar and A. Datta, *J. Phys. Chem. C*, 2017, **121**, 7615–7624.
 - 30 R. Jain, R. Narayan, S. P. Sasikala, K. E. Lee, H. J. Jung and S. O. Kim, *2D Mater*, 2017, **4**, 042006–042089.
 - 31 H. Liu, A. T. Neal, Z. Zhu, X. Xu, D. Tomanek, P. D. Ye and Z. Luo, *ACS Nano*, 2014, **8**, 4033–4041.
 - 32 V. Tran, R. Soklaski, Y. Liang and L. Yang, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2014, **89**, 235319.
 - 33 L. Li, J. Kim, C. Jin, G. J. Ye, D. Y. Qiu, F. H. D. Jornada, Z. Shi, L. Chen, Z. Zhang, F. Yang, K. Watanabe, T. Taniguchi, W. Ren, S. G. Louie, X. H. Chen, Y. Zhang and F. Wang, *Nat. Nanotechnol.*, 2017, **12**(1), 21–25.
 - 34 S. Larentis, B. Fallahazad and E. Tutuc, *Appl. Phys. Lett.*, 2012, **101**, 223104.
 - 35 H. Liu, A. T. Neal, Z. Zhu, D. Tomanek and P. D. Ye, *ACS Nano*, 2014, **8**, 4033–4041.
 - 36 Y. L. Lu, S. Dong, J. Li, Y. Wu, L. Wang and H. Zhao, *Phys. Chem. Chem. Phys.*, 2020, **22**, 13713–13720.
 - 37 K. Zhang, B. Jin, C. Park, Y. Cho, X. Song, X. Shi, S. Zhang, W. Kim, H. Zeng and J. H. Park, *Nat. Commun.*, 2019, **10**, 1–10.
 - 38 L. Zhang, H. Huang, B. Zhang, M. Gu, D. Zhao, X. Zhao, L. Li and Y. Cheng, *Angew. Chem., Int. Ed.*, 2020, **132**(3), 1090–1096.
 - 39 T. Kaewmaraya, L. Ngamwongwan, P. Moontragoon, W. Jareboon, D. Singh, R. Ahuja, A. Karton and T. Hussain, *J. Hazard. Mater.*, 2021, **401**, 123340.
 - 40 M. Rahman, C. Kwong, K. Davey and S. Qiao, *Energy Environ. Sci.*, 2016, **9**, 709–728.
 - 41 S. Kaur, A. Kumar, S. Srivastava, K. Tankeshwar and R. Pandey, *J. Phys. Chem. C*, 2018, **122**, 26032–26038.
 - 42 N. Jiao, P. Zhou, L. Xue, C. He and L. Sun, *J. Phys.: Condens. Matter*, 2019, **31**, 075702.
 - 43 Y. Lu, S. Dong, W. Zhou, S. Dai, B. Zhou, H. Zhao and P. Wu, *Phys. Chem. Chem. Phys.*, 2018, **20**, 11967–11975.
 - 44 J. L. Zhang, S. Zhao, C. Han, Z. Wang, S. Zhong, S. Sun, R. Guo, X. Zhou, C. D. Gu and K. D. Yuan, *Nano Lett.*, 2016, **16**, 4903–4908.
 - 45 J. Zeng, P. Cui and Z. Zhang, *Phys. Rev. Lett.*, 2017, **118**, 046101.
 - 46 Z. Zhu and D. Tomanek, *Phys. Rev. Lett.*, 2014, **112**, 176802.
 - 47 J. Li, X. Sun, C. Xu, X. Zhang, Y. Pan, M. Ye, Z. Song, R. Quhe, Y. Wang, H. Zhang, Y. Guo, J. Yang, F. Pan and J. Lu, *Nano Res.*, 2018, **11**, 1834–1849.
 - 48 E. Montes and U. Schwingenschlög, *J. Mater. Chem. C*, 2017, **5**, 5365–5371.
 - 49 F. Safari, M. Moradinasab, M. Fathipour and H. Kosina, *Appl. Surf. Sci.*, 2018, **464**, 153–161.
 - 50 L. Ju, Y. Dai, W. Wei, Y. Liang and B. Huang, *J. Mater. Chem. A*, 2018, **6**, 21087.
 - 51 B. Wang, X. Li, X. Cai, W. Yu, L. Zhang, R. Zhao and S. Ke, *J. Phys. Chem. C*, 2018, **122**, 7075–7080.
 - 52 Y. Cheng, Y. Song and Y. Zhang, *Phys. Chem. Chem. Phys.*, 2019, **21**, 24449.
 - 53 C. Li, Y. Xu, W. Sheng, W. Yin, G. Nie and Z. Ao, *Phys. Chem. Chem. Phys.*, 2020, **22**, 615.
 - 54 B. Wang, X. Li, R. Zhao, X. Cai, W. Yu, W. Li, Z. Liu, L. Zhang and S. Ke, *J. Mater. Chem. A*, 2018, **6**, 8923.
 - 55 B. Ghosh, S. Nahas, S. Bhowmick and A. Agarwal, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2015, **91**, 115433.
 - 56 Y. Mogulkoc, M. Modarresi, A. Mogulkoc and Y. O. Ciftci, *Comput. Mater. Sci.*, 2016, **124**, 23–29.
 - 57 R. B. Pontes, R. H. Miwa, A. J. R. da Silva, A. Fazzio and J. E. Padilha, *Phys. Rev. B*, 2018, **97**, 235419.



- 58 L. Zhu, S. Wang, S. Guan, Y. Liu, T. Zhang, G. Chen and S. A. Yang, *Nano Lett.*, 2016, **16**, 6548–6554.
- 59 G. Yang, Z. Xu, Z. Liu, S. Jin, H. Zhang and Z. Ding, *J. Phys. Chem. C*, 2017, **121**(23), 12945–12952.
- 60 M. Sun, S. Wang, J. Yu and W. Tang, *Appl. Surf. Sci.*, 2017, **392**, 46–50.
- 61 M. Sun, W. Tang, Q. Ren, S.-K. Wang, J. Yu and Y. Du, *Appl. Surf. Sci.*, 2015, **356**, 110–114.
- 62 M. Sun, Y. Hao, Q. Ren, Y. Zhao, Y. Du and W. Tang, *Solid State Commun.*, 2016, **242**, 36–40.
- 63 W. X. Zhang, J. W. Zhao, W. H. He, L. J. Luan and C. He, *Chem. Phys. Lett.*, 2017, **675**, 20–26.
- 64 H. Schmidt, F. Giustiniano and G. Eda, *Chem. Soc. Rev.*, 2015, **44**, 7715–7736.
- 65 Y. D. Zhao, K. Xu, F. Pan, C. J. Zhou, F. C. Zhou and Y. Chai, *Adv. Funct. Mater.*, 2017, **27**, 1603484.
- 66 F. Wang, Z. X. Wang, C. Jiang, L. Yin, R. Q. Cheng, X. Y. Zhan, K. Xu, F. M. Wang, Y. Zhang and J. He, *Small*, 2017, **13**, 1604298.
- 67 B. Li, T. Xing, M. Z. Zhong, L. Huang, N. Lei, J. Zhang, J. B. Li and Z. M. A. Wei, *Nat. Commun.*, 2017, **8**, 1958.
- 68 V. Kochat, A. Apte, J. A. Hachtel, H. Kumazoe, A. Krishnamoorthy, S. Susarla, J. C. Idrobo, F. Shimojo, P. Vashishta, R. Kalia, A. Nakano, C. S. Tiwary and P. M. Ajayan, *Adv. Mater.*, 2017, **29**, 1703754.
- 69 Z. Chen, K. Leng, X. Zhao, S. Malkhandi, W. Tang, B. Tian, L. Dong, L. Zheng, M. Lin, B. S. Yeo and K. P. Loh, *Nat. Commun.*, 2017, **8**, 14548.
- 70 G. L. Liu, A. W. Robertson, M. M. J. Li, W. C. H. Kuo, M. T. Darby, M. H. Muhieddine, Y. C. Lin, K. Suenaga, M. Stamatakis, J. H. Warner and S. C. E. Tsang, *Nat. Chem.*, 2017, **9**, 810–816.
- 71 Y. J. Yu, F. Y. Yang, X. F. Lu, Y. J. Yan, Y. H. Cho, L. G. Ma, X. H. Niu, S. Kim, Y. W. Son, D. L. Feng, S. Y. Li, S. W. Cheong, X. H. Chen and Y. B. Zhang, *Nat. Nanotechnol.*, 2015, **10**, 270–276.
- 72 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.
- 73 V. V. Kulish, O. I. Malyi, C. Persson and P. Wu, *Phys. Chem. Chem. Phys.*, 2015, **17**, 992–1000.
- 74 R. Zhang, B. Li and J. Yang, *J. Phys. Chem. C*, 2015, **119**(5), 2871–2878.
- 75 G. Wang, R. Pandey and S. P. Karna, *Appl. Phys. Lett.*, 2015, **106**, 173104.
- 76 W. Yu, Z. Zhu, C. Y. Niu, C. Li, J. H. Cho and Y. Jia, *Phys. Chem. Chem. Phys.*, 2015, **17**, 16351–16358.
- 77 G. Wang, R. Pandey and S. P. Karna, *Nanoscale*, 2015, **7**, 524–531.
- 78 M. A. Hughes, Y. Fedorenko, B. Gholipour, J. Yao, T. H. Lee, R. M. Gwilliam, K. P. Homewood, S. Hinder, D. W. Hewak, S. R. Elliott and R. J. Curry, *Nat. Commun.*, 2014, **5**, 5346.
- 79 V. P. Pham and G. Y. Yeom, *Adv. Mater.*, 2016, **28**, 9024–9059.
- 80 J. Suh, T. L. Tan, W. J. Zhao, J. Park, D. Y. Lin, T. E. Park, J. Kim, C. H. Jin, N. Saigal, S. Ghosh, Z. M. Wong, Y. B. Chen, F. Wang, W. Walukiewicz, G. Eda and J. Q. Wu, *Nat. Commun.*, 2018, **9**, 199.
- 81 J. Gao, Y. D. Kim, L. B. Liang, J. C. Idrobo, P. Chow, J. W. Tan, B. C. Li, L. Li, B. G. Sumpter, T. M. Lu, V. Meunier, J. Hone and N. Koratkar, *Adv. Mater.*, 2016, **28**, 9735–9743.
- 82 X. Zhang, Z. Shao, X. Zhang, Y. He and J. Jie, *Adv. Mater.*, 2016, **28**, 10409–10442.
- 83 J. Y. Wan, S. D. Lacey, J. Q. Dai, W. Z. Bao, M. S. Fuhrer and L. B. Hu, *Chem. Soc. Rev.*, 2016, **45**, 6742–6765.
- 84 Y. Jung, Y. Zhou and J. J. Cha, *Inorg. Chem. Front.*, 2016, **3**, 452–463.
- 85 J. Wang, H. Fang, X. Wang, X. Chen, W. Lu and W. Hu, *Small*, 2017, **13**, 1700894.
- 86 M. O. Valappil, M. Ahlawat, K. V. Pillai and S. Alwarappan, *Chem. Commun.*, 2018, **54**, 11733–11736.
- 87 S. Mukherjee, L. Kavalsky, K. Chattopadhyay and C. V. Singh, *Nanoscale*, 2018, **10**, 21335–21352.
- 88 J. Wang, S. Li, F. Yun, X. Zhang and Q. Li, *Int. J. Quantum Chem.*, 2020, **13**, e26230.
- 89 F. Safari, M. Fathipour and A. Y. Goharrizi, *J. Comput. Electron.*, 2018, **17**, 499–513.
- 90 J. D. Correa, *Superlattices Microstruct.*, 2019, **30**, 401–408.
- 91 H. Li, L. Zhang, X. Cai, X. Li, B. Wang, W. Yu and R. Zhao, *Mater. Res. Express*, 2018, **5**, 055007.
- 92 B. Meshgingalam and J. Baryestani, *Appl. Surf. Sci.*, 2020, **526**, 146692.
- 93 E. A. Z. Hernandez, E. Florez, L. Dorkis, E. M. Mora-Ramos and J. E. Correa, *Int. J. Quantum Chem.*, 2020, **120**, e26075.
- 94 Y. Xiao, J. Wang, Y. Wang and W. Zhang, *Appl. Surf. Sci.*, 2019, **488**, 620–628.
- 95 K. Liu, J. Fu, L. Zhu, X. Zhang, H. Li, H. Liu, J. Hu and M. Liu, *Nanoscale*, 2020, **12**, 4903–4908.
- 96 Y. Cheng, Y. Song and Y. Zhang, *Phys. Chem. Chem. Phys.*, 2019, **21**, 24449–24457.
- 97 W. Zhang, J. Wang and Y. Xiao, *Appl. Surf. Sci.*, 2019, **488**, 620–628.
- 98 L. Ju, Y. Dai, W. Wei, Y. Liang and B. Huang, *J. Mater. Chem. A*, 2018, **6**, 21087–21097.
- 99 B. J. Wang, X. H. Li, R. Zhao, X. L. Cai, W. Y. Yu, W. B. Li, Z. S. Liu, L. W. Zhang and S. H. Ke, *J. Mater. Chem. A*, 2018, **6**, 8923–8929.
- 100 J. D. Correa, *Superlattices Microstruct.*, 2019, **130**, 401–408.
- 101 W. Lv, B. Yang, B. Wang, W. Wan, Y. Ge, R. Yang, C. Hao, J. Xiang, B. Zhang, Z. Zeng and Z. Liu, *ACS Appl. Mater. Interfaces*, 2018, **10**, 9663–9668.
- 102 P. Luo, F. Zhuge, Q. Zhang, Y. Chen, L. Lv, Y. Huang, H. Li and T. Zhai, *Nanoscale Horiz.*, 2019, **4**, 26–51.
- 103 M. Z. Rahman, C. Kwong, K. Davey and S. Z. Qiao, *Energy Environ. Sci.*, 2016, **9**, 709–728.
- 104 A. Singh, H. Bae, S. Lee, K. Shabbiri, T. Hussain and H. Lee, *Appl. Surf. Sci.*, 2020, **512**, 145641.
- 105 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G. A. Petersson, H. Nakatsuji, M. Caricato, X. Li, H. P. Hratchian, A. F. Izmaylov, J. Bloino, G. Zheng, J. L. Sonnenberg, M. Hada, M. Ehara, K. Toyota,



- R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, J. J. A. Montgomery, J. E. Peralta, F. Ogliaro, M. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, R. Kobayashi, J. Normand, K. Raghavachari, A. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, N. Rega, N. J. Millam, M. Klene, J. E. Knox, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, R. L. Martin, K. Morokuma, V. G. Zakrzewski, G. A. Voth, P. Salvador, J. J. Dannenberg, S. Dapprich, A. D. Daniels, Ö. Farkas, J. B. Foresman, J. V. Ortiz, J. Cioslowski and D. J. Fox, *Gaussian 09, Revision D.01*, Gaussian, Inc., Wallingford CT, 2009.
- 106 J. M. Soler, E. Artacho, J. D. Gale, A. García, J. Junquera, P. Ordejón and D. Sanchez-Portal, *J. Phys.: Condens. Matter*, 2002, **14**, 2745–2779.
- 107 N. Troullier and J. L. Martins, *Phys. Rev. B: Condens. Matter Phys.*, 1991, **43**, 1993–2006.
- 108 J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **77**, 3865–3868.
- 109 J. Guan, Z. Zhu and D. Tomanek, *Phys. Rev. Lett.*, 2014, **113**, 046804.
- 110 J. Bao, L. Zhu, H. Wang, S. Han, Y. Jin, G. Zhao, Y. Zhu, X. Guo, J. Hou, H. Yin and J. Tian, *J. Phys. Chem. C*, 2018, **122**(41), 23329–23335.
- 111 B. Ghosh, S. Nahas, S. Bhowmick and A. Agarwal, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2015, **91**, 115433.
- 112 J. D. Correa, *Superlattices Microstruct.*, 2019, **130**, 401–408.
- 113 Z. Q. Guo, J. P. Zhou, J. Z. Wang, Q. U. Hassan, J. Yang and Y. Ma, *APL Mater.*, 2017, **5**, 026104.
- 114 Z. Zhu and D. Tománek, *Phys. Rev. Lett.*, 2014, **112**, 176802.
- 115 J. Zhang, P. Zhou, J. Liu and J. Yu, *Phys. Chem. Chem. Phys.*, 2014, **16**, 20382–20386.
- 116 M. Sun, W. Tang, Q. Ren, S. K. Wang, J. Yu and Y. Du, *Appl. Surf. Sci.*, 2015, **356**, 110–114.
- 117 J. Li, Q. Pei, R. Wang, Y. Zhou, Z. Zhang, Q. Cao, D. Wang, W. Mi and Y. Du, *ACS Nano*, 2018, **12**(4), 3351–3359.
- 118 W. Gao, J. Lu, S. Zhang, X. Zhang, Z. Wang, W. Qin, J. Wang, W. Zhou, H. Liu and Y. Sang, *Adv. Sci.*, 2019, **6**, 1901244.
- 119 C. Chen, B. Huang and J. Wu, *AIP Adv.*, 2018, **8**, 105105.
- 120 C. Kittel, *Introduction to Solid State Physics*, John Wiley & Sons, New York, 8th edn, 2005.
- 121 P. Bhatia, R. Swaroop and A. Kumar, *RSC Adv.*, 2016, **6**, 101835–101845.
- 122 U. Gupta and C. N. R. Rao, *Nano Energy*, 2017, **41**, 49–65.
- 123 V. Chakrapani, J. C. Angus, A. B. Anderson, S. D. Wolter, B. R. Stoner and G. U. Sumanasekera, *Science*, 2007, **318**, 1424–1430.
- 124 B. Sa, Y.-L. Li, J. Qi, R. Ahuja and Z. Sun, *J. Phys. Chem. C*, 2014, **118**, 26560–26568.
- 125 P. Kumar, R. Boukherroub and K. Shankar, *J. Mater. Chem. A*, 2018, **6**, 12876–12931.
- 126 S. N. Habisreutinger, L. Schmidt-Mende and J. K. Stolarczyk, *Angew. Chem., Int. Ed.*, 2013, **52**, 7372–7408.
- 127 G. Rothenberger, D. Fitzmaurice and M. Graetzel, *J. Phys. Chem.*, 1992, **96**, 5983–5986.
- 128 W. Hu, Z. Li and J. Yang, *Nano Res.*, 2017, **10**, 2223–2233.

