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Unraveling energetics and states of adsorbing oxygen species with MoS₂ for modulated work function†

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MoS₂ and related transition metal dichalcogenides (TMDs) have recently been reported as having extensive applications in nanoelectronics and catalysis because of their unique physical and chemical properties. However, one practical challenge for MoS₂-based applications arises from the easiness of oxygen contamination, which is likely to degrade performance. To this end, understanding the states and related energetics of adsorbed oxygen is critical. Herein, we identify various states of oxygen species adsorbed on the MoS₂ surface with first-principles calculations. We reveal a “dissociative” mechanism through which a physisorbed oxygen molecule trapped at a sulfur vacancy can split into two chemisorbed oxygen atoms, namely a top-anchoring oxygen and a substituting oxygen, both of which show no adsorbate induced states in the bandgap. The electron and hole masses show an asymmetric effect in response to oxygen species with the hole mass being more sensitive to oxygen content due to a strong hybridization of oxygen states in the valence band edge of MoS₂. Alteration of oxygen content allows modulation of the work function up to 0.5 eV, enabling reduced Schottky barriers in MoS₂/metal contact. These results show that oxygen doping on MoS₂ is a promising method for sulfur vacancy healing, carrier mass controlling, contact resistance reduction, and anchoring of surface electron dopants. Our study suggests that tuning the chemical composition of oxygen is viable for modulating the electronic properties of MoS₂ and likely other chalcogen-incorporated TMDs, which offers promise for new optoelectronic applications.

Introduction

Molybdenum disulfide (MoS₂), an emerging two-dimensional (2D) material consisting of a Mo monolayer sandwiched between two S monolayers, has recently attracted extensive

New concepts

MoS₂ has been extensively explored for integration into advanced electronics and is leading the trend of next-generation semiconductor devices. However, the low formation energy of sulfur vacancies leads to intrinsic n-type doping character and degrades the original material performance. Furthermore, the existence of ubiquitous oxygen species, likely to be introduced during manufacturing, processing, or ambient exposure, emphasizes the need for investigation of the atomic-scale mechanism of oxygen doping. Here, we discover a “dissociative” mechanism of molecular oxygen trapped at sulfur vacancies to facilitate single-atomic oxygen doping in MoS₂, setting the foundation for forming molybdenum oxy-sulfide. Density-functional theory calculations show that there exist two favorable states of single-atomic oxygen: the top stacking and bottom substituting states. Interestingly, the oxygen doping induces a lattice strain field which competes with the stoichiometry to tune the carrier mass. Furthermore, we demonstrate modulation of the work function and accordingly the Schottky barrier in MoS₂/metal contact with varying oxygen concentrations. This sets the foundation of oxygen doping engineering of MoS₂ for sulfur vacancy healing, contact resistance reduction, and anchoring of surface electron dopants for a plethora of applications.

attention due to its high mechanical flexibility and intriguing electronic properties.^{1–3} Monolayer and few-layer MoS₂ are being intensively explored for field effect transistors,^{4,5} sensors,^{6,7} photocatalysis,⁸ energy harvesting,^{9–11} and other advanced optoelectronics.^{12–14} Extraordinary performance was obtained by depositing MoS₂ on high-dielectric oxides leading to carrier mobility up to 200 cm² V⁻¹ s⁻¹ and a high switching ratio of ~10⁸, making it a substitute for classical silicon semiconductors for miniaturization.¹⁵

Thus far, extension of Moore’s law by continually scaling down silicon-based integrated circuits has encountered a dilemma.¹⁶ As a 2D alternative, MoS₂ is a promising material for balancing device minimization and performance optimization.¹⁷ Its atomic level thickness and moderate carrier mobility are favorable for gate controllability.^{18,19} Nevertheless, the semiconducting character and weak dielectric screening make monolayer MoS₂ extremely sensitive to intrinsic defects, extrinsic adsorbates, and strain. Indeed, as the most common natural defect, sulfur vacancy (V_S),

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at a typical defect density of 10^{13} cm^{-2} , degrades carrier mobility due to defective bandgap states and lifting of Schottky barriers through Fermi-level pinning.^{20,21}

Owing to their atomically thin character and high surface-to-volume ratio, MoS₂ nanostructures are subjected to environmental doping by molecular adsorbates which has a pronounced effect on their electronic and optoelectronic properties.^{22,23} On the other hand, it has been widely recognized that molecular adsorbate doping is an effective technique for defect passivation in 2D electronics.^{24,25} Among various possible adsorbates, oxygen species must be the most prevalent impurity source after or during growth of 2D materials. Unintentional uptake of O species in a MoS₂ layer may arise from (1) synthesis due to incomplete sulfurization of MoO₃ or the presence of oxygen using chemical vapor deposition;^{26–30} (2) introducing oxide substrates, normally Al₂O₃ or SiO₂, and high-dielectric capping layers like HfO₂ or ZrO₂ during device integration, where oxygen vacancies in the oxides, generated by gate voltage or temperature gradient, can be formed and the depleted O atoms are susceptible to recapture by the MoS₂ layer;^{31–34} (3) exposing MoS₂ layers to atmospheric adsorbates in ambient conditions during application.³⁵ Other processes with a high possibility of introducing O impurities include adopting an ionic-liquid gate, liquid-phase exfoliation, plasma-assisted

nano-printing, and synthesizing MoS₂-based nanohybrids.^{36–40} Compared with a perfect surface, the adhesion and effect of O species should be more prominent if the layer is partially reduced by forming V_S in the basal plane or dissected for ribbons or nanoflakes due to abundant dangling bonds. In contrast to the well-understood effects of O on graphene (bandgap opening and formation of graphene oxide) and boron nitride (BN), a clear picture of O species evolution in MoS₂ is still absent; to be specific, the energetics and kinetics of O species and the impact of adsorbed O species on structural modification and electronic properties of MoS₂ are still unclear,^{41,42} although their pronounced effects on healing of vacancy sites and reducing the MoS₂/metal contact barrier have been recognized.^{43,44}

In this work, we present a theoretical investigation of the surface affinity of MoS₂ toward O species based on a first-principles study. We identify the possible adsorbing states and types of oxygen atoms on MoS₂. A dissociative mechanism of molecular oxygen adsorbing at a V_S into two atomic species, a substituting oxygen and a top oxygen, is revealed. We demonstrate substantial atomic displacements and gradient strains surrounding each single oxygen dopant. Through the uptake of more oxygen atoms, lattice strain is developed and scaled with the oxygen content. We show an opposite effect on electron and hole mass with this chemical functionalization, but the electron mass is less sensitive to the oxygen content than the hole

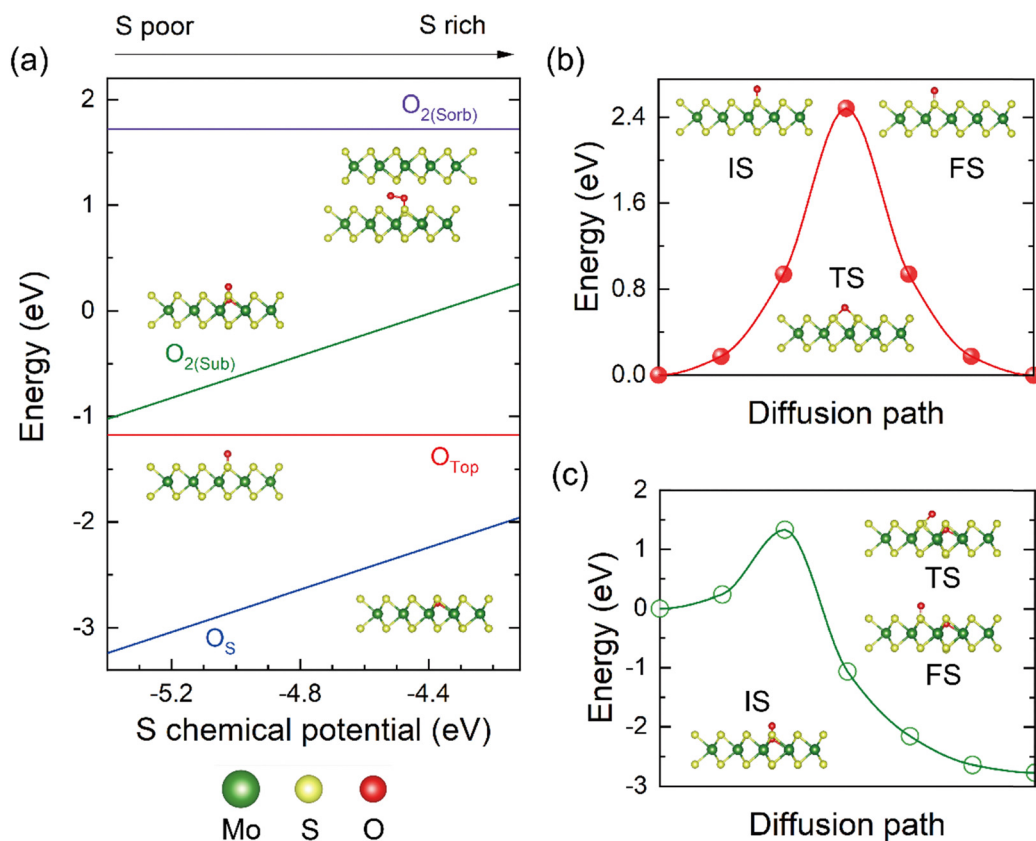


Fig. 1 (a) Sulfur chemical potential (μ_s) determined the formation energy for various O adsorbates on MoS₂. The μ_s in the S-rich condition is equal to that in the S₈ molecular crystal, and in the extreme S-poor limit μ_s is bounded by the formation of Mo in the film. (b) Activation energy for O_{Top} diffusing above the MoS₂ layer. (c) Activation energy for O₂(Sub) decomposing into a (O_{Top}, O_S) pair. The green, yellow, and red spheres represent Mo, S, and O atoms, respectively. The insets in (b) and (c) are the structure of the initial state (IS), transition state (TS), and final state (FS).



mass. Finally, a surface decoration modulated work function is also demonstrated with implication for reducing the Schottky barriers.

Results and discussion

Energetics, dynamics and states of adsorbed oxygen species

In the graphene case, more than one type of oxygen species is involved in modifying the properties.⁴⁵ Here, energetics and stability of O species on a MoS₂ monolayer are quantitatively evaluated by considering the following potential adsorption configurations (Fig. 1(a)): oxygen adsorption above the anionic plane sitting atop a S atom (O_{Top}), oxygen atom substituting a S atom (O_S) which is equivalent to the oxygen atom occupying the S vacancy, and molecular O₂ substituting a S atom which is equivalent to formation of peroxide species or oxygen molecule trapped by a S vacancy (O_{2(Sub)}). Other possible adsorption states, like O occupying above the hollow position defined by the equilateral triangle formed by three S atoms, are unstable (Fig. S1 and Table S1, ESI†). The formation energy (E_f) for the O_{Top} is -1.2 eV with an O–S bond length of 1.483 Å. For the O_S case, this state shows the lowest E_f of -3.25 eV under the S-poor condition. This indicates that the substituting O species is very likely to occur in the MoS₂ layer, especially for films after reducible gas treatment, *e.g.*, a hydrogen plasma increases the V_S on the MoS₂ basal plane.^{46,47} The substituting O atom relaxes about 0.412 Å toward the middle Mo layer with an O–Mo bond length of 2.052 Å. For the O_{2(Sub)} structure, E_f is still negative but higher than that of the O_S type. The O–O bond length is elongated to 1.374 Å compared with 1.211 Å of a free O₂ molecule, implying a strong activation of the O–O bond by populating the 2π* antibonding orbital of the O₂ molecule.

Binding of an O₂ molecule above the anionic basal plane of perfect MoS₂ is unstable and shows very weak physisorption on the monolayer. However, for multi-layer or bulk MoS₂, oxygen molecules may be trapped inside the van der Waals (vdWs) gap, denoted as O_{2(Sorb)}. As shown in Fig. 1(a), the adsorption is an endothermic process with a positive E_f of 1.7 eV. In this metastable configuration, one of the O atoms forms a chemical bond with a S atom in one MoS₂ layer, while the other is tilted toward the gap above the center of a triangle formed by S atoms of the same MoS₂ layer. The O–O and the O–S bond lengths are 1.465 Å and 1.591 Å, respectively. Additionally, the much longer O–O bond in O_{2(Sorb)} within the vdWs gap than that in O_{2(Sub)} trapped at the vacancy suggests higher amounts of electrons transferring to occupy the 2π* antibonding orbital of the O₂ molecule, probably due to a denser electron density in the vdWs gap. Therefore, through thermal activation under finite temperature, we can still expect the entry and adsorption of O₂ molecules in bulk or multi-layer MoS₂ despite the ultraweak adsorption above the basal plane of the surface. Fig. 1(b) shows activation barriers calculated by the climbing image nudged elastic band (CI-NEB) method for the O species diffusing on the surface. For O_{Top} species, the barrier for O jumping along the neighboring lattice is around 2.4 eV. For O_{2(Sub)}, there is likely to be a dissociative process *via* O_{2(Sub)} → O_{Top} + O_S, where the

protruding O atom moves to the top of its neighboring S atom simultaneously breaking the O–O bonding and the activation energy is around 1.2 eV, much lower than that of the O_{Top} species. This indicates a viable way of filling V_S through molecular oxygen while forming oxygen decoration. Notably, the low-temperature preparation of MoS₂ below 100 °C is plagued by the increased adsorption of water, serving as an additional oxygen contamination source, and the related H₂O molecule adsorption and dissociation on MoS₂ have been reported.^{48,49} Here, the energy barriers for O₂ dissociation in bulk-phase MoS₂ are also considered (Fig. S2, ESI†). This barrier height is correlated with the stability of bulk-phase MoS₂, as the diffusion and dissociation of O₂ molecules change the interlayer vdWs interactions and promote the exfoliation of the MoS₂ monolayer.

Atomic displacement and gradient strain surrounding oxygen dopants

Fig. 2(a) and (b) illustrate the relaxation of periphery S/Mo atoms around the single O_{Top} and O_S dopant centers in the supercell, respectively. The introduction of O species in the MoS₂ monolayer causes gradient strains, leading to the host lattice expansion (for O_{Top}) and contraction (for O_S). For the O_{Top} species, the three nearest Mo atoms relaxed outward by about 0.029 Å. The magnitude of the strain field, indicated by the length of orange arrows, decays rapidly away from the oxygen species; for instance, the second-nearest-neighbor S atoms relax only outwardly by about 0.003 Å. For the O_S, there is a strong compressive field around the substituting dopant center with stronger O–Mo binding and a shorter O–Mo bond length, due to a smaller atomic radius and a lower electronegativity of the O atom than that of the S atom. The three neighboring Mo atoms move inward 0.129 Å, and the next-nearest S species relaxed inward 0.065 Å with appreciable strain even extends to about 4 or 5 nearest neighbors. In a word, we identify a sizable strain field associated with a substitutional oxygen dopant in MoS₂. This is expected to be true in other 2D materials. With the lack of a continual lattice in 3D crystal, 2D materials are prone to generate a larger area of buffering region with the formation of gradient strain around the dopant.

We next examine the effect of varying the content of oxygen dopants. To simulate the ambient condition where oxygen sources largely access one surface of MoS₂, the other side would be blocked due to the substrate. In this extreme condition, all O atoms are modeled to adsorb on one facet of the S–Mo–S layer. As for the case of O adsorbing on both facets, all the O species are uniformly distributed with maximized separation, and the more complicated oxygen cluster formation on the surface is not considered (see Fig. S3 and S4 for the optimized structures, ESI†). Notably, the uptake of O atoms does not induce any dangling bonds, which is favorable for keeping planeness and speed of transporting carriers.⁵⁰

Highly concentrated O-doped MoS₂ becomes molybdenum oxy-sulfide (MoS_yO_x), and here the evolution of the lattice constant of two types of molybdenum oxy-sulfide (MoS₂O_x containing O_{Top} and MoS_{2–x}O_x containing O_S with x being the content of oxygen normalized to normal chemical formula)



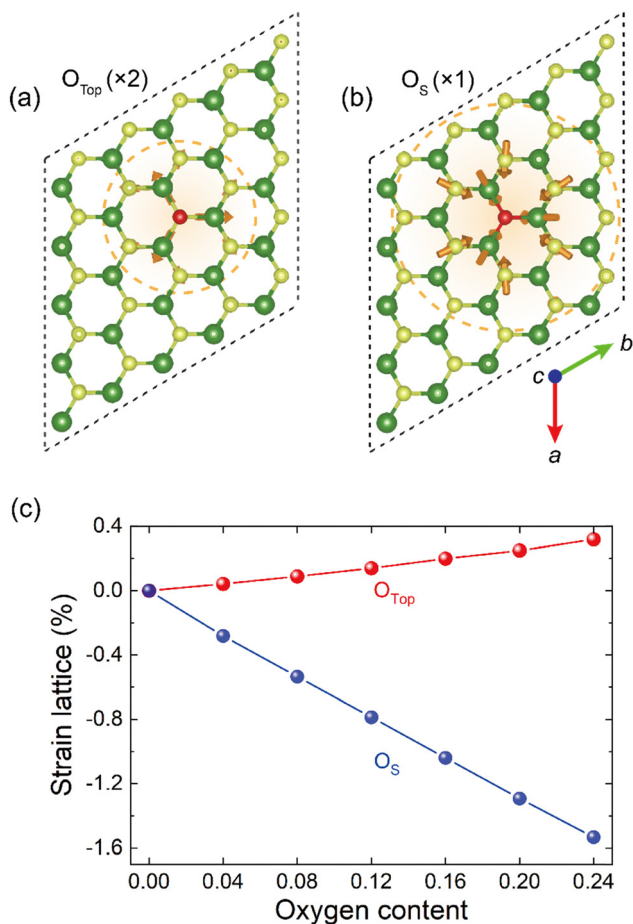


Fig. 2 Doping-induced strain field by single O_{Top} (a) and O_{S} (b). The orange arrows denote displacements of atoms from their original positions in a pristine lattice to positions in a defective lattice, with the arrow lengths representing the magnitudes of the adjustments. For visualization purposes, the length of the displacement vectors in (a) is doubled. The dashed black line represents the simulation supercell box. (c) Lattice strain intensity (expansion for O_{Top} and shrinkage for O_{S}) as a function of oxygen content. The $5 \times 5 \times 1$ supercell of molybdenum oxy-sulfide is used to account for the lattice evolution.

layer with different stoichiometry is calculated and shown in Fig. 2(c). For the O_{Top} case, the lattice constant increases gradually, and the strain (δ)-O content (x) relation scales as $\delta = C \times x$ with C fitted to be 1.32, consistent with the previous expansive strain field around the dopant center shown in Fig. 2(a). For the O_{S} case, introducing O_{S} causes significant compressive strain on the lattice. The δ - x relation also follows $\delta = C \times x$ with C fitted to -6.36 , showing a much larger distortion of the MoS_2 host lattice than the O_{Top} case. For instance, for the O content with x of 0.24, the induced lattice strain is 0.3% for O_{Top} and -1.5% for O_{S} . While here the results are only calculated for single-layer MoS_2 , similar scaling behavior of the lattice constant with oxygen dopant would be true for multi-layer MoS_2 and for adsorption evenly at both basal planes of MoS_2 (Tables S2 and S3, ESI[†]). For the latter, due to the intercalation of O or O_2 species (Fig. 1) in the vdWs gap, the lattice constant normal to the atomic plane would also increase

with the O content. From an experimental perspective, Burns *et al.* used high-resolution transmission electron microscopy (HR-TEM) to detect local strain fields induced by vacancies in MoS_2 . They suggest that O atoms adsorbed on a vacant S site, denoted as O_{S} in our work, lead to the lattice contraction at high irradiation fluences.⁵¹ This phenomenon aligns with the trend in Fig. 2(b). Recently, Kumar *et al.* found the combination of tensile strain and the absence of charged oxygen adatoms, denoted as O_{Top} in our work, triggering a drastic photoluminescence (PL) redshift of ~ 130 meV for MoS_2 .⁵² These findings suggest that surface oxygen dopants and induced lattice strain field have profound effects on modifying the electronic properties.

Modulation of band structure, effective mass and work function

We next show the evolution of the electronic structures of MoS_2 after surface oxygen treatment. Fig. 3(a) shows band structures of the O_{Top} and O_{S} cases, where the contribution of O atoms is projected out and their components are scaled by the size of red circle markers at each k point and band. In contrast to the bandgap opening of graphene oxide,⁵³ there is no strong modification of bandgap with mild O decoration. The bandgap of the oxygen-decorated MoS_2 monolayer decreases almost linearly with the oxygen content varying from 0.04 to 0.24 for O_{Top} and O_{S} oxygen species (Fig. 3(c)). As an iso-valent dopant, the addition of oxygen, either the substitutional O_{S} or the top O_{Top} , does not induce any defective levels in the bandgap, thus trapping of carriers by these two types of O species is unlikely to be prominent. However, there is evident modification of the valence band edge through orbital hybridization, while modulation of the conduction band edge is tiny as there is negligible O hybridization with Mo atoms.

Since band edges are critically important in terms of effective mass of carriers and deformation energy of the frontier orbitals, the adsorption of O species may influence the performance of the MoS_2 layer such as mobility of carriers, especially for holes. Fig. 3(b) presents the partial density of states (PDOS) for O_{Top} , O_{S} , and $\text{O}_{2(\text{Sub})}$. For O_{Top} and O_{S} , the O states spread over the whole valence band and no defective states occur in the gap. In contrast, for $\text{O}_{2(\text{Sub})}$ adsorption, the clamped O_2 at the S vacancy site induces new states above the valence top. From the plot of surface charge density of valence band maximum (VBM) and conduction band minimum (CBM) shown in Fig. S5a and c (ESI[†]), these O states are strongly localized, suggesting relatively weak hybridization of the molecule with the MoS_2 substrate. The additional peak located ~ 0.3 eV above the VBM corresponds to the π orbital of surface O_2 . According to these results, the shallow defective level would have a weak effect on carrier mobility, as $\text{O}_{2(\text{Sub})}$ is metastable and has a tendency to transfer into the O_{Top} and O_{S} states.

Fig. 4 shows the evolution of electron/hole effective mass with the content of O species. Electron (hole) mass of pristine MoS_2 monolayer is $0.46 m_e$ ($-0.60 m_e$), consistent with a previous report of $0.47 m_e$ ($-0.60 m_e$).⁵⁴ Upon uptake of O species, for O_{Top} cases (Fig. 4(a)), the magnitude of hole mass increases steadily from $-0.60 m_e$ for zero oxygen content to $-0.75 m_e$ with O content x of 0.24. In contrast, the trend for the



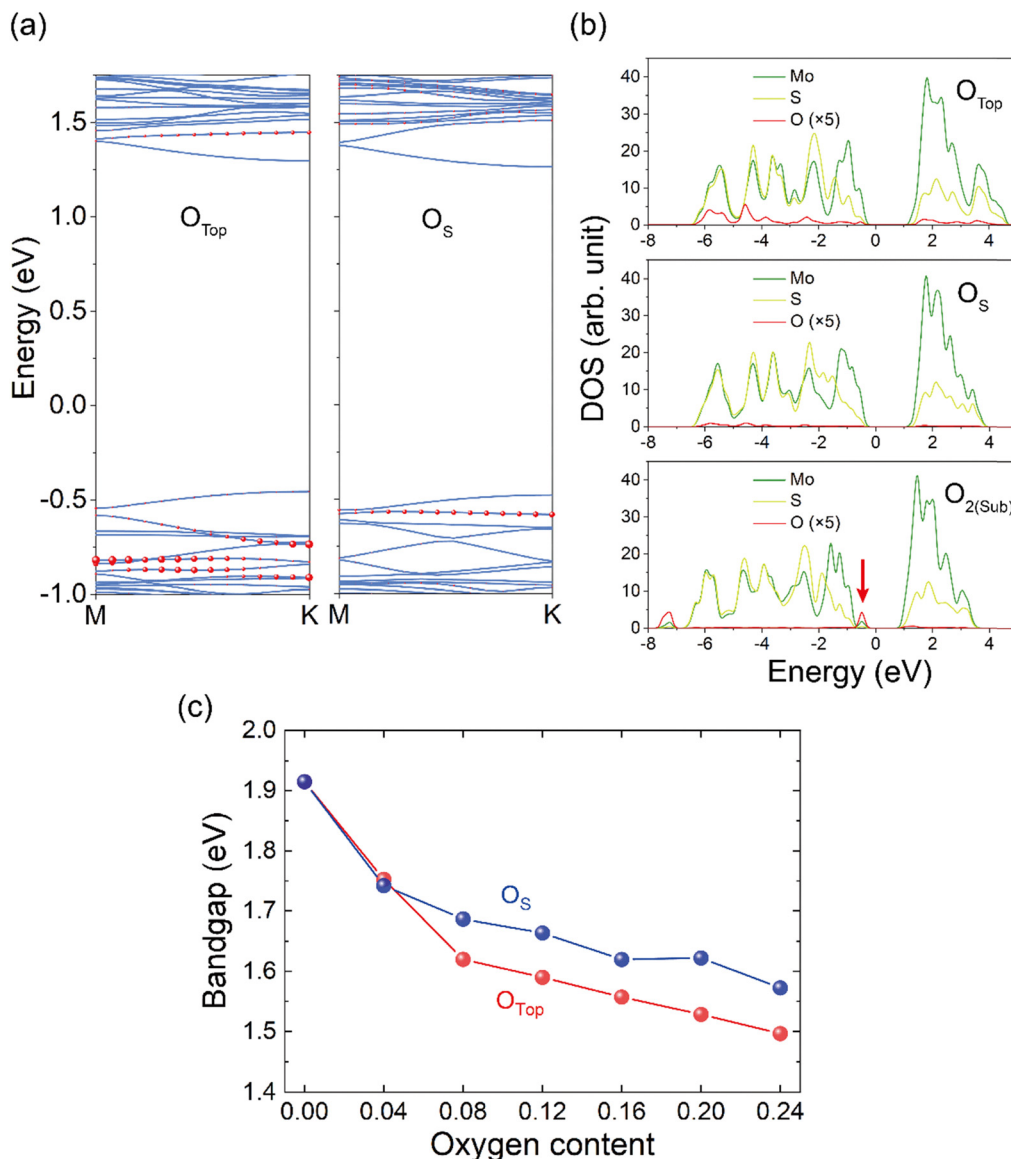


Fig. 3 Electronic structures in oxygen-decorated MoS_2 . (a) Oxygen contribution projected band structure of O_{Top} and O_S . The size of the red circle markers indicates the contribution from oxygen orbitals. (b) Partial density of states (PDOS) of O_{Top} , O_S , and $O_{2(Sub)}$. The red arrow denotes the in-gap states in $O_{2(Sub)}$. (c) Bandgap as a function of oxygen content.

O_S case (Fig. 4(b)) starts with a plateau occurring at the onset corresponding to the dilute doping limit, the hole mass then decreases with increasing O content and levels off at $-0.22 m_e$ at x equal to $0.12 m_e$. For both cases, the value of electron mass fluctuates around $\sim 0.50 m_e$, and the adsorption of O species has a weaker effect on electron mass than on hole mass. According to Fig. 3(a) and (b), this is due to a weaker hybridization of O states with Mo orbitals at the CBM while a strongly coupled O component at the VBM (also see Fig. S6 and S7, ESI[†]). Similar results also hold true for oxygen species distributing on both facets of sulfur in MoS_2 (Fig. S8 and S9, ESI[†]).

In fact, such doping-level-controlled carrier mass is a synergistic effect of chemical variation and strain modification after introduction of O species. For the O_{Top} case, the adsorption only leads to a tiny strain within the range of O content

$x \leq 0.24$; thus the increasing hole mass is solely attributed to the chemical variation of the top O atoms above the S atoms. However, for O_S , as shown in Fig. 2(b), it induces significant strain on the host lattice. To decouple the effect of the chemical modification of O atoms from strain field and identify dominant factors affecting the carrier mass, we performed additional calculations with a frozen lattice where the lattice constant is kept the same for all the O content to exclude the strain effect. The lattice constants of the O_{Top} and O_S models with varied oxygen content are fixed to the same lattice values as their 0.04 oxygen content cases, respectively (see Table S2 and S3 for details, ESI[†]). The x -electron/hole mass relationship for the frozen lattice is represented by the dashed lines in Fig. 4(a) and (b). For O_{Top} , the lines associated with holes for the frozen lattice are nearly overlapped with those for the



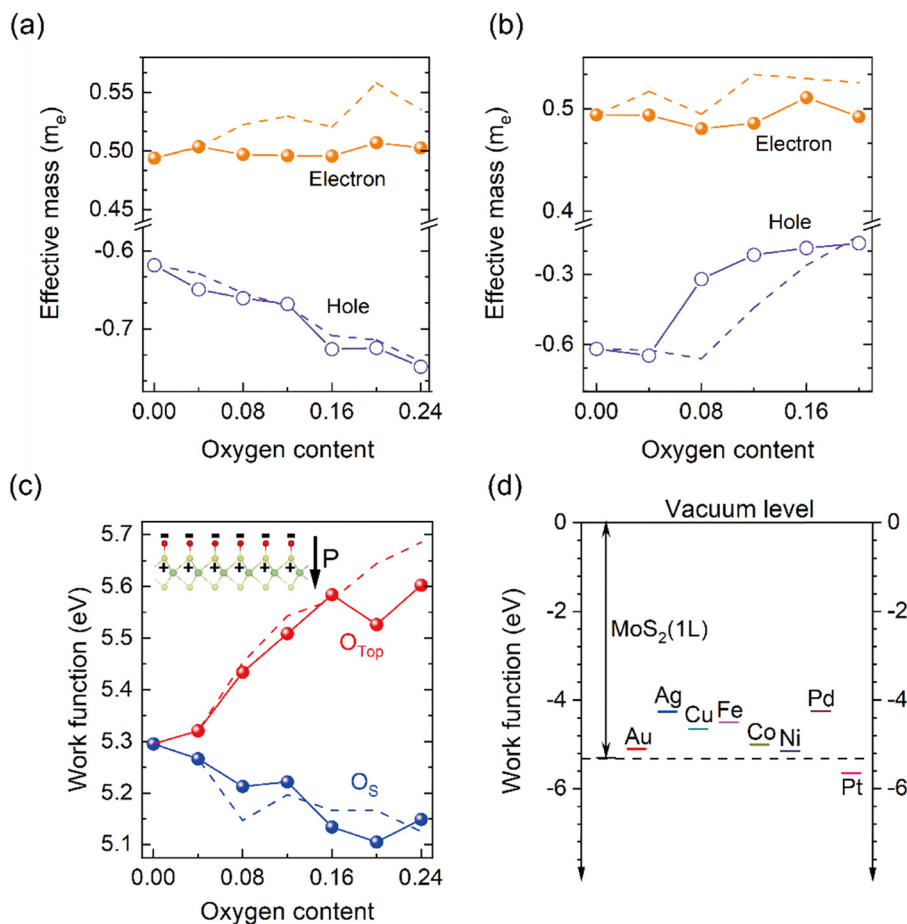


Fig. 4 Oxygen content-mediated effective mass for the single O_{Top} (a) and O_{S} (b) species. (c) Work function of O_{Top} and O_{S} as a function of oxygen content. The solid and dashed lines correspond to calculations with structures of relaxed lattice of oxygen-doped MoS₂ and frozen lattice of pristine MoS₂, respectively. The inset shows the inward surface dipole associated with surface top-type oxygen, O_{Top} , accounting for the increase of work function. (d) Work function alignment of monolayer MoS₂ and normal metal electrodes.

relaxed lattice. This is reasonable considering the moderate structural relaxation of the host layer with O uptake. For O_{S} , both trends of with/without strain are similar, showing a lighter hole mass with greater oxygen incorporation. However, there is a clear deviation of the hole mass between the relaxed and fixed cases. For x between 0.08 and 0.2, the hole mass of frozen lattice is linearly correlated with x indicating the substituting O decreases the hole mass (violet dashed line in Fig. 4(b)). While for the relaxed lattice, the slope of the line continually decreases, and the hole mass levels off to $-0.22m_e$ at a doping content of 0.12 (violet solid line in Fig. 4(b)). Therefore, compressive strain induced by dense O_{S} tends to promote the hole mass.

A similar strain effect has also been found in other types of sulfide semiconductor.⁵⁵ In a word, upon introduction of O, while the chemical modification decreases the hole mass, there is a concomitant increase of hole mass due to the compressive strain, thus decreasing the slope of the x -hole mass profile in the relaxed lattice case. This inherent strain accompanied by the O doping would play an important role in affecting the electronic properties and accordingly carrier transport behavior in field effect transistors (FETs). Our study shows that creating

O_{S} species with an appropriate O content may effectively decrease the hole mass thus promoting the hole mobility, whereas the electron mass is less affected. Compared with the V_{S} -induced flat defect states (Fig. S10b, ESI†) for trapping and scattering charge carriers, the adsorbed oxygen atoms somehow heal the vacancies and mitigate the defect effect on carrier mobilities. Indeed, Wei *et al.* fabricated oxygen-doped MoS₂ monolayer at the wafer scale, and the final FET device shows high mobilities with an average of $78 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$, being greater than that of intrinsic MoS₂ FETs of $48 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$.⁵⁶ Tang *et al.* also suggested that moderate oxygen doping significantly improves the carrier mobilities of MoS₂ by removing V_{S} .⁵⁷

Next, we demonstrate that by controlling the type and content of O species, the work function (W) of the hybrid MoS _{y} O _{x} layer can be tuned, which is highly desired for electronic devices to control band alignment with metal electrodes. Several metals have been proven to form a Schottky contact at the MoS₂/metal interface, and an effective strategy for reducing the Schottky barrier height (SBH) is the shifting of W of MoS₂.^{58–60} In Fig. 4(c), W is calculated by subtracting the Fermi energy from the electrostatic potential in the middle of the vacuum layer (eqn (5)).^{61,62} W of the MoS _{y} O _{x} layer increases



monotonically from 5.3 to 5.6 eV for O_{Top} species with x increasing from 0 to 0.24. In contrast, W decreases gradually with increasing O_{S} content, implying overall effective surface positive charges with the presence of O_{S} species. The modification from 5.6 eV (for O_{Top}) to 5.15 eV (for O_{S}) at an O content of 0.24 indicates a large modulation space of W (around 0.5 eV) with the addition and control of O species in the film. Due to the larger electronegativity of an oxygen atom than that of a sulfur atom, the oxygen is negatively charged, and the sulfur is positively charged. A dipole is induced associated with the oxygen–sulfur pair with the dipole direction pointing inwardly from the adsorbing oxygen atom to its bonded sulfur atom (see the inset in Fig. 4(c)). On the O_{Top} surface, the exterior negatively charged oxygen inhibits electrons escaping from the surface and promotes W . While for the O_{S} surface, the reversed dipole direction is favorable for surface electron escape, hence reducing W . Fig. 4(d) shows the comparison and alignment of W of pristine MoS_2 monolayer and common metal electrodes. By considering the variance of W induced by different types and varying content of oxygen as shown in Fig. 4(c), Au, Co, Ni, and Pt are predicted to be suitable for contact with the MoS_2O_x layer due to a low SBH. For comparison, Gupta *et al.* used Kelvin probe force microscopy (KPFM) to measure W of MoS_2 , for which the existence of MoO_{3-x} phase alters W from 4.98 to 5.56 eV.⁶³ In fact, such dopant-controlled electronic structures were also observed in other TMDs. Oberoi *et al.* adopted a self-limiting oxidation method to convert a

top-layer WSe_2 channel to WO_xSe_y alloy, achieving high-performance p-type FETs with a 5-fold improvement in the on-state performance and a 9-fold reduction in the contact resistance.⁶⁴

Due to the relatively wide bandgap of MoS_2 and thus an enlarged SBH, the contact resistance of the MoS_2 /metal junction is supposed to be much larger than that of the graphene/metal junction.⁶⁵ A variable W of the MoS_2 layer is highly important for tuning the SBH. The demonstrated tunable range of W of ~ 0.5 eV by O doping is large compared to the intrinsic bandgap of the monolayer (1.8 eV). Therefore, surface modification can significantly modulate the barrier of carrier injection at a metal/ MoS_2 contact and may even allow control of carrier polarity in the film by tuning the content and the type of O species in the monolayer. For silicon or other oxide as the channel material, the energetic alignment at the interface can be controlled by dopants or defects.⁶⁶ However, unlike graphene, the presence of a finite gap of MoS_2 may create in-gap states upon doping which will trap the charges and affect the carrier mobility. Here we prove that the introduction of iso-chemical species like substituting O, being free of in-gap levels, is highly important for tuning the properties of MoS_2 for various applications. The strain-chemical-property relationships inherent in the O modification can be tuned by controlling the adsorbing states of O species on the MoS_2 monolayer, enabling a synergetic effect of strain and chemical functionalization. Moreover, due to the more electronegative O species,

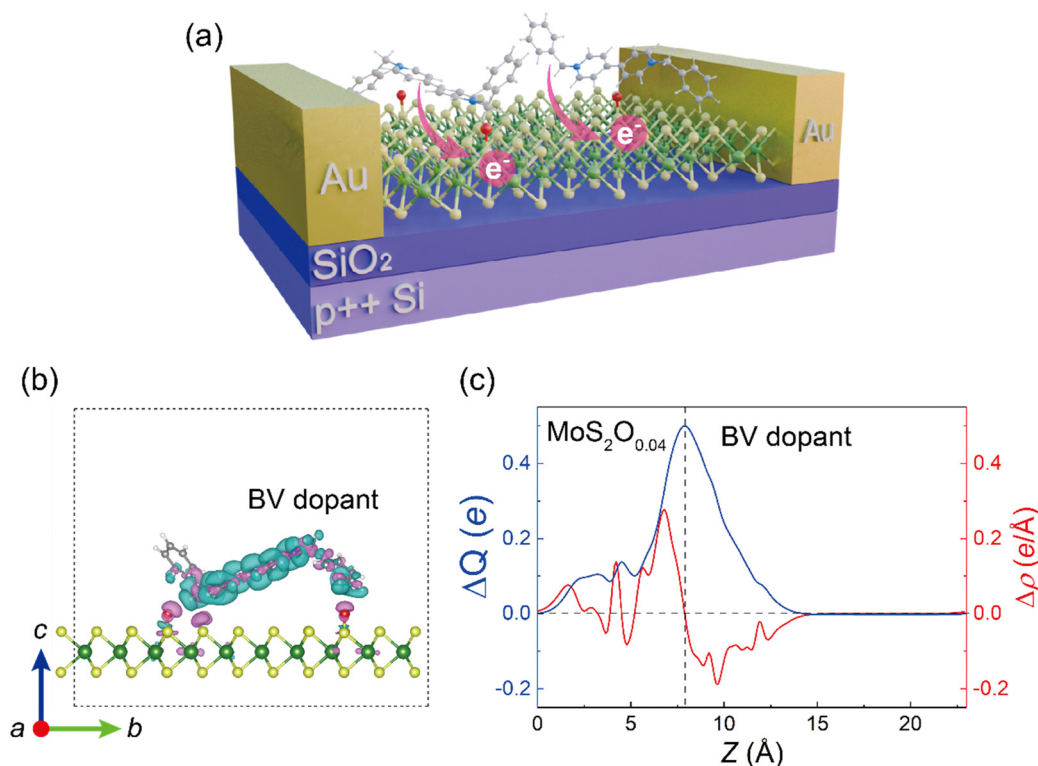


Fig. 5 (a) Schematic of device setup of MoS_2O_x FETs after surface BV doping. The monolayer $\text{MoS}_2\text{O}_{0.04}$ serves as a gate layer and contact with Au electrode. (b) Charge redistribution across BV/ $\text{MoS}_2\text{O}_{0.04}$ interfaces by DFT-calculated differential charge density with an iso-surface of 7×10^{-3} e. The charge accumulation and dissipation regions are represented by pink and cyan colors, respectively. (c) Planar-averaged differential charge density $\Delta\rho$ and transferred charge ΔQ along the out-of-plane direction (c axis). The dashed black line in (c) is used as a guide to the eye.



this 2D MoS₂-O hybrid layer can act as a host for assembling with positively charged nanoclusters or particles.³⁷

Finally, we demonstrate a proof-of-concept device based on the physicochemical properties of the MoS_yO_x layer in this work. Fig. 5(a) shows the device structure of benzyl viologen molecule (BV)-doped MoS₂O_x FETs with Au layer as contact metal. BV is a common n-type dopant with strong reductant ability for nanomaterials.^{22,67,68} Fig. 5(b) shows the MoS₂O_{0.04}/BV interface model, where a BV molecule interacts with MoS₂O_{0.04} through the stretched oxygen site, and it can be seen that the BV molecule loses electrons (cyan surface for electron loss) which transfer to the MoS₂O_{0.04} side (pink surface for electron gain). To quantify the interaction between the surface O species and BV molecules, the planar-averaged differential charge density $\Delta\rho$ and transferred charge ΔQ are considered, defined as:

$$\Delta\rho = \rho_{\text{MoS}_2\text{O}_{0.04}/\text{BV}} - \rho_{\text{MoS}_2\text{O}_{0.04}} - \rho_{\text{BV}} \quad (1)$$

$$\Delta Q(z) = \int_0^z \Delta\rho(z) dz \quad (2)$$

where $\rho_{\text{MoS}_2\text{O}_{0.04}/\text{BV}}$, $\rho_{\text{MoS}_2\text{O}_{0.04}}$, and ρ_{BV} are the planar-averaged charge density of the entire interface system, isolated MoS₂O_{0.04} flake, and isolated BV molecule, respectively, and z is the vertical distance from the bottom. In Fig. 5(c), the maximum ΔQ value of $\sim 0.5e$ at the vdWs gap denotes the transferred electrons from each BV molecule to MoS₂O_{0.04}. This large transfer value indicates the strong anchoring of BV molecules on the oxygen site (O_{Top}).

Conclusions

To summarize, the ubiquitous presence of oxygen species in the environment and synthesis processes renders a high possibility of oxygen uptake in MoS₂ samples. Here based on DFT calculations, we reveal the energetics, forms, and effects of oxygen adsorbates and dopants in MoS₂. We identify the two most stable forms of oxygen, surface atomic adsorbate O_{Top} and substituting oxygen O_S, both of which are electronically inert and free of in-gap dopant-induced states despite appreciable local strains around the dopants. These dopant states would occur through a dissociative mechanism of O₂ trapped at the V_S overcoming an energy barrier of ~ 1.2 eV. We predict that the hole mass is more sensitive to the oxygen content than the electron mass, due to less hybridization of oxygen with CBM. Controlling the oxygen stoichiometry allows modulation of the work function up to ~ 0.5 eV, which is favorable for reducing the SBH and contact resistance with metal electrodes. Our study elucidates the mechanism for controlling oxygen dopants on the MoS₂ monolayer to achieve modulated carrier mass and work function, motivating further study of emerging optoelectronic devices based on 2D materials.

Methods

The Vienna *ab initio* simulation package (VASP) with the projected augmented wave (PAW) potential is adopted for density functional theory (DFT) calculations.⁶⁹ Exchange and

correlation energy of electrons are considered through the generalized gradient approximation (GGA) with the Perdew–Burke–Ernzerhof (PBE) functional.⁷⁰ The DFT-D2 method of Grimme is used to describe the vdWs interactions, this correction having been widely adopted for multiple low-dimensional materials.⁷¹ The energy cutoff is set to 450 eV and sampling of the Brillouin zone is performed by using a uniform $3 \times 3 \times 1$ dense k -point grid. Energy and forces are converged until 10^{-5} eV and 0.01 eV \AA^{-1} , respectively. A slab model is based on a $5 \times 5 \times 1$ supercell with a vacuum space with a thickness of 15 \AA to avoid interactions from periodic replicas. The vacuum layer thickness is tested for the coverage of work function (see Fig. S11, ESI†). Spin polarization is considered in all calculations. The lattice constraint values are calculated by comparing the lattice constants of oxygen-adsorbed systems and those of the pristine MoS₂ monolayer. The chemical potential (μ) should be considered for the formation of MoS₂ film. Here, the energy values based on DFT results are used for estimating the formation energy of O adsorbates on MoS₂ within the allowed crystal growth conditions with the chemical potential of sulfur species (μ_s) varying from S-poor to S-rich conditions. The stability of these O adsorbates is estimated by judging E_f defined as:

$$E_f = E(\text{O:MoS}_2) - E(\text{MoS}_2) - \sum_i n_i \mu_i \quad (3)$$

where $E(\text{O:MoS}_2)$ and $E(\text{MoS}_2)$ are the total energy of the O-adsorbed system and the equivalent MoS₂ supercell, respectively. The doped system is formed by either adding (positive) or removing (negative) n_i atoms for the i type of atom with a chemical potential of μ_i , with i being S (sulfur) or O (oxygen). By assuming the chemical potential of Mo and S are in thermal equilibrium with MoS₂, μ_{Mo} and μ_s satisfy the relation:

$$\mu_{\text{MoS}_2} = 2\mu_s + \mu_{\text{O}} \quad (4)$$

where μ_{MoS_2} represents total energy of MoS₂ per formula unit. For the S-rich limit condition, the source of sulfur is from gaseous S₈ molecules during chemical vapor growth; therefore S₈ is chosen as the reference state for determining $\mu_{\text{Mo,max}} = E_{\text{S}_8}$, where E_{S_8} is the total energy of molecular crystal S₈ per atom obtained from DFT calculation. Under the Mo-rich limit (in the meanwhile S-poor limit) condition, $\mu_{\text{Mo,max}} = E_{\text{Mo}}$, and E_{Mo} is the energy of bulk Mo bcc phase metal crystal per atom from DFT results, while $\mu_{\text{s,min}} = (\mu_{\text{MoS}_2} - \mu_{\text{Mo,max}})/2$ as required by the assumption of equilibrium-growth condition. The chemical potential of O atom (μ_{O}) and O₂ molecule (μ_{O_2}) are the energy of isolated O atom and O₂ molecule, respectively. Dipole correction is added for all structures along the out-of-plane direction to avoid deviation of the total energy, atomic forces, and electrostatic potential.⁷² W is extracted from the following formula:

$$W = E_{\text{vac}} - E_{\text{F}} \quad (5)$$

where E_{vac} is the vacuum energy level and E_{F} refers to the Fermi level. W of metal electrodes are referred to experimental values.⁷³ The diffusion energy barriers for oxygen atom diffusion along certain paths over the MoS₂ surface are calculated though the CI-NEB method.⁷⁴ Part of data postprocessing is



conducted by the Vaspkit code, and Vesta software is used for atomic structure visualization.^{75,76}

Author contributions

Hejin Yan and Yongqing Cai: conceptualization, methodology, and writing – original draft. Hongfei Chen and Xiangyue Cui: software and validation. Qiye Guan: data curation and validation. Bowen Wang: visualization and investigation. Yongqing Cai: supervision. All authors participated in general discussion and manuscript revision.

Data availability

All relevant data are available from the authors on reasonable request and/or are included within the article and the ESI.†

Conflicts of interest

The authors declare that they have no conflict of interest.

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References

- A. Daus, S. Vaziri, V. Chen, Ç. Köroğlu, R. W. Grady, C. S. Bailey, H. R. Lee, K. Schauble, K. Brenner and E. Pop, *Nat. Electron.*, 2021, **4**, 495–501.
- J. Tang, Q. Wang, J. Tian, X. Li, N. Li, Y. Peng, X. Li, Y. Zhao, C. He and S. Wu, *Nat. Commun.*, 2023, **14**, 3633.
- A. Liu, X. Zhang, Z. Liu, Y. Li, X. Peng, X. Li, Y. Qin, C. Hu, Y. Qiu and H. Jiang, *Nano-Micro Lett.*, 2024, **16**, 119.
- S. Das, A. Sebastian, E. Pop, C. J. McClellan, A. D. Franklin, T. Grasser, T. Knobloch, Y. Illarionov, A. V. Penumatcha and J. Appenzeller, *Nat. Electron.*, 2021, **4**, 786–799.
- Y. Zou, P. Li, C. Su, J. Yan, H. Zhao, Z. Zhang and Z. You, *ACS Nano*, 2024, **18**, 9627–9635.
- S. Aftab, M. Z. Iqbal, S. Hussain, H. H. Hegazy, F. Kabir, S. H. A. Jaffery and G. Koyyada, *Chem. Eng. J.*, 2023, 144039.
- Y. Kim, I. Sohn, D. Shin, J. Yoo, S. Lee, H. Yoon, J. Park, S.-M. Chung and H. Kim, *Adv. Eng. Mater.*, 2024, **26**, 2301063.
- D. Panchal, A. Sharma and S. Pal, *Mater. Today Sustain.*, 2023, **21**, 100264.
- T. Wu, Y. Song, Z. Shi, D. Liu, S. Chen, C. Xiong and Q. Yang, *Nano Energy*, 2021, **80**, 105541.
- D. Guo, X.-x. Xue, M. Jiao, J. Liu, T. Wu, X. Ma, D. Lu, R. Zhang, S. Zhang, G. Shao and Z. Zhou, *Chem. Sci.*, 2024, **15**, 16281–16290.
- G. Shao, J. Xu, S. Gao, Z. Zhang, S. Liu, X. Zhang and Z. Zhou, *Carbon Energy*, 2024, **6**, e417.
- X. Huang, C. Liu and P. Zhou, *npj 2D Mater. Appl.*, 2022, **6**, 51.
- S. Aftab, H. H. Hegazy and F. Kabir, *Adv. Mater. Technol.*, 2023, **8**, 2201897.
- Y. Jing, X. Tan, Z. Zhou and P. Shen, *J. Mater. Chem. A*, 2014, **2**, 16892–16897.
- B. Radisavljevic and A. Kis, *Nat. Mater.*, 2013, **12**, 815–820.
- Y. Shen, Z. Dong, Y. Sun, H. Guo, F. Wu, X. Li, J. Tang, J. Liu, X. Wu and H. Tian, *Adv. Mater.*, 2022, **34**, 2201916.
- F. Wu, H. Tian, Y. Shen, Z. Hou, J. Ren, G. Gou, Y. Sun, Y. Yang and T.-L. Ren, *Nature*, 2022, **603**, 259–264.
- J. Xu, L. Chen, Y.-W. Dai, Q. Cao, Q.-Q. Sun, S.-J. Ding, H. Zhu and D. W. Zhang, *Sci. Adv.*, 2017, **3**, e1602246.
- K. Santosh, R. C. Longo, R. Addou, R. M. Wallace and K. Cho, *Nanotechnology*, 2014, **25**, 375703.
- J. Hong, Z. Hu, M. Probert, K. Li, D. Lv, X. Yang, L. Gu, N. Mao, Q. Feng and L. Xie, *Nat. Commun.*, 2015, **6**, 6293.
- C. Kim, I. Moon, D. Lee, M. S. Choi, F. Ahmed, S. Nam, Y. Cho, H.-J. Shin, S. Park and W. J. Yoo, *ACS Nano*, 2017, **11**, 1588–1596.
- J. Jang, J.-K. Kim, J. Shin, J. Kim, K.-Y. Baek, J. Park, S. Park, Y. D. Kim, S. S. Parkin and K. Kang, *Sci. Adv.*, 2022, **8**, eabn3181.
- M. Yarali, Y. Zhong, S. N. Reed, J. Wang, K. A. Ulman, D. J. Charboneau, J. B. Curley, D. J. Hynek, J. V. Pondick and S. Yazdani, *Adv. Electron. Mater.*, 2021, **7**, 2000873.
- E. Mitterreiter, B. Schuler, A. Micevic, D. Hernangómez-Pérez, K. Barthelmi, K. A. Cochran, J. Kiemle, F. Sigger, J. Klein and E. Wong, *Nat. Commun.*, 2021, **12**, 3822.
- X. Zhang, Z. Shao, X. Zhang, Y. He and J. Jie, *Adv. Mater.*, 2016, **28**, 10409–10442.
- H. Liu, M. Si, S. Najmaei, A. T. Neal, Y. Du, P. M. Ajayan, J. Lou and P. D. Ye, *Nano Lett.*, 2013, **13**, 2640–2646.
- Y.-H. Lee, X.-Q. Zhang, W. Zhang, M.-T. Chang, C.-T. Lin, K.-D. Chang, Y.-C. Yu, J. T.-W. Wang, C.-S. Chang and L.-J. Li, *Adv. Mater.*, 2012, **24**, 2320–2325.
- J. V. Pondick, J. M. Woods, J. Xing, Y. Zhou and J. J. Cha, *ACS Appl. Nano Mater.*, 2018, **1**, 5655–5661.
- X. Wang, H. Feng, Y. Wu and L. Jiao, *J. Am. Chem. Soc.*, 2013, **135**, 5304–5307.
- Y.-H. Lee, L. Yu, H. Wang, W. Fang, X. Ling, Y. Shi, C.-T. Lin, J.-K. Huang, M.-T. Chang and C.-S. Chang, *Nano Lett.*, 2013, **13**, 1852–1857.
- J.-K. Huang, Y. Wan, J. Shi, J. Zhang, Z. Wang, W. Wang, N. Yang, Y. Liu, C.-H. Lin and X. Guan, *Nature*, 2022, **605**, 262–267.
- Y. Y. Illarionov, T. Knobloch, M. Jech, M. Lanza, D. Akinwande, M. I. Vexler, T. Mueller, M. C. Lemme, G. Fiori and F. Schwierz, *Nat. Commun.*, 2020, **11**, 3385.
- C. Zhang, T. Tu, J. Wang, Y. Zhu, C. Tan, L. Chen, M. Wu, R. Zhu, Y. Liu and H. Fu, *Nat. Mater.*, 2023, **22**, 832–837.
- Y. Jin, J. Sun, L. Zhang, J. Yang, Y. Wu, B. You, X. Liu, K. Leng and S. Liu, *Adv. Mater.*, 2023, **35**, 2212079.
- Y. Jiang, Z. Liu, H. Zhou, A. Sharma, J. Deng and C. Ke, *Nanotechnology*, 2023, **34**, 405701.



- 36 M. M. Perera, M.-W. Lin, H.-J. Chuang, B. P. Chamlagain, C. Wang, X. Tan, M. M.-C. Cheng, D. Tománek and Z. Zhou, *ACS Nano*, 2013, 7, 4449–4458.
- 37 S. Kc, R. C. Longo, R. M. Wallace and K. Cho, *J. Appl. Phys.*, 2015, 117, 135301.
- 38 S. Wang, X. Zeng, Y. Zhou, J. Lu, Y. Hu, W. Wang, J. Wang, Y. Xiao, X. Wang and D. Chen, *ACS Appl. Electron. Mater.*, 2022, 4, 955–963.
- 39 H. Nam, S. Wi, H. Rokni, M. Chen, G. Priessnitz, W. Lu and X. Liang, *ACS Nano*, 2013, 7, 5870–5881.
- 40 Y. A. Kabachii, A. S. Golub, S. Y. Kochev, N. D. Lenenko, S. S. Abramchuk, M. Y. Antipin, P. M. Valetsky, B. D. Stein, W. E. Mahmoud and A. A. Al-Ghamdi, *Chem. Mater.*, 2013, 25, 2434–2440.
- 41 L. Wang, Z. Chen, C. R. Dean, T. Taniguchi, K. Watanabe, L. E. Brus and J. Hone, *ACS Nano*, 2012, 6, 9314–9319.
- 42 Q. He, Z. Zeng, Z. Yin, H. Li, S. Wu, X. Huang and H. Zhang, *Small*, 2012, 8, 2994–2999.
- 43 P.-C. Shen, Y. Lin, C. Su, C. McGahan, A.-Y. Lu, X. Ji, X. Wang, H. Wang, N. Mao and Y. Guo, *Nat. Electron.*, 2022, 5, 28–36.
- 44 I. Lee, M. Kang, S. Park, C. Park, H. Lee, S. Bae, H. Lim, S. Kim, W. Hong and S. Y. Choi, *Small*, 2024, 20, 2305143.
- 45 H. Shin, Y. Luo, A. Benali and Y. Kwon, *Phys. Rev. B*, 2019, 100, 075430.
- 46 A. Soman, R. A. Burke, Q. Li, M. D. Valentin, T. Li, D. Mao, M. Dubey and T. Gu, *ACS Appl. Mater. Interfaces*, 2020, 12, 37305–37312.
- 47 B. H. Kim, M. Park, M. Lee, S. J. Baek, H. Y. Jeong, M. Choi, S. J. Chang, W. G. Hong, T. K. Kim and H. R. Moon, *RSC Adv.*, 2013, 3, 18424–18429.
- 48 T. Jurca, M. J. Moody, A. Henning, J. D. Emery, B. Wang, J. M. Tan, T. L. Lohr, L. J. Lauhon and T. J. Marks, *Angew. Chem., Int. Ed.*, 2017, 56, 4991–4995.
- 49 C. Ataca and S. Ciraci, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2012, 85, 195410.
- 50 F.-X. R. Chen, N. Kawakami, C.-T. Lee, P.-Y. Shih, Z.-C. Wu, Y.-C. Yang, H.-W. Tu, W.-B. Jian, C. Hu and C.-L. Lin, *Appl. Phys. Lett.*, 2022, 121, 151601.
- 51 K. Burns, A. M. Z. Tan, H. Gordon, T. Wang, A. Gabriel, L. Shao, R. G. Hennig and A. Aitkaliyeva, *Phys. Rev. B*, 2020, 102, 085421.
- 52 K. S. Kumar, A. K. Dash, H. Sabreen H, M. Verma, V. Kumar, K. Watanabe, T. Taniguchi, G. S. Gautam and A. Singh, *2D Mater.*, 2024, 12, 045003.
- 53 A. Nourbakhsh, M. Cantoro, T. Vosch, G. Pourtois, F. Clemente, M. H. Van Der Veen, J. Hofkens, M. M. Heyns, S. De Gendt and B. F. Sels, *Nanotechnology*, 2010, 21, 435203.
- 54 Y. Cai, G. Zhang and Y.-W. Zhang, *J. Am. Chem. Soc.*, 2014, 136, 6269–6275.
- 55 B. Hou, Y. Zhang, H. Zhang, H. Shao, C. Ma, X. Zhang, Y. Chen, K. Xu, G. Ni and H. Zhu, *J. Phys. Chem. Lett.*, 2020, 11, 3116–3128.
- 56 Z. Wei, J. Tang, X. Li, Z. Chi, Y. Wang, Q. Wang, B. Han, N. Li, B. Huang and J. Li, *Small Methods*, 2021, 5, 2100091.
- 57 J. Tang, Z. Wei, Q. Wang, Y. Wang, B. Han, X. Li, B. Huang, M. Liao, J. Liu and N. Li, *Small*, 2020, 16, 2004276.
- 58 X. Zheng, A. Calò, E. Albisetti, X. Liu, A. S. M. Alharbi, G. Arefe, X. Liu, M. Spieser, W. J. Yoo and T. Taniguchi, *Nat. Electron.*, 2019, 2, 17–25.
- 59 X. Liu, M. S. Choi, E. Hwang, W. J. Yoo and J. Sun, *Adv. Mater.*, 2022, 34, 2108425.
- 60 X. Tong, E. Ashalley, F. Lin, H. Li and Z. M. Wang, *Nano-Micro Lett.*, 2015, 7, 203–218.
- 61 H. Gao, M. Hu, J. Ding, B. Xia, G. Yuan, H. Sun, Q. Xu, S. Zhao, Y. Jiang and H. Wu, *Adv. Funct. Mater.*, 2023, 33, 2213410.
- 62 Q. Guan, H. Yan and Y. Cai, *Adv. Mater. Interfaces*, 2022, 9, 2101838.
- 63 S. Gupta, A. Johnston and S. Khondaker, *J. Appl. Phys.*, 2022, 131, 164303.
- 64 A. Oberoi, Y. Han, S. P. Stepanoff, A. Pannone, Y. Sun, Y.-C. Lin, C. Chen, J. R. Shallenberger, D. Zhou and M. Terrones, *ACS Nano*, 2023, 17, 19709–19723.
- 65 S. Das, H.-Y. Chen, A. V. Penumatcha and J. Appenzeller, *Nano Lett.*, 2013, 13, 100–105.
- 66 S. Nagarajan, D. Hiller, I. Ratschinski, D. König, S. C. Smith, T. Mikolajick and J. Trommer, *Adv. Mater. Interfaces*, 2024, 11, 2300600.
- 67 D. Kiriya, M. Tosun, P. Zhao, J. S. Kang and A. Javey, *J. Am. Chem. Soc.*, 2014, 136, 7853–7856.
- 68 X. Tong, E. Ashalley, F. Lin, H. Li and Z. M. Wang, *Nano-Micro Lett.*, 2015, 7, 203–218.
- 69 G. Kresse and J. Furthmüller, *Comput. Mater. Sci.*, 1996, 6, 15–50.
- 70 G. Kresse and D. Joubert, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1999, 59, 1758.
- 71 S. Grimme, *J. Comput. Chem.*, 2006, 27, 1787–1799.
- 72 L. Bengtsson, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1999, 59, 12301.
- 73 W. M. Haynes, *CRC handbook of chemistry and physics*, CRC press, 2016.
- 74 G. Henkelman, B. P. Uberuaga and H. Jónsson, *J. Chem. Phys.*, 2000, 113, 9901–9904.
- 75 V. Wang, N. Xu, J.-C. Liu, G. Tang and W.-T. Geng, *Comput. Phys. Commun.*, 2021, 267, 108033.
- 76 K. Momma and F. Izumi, *J. Appl. Crystallogr.*, 2008, 41, 653–658.

