

Cite this: *Chem. Sci.*, 2018, 9, 4769

Transition metal atom doping of the basal plane of MoS₂ monolayer nanosheets for electrochemical hydrogen evolution†

Thomas H. M. Lau,^a XiaoWei Lu,^a Jiří Kulhavý,^a Simson Wu,^a Lilin Lu,^b Tai-Sing Wu,^c Ryuichi Kato,^d John S. Foord,^a Yun-Liang Soo,^c Kazu Suenaga^d and Shik Chi Edman Tsang^{id}*^a

Surface sites of extensively exposed basal planes of MoS₂ monolayer nanosheets, prepared via BuLi exfoliation of MoS₂, have been doped with transition metal atoms for the first time to produce 2D monolayer catalysts used for the electrochemical hydrogen evolution reaction (HER). Their HER activity is significantly higher than the corresponding thin and bulk MoS₂ layers. HAADF-STEM images show direct proof that single transition metal atoms reside at the surface basal sites, which subtly modify the electro-catalytic activity of the monolayer MoS₂, dependent on their electronic and stereospecific properties. It is found that these dopants play an important role in tuning the hydrogen adsorption enthalpies of the exposed surface S atoms and Mo atoms in HER. We report electrochemical testing, characterization and computational modelling and demonstrate that Co can significantly enhance the HER activity by the dominant Co–S interaction, whereas Ni substantially lowers the HER rate due to the Ni–Mo interaction at the same basal site. The two transition metal dopants show opposite doping behavior despite the fact that they are neighbors in the periodic table.

Received 9th March 2018

Accepted 29th April 2018

DOI: 10.1039/c8sc01114a

rsc.li/chemical-science

Introduction

There is an increasing demand for hydrogen to play a larger role in enabling the use of renewable energy sources (*i.e.* solar, wind, hydropower *etc.*) to reduce the carbon emissions from various industries in the near future. Particularly, the progressive decrease in the cost of electrolyzers and the possible implementation of carbon taxation may justify large scale H₂ production from the electrolysis of water in centralised installations. However, noble metals are commonly used as catalysts for both the anode and cathode in the electrolytic production of hydrogen from water. Their high cost is currently prohibitive in scaling up green hydrogen production. Therefore, developing inexpensive and Earth-abundant catalytic materials for the electrolytic hydrogen production reaction (HER) is of great interest. The discovery of graphene demonstrated that the material properties of an atomically thin material can be fairly

distinct from the bulk counterpart.¹ This has stimulated the extensive exploration of different two-dimensional (2D) materials. This research therefore stems from the current interest in layered materials, which represent a diverse and largely untapped source of 2D systems. As one of the emerging layered materials, 2D molybdenum disulphide (MoS₂) has drawn growing research attention in recent years due to its novel electronic, optical, mechanical, and electrochemical properties that are important for sensing,^{2,3} catalysis,^{4,5} and energy storage applications.^{6,7}

A single tri-atomic MoS₂ layer has a chemical structure of a hexagonal plane of molybdenum atoms sandwiched between two other hexagonal planes of sulphur atoms *via* strong in-plane covalent bonding. Adjacent layers are then held by weak out-of-plane van der Waals interactions.⁸ It is well accepted that the Mo edges (10 $\bar{1}$ 0) and the S edges ($\bar{1}$ 010) of the MoS₂ particles provide the two main active sites for most electrochemical reactions, including HER.⁹ They are preferentially exposed for reaction, which is one of the reasons why 2D MoS₂ shows overall better performance in electrochemical catalysis. In contrast, the basal plane (0001) has been verified to be electrochemically inert¹⁰ and thus is usually ignored as a contributor to the catalytic activity. In an attempt to enhance the activity of MoS₂ for HER, the general strategies are to increase the density of these active edge sites by tuning and exposing more edge planes to the surface^{11–14} or using doper promoters¹⁵ to improve the conductivity and binding energy to hydrogen of the active sites. It

^aDepartment of Chemistry, University of Oxford, Oxford, OX1 3QR, UK. E-mail: Edman.tsang@chem.ox.ac.uk

^bCollege of Chemistry and Chemical Engineering, Wuhan University of Science and Technology, China

^cDepartment of Physics, National Tsing Hua University, Hsinchu, Taiwan

^dNational Institute of Advanced Industrial Science and Technology (AIST), Central 5, 1-1-1 Higashi, Tsukuba, Ibaraki 305-8565, Japan

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8sc01114a



wasn't until recently that the presence of S vacancies as surface defects was shown to be able to activate the basal plane of MoS₂, which rendered the material highly active for hydrogen evolution reaction (HER).¹⁶ To further enhance the exposure of the basal planes, exfoliation of bulk MoS₂ may be performed to decrease the thickness of the planar layers by solvent exfoliation^{8,17,18} and Li exfoliation^{19,20} such that the interlayer van der Waals forces are partially or totally overcome, respectively. This opens up a new research area of studying and engineering the basal plane active sites of MoS₂.^{21–23}

There have been some testing and theoretical studies using bulk or thin MoS₂ layers with and without metal promotion for HER, but not much work on employing a single MoS₂ layer for modification at the atomic level for the HER reaction.^{24–28} Thus, the direct experimental proof of high quality samples with a single transition atom doped on monolayer MoS₂ and correlation to HER activity is not yet established, despite some excellent images of single atoms on thin MoS₂ layers.²⁸ Additionally, the fundamental modification mechanisms of the single transition metal dopants on monolayer MoS₂ for HER are not yet clear. Herein, we report our systematic experimental and modelling approaches to investigate high quality exfoliated single layer ⁵MoS₂ samples, with and without transition metal doping, as 2D catalysts for electrochemical hydrogen production, which we believe can offer comprehensive insight into the structure–activity relationship of the 2D monolayer type of catalyst. Particularly, the bonding sites of Co and Ni on the basal planes of the MoS₂ nanosheets (Co-⁵MoS₂ and Ni-⁵MoS₂) are directly imaged by HADDF-STEM and probed by XAS spectroscopy. It is evident that the use of different transition metal atoms as promoters can greatly influence the electronic properties and HER catalytic activity of the metal doped MoS₂ nanosheets due to the geometric characteristics and the characteristic binding affinities of the transition metals to the surface sites. Typically, it is found that Co displays a high affinity to bind on exposed surface S sites but is unable to bind on Mo sites (no Co–Mo interaction), whereas Ni can give rise to an Ni–Mo interaction due to stereo-specificity in the metal–metal bond formation. Density functional theory calculations were also employed to assess the thermodynamic favourability of the doping processes. It is believed that through this investigation, more insight can be provided into elucidating the effect of single atomic dopants on 2D layered materials to enable new catalytic or electrocatalytic reactions.

Results and discussion

Catalytic performance and electronic properties

The monolayer molybdenum disulfide nanosheets (⁵MoS₂) were prepared *via* lithium intercalation from a bulk molybdenum disulfide precursor.^{29,30} (Fig. 1, synthetic details are provided in the ESI†). The sheet-like morphology of exfoliated MoS₂ with regions of 1–3 molecular layers was characterized by high resolution transmission electron microscopes. Atomic force microscopy enabled the statistical analysis of 100 flakes produced by the lithium exfoliation method, which showed that 56% of the flakes were monolayer, 28% had two layers and 13% had three layers,

with a lower concentration of the thicker flakes. (ESI, Fig. S1 and S2†). Different single transition metal atom doped MoS₂ nanosheets (M-⁵MoS₂, where M = Fe, Co, Ni and Ag) were then synthesised from ⁵MoS₂ through a hydrothermal doping method. The details of the preparation and material characterization are summarized in S1 and S2 in the ESI.†

The electrochemical HER activity of the M-⁵MoS₂ nanosheets was measured with pre-calibration (ESI, Fig. S3a†) and is shown in Fig. 2a. From the linear sweep voltammogram (LSV), the ⁵MoS₂ nanosheets exhibit a distinctive HER catalytic activity when decorated with different single metal atoms. Among the tested samples, Co-⁵MoS₂ appears to significantly promote the activity of the ⁵MoS₂ nanosheets with the lowest onset potential of 220 mV at 10 mA cm⁻². Taking this sample as a reference, Co atom doping on ⁵MoS₂ nanosheets prepared by our Li intercalation route is significantly more active (lowest onset potential) than corresponding few layer ⁵MoS₂ and bulk MoS₂ samples (Fig. S3b†). This suggests that the activity is highly dependent on the quality of the MoS₂ used. Previous studies in the literature using thin or few MoS₂ layers with and without metal promotion for HER might have underestimated the metal promotion effect on single basal MoS₂ with defects on the terrace surface. Interestingly, other transition metal doped nanosheets follow the order Ag < Fe < Ni, and show a higher onset potential than pristine ⁵MoS₂ (300 mV at 10 mA cm⁻²). Typically, Ni-⁵MoS₂ has an onset potential of 353 mV at 10 mA cm⁻². The data clearly suggests that the choice of transition metal dopant greatly influences the HER catalytic performance of the basal ⁵MoS₂ materials. It is known that the effect of the metal dopant is to modify the H adsorption enthalpy of the surface edge S sites and Mo sites in HER.³¹ It would be interesting to see whether the basal sites can be modified by the metal dopants and their effect on the HER. In addition, Ni and Co are two neighbouring transition metal elements in the periodic table. One would expect their electronic effects to be very similar to each other.^{32,33} Clearly, our HER measurements



Fig. 1 Synthesis of single-layered MoS₂ (⁵MoS₂) and single transition-metal (TM) atom doped ⁵MoS₂/⁵MoS₂SO₄²⁻ groups.





Fig. 2 HER activity analysis by linear sweep voltammography, LSV. (a) LSV of the M-SMoS₂ nanosheets and the reference 20% Pt/C in 0.5 M H₂SO₄ at a scan rate of 2 mVs⁻¹. (b) Tafel plot of Co-SMoS₂, S-MoS₂, Ni-SMoS₂ and 20%Pt/C. (c) LSV of Co-SMoS₂ and Ni-SMoS₂ before and after 1000 repeat scans at a scan rate of 50 mV s⁻¹.

cannot be simply accounted for by a volcano plot,³⁴ indicating that the HER activity may also be affected by other factors.

To elucidate this relationship, Ni-SMoS₂ and Co-SMoS₂ were purposefully selected for comparison. Tafel analysis was performed on both materials to understand the inherent electrochemical HER reaction mechanism, as shown in Fig. 2b. The Tafel slopes of Ni-SMoS₂ and Co-SMoS₂ have close values of 103 mV dec⁻¹ and 92 mV dec⁻¹, respectively. Since both values are similar to that of pristine S-MoS₂ (94 mV dec⁻¹), this suggests that all three MoS₂ nanosheets follow the same HER reaction mechanism, *i.e.* the Volmer-Heyrovsky mechanism.³⁵ An electrochemical durability test was also carried out to analyse the doping stability of the two nanosheet samples. Fig. 2c shows a comparison of the catalytic performances of Co-SMoS₂ and Ni-SMoS₂ before and after 1000 scans. Both nanosheets retain

most of their HER activity, which indicates that the metal atoms were firmly attached onto the S-MoS₂.

Chemical and structural characterisation of M-SMoS₂

In general, doping metals on a MoS₂-based nanosheet increases the HER activity.^{35,36} For example, Ni nanoparticle doped MoS₂ films were found to perform (onset potential: 310 mV at 10 A cm⁻²) better than the pristine film (onset potential: 350 mV at 10 A cm⁻²).³⁷ However, the metal dopants are in the form of clusters or nanoparticles. They can give excellent electrocatalytic activity on their own while the 2D MoS₂ this time serves as a supporting substrate.

As seen from the TEM images (ESI[†]), we did not see any Co or Ni containing nanoparticles on the MoS₂ sheets. Inductively



Fig. 3 (a) and (b) HAADF-STEM images of Co-SMoS₂.²⁹ Simultaneous acquisition of (c) Co at the Mo-atop site model and (d) Co at the S vacancy site model (e) ADF and (f) EELS acquired along the line in (a). (g) ADF intensity line profiles taken along the numbered lines 1 and 2 shown in (b) ($\llcorner 110$) direction). The red arrows in the plot indicate sample drift during image acquisition.



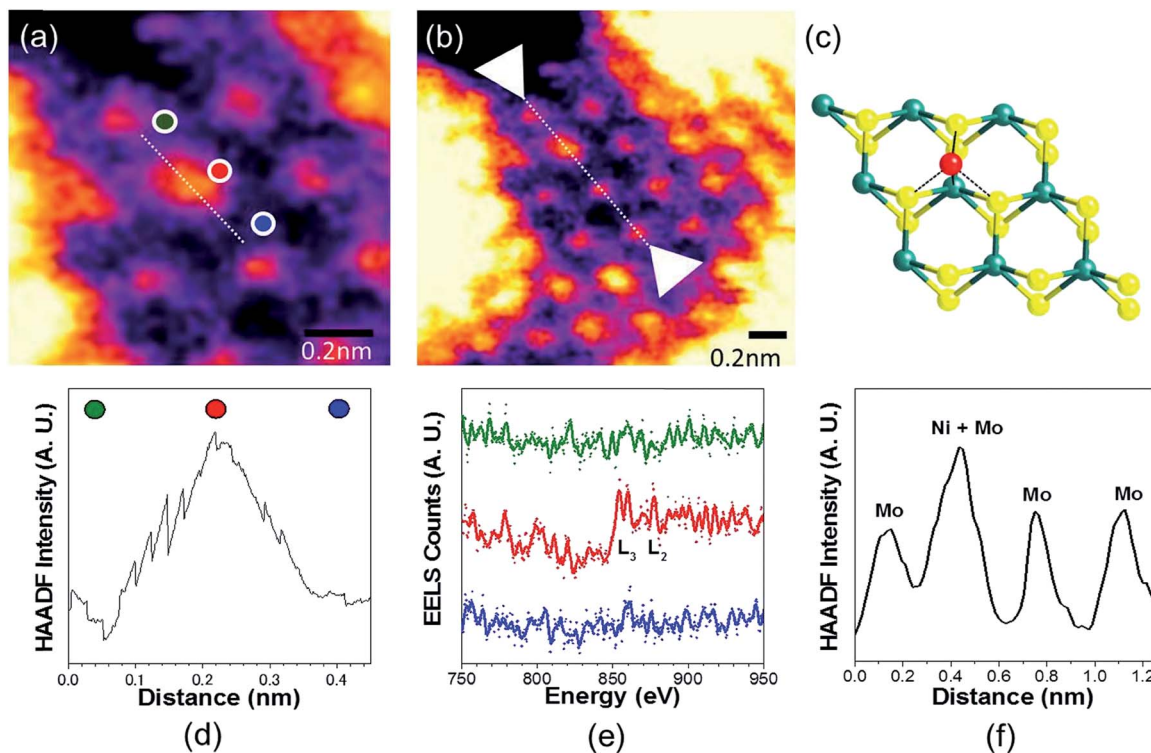


Fig. 4 (a) and (b) HAADF-STEM images of Ni-⁵MoS₂. (c) Ni at Mo-atop site model (d) ADF and (e) EELS acquired along the line in b. (f) ADF intensity line profile acquired along the line in b of the Mo sites in $\langle 100 \rangle$ direction.

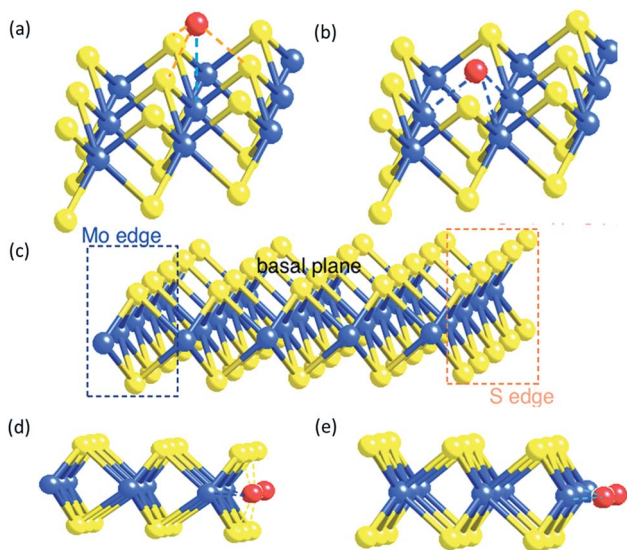


Fig. 5 (a) A metal atom on the Mo top site of ^MMoS₂ (b) a metal atom on the S vacancy site of ^MMoS₂ (c) the monolayer of perfect triatomic MoS₂ (d) metal doped S edge of ^MMoS₂ (e) metal-doped Mo edge of ^MMoS₂.

coupled plasma mass spectrometry (ICP-MS) was employed to analyse the concentration of the M-⁵MoS₂ surface (ESI, Table S1[†]). A typical result shows a Co metal content (mg mg⁻¹) of around 3.0 wt% and a corresponding calculated [Co/Mo] ratio of 11 can thus be deduced for M-⁵MoS₂. Thus, the hydrothermal

synthesis process employed using transition metal precursors in thiourea is proven to be successful in immobilizing metal dopants on the ⁵MoS₂ surface.

To analyse the specific location of the two transition metal Co and Ni dopants, which show contrasting electrocatalytic properties on the basal plane of ⁵MoS₂, High-Angle Annular Dark-Field (HAADF) imaging and Electron Energy Loss Spectroscopy (EELS) mapping of Co-⁵MoS₂ (ref. 29) (Fig. 3) and Ni-⁵MoS₂ (Fig. 4) were performed by High-Resolution Scanning Transmission Electron Microscopy (HR-STEM) and simulation (ESI, Fig. S4[†]). Both STEM analyses of the two samples show a characteristic and well-ordered hexagonal pattern of the honey-comb like structure which corresponds to monolayer ⁵MoS₂. Due to the Z contrast nature of ADF imaging, Mo atoms (blue, Fig. 3a) would appear to be brighter than S atoms (yellow, Fig. 3a). In Fig. 3b, at least two major types of Co dwelling sites are clearly observed. First, a very bright image feature appears on top of the Mo position sandwiched by S atoms (white arrow). This is attributed to the single Co atom sitting on top of the trigonal prismatic Mo atom (Mo atop site).²⁹ Similar features can also be observed at S positions (yellow arrow) (Fig. 3c and d), which can be attributed to the incorporation of a Co atom at the S vacant position (Co substituted in the S vacant site). For both samples, intensity profiles of the ADF image were taken across the doped sites in the $\langle 110 \rangle$ direction and the corresponding atomic composition of the peaks were assigned (Fig. 3e and f). ADF-EELS simultaneous acquisition line-scans were taken across a line containing either the single Co or Ni metal dopants



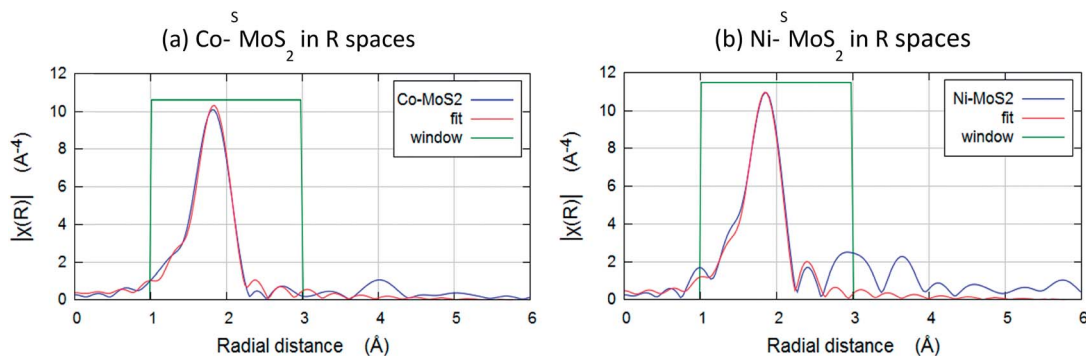


Fig. 6 Fourier transforms of the k^3 -weighted Co and Ni K-edge of extended X-ray absorption fine structure spectroscopy (EXAFS) spectra of (a) Co-SMoS_2 and (b) Ni-SMoS_2 .

Table 1 EXAFS scattering path analysis of Co-SMoS_2 and Ni-SMoS_2

Scattering path	Bond length (Å)	Coordination number	σ^2
$R = 0.9\%$, $K_{\text{wt}} = 1, 2, 3$; k range 3–12; R range 1–3; $E_{\text{not}} 0.23^a$			
Co-S	2.27 ± 0.01	3.9 ± 0.3	0.007
$R = 1.8\%$, $K_{\text{wt}} = 1, 2, 3$; k range 3–12; R range 1–3; $E_{\text{not}} 0.1^a$			
Ni-S	2.29 ± 0.01	6.5 ± 0.3	0.008
Ni-Mo	2.56 ± 0.04	1.2 ± 0.3	0.011

^a E_{not} is the difference in absorption energy between the experimental value and the calculated value.

(Co: Fig. 3e and f and Ni: Fig. 4d and e). For Co-SMoS_2 , the EELS extracted from before (green), on (dark blue), and after (khaki) the Co atom was doped on the Mo top site are shown in Fig. 3f. The characteristic $L_{3,2}$ edges occur at the expected energy in the EELS extracted from Co on top of the Mo site, and are absent in the two other EELS. Both atop (Fig. 4c, e and f) and substituted S vacant sites can also be observed for Ni-SMoS_2 (ESI, Fig. S5a and b†). The appearance of corresponding characteristic $L_{3,2}$ edges at the original Mo position (red circle), compared to before (green) and after (blue), confirms the presence of the Ni atom (Fig. 4e and f). This is further supported by the ADF analysis along the Mo sites in the $\langle 100 \rangle$ direction, where the larger peak (red) is contributed by Ni doping on top of the Mo atom (Mo atop site). Although we occasionally detected the presence of the two metals by EELS on the edges of layers (see ESI, Fig. S6a, b and S7a, b†), the large quantity and increasing number of basal sites with a significant increase in surface area compared to the peripheral edge sites of the reduced MoS_2 slab size due to exfoliation make the metal doped basal sites a more dominant feature. Thus, the imaging results provide direct experimental evidence of the successful doping of a Co atom and a Ni atom onto the basal planes of our exfoliated samples of SMoS_2 . Particularly, the chemisorption of the Co or Ni precursor directly on the abundant atop sites of the intact basal plane of SMoS_2 is envisaged to offer the accommodation of these transition metal atoms/ions in a significant quantity. It is also logical to assume that the Co or Ni precursor can occupy some surface S vacancies of SMoS_2 generated *via* the *n*BuLi exfoliation

to establish Co-Mo or Ni-Mo interactions with the exposed Mo sites on the ruptured tri-atomic MoS_2 layer. However, our STEM-EELS characterization so far was not able to offer any quantitative assessment of the main rooting sites for the two transition metals.

According to our molecular model of the single tri-atomic MoS_2 layer, metal M on the Mo atop site of the basal plane of SMoS_2 (Fig. 5a) exerts somewhat similar geometric characteristics to form M-S interactions to that of the M doped S edge (Fig. 5d), as described in the S edge model (Fig. 5c). Similarly, when M substitutes at the S vacant site of the partially damaged basal plane of SMoS_2 (Fig. 5b), M-Mo interactions can be formed. This is akin to that of the M-doped Mo edge structure (Fig. 5e). Thus, the S vacancies on the monolayer MoS_2 structure allow free access to the interior Mo sites modified by the chemisorbed M sulphide complex (akin to the Mo edge sites), which functions very similarly to the edge structure modification.

The chemical environments of Co and Ni were then examined by Extended X-ray Absorption Fine Structure Spectroscopy (EXAFS). Fig. 6a and b show the result of Fourier transforms of the Co and Ni K-edge of Co-SMoS_2 and Ni-SMoS_2 . All the parameters are fitted with acceptable Debye-Waller factors as listed in Table 1a and b. Both EXAFS results show there is no first-shell Co-Co or Ni-Ni contribution, suggesting that the major species of Co and Ni on the basal plane of the nanosheets are indeed not in the form of metal clusters nor nanoparticles but are single metal atoms/ions as suggested from the HADDF-STEM images (see Fig. 3 and 4). It is interesting to note that there is only a first shell Co-S contribution with a coordination number of about 4 (tetrahedron) at 2.27 ± 0.01 Å for Co-SMoS_2 and a first shell Ni-S contribution with a coordination number of about 6 (octahedron) at 2.29 ± 0.01 Å for Ni-SMoS_2 (Table 1). There is no first shell Co-Mo or Ni-Mo contribution, suggesting the majority of the Co and Ni species are indeed mainly on the atop sites (chemisorbed sites) of SMoS_2 , since doping of M (Co or Ni) on the S vacancies or Mo edge sites to a significant extent would generate the corresponding first shell M-Mo interaction. Part of the first shell sulphide ligands of these two immobilized metal sulphides must be derived from the surface sulphide groups, presumably from the higher degree of exposure of the





Fig. 7 Calculated Density of States (DoS) of (a) Mo and (b) S in $^{\text{S}}\text{MoS}_2$, $\text{Co-}^{\text{S}}\text{MoS}_2$ and $\text{Ni-}^{\text{S}}\text{MoS}_2$ (c) simplified molecular orbital diagram of MoS_2 (d) downshift of the conduction band (CB) upon addition of a single Co and Ni metal atom at the Mo atop site, respectively.

basal MoS_2 plane during the synthesis. The higher affinity for octahedral coordinated $3d^8 \text{Ni}^{2+}$ over tetrahedral coordinated $3d^7 \text{Co}^{2+}$ with sulphide ligands due to the greater ligand field stabilization energy accounts for the fundamental difference in coordination environment.³⁸ However, it is very interesting to find that the best fit data for $\text{Ni-}^{\text{S}}\text{MoS}_2$ depicts the presence of an Ni–Mo interaction at the second shell of $2.56 \pm 0.04 \text{ \AA}$ with CN of 1.2 ± 0.3 , whereas absolutely no Co–Mo interaction is detected in $\text{Co-}^{\text{S}}\text{MoS}_2$. Considering the larger size of Co (152 pm) and its tetrahedral coordination, the ‘Co–Mo’ distance between the shared triangular face with prismatic Mo (octahedron) is shorter than a typical Co–Mo bond from a simple geometric model. It therefore destabilizes the close face sharing for Co–Mo formation. On the other hand, the smaller size of Ni (149 pm) and its octahedral coordination can accommodate well for the stereospecific Ni–Mo interaction with the surface Mo site. The rigid close packed face-sharing model of chemisorbed transition metal sulphides can account for the absence of the Co–Mo interaction but facilitates the Ni–Mo interaction for the same atop Mo site.

Density functional theory calculations

The favourability of incorporating a single metal atom M onto $^{\text{S}}\text{MoS}_2$ is believed to cause the distinctive HER activity for electrochemical hydrogen production.

There have also been previous attempts to model the activity by doping single transition metal atoms onto the edge sites.^{39–41}

The Gibbs free energy of hydrogen adsorption (ΔG_{H}) is generally used as a descriptor for HER activity. It is believed that ΔG_{H} plays an important role in the electrochemical production of hydrogen gas: too weak H adsorption on the surface will not favour the electrochemical reaction and too strong H adsorption would also not facilitate H_2 recombination and desorption. Thus, this theoretical value should be close to 0 to maximise the overall thermodynamic enthalpy value for the H surface adsorption, surface recombination and desorption processes in HER as previously discussed.⁴¹

The relationship between the rate of HER and the doping effect of the metal atoms at the Mo atop site on the basal plane can be rationalised with the help of the calculated density of states (DoS) spectra of the three MoS_2 samples ($^{\text{S}}\text{MoS}_2$, $\text{Co-}^{\text{S}}\text{MoS}_2$, $\text{Ni-}^{\text{S}}\text{MoS}_2$) analysis (the Fermi energy level is arbitrarily set as 0, see ESI S1†). In general, the filled valence bands (VB) and the empty conduction bands (CB) in the DoS of MoS_2 are composed of both S-3p and Mo-4d states (Fig. 7a and b). From Fig. 2b, we understand that hydrogen generation for all three MoS_2 samples will progress through the Volmer–Heyrovsky mechanism. H^+ reduction, as the rate determining step, takes place by promoting electrons from the VB to the empty CB by electrical means. For unmodified MoS_2 , the H^+ reduction energy level (H^+/H_2) is close to the lower part of the CB, which is majorly occupied by the empty Mo- $4d_{x^2-y^2}$ and Mo- $4d_{z^2}$ bands (ESI Fig. S8a to f†). Therefore, hydrogen will be mainly generated from the Mo sites with a small contribution from the S sites



in pristine MoS₂, as also described by the edge model.³⁹ The value of the Gibbs free energy of hydrogen adsorption by the Mo component ($\Delta G_{\text{H-Mo}}$) in this case will be small and close to zero (Fig. 7d). It is expected that the addition of a 3d transition metal atom (Co and Ni) onto the Mo atop site will both cause a downshift on both the empty S-3p and Mo-4d CB bands (Fig. 7c). From our calculated DoS spectra, Co doping will induce a more remarkable downshift of the empty CB than that of Ni doping (Fig. 7a and b). Due to the absence of the Co–Mo interaction shown in our EXAFS results, it can be predicted that the H⁺ reduction energy level in Co–⁵MoS₂ will now be very close to the upper level of the empty CB occupied by S-3p, thus the value of the Gibbs free energy of hydrogen adsorption by the S component ($\Delta G_{\text{H-S}}$) is close to zero (Fig. 7d). H⁺ reduction therefore mainly takes place over the S sites. For Ni–⁵MoS₂, due to the smaller downshift of the CB as seen from our calculated bands (Fig. 7a and b), the H⁺ reduction level this time should be near the centre of the CB. While the contribution by both Mo and S to the overall H₂ production are similar but not close to their individual S or Mo component centres, they give a smaller hydrogen contribution from both sites (Fig. 7d). This above model is consistent with the computational analysis of the change in ΔG_{H} due to edge site modification by Wang *et al.*⁴¹ Overall, due to the different degree of downshifting on both the S-3p and Mo-4d components in the CB as compared to ⁵MoS₂, the Co atom selectively sitting at the Mo atop sites will enhance the HER activity while the Ni atom located on the same site with the stereospecifically allowed Ni–Mo interaction will show a much lower HER performance compared to undoped MoS₂.

Conclusions

In conclusion, we have provided direct experimental evidence that transition metal atoms can be chemically attached to multiple surface sites of basal plane of exfoliated ⁵MoS₂. From the HADDF-STEM image and the EXAFS analysis the data suggests that the majority of the metal dopants are attached to the Mo atop sites on the basal ⁵MoS₂ plane as chemisorbed sulphide complexes. For structural influence, it is difficult to use XRD to assess whether there is any change in the monolayer MoS₂ structure due to the doped atoms since our samples did not give good quality peaks for the structural analysis. Instead, our STEM images, EXAFS analysis and Raman spectra (given in the revised ESI[†]) suggested that there was no significant change in the crystallographic parameters of the underneath monolayer of MoS₂ (*i.e.* the Mo–S, Mo–Mo, and S–S distances). However, the anchored tetrahedral coordinated Co can affect the surface neighbour S sites through a Co–S interaction on the basal plane in attenuating ΔG_{H} towards a value of zero at high hydrogen coverage hence enhancing the HER activity. On the other hand, the geometric constraints of the large sized Co atom in the tetrahedral arrangement deny the direct influence of the Mo site on ⁵MoS₂ with no Co–Mo interaction. In contrast, the chemisorbed octahedral Ni sulphide on the same atop site allows the direct electronic modification of the Mo site by establishing a Ni–Mo interaction, which causes the deviation of ΔG_{H} from zero at the surface exposed Mo sites at high hydrogen

coverage hence greatly attenuating the HER activity. As seen from the DFT calculations, we have also attributed their difference in HER activity to the electronic modifications of the MoS₂ structure by the doped atoms/ions. Thus, both the structural and electronic factors are expected to influence their resulting activity over the single atom doped monolayer MoS₂ samples. It is believed that both the geometric and electronic factors exerted by the transition metal dopants are therefore important parameters in further tuning the 2D MoS₂ structure for the rational design of composite materials for more efficient electrochemical hydrogen production from water.

Contributions

THML, XL and JK prepared the samples and carried out testing; SW, TW and YS worked on the XAS (EXAFS); RK and KS did the STEM and EELS; LL carried out the DFT calculations; JSF advised on the electrochemical measurements; THML and SCET wrote the paper, SCET planned and supervised this project. All contributed toward the preparation of the manuscript.

Conflicts of interest

There is no conflict of interest to declare.

Acknowledgements

The financial support of this project from the EPSRC research council of the UK is acknowledged. The authors wish to thank the National Synchrotron Radiation Center, Hsinchu, Taiwan for access to EXAFS facilities. RK and KS acknowledge JSPS KAKENHI (JP16H06333) for support. The authors also acknowledge the use of the Computing Facilities of Wuhan University of Science and Technology in the completion of the theoretical part of this work and support from National Natural Science Foundation of China (No. 21671154).

Notes and references

- 1 J. C. Meyer, A. K. Geim, M. I. Katsnelson, K. S. Novoselov, T. J. Booth and S. Roth, *Nature*, 2007, **446**, 60–63.
- 2 F. K. Perkins, A. L. Friedman, E. Cobas, P. M. Campbell, G. G. Jernigan and B. T. Jonker, *Nano Lett.*, 2013, **13**, 668–673.
- 3 C. Zhu, Z. Zeng, H. Li, F. Li, C. Fan and H. Zhang, *J. Am. Chem. Soc.*, 2013, **135**, 5998–6001.
- 4 F. M. Pesci, M. S. Sokolikova, C. Grotta, P. C. Sherrell, F. Reale, K. Sharda, N. Ni, P. Palczynski and C. Mattevi, *ACS Catal.*, 2017, **7**, 4990–4998.
- 5 T. Jia, A. Kolpin, C. Ma, R. C.-T. Chan, W.-M. Kwok and S. C. E. Tsang, *Chem. Commun.*, 2014, **50**, 1185–1188.
- 6 H. Hwang, H. Kim and J. Cho, *Nano Lett.*, 2011, **11**, 4826–4830.
- 7 J. Xiao, D. Choi, L. Cosimbescu, P. Koech, J. Liu and J. P. Lemmon, *Chem. Mater.*, 2010, **22**, 4522–4524.
- 8 Q. H. Wang, K. Kalantar-Zadeh, A. Kis, J. N. Coleman and M. S. Strano, *Nat. Nanotechnol.*, 2012, **7**, 699–712.



- 9 T. F. Jaramillo, K. P. Jørgensen, J. Bonde, J. H. Nielsen, S. Horch and I. Chorkendorff, *Science*, 2007, **317**, 100–102.
- 10 H. Tang and S. Roy Morrison, *Thin Solid Films*, 1993, **227**, 90–94.
- 11 J. Kibsgaard, Z. Chen, B. N. Reinecke and T. F. Jaramillo, *Nat. Mater.*, 2012, **11**, 963–969.
- 12 S. Li, S. Wang, M. M. Salamone, A. W. Robertson, S. Nayak, H. Kim, S. C. E. Tsang, M. Pasta and J. H. Warner, *ACS Catal.*, 2017, **7**, 877–886.
- 13 G. Ye, Y. Gong, J. Lin, B. Li, Y. He, S. T. Pantelides, W. Zhou, R. Vajtai and P. M. Ajayan, *Nano Lett.*, 2016, **16**, 1097–1103.
- 14 C. Tsai, F. Abild-Pedersen and J. K. Nørskov, *Nano Lett.*, 2014, **14**, 1381–1387.
- 15 Z. He and W. Que, *Applied Materials Today*, 2016, **3**, 23–56.
- 16 M. Chhowalla, H. S. Shin, G. Eda, L.-J. Li, K. P. Loh and H. Zhang, *Nat. Chem.*, 2013, **5**, 263–275.
- 17 G. Cunningham, M. Lotya, C. S. Cucinotta, S. Sanvito, S. D. Bergin, R. Menzel, M. S. P. Shaffer and J. N. Coleman, *ACS Nano*, 2012, **6**, 3468–3480.
- 18 R. Bhandavat, L. David and G. Singh, *J. Phys. Chem. Lett.*, 2012, **3**, 1523–1530.
- 19 G. Eda, H. Yamaguchi, D. Voiry, T. Fujita, M. Chen and M. Chhowalla, *Nano Lett.*, 2011, **11**, 5111–5116.
- 20 H. Li, M. Du, M. J. Mleczko, A. L. Koh, Y. Nishi, E. Pop, A. J. Bard and X. Zheng, *J. Am. Chem. Soc.*, 2016, **138**, 5123–5129.
- 21 Y. Xu, L. Wang, X. Liu, S. Zhang, C. Liu, D. Yan, Y. Zeng, Y. Pei, Y. Liu and S. Luo, *J. Mater. Chem. A*, 2016, **4**, 16524–16530.
- 22 X. Chen, N. C. Berner, C. Backes, G. S. Duesberg and A. R. McDonald, *Angew. Chem. Int. Ed.*, 2016, **55**, 5803–5808.
- 23 Y. Yin, J. Han, Y. Zhang, X. Zhang, P. Xu, Q. Yuan, L. Samad, X. Wang, Y. Wang, Z. Zhang, P. Zhang, X. Cao, B. Song and S. Jin, *J. Am. Chem. Soc.*, 2016, **138**, 7965–7972.
- 24 Y. Xu, L. Wang, X. Liu, S. Zhang, C. Liu, D. Yan, Y. Zeng, Y. Pei, Y. Liu and S. Luo, *J. Mater. Chem. A*, 2016, **4**, 16524–16530.
- 25 Y. Shi, Y. Zhou, D.-R. Yang, W.-X. Xu, C. Wang, F.-B. Wang, J.-J. Xu, X.-H. Xia and H.-Y. Chen, *J. Am. Chem. Soc.*, 2017, **139**(43), 15479–15485.
- 26 H. Li, L. Wang, Y. Dai, Z. Pu, Z. Lao, Y. Chen, M. Wang, X. Zheng, J. Zhu, W. Zhang, R. Si, C. Ma and J. Zeng, *Nat. Nanotechnol.*, 2018, **13**, 411–417.
- 27 Y. Yan, B. Xia, X. Ge, Z. Liu, J. Y. Wang and X. Wang, *ACS Appl. Mater. Interfaces*, 2013, **5**, 12794–12798.
- 28 Y. Shi, J. Wang, C. Wang, T.-T. Zhai, T.-J. Bao, J.-J. Xu, X.-H. Xia, H.-Y. Chen and J. Am, *Chem. Soc.*, 2015, **137**, 7365–7370.
- 29 G. 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.
- 30 L. Coulier, V. H. J. de Beer, J. A. R. van Veen and J. W. Niemantsverdriet, *Top. Catal.*, 2000, **13**, 99–108.
- 31 J. Kibsgaard, A. Tuxen, K. G. Knudsen, M. Brorson, H. Topsøe, E. Lægsgaard, J. V. Lauritsen and F. Besenbacher, *J. Catal.*, 2010, **272**, 195–203.
- 32 P. B. Armentrout, L. F. Halle and J. L. Beauchamp, *J. Am. Chem. Soc.*, 1981, **103**, 6501–6502.
- 33 J. A. M. Simões and J. L. Beauchamp, *Chem. Rev.*, 1990, **90**, 629–688.
- 34 S. Trasatti, *J. Electroanal. Chem.*, 1972, **39**, 163–184.
- 35 Y. Shi, J.-K. Huang, L. Jin, Y.-T. Hsu, S. F. Yu, L.-J. Li and H. Y. Yang, *Sci. Rep.*, 2013, **3**, 1839.
- 36 D. Merki, H. Vrubel, L. Rovelli, S. Fierro and X. Hu, *Chem. Sci.*, 2012, **3**, 2515.
- 37 D. Wang, X. Zhang, Y. Shen and Z. Wu, *RSC Adv.*, 2016, **6**, 16656–16661.
- 38 U. Müller, in *Inorganic Structural Chemistry*, John Wiley & Sons, Ltd, 2007, pp. 73–84.
- 39 J. V. Lauritsen, J. Kibsgaard, G. H. Olesen, P. G. Moses, B. Hinnemann, S. Helveg, J. K. Nørskov, B. S. Clausen, H. Topsøe, E. Lægsgaard and F. Besenbacher, *J. Catal.*, 2007, **249**, 220–233.
- 40 H. Schweiger, P. Raybaud and H. Toulhoat, *J. Catal.*, 2002, **212**, 33–38.
- 41 H. Wang, C. Tsai, D. Kong, K. Chan, F. Abild-Pedersen, J. K. Nørskov and Y. Cui, *Nano Res.*, 2015, **8**, 566–575.

