

## MINIREVIEW

[View Article Online](#)  
[View Journal](#) | [View Issue](#)Cite this: *Nanoscale*, 2020, **12**, 10447

## 2D molybdenum disulphide nanosheets incorporated with single heteroatoms for the electrochemical hydrogen evolution reaction

Thomas H. M. Lau, John S. Foord and S. C. Edman Tsang \*

2D nanosheets give enhanced surface area to volume ratios in particle morphology and they can also provide defined surface sites to disperse foreign atoms. Placing atoms of catalytic interest on 2D nanosheets as Single Atom Catalysts (SAC) represents one of the novel approaches due to their unique but tunable electronic and steric characteristics. Here in this mini-review, we particularly highlight some recent and important developments on heteroatom doped MoS<sub>2</sub> nanosheets (SAC-MoS<sub>2</sub>) as catalysts for the electrochemical hydrogen evolution reaction (HER) from water, which could lead to opening up to a flagship of important renewable technologies in future. It is shown that the nature of dopants, doping positions and the polytypes of MoS<sub>2</sub> nanosheets are the determining factors in the overall catalytic abilities of these functionalised nanosheets. This may serve to obtain atomic models which lead to further understanding of the 'metal-support interaction' in catalysis.

Received 15th February 2020,  
Accepted 15th April 2020  
DOI: 10.1039/d0nr01295e  
[rsc.li/nanoscale](http://rsc.li/nanoscale)

### 1. Introduction

The subtle climate change due to excessive carbon emissions to the atmosphere has prompted our community to switch energy sources from oil, gas and coal to renewables such as solar and wind power, tidal waves, *etc.* Green H<sub>2</sub> manufacture from electrolysis of water over an efficient electrocatalyst is the core technology to harness these renewables. The produced H<sub>2</sub> with zero carbon emission can be directly used in combustion engines, and fuel cells, for reduction of CO<sub>2</sub>, *etc.* Alternatively, it can be stored as chemicals such as ammonia or methanol to facilitate transport and for use in other areas. Although noble metal catalysts give satisfactory performance in electrolyzers, their high cost and scarcity have compelled chemical manufacturers to find greener and more cost-effective alternatives. Thanks to the advances in nanotechnology, two-dimensional (2D) metallic or semi-conductive materials such as graphite based nanostructures,<sup>1,2</sup> transition metal dichalcogenides (TMDC),<sup>3–5</sup> transition metal oxides (TMO),<sup>6,7</sup> and transition metal nitrides (TMN)<sup>8,9</sup> have been thoroughly studied and characterised. These atomically thick 2D nanosheets possess a much higher surface-to-volume-ratio than three-dimensional (3D) nanomaterials. Since more active sites located at the surface can be accessed for the desired reactions, they generally exhibit better catalytic performances than their 3D

counterparts. However, they are still incomparable to conventional noble metal catalysts such as Pt and Ru. Thus, further modifications have been attempted to improve the material properties of these 2D catalysts. The general idea is to either increase the number of active sites available or enhance the intrinsic activity of each active site. For example, the former approach can be achieved by exposing more edge sites to terrace sites by introducing surface defects,<sup>10–14</sup> while the latter one can involve changing the electronic structure of the responsible atoms at the active sites *via* strain engineering.<sup>15–17</sup>

One of the newest modification methods to functionalise 2D nanosheets is to incorporate single heteroatoms onto the large basal surface of 2D catalysts. Generally, single-atom catalysts (SAC) refers to foreign single atoms deposit on support materials, the atomic dopants act as the active sites and contribute primarily towards the catalytic activities in the reactions of interest.<sup>18–20</sup> The dispersed atomic dopants maximise metal utilisation, which makes the catalyst excel in small molecule activation. Such a combination of SAC and 2D catalysts creates a new group of materials, which is multifunctional in catalysis due to the coexistence of two types of active sites on the surface, potentially making more complicated reaction mechanisms plausible. While this new type of catalyst generates much interest as it combines the knowledge of the two hottest research areas in catalysis, the research field has not yet been well explored. Several excellent review articles have been published to highlight some recent research work in the area of SAC.<sup>21–23</sup> Similarly, there are some good reviews focus-

Department of Chemistry, University of Oxford, Oxford, OX1 3QR, UK.  
E-mail: [edman.tsang@chem.ox.ac.uk](mailto:edman.tsang@chem.ox.ac.uk)

ing on using 2-D materials such as graphene, molybdenum disulphide ( $\text{MoS}_2$ ), graphitic carbon nitride ( $g\text{-C}_3\text{N}_4$ ), *etc.* as catalysts in electrochemical reactions including the HER when doped with heteroatoms. The challenges of synthesising and analysing SAC-2D catalysts have also been reviewed therein. The readers should refer to these articles for reference without being repeatedly discussed in this article. In an attempt to provide insights for future research in the field of SAC and 2D materials in combination, this mini-review will particularly examine some selected key publications on single atom doped  $\text{MoS}_2$  nanosheets (SAC- $\text{MoS}_2$ ) in the HER as a short but sharply focused review. The synthetic approaches and the electrocatalytic activities of these materials will be thoroughly analysed with the aim to better understand the structure-activity relationship of SAC- $\text{MoS}_2$ .

## 2. Polytypes of 2D $\text{MoS}_2$

$\text{MoS}_2$  is a layered structure with each triatomic monolayer consisting 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 stacked together by weak out-of-plane van der Waals interactions.<sup>24</sup> There are multiple polytypes of  $\text{MoS}_2$ , 2H- $\text{MoS}_2$ , 3R- $\text{MoS}_2$ , and 1T- $\text{MoS}_2$ , which give distinctive electronic properties. The digit refers to the number of layers in the unit cell while the letters “H”, “R”, and “T” indicate hexagonal ( $D_{3h}$  group), rhombohedral ( $C_{3v}$  group), and tetragonal ( $D_{3d}$  group) symmetry, respectively. Both 2H- $\text{MoS}_2$  and 3R- $\text{MoS}_2$  are semiconducting while 1T- $\text{MoS}_2$  displays a metallic behaviour. 1T- $\text{MoS}_2$  is found to possess an electrical conductivity of around  $10^7$  times higher than that of 2H- $\text{MoS}_2$ .<sup>32</sup> If 2H- and 3R- $\text{MoS}_2$  crystals are exfoliated into monolayers, the polytype will then be referred to as 1H to denote a monolayer structure. 1T- $\text{MoS}_2$  is metastable as its unit cell is the distorted form of 2H- $\text{MoS}_2$ . Thus, it needs to be stabilised or else it will be converted back to 2H- $\text{MoS}_2$  upon heating above 95 °C.<sup>33</sup> For

example, computational calculations suggested that the 1T polytype can be distorted into a lower energy 1T' structure due to the stabilisation of hydrogen adsorption in the HER.<sup>34,35</sup>

The choice of the  $\text{MoS}_2$  polytype not only affects the overall electrical conductivity but also the density state of the active sites. During catalysis, the Mo edges (1010) and the S edges (1010) of  $\text{MoS}_2$  can accept and donate electrons like a metal surface.<sup>36</sup> The contribution by the edge sites however is far less important for 2D  $\text{MoS}_2$  due to the presence of a larger basal surface. The basal plane (0001) of semiconducting  $\text{MoS}_2$  has been verified to be chemically inert both experimentally<sup>15</sup> and computationally<sup>37</sup> as further modification is required to activate it. In contrast, the metallic feature of 1T- $\text{MoS}_2$  has made the basal sites highly active for electrocatalysis.<sup>38,39</sup> Top-down and bottom-up methods are two major approaches to obtain 2D  $\text{MoS}_2$  nanosheets. The former method is to isolate monolayers from bulk  $\text{MoS}_2$  crystals by weakening the inter-layer van der Waals interactions. Lithium intercalation is one example to create monolayer 1T- $\text{MoS}_2$  with rich S-defects.<sup>40–42</sup> The latter method involves reacting Mo precursors with S sources to grow  $\text{MoS}_2$  unit by unit. Chemical vapour deposition (CVD)<sup>43–46</sup> and one-pot hydrothermal/solvothermal synthesis<sup>47–49</sup> are often used to grow high-quality ultrathin 2H- $\text{MoS}_2$  or monolayer 1H- $\text{MoS}_2$ .

## 3. Synthetic methods of SAC- $\text{MoS}_2$

Table 1 lists the synthetic approaches of selected publications that were adopted in preparing different SAC- $\text{MoS}_2$  for the electrochemical HER. It has been shown that the synthesis of SAC- $\text{MoS}_2$  nanosheets is predominantly achieved by two different synthetic routes: (1) incorporating single heteroatoms onto the as-prepared 2D  $\text{MoS}_2$  nanosheets with the desired polytypes and (2) directly growing the catalysts through the hydrothermal/solvothermal reaction. The synthetic route will largely influence the doping position of the heteroatoms adopted in the SAC- $\text{MoS}_2$  catalysts. It was revealed that there

**Table 1** Highlighted publications showing synthetic methods, polytypes of 2D  $\text{MoS}_2$  nanosheets, metal doping concentrations and their preferable doping position(s)

Entry	Catalyst	Synthetic method	Polytype	Doping concentration and preferable doping position	Ref.
1A	$\text{MoS}_2$	Mechanical exfoliation under sonication	2H	—	25
1B	$\text{MoS}_2$	Lithium intercalation	1T	—	25
2A	Co- $\text{MoS}_2$	One pot solvothermal reaction at 400 °C, 4 h	2H	1.7% Co, Mo substitution sites at edge sites	26
2B	Pt- $\text{MoS}_2$	One pot solvothermal reaction at 400 °C, 4 h	2H	1.7% Pt, Mo substitution sites at edge sites	26
3	Co- $\text{MoS}_2$ with S defects	Lithium intercalation, hydrothermal doping at 180 °C, 12 h	1H	3% Co, Mo atop sites at the basal plane	27
4	Pd- $\text{MoS}_2$	One pot solvothermal reaction at 180 °C, 24 h	1T	1% Pd, Mo substitution sites at the basal plane	28
5A	Pt- $\text{MoS}_2$	Lithium intercalation, sonochemical doping at room temperature, 24 h	1T	3% Pt, Mo atop sites at the basal plane	29
5B	Pd- $\text{MoS}_2$	Lithium intercalation, sonochemical doping at room temperature, 24 h	1T	3% Pd, Mo atop sites at the basal plane	29
6	P- $\text{MoS}_2$	One pot solvothermal reaction at 200 °C, 24 h	1T	5% P, S substitution at the basal plane	30
7	N- $\text{MoS}_2$	One pot solvothermal reaction at 200 °C, 24 h	1T	5.7% N, Mo substitution sites at the basal plane	31

are four general doping sites for single molecular layer  $\text{MoS}_2$ , namely the Mo atop site, S substitution site, Mo substitution site, and hollow site.<sup>50</sup> For the first route,  $\text{MoS}_2$  nanosheets are first synthesised beforehand usually by lithium intercalation (Method 1) or by one-pot synthesis (Method 2). As mentioned, lithium intercalation can be applied to exfoliate bulk  $\text{MoS}_2$  into 1T- $\text{MoS}_2$ . However, a prolonged heating above the 95 °C threshold using hydrothermal techniques on non-stabilised 1T- $\text{MoS}_2$  would reconvert the structure from metallic 1T- to semiconducting 2H- $\text{MoS}_2$ . For the one-pot hydrothermal/solvothermal synthesis, nanosheets are being built-up unit by unit through reactions between the Mo and S precursors. The heteroatoms are then incorporated onto the as-prepared nanosheets afterwards. Since the six-coordinated Mo atoms are well located inside the intact  $\text{MoS}_2$  unit cells, the incoming heteroatom dopants during the incorporation process are less likely to penetrate into the outer S layer and substitute the Mo atom. Therefore, they are placed at the Mo atop sites or S substitution sites unless extra treatment such as high temperature annealing is applied.

For the second route, the one pot bottom-up synthetic approach can also be adopted to directly synthesise SAC- $\text{MoS}_2$  through the reactions between Mo, S, and the dopant precursors. The heteroatoms can be embedded inside the  $\text{MoS}_2$  nanosheets as a result due to their presence during the unit by unit construction of  $\text{MoS}_2$  nanosheets. However, it should be noted that a precise control of the molar concentration for the three precursors is vital for obtaining monolayer nanosheets

confined with single heteroatoms within the tolerance of the thermodynamic maximum limits. An excess amount of single metal atom dopants may result in the formation of clusters or nanoparticles<sup>28</sup> while an insufficient amount of Mo or S sources can lead to the creation of surface defects.<sup>30</sup>

Several selected publications listed in Table 1 (entries 1, 3 and 5) adopted the first route to synthesise SAC- $\text{MoS}_2$  (Fig. 1, Method 1). Lithium intercalation was applied to obtain 1T- $\text{MoS}_2$  nanosheets. The delicate 1T polytype was then preserved when the as-prepared nanosheets underwent doping in a sonochemical reaction under ambient conditions to prevent the conversion back to 2H.<sup>29</sup> The same hydrothermal/solvothermal techniques are also being applied in the one pot synthesis of SAC- $\text{MoS}_2$  (Table 1, entries 2, 3, 4, 6 and 7) (Fig. 1, Method 3). While the reaction temperature in all cases is at least above 180 °C, the metal dopants embedded into the as-grown  $\text{MoS}_2$  can provide extra kinetic stability to prevent the conversion of 1T polytypes. However, this only applies to a certain extent as the  $\text{MoS}_2$  nanosheets are grown at a high temperature of 400 °C, which will still drive to the 2H polytype with a multiple-layer thickness. Therefore, the synthetic condition also plays a crucial role in determining the resulting polytypes. Overall, the doping concentration, doping position, and polytypes largely depend on the synthetic route and condition. Since there are pros and cons for each discussed synthetic method, the choice should depend on the structure of the SAC- $\text{MoS}_2$  material desired as it ultimately determines the catalytic performance.



**Fig. 1** Summary of the synthetic routes of SAC- $\text{MoS}_2$  adopted in recent highlighted publications. Method 1 uses the top-down exfoliation of bulk  $\text{MoS}_2$  to create  $\text{MoS}_2$  nanosheets, while method 2 uses the bottom-up hydrothermal/solvothermal reactions to make the nanosheets directly from Mo and S sources. The pure  $\text{MoS}_2$  created with method 1 and method 2 will then be doped with the heteroatoms to form SAC- $\text{MoS}_2$ . Method 3 uses one-pot synthesis of the SAC- $\text{MoS}_2$  via hydrothermal/solvothermal reactions. For all the  $\text{MoS}_2$  structural models shown in the above figure, the green, yellow, and red spheres represent Mo atoms, S atoms, and dopant atoms, respectively.

#### 4. Catalytic performance of SAC-MoS<sub>2</sub>

The reaction then proceeds with either the Tafel step or the Heyrovsky step, which is usually dependent on the amount of

To understand the effect of single metal dopants on MoS<sub>2</sub> for the HER, Deng *et al.* were among the pioneer groups to conduct electrochemical scan tests of 2H-MoS<sub>2</sub> doped with several types of single transition metal atoms (Pt, Co, and Ni) at Mo substitution sites for the HER (Table 2, entry 2).<sup>26</sup>

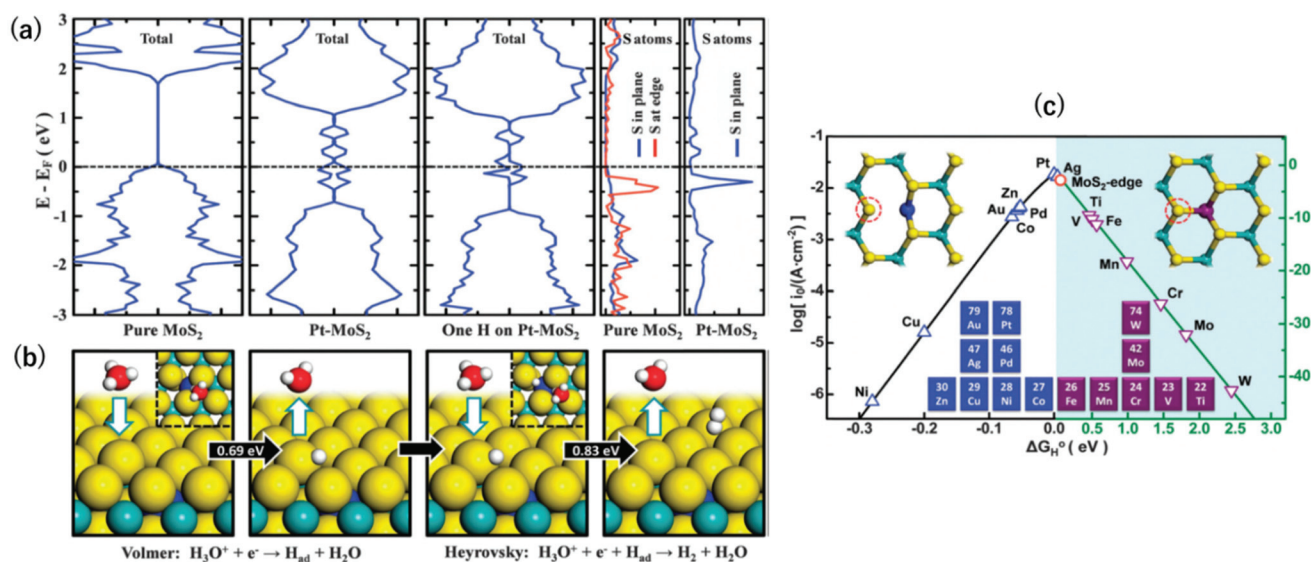


Fig. 2 (a) DOS of MoS<sub>2</sub>, Pt-MoS<sub>2</sub>, and H adsorbed Pt-MoS<sub>2</sub>. The dashed line represents the Fermi level. (b) HER process through the Volmer-Heyrovsky pathway on a Pt-MoS<sub>2</sub> catalyst. (c) Volcano plot of 2H-MoS<sub>2</sub> doped with different transition metals at the Mo substitution site.<sup>26</sup>



**Table 2** Electrocatalytic production of H<sub>2</sub> over various MoS<sub>2</sub> systems

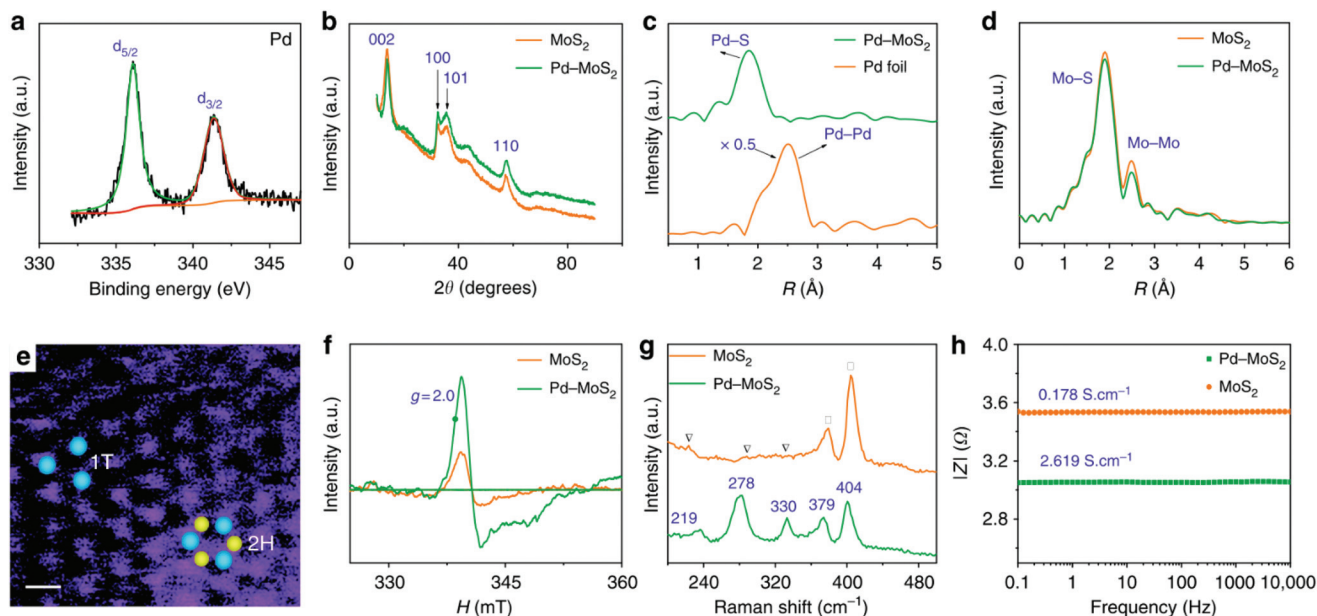
Entry	Catalyst	Overpotential ( $\eta$ ) (mV@10 mA cm <sup>-2</sup> ) vs. RHE	Tafel slope (mV per decade)	Ref.
1A	MoS <sub>2</sub>	~-375 mV	186	25
1B	MoS <sub>2</sub>	~-250 mV	40	25
2A	Co-MoS <sub>2</sub>	~-285 mV	N/A	26
2B	Pt-MoS <sub>2</sub>	~-185 mV	96	26
3	Co-MoS <sub>2</sub> with S defects	-300 mV	92	27
4	Pd-MoS <sub>2</sub>	-89 mV	62	28
5A	Pt-MoS <sub>2</sub>	-223 mV	57	29
5B	Pd-MoS <sub>2</sub>	-140 mV	50	29
6	P-MoS <sub>2</sub>	-43 mV	34	30
7	N-MoS <sub>2</sub>	-168 mV	41	31

According to the experimental results, the Pt doped one exhibited the best catalytic performance among three with the smallest overpotential of -185 mV at 10 mA cm<sup>-2</sup>. The Density of States (DOS) simulation provided an explanation that the introduction of Pt atoms into the Mo substitution site induced more hybridized electronic states to occur near the Fermi level, which enhanced the H adsorption ability of basal S atoms and thus the catalytic performance for the HER (Fig. 2a and b).

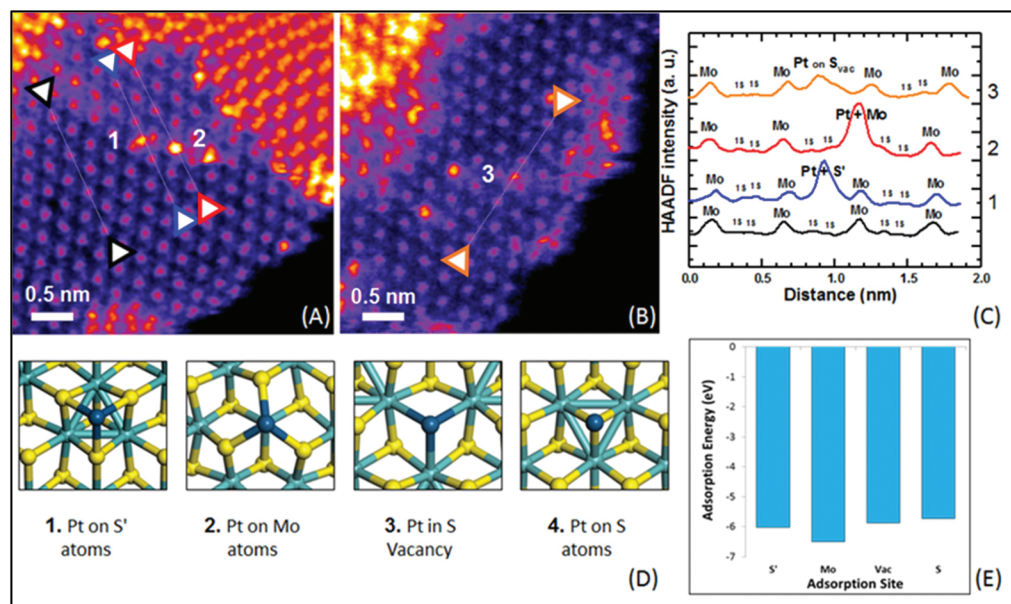
Density functional theory (DFT) calculations are also applied to calculate the Gibbs energy for hydrogen adsorption ( $\Delta G_{\text{H}^*}$ ) of the catalytic or electrocatalytic system when doped with different single transition metal atoms in order to predict the overall trend. According to the Sabatier principle, the H adsorption process is optimal when  $\Delta G_{\text{H}^*}$  is close to zero.<sup>54</sup> When the value of  $\Delta G_{\text{H}^*}$  is too negative, the hydrogen atom will bind too strongly onto the surface, which inhibits the subsequent desorption process for the formation of H<sub>2</sub> gas until thermal or electrical energy is supplied. In contrast, if the value of  $\Delta G_{\text{H}^*}$  is too positive, the catalyst surface will not favour the adsorption process for either the catalytic or electrocatalytic processes. It should be noted that this principle offers good predictive values to metallic phases with no impedance or diffusional limitations imposed between the active sites. Pt is known to be at the apex of the Sabatier volcano for H species. On the other hand, for composite materials such as SAC-MoS<sub>2</sub> one should consider the composite surface for the H interactions. For unmodified single molecular 2H-MoS<sub>2</sub> monolayers, the value of the Tafel slope (96 mV per decade) indicated that H adsorption is predominately the RDS for the overall HER. A plot of the reaction rate of 2H-MoS<sub>2</sub> when doped with different metal atoms (in terms of exchange current density) against their  $\Delta G_{\text{H}^*}$  in the HER was shown to have a volcano shape with Pt at the apex, indicating the dominant role of Pt compared to the 2H-MoS<sub>2</sub><sup>26</sup> (Fig. 2c). The excellent experimental results of Co-doped MoS<sub>2</sub> compared with other metals further supported the volcano plot. This suggests that this inexpensive non-noble metal can be effectively used without much activity compromise compared to using Pt. However, this work did not give a satisfactory explanation for the surprisingly poor Ni performance, which is electronically similar to Co atoms as they are neighbouring elements to each

other. Another independent work also studied how single transition metal atoms (Co, Ni, Ag, Cu) doped at the Mo atop sites would affect the semiconducting 2H-MoS<sub>2</sub>, but with a somewhat different conclusion drawn. Co-MoS<sub>2</sub> was confirmed to be the best HER catalyst among four transition metals with a Tafel slope value of 92 mV per decade and an overpotential of -300 mV at 10 mA cm<sup>-2</sup> (Table 2, entry 3).<sup>27</sup> The DOS calculations suggested that Co dopants help shifting the conduction band close to the Fermi level, which ultimately makes the surrounding S atoms at the basal planes better active sites for H adsorption. Also, no nice volcano relationship was noted when considering theoretical electronic factors of the metal dopers. The extended X-ray absorption fine structure (EXAFS) analysis was in fact conducted to study the chemical environment of the dopants. Interestingly, it was found that Co atoms anchored at the Mo atop site did not form a Co-Mo bond due to geometric constraint whereas the Ni-Mo formed negatively influenced the local S atoms in terms of the H adsorption ability. Thus, this clearly demonstrates that the metal doper can have different interactions with the single molecular layer MoS<sub>2</sub>. Therefore, basing solely on the electronic factor it is difficult to predict the trend without considering the precise position of the dopers and the steric constraints of the atoms in the composite structures. Regardless, the two pieces of research have interestingly shown that even when the same metal atoms are doped, the different doping position (Mo substitution site and Mo atop site) will largely affect the electronic properties of the resulting SAC-MoS<sub>2</sub> with 1H/2H polytypes.

For SAC-MoS<sub>2</sub> with 1T polytypes, Luo *et al.* suggested a significantly much stronger interaction between the transition metal doper and the 1T-MoS<sub>2</sub> support. They showed that the inclusion of 1% Pd atoms into the 1T structure can dramatically boost the HER activity (Table 2, entry 4).<sup>28</sup> It is confirmed by X-ray photoelectron spectroscopy (XPS), X-ray powder diffraction and EXAFS analysis that Pd atoms were successfully grown at the Mo substitution sites (Fig. 3). It is also possible to help create S defects during the synthesis. DFT calculations suggested that the  $\Delta G_{\text{H}}$  of the S adjacent to Pd (Pd-S\*-Mo) exhibited an almost thermoneutral value of -0.02 eV. Similar to the aforementioned metal atoms doped on 2H-MoS<sub>2</sub>, the major function of Pd atoms helps activating the adjacent S atoms (Pd-S\*-Mo) at the basal plane of 1T-MoS<sub>2</sub>. As suggested by the durability test, the highly stable Pd-S bonds also provide overall stability, which is another important criterion for a good heterocatalyst. However, it also should be noted that since the metastable 1T polytype is just a distorted form of 2H polytypes, both 1T and 2H polytypes will always coexist upon the incorporation of heteroatoms. Most key research studies in the literature have tended to neglect this important factor as the degree of coexistence (1T:2H ratios) was usually not provided nor addressing their correlation with activity. Another independent research work also studied the catalytic performance of Pd doped on 1T-MoS<sub>2</sub>. High-angle annular dark-field scanning transmission electron microscopy (HAADF-STEM, Fig. 4) combined with other structural characterisation techniques and computational calculations suggested that the Pd



**Fig. 3** Structural analysis of 1%Pd-MoS<sub>2</sub> and MoS<sub>2</sub>. (A) Pd-3d XPS and (B) XRD spectrum of the 1%Pd-MoS<sub>2</sub>. Fourier transform of the  $k^3$ -weighted (C) Pd K-edge and (D) Mo K-edge of the EXAFS spectra. (E) Dark-field scanning transmission electron microscopy image of the 1%Pd-MoS<sub>2</sub>. Blue and yellow dots represent Mo and S atoms respectively. The white line is the scale bar of 1 nm. (F) ESR spectra and (G) Raman spectra of 1%Pd-MoS<sub>2</sub> and MoS<sub>2</sub>. (H) Bode spectra with a frequency range from 0.1 Hz to 10 kHz and an amplitude of 5.0 mV.<sup>28</sup>



**Fig. 4** (a) and (b) HAADF-STEM of Pt-1T-MoS<sub>2</sub>. (c) ADF intensity line profiles taken along the corresponding coloured line of black, blue, red, and orange in the  $\langle 110 \rangle$  direction. (d) The proposed model of the Pt atom doping at 4 different adsorption sites. Numbers 1, 2 and 3 correspond to the experimentally observed Pd position shown in (a) and (b). (e) Adsorption energy for Pt atoms on each proposed adsorption site.<sup>29</sup>

atoms are doped at the Mo atop site rather than the Mo substitution site.<sup>29</sup> The Tafel slope analysis and DFT modelling reconfirmed that the H adsorption was readily facilitated by the 1T surface and thus the Volmer step is not the RDS. More importantly, the incorporated Pd atoms here could act as the active sites to accelerate the rate limiting recombination step

(Heyrovsky step or Tafel step) to form hydrogen gas. Overall, it was suggested that H adsorption would preferably take place on the 1T surface. The  $H_{ads}^*$  then hopped to the single Pd atoms to be recombined to H<sub>2</sub>. The difference in the roles of the Pd atoms for the HER from these two different studies probably originates from the two different doping positions due

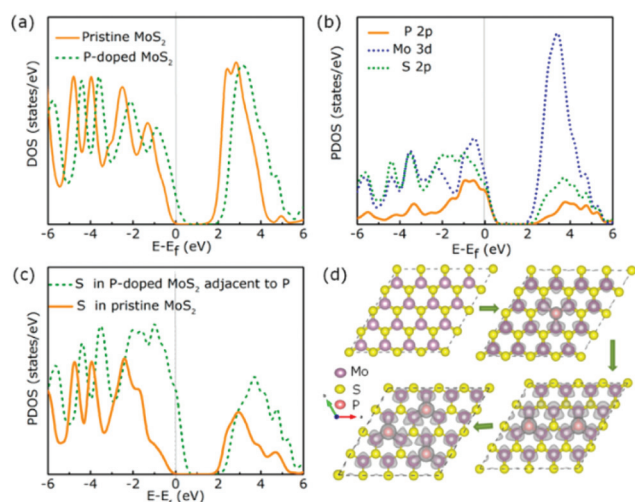
to their difference in synthetic methods used. The Pd atoms substituting Mo atoms usually formed stable Pd-S bonds which may offer steric hindrance to prevent any interactions with the  $H_{ads}^+$  while uncoordinated Pd atoms at the Mo atop site are electronically and spatially free to undergo redox reactions, making Pd-1T-MoS<sub>2</sub> a bifunctional catalyst for the HER.

It is interesting to note that non-metal P atom doped 1T MoS<sub>2</sub> nanosheets prepared by Liu and co-workers exhibited a small Tafel slope of 34 mV per decade with an excellent overpotential of 43 mV at 10 mA cm<sup>-2</sup> (Table 2, entry 6). Their catalytic performance is comparable to that of Pt doped graphene (with an overpotential of 45 mV and a Tafel slope of 29 mV per decade) which is considered to be one of the best SAC electrocatalysts reported.<sup>55</sup> The DFT calculation suggested the possibility that the P atoms incorporated at S substitution sites (S vacancy sites) could both act as new main active sites for H adsorption and activate the basal S atoms of MoS<sub>2</sub> at the same time. This offers a new opportunity to develop a non-metal based catalyst with no metal leaching issues during electrolysis of water at high potentials. The DOS and partial DOS simulations also suggested that more electronic states are shifted near the Fermi level after introducing P atoms (Fig. 5), which ultimately increased the electrical conductivity at the basal plane for an enhanced HER. In contrast, non-metal N atoms doped at the Mo substitution site once again did not directly participate in the HER but helped activate the surrounding S atoms to be more active for the H adsorption (Table 2, entry 7). Similarly, more DOS can be observed at the Fermi level when N atoms were embedded into MoS<sub>2</sub>, which also increases the electrical conductivity of basal sites. This once again reinforces the emphasis on how important the nature of the doper and the doping position can be when tailoring the SAC-MoS<sub>2</sub> for hydrogen reduction.

## 5. Conclusions

The effective use of catalyst components to convert substrate molecules to a product is important in catalysis. As stated, the use of SACs on 2D materials is a hot topic of research due to their high characteristic surface area to volume ratios and the electronic and steric influence on catalytically active atoms can be made clearer over these defined nanostructures. However, at this early stage, there are still limited amounts of high-quality data with corresponding details of experimental analysis, and computational calculations to lead to the understanding of the catalyst structure and activity relationship at the atomic level.

Here, through reviewing some selected key publications of SAC-MoS<sub>2</sub> for the HER, we can draw some conclusions with new insights into the otherwise commonly used term 'metal-support interactions' in heterogeneous catalysis. Also, understanding the HER over the catalyst surface for H<sub>2</sub> production from water is also very important in developing new catalytic technologies to harness renewable energies. It is shown that the polytypes of 2D MoS<sub>2</sub> (the specific support material with characteristic structure and properties), the choice of single heteroatoms (*i.e.* both metal or non-metal atoms), and the doping position (defined site for specific electronic and steric perturbation) are the three most important factors for deciding the catalytic performance of the resulting SAC-MoS<sub>2</sub> in the HER. Conventional electrochemical scan tests and volcano plots can be used to gauge which metal dopants on the 2H-MoS<sub>2</sub> support could potentially help enhancing the overall HER performance. DFT modelling is important to characterise the electronic and steric properties of the atoms as the catalyst as well as the influence from the support. However, such a trend can subtly be changed if the chemical nature of dopers, relative sizes, and dwelling positions are altered. As a result, atomic positions on defined structures with defined interactions are required to obtain further understanding of structure-activity relationships. Specifically to the HER, the heteroatoms that replace Mo atoms can activate the adjacent in-plane S atoms for H adsorption which ultimately enhance the ability for H adsorption. They are anticipated to be more stable and durable in position since they are stably embedded into the MoS<sub>2</sub> structure, whereas heteroatom doping at the Mo atop sites or S vacancy sites through interacting with the outer S layers can act as the active sites themselves and also activate the surrounding S through electronic perturbation for the HER. In the case where the basal S atoms have already been activated for H adsorption by other reasons, for example the heteroatoms are doped on a S defect rich 1T-MoS<sub>2</sub>, the heteroatoms may become the active sites for recombination of  $H_{ads}^+$  to H<sub>2</sub> if preferable. Overall, all these parameters could be tailored by choosing the correct synthetic methods, which makes SAC-MoS<sub>2</sub> an interesting material for various electrochemical applications. This also provides insight into developing other 2D nanomaterials confined with different heteroatoms for various catalytic uses.



**Fig. 5** (a) Total DOS of pure MoS<sub>2</sub> and P doped MoS<sub>2</sub>. (b) Partial DOS of P-MoS<sub>2</sub>. (c) Partial DOS for S atoms. The vertical grey lines represent the Fermi level. (d) Partial charge density of P-MoS<sub>2</sub> when doped with 0–3 P atoms.



## Conflicts of interest

There are no conflicts to declare.

## Notes and references

- W. Choi, I. Lahiri, R. Seelaboyina and Y. S. Kang, *Crit. Rev. Solid State Mater. Sci.*, 2010, **35**, 52–71.
- D. Akinwande, C. J. Brennan, J. S. Bunch, P. Egberts, J. R. Felts, H. Gao, R. Huang, J.-S. Kim, T. Li, Y. Li, *et al.*, *Extreme Mech. Lett.*, 2017, **13**, 42–77.
- M. Chhowalla, H. S. Shin, G. Eda, L. J. Li, K. P. Loh and H. Zhang, *Nat. Chem.*, 2013, **5**, 263–275.
- H. Pan, *Sci. Rep.*, 2014, **4**, 1–6.
- M. Chhowalla, Z. Liu and H. Zhang, *Chem. Soc. Rev.*, 2015, **44**, 2584–2586.
- A. Yusuf, C. Snape, J. He, H. Xu, C. Liu, M. Zhao, G. Z. Chen, B. Tang, C. Wang, J. Wang and S. N. Behera, *Catal. Rev. - Sci. Eng.*, 2017, **59**, 189–233.
- S. Royer and D. Duprez, *ChemCatChem*, 2011, **3**, 24–65.
- M. Khazaei, M. Arai, T. Sasaki, C. Y. Chung, N. S. Venkataramanan, M. Estili, Y. Sakka and Y. Kawazoe, *Adv. Funct. Mater.*, 2013, **23**, 2185–2192.
- J. Xie and Y. Xie, *Chem. – Eur. J.*, 2016, **22**, 3588–3598.
- D. Kong, H. Wang, J. J. Cha, M. Pasta, K. J. Koski, J. Yao and Y. Cui, *Nano Lett.*, 2013, **13**, 1341–1347.
- Y. Zhang, Q. Ji, G. F. Han, J. Ju, J. Shi, D. Ma, J. Sun, Y. Zhang, M. Li, X. Y. Lang, Y. Zhang and Z. Liu, *ACS Nano*, 2014, **8**, 8617–8624.
- H. Nan, Z. Wu, J. Jiang, A. Zafar, Y. You and Z. Ni, *J. Phys. D: Appl. Phys.*, 2017, **50**, 154001.
- M. R. Islam, N. Kang, U. Bhanu, H. P. Paudel, M. Erementchouk, L. Tetard, M. N. Leuenberger and S. I. Khondaker, *Nanoscale*, 2014, **6**, 10033–10039.
- L. Tao, X. Duan, C. Wang, X. Duan and S. Wang, *Chem. Commun.*, 2015, **51**, 7470–7473.
- H. Li, C. Tsai, A. L. Koh, L. Cai, A. W. Contryman, A. H. Fragapane, J. Zhao, H. S. Han, H. C. Manoharan, F. Abild-Pedersen, J. K. Nørskov and X. Zheng, *Nat. Mater.*, 2015, **15**, 48–53.
- H. Li, A. W. Contryman, X. Qian, S. M. Ardakani, Y. Gong, X. Wang, J. M. Weisse, C. H. Lee, J. Zhao, P. M. Ajayan, J. Li, H. C. Manoharan and X. Zheng, *Nat. Commun.*, 2015, **6**, 7381.
- Y. Tan, P. Liu, L. Chen, W. Cong, Y. Ito, J. Han, X. Guo, Z. Tang, T. Fujita, A. Hirata and M. W. Chen, *Adv. Mater.*, 2014, **26**, 8023–8028.
- B. Qiao, A. Wang, X. Yang, L. F. Allard, Z. Jiang, Y. Cui, J. Liu, J. Li and T. Zhang, *Nat. Chem.*, 2011, **3**, 634–641.
- G. Vilé, D. Albani, M. Nachtegaal, Z. Chen, D. Dontsova, M. Antonietti, N. López and J. Pérez-Ramírez, *Angew. Chem., Int. Ed.*, 2015, **54**, 11265–11269.
- X.-F. Yang, A. Wang, B. Qiao, J. Li, J. Liu and T. Zhang, *Acc. Chem. Res.*, 2013, **46**, 1740–1748.
- A. Alarawi, V. Ramalingam and J. H. He, *Mater. Today Energy*, 2019, **11**, 1–23.
- C. Zhu, S. Fu, Q. Shi, D. Du and Y. Lin, *Angew. Chem., Int. Ed.*, 2017, **56**, 13944–13960.
- K. C. Kwon, J. M. Suh, R. S. Varma, M. Shokouhimehr and H. W. Jang, *Small Methods*, 2019, **3**, 1–20.
- Q. H. Wang, K. Kalantar-Zadeh, A. Kis, J. N. Coleman and M. S. Strano, *Nat. Nanotechnol.*, 2012, **7**, 699–712.
- D. Voiry, M. Salehi, R. Silva, T. Fujita, M. Chen, T. Asefa, V. B. Shenoy, G. Eda and M. Chhowalla, *Nano Lett.*, 2013, **13**, 6222–6227.
- J. Deng, H. Li, J. Xiao, Y. Tu, D. Deng, H. Yang, H. Tian, J. Li, P. Ren and X. Bao, *Energy Environ. Sci.*, 2015, **8**, 1594–1601.
- T. H. M. Lau, X. Lu, J. Kulhavy, S. Wu, L. Lu, T. S. Wu, R. Kato, J. S. Foord, Y. L. Soo, K. Suenaga and S. C. E. Tsang, *Chem. Sci.*, 2018, **9**, 4769–4776.
- Z. Luo, Y. Ouyang, H. Zhang, M. Xiao, J. Ge, Z. Jiang, J. Wang, D. Tang, X. Cao, C. Liu and W. Xing, *Nat. Commun.*, 2018, **9**, 1–8.
- T. H. M. Lau, S. Wu, R. Kato, T. S. Wu, J. Kulhavy, J. Mo, J. Zheng, J. S. Foord, Y. L. Soo, K. Suenaga, M. T. Darby and S. C. E. Tsang, *ACS Catal.*, 2019, **9**, 7527–7534.
- P. Liu, J. Zhu, J. Zhang, P. Xi, K. Tao, D. Gao and D. Xue, *ACS Energy Lett.*, 2017, **2**, 745–752.
- R. Li, L. Yang, T. Xiong, Y. Wu, L. Cao, D. Yuan and W. Zhou, *J. Power Sources*, 2017, **356**, 133–139.
- M. Acerce, D. Voiry and M. Chhowalla, *Nat. Nanotechnol.*, 2015, **10**, 313–318.
- F. Wypych and R. Schollhorn, *J. Chem. Soc., Chem. Commun.*, 1992, 1386–1388.
- D. B. Putungan, S. Lin and J. Kuo, *Phys. Chem. Chem. Phys.*, 2015, **17**, 21702–21708.
- J. C. Torres-Guzmán, G. Martínez-Mekler and M. F. Müller, *J. Chem. Phys.*, 2016, 174702.
- T. F. Jaramillo, K. P. Jørgensen, J. Bonde, J. H. Nielsen, S. Hørch and I. Chorkendorff, *Science*, 2007, **317**, 100–102.
- H. Tang and S. Roy Morrison, *Thin Solid Films*, 1993, **227**, 90–94.
- M. A. Lukowski, A. S. Daniel, F. Meng, A. Forticaux, L. Li and S. Jin, *J. Am. Chem. Soc.*, 2013, **135**, 10274–10277.
- D. Voiry, M. Salehi, R. Silva, T. Fujita, M. Chen, T. Asefa, V. B. Shenoy, G. Eda and M. Chhowalla, *Nano Lett.*, 2013, **13**, 6222–6227.
- C. A. Papageorgopoulos and W. Jaegermann, *Surf. Sci.*, 1995, **338**, 83–93.
- Y. Wang, J. Z. Ou, S. Balendhran, A. F. Chrimes, M. Mortazavi, D. D. Yao, M. R. Field, K. Latham, V. Bansal, J. R. Friend, S. Zhuiykov, N. V. Medhekar, M. S. Strano and K. Kalantar-Zadeh, *ACS Nano*, 2013, **7**, 10083–10093.
- X. Fan, P. Xu, D. Zhou, Y. Sun, Y. C. Li, M. A. T. Nguyen, M. Terrones and T. E. Mallouk, *Nano Lett.*, 2015, **15**, 5956–5960.
- X. Ma, P. Zhang, Z. Wang, R. Wu, T. Jiang, H. Wang, Y. Zhao, S. Wu, Y. Xie, Y. Zhan and T. Nan, *Nanotechnology*, 2016, **28**, 084001.



- 44 T. Payam, W. Jieqiong, X. Hui, F. D. Joel, A. Mumtaz Murat, Z. Chuan, K. Kaifei, B. Brett, Z. Lijie, Z. Puqin, H. Shaoming, Y. Sen, V. B. Frank, C. John and Z. Hao, *Mater. Res. Express*, 2016, **3**, 75009.
- 45 Z. He and W. Que, *Appl. Mater. Today*, 2016, **3**, 23–56.
- 46 L. Samad, S. M. Bladow, Q. Ding, J. Zhuo, R. M. Jacobberger, M. S. Arnold and S. Jin, *ACS Nano*, 2016, **10**, 7039–7046.
- 47 X. Feng, Q. Tang, J. Zhou, J. Fang, P. Ding, L. Sun and L. Shi, *Cryst. Res. Technol.*, 2013, **48**, 363–368.
- 48 C. Perumal Veeramalai, F. Li, H. Xu, T. W. Kim and T. Guo, *RSC Adv.*, 2015, **5**, 57666–57670.
- 49 X. Feng, Q. Tang, J. Zhou, J. Fang, P. Ding, L. Sun and L. Shi, *Cryst. Res. Technol.*, 2013, **48**, 363–368.
- 50 G. Liu, A. W. Robertson, M. M.-J. J. Li, W. C. H. H. Kuo, M. T. Darby, M. H. Muhieddine, Y.-C. C. Lin, K. Suenaga, M. Stamatakis, J. H. Warner and S. C. E. Tsang, *Nat. Chem.*, 2017, **9**, 810–816.
- 51 R. G. Compton and C. E. Banks, *Understanding voltammetry*, World Scientific, 2011.
- 52 S. Park, Y. Shao, J. Liu and Y. Wang, *Energy Environ. Sci.*, 2012, **5**, 9331–9344.
- 53 Y. Li, H. Wang, L. Xie, Y. Liang, G. Hong and H. Dai, *J. Am. Chem. Soc.*, 2011, **133**, 7296–7299.
- 54 S. Trasatti, *J. Electroanal. Chem.*, 1972, **39**, 163–184.
- 55 N. Cheng, S. Stambula, D. Wang, M. N. Banis, J. Liu, A. Riese, B. Xiao, R. Li, T. K. Sham, L. M. Liu, G. A. Botton and X. Sun, *Nat. Commun.*, 2016, **7**, 1–9.