

PAPER

[View Article Online](#)
[View Journal](#) | [View Issue](#)Cite this: *J. Mater. Chem. A*, 2024, **12**, 8438Origin of electrocatalytic nitrogen reduction activity over transition metal disulfides: critical role of *in situ* generation of S vacancy†Tianyi Wang,^a Zhongyuan Guo,^b Hirofumi Oka,^a Akichika Kumatani,^c Chuangwei Liu^{*f} and Hao Li^{*a}

The electrochemical nitrogen reduction reaction (ENRR) is a promising and sustainable alternative to conventional Haber–Bosch ammonia (NH₃) synthesis. Pursuing high-performance and cost-effective ENRR catalysts is an open challenge for achieving commercial-scale ambient NH₃ production. Less-precious transition metal disulfides (TMS₂) are a class of promising catalysts that can be highly active for ENRR. However, the origin of their high ENRR performance is not well understood. Herein, we analyze the origin of their activity by probing their electrochemistry-induced surface states. Starting with a typical ENRR TMS₂ catalyst, iron disulfide (FeS₂), from our calculated surface Pourbaix diagrams we found that S-vacancies can be easily generated under an ENRR potential. Our subsequent spin-polarized density functional theory (DFT) calculations show that this electrochemistry-driven “*in situ*” generation of S-vacancies shows significantly higher ENRR activity than a stoichiometric pristine FeS₂ surface due to the stronger N–N adsorption and activation capacity of a lower-coordination-number S-vacancy site. This finding is in excellent agreement with experimental observations published in recent years regarding potential windows reaching the maximum faradaic efficiency. We then expanded our analysis to other typical TMS₂ that had shown promising ENRR performance in recent experimental literature (SnS₂, MoS₂, NiS₂, and VS₂), and found that such an “*in situ*” S-vacancy generation phenomenon is universal under ENRR potentials, with results in good agreement with many experimental observations reported to date. We conclude that, though S-vacancy engineering during synthesis is a promising strategy to enhance the ENRR performance on TMS₂ catalysts, the “*in situ*” generation of S-vacancies will also endow pristine TMS₂ with a measurable ENRR performance. This study shows that the surface states of ENRR catalysts should not be dismissed before analyzing the activity of an ENRR catalyst. Most importantly, we found that when designing a promising TMS₂ catalyst for ENRR, its capacity to form S-vacancies is a key performance indicator that needs to be analyzed.

Received 14th January 2024
Accepted 21st February 2024

DOI: 10.1039/d4ta00307a

rsc.li/materials-a

Introduction

Ammonia (NH₃), an indispensable chemical feedstock, plays a pivotal role in facilitating global economic development and

demographic expansion. Its versatile utility spans pharmaceutical synthesis, fertilizer production, and compound fabrication.¹ NH₃ also emerges as a prospective carbon-neutral medium for realizing the hydrogen economy, functioning as a liquid hydrogen carrier without carbon emissions.² Currently, the Haber–Bosch (H–B) process prevails as the industrial framework for NH₃ synthesis, but harsh reaction conditions (*c.a.* 450 °C and 300 bar) demand substantial energy consumption and emit enormous amounts of greenhouse gases.³ As an alternative, the electrochemical nitrogen reduction reaction (ENRR) offers an ecologically benign route to synthesize NH₃ under ambient reaction conditions. In this context, the N₂ is fixed and hydrogenated by protons and electrons in an aqueous solution instead of gaseous H₂. However, the competing hydrogen evaluation reaction (HER) and insertion of N₂ fixation are two significant challenges for industrial-scale ENRR.^{4–6} Specifically, optimal catalysts maintain a delicate equilibrium between N₂ adsorption and NH₃ desorption, reducing formidable energy barriers. Additionally, HER

^aAdvanced Institute for Materials Research (WPI-AIMR), Tohoku University, Sendai, 980-8577, Japan. E-mail: li.hao.b8@tohoku.ac.jp^bCollege of Environmental and Resource Sciences, Zhejiang University, Hangzhou, 310058, China^cInstitute of Engineering Innovation (IEI), School of Engineering, The University of Tokyo, Tokyo, 113-8656, Japan. E-mail: kumatani@g.ecc.u-tokyo.ac.jp^dPrecursory Research for Embryonic Science and Technology (PRESTO), Japan Science and Technology Agency (JST), Saitama 332-0012, Japan^eGraduate School of Environmental Studies, Center for Science and Innovation in Spintronics (CSIS), Tohoku University, Sendai, Miyagi 980-8579, Japan^fKey Lab for Anisotropy and Texture of Materials, School of Materials Science and Engineering, Northeastern University, Shenyang, 110819, China. E-mail: liucw@mail.neu.edu.cn† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4ta00307a>

remains the key obstacle to improving the faradaic efficiency (FE). Comparatively, HER is characterized by a two-step hydrogenation pathway, which is thermodynamically more favorable than the complicated six-step hydrogenation ENRR process.^{7,8}

Some noble-metal-based catalysts, such as those based on Ru, Au, Pd, and Rh, have demonstrated exceptional activities and FEs in ENRR. However, their prohibitive price and shortage impede their widespread industrial implementation.^{3,9–11} Therefore, the pursuit of cost-effective and high-performance catalysts has become an intensified research focus in recent years. Fe-based catalysts have garnered significant attention as promising candidates to get close to or even surpass existing benchmarks in ENRR. For example, Fe-related proteins can serve as intrinsic active sites within enzymes in biological N₂ fixation.¹² Fe is also the central constituent within catalysts deployed in the H-B method.¹³ Drawing inspiration from biological and H-B approaches, FeS₂, as a typical transition metal disulfide (TMS₂), has emerged as a promising candidate for ENRR with structural alignment with the Fe–S ligand arrangement characteristic of nitrogenase.¹⁴ Additionally, S-containing Fe materials exhibit notable hydrophobic properties attributed to the inherent hydrophobic nature of S, thereby mitigating the adverse effect of the competing HER.^{15–17} In recent studies, FeS₂ nanoparticles embedded in graphene oxide (FeS₂@GO) could generate low-coordination Fe active sites to facilitate N₂ fixation and ENRR, exhibiting an FE of 6.80% and an NH₃ yield of 27.9 μg h^{−1} mg_{cat}^{−1} at −0.3 V vs. reversible hydrogen electrode (V_{RHE}) under acidic and neutral conditions.¹⁸ In addition, doping is a common strategy to improve the catalytic performance of FeS₂-based catalysts. For example, the synergy between Cr and S-vacancies (S_V) of FeS₂ can achieve an NH₃ yield of 11.5 μg h^{−1} mg_{cat}^{−1} and an FE of 14.6% at −0.2 V_{RHE}.¹⁹ The synergistic effects among FeS₂, MoS₂, and reduced GO may also expedite the kinetics of N₂ hydrogenation in pH-universal aqueous solutions, elevating the NH₃ yield and FE to 41.1 μg h^{−1} mg_{cat}^{−1} and 38.6% at −0.2 V_{RHE}, respectively.²⁰ Moreover, FeS₂–MoS₂ supported on Fe foam exhibited a considerable FE of 4.6% at −0.5 V_{RHE} and an NH₃ production rate of 7.1 × 10^{−10} mol s^{−1} cm^{−2} at −0.3 V_{RHE}. This was attributed to the synergistic effects and interface interaction between FeS₂ and MoS₂ phases, which furnish favourable electron transfer pathways with abundant active sites.²¹ Doping Mo into FeS₂ also achieved an FE of 14.41% at −0.2 V_{RHE}, which was attributed to the joint factors of the superior activity of Mo and HER suppression of FeS₂.²² Besides, other TMS₂, such as VS₂, NiS₂, and SnS₂, show great promise in ENRR.^{23–25} Fig. 1 and Table S1† summarize the ENRR performance in terms of FE of typical TMS₂ reported after 2018. It can be seen that, in general, these materials preferentially achieve the highest FE or experience a rapid growth in FE at a relatively “early” potential (*c.a.* −0.4 V_{RHE}). For the better design and modification of TMS₂ catalysts for ENRR, it is particularly important to provide a deep understanding of the superior ENRR performance of TMS₂. Furthermore, it is worth noting that electrochemical nitrate reduction is another efficient approach for NH₃ synthesis. The synergy achieved through the formation of the Sn–Fe pair site, where Sn atoms are dispersed on FeS₂, can also promote the NO₃ protonation

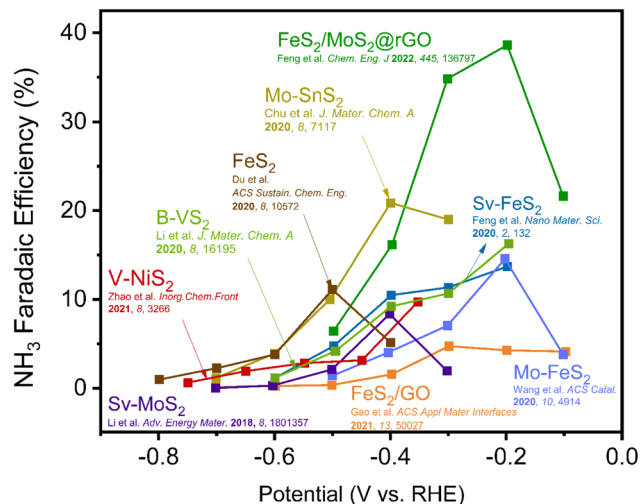


Fig. 1 Statistics of ENRR faradaic efficiencies (FEs) of transition metal disulfides (TMS₂) with varying applied potentials. Data were extracted from typical experimental reports after 2018.^{18–20,22–25,27,28}

process, reaching a maximum FE of 96.7% and an NH₃ yield of 15.8 mg h^{−1} cm^{−2} at −0.5 V_{RHE}.²⁶ All above pioneering experimental results suggest that TMS₂ catalysts, especially FeS₂, are promising materials for electrocatalytic NH₃ synthesis.

Due to the electrochemistry-driven water-adsorbate equilibrium,²⁹ the surface states (*e.g.*, the electrochemistry-induced surface coverage and vacancy formation) of an electrocatalyst should be considered before the analysis of electrocatalytic activity. Recent combined experimental and theoretical studies found that under a moderate or high potential (*e.g.*, oxygen reduction and evolution potentials), many transition metal X-ide (TMX) surfaces are precovered by O* or HO*, resulting in a very different electronic structure of a surface or the poisoning of too-reactive sites.^{30–32} This makes TMXs generally behave very differently from a stoichiometric pristine surface under electrocatalytic conditions. However, surface state analysis is a largely dismissed part of many previous theoretical analyses for electrocatalysis.³³ Very recently, we found that only when we consider a more realistic surface coverage of ZrN can we fully understand its superior electrocatalytic oxygen reduction performance with good agreement with experiments.³⁴ Therefore, for ENRR analysis, it is reasonable to hypothesize that in the presence of a low potential, anion vacancies may be directly formed on some TMXs (*e.g.*, TMS₂) because protons and electrons can easily combine with the surface and lead to the leaching out of the anions of the material. This may lead to the formation of vacancies, which would significantly change the type of active sites and electronic structures of an ENRR catalyst. Unfortunately, this phenomenon was usually dismissed in previous analyses of either TMX materials or ENRR processes.

Motivated by current stages, herein, we analyze the ENRR performance of TMS₂ starting from probing electrochemistry-induced surface states, based on spin-polarized *ab initio* calculations with van der Waals corrections. Using FeS₂ as an initial example, we found an interesting “*in situ*” generation of S-vacancies under ENRR operating potentials. This potential



window is in good agreement with the potential where the highest experimental ENRR FE is located. This suggests that, even if starting with a stoichiometric FeS_2 for ENRR, it is still more likely to result in an S-vacancy-containing surface. Furthermore, we found that these “*in situ*” generated S-vacancies are more active than an original FeS_2 surface due to the lower coordination number of an Fe site that leads to stronger N–N adsorption and activation capacity. Finally, we extended this analysis to other TMS₂ catalysts (SnS_2 , MoS_2 , NiS_2 , and VS_2) and found a similar conclusion. All these analyses lead to good agreement with the experimentally highest ENRR FE-windows of TMS₂. Therefore, a comprehensive assessment of the more realistic surface states of TMS₂-based catalysts should be considered before theoretical and experimental studies of ENRR. Most importantly, this study proposes that, when designing a promising TMS₂ catalyst for ENRR, whether it can form an S-vacancy easily is a key performance indicator that should be carefully evaluated.

Results and discussion

Surface properties are of paramount importance in the comprehension and design of catalysts, yielding characteristics distinctive to those of bulk materials. Grazing-incidence X-ray diffraction verified the prevalence of the FeS_2 crystallographic phase (namely, (111), (200), (210), (211), and (311)) exhibiting higher peak intensities (Fig. S1†).³⁵ Furthermore, we analysed the surface energies of these facets. Table S2† shows a comparison of their surface energies, highlighting that the (111) facet has higher thermodynamic stability owing to its lower surface energy, in good agreement with previous relevant experimental observations.^{27,36,37}

The calculated surface Pourbaix diagram is a key tool for simulating and describing catalytic surface states under electrocatalytic operating conditions as a function of pH and potential, which indicates the thermodynamic equilibrium surface structure in a certain aqueous environment.³⁸ Under electrocatalytic conditions, pre-coverage with H^* , O^* , or HO^* species generated through water activation can possibly occur on the surface under an oxidizing potential. These coverages exert a substantial influence on the reaction overpotential and surface configuration.^{39,45} The calculated surface Pourbaix diagram provides a visual representation of thermodynamically stable FeS_2 (111) (Fig. 2), and the lowest-energy line represents the surface with the lowest free energy under given operating conditions.³⁸ It illustrates the diverse configurations of H^* , HO^* , and O^* coverages across a range of pH and applied potentials, which also includes a stability assessment of various S_v coverages. The observed trend reveals the transformation of pristine FeS_2 (111) into an O-terminated state at highly positive potentials exceeding $1.10 V_{\text{RHE}}$. Additionally, the single S-vacancy decorated FeS_2 has the lowest free energy in the ENRR-preferred potential window (e.g., around and above $-0.5 V_{\text{RHE}}$), indicating that S-vacancies are prone to occur spontaneously. Therefore, electrochemistry can drive the formation of S-vacancies during ENRR. Based on experimental data extracted from previous reports (Fig. 2b), it is easy to find that these

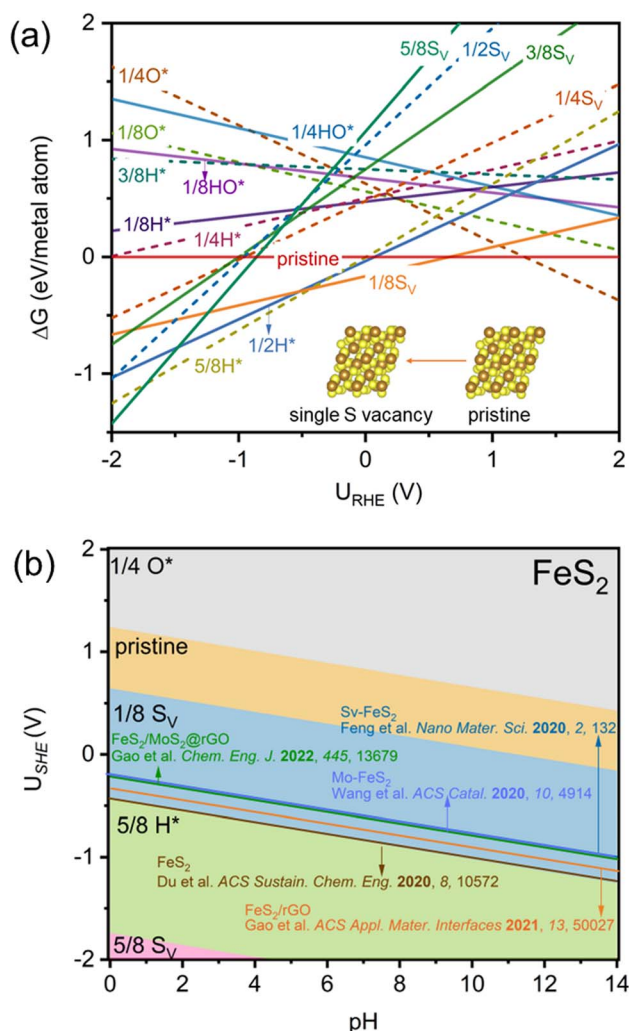


Fig. 2 Calculated (a) 1D and (b) 2D surface Pourbaix diagrams of FeS_2 (111) considering different coverages of S_v , O^* , H^* , and HO^* . The experimental potentials at the highest faradaic efficiencies of reported FeS_2 -based catalysts are plotted for a direct comparison. These experimental data were extracted from ref. 18–20, 22 and 27.

experimentally identified high-FE potentials are located in the S-vacancy formation potential range, which means that *in situ* formed S-vacancy generation is prone to occur on the surface in experiments. The S-vacancy formation window and high activity of ENRR identified on the S-vacancy sites are linked to the experimental potential where a high ENRR FE is located.^{9,10,12,16,17} The H-coverage subsequently occurs at a potential more negative than $-0.5 V_{\text{RHE}}$, suggesting that the competing HER or H^* -poisoning may gradually predominate when the potential becomes too negative. This is also in good agreement with the experimental observation that ENRR-FEs of FeS_2 -based catalysts will gradually drop when the potential becomes more negative than $-0.5 V_{\text{RHE}}$. These results suggest that the S_v -containing surface is more favourable than a pristine FeS_2 surface during ENRR, and therefore, it is vital to explore electrocatalytic surface states for guiding and designing superior catalysts. As illustrated in Fig. S2,† AIMD simulations



further suggest a high stability of $\text{FeS}_{2-x}(111)\text{-}1\text{S}_\text{V}$ without significant structural deformation.

Next, we analyzed the free energy pathways of the ENRR reaction over pristine FeS_2 and FeS_{2-x} surfaces (Fig. 3 and S3†). The ENRR is commonly divided into associative and dissociative mechanisms. In the dissociative mechanism, $\text{N}\equiv\text{N}$ cleavage occurs first, followed by the subsequent hydrogenation of the N atom. The direct N_2 dissociation is energetically unfavorable due to the N_2 endoergicity of 1.73 eV, and thus, it is not discussed in this study.³⁹ Fig. 3 displays the ENRR elementary steps of pristine $\text{FeS}_2(111)$ and $\text{FeS}_{2-x}(111)\text{-}1\text{S}_\text{V}$ via the associative mechanism, including alternating and distal pathways. When ENRR follows an alternating pathway, the protons attack two N atoms alternately, and the reaction continuously releases two NH_3 molecules. On stoichiometric pristine $\text{FeS}_2(111)$, Fe serves as the reaction site, and the first hydrogenation step ($\text{N}_2^* + \text{H}^* \rightarrow \text{NNH}^*$) is the potential-determining step (PDS) with a very endothermic energy ($\Delta G_{\text{max}} = 2.76$ eV). For $\text{FeS}_{2-x}(111)\text{-}1\text{S}_\text{V}$, Fe and S_V can potentially serve as active sites. The protonation of N_2^* to NNH^* is also the PDS along the Fe site, while the energy barrier is much lower ($\Delta G_{\text{max}} = 1.02$ eV). The transition state energy of the first coupled proton–electron transfer is 0.19 eV (Fig. S5†). Notably, the S_V site has an excellent ability to fix N_2 ($\Delta G_{\text{N}_2^*} = -1.14$ eV), but release of the second NH_3 molecule is a significantly endothermic step ($\Delta G_{\text{max}} = 1.73$

eV). In the distal pathway, protons are continuously added to the distal N atom of adsorbed N_2 , and later, the first NH_3 molecule is released. Afterward, the other N atom is hydrogenated to generate the second NH_3 molecule. For pristine $\text{FeS}_2(111)$ and $\text{FeS}_{2-x}(111)\text{-}1\text{S}_\text{V}$, the PDS and associated ΔG_{max} remain consistent with those identified in the alternating pathway. However, the reaction $\text{NNH}^* + \text{H}^* \rightarrow \text{NNH}_2^*$ occurs at the S_V site of $\text{FeS}_{2-x}(111)\text{-}1\text{S}_\text{V}$, featuring a relatively higher ΔG (0.99 eV). Therefore, the Fe active site of $\text{FeS}_{2-x}(111)\text{-}1\text{S}_\text{V}$ yields a smoother and lower energy diagram in the alternating pathway. Note that in this study, we followed the reaction mechanisms discussed in some previous studies (e.g., studies by Nørskov and colleagues)⁴⁰ that the formation and desorption of NH_3^* may be coupled into one step. Therefore, based on this mechanism, the rate-determining step may still be the formation of NNH^* instead of NH_3^* desorption on S_V . After the formation of NH_3 , NH_4^+ can be further formed easily, which will provide a further driving force to make NH_3 less likely to get stuck on an S_V . In addition, as a supplementary calculation, we also tabulated the NH_3 desorption energy on the S_V site (Table S3†), showing that the energetics of NH_3 desorption from NH_3^* are lower than for the formation of NNH^* . In addition, Table S4† directly compares the adsorption energies of N_2 and NH_3 , elucidating the feasibility of NH_3 desorption in the vicinity of the S-vacancy. Besides, the markedly positive ΔG_{H^*} value (1.16 eV) indicates the thermodynamic infeasibility of the competing HER. In addition, solvation effects were tested by developing an explicit model with three water molecules (Fig. S6†). After considering the solvation effect, the energy barrier of the PDS ($\text{NN}^* + \text{H}^* \rightarrow \text{NNH}^*$) became 0.98 eV, which is only slightly more negative than that without a solvent (1.02 eV). Therefore, we consider that the solvation effect has no significant influence on the ENRR activity on FeS_{2-x} , which is consistent with a previous study.⁴¹ Previous studies considering explicit solvent effects on ENRR suggested that the solvation effects of ENRR may lower theoretical overpotentials by less than 0.1 eV.⁴¹ The PDS and energy profiles remain largely unchanged following the interactions exclusively involving H_2O . Furthermore, an implicit model is unable to provide accurate results because it cannot consider the H-bonding effect between the adsorbate and solvent; therefore, it was not considered in our study. To further assess the Hubbard- U correction, U values were implemented to investigate the energy barriers to N_2 adsorption and PDS ($\text{NN}^* + \text{H}^* \rightarrow \text{NNH}^*$) on the identified active site.⁴² As shown in Table S5†, standard RPBE and RPBE + U led to similar results. In addition, the Material Project database suggests that U correction is not necessary for FeS_2 .^{43,44}

Based on Fig. 2 and 3, the electrochemical surface states (i.e., the electrochemistry-induced surface coverage) of FeS_2 under different pH conditions and applied potentials indicate that highly active S-vacancies are formed under ENRR operating conditions (e.g., around and above -0.50 V_{RHE}). Meanwhile, the experimental potentials of Fe-based catalysts in ENRR at their highest FEs are located in the S-vacancy formation potential range, which means that the *in situ* generation of S-vacancies is prone to occurring on the surface during ENRR experiments. The DFT-calculated energy profiles further proved that S-

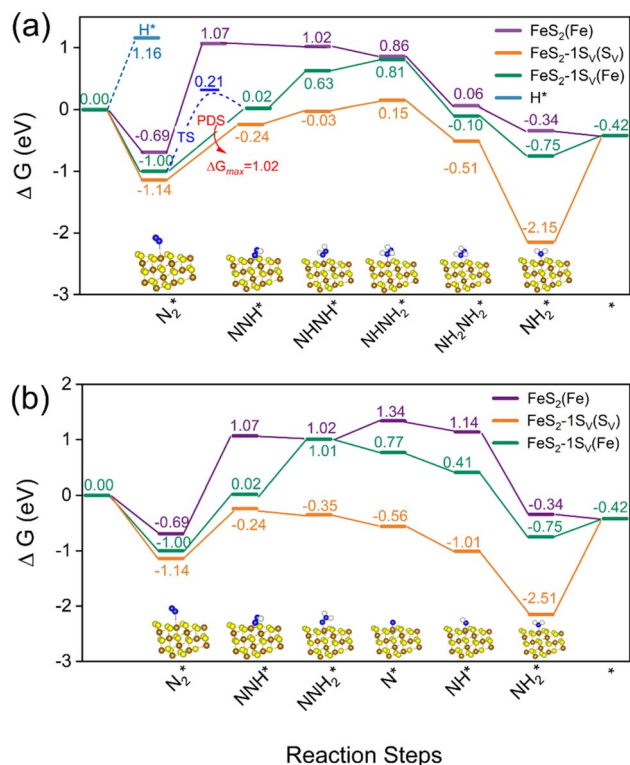


Fig. 3 Calculated Gibbs free energy diagrams of ENRR over $\text{FeS}_2(111)$ and $\text{FeS}_{2-x}(111)\text{-}1\text{S}_\text{V}$ surfaces. (a) and (b) demonstrate the energy profiles through the alternating and distal pathways, respectively. The insets show the optimized geometries of the key reaction intermediates. Brown, yellow, blue, and white spheres represent Fe, S, N, and H, respectively.



vacancies are highly active for ENRR, in contrast to the low theoretical activity on a stoichiometric pristine FeS_2 surface. Because N_2 fixation is the pivotal step in the entire ENRR process, herein, the projected density of states (PDOS) and charge density difference upon N_2 adsorption were calculated to analyze the electronic structure of the N_2^* intermediate (Fig. 4). N_2^* has a strong peak in the deep valence band of N_2 on pristine $\text{FeS}_2(111)$, while the S_V formation encourages electron transfer between N_2 and $\text{FeS}_{2-x}(111)-1\text{S}_\text{V}$. The robust N_2 bonding is weakened as it serves as an electron donor, which in turn can promote N_2 fixation and hydrogenation.

Currently, TMS_2 catalysts have become a class of popular components for ENRR, due to their excellent electrochemistry and cost-effectiveness. In addition to $\text{FeS}_2(111)$, other TMS_2 catalysts, including $\text{MoS}_2(100)$, $\text{NiS}_2(210)$, $\text{VS}_2(001)$, $\text{SnS}_2(111)$, and $\text{VS}_2(011)$, were observed in experiments as primary materials and facets in effective electrochemical NH_3 synthesis.^{23–25,28} Therefore, we further expanded our analysis to these catalysts. Interestingly, as depicted in Fig. 5a, *in situ* S_V generation on all these TMS_2 catalysts is energetically favorable within the ENRR potential range, which may regulate the electronic structure and enhance catalytic activity. In either a direct or an indirect way, various theoretical and experimental studies can support the idea that S_V sites of TMS_2 generally show better ENRR performance than a pristine surface. For example, Chu *et al.*²⁴ found that N_2 adsorption is typically weak on pristine SnS_2 , while S_V formation encourages Mo doping to form an Mo–Sn–Sn trimer site. Such an active site exhibits strong cleavage of the inert $\text{N}\equiv\text{N}$ bond, and the energy barrier is conspicuously reduced with an optimal FE of 20.8% at $-0.40 V_{\text{RHE}}$. Similarly, introducing S_V on the VS_2 basal plane can diminish the high N_2 adsorption barrier on the pristine surface, reducing it from 0.25 to -0.07 eV. Additionally, B-doping could further elevate the activity of S_V -enriched surfaces *via* a synergistic effect. The

electron-deficient B-adjacent-unsaturated-V sites achieve an excellent NH_3 yield of $55.7 \mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$ at $-0.4 V_{\text{RHE}}$ and an FE of 16.4% at $-0.2 V_{\text{RHE}}$, with a reduced N_2 adsorption energy of -0.59 eV.²³ It should be noted that MoS_2 is a prominent HER electrocatalyst, while its similar Mo–S linkages with nitrogenases are expected to be active for ENRR. Sun and co-workers showed that MoS_2 has high ENRR activity and selectivity even under acid conditions;⁴⁵ they further reported a remarkable study in which defect- MoS_2 attains a significantly high FE of 8.34% and NH_3 yield of $29.28 \mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$ at $-0.4 V_{\text{RHE}}$, which are significantly higher than its less-defected counterpart under the same potential (FE: 2.18% and NH_3 yield: $13.42 \mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$).²⁸ Recent studies have further shown that S_V -rich MoS_2 can effectively facilitate N_2 fixation and hydrogenation. For instance, the in-plane defect cluster of MoS_2 obtained a remarkable performance, achieving optimal FE and NH_3 yield of 16.8% and $13.42 \mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$ at $-0.3 V_{\text{RHE}}$, respectively.⁴⁶ Chen and co-workers also demonstrated a significant enhancement in ENRR performance in MoS_2 through S_V introduction (NH_3 yield: $23.38 \mu\text{g h}^{-1} \text{mg}_{\text{cat}}^{-1}$ and FE: 17.9% at $-0.35 V_{\text{RHE}}$). These values represent an approximately twofold improvement compared to pristine MoS_2 .⁴⁷

Notably, Fig. 5b and Table S6† demonstrate the operating potentials of TMS_2 -based catalysts at their highest ENRR-FEs (with data extracted from previous experimental results^{23–25,28,46–54}), which are all consistent with our predicted potential windows for S_V formation (Fig. 5a). Note that six cited reports in Fig. 5b employed isotope labeling experiments to confirm the N-source in their NH_3 synthesis processes.⁴⁸ These findings can be further supported by the experimental evidence that SnS_2 was significantly reduced after ENRR,⁵⁰ and some experiments on metal sulfides showed an increased ENRR performance after the first few cycles.^{51,54,55} The occupations of H_2O , HO , and O on the S-vacancy sites were further analyzed.

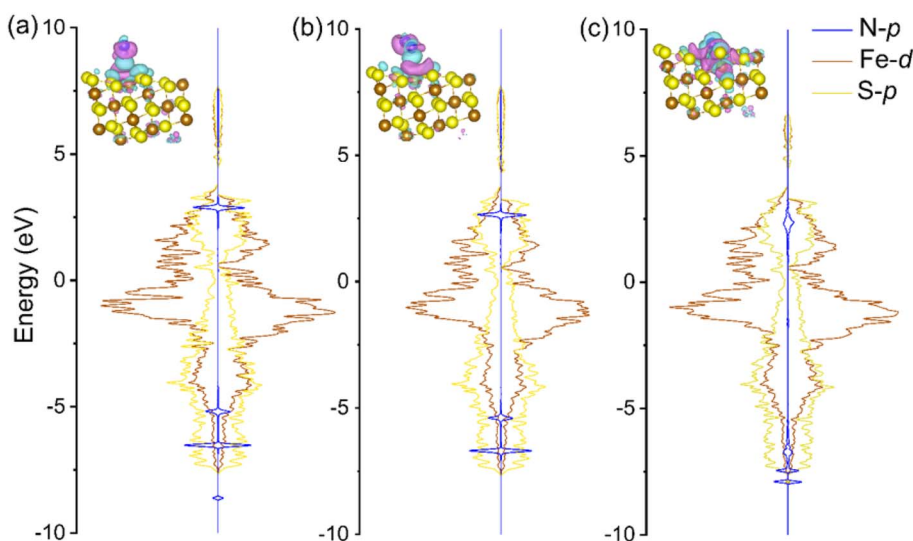


Fig. 4 Calculated projected density of states (PDOS) of N_2^* intermediates on (a) pristine $\text{FeS}_2(111)$, (b) the Fe site on $\text{FeS}_{2-x}(111)-1\text{S}_\text{V}$, and (c) the S_V site on $\text{FeS}_{2-x}(111)-1\text{S}_\text{V}$. The charge density differences upon N_2^* adsorption are shown in insets. Brown, yellow, and blue spheres represent Fe, S, and N, respectively. Pink and cyan isosurfaces indicate electron charge accumulation and depletion, respectively.



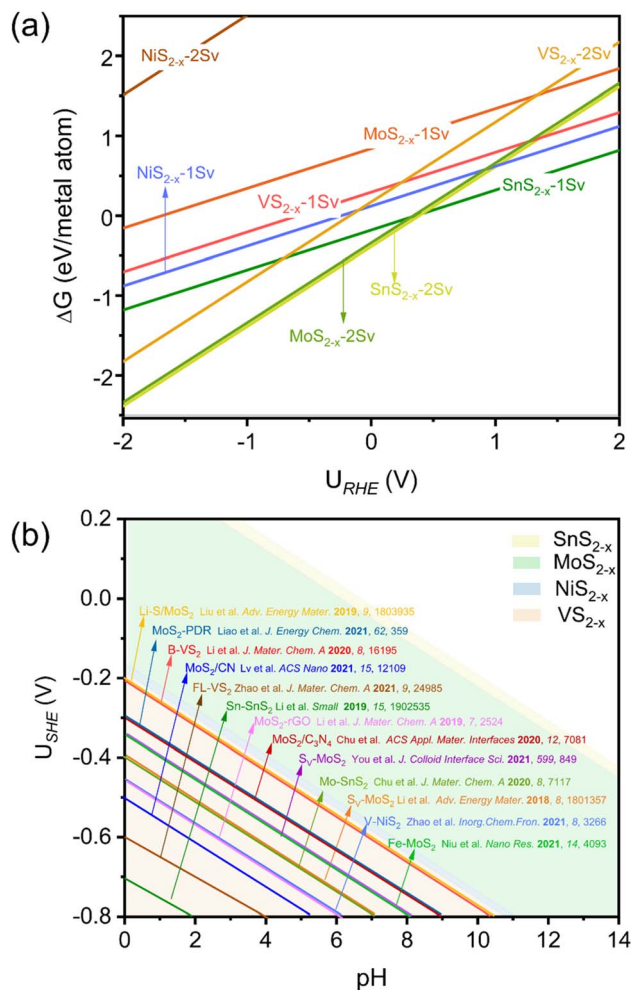


Fig. 5 (a) 1D surface Pourbaix diagrams of different transition metal disulfide (TMS₂) surfaces. (b) 2D surface Pourbaix diagram of S_v formation as a function of pH and potential. The color-coded segments demarcate the potential windows of S_v generation on the respective instances of TMS₂. The reported experimental operating potentials at their highest NH₃ faradaic efficiencies are drawn on the diagram.

Fig. S6† shows that S_v is still the most stable surface state, which is less likely to be further occupied by H₂O, OH, or O. Table S7† further confirms that the adsorption free energy of N₂ is much more negative than the formation free energies of H₂O*, HO*, and O* on FeS_{2-x}, indicating the stronger adsorption capacity of N₂ for ENRR. All in all, our analyses prove that the *in situ* generation of S_v is facile on pristine TMS₂ catalysts under ENRR operating conditions, which holds great promise for facilitating N₂ fixation and hydrogenation in ENRR. Specifically, even if there are no S-vacancies after the synthesis of TMS₂, these highly active S-vacancies will still be generated simultaneously under ENRR operating conditions. Based on our surface Pourbaix diagrams (Fig. 2 and 5), surface S-vacancies are formed on aforementioned TMSs under operating conditions that are favorable for ENRR, which are more realistic surfaces under electrochemical conditions. Currently, most theoretical studies have either considered a pristine surface as

the active site of TMS₂ or have directly created artificial S-vacancies for theoretical activity analysis, which might lead to high uncertainty in the analysis because they generally dismiss the fact that S-vacancies can easily be generated under ENRR operating conditions. Therefore, it is crucial to investigate a more realistic surface state for a TMS₂ ENRR catalyst, which will ensure a deeper understanding of catalytic performance and reaction mechanisms. Our study also provides guidance for designing promising TMS₂ catalysts for ENRR, where the capacity to form S-vacancies is a key performance indicator. TMS₂ surfaces are prone to forming surface S-vacancies spontaneously under ENRR operating conditions, and better N₂ adsorption and reduction capacity is observed. However, it should be noted that, when designing a high-performance TMS₂ catalyst for ENRR, the bonding strength of its surface-S should neither be too strong nor too weak. S-binding that is too strong will lead to difficulty in forming highly active S-vacancies, while S-binding that is too weak will mean the materials are not stable under the operating conditions.

Conclusion

In summary, using *ab initio* calculations, we have analyzed the origin of the ENRR activity of pristine TMS₂ catalysts, starting by probing their electrochemistry-induced surface states. Our calculated surface Pourbaix diagrams found that S-vacancies can be generated directly over a pristine TMS₂ surface under ENRR operating conditions. Using FeS₂ as a typical example, we found that this electrochemistry-driven “*in situ*” generation of S-vacancies possesses a significantly higher ENRR activity than stoichiometric pristine FeS₂ due to the stronger N–N adsorption and activation capacity of a lower-coordination-number S-vacancy site. This finding is in excellent agreement with the experimental observations published in recent years in terms of potential windows reaching the maximum FES. Expanding the analysis to other typical TMS₂ which had shown promising ENRR performance in recent experimental literature (e.g., SnS₂, MoS₂, NiS₂, and VS₂), we found that such an “*in situ*” S-vacancy generation phenomenon is universal under ENRR potentials, with the results generally in good agreement with the experimental observations reported to date. We concluded that, though S-vacancy engineering during synthesis is a promising strategy to enhance the ENRR performance of TMS₂ catalysts, the “*in situ*” generation of S-vacancies will also endow pristine TMS₂ with a measurable ENRR performance. This study shows that the electrochemistry-induced surface states of ENRR catalysts should not be dismissed before analyzing the activity of an ENRR catalyst. Most importantly, when designing a promising TMS₂ catalyst for ENRR, its capacity to form S-vacancies is a key performance indicator that needs to be analyzed.

Methods

Spin-polarized DFT calculations were employed to optimize structural geometries and calculate free energy changes *via* the Vienna *ab initio* simulation package.⁵⁶ The generalized gradient approximation (GGA) method with the revised Perdew–Burke–



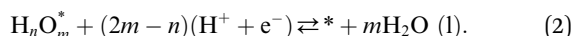
Ernzerhof (RPBE) functional was used to treat exchange–correlation interactions.⁵⁷ The projected augmented-wave method was applied to describe core electrons, and valence electrons were described *via* Kohn–Sham wave functions.^{58,59} A 400 eV energy cutoff was adopted, and *k*-points in the Brillouin zone were sampled with a $3 \times 3 \times 1$ grid by the Monkhorst–Pack scheme.⁶⁰ A 15 Å vacuum space was built to prohibit interactions between periodic units. All geometries were optimized when the residual force was lower than 0.02 eV Å^{-1} , and the energy convergence was set to $1 \times 10^{-4} \text{ eV}$. The binding interactions between nitrogen-containing species and the catalyst were corrected by the DFT-D3 method.⁶¹ *Ab initio* molecular dynamics (AIMD) simulations were employed to analyse the stability of the vacancy-containing surface at 300 K with a time step of 0.2 fs and an overall time of 10 ps. The climbing-image nudged elastic band (CI-NEB) method was employed to calculate transition states.⁶²

Surface energy (E_{surf}) serves as a critical metric for stability assessment, where a lower E_{surf} is indicative of higher stability. The following equation was employed to calculate E_{surf} .^{63,64}

$$E_{\text{surf}} = \frac{E_{\text{slab}} - aE_{\text{bulk}}}{2S}, \quad (1)$$

where E_{slab} and E_{bulk} are the free energies of the slab and bulk, respectively, a is the atomic ratio between the slab and bulk, and S is the surface area.

In this study, the computational hydrogen electrode (CHE) method was employed to establish the surface Pourbaix diagram as a function of pH and potential.⁵⁷ The water dissociation equilibrium during electrochemical reaction conditions is described by eqn (2) because a pristine surface can be covered by O^* , HO^* , and H^* .^{30,38,48,65,66}



For the surface Pourbaix diagrams, the free energy of each surface state was calculated *via* eqn (3):

$$E_{\text{SP}} = E_{\text{slab}} + mE_{\text{H}_2\text{O}} - E_{\text{total}} - (2m - n)\left(\frac{1}{2}E_{\text{H}_2} - U_{\text{SHE}} - 0.2303k_{\text{B}}T \times \text{pH}\right), \quad (3)$$

where n and m are the numbers of adsorbed H^* , O^* , and HO^* , respectively, on a pristine surface. E_{slab} , $E_{\text{H}_2\text{O}}$, E_{H_2} , and E_{total} are the energies of the pristine surface, H_2O molecule, H_2 molecule, and surfaces with adsorbed species, respectively. U_{SHE} , k_{B} , and T are the potential under the scale of standard hydrogen electrode (SHE), the Boltzmann constant, and temperature, respectively.

The stability of the S_{V} -containing surface was analyzed in the surface Pourbaix diagram *via* eqn (4):

$$E_{\text{SV}} = E_{\text{bare}} + l\left(\frac{1}{2}E_{\text{H}_2} - U_{\text{SHE}} - 0.2303k_{\text{B}}T \times \text{pH}\right) - E_{\text{total}} - lE_{\text{H}_2\text{S}}, \quad (4)$$

where l , $E_{\text{H}_2\text{S}}$, and E_{SV} are the number of S_{V} , and the total energies of the H_2S molecule and S_{V} -containing surface, respectively.

The Gibbs free energy change of each hydrogenation step in ENRR was calculated *via* the following equation:

$$\Delta G = \Delta E_{\text{DFT}} + \Delta E_{\text{ZPE}} - T\Delta S, \quad (5)$$

where ΔE_{DFT} is the binding energy difference of an adsorbed molecule obtained from DFT calculations, ΔE_{ZPE} is the zero-point energy difference, and $T\Delta S$ is the entropy correction acquired from a database.⁶⁷

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the JSPS KAKENHI (No. JP23K13703, JP23KF0102, JP19H05814, and JP23H00301), AIMR Fusion Research & Advanced Target Project, Ensemble Grant for Early Career Researchers 2023, JST PRESTO Grant Number: JPMJPR2274, and Tanaka Kikinzoku Memorial Foundation. We acknowledge the Center for Computational Materials Science, Institute for Materials Research, Tohoku University for the use of MASAMUNE-IMR (202212-SCKXX-0204) and Institute for Solid State Physics (ISSP) at the University of Tokyo for the use of their supercomputers.

References

- 1 D. Bao, Q. Zhang, F. L. Meng, H. X. Zhong, M. M. Shi, Y. Zhang, J. M. Yan, Q. Jiang and X. B. Zhang, *Adv. Mater.*, 2017, **29**, 1604799.
- 2 S. Wu, N. Salmon, M. M. J. Li, R. Bañares-Alcántara and S. C. E. Tsang, *ACS Energy Lett.*, 2022, **7**, 1021–1033.
- 3 D. Wang, L. M. Azofra, M. Harb, L. Cavallo, X. Zhang, B. H. R. Suryanto and D. R. MacFarlane, *ChemSusChem*, 2018, **11**, 3416–3422.
- 4 C. Liu, Q. Li, J. Zhang, Y. Jin, D. R. MacFarlane and C. Sun, *J. Phys. Chem. C*, 2018, **122**, 25268–25273.
- 5 B. Yu, H. Li, J. White, S. Donne, J. Yi, S. Xi, Y. Fu, G. Henkelman, H. Yu, Z. Chen and T. Ma, *Adv. Funct. Mater.*, 2019, **30**, 1905665.
- 6 Q. Li, C. Liu, S. Qiu, F. Zhou, L. He, X. Zhang and C. Sun, *J. Mater. Chem. A*, 2019, **7**, 21507–21513.
- 7 T. Wang, Z. Guo, X. Zhang, Q. Li, A. Yu, C. Wu and C. Sun, *J. Mater. Sci. Technol.*, 2023, **140**, 121–134.
- 8 Y. Fu, Y. Liao, P. Li, H. Li, S. Jiang, H. Huang, W. Sun, T. Li, H. Yu, K. Li, H. Li, B. Jia and T. Ma, *Coord. Chem. Rev.*, 2022, **460**, 214468.
- 9 Y. Yao, S. Zhu, H. Wang, H. Li and M. Shao, *J. Am. Chem. Soc.*, 2018, **140**, 1496–1501.
- 10 J. Wang, L. Yu, L. Hu, G. Chen, H. Xin and X. Feng, *Nat. Commun.*, 2018, **9**, 1795.
- 11 H. M. Liu, S. H. Han, Y. Zhao, Y. Y. Zhu, X. L. Tian, J. H. Zeng, J. X. Jiang, B. Y. Xia and Y. Chen, *J. Mater. Chem. A*, 2018, **6**, 3211–3217.
- 12 O. Einsle and D. C. Rees, *Chem. Rev.*, 2020, **120**, 4969–5004.



- 13 J. Humphreys, R. Lan and S. Tao, *Adv. Energy Sustainability Res.*, 2020, **2**, 2000043.
- 14 M. I. Ahmed, L. J. Arachchige, Z. Su, D. B. Hibbert, C. Sun and C. Zhao, *ACS Catal.*, 2022, **12**, 1443–1451.
- 15 F. He, Z. Li, S. Shi, W. Xu, H. Sheng, Y. Gu, Y. Jiang and B. Xi, *Environ. Sci. Technol.*, 2018, **52**, 8627–8637.
- 16 J. Xu, Y. Wang, C. Weng, W. Bai, Y. Jiao, R. Kaegi and G. V. Lowry, *Environ. Sci. Technol.*, 2019, **53**, 5936–5945.
- 17 H. Li, W. Yang, C. Wu and J. Xu, *Phys. Chem. Chem. Phys.*, 2021, **23**, 13971–13976.
- 18 L. Gao, C. Guo, M. Zhao, H. Yang, X. Ma, C. Liu, X. Liu, X. Sun and Q. Wei, *ACS Appl. Mater. Interfaces*, 2021, **13**, 50027–50036.
- 19 D. Feng, X. Zhang, Y. Sun and T. Ma, *Nano Mater. Sci.*, 2020, **2**, 132–139.
- 20 Z. Feng, G. Li, X. Wang, C. J. Gómez-García, J. Xin, H. Ma, H. Pang and K. Gao, *Chem. Eng. J.*, 2022, **445**, 136797.
- 21 M. Yang, Z. Jin, C. Wang, X. Cao, X. Wang, H. Ma, H. Pang, L. Tan and G. Yang, *ACS Appl. Mater. Interfaces*, 2021, **13**, 55040–55050.
- 22 H. B. Wang, J. Q. Wang, R. Zhang, C. Q. Cheng, K. W. Qiu, Y. J. Yang, J. Mao, H. Liu, M. Du, C. K. Dong and X. W. Du, *ACS Catal.*, 2020, **10**, 4914–4921.
- 23 Q. Li, Y. Guo, Y. Tian, W. Liu and K. Chu, *J. Mater. Chem. A*, 2020, **8**, 16195–16202.
- 24 K. Chu, J. Wang, Y. P. Liu, Q. Q. Li and Y. L. Guo, *J. Mater. Chem. A*, 2020, **8**, 7117–7124.
- 25 M. Zhao, C. Guo, L. Gao, X. Kuang, H. Yang, X. Ma, C. Liu, X. Liu, X. Sun and Q. Wei, *Inorg. Chem. Front.*, 2021, **8**, 3266–3272.
- 26 G. Zhang, F. Wang, K. Chen, J. Kang and K. Chu, *Adv. Funct. Mater.*, 2023, 2305372.
- 27 H. Du, C. Yang, W. Pu, L. Zeng and J. Gong, *ACS Sustain. Chem. Eng.*, 2020, **8**, 10572–10580.
- 28 X. Li, T. Li, Y. Ma, Q. Wei, W. Qiu, H. Guo, X. Shi, P. Zhang, A. M. Asiri, L. Chen, B. Tang and X. Sun, *Adv. Energy Mater.*, 2018, **8**, 1801357.
- 29 V. Sumaria, D. Krishnamurthy and V. Viswanathan, *ACS Catal.*, 2018, **8**, 9034–9042.
- 30 H. Liu, X. Jia, A. Cao, L. Wei, C. D'agostino and H. Li, *J. Chem. Phys.*, 2023, **12**, 158.
- 31 S. Pan, H. Li, D. Liu, R. Huang, X. Pan, D. Ren, J. Li, M. Shakouri, Q. Zhang, M. Wang, C. Wei, L. Mai, B. Zhang, Y. Zhao, Z. Wang, M. Graetzel and X. Zhang, *Nat. Commun.*, 2022, **13**, 2294.
- 32 R. Zhao, Z. Chen, Q. Li, X. Wang, Y. Tang, G. Fu, H. Li, J. M. Lee and S. Huang, *Chem Catal.*, 2022, **2**, 3590–3606.
- 33 W. Yang, Z. Jia, B. Zhou, L. Wei, Z. Gao and H. Li, *Commun. Chem.*, 2023, **6**, 6.
- 34 H. Liu, D. Zhang, S. M. Holmes, C. D'Agostino and H. Li, *Chem. Sci.*, 2023, **14**, 9000–9009.
- 35 S. Kment, H. Kmentova, A. Sarkar, R. J. Soukup, N. J. Ianno, D. Sekora, J. Olejnicek, P. Ksirova, J. Krysa, Z. Remes and Z. Hubicka, *J. Alloys Compd.*, 2014, **607**, 169–176.
- 36 M. Cabán-Acevedo, D. Liang, K. S. Chew, J. P. DeGrave, N. S. Kaiser and S. Jin, *ACS Nano*, 2013, **7**, 1731–1739.
- 37 A. B. Puthirath, A. P. Balan, E. F. Oliveira, V. Sreepal, F. C. R. Hernandez, G. Gao, N. Chakingal, L. M. Sassi, P. Thibeorchews, G. Costin, R. Vajtai, D. S. Galvao, R. R. Nair and P. M. Ajayan, *J. Phys. Chem. C*, 2021, **125**, 18927–18935.
- 38 H. A. Hansen, J. Rossmeisl and J. K. Nørskov, *Phys. Chem. Chem. Phys.*, 2008, **10**, 3722–3730.
- 39 C. Yang, Y. Zhu, J. Liu, Y. Qin, H. Wang, H. Liu, Y. Chen, Z. Zhang and W. Hu, *Nano Energy*, 2020, **77**, 105126.
- 40 A. R. Singh, B. A. Rohr, M. J. Statt, J. A. Schwalbe, M. Cargnello and J. K. Nørskov, *ACS Catal.*, 2019, **9**, 8316–8324.
- 41 J. H. Montoya, C. Tsai, A. Vojvodic and J. K. Nørskov, *ChemSusChem*, 2015, **8**, 2180–2186.
- 42 X. Wen, Y. Liang, P. Bai, B. Luo, T. Fang, L. Yue, T. An, W. Song and S. Zheng, *Phys. B*, 2017, **525**, 119–126.
- 43 A. Kjekshus and T. Rakke, *Acta Chem. Scand., Ser. A*, 1975, **29**, 443–452.
- 44 P. Bayliss, *Am. Mineral.*, 1977, **62**, 1168–1172.
- 45 L. Zhang, X. Ji, X. Ren, Y. Ma, X. Shi, Z. Tian, A. M. Asiri, L. Chen, B. Tang and X. Sun, *Adv. Mater.*, 2018, **30**, e1800191.
- 46 W. Liao, K. Xie, L. Liu, X. Wang, Y. Luo, S. Liang, F. Liu and L. Jiang, *J. Energy Chem.*, 2021, **62**, 359–366.
- 47 M. You, S. Yi, X. Hou, Z. Wang, H. Ji, L. Zhang, Y. Wang, Z. Zhang and D. Chen, *J. Colloid Interface Sci.*, 2021, **599**, 849–856.
- 48 Y. Liu, M. Han, Q. Xiong, S. Zhang, C. Zhao, W. Gong, G. Wang, H. Zhang and H. Zhao, *Adv. Energy Mater.*, 2019, **9**, 1803935.
- 49 K. Chu, Y. P. Liu, Y. B. Li, Y. L. Guo and Y. Tian, *ACS Appl. Mater. Interfaces*, 2020, **12**, 7081–7090.
- 50 P. Li, W. Fu, P. Zhuang, Y. Cao, C. Tang, A. B. Watson, P. Dong, J. Shen and M. Ye, *Small*, 2019, **15**, e1902535.
- 51 X. Li, X. Ren, X. Liu, J. Zhao, X. Sun, Y. Zhang, X. Kuang, T. Yan, Q. Wei and D. Wu, *J. Mater. Chem. A*, 2019, **7**, 2524–2528.
- 52 X. W. Lv, X. L. Liu, Y. J. Suo, Y. P. Liu and Z. Y. Yuan, *ACS Nano*, 2021, **15**, 12109–12118.
- 53 L. Niu, D. Wang, K. Xu, W. Hao, L. An, Z. Kang and Z. Sun, *Nano Res.*, 2021, **14**, 4093–4099.
- 54 L. Zhao, R. Zhao, Y. Zhou, X. Wang, X. Chi, Y. Xiong, C. Li, Y. Zhao, H. Wang, Z. Yang and Y. M. Yan, *J. Mater. Chem. A*, 2021, **9**, 24985–24992.
- 55 J. Wang, H. Nan, Y. Tian and K. Chu, *ACS Sustain. Chem. Eng.*, 2020, **8**, 12733–12740.
- 56 G. Kresse and J. Furthmüller, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1996, **16**, 11169.
- 57 J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **18**, 3865.
- 58 W. Kohn and L. J. Sham, *Phys. Rev.*, 1965, **140**, A1133–A1138.
- 59 P. E. Blochl, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **50**, 17953–17979.
- 60 H. J. Monkhorst and J. D. Pack, *Phys. Rev. B: Solid State*, 1976, **13**, 5188–5192.
- 61 S. Grimme, J. Antony, S. Ehrlich and H. Krieg, *J. Chem. Phys.*, 2010, **132**, 154104.



- 62 G. Henkelman, B. P. Uberuaga and H. Jónsson, *J. Chem. Phys.*, 2000, **113**, 9901–9904.
- 63 M. Li, H. Li, X. Jiang, M. Jiang, X. Zhan, G. Fu, J. M. Lee and Y. Tang, *J. Mater. Chem. A*, 2021, **9**, 2999–3006.
- 64 W. Xiong, M. Zhou, X. Huang, W. Yang, D. Zhang, Y. Lv and H. Li, *Chemistry*, 2022, **28**, e202200779.
- 65 A. E. Russell, *Phys. Chem. Chem. Phys.*, 2008, **10**, 3607–3608.
- 66 J. K. Nørskov, J. Rossmeisl, A. Logadottir, L. Lindqvist, J. R. Kitchin, T. Bligaard and H. Jónsson, *J. Phys. Chem. B*, 2004, **108**, 17886–17892.
- 67 R. D. Johnson III, *NIST 101. Computational Chemistry Comparison and Benchmark Database*, 1999.

