

Cite this: *J. Mater. Chem. A*, 2020, **8**, 7117Received 12th February 2020
Accepted 24th March 2020

DOI: 10.1039/d0ta01688h

rsc.li/materials-a

Mo-doped SnS₂ with enriched S-vacancies for highly efficient electrocatalytic N₂ reduction: the critical role of the Mo–Sn–Sn trimer†

Ke Chu,[✉] Jing Wang, Ya-ping Liu,[✉] Qing-qing Li and Ya-li Guo

Vacancy engineering and heteroatom doping are two effective approaches to tailor the electronic structures of catalysts for improved electrocatalytic activity. Herein, these two approaches were rationally combined to modulate the structure of SnS₂ toward the N₂ reduction reaction (NRR) by means of Mo-doping, which simultaneously induced the generation of enriched S-vacancies (V_s). The developed Mo-doped SnS₂ nanosheets with enriched V_s presented a conspicuously enhanced NRR activity with an NH₃ yield of 41.3 μg h⁻¹ mg⁻¹ (-0.5 V) and a faradaic efficiency of 20.8% (-0.4 V) and are among the best SnS₂-based NRR catalysts to date. Mechanistic studies revealed that the co-presence of the Mo dopant and V_s enabled the creation of Mo–Sn–Sn trimer catalytic sites, capable of strongly activating N₂ even for the cleavage of the N≡N triple bond to the N=N double bond at the N₂ adsorption stage, consequently leading to a downhill process of the first hydrogenation step and a largely reduced energy barrier.

Dinitrogen (N₂) fixation to ammonia (NH₃) is an essential chemical process for the modern industrial society because NH₃ is a widely used feedstock for various chemical products and is also a promising carbon-free energy carrier.¹ The electrochemical N₂ reduction reaction (NRR) under ambient conditions holds great promise for green and sustainable NH₃ production,² in stark contrast to the industrial Haber–Bosch process which involves huge energy consumption and enormous CO₂ emission. However, the NRR process is largely limited by the unsatisfactory NH₃ production rate and low faradaic efficiency (FE), stemming from the extremely stable N≡N bond, poor N₂ adsorption/activation on the catalyst surface and the competing hydrogen evolution reaction (HER).³ Exploring highly active electrocatalysts may hold the key to mitigating these limitations and realizing high NRR efficiency. Over the past three years, an increasingly large number of noble metals,^{4–7} transition metal-based compounds,^{8–14} and metal-free

materials^{15–19} have been designed as promising candidates toward the NRR.

SnS₂-based materials have shown great potential in electrolysis applications, owing to their fascinating catalytic performance, good stability, tunable electronic structures, and 2D morphology with largely exposed active edges.²⁰ Recently, SnS₂ has been proved to be an attractive catalyst with favorable NRR activity as well.^{21,22} Nonetheless, the reported NRR performance of SnS₂ materials is still far from satisfactory and competitive with that of the state-of-the-art NRR catalysts. The main bottlenecks are the poor NRR activation, *i.e.*, the high energy barrier for the first hydrogenation step,²² and the low intrinsic electrical conductivity, *i.e.*, the inefficient proton-coupled electron transfer process, even when employing approaches such as growth of amorphous Sn on crystalline SnS₂ nanosheets (FE: 6.5%)²¹ and construction of well-aligned SnS₂ nanoarrays on porous Ni foam (FE: 11.2%).²² Thus, more efficient methods are still needed to further enhance the intrinsic NRR activity of SnS₂.

As an effective strategy for electronic structure modulation of the catalysts, vacancy engineering by introducing anion vacancies into the catalyst lattice,²³ such as oxygen vacancies (TiO₂,²⁴ WO₃ (ref. 25) and MnO₂ (ref. 26)), and nitrogen vacancies (C₃N₄,²⁷ VN²⁸ and W₂N₃ (ref. 29)), provides unique active sites for the effective adsorption and activation of dinitrogen. In addition, heteroatom doping is another widely used method to optimize the adsorption of NRR intermediates on catalysts and promote the NRR reaction kinetics,^{30–33} which have been demonstrated in V-doped TiO₂,³⁴ Fe-doped Ni₂P³⁵ and Mo-doped MnO₂.³⁶ More importantly, heteroatom doping has been recently confirmed to facilitate the generation of vacancies, while the synergistic effect of dopants and vacancies leads to a conspicuously enhanced NRR performance.^{37,38} For instance, introducing Zr-dopants in TiO₂ nanotubes has been reported to favorably generate both enriched oxygen vacancies and bi-Ti³⁺ pairs,³⁸ and the bi-Ti³⁺ pairs serve as the dominant active sites in facilitating the catalytic performance of the NRR with an FE of 17.3%. Similar results and a more improved NRR

School of Materials Science and Engineering, Lanzhou Jiaotong University, Lanzhou 730070, China. E-mail: chukelut@163.com

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0ta01688h

performance (FE: 25.6%) have been presented in Fe-doped TiO₂ nanoparticles,³⁷ where the combined effect of bi-Ti³⁺ pairs and oxygen vacancies contributes to the substantially boosted NRR activity. Inspired by these studies, it is anticipated that the NRR performance of SnS₂ may be greatly enhanced by the combination of vacancy engineering and heteroatom doping.

In this study, we rationally combined vacancy engineering and heteroatom doping to regulate the electronic structure of SnS₂ by Mo-doping, which enabled the spontaneous generation of enriched S-vacancies (V_s) as well. The developed Mo-doped SnS₂ nanosheets with enriched V_s exhibited a significantly enhanced NRR activity with an NH₃ yield of 41.3 μg h⁻¹ mg⁻¹ (-0.5 V) and an FE of 20.8% (-0.4 V) and were among the best SnS₂-based NRR catalysts to date. Density functional theory (DFT) calculations revealed that the co-presence of the Mo dopant and V_s enabled the creation of Mo-Sn-Sn trimer active sites that could strongly activate N₂ and lower the reaction energy barrier.

Mo-doped SnS₂ (Mo-SnS₂) nanosheets were directly grown on CC *via* a facile hydrothermal method. Pristine SnS₂/CC was also prepared as a reference. The synthesis details are provided in the ESI.† The obtained SnS₂ and Mo-SnS₂ nanosheets scraped from CC were first investigated by X-ray photoelectron spectroscopy (XPS). As shown in Fig. 1a, the deconvolution of the Mo3d spectrum results in three well-resolved peaks of S2s (226.4 eV), Mo⁴⁺3d_{3/2} (229.3 eV) and Mo⁴⁺3d_{1/2} (232.7 eV) for Mo-SnS₂ nanosheets, which are absent in pristine SnS₂ nanosheets, indicating the successful Mo-doping in the Mo-SnS₂ nanosheets. Fig. 1b shows the XRD patterns of SnS₂/CC and Mo-SnS₂/CC. Excluding the peaks of CC, both samples display a pure hexagonal SnS₂ phase (JCPDS no. 23-677) with no detectable impurities. Obviously, all the peaks of Mo-SnS₂/CC are much weaker than those of SnS₂/CC, indicating the lowered crystallinity of Mo-SnS₂ caused by Mo-doping. The morphologies of SnS₂ and Mo-SnS₂ nanosheets were further investigated by scanning electron microscopy (SEM), transmission electron microscopy (TEM) and high-resolution TEM (HRTEM). The SEM images show that bushy nanosheets are densely and vertically anchored on the CC for both SnS₂/CC (Fig. 1c) and Mo-SnS₂/CC (Fig. 1d). The nanosheet features can be further revealed by TEM with the observation of clear wrinkles and corrugations on SnS₂ (Fig. 1e) and Mo-SnS₂ (Fig. 1f) nanosheets.³⁹⁻⁴¹ The HRTEM image shows that the pristine SnS₂ nanosheets (Fig. 1g) possess a high crystallinity as evidenced by the sharp lattice fringes with a spacing value of 0.59 nm, assigned to the (001) facet of SnS₂. In contrast, blurred and discontinuous lattice fringes can be observed in Mo-SnS₂ nanosheets (Fig. 1h), indicating that Mo-doping can remarkably reduce the crystallinity of Mo-SnS₂ nanosheets, consistent with the XRD analysis (Fig. 1b). The selected areas in Fig. 1h (A and B squares), when applying an Inverse Fast Fourier Transform (IFFT) mask, display obvious distortions and dislocations in the lattice fringes,⁴²⁻⁴⁴ suggesting the existence of plentiful defects (*i.e.*, vacancies) in Mo-SnS₂ nanosheets, which rationalizes the reduced crystallinity of Mo-SnS₂. The scanning TEM (STEM) elemental mapping images (Fig. 1i) reveal that considerable Mo dopants are uniformly distributed on the entire surface of Mo-SnS₂ nanosheets.

The vacancies existing in Mo-SnS₂ nanosheets were further investigated by a combination of XPS, electron paramagnetic resonance (EPR) spectroscopy and DFT analysis. For both SnS₂ and Mo-SnS₂ nanosheets, the XPS Sn3d spectra (Fig. 2a) reveal Sn3d_{5/2} (~486 eV) and Sn3d_{3/2} (~495 eV) states of Sn⁴⁺, and the S2p spectra (Fig. 2b) reveal S2p_{3/2} (~161 eV) and S2p_{1/2} (~163 eV) states of S²⁻. With respect to those of the pristine SnS₂ nanosheets, the Sn3d and S2p spectra of Mo-SnS₂ nanosheets are negatively shifted by 0.4 and 0.3 eV, respectively, suggesting the decreased valence state and increased electrons in Mo-SnS₂ nanosheets caused by the presence of V_s.⁴⁵ The existence of enriched V_s in Mo-SnS₂ nanosheets can also be directly verified from the remarkably reduced atomic ratio of S/Mo from 1.88 (SnS₂) to 1.62 (Mo-SnS₂) derived from the XPS element analysis. In the EPR spectra (Fig. 2c), the signal at *g* = 2.001 represents the electrons trapped in V_s. Compared to SnS₂ nanosheets, Mo-SnS₂ nanosheets exhibit a much stronger EPR signal intensity, suggesting a higher concentration of V_s,²³ consistent with the XPS results. To gain further insight, DFT was applied to predict the V_s formation energy (*E_f*). As shown in Fig. 2d, the formation of isolated V_s in SnS₂-V_s requires a largely positive *E_f* (5.25 eV) and thus is thermodynamically unfavorable. With the introduction of the Mo dopant (Fig. 2e), V_s can be favorably formed at the Mo-adjacent site of Mo-SnS₂-V_s with a significantly reduced *E_f* (-0.78 eV). Therefore, the DFT results concur well with the XPS/EPR measurements and demonstrate that Mo-doping can considerably lower the V_s formation energy and enable the spontaneous generation of V_s in Mo-SnS₂ nanosheets.

DFT calculations were used to further investigate the electronic structure of Mo-SnS₂-V_s. From the electron contour maps sliced along the (001) plane, it can be seen that more electrons are accumulated in the Mo/V_s region in Mo-SnS₂-V_s (Fig. 2g) as compared to SnS₂ (Fig. 2f), which is also confirmed by the differential charge density (Fig. S1†). Upon N₂ adsorption, it is believed that these Mo/V_s-induced abundant accumulated electrons can be easily transferred into the anti-bonding orbitals of N₂ molecules for the weakening and dissociation of the N≡N bond.⁴⁶ Projected density of states (PDOS, Fig. 2h) analysis indicates that SnS₂ possesses a 0.64 eV bandgap indicative of its semiconducting character. In contrast, the co-presence of the Mo-dopant and V_s creates noticeable electronic states crossing the Fermi level, leading to the metallic characteristics of Mo-SnS₂-V_s and thus higher conductivity relative to that of SnS₂, which is favorable for the proton-coupled electron-transfer process to boost the NRR kinetics. Meanwhile, the calculated work function (Fig. 2i & S2†) indicates that Mo-SnS₂-V_s (5.513 eV) possesses a lower work function than SnS₂ (6.758 eV), which suggests the higher capability of Mo-SnS₂-V_s for electron back-donation from its active sites to the absorbed N₂ and NRR intermediates,⁴⁷⁻⁴⁹ thus facilitating N₂ adsorption, activation, and hydrogenation.

The electrocatalytic NRR performance of Mo-SnS₂/CC as a self-standing electrode was examined in N₂-saturated 0.5 M LiClO₄ using a gas-tight two-compartment cell, as displayed in Fig. S3.† An absorber was placed at the end of the cell to prevent the loss of produced NH₃ by N₂ flow during the NRR test.⁵⁰ All potentials were converted into values *versus* the reversible

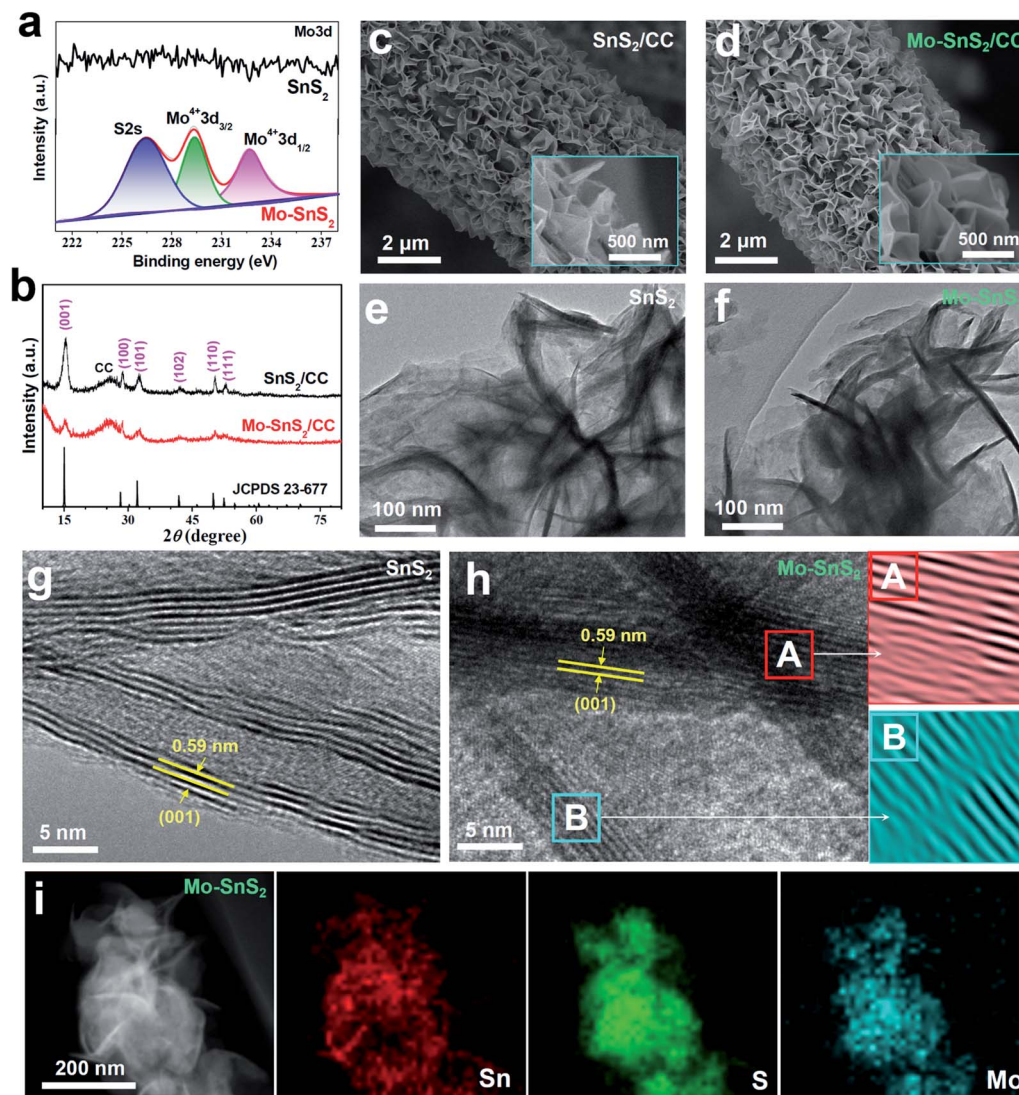


Fig. 1 (a) XPS Mo3d spectrum of SnS₂ and Mo-SnS₂ nanosheets scraped from CC. (b) XRD patterns for SnS₂/CC and Mo-SnS₂/CC. (c and d) SEM images of (c) SnS₂/CC and (d) Mo-SnS₂/CC. (e and f) TEM images of (e) SnS₂ and (f) Mo-SnS₂ nanosheets. (g and h) HRTEM images of (g) SnS₂ and (h) Mo-SnS₂ nanosheets (A and B are the corresponding IFFT images recorded from regions A and B in (h)). (i) STEM element mapping images for Mo-SnS₂ nanosheets.

hydrogen electrode (*vs.* RHE), and the standard RHE was experimentally calibrated using cyclic voltammetry curves in a high-purity hydrogen saturated solution (Fig. S4†).³⁶ The concentration of generated NH₃ was experimentally determined by the indophenol blue method,⁵¹ while the concentration of the possible N₂H₄ as a byproduct was measured by the Watt-Chrisp approach.⁵² Their standard calibration curves are shown in Fig. S5 and S6.† As displayed in Fig. S7,† N₂H₄ can hardly be detected, implying a high NRR selectivity for Mo-SnS₂/CC for N₂-to-NH₃ conversion.

The polarization curves recorded by linear sweep voltammetry (LSV, Fig. 3a) show a higher current density in N₂-saturated solution than in Ar-saturated solution, suggesting the feasibility of the NRR on Mo-SnS₂/CC. Before the quantitative evaluation of the NRR performance, a series of control tests were preliminarily carried out to exclude the possible influences

from any nitrogen contaminants.⁵³ The isotopic labeling measurements based on ¹H nuclear magnetic resonance (NMR) were first utilized to trace the origin of the N source, as shown in Fig. 3b. After NRR electrolysis using ¹⁴N₂ or ¹⁵N₂ as the feed gas, a triplet for ¹⁴NH₄⁺ (¹J_{N-H} = 52 Hz) or a doublet for ¹⁵NH₄⁺ (¹J_{N-H} = 72 Hz) can be distinguished, whereas no labeled ¹⁴NH₄⁺ or ¹⁵NH₄⁺ can be found when using Ar as the feed gas. In addition, UV-vis analysis (Fig. S8†) in an Ar-saturated solution, or at open circuit, or on pristine CC does not produce a detectable amount of NH₃.^{54–56} Furthermore, the time-dependent test (Fig. S9†) shows that the produced NH₃ increases linearly with the electrolysis time, indicating that NH₃ can be continuously generated by NRR catalysis over Mo-SnS₂/CC. We also employed the ion chromatography (IC) technique to quantitatively determine the concentration of NRR-derived NH₃. As depicted in Fig. 3c and S10,† the IC measurement (1.27 μg mL⁻¹) is very



Fig. 2 (a and b) XPS spectra of SnS₂ and Mo-SnS₂ nanosheets: (a) Sn3d and (b) S2p. (c) EPR spectra of SnS₂ and Mo-SnS₂ nanosheets. (d and e) Optimized SnS₂ structures with isolated V_s (SnS₂-V_s) and Mo-adjacent V_s (Mo-SnS₂-V_s), and the corresponding formation energies (E_f). (f and g) Electron contour maps of the (001) slice for (f) SnS₂ and (g) Mo-SnS₂-V_s. The blue and red regions represent electron depletion and accumulation, respectively. (h) PDOS of SnS₂ and Mo-SnS₂-V_s. (i) Calculated work functions of SnS₂ and Mo-SnS₂-V_s (Fig. S2†).

close to the 1.15 $\mu\text{g mL}^{-1}$ attained by the indophenol blue method with a reasonable margin of experimental error. Therefore, all these control experiments convincingly demonstrate that the produced NH₃ originates from the NRR.

The NRR performance of Mo-SnS₂/CC was then quantitatively examined by the combination of chronoamperometry testing with each cycle for 2 h of electrolysis (Fig. 3d) and UV-vis analysis (average of three measurements) at various potentials. The corresponding data of NH₃ yield and FE are shown in Fig. 3e. As observed, the NRR activity of Mo-SnS₂/CC exhibits optimum NRR performance with an NH₃ yield of 41.3 $\mu\text{g h}^{-1} \text{mg}^{-1}$ at -0.5 V and a faradaic efficiency of 20.8% at -0.4 V. As shown in Table S1,† Mo-SnS₂/CC is the best SnS₂-based NRR catalyst reported to date and compares favorably to most state-of-the-art NRR catalysts. Nonetheless, the NRR performance shows a sharp decline at more negative potentials, which is attributed to the significantly enhanced HER.⁵⁷

We also assessed the NRR performance of undoped SnS₂/CC for comparison under identical conditions. As shown in Fig. 3f, SnS₂/CC possesses a more inferior NRR activity with the highest NH₃ yield of 10.6 $\mu\text{g h}^{-1} \text{mg}^{-1}$ at -0.5 V, which is about one-quarter that of Mo-SnS₂/CC (41.3 $\mu\text{g h}^{-1} \text{mg}^{-1}$), demonstrating that Mo-doping is able to dramatically promote the NRR activity of SnS₂ nanosheets. To elucidate the NRR

enhancement observed in Mo-SnS₂/CC, we determined the electrochemically active surface area (ECSA) by measuring the double-layer capacitance (C_{dl}). As shown in Fig. S11,† Mo-SnS₂/CC exhibits only a 1.3 times higher C_{dl} , but a ~3.9 times higher NH₃ yield than SnS₂/CC, suggesting that the ECSA is not the primary factor and Mo-SnS₂/CC is intrinsically more active than SnS₂/CC. In addition, as depicted in the electrochemical impedance spectra (EIS, Fig. S12†), Mo-SnS₂/CC delivers a smaller charge-transport resistance than SnS₂/CC, indicating faster electron-transfer and enhanced NRR reaction kinetics of Mo-SnS₂/CC. This can be attributed to the synergistic role of the Mo-dopant and V_s in improving the conductivity of Mo-SnS₂, as revealed by the DFT results (Fig. 2h). Therefore, the combined Mo-dopant and V_s can bring about an improved conductivity and elevated intrinsic NRR activity, resulting in significantly enhanced NRR performance of Mo-SnS₂/CC.

We further evaluate the NRR stability which is another critical factor for practical applications.^{58,59} Chronopotentiometric response measurements (Fig. 3g) reveal that the current density over Mo-SnS₂/CC presents a negligible degeneration for at least 20 h of continuous electrolysis, verifying the excellent long-term stability. Besides, when conducting seven chronoamperometric runs for Mo-SnS₂/CC, there is no remarkable change in the UV-vis spectra of the resultant electrolytes (Fig. S13†), nor in the

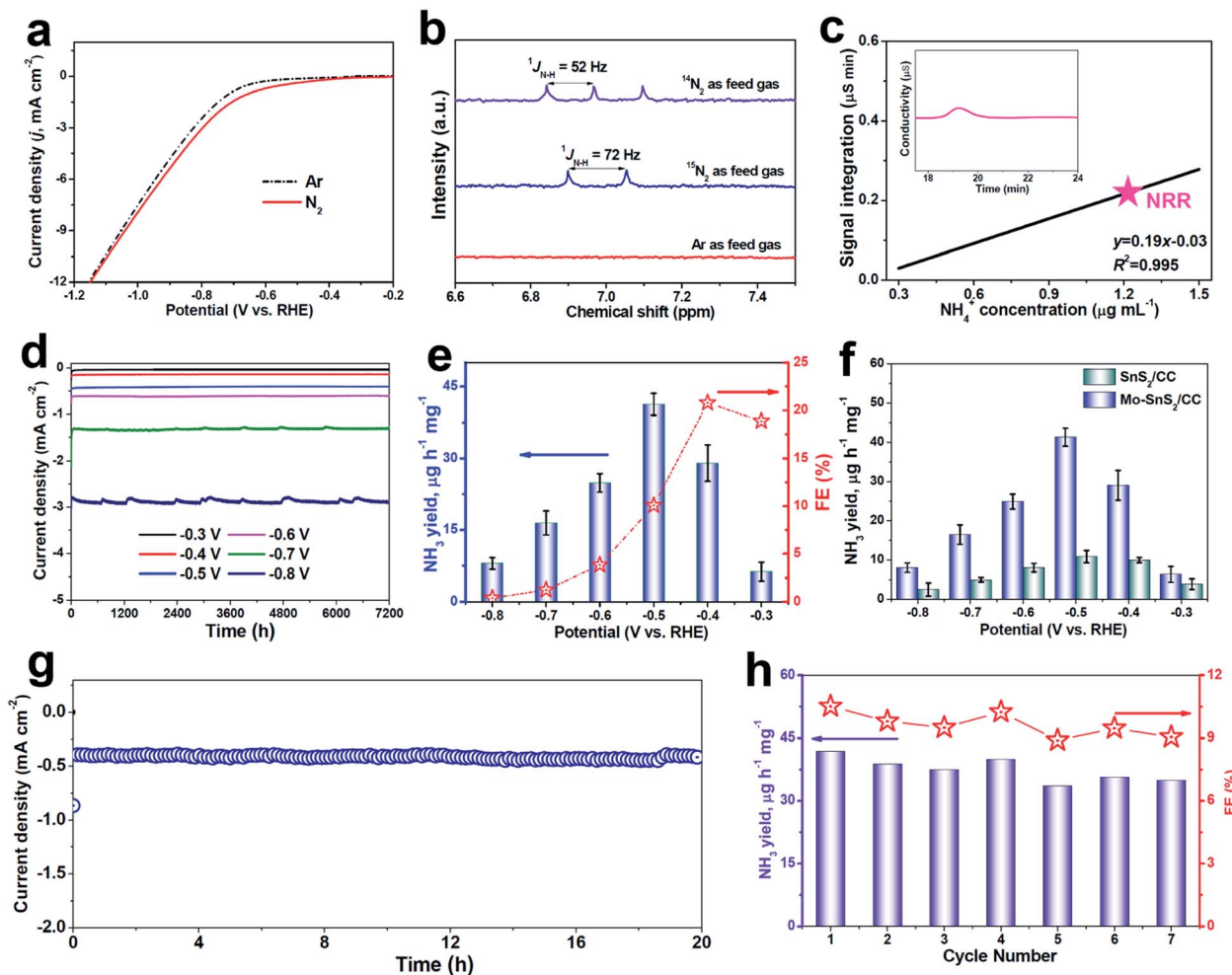


Fig. 3 (a) LSV curves of Mo-SnS₂/CC in Ar- and N₂-saturated solutions. (b) ¹H NMR spectra of electrolytes after NRR electrolysis for 2 h on Mo-SnS₂/CC using ¹⁴N₂, ¹⁵N₂ or Ar as feed gases. (c) IC spectra of the electrolyte after NRR electrolysis on Mo-SnS₂/CC for 2 h at −0.5 V (inset), and the determined NH₄⁺ concentration of the electrolyte by referring to the calibration curve (Fig. S10†). (d) Chronoamperometry test results of Mo-SnS₂/CC for 2 h of NRR electrolysis at various potentials, and (e) the corresponding NH₃ yields and FEs. (f) NH₃ yields of SnS₂/CC and Mo-SnS₂/CC at various potentials. (g) Chronoamperometry test results of Mo-SnS₂/CC for 20 h of NRR electrolysis. (h) NH₃ yields and FEs of Mo-SnS₂/CC for seven cycles (each for 2 h of electrolysis at −0.5 V).

resulting NH₃ yield and FE data (Fig. 3h), confirming the good cycling stability. Further, the morphology, crystal phase and chemical bonding states of Mo-SnS₂ nanosheets can be well preserved after the stability test, as evidenced by SEM/TEM (Fig. S14†), XRD (Fig. S15†) and XPS (Fig. S16†) measurements. The outstanding stability of Mo-SnS₂/CC is believed to originate from the strong atomic and electronic bonding of Mo-dopants, 2D confinement effect of the nanosheet structure,²⁹ and direct nanosheet growth on CC with tight catalyst attachment.³¹ Therefore, Mo-SnS₂/CC has great potential as a promising catalyst for electroreduction of N₂ to NH₃ with a favorable NH₃ production rate and robust stability.

DFT calculations based on the energetically stable Mo-SnS₂-V_s structure (Fig. 2e) were further performed to gain deep insights into the synergistic role of the Mo-dopant and V_s in facilitating the NRR. In view of N₂ adsorption as the critical step to initialize the NRR, N₂ adsorption behaviors over pristine SnS₂ and Mo-SnS₂-V_s were first analyzed. For pristine SnS₂ (Fig. 4a

and b), the N₂ molecule barely adsorbs on either the central Sn site (Fig. 4a) or the edge Sn site (Fig. 4b), as evidenced by the negligible charge transfer and much less N≡N bond elongation (1.105 Å for the original N₂ gas), suggesting that pristine SnS₂ is almost inactive for the NRR. In sharp contrast, after N₂ adsorption on the V_s site of Mo-SnS₂-V_s (Fig. 4c), the N≡N bond is considerably elongated to 1.211 Å and 0.55|e| is injected into *N₂, implying the greatly enhanced N₂ adsorption on Mo-SnS₂-V_s. The enlarged view (Fig. 4d) reveals a unique N₂ adsorption mode on the V_s site of Mo-SnS₂-V_s, that is, the N₂ molecule is fixed to the Mo-Sn-Sn trimer center *via* a side-on configuration. Meanwhile, the N≡N triple bond is even cleaved to the N=N double bond without hydrogenation, indicating that the Mo-Sn-Sn trimer can strongly activate N₂ through an analogous dissociative pattern. As shown in the PDOS of the Mo-Sn-Sn trimer after N₂ adsorption (Fig. 4e), the Mo4d and Sn5p orbitals are both considerably hybridized with the N2p orbitals both below and above the Fermi level, suggesting the efficient back-

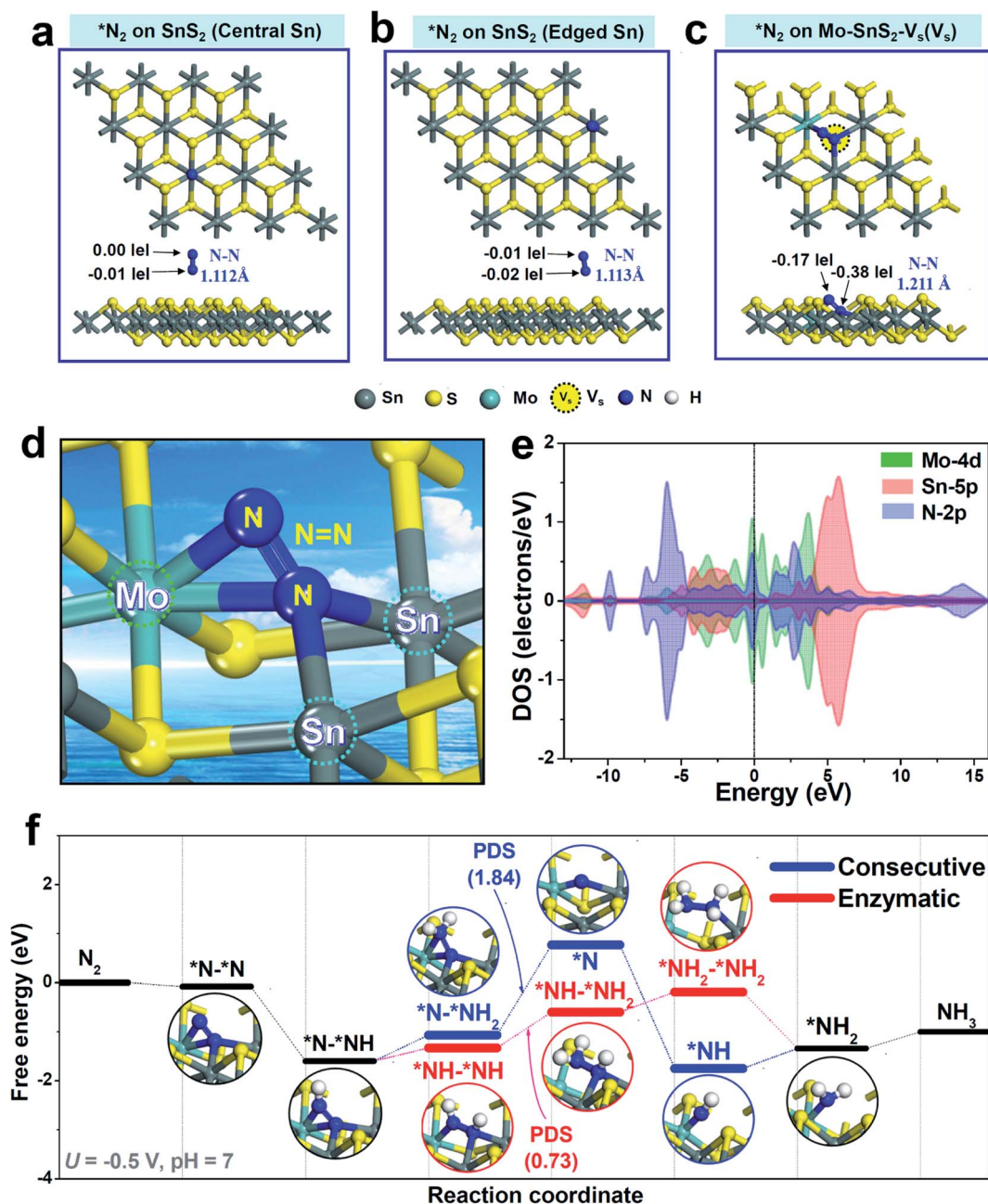


Fig. 4 (a–c) Optimized structures of N_2 adsorption on (a) SnS₂ (central Sn site), (b) SnS₂ (edge Sn site) and (c) Mo-SnS₂-V_s (V_s site). (d) Enlarged view of the N_2 adsorption mode on a V_s site of Mo-SnS₂-V_s (Mo-Sn-Sn trimer). (e) PDOS of the Mo-Sn-Sn trimer after N_2 adsorption. (f) Free energy diagrams of NRR consecutive/enzymatic reaction pathways over the Mo-Sn-Sn trimer at $U = -0.5$ V, pH = 7.

donation of electrons from the Mo-Sn-Sn trimer to $*N_2$.^{60,61} The prominent back-donation of electrons from the Mo-Sn-Sn trimer to $*N_2$ can be directly visualized from the differential charge density (Fig. S17[†]), showing the pronouncedly accumulated electrons on $*N_2$. Mulliken charge analysis (Fig. S18[†]) reveals that the Mo-Sn-Sn trimer back-donates a total of $0.55|e|$ to $*N_2$, with $0.37|e|$ from the Mo dopant and $0.09|e|$ from each of the two Sn atoms. Hence, the co-presence of the Mo dopant and V_s in Mo-SnS₂-V_s enables the creation of Mo-Sn-Sn trimer catalytic sites that can strongly activate the N_2 molecule even for the cleavage of the $N\equiv N$ triple bond to the $N=N$ double bond at the N_2 adsorption stage.

The hydrogenation pathway of Mo-SnS₂-V_s was then evaluated using Gibbs free energy profiles through the consecutive (analogous to the distal pathway but *via* side-on configuration⁶²) and enzymatic associative mechanisms, as shown in Fig. 4f. Optimized structures of all the NRR intermediates are presented in Fig. S19[†] (consecutive) and Fig. S20[†] (enzymatic). Notably, with the strong N_2 activation by the Mo-Sn-Sn trimer, the free energy of $*N-NH$ even becomes negative, making the first hydrogenation step ($*N_2 \rightarrow *N-NH$) occur spontaneously. This is in stark contrast to the largely positive free energy of $*N-NH$ (2.19 eV) observed in pristine SnS₂,²² which has the first hydrogenation step ($*N_2 \rightarrow *N-NH$) as the potential

determining step (PDS). After the first hydrogenation, $*N-*NH$ can be further hydrogenated through a consecutive or enzymatic pathway. For the consecutive pathway, $*N-*NH_2 \rightarrow *N$ is the PDS with a high energy barrier of 1.84 eV, while the PDS energy barrier ($*NH-*NH \rightarrow *NH-*NH_2$) is substantially reduced to 0.73 eV for the enzymatic pathway. Hence, the hydrogenation of Mo-SnS₂-V_s prefers to proceed *via* the enzymatic pathway with an overpotential of 0.57 V,⁶³ which is theoretically lower than that of most reported NRR catalysts.^{64–67} These results demonstrate that the highly active Mo-Sn-Sn trimer provides Mo-SnS₂-V_s with the downhill process of the first hydrogenation step, leading to the significantly decreased energy barrier and largely enhanced NRR activity.

On the other hand, as reported in the literature,^{68,69} the surface charge and hydrogen bonding can affect the electrocatalytic calculations. As shown in Fig. S21,† after considering the effects of surface charge (adding one charge e⁻ in the catalyst system) and hydrogen bonding (adding two H₂O molecules on the catalyst surface), we find that these two factors can make the free energies of $*N_2$ and $*N-*NH$ become more negative but the downhill trend of the $*N_2 \rightarrow *N-*NH$ process still remains. Therefore, our main conclusion is not affected by the effects of surface charge and hydrogen bonding.

In conclusion, through combined experimental and theoretical investigations, we demonstrated that the synergistic modulation of vacancy engineering and heteroatom doping has been successfully achieved in Mo-SnS₂ nanosheets with enriched V_s, which showed greatly enhanced NRR performance with an NH₃ yield of 41.3 μg h⁻¹ mg⁻¹ (-0.5 V) and an FE of 20.8% (-0.4 V). DFT calculations revealed the unique NRR mechanism of Mo-SnS₂-V_s, in which the created Mo-Sn-Sn trimer active sites could strongly activate N₂ for even the cleavage of the N≡N triple bond to the N=N double bond at the N₂ adsorption stage, consequently resulting in a downhill process of the first hydrogenation step and a largely reduced energy barrier. This work not only offers an efficient strategy towards the design of SnS₂-based catalysts for highly efficient electrosynthesis of NH₃, but also provides new insights into the synergistic role of vacancies and dopants in regulating the NRR activity.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

This work is supported by the National Natural Science Foundation of China (51761024), “Feitian Scholar” Program of Gansu Province, CAS “Light of West China” Program, and Foundation of A Hundred Youth Talents Training Program of Lanzhou Jiaotong University.

References

- 1 J. A. Brandes, N. Z. Boctor, G. D. Cody, B. A. Cooper, R. M. Hazen and H. S. Yoder Jr, *Nature*, 1998, **395**, 365.

- 2 X. Zhu, S. Mou, Q. Peng, Q. Liu, Y. Luo, G. Chen, S. Gao and X. Sun, *J. Mater. Chem. A*, 2020, **8**, 1545–1556.
- 3 G. F. Chen, S. Y. Ren, L. L. Zhang, H. Cheng, Y. R. Luo, K. H. Zhu, L. X. Ding and H. H. Wang, *Small Methods*, 2019, **3**, 1800337.
- 4 S. J. Li, D. Bao, M. M. Shi, B. R. Wulan, J. M. Yan and Q. Jiang, *Adv. Mater.*, 2017, **29**, 1700001.
- 5 H. Xie, Q. Geng, X. Zhu, Y. Luo, L. Chang, X. Niu, X. Shi, A. M. Asiri, S. Gao, Z. Wang and X. Sun, *J. Mater. Chem. A*, 2019, **7**, 24760–24764.
- 6 R. Zhao, C. Liu, X. Zhang, X. Zhu, P. Wei, L. Ji, Y. Guo, S. Gao, Y. Luo, Z. Wang and X. Sun, *J. Mater. Chem. A*, 2020, **8**, 77–81.
- 7 G. Deng, T. Wang, A. A. Alshehri, K. A. Alzahrani, Y. Wang, H. Ye, Y. Luo and X. Sun, *J. Mater. Chem. A*, 2019, **7**, 21674–21677.
- 8 H. Xian, Q. Wang, G. Yu, H. Wang, Y. Li, Y. Wang and T. Li, *Appl. Catal., A*, 2019, **581**, 116–122.
- 9 G. Yu, H. Guo, S. Liu, L. Chen, A. A. Alshehri, K. A. Alzahrani, F. Hao and T. Li, *ACS Appl. Mater. Interfaces*, 2019, **11**, 35764–35769.
- 10 H. Xian, H. Guo, Z. Chen, G. Yu, A. A. Alshehri, K. A. Alzahrani, F. Hao, R. Song and T. Li, *ACS Appl. Mater. Interfaces*, 2020, **12**, 2445–2451.
- 11 Y. R. Luo, G. F. Chen, L. Ding, X. Z. Chen, L. X. Ding and H. H. Wang, *Joule*, 2019, **3**, 279–289.
- 12 H. Cheng, L. X. Ding, G. F. Chen, L. L. Zhang, J. Xue and H. H. Wang, *Adv. Mater.*, 2018, **30**, 1803694.
- 13 K. Chu, Q. Q. Li, Y. H. Cheng and Y. P. Liu, *ACS Appl. Mater. Interfaces*, 2020, **12**, 11789–11796.
- 14 H. Cheng, P. X. Cui, F. R. Wang, L. X. Ding and H. H. Wang, *Angew. Chem., Int. Ed.*, 2019, **58**, 15541–15547.
- 15 X. Zhu, T. Wu, L. Ji, C. Li, T. Wang, S. Wen, S. Gao, X. Shi, Y. Luo, Q. Peng and X. Sun, *J. Mater. Chem. A*, 2019, **7**, 16117–16121.
- 16 L. Xia, X. Wu, Y. Wang, Z. Niu, Q. Liu, T. Li, X. Shi, A. M. Asiri and X. Sun, *Small Methods*, 2018, **3**, 1800251.
- 17 K. Chu, Q. Li, Y. Liu, J. Wang and Y. Cheng, *Appl. Catal., B*, 2020, **267**, 118693.
- 18 L. L. Zhang, L. X. Ding, G. F. Chen, X. F. Yang and H. H. Wang, *Angew. Chem., Int. Ed.*, 2019, **131**, 2638–2642.
- 19 L. Xia, J. Yang, H. Wang, R. Zhao, H. Chen, W. Fang, A. M. Asiri, F. Xie, G. Cui and X. Sun, *Chem. Commun.*, 2019, **55**, 3371–3374.
- 20 F. Li, L. Chen, G. P. Knowles, D. R. MacFarlane and J. Zhang, *Angew. Chem., Int. Ed.*, 2017, **56**, 505–509.
- 21 P. Li, W. Fu, P. Zhuang, Y. Cao, C. Tang, A. B. Watson, P. Dong, J. Shen and M. Ye, *Small*, 2019, **15**, 1902535.
- 22 X. Chen, Y.-T. Liu, C. Ma, J. Yu and B. Ding, *J. Mater. Chem. A*, 2019, **7**, 22235–22241.
- 23 Y. B. Li, Y. P. Liu, J. Wang, Y. L. Guo and K. Chu, *Inorg. Chem. Front.*, 2020, **7**, 455–463.
- 24 L. Yang, T. Wu, R. Zhang, H. Zhou, L. Xia, X. Shi, H. Zheng, Y. Zhang and X. Sun, *Nanoscale*, 2019, **11**, 1555–1562.
- 25 Z. Sun, R. Huo, C. Choi, S. Hong, T.-S. Wu, J. Qiu, C. Yan, Z. Han, Y. Liu, Y.-L. Soo and Y. Jung, *Nano Energy*, 2019, **62**, 869–875.

- 26 L. Zhang, X.-Y. Xie, H. Wang, L. Ji, Y. Zhang, H. Chen, T. Li, Y. Luo, G. Cui and X. Sun, *Chem. Commun.*, 2019, **55**, 4627–4630.
- 27 C. Lv, Y. Qian, C. Yan, Y. Ding, Y. Liu, G. Chen and G. Yu, *Angew. Chem., Int. Ed.*, 2018, **57**, 10246–10250.
- 28 X. Yang, J. Nash, J. Anibal, M. Dunwell, S. Kattel, E. Stavitski, K. Attenkofer, J. G. Chen, Y. Yan and B. Xu, *J. Am. Chem. Soc.*, 2018, **140**, 13387–13391.
- 29 H. Jin, L. Li, X. Liu, C. Tang, W. Xu, S. Chen, L. Song, Y. Zheng and S.-Z. Qiao, *Adv. Mater.*, 2019, **31**, 1902709.
- 30 X. H. Wang, J. Wang, Y. B. Li and K. Chu, *ChemCatChem*, 2019, **11**, 4529–4536.
- 31 Y. P. Liu, Y. B. Li, H. Zhang and K. Chu, *Inorg. Chem.*, 2019, **58**, 10424–10431.
- 32 K. Chu, Y. Liu, Y. Chen and Q. Li, *J. Mater. Chem. A*, 2020, **8**, 5200–5208.
- 33 K. Chu, Y. H. Chen, Q. Q. Li, Y. P. Liu and Y. Tian, *J. Mater. Chem. A*, 2020, **8**, 5865–5873.
- 34 T. Wu, W. Kong, Y. Zhang, Z. Xing, J. Zhao, T. Wang, X. Shi, Y. Luo and X. Sun, *Small Methods*, 2019, 1900356.
- 35 C. Guo, X. Liu, L. Gao, X. Kuang, X. Ren, X. Ma, M. Zhao, H. Yang, X. Sun and Q. Wei, *Appl. Catal., B*, 2020, **263**, 118296.
- 36 K. Chu, Y. P. Liu, Y. B. Li, Y. L. Guo, Y. Tian and H. Zhang, *Appl. Catal., B*, 2020, **264**, 118525.
- 37 T. Wu, Z. Xing, S. Mou, C. Li, Y. Qiao, Q. Liu, X. Zhu, Y. Luo, X. Shi, Y. Zhang and X. Sun, *Angew. Chem., Int. Ed.*, 2019, **58**, 18449–18453.
- 38 N. Cao, Z. Chen, K. Zang, J. Xu, J. Zhong, J. Luo, X. Xu and G. Zheng, *Nat. Commun.*, 2019, **10**, 1–12.
- 39 K. Chu, X. H. Wang, Y. B. Li, D. J. Huang, Z. R. Geng, X. L. Zhao, H. Liu and H. Zhang, *Mater. Des.*, 2018, **140**, 85–94.
- 40 K. Chu, J. Wang, Y. P. Liu and Z. R. Geng, *Carbon*, 2018, **140**, 112–123.
- 41 K. Chu, X. H. Wang, F. Wang, Y. B. Li, D. J. Huang, H. Liu, W. L. Ma, F. X. Liu and H. Zhang, *Carbon*, 2018, **127**, 102–112.
- 42 K. Chu, J. Wang, Y. P. Liu, Y. B. Li, C. C. Jia and H. Zhang, *Carbon*, 2019, **143**, 85–96.
- 43 K. Chu, F. Wang, X. H. Wang, Y. B. Li, Z. R. Geng, D. J. Huang and H. Zhang, *Mater. Des.*, 2018, **144**, 290–303.
- 44 K. Chu, F. Wang, Y. B. Li, X. H. Wang, D. J. Huang and H. Zhang, *Carbon*, 2018, **133**, 127–139.
- 45 L. Meng, S. Wang, F. Cao, W. Tian, R. Long and L. Li, *Angew. Chem., Int. Ed.*, 2019, **58**, 6761–6765.
- 46 M.-A. Légaré, G. Bélanger-Chabot, R. D. Dewhurst, E. Welz, I. Krummenacher, B. Engels and H. Braunschweig, *Science*, 2018, **359**, 896–900.
- 47 J. Wang, Y. P. Liu, H. Zhang, D. J. Huang and K. Chu, *Catal. Sci. Technol.*, 2019, **9**, 4248–4254.
- 48 Y. P. Liu, Y. B. Li, D. J. Huang, H. Zhang and K. Chu, *Chem.–Eur. J.*, 2019, **25**, 11933–11939.
- 49 K. Chu, Y. P. Liu, Y. B. Li, J. Wang and H. Zhang, *ACS Appl. Mater. Interfaces*, 2019, **11**, 31806–31815.
- 50 L. Li, C. Tang, B. Xia, H. Jin, Y. Zheng and S.-Z. Qiao, *ACS Catal.*, 2019, **9**, 2902–2908.
- 51 D. Zhu, L. Zhang, R. E. Ruther and R. J. Hamers, *Nat. Mater.*, 2013, **12**, 836.
- 52 G. W. Watt and J. D. Chrisp, *Anal. Chem.*, 1952, **24**, 2006–2008.
- 53 K. Chu, Y. P. Liu, Y. B. Li, Y. L. Guo and Y. Tian, *ACS Appl. Mater. Interfaces*, 2020, **12**, 7081–7090.
- 54 J. Yu, C. Li, B. Li, X. Zhu, R. Zhang, L. Ji, D. Tang, A. M. Asiri, X. Sun, Q. Li, S. Liu and Y. Luo, *Chem. Commun.*, 2019, **55**, 6401–6404.
- 55 R. Zhang, L. Ji, W. Kong, H. Wang, R. Zhao, H. Chen, T. Li, B. Li, Y. Luo and X. Sun, *Chem. Commun.*, 2019, **55**, 5263–5266.
- 56 C. Li, J. Yu, L. Yang, J. Zhao, W. Kong, T. Wang, A. M. Asiri, Q. Li and X. Sun, *Inorg. Chem.*, 2019, **58**, 9597–9601.
- 57 A. R. Singh, B. A. Rohr, J. A. Schwalbe, M. Cargnello, K. Chan, T. F. Jaramillo, I. Chorkendorff and J. K. Nørskov, *ACS Catal.*, 2016, **7**, 706–709.
- 58 P. Wang, Q. Q. Li, Y. H. Cheng and K. Chu, *J. Mater. Sci.*, 2020, **55**, 4624–4632.
- 59 F. Wang, Y. P. Liu, H. Zhang and K. Chu, *ChemCatChem*, 2019, **11**, 1441–1447.
- 60 Q. Li, L. He, C. Sun and X. Zhang, *J. Phys. Chem. C*, 2017, **121**, 27563–27568.
- 61 Y.-C. Hao, Y. Guo, L.-W. Chen, M. Shu, X.-Y. Wang, T.-A. Bu, W.-Y. Gao, N. Zhang, X. Su, X. Feng, J.-W. Zhou, B. Wang, C.-W. Hu, A.-X. Yin, R. Si, Y.-W. Zhang and C.-H. Yan, *Nat. Catal.*, 2019, **2**, 448–456.
- 62 L. Shi, Q. Li, C. Ling, Y. Zhang, Y. Ouyang, X. Bai and J. Wang, *J. Mater. Chem. A*, 2019, **7**, 4865–4871.
- 63 J. Zhao and Z. Chen, *J. Am. Chem. Soc.*, 2017, **139**, 12480–12487.
- 64 K. Chu, Y. Liu, J. Wang and H. Zhang, *ACS Appl. Energy Mater.*, 2019, **2**, 2288–2295.
- 65 K. Chu, Y. Liu, Y. Li, H. Zhang and Y. Tian, *J. Mater. Chem. A*, 2019, **7**, 4389–4394.
- 66 Z. Wang, F. Gong, L. Zhang, R. Wang, L. Ji, Q. Liu, Y. Luo, H. Guo, Y. Li, P. Gao, X. Shi, B. Li, B. Tang and X. Sun, *Adv. Sci.*, 2018, 1801182.
- 67 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.
- 68 X. Zhao and Y. Liu, *J. Am. Chem. Soc.*, 2020, **142**, 5773–5777.
- 69 D. Kim, J. Shi and Y. Liu, *J. Am. Chem. Soc.*, 2018, **140**, 9127–9131.