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Atomic modulation and phase engineering of MoS₂ for boosting N₂ reduction

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Abstract

Electrochemical nitrogen reduction reaction (ENRR) has emerged as a potential alternative to the conventional Haber-Bosch process for ammonia production. However, ENRR technology is still restricted by the limited Faradaic efficiency due to the hard-to-break N-N triple bond. Herein, inspired by the biomimetic catalyst, we developed a Fe-modulated MoS₂ catalyst (named Fe@MoS₂) as an efficient ENRR catalyst. Raman spectra, coupled with the X-ray absorption spectroscopy, demonstrate the introduction of Fe into the MoS₂ lattice and achieve partial 2H to 1T phase conversion. The presence of S-vacancies on MoS₂ substrates was observed on scanning transmission electron microscopy images. Operando infrared absorption spectroscopy confirms that the constructed catalytic site significantly reduces barriers to nitrogen activation. The synthesized Fe@MoS₂, with its superior geometric and electronic structures, exhibits a remarkable Faradaic efficiency of 19.7 ± 5.5% at -0.2 V vs. Reversible Hydrogen Electrode and a high yield rate of 20.2 ± 5.3 μg h⁻¹ mg⁻¹ at -0.8 V vs. Reversible Hydrogen Electrode. Therefore, this work provides a fresh direction for designing novel catalysts, eventually boosting the nitrogen reduction reaction



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kinetics and accelerating the ENRR application.

Keywords: Nitrogen reduction, heteroatom doping, molybdenum disulfide, phase engineering

INTRODUCTION

Ammonia (NH_3) ranks among the most demanded chemical products in the world, playing an essential role in producing fertilizers, plastics, and medicines^[1,2]. Furthermore, it is also an important carbon-free energy carrier, benefiting from its high hydrogen content (17.6 wt%) and high gravimetric energy density (3 kWh kg^{-1})^[3,4]. The lower liquefaction difficulty makes it more convenient for storage and transportation^[5]. Nowadays, the artificial synthesis of NH_3 heavily depends on the Haber-Bosch (H-B) method. The N_2 and H_2 are transformed into NH_3 on the catalyst surface under severe reaction conditions ($400\text{--}600^\circ\text{C}$, $\sim 60 \text{ bar}$)^[6–8]. However, the H-B process is energy-intensive and has high carbon emissions and high cost of raw materials (H_2)^[9,10]. Therefore, it is necessary to search for a green NH_3 synthesis technology. Recently, an electrochemical nitrogen reduction reaction (ENRR) technology has been developed and can realize the N_2 to NH_3 conversion at ambient temperature, with H_2O as the hydrogen source^[11,12]. The total cost and carbon footprint of NH_3 synthesis will be effectively reduced by driving ENRR using renewable energy sources^[13]. Moreover, the simple process equipment makes flexible and decentralized NH_3 preparation possible. Nevertheless, the ENRR process is still restricted by low yield rates and Faradaic efficiency (FE). Thus, the pursuit of highly efficient electrocatalysts for ENRR is greatly desirable but challenging.

Mo-based nitrogenase enables N_2 fixation under ambient conditions^[14,15], which inspired researchers to investigate biomimetic catalysts with similar elemental compositions. MoS_2 , as a layered two-dimensional (2D) semiconductor material, possesses a large surface area and crystal phase^[16–19]. Multi-variable interlayer stacking forms and tunable electronic structure make it a promising candidate in the field of electrocatalytic synthesis of NH_3 ^[20–22]. Due to the difficulty of activating the non-polar N-N triple bond and the high adsorption barrier of N_2 , the NH_3 yield rate and energy transfer efficiency in ENRR are limited. Moreover, the hydrogen evolution reaction (HER) has a close reaction equilibrium potential to that of ENRR, affecting the selectivity of ENRR in the aqueous phase and leading to unsatisfactory FE^[23–25]. Therefore, several strategies have been reported to optimize the MoS_2 -based catalysts for improving ENRR performance, including vacancy engineering^[26,27], heteroatom doping^[28,29], phase engineering^[30,31], *etc.*

The catalytic activity and stability of MoS_2 with different phases vary greatly in ENRR. Lin *et al.*^[30] compared the ENRR performance of the metastable phase (1T'- and 1T'''- MoS_2) and the stable phase (2H- MoS_2). Thanks to the high localized electron density around Mo-Mo, the NH_3 yield and FE of 1T'''- MoS_2 are approximately nine and 12 times higher than those of 2H- MoS_2 , respectively. Besides, constructing defects on the MoS_2 basal plane can expose more active sites for N_2 adsorption. You *et al.*^[32] successfully introduced S vacancies in natural molybdenite through a one-step annealing method. The electron structure at the Mo edge was altered, which optimized the free energy barrier of the rate-limiting step. The prepared rich S-vacancy MoS_2 exhibited significantly improved ENRR performance compared to the raw material. In addition, benefiting from the synergistic effect among different elements, superior ENRR activity can be achieved by doping heteroatoms in MoS_2 . Zhao *et al.* synthesized Fe-doped MoS_2 nanosheets (Fe- MoS_2 /CC) using a chemical reduction method^[33]. The introduction of Fe induced the redistribution of charges on the MoS_2 basal plane and generated new active sites, significantly improving the NH_3 synthesis performance. The NH_3 yield was $12.5 \mu\text{g h}^{-1} \text{ cm}^{-2}$ with FE of 10.8% at -0.1 V vs. Reversible Hydrogen Electrode (RHE). Moreover, the average size and chemical valence state of Fe nanodots did not change significantly after long-term electrolysis, which demonstrated the excellent stability of Fe- MoS_2 /CC.

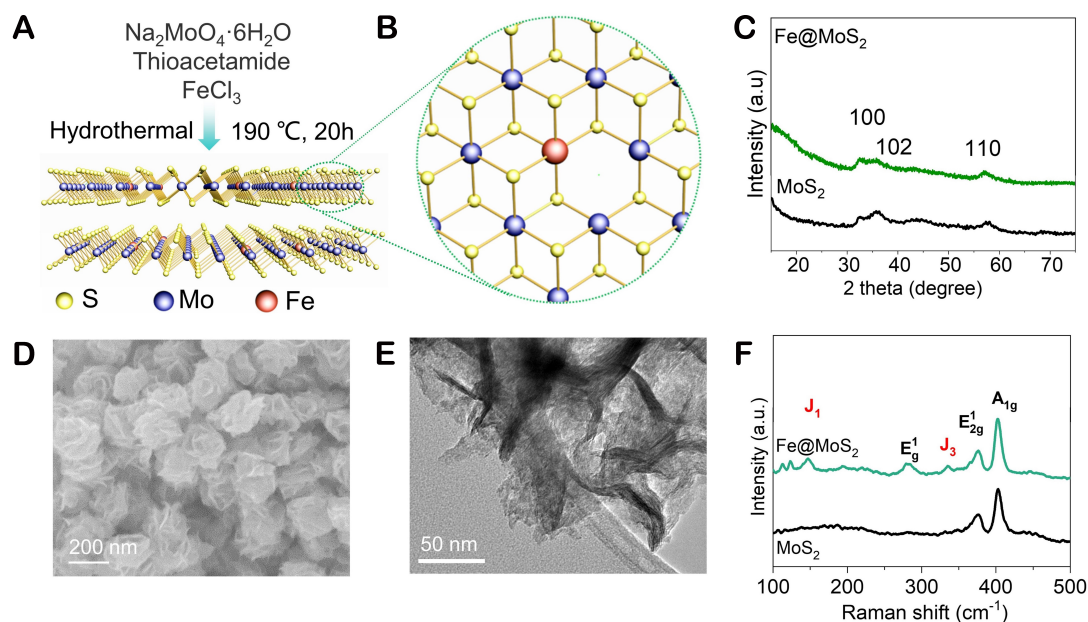


Figure 1. Schematic illustration of (A) synthesis process and (B) atomic structure; (C) XRD patterns of MoS₂ and Fe@MoS₂. (D) SEM image of Fe@MoS₂, (E) TEM image of Fe@MoS₂, (F) Raman spectra of MoS₂ and Fe@MoS₂.

In this work, we developed a synergetic optimization strategy to enhance the ENRR activity of MoS₂, coupling vacancy engineering, heteroatom doping, and phase engineering. Biomimetic Fe@MoS₂ electrocatalysts were prepared using a one-pot hydrothermal method. The partial MoS₂ converts from 2H to 1T phase after Fe atom doping. Furthermore, the unique hydrangea-like morphology and defects on the MoS₂ basal plane promote the accessibility of the active site. *In-situ* and electrochemical characterizations demonstrated that Fe atoms had been inserted into the lattice of MoS₂, and its unique coordination structure was analyzed which accelerates the reaction kinetics of N₂ to NH₃, thus achieving a promising ENRR performance. The maximum FE of 19.7% ± 5.5% and the highest NH₃ yield of 20.2 ± 5.3 μg h⁻¹ mg⁻¹ were achieved by Fe@MoS₂ at -0.2 V vs. RHE and -0.8 V vs. RHE, respectively.

MATERIALS AND METHODS

Synthesis of materials

One-step hydrothermal synthesis was used to prepare MoS₂ [Figure 1A]. Briefly, the mixed solution after sonication (30 mL deionized water (D.I water)) with 0.4 mmol Na₂MoO₄·6H₂O and 0.82 mmol thioacetamide) was transferred into a stainless steel autoclave (50 mL). The heating parameter was set to 190 °C for 20 h. After cooling, filtration, washing and drying at room temperature, MoS₂ was collected. The synthesis process of Fe@MoS₂ was the same as that of MoS₂, except for adding 0.004 mmol FeCl₃.

Electrochemical measurement

ENRR electrochemical performance was evaluated using a three-electrode system (0.25 M LiClO₄) under an ambient environment (25 °C, 1 atm). The working, reference, and counter electrodes are ENRR catalyst, Ag/AgCl (saturated KCl solution) electrode, and Pt wire, respectively. Firstly, 2 mg of catalyst was dispersed into 1 mL Nafion/ISO/D.I water solution (with the volume ratio of 1:9:40) with the sonication for 30 min. Then, 100 μL catalyst inks (MoS₂ and Fe@MoS₂) were drop-casted on carbon paper (1 cm²) with 0.2 mg cm⁻² and dried in the ambient environment. Before the ENRR test, the electrolyte was bubbled with N₂ gas (purity, 99.99%) for 20 min. During the ENRR test, the N₂ flow was continuously inputted with a flow rate of 20 sccm.

Determination of NH₃

The quantification of NH₃ was determined using the salicylic acid-based colorimetric method^[34]. To obtain the calibration curve, standard NH₄Cl solutions with different concentrations of 0.0, 0.1, 0.2, 0.5, 1.0, and 2.0 g mL⁻¹ in the electrolyte were first prepared. Then, 50 µL catalytic solution (10 mL D.I water with 0.1 g sodium nitroprusside dihydrate), 500 µL coloring solution (0.32 M NaOH and 0.4 M sodium salicylate), and 50 µL oxidation solution (NaClO solution (pCl = 4 - 4.9) with 0.75 M NaOH) were mixed with the 4 mL standard NH₄Cl solution, and standing for 1h for color development. Next, the ultraviolet-visible (UV-Vis) spectrophotometer was used to measure the absorbance (λ = 655 nm) with different concentrations of NH₄Cl. Finally, we got the calibration curve with a good linear relationship [Supplementary Figure 1]. The NH₃ concentration in the reaction electrolyte is calculated based on the standard curve.

The NH₃ formation rate can be calculated by:

$$R_{NH_3}(\mu g h^{-1} mg^{-1}) = \frac{x(ppm) \times 10^3 (\mu g/mg) \times V(L)}{t(h) \times m(mg)}$$

where:

x (ppm): calculated concentration of ammonia.

V (L): volume of reacted electrolyte.

t (h): reaction time in hours.

m (mg): mass of catalyst on the carbon paper.

R_{NH_3} $\mu g h^{-1} mg^{-1}$

The FE is calculated using:

$$FE_{NH_3}(\%) = \frac{3 \times R_{NH_3}(\mu g h^{-1} mg^{-1}) \times t(h) \times 10^{-6} (g/\mu g) \times m(mg) \times F}{Mr_{NH_4}(g/mol) \times I(A) \times t(s)} \times 100\%$$

where:

x (ppm): calculated concentration of ammonia.

Mr (NH₄⁺): 18 (g/mol).

V (L): the volume of reacted electrolyte.

I (A): the average current during the reaction.

F : the Faraday constant $96,485 \text{ mol}^{-1}$.

Operando attenuated total reflection surface enhanced infrared absorption spectroscopy

Operando attenuated total reflection surface enhanced infrared absorption spectroscopy (ATR-SEIRAS) utilizes the same three-electrode system as the electrochemical test, and the electrolyte is $0.1 \text{ M Li}_2\text{SO}_4$ (N_2 saturated). Before each measurement, the background spectrum of the catalyst electrode needs to be collected at the open circuit potential. Afterward, the spectra at different potentials were collected. All collected spectra were given by the absorbance ($-\log(R/R_0)$) with a spectral resolution of 4 cm^{-1} .

RESULTS AND DISCUSSION

Material characterization

The preparation process of Fe@MoS_2 and the atomic structure are illustrated in [Figure 1A](#) and [1B](#). X-ray diffraction (XRD) patterns of MoS_2 and Fe@MoS_2 are shown in [Figure 1C](#). The diffraction peaks at 32.5° , 35.8° , and 57.2° could be indexed to (100), (102), and (110) planes of the standard MoS_2 phase (PDF #37-1492)^[35-37], respectively. The diffraction peaks of Fe@MoS_2 do not change significantly compared to MoS_2 , and no FeS or FeS_2 phases were observed, which indicates that traces of Fe element did not form new sulfide. A scanning electron microscopy (SEM) image [[Figure 1D](#)] shows that the synthesized Fe@MoS_2 is assembled from multiple 2D nanosheets and exhibits a hydrangea-like morphology. Moreover, Brunauer-Emmett-Teller (BET) test was carried out to characterize the specific surface area of catalysts. [Supplementary Figure 2](#) shows the N_2 adsorption-desorption isotherm curve of MoS_2 and Fe@MoS_2 . Fe@MoS_2 has a higher BET specific surface area of $54.034 \text{ m}^2 \text{ g}^{-1}$ and a higher pore volume of $0.32 \text{ cm}^3 \text{ g}^{-1}$ than MoS_2 ($22.63 \text{ m}^2 \text{ g}^{-1}$ and $0.09 \text{ cm}^3 \text{ g}^{-1}$, respectively). The catalytic activity was improved due to more exposed active sites [[Supplementary Table 1](#)]. The high specific surface area exposes more active sites, which benefits its catalytic activity. Electrochemically active surface area (ECSA) can be used to evaluate the number of active sites in a catalyst, which is proportional to the double-layer capacitance (C_{dl}). [Supplementary Figure 3](#) presents the C_{dl} values obtained after linear fitting. Notably increased C_{dl} values were observed in Fe@MoS_2 (17.85 mF cm^{-2}), compared to MoS_2 (0.35 mF cm^{-2}), indicating the beneficial effect of Fe doping on the number of active sites.

No recognizable nanoparticles or clusters are observed in the transmission electron microscopy (TEM) image [[Figure 1E](#)]. This suggests that Fe was more likely to be inserted into the MoS_2 plane rather than deposited on the surface, which requires further analysis in combination with spectra. Raman spectra [[Figure 1F](#)] were employed to further investigate the effect of Fe insertion on MoS_2 . The results suggest that synthesized MoS_2 exhibits distinct 2H phase signals, with two peak positions at 376.5 and 403 cm^{-1} corresponding to the E_{2g}^1 vibrational peak and A_g^1 vibrational peak of 2H- MoS_2 , respectively^[35,38,39]. However, Fe@MoS_2 is a mixed 1T/2H- MoS_2 phase; two new 1T phase characteristic peaks of J_1 (146 cm^{-1}) and J_3 (334 cm^{-1}) are also clearly observed, indicating that the doping of Fe atoms triggers a phase transition from 2H- MoS_2 to 1T- MoS_2 ^[40-42].

A high-resolution TEM (HRTEM) image [[Figure 2A](#)] shows the same results as those obtained from TEM images, with no impurities or other Fe-based structures observed in Fe@MoS_2 . Further, with the help of dark field scanning TEM (STEM) characterization [[Figure 2B](#) and [Supplementary Figure 4](#)], it is seen that defects are present on the surface of the 2D structure. Atomic-level Fe (weak Z contrast intensity) replaces some Mo atoms and dopes in the panel of the 2D MoS_2 plane. To explore the distribution of different elements in the catalysts, corresponding energy dispersive spectroscopy (EDS) elemental mapping was

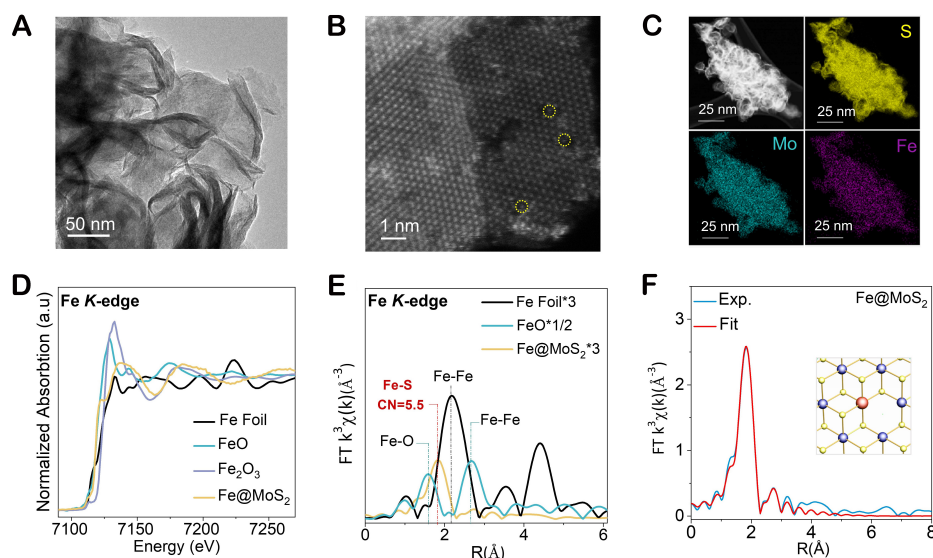


Figure 2. Material Characterization. (A) HRTEM image of Fe@MoS₂, (B) STEM image, and (C) the corresponding EDS elemental mappings of Fe@MoS₂, (D) XANES spectra of the Fe K-edge in Fe foil, FeO, Fe₂O₃, and Fe@MoS₂, (E) FT-EXAFS spectra of the Fe K-edge in Fe foil, FeO, and Fe@MoS₂, (F) XAFS fitting result of Fe@MoS₂ at R space. Inset: Atomic illustration of Fe@MoS₂ (Mo atoms: purple; S atoms: yellow; Fe atoms: red).

conducted to characterize Fe@MoS₂. **Figure 2C** demonstrates that Mo, S, and Fe are uniformly distributed in Fe@MoS₂.

X-ray absorption near edge structure (XANES) and Fourier transform-extended X-ray absorption fine structure (FT-EXAFS) spectra further revealed the coordination structure and coordination number of Fe atom in Fe@MoS₂. As shown in **Figure 2D**, the absorption edge of Fe@MoS₂ is close to the curve of FeO, indicating that Fe of Fe@MoS₂ is dominated by a chemical state of +2. In the FT-EXAFS spectra [**Figure 2E**], a peak can be observed at 1.87 Å for Fe@MoS₂, corresponding to the first coordination^[43,44]. No Fe-O and Fe-Fe are observed in Fe@MoS₂, indicating that the atomically dispersed Fe may dope within Mo-based vacancy and bond with S atoms. Fitting curves [**Figure 2F**, **Supplementary Figure 5**, **Supplementary Table 2**] of the Fe K-edge EXAFS spectra of Fe@MoS₂ indicate that the Fe-S coordination number is approximately 5.5. This suggests that Fe primarily replaces the Mo atom and coordinates with five S atoms near the S vacancy in MoS₂^[45]. X-ray Photoelectron Spectroscopy (XPS) reveals the valence changes of the Mo and S elements after Fe doping. **Figure 3A** and **B** shows the S 2*p* and Mo 3*d* spectra of MoS₂ and Fe@MoS₂. Two characteristic peaks of 162.35 and 160.8 eV for MoS₂ were assigned to S 2*p*_{1/2} and S 2*p*_{3/2}, respectively^[46,47]. The introduction of Fe resulted in a positive shift of the peak position with altered binding energies of 162.70 and 161.20 eV, which may be related to the Fe-S bond formation. In addition, the peaks with binding energies at 230.9 and 227.7 eV correspond to Mo 3*d*_{3/2} and Mo 3*d*_{5/2}^[48]. A positive shift of 1.1 eV was observed in Fe@MoS₂, indicating an elevated valence state for Mo as well, which is well in line with the previously reported research^[49,50].

Electrochemical measurements

The ENRR performance of the Fe@MoS₂ is evaluated in an H-type cell equipped with a three-electrode system [**Figure 4A**]. The linear scan voltammetry (LSV) was first performed to compare the response current density under Ar- or N₂-saturated electrolyte of Fe@MoS₂. As shown in **Figure 4B**, higher current density in N₂ indicates the possibility of the Fe@MoS₂ towards N₂ reduction. The subsequent ENRR

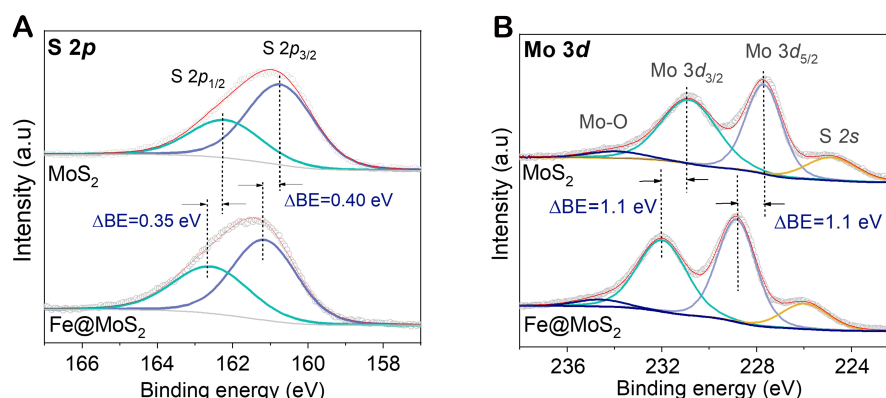


Figure 3. (A) S 2p and (B) Mo 3d XPS of MoS₂ and Fe@MoS₂.

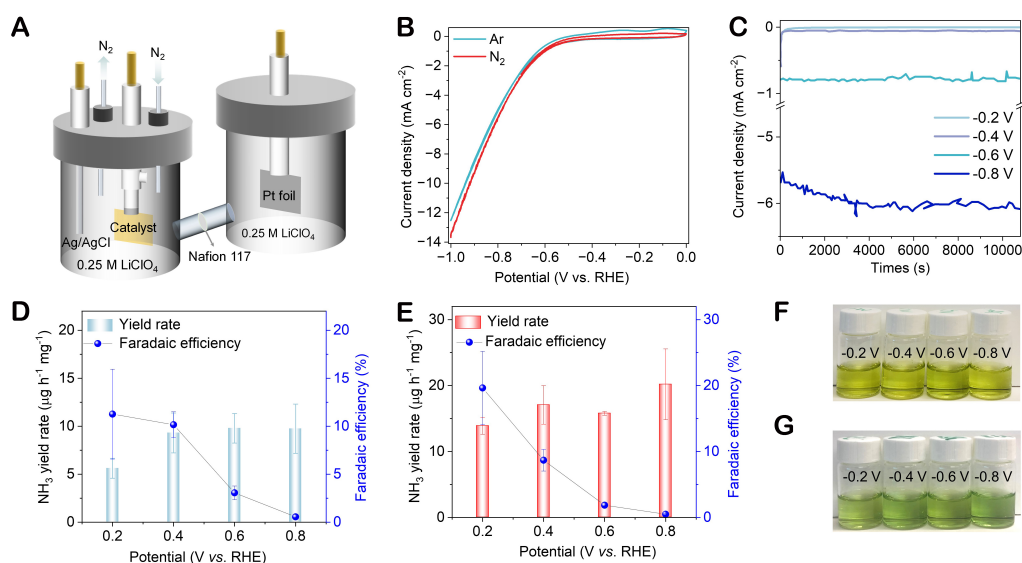


Figure 4. Electrochemical performance evaluation. (A) Schematic illustration of the reactor for electrochemical N₂ reduction. (B) LSV curves of Fe@MoS₂ in Ar and N₂-saturated 0.25 M LiClO₄ at a scan rate of 10 mV s⁻¹. (C) Chronoamperometric curves of Fe@MoS₂ at different applied potentials. NH₃ yield rate and FE of (D) MoS₂ and (E) Fe@MoS₂. The picture of the reacted solution of (F) MoS₂ and (G) Fe@MoS₂.

performance was evaluated using a chronoamperometry method in N₂-saturated electrolyte for 3 h, and the response current densities at different potentials were recorded [Figure 4]. The reacted electrolyte was collected and analyzed for the NH₃ concentration^[34]. The relevant equations have been mentioned above.

The yield rate and FE of NH₃ over the electrocatalyst Fe@MoS₂ are higher than those of the pristine MoS₂ electrocatalyst, which indicates the optimization of the introduction of Fe element in the pristine MoS₂ plane. As shown in Figure 4D, the MoS₂ can achieve the yield rate of NH₃ less than 10 μg h⁻¹ mg⁻¹ between the potential of -0.2 to -0.8 V vs. RHE, while the Fe@MoS₂ can achieve the yield rate of NH₃ larger than 13.9 μg h⁻¹ mg⁻¹. The highest yield rate of 20.2 ± 5.3 μg h⁻¹ mg⁻¹ is obtained at -0.8 V vs. RHE, about two times higher than that of MoS₂ [Figure 4E]. As shown in Figure 4F and G, the colors of the reacted electrolytes of Fe@MoS₂ and MoS₂ are green and yellow, respectively, suggesting that more NH₃ is generated with Fe@MoS₂ than that of MoS₂. The highest FE of ~19.7% ± 5.5% is obtained at -0.2 V vs. RHE. The FE

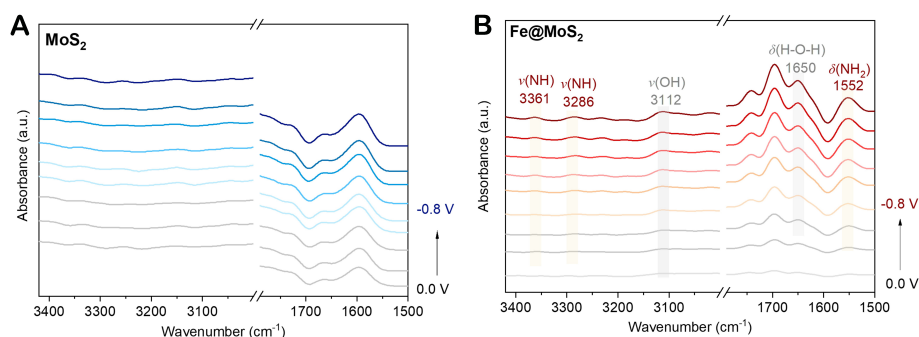


Figure 5. Mechanism study. Operando ATR-SEIRAS of (A) MoS₂ and (B) Fe@MoS₂.

dramatically decreases as the applied potentials increase, suggesting that the reaction is dominated by side-reaction HER. Compared with previously reported MoS₂-based catalysts, [Supplementary Table 3](#) shows that Fe@MoS₂ exhibits a relatively high FE and yield rate of NH₃. In addition, Fe@MoS₂ demonstrates reliable stability in long-term (10 h) stability tests. [Supplementary Figure 6](#) illustrates no significant deterioration of the current density of Fe@MoS₂.

Operando ATR-SEIRAS was performed to characterize the reaction intermediates and study the reaction mechanism. [Figure 5](#) recorded the ATR-SEIRAS spectra of MoS₂ and Fe@MoS₂ under different potentials during the ENRR process. As shown in [Figure 5A](#), no obvious peak can be observed for pristine MoS₂, indicating no intermediates generated. In contrast, several peaks gradually increase with the increasing potentials for Fe@MoS₂ [[Figure 5B](#)]. Some peaks attributed to N-H stretching are located at 3,361 and 3,286 cm⁻¹^[51,52]. The peaks at 3,112 and 1,650 cm⁻¹ correspond to the O-H stretching and bending mode of H₂O, respectively^[53,54]. The bending vibration absorption peak of -N-H at 1,552 cm⁻¹ gradually strengthened^[55], indicating a strong N₂ reduction reaction over the Fe@MoS₂. These results demonstrate that introducing Fe can accelerate the ENRR kinetics.

CONCLUSIONS

In conclusion, an inspired biomimetic electrocatalyst of Fe@MoS₂ is designed and successfully prepared using a one-step hydrothermal method. The introduction of atomic-level Fe achieves the geometric and electronic structure modulation of MoS₂ and triggers the phase transition of the 2H-phase MoS₂ into 1T-phase MoS₂. Thanks to the improved atomic structure and increased electrical conductivity of Fe@MoS₂, the kinetics are greatly accelerated, thus enhancing ENRR performance. Compared to the pristine 2H MoS₂, the prepared Fe@MoS₂ boosts the ENRR kinetics and exhibits superior ENRR performance with a yield rate of 20.2 ± 5.3 μg h⁻¹ mg⁻¹ at -0.8 V vs. RHE and a FE of ~19.7% ± 5.5% at -0.2 V vs. RHE. This work provides a new insight for designing efficient ENRR catalysts by synergistic doping and phase engineering.

DECLARATIONS

Authors' contributions

Synthesis and testing of materials, data collection, and original manuscript writing: Jia Y, Shao G, Li Y, Yang R

Validation and original manuscript revision: Huang M, Huang H, Liu M, Huang G, Lu Q, Gu C

Data collection: Jia Y, Shao G, Li Y, Yang R

Data analysis, writing, review, and editing: Jia Y, Li Y, Shao G

Review and discussion: Jia Y, Li Y, Shao G, Huang H, Huang G, Gu C

Supervision, funding acquisition: Li Y, Huang M

All authors have read and agreed to the published version of the manuscript.

Availability of data and materials

According to reasonable requirements, all of the data examined in this research can be obtained from the correspondents.

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Conflicts of interest

Liu M is affiliated with “State Grid Zhejiang Electric Power CO., LTD Research Institute, Hangzhou, China”. While the other authors have declared that they have no conflicts of interest.

Ethical approval and consent to participate

Not applicable.

Consent for publication

Not applicable.

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