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Three-dimensional Nitrogen-Doped Graphene Supported Molybdenum Disulfide Nanoparticles as an Advanced Catalyst for Hydrogen Evolution Reaction

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An efficient three-dimensional (3D) hybrid material of nitrogen-doped graphene sheets (N-RGO) supporting molybdenum disulfide (MoS₂) nanoparticles with high-performance electrocatalytic activity for hydrogen evolution reaction (HER) is fabricated by using a facile hydrothermal route. Comprehensive microscopic and spectroscopic characterizations confirm the resulting hybrid material possesses a 3D crumpled few-layered graphene network structure decorated with MoS₂ nanoparticles. Electrochemical characterization analysis reveals that the resulting hybrid material exhibits efficient electrocatalytic activity toward HER under acidic conditions with a low onset potential of 112 mV and a small Tafel slope of 44 mV per decade. The enhanced mechanism of electrocatalytic activity has been investigated in detail by controlling the elemental composition, electrical conductance and surface morphology of the 3D hybrid as well as Density Functional Theory (DFT) calculations. This demonstrates that the abundance of exposed active sulfur edge sites in the MoS₂ and nitrogen active functional moieties in N-RGO are synergistically responsible for the catalytic activity, whilst the distinguished and coherent interface in MoS₂/N-RGO facilitates the electron transfer during electrocatalysis. Our study gives insights into the physical/chemical mechanism of enhanced HER performance in MoS₂/N-RGO hybrids and illustrates how to design and construct a 3D hybrid to maximize the catalytic efficiency.

Hydrogen has emerged as an effective alternative to fossil fuels because it is environment-friendly energy with water as exhaust¹. Hydrogen formation from water has long been considered as a promising approach for solar energy storage². To improve the hydrogen transition efficiency, advanced catalysts are continuously explored to reduce the overpotential of the hydrogen evolution reaction (HER), a basic

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step involved in electrochemical water splitting ($2\text{H}^+ + 2\text{e}^- \rightarrow \text{H}_2$)^{2–4}. The widespread application of the most effective catalysts (drawn from the Pt group of metals) are limited by their low natural abundance and high cost⁵. Therefore, there is an urgent need to develop sustainable well-performing alternatives for HER electrocatalysis with high natural abundance and low cost^{6,7}. Various transition metals including Co, Fe, Ni and their derivatives, and metal-free catalysts like graphitic-carbon nitride coupled with nitrogen-doped graphene have been widely reported for HER^{8–13}. However, these non-precious-metal catalysts suffer from instability and low catalytic activity in acidic conditions^{13,14}. Empirical results and theoretical predications have demonstrated that molybdenum disulphide (MoS_2) and its derivatives possess high-performance electrocatalytic activity towards HER^{15,16}. It has been identified that the edge site of MoS_2 can act as a promising hydrogen evolution catalyst, leading to a lateral dimension size-dependent catalytic activity⁵. However, the deficiencies of aggregation and low conductivity of MoS_2 are still challenging problems which limit their widespread application. Monolayer metallic MoS_2 with high conductivity have been explored for HER catalyst^{17,18}. For example, metallic nanosheets of 1T- MoS_2 chemically exfoliated via lithium intercalation from semiconducting 2H- MoS_2 nanostructures grown directly on graphite was reported as a highly competitive earth-abundant catalyst due to favorable kinetics, metallic conductivity, and proliferation of active sites¹⁷. Meanwhile, the functionalization of MoS_2 materials with conductive templates or supports to form a multidimensional structure has been recently presented as an effective approach to improve the catalytic activity¹⁹. Carbon materials are promising candidates to support catalysts due to their unique physicochemical properties. Increasing evidence has demonstrated that the significantly improved HER performance of MoS_2 can be implemented by incorporating MoS_2 with carbon nanotube (CNTs), graphene, graphite or other mesoporous carbon to form a MoS_2 -based hybrid or composite^{20–24}. For example, Dai *et al.* developed an HER advanced catalyst of MoS_2 nanoparticles grown on graphene via a facile solvothermal approach, the resulted highly exposed edges and excellent electrical coupling to the underlying graphene sheets endowed the hybrid catalyst with excellent HER activity²⁵. The high electrical conductance of the carbon architecture facilitates the electron charge transfer leading to enhanced performance, whilst the active site introduced in the supports is another effective method to enhance the activity of the catalyst. The incorporation of electron-rich nitrogen atoms into the carbon architecture promotes the interaction between neighboring carbons and electrons, providing a superior heteroatom-doped catalyst^{23,26}. Nitrogen-doped graphene is one of the most promising candidates owing to its high chemical stability, good electrical conductivity and intrinsic moderate catalytic activity for hydrogen evolution²⁷. For instance, Qiao's group has obtained porous C_3N_4 nanolayers@N-graphene films by integrating porous C_3N_4 nanolayers with nitrogen-doped graphene sheets, displaying an unbeatable HER performance which stems from highly exposed active sites, hierarchical porous structure and 3D conductive graphene network²⁸. Furthermore, it has been reported that the p-type MoS_2 nanoplatelets grown on the n-type nitrogen-doped reduced graphene oxide (N-RGO) can form p–n junctions on nanoscale and act as an outstanding catalyst in photocatalytic HER²⁹. However, to the best of our knowledge, little work on the influence of the composition and morphology of the nitrogen-doped graphene support to the HER has been reported.

Herein, we report MoS_2 nanoparticles decorated nitrogen-doped reduced graphene oxide (N-RGO), a three-dimensional (3D) hybrid structure with macro-porosity by a hydrothermal route, which can efficiently catalyze HER under acidic conditions with low overpotential and small Tafel slope. The resulting MoS_2 /N-RGO hybrid materials display high-performance HER catalytic activity. By controlling the elemental composition, electrical conductance and morphology of the as-prepared 3D MoS_2 /N-RGO hybrid, the enhanced electrocatalytic activity mechanism toward hydrogen evolution was investigated in detail.

Results

Structural and compositional characterization. Firstly, graphene oxide (GO) and $(\text{NH}_4)_2\text{MoS}_4$ were sonicated in N, N-dimethylformamide (DMF) to generate a highly dispersed suspension. Hydrazine and polypyrrole (PPy) were then slowly added into the suspension, followed by hydrothermal assembly at 180 °C for 12 h. Besides as a reductant, hydrazine could work as a nitrogen source as PPy during the assembly process. Regarding to PPy, it acted not only as the nitrogen source, but also was used to fabricate 3D network structure which promoted the BET specific surface and conductivity of the final product. The hydrothermal environment plays a significant role in the nitrogen doping because in the absence of hydrothermal environment, the GO could be reduced directly without the incorporation of nitrogen into the graphene network. During the hydrothermal process, $(\text{NH}_4)_2\text{MoS}_4$ and GO were reduced to MoS_2 nanoparticles and RGO respectively by hydrazine; and MoS_2 nanoparticles were grown on the RGO accompanied by incorporation of nitrogen species into the graphene lattice²². In order to sustain the perfect 3D structure, the as-prepared hybrid was directly dehydrated by a vacuum freeze drier and then heated at 600 °C for 3 h under nitrogen (Fig. 1). Other samples were also fabricated in the same way for comparison: N-RGO where $(\text{NH}_4)_2\text{MoS}_4$ is removed from the starting materials; MoS_2 /N-RGO where nitrogen-doped graphene sheets decorated with half quantity of MoS_2 ; MoS_2 /N'-RGO where the addition order of hydrazine and PPy is exchanged; MoS_2 /N-RGO' is the sample fabricated by GO of higher oxidation degree.

The 3D MoS_2 /N-RGO synthesized was comprehensively characterized by a number of microscopic and spectroscopic tools. Scanning electron microscopic (SEM) images revealed a 3D network structure

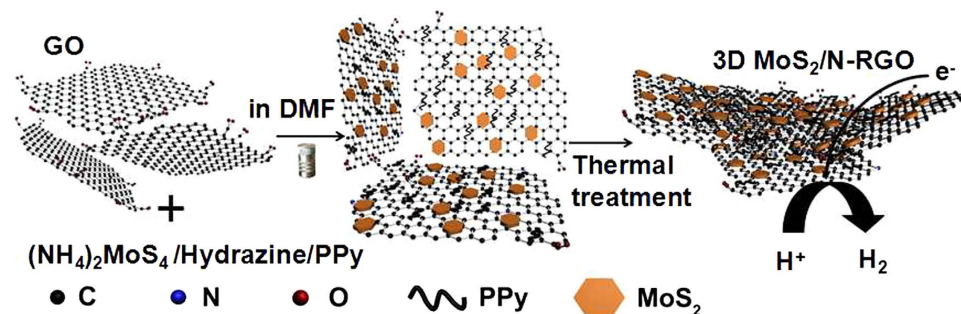


Figure 1. Schematic present of synthetic method and resultant 3D $\text{MoS}_2/\text{N-RGO}$ hybrid for hydrogen evolution reaction.

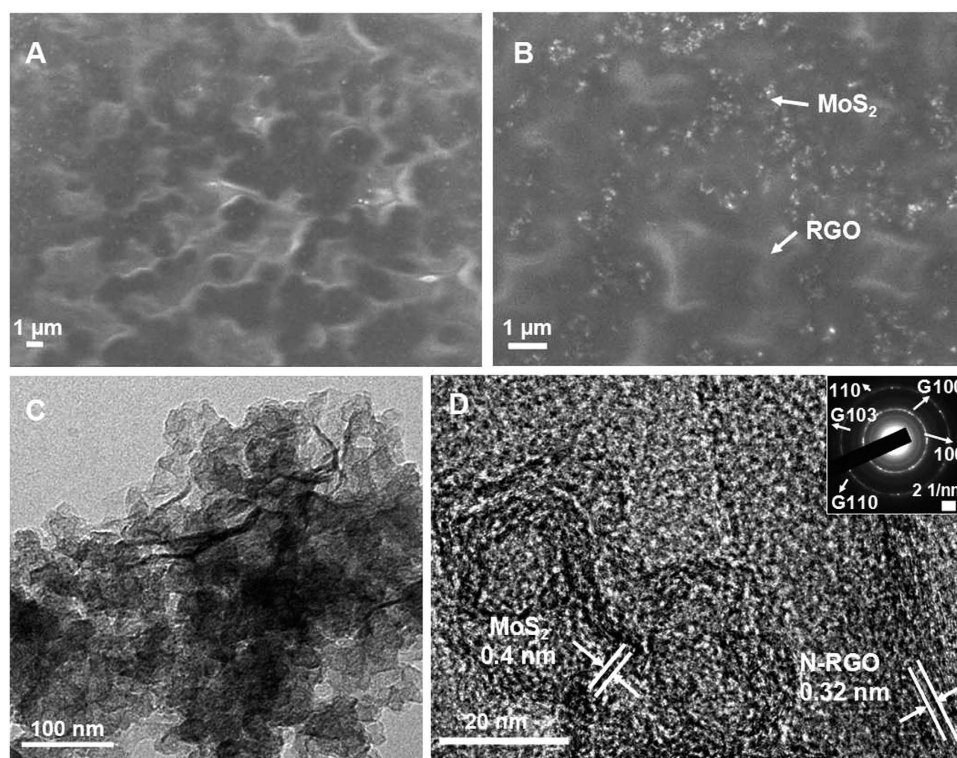


Figure 2. (A) SEM and (B) magnified SEM images of 3D $\text{MoS}_2/\text{N-RGO}$ hybrid. (C) TEM images and (D) high-resolution TEM images of the 3D $\text{MoS}_2/\text{N-RGO}$ hybrid.

with crumpled few-layered graphene sheets with plenty of folded edges conglutinated by PPy and decorated by numerous MoS_2 nanoparticles with a lateral dimension of about 35 nm (Fig. 2A,B). The large area of graphene provided a large attachment surface for MoS_2 nanoparticle growth and created an efficient template to restrain the aggregation of particles. The information presented by the transmission electron microscopic (TEM) images was consistent with the SEM in that there were laminar structures of graphene in the catalyst and MoS_2 nanoparticles were evenly distributed on N-RGO sheets, suggesting successful assembly between the nanoparticles and the graphene sheets (Fig. 2C). The high resolution TEM (HRTEM) images of the 3D $\text{MoS}_2/\text{N-RGO}$ structure showed the MoS_2 with a lattice spacing of 0.4 nm, consistent with the spacing between the two sulfur atoms, which indicates the presence of MoS_2 . The thickness of N-RGO was 0.34 nm, in accordance with the theoretical prediction³⁰ (Fig. 2D). The corresponding selected area electron diffraction (SAED) revealed several sets of diffraction signals assigned to planes of hexagonal-phase of MoS_2 and N-RGO (inset in Fig. 2D). The two separated diffraction rings can be indexed to the (100) and (110) planes of MoS_2 , and the diffraction of the (103) assigned to the N-RGO can also be observed. Notably, a distinguished and coherent interface was observed in $\text{MoS}_2/\text{N-RGO}$, which is known to facilitate the electron transfer to enhance catalytic activity³⁰.

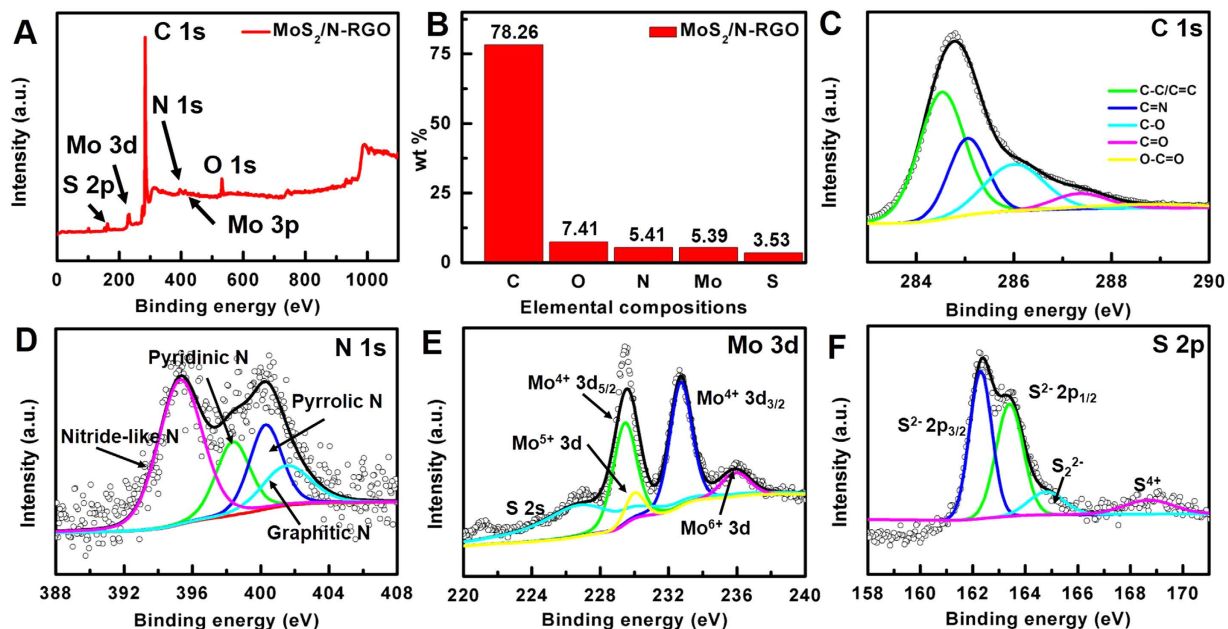


Figure 3. (A) The survey XPS spectrum and (B) corresponding elemental compositions of 3D MoS₂/N-RGO catalyst. High-resolution (C) C 1s, (D) N 1s, (E) Mo 3d and (F) S 2p XPS spectra of 3D MoS₂/N-RGO (the red curve: background; the black curve: fitting line; the open circle: raw data).

X-ray photoelectron spectroscopy (XPS) measurements were performed to investigate the components of the prepared 3D MoS₂/N-RGO hybrid. As shown in Fig. 3A, the XPS survey spectrum of the 3D MoS₂/N-RGO hybrid shows the characteristic peaks of N-RGO; including a prominent graphitic C 1s peak at 284 eV, a strong O 1s peak at 532 eV and an N 1s peak located at 402 eV. The peaks assigned to Mo 3d and S 2p in MoS₂ were also present in the survey spectrum, which confirmed the successful assembly between the nanoparticles and the graphene sheets. The detailed information of elemental composition is presented in Fig. 3B. The corresponding element analysis revealed that the atomic ratio of Mo (0.75%) to S (1.46%) was about 1:2, consistent with the theoretical value of MoS₂. It is worthy mentioning that the N/C atomic ratio of the 3D MoS₂/N-RGO hybrid was calculated up to be 5.9%, much higher than that of N-doped graphene from previous reports^{31–33}. The higher nitrogen element included in the 3D MoS₂/N-RGO hybrid could lead to a more efficient catalytic activity³⁰.

In the high resolution XPS of C 1s (Fig. 3C), a prominent C=N peak (285.2 eV) can be observed, confirming the successful incorporation of N atoms into the 3D MoS₂/N-RGO hybrid. The binding energy of C-C/C=C (284.5 eV) in the 3D MoS₂/N-RGO hybrid was downshifted by ~0.5 eV compared to the graphitized carbon peak of pristine GO (285.0 eV)³⁴. This implies a remarkable amount of charge transfer from graphene to MoS₂, which can improve both the current density and catalytic activity of the catalyst. The N 1s region shown in Fig. 3D can be divided into four peaks: (1) the peak at 395.7 ± 0.3 eV is attributed to nitride-like species or aromatic N-imines; the pyridine-like peak located at 398.7 ± 0.3 eV; (3) the pyrrolic or amine moieties peak at 400.3 ± 0.3 eV; and (4) the graphitic nitrogen peak at 401.4 ± 0.3 eV^{35,36}. The high content of nitride-like peak was attributed to the adsorbed N₂H₄³⁷. It was reported that hydrazine were reducing reagents³⁷, the nitride-like peak located at 395.7 eV should be ascribed to the adsorbed hydrazine, even samples experienced several-times washing and dialysis³⁷. PPy was an efficient nitrogen precursor to fabricate nitrogen-doped graphene due to its excellent catalytic activity and high durability, which mainly existed in forms of pyrrolic N (401.0 ± 0.2 eV) and pyridinic N (398.1 ± 0.2 eV) in NGO name³⁸. The characteristic peaks of nitride-like peak, pyrrolic and pyridinic N in the resulting hybrid materials indicated that hydrazine and PPy were both nitrogen source.

As shown in Fig. 3E, five characteristic peaks located at 228.8, 232.7, 230.0, 235.9 and 226.7 eV, corresponding to Mo⁴⁺ 3d_{5/2}, Mo⁴⁺ 3d_{3/2}, MoO₃ or MoO₄²⁻, Mo⁵⁺ and S 2s, respectively, are displayed in the high resolution XPS of Mo 3d³⁹. The two main Mo 3d_{5/2} and Mo 3d_{3/2} peaks are close to the meta-stable 1T phase of MoS₂ (associating with better HER performance)^{40,41}, and the peak of Mo⁶⁺ results from the oxidation of the catalyst in air to form MoO₃ or MoO₄²⁻³⁴ and the partial reduction of Mo⁶⁺ results in the formation of Mo⁵⁺³⁹. The deconvolution of S 2p spectra (Fig. 3F) has yielded four main peaks located at 162.3, 163.4, 164.8 and 168.7 eV, which are assigned to S 2p_{3/2}, S 2p_{1/2}, S₂²⁻ (or S²⁻) ligands and the S⁴⁺ of sulfate groups, respectively^{40,43,44}. The existence of S₂²⁻ and/or S²⁻ is likely to be caused by the doping of S in the graphene sheet or formation of an S-rich MoS₂ structure. This facilitates the development of high HER activity at the active sites for HER on the edge of MoS₂ where the unsaturated S atoms are exposed to adsorb H⁺^{40,43,44}.

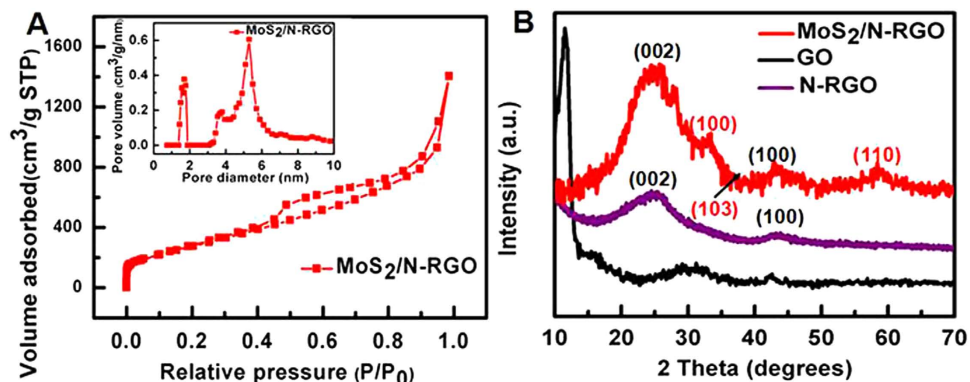


Figure 4. (A) Nitrogen adsorption-desorption isotherm and pore-size distribution (inset) of 3D MoS₂/N-RGO. (B) XRD patterns of GO, N-RGO and 3D MoS₂/N-RGO.

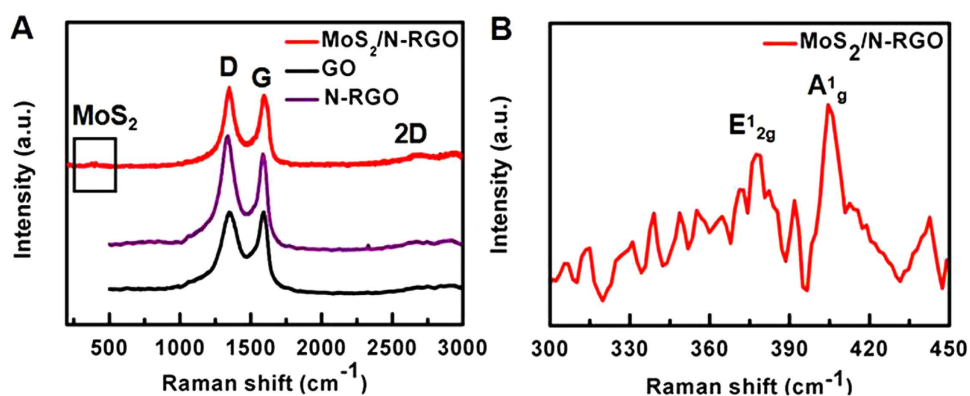


Figure 5. (A) Raman spectra of 3D MoS₂/N-RGO, GO and N-RGO. (B) Raman spectra of 3D MoS₂/N-RGO in the range of 300–450 cm⁻¹.

The porosity structures of the 3D MoS₂/N-RGO hybrid were characterized using the nitrogen isotherm (Fig. 4A). The N₂ adsorption-desorption isotherm of 3D MoS₂/N-RGO hybrid showed an intermediate shape between types II and IV (in the IUPAC classification) with a small H3 hysteresis loop extending from P/P₀ = 0.45 to 0.95, demonstrating the presence of mesoporous structure and slit-shaped pores⁴⁵. The N₂ adsorption-desorption measurement revealed the as-prepared 3D MoS₂/N-RGO hybrid had a Brunauer-Emmett-Teller (BET) specific surface area of 1066.6 m² g⁻¹, which was significantly larger than many reported porous carbon nanomaterials^{46–49}. The superior BET specific surface area was resulted from that unique 3D structure of MoS₂/N-RGO hybrid materials and formation of numerous porous structures during the sintering and elimination process in the presence of PPy. The pore-size distribution of the 3D MoS₂/N-RGO hybrid calculated by the DFT method exhibited one dominant peak at 5.30 nm and another two weak peaks at ~1.56 and 3.79 nm, respectively (Inset of Fig. 4A), suggesting that micropores and mesopores co-exist within the 3D MoS₂/N-RGO hybrid. Figure 4B showed XRD patterns of the 3D MoS₂/N-RGO hybrid, N-RGO and GO. The sharp oxidation peak at 11.7° distinctly exhibited in GO but had completely disappeared in MoS₂/N-RGO and N-RGO. However, for the latter a broad peak centered at around 24° which was associated with a graphitic crystal structure, implying that the GO was efficiently deoxidized during the hydrothermal process. In comparison with N-RGO, the diffraction peaks at 33.5°, 39.4° and 58.9° corresponding to (100), (103) and (110) of MoS₂ could be observed in MoS₂/N-RGO, indicating the MoS₂ nanoparticles have been successfully grown on the N-RGO surface. The disappearance of the typical (002) peak of MoS₂ located at 14.1° resulted from the small size of the MoS₂ nanoparticles preferring to attach to the surface of the silicon supports^{50–52}.

The successful fabrication of the 3D MoS₂/N-RGO hybrid was further confirmed by Raman spectra analysis. As shown in Fig. 5A, the characteristic peaks located at ~1346 and 1599 cm⁻¹ associated with the D and G bands of graphene can be observed in GO, N-RGO and MoS₂/N-RGO. The reduction of GO induces the recovery of the graphene structure, which leads to the increase of 2D bands in N-RGO and MoS₂/N-RGO. Notably, compared to the GO with a G band at 1599 cm⁻¹, the G band of N-RGO (1586 cm⁻¹) and MoS₂/N-RGO (1593 cm⁻¹) showed a downshift due to the incorporation of N heteroatoms⁴⁹. The characteristic peaks of MoS₂ located at 375.5 and 403.5 cm⁻¹ (and assigned to the in-plane E¹_{2g}

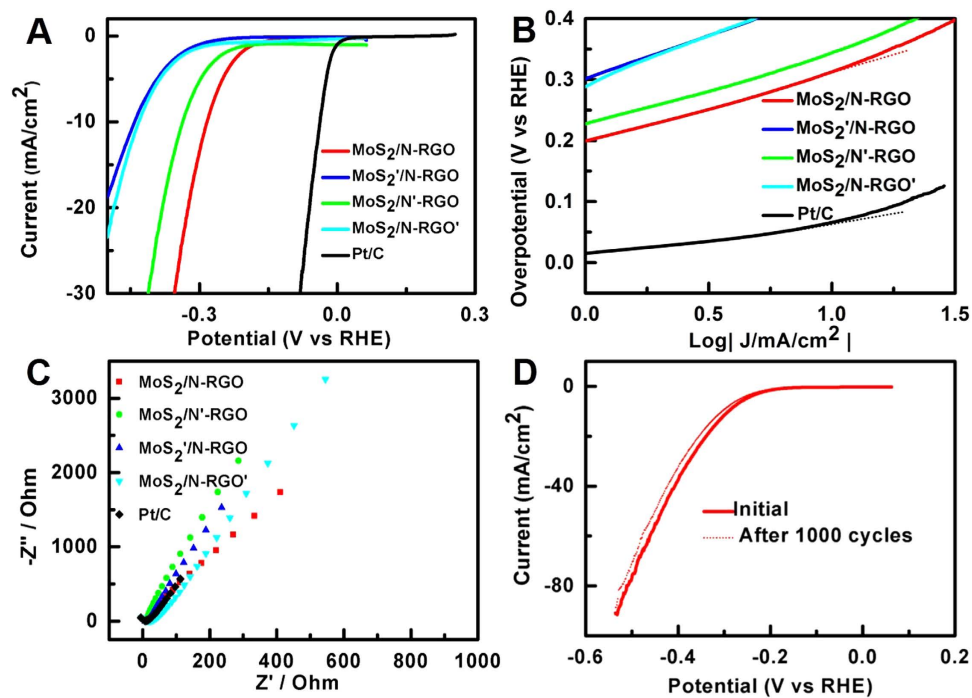


Figure 6. (A) Polarization curves obtained of catalysts as indicated and (B) corresponding Tafel plots recorded on glassy carbon electrodes with a catalyst loading of 0.28 mg/cm^2 (solid lines), and fitted Tafel plots (dashed dot). (C) Nyquist plots of the 3D $\text{MoS}_2/\text{N-RGO}$ catalyst recorded in nitrogen-purged $0.5 \text{ M H}_2\text{SO}_4$ from 10^{-2} to 10^6 Hz with an AC amplitude of 5 mV . (D) Stability test for the 3D $\text{MoS}_2/\text{N-RGO}$ catalyst. Negligible current was lost after 1000 cycles from -0.2 to -0.8 V at 10 mV/s .

and out-of-plane A_g^1 vibrational modes of the hexagonal MoS_2) were observed in the Raman spectrum of $\text{MoS}_2/\text{N-RGO}$ ranging from 300 to 450 cm^{-1} (Fig. 5B)⁵³. In comparison with MoS_2 , the decreasing difference between the peak frequencies of E_{2g}^1 and A_g^1 for $\text{MoS}_2/\text{N-RGO}$ (27 cm^{-1}) suggested that the interaction between the Mo precursors and graphene could efficiently avoid the aggregation of MoS_2 ³⁹.

Electrochemical activity and durability. The electrocatalytic activity of the as-prepared 3D $\text{MoS}_2/\text{N-RGO}$ hybrid toward the HER was investigated in $0.5 \text{ M H}_2\text{SO}_4$ solution with commercial Pt catalyst ($10 \text{ wt } \% \text{ Pt}$ on carbon black) as the reference. N-RGO, $\text{MoS}_2'/\text{N-RGO}$, $\text{MoS}_2/\text{N'-RGO}$, $\text{MoS}_2/\text{N'-RGO}$ and $\text{MoS}_2/\text{N-RGO}'$ were also prepared to investigate in detail the influence of composition and structure to the HER catalytic activity. XPS (Fig. S1), XRD (Fig. S2) and SEM (Fig. S3) confirmed the successful synthesis of the three catalysts. As shown in Fig. 6A, the polarization curves revealed that the as-generated 3D $\text{MoS}_2/\text{N-RGO}$ displayed a small onset potential of 112 mV for HER activity, after which the cathodic current rose rapidly. In contrast, the $\text{MoS}_2'/\text{N-RGO}$ (210 mV), $\text{MoS}_2/\text{N'-RGO}$ (155 mV) and $\text{MoS}_2/\text{N-RGO}'$ (200 mV) showed more negative onset potential, indicating lower HER activity. The current intensity of the 3D $\text{MoS}_2/\text{N-RGO}$ hybrid was higher than the other catalysts along the whole potential region. To further investigate the HER activity, the linear portions of the Tafel plots were fitted to the Tafel equation ($\eta = a + b \log |j|$), where j is the current density and b is the Tafel slope. In Fig. 6B, the generated $\text{MoS}_2/\text{N-RGO}$ hybrid showed a Tafel slope of 44 mV per decade, which outperformed $\text{MoS}_2'/\text{N-RGO}$ of 114 mV per decade, $\text{MoS}_2/\text{N'-RGO}$ of 71 mV per decade and $\text{MoS}_2/\text{N-RGO}'$ of 57 mV per decade. The slight higher Tafel slope compared to MoS_2 nanoparticle on graphene with a value of 41 mV ²⁵ and metallic MoS_2 nanosheets on graphite with a value of 43 mV ¹⁷ may be resulted from the reduce of the conductivity of nitrogen dopants. Notably, the lower HER activity of other catalysts prepared led to higher Tafel slopes than the $\text{MoS}_2/\text{N-RGO}$ hybrid. Electrochemical impedance spectroscopy (EIS) was measured to evaluate the HER catalytic activity of these catalysts. The $\text{MoS}_2/\text{N-RGO}$ displayed the lowest faradaic impedance among these prepared catalysts (Fig. 6C) and was comparable with Pt/C, suggesting a small charge transfer resistance in the $\text{MoS}_2/\text{N-RGO}$ ⁵⁴. Such a low charge transfer resistance resulted both from a distinguished and coherent interface between the $\text{MoS}_2/\text{N-RGO}$ hybrid and from the good conductivity of N-RGO, leading to efficient electrical communication between the catalytic edge sites and underlying electrodes to facilitate the kinetic response of HER^{55,56}. Stability is a significant criterion to evaluate the practicality of a HER catalyst. After a long period of 1000 potential-cycling between -0.4 and 0 V , the as-generated $\text{MoS}_2/\text{N-RGO}$ hybrid exhibited a negligible decrease in the current density, indicating outstanding electrochemical stability (Fig. 6D). The unique three-dimensional structure of the MoS_2 nanoparticles grown on N-doped graphene may account for the good stability

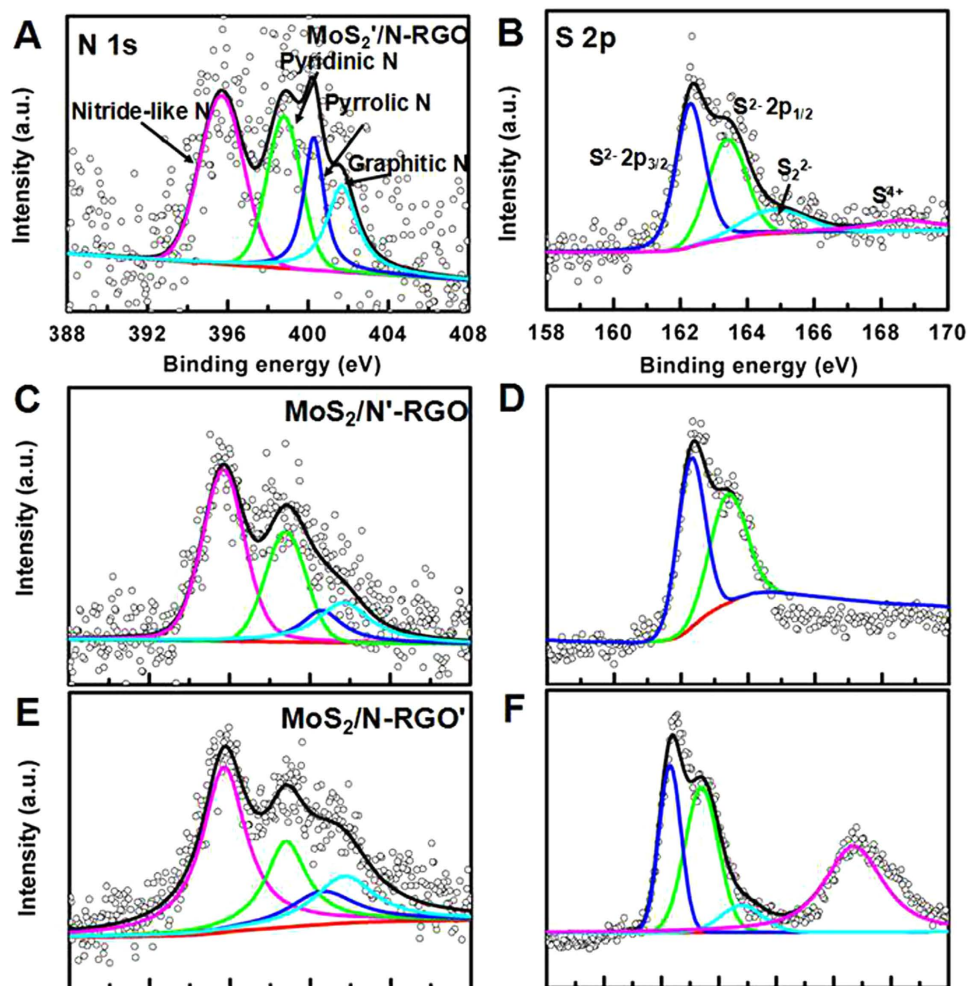


Figure 7. High-resolution XPS spectra of (A), (C), (E) N 1s and (B), (D), (F) S 2p of the MoS₂/N-RGO, MoS₂/N'-RGO and MoS₂/N-RGO' (the red curve: background; the black curve: fitting line; the open circle: raw data).

that was observed. The supports of the N-doped graphene restrain the agglomeration of MoS₂ nanoparticles and facilitate the transport of electrolyte ions^{56,57}. In addition, the intercalation of sulfur into the graphene structure to form a stable covalent bond also contributes to the excellent stability (Fig. 3F).

Discussion

These results demonstrated that the 3D MoS₂/N-RGO hybrid exhibited superior electrochemical catalytic activity to MoS₂/N-RGO, MoS₂/N'-RGO and MoS₂/N-RGO', the reasons for the enhanced mechanism of MoS₂/N-RGO hybrid toward HER activity can be explained as followed. Firstly, structure analysis derived from Nitrogen Sorption date (Fig. S4) showed that the 3D MoS₂/N-RGO hybrid had an 87% mesoporous surface area ratio (the mesopore surface area to BET total surface area), which was larger than MoS₂/N-RGO and MoS₂/N'-RGO, which lead to better reactant accessibility and superior efficiency of build of the triple-phase boundaries (gas–electrode–electrolyte)^{58,59}. Secondly, both of the N-RGO without decorated MoS₂ (Fig. S5) and MoS₂/N-RGO with less decorated MoS₂ (Fig. 6A) displayed a little HER activity compared to the proposed MoS₂/N-RGO hybrid, indicating the composition of MoS₂ nanoparticles played a critical role in the active HER catalytic performance¹³.

Furthermore, the proposed hybrid contains effective catalytic sites originated from sulfur in MoS₂ and nitrogen impurities in RGO boost the activity of the catalyst. The relative content (RC) of S₂²⁻ and/or apical S²⁻ related to active HER catalytic activity, where RC is mean that sulfur elemental compositions (wt %) (Table S1) multiply sulfur distribution (at. %) (Table S2), in MoS₂/N-RGO hybrid is 3.5-folds and 1.9-folds higher than MoS₂/N-RGO (Fig. 7B), MoS₂/N-RGO' (Fig. 7F), while the distribution of S₂²⁻ and/or apical S²⁻ in MoS₂/N'-RGO (Fig. 7) was not observed. For the nitrogen impurities, it was found the nitrogen hybrid species are sensitive to the graphene oxidation degree, MoS₂, PPy and hydrazine (Fig. 7). The MoS₂/N-RGO hybrid contains more pyrrolic N and less pyridinic N than MoS₂/N-RGO, MoS₂/N'-RGO and MoS₂/N-RGO' (Fig. 7A,C,E, Table S3), it was speculated that the pyrrolic N instead

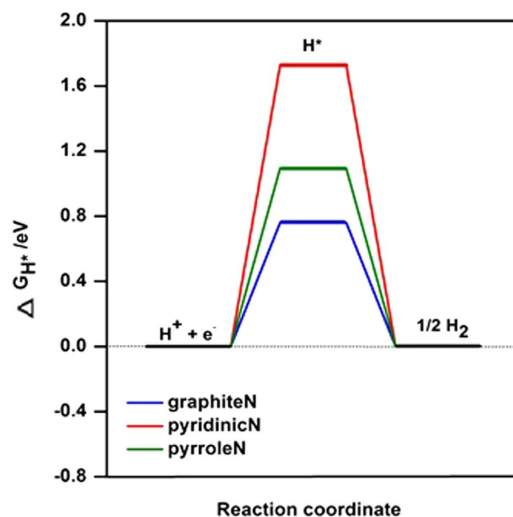


Figure 8. The calculated free-energy diagram of HER for graphite N, pyridinic N and pyrrole N catalysts.

of pyridinic N is more active towards HER catalysis. The overall HER pathway can be described by a three-state diagram comprising an initial state $H^+ + e^-$, an intermediate adsorbed H^* , and a final product $1/2H_2$ ⁶⁰. The Gibbs free-energy of the intermediate state, $|\Delta G_{H^*}|$, has been considered as a major descriptor of the HER activity for a wide variety of metal catalysts. The optimum value of $|\Delta G_{H^*}|$ should be zero. Density Functional Theory (DFT) calculations showed that graphite N shows the smallest Gibbs free-energy ($|\Delta G_{H^*}|$) value of 0.76 eV, followed by pyrrole N of 1.10 eV and pyridinic N of 1.73 eV, indicating graphite N was the best electrocatalytic activity from the viewpoint of thermodynamics and pyrrole N also possessed the enhanced electrocatalytic activity than pyridinic N toward HER (Fig. 8). Additionally, the higher oxidation degree of the RGO can induce the more nitrogen hybrid (Table S1), larger mesopore surface area but larger faradaic impedance (Fig. 6C), which led to a lower HER performance. Therefore, it was demonstrated that the efficient active nitrogen and sulfur species combined with the good conductance and large mesoporous surface area ratio of 3D MoS_2/N -RGO hybrid were all beneficial to the enhanced HER catalytic activity.

Conclusion

In summary, we have developed an efficient hydrothermal route to successfully synthesize a 3D hybrid material of nitrogen-doped graphene oxide sheets-supported molybdenum disulfide nanoparticles with high capability of hydrogen evolution. The morphology and element of the as-generated 3D MoS_2/N -RGO hybrid were characterized by comprehensively microscopic and spectroscopic methods including SEM, TEM, HRTEM, BET, XRD, XPS and Raman spectroscopy. Electrochemical characterization data including low onset potential, small Tafel slope, low charge transfer resistance and high electrochemical stability revealed the high performance of the MoS_2/N -RGO hybrid toward HER catalytic activity compared to analogous composites. Detailed experimental analysis implies that the abundant active S_2^{2-} and/or S^{2-} ligand species and pyrrolic nitrogen and graphene N of the 3D MoS_2/N -RGO hybrid and its good conductance were all beneficial to its enhanced HER catalytic activity. The exploration of the enhanced HER mechanism of the MoS_2/N -RGO hybrid provides guidelines to design and construct 3D hybrids to maximize their catalytic efficiency.

Methods

Materials synthesis. Graphene oxide (GO, made from graphite flake, GO', small flakes and dry platelets) was obtained from XF NANO, INC and graphene supermarket, respectively. Hydrazine monohydrate ($N_2H_4 \cdot H_2O$, 80%) was purchased from Guangdong Guanghua Sci-Tech Co., Ltd (JHD). $(NH_4)_2MoS_4$, Polypyrrole (PPy), and Pt/C (10% Pt) were obtained from Sigma-Aldrich. N, N-dimethylformamide (DMF, $\geq 99.5\%$) and KOH were purchased from Sinopharm Chemical Reagent Co., Ltd. Sulfuric acid (H_2SO_4 , 95–98%) and ethanol (99.9%) was acquired from Beijing Chemical Works. All aqueous solutions were prepared with doubly distilled water.

Materials preparation. 22 mg of $(NH_4)_2MoS_4$ and 10 mg of GO was dispersed in 10 ml of DMF by 10 min of sonication at room temperature. Then 0.1 ml of $N_2H_4 \cdot H_2O$ was added to the above solution followed by 30 min of sonication. 100 mg PPy was then added, followed by sonication for 10 min to form a stable complex solution. The mixture solution was transferred to a 40 mL Teflon-lined autoclave and heated in an oven at 180 °C for 12 h with no intentional control of ramping or cooling rate.

The product was collected by centrifugation at 10000 rpm for 5 min and washed with DI water several times to remove most of the DMF. Subsequently, the product was redispersed in 3 ml of DI water and freeze-dried overnight, followed by thermal treatment at 600 °C for 3 h in N₂ gas with 400 standard cubic centimeters per minute (sccm) to remove the organic species (DMF) and improving crystallinity. The obtained product was denoted as 3D MoS₂/N-RGO. Other samples were also fabricated in the same way for comparison: N-RGO where (NH₄)₂MoS₄ is removed from the starting materials; MoS₂/N-RGO where Nitrogen-doped graphene sheets decorated with half quantity of MoS₂; MoS₂/N'-RGO where the addition order of hydrazine and PPy is exchanged; MoS₂/N-RGO' is the sample fabricated by GO of higher oxidation degree.

Characterizations. X-ray diffraction (XRD) was performed with a Rigaku X-ray diffractometer with Cu KR target. The porosity was measured with a nitrogen adsorption-desorption isotherm using a surface area analyzer (QuadraSorb SI 2000-08, Quantachrome Instruments). The structure of products was observed under a field-emission scanning electron microscope (SEM; JEOL-6300F, 3 kV) and a transmission electron microscope (TEM; JEM-2010, 200 kV). X-ray photoelectron spectroscopy (XPS) spectra were obtained using an AXIS ULTRA^{DLD} instrument equipped with an Al K α X-ray source. Raman spectrum of powder samples were recorded on an InVia-Reflex Raman microscope with a laser excitation wavelength of 532 nm.

Electrochemical measurement. All electrochemical studies were performed using a CHI 852C electrochemical workstation (Shanghai Chenhua Instrument Co., China) in a standard three-electrode setup. A three-electrode configuration consisting of a saturated calomel electrode (SCE) as the reference electrode, a graphite rod as the counter electrode and a glass carbon RDE after loading the catalyst as the working electrode was employed. Typically, 0.5 mg of catalyst was dispersed in 500 μ l of DI water by sonication to form a homogeneous ink. Then, 20 μ l of the catalyst ink (containing 20 μ g of catalyst) was loaded onto a glassy carbon electrode (GCE) of 3 mm in diameter (loading \sim 0.283 mg/cm²). After the catalyst ink dried, 5 μ l of 1 wt% Nafion solution was dropped onto the GCE and the working electrode was prepared. Linear sweep voltammetry (LSV) was conducted in nitrogen-purged 0.5 M H₂SO₄ at a scan rate of 3 mV s⁻¹ at 1400 rpm. Electrochemical impedance spectroscopy (EIS) was carried out in the same configuration in nitrogen-purged 0.5 M H₂SO₄ from 10⁻² to 10⁶ Hz with a modulation amplitude of 5 mV. SCE was calibrated with respect to reversible hydrogen electrode (RHE). The calibration was performed in the high purity H₂ saturated electrolyte with a Pt wire as the working electrode and the counter electrode. LSV were run at a scan rate of 0.1 mV s⁻¹, and the potential at which the current crossed zero was taken to be the thermodynamic potential for the hydrogen electrode reactions. In 0.5 M H₂SO₄, E (RHE) = E (SCE) + 0.314 V. All the potentials reported in our manuscript are against RHE.

Density Functional Theory (DFT) calculations. The computations for DFT calculations were performed using the *ab initio* density functional code VASP with the PBE exchange correlation functional and PAW potentials. The PBE functional was chosen in order to obtain reasonable adsorption energies. The Brillouin zone is sampled using a 1 \times 1 \times 1 Monkhorst-Pack grid for the geometry optimizations and a 9 \times 9 \times 1 Monkhorst-Pack grid for the calculations of electronic properties. A basis set with the cut-off energy of 400 eV was chosen. The convergence criteria for energies and forces are set to 1.0 \times 10⁻⁴ eV and -0.05 eV/Å, respectively. All calculations were performed using spin unrestricted method.

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Author Contributions

H.D., C.L. and X.Z. conceived the project and designed the experiment; C.L. synthesized the material; W.D., Y.C. and H.L. performed the microscopic and spectroscopic study; C.L. and B.F. performed the electrochemistry experiments; L.H., X.Q. and H.Y. performed Density Functional Theory (DFT) calculations; H.D. and C.L. analyzed the data and wrote the main manuscript text; H.Y., B.F. and X.Z. modified the manuscript. All authors have given approval to the final version of the manuscript.

Additional Information

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