

Boosting hydrogen evolution on MoS₂ via co-confining selenium in surface and cobalt in inner layer

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The lack of highly efficient, inexpensive catalysts severely hinders large-scale application of electrochemical hydrogen evolution reaction (HER) for producing hydrogen. MoS₂ as a low-cost candidate suffers from low catalytic performance. Herein, taking advantage of its tri-layer structure, we report a MoS₂ nanofoam catalyst co-confining selenium in surface and cobalt in inner layer, exhibiting an ultra-high large-current-density HER activity surpassing all previously reported heteroatom-doped MoS₂. At a large current density of 1000 mA cm⁻², a much lower overpotential of 382 mV than that of 671 mV over commercial Pt/C catalyst is achieved and stably maintained for 360 hours without decay. First-principles calculations demonstrate that inner layer-confined cobalt atoms stimulate neighbouring sulfur atoms while surface-confined selenium atoms stabilize the structure, which cooperatively enable the massive generation of both in-plane and edge active sites with optimized hydrogen adsorption activity. This strategy provides a viable route for developing MoS₂-based catalysts for industrial HER applications.

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The electrochemical splitting of water for producing hydrogen via renewable energy supply is a significant, carbon-neutral technology for clean energy generation^{1–3}. However, the large-scale implementation of the reaction process is severely limited by scarcity of traditional platinum-based catalysts^{4,5}. Two-dimensional MoS₂, an earth-abundant material with unique structural and chemical properties, has shown attractive catalytic activity for the reaction and is considered as a potential alternative to the precious platinum-based catalysts in acidic medium^{6–14}. However, only the edge S sites of pure MoS₂ are catalytically active for the HER and the vast amount of S sites in the basal plane are quite inert and not sufficiently utilized^{8,15–18}. Creating abundant edges is straight-forward but leads to high surface energy and low stability^{15,19,20}. Doping metallic heteroatoms into the MoS₂ lattice, on the one hand can effectively activate the basal plane S atoms and introduce in-plane active sites for the HER^{10,21–25}, whereas on the other hand it may also unavoidably lead to over-activation of the edges, resulting in overly strong adsorption of hydrogen at the edge sites which is detrimental to the HER activity. Thus, selectively activating the inert basal plane combined with stabilizing the edges without quenching the activity can maximally increase active sites of MoS₂ for the HER, but is highly challenging owing to the difficulty in balancing the activity and stability.

MoS₂ possesses a tri-layer atomic structure, which allows multidimensional manipulation of atomic composition to modulate the catalytic activity. Herein, we report a strategy of co-confining Se in the surface and Co in the inner layer of the MoS₂ lattice, which enables simultaneously activation of the basal plane and stabilization of the edges and optimization of the hydrogen adsorption activity. Combined with a morphology-controlling method of fabricating three-dimensionally nanofoam architecture to promote edge formation, the massive generation of both in-plane and edge active sites are achieved in the synthesized MoS₂ nanofoam with co-confined Co and Se atoms (Co/Se-MoS₂-NF). Such a catalyst exhibits a high HER activity at large current densities, which, to the best of our knowledge, surpasses those of all previously reported heteroatom-doped MoS₂ catalysts and even the commercial precious Pt/C catalyst in acidic electrolyte. At a large current density of 1000 mA cm⁻², the optimized Co/Se-MoS₂-NF presents a much lower overpotential of 382 mV than that of 671 mV over the commercial Pt/C, and works stably for over 360 h without any activity decay. Density functional theory (DFT) studies demonstrate a synergy between the activating effect of confining Co in the inner Mo-layer and the stabilizing effect of confining Se in the surface S-layer, which promotes the formation of both in-plane and edge active sites and meanwhile optimizes the adsorption free energies of hydrogen. This strategy opens up a new prospect of tailoring the catalytic performance of MoS₂ toward large-scale HER applications through confining multi-elements in different layers.

Results

Se-doping promotes edge formation. The terminating atoms of the MoS₂ edges determine the stability and reactivity of the edges. Selenium being in the same group with S was introduced into MoS₂ to modify the edge properties. A series of Se-doped MoS₂ nanofoam (Se-MoS₂-NF) with different Se content were synthesized via a one-pot solvothermal method by using (NH₄)₆Mo₇O₂₄·4H₂O, CS₂, and Se powder as precursors. Monodispersed SiO₂ nanospheres were used as templates to fabricate a foam-like morphology. As shown in scanning electron microscopy (SEM) (Fig. 1a) and transmission electron microscopy (TEM) (Supplementary Fig. 1) images, the Se-MoS₂-NF possesses a nanofoam morphology with abundant spherical

cavities (average pore size ~100 nm), which will favor the mass transportation of reactants to access more active sites of the catalyst^{15,19,21,26}.

Comprehensive spectroscopic characterizations of the Se-MoS₂-NF samples were performed to investigate the geometric distribution and electronic states of the Se atoms. Energy-dispersive X-ray (EDX) analysis shows that the Mo, S, and Se elements are homogeneously distributed in the nanofoam framework (Fig. 1b). X-ray diffraction (XRD) analysis show no obvious difference in the crystallographic structure and lattice parameters of the Se-MoS₂-NF compared with those of the pure MoS₂-NF, which excludes the formation of new crystal phases after the Se doping (Supplementary Fig. 2 and Table 1). Raman spectroscopy of a Se-MoS₂-NF sample displays a new satellite peak at the lower frequency side of the Mo-S vibrational peaks (Supplementary Fig. 3), which can be assigned to the A_{1g} mode of Mo-Se vibrations and thus indicates that Se atoms are embedded into the MoS₂ lattice replacing the S atoms^{27–30}. In the X-ray absorption near-edge structure (XANES) spectra of Se K-edge, the Se-MoS₂-NF shows higher absorption edge than that in the MoSe₂ (Supplementary Fig. 4), denoting a higher oxidation state of the Se atoms in the Se-MoS₂-NF. It may be owing to the lower electronegativity of Se atom than that of S atom so that the Se atoms possess less charges in the Se-MoS₂-NF^{31–33}. These results demonstrate that the Se atoms are covalently doped into the MoS₂ lattice rather than forming a new phase of MoSe₂ or being adsorbed on the surface.

Se-doping combined with fabrication of nanofoam morphology leads to massive generation of edges. High resolution TEM (HRTEM) images show that the Se-MoS₂-NF possesses abundant edges compared with the MoS₂ nanofoam (MoS₂-NF) and few-layer MoS₂ (MoS₂-FL) (Fig. 1c and Supplementary Fig. 5). The clear crystal lattice fringes with an interplanar spacing of 0.66 nm denotes the (002) planes of the 2H phase of MoS₂. X-ray photoelectron spectroscopy (XPS, Fig. 1d) characterizations show that a new doublet peak arises at around 164.9 and 163.7 eV for the Se-MoS₂-NF and MoS₂-NF samples, which can be assigned to the bridging disulfide (S₂²⁻) ligands at the edges^{34–37}. This confirms that establishing a nanofoam morphology is favorable for the formation of more edges. In addition, the S₂²⁻ content of 23% in the Se-MoS₂-NF sample obtained from the peak area is higher than that of 18% in the MoS₂-NF, thus implying that Se-doping further promotes the edge formation. This is also reflected by the extended X-ray absorption fine structure (EXAFS) spectroscopy of the Mo K-edge of these samples, which exhibits reduced Mo-Mo coordination of the Se-MoS₂-NF compared with those of the MoS₂-NF and MoS₂-FL (Fig. 1e), thus indicating a decreased lateral sizes and increased exposure of edges in the Se-MoS₂-NF^{21,26,38}. DFT calculations also show that Se-saturated edge is more stable than S-saturated edge by 0.16 eV per Se replacing a S atom (Fig. 1f). These results demonstrate that Se-doping can promote the formation of abundant edges via stabilizing the edges. The Se-dopants were directly observed in the atomic-resolution high-angle annular dark field scanning transmission electron microscopy (HAADF-STEM) images of the Se-MoS₂-NF, showing the homogeneous dispersion of the confined Se atoms in the basal plane and at the edge of the Se-MoS₂-NF (Fig. 1g). The magnified image in Fig. 1h clearly illustrates that the S atoms are substituted by the Se atoms in accordance with the simulated image of the atomic model.

HER performance of the Se-MoS₂-NF. The Se-MoS₂-NF samples present obviously enhanced HER activity compared with those of bulk MoS₂, MoS₂-FL, and MoS₂-NF (Fig. 2a). The sample with Se doping content of 9.1% (Se(9.1)-MoS₂-NF)

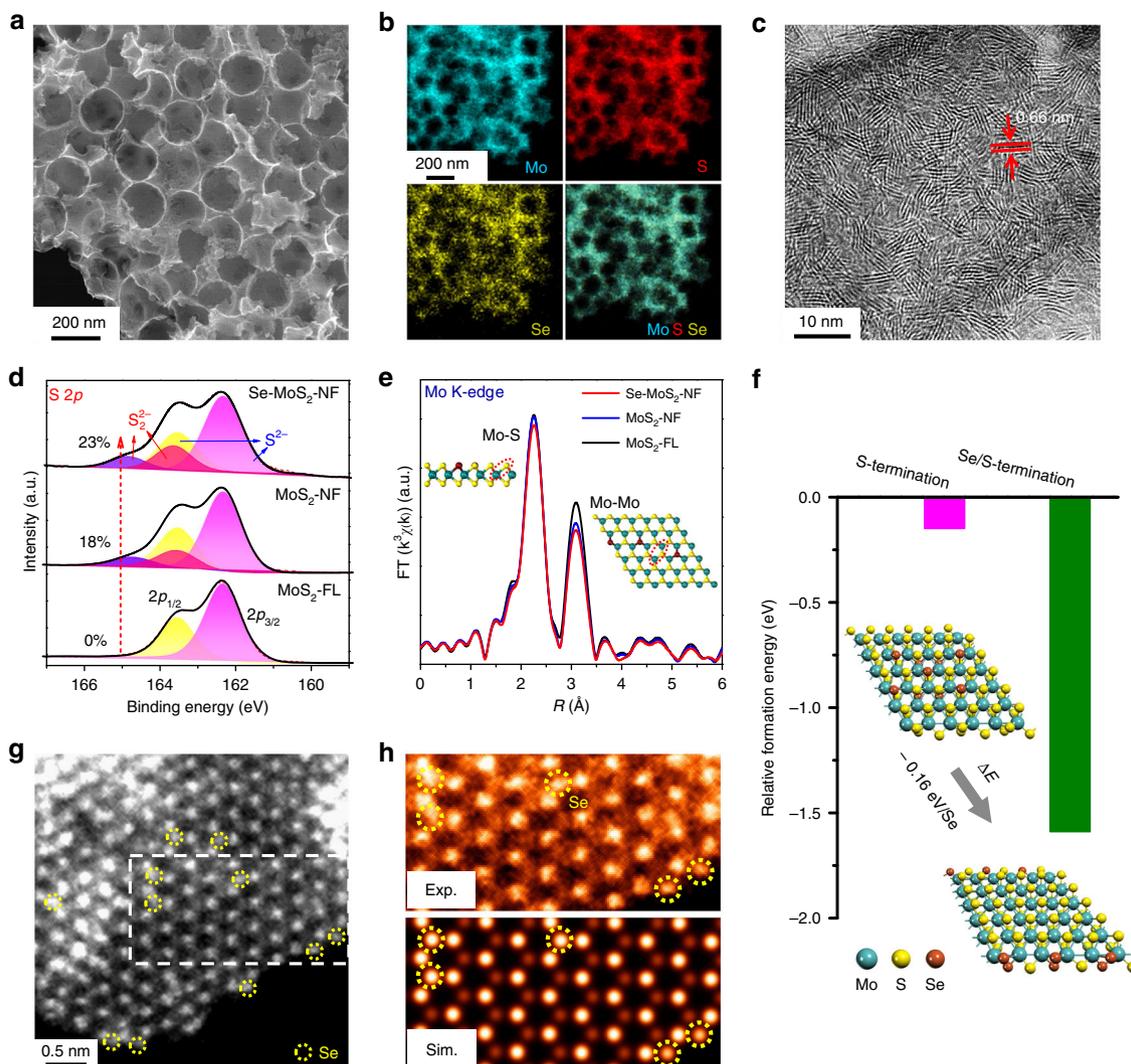


Fig. 1 Structural and electronic properties of the Se-doped MoS₂ nanofoam (Se-MoS₂-NF). SEM image (a), EDX mappings (b), and HRTEM image (c) of the Se-MoS₂-NF. Comparison in the decomposed XPS spectrum of the S 2p (d) and k³-weighted EXAFS spectra of Mo K-edge (e) between the Se-MoS₂-NF, MoS₂-NF and MoS₂-FL catalysts. **f** Comparison in DFT calculated formation energies of the S-terminated and Se-terminated edges. **g** Low pass filtered large-scale atomic-resolution HAADF-STEM image of Se-MoS₂-NF. The yellow dotted circles denote the Se dopants. **h** Enlarged dotted region in **g** and the corresponding simulated image. The Se-MoS₂-NF sample with 9.1% Se-doping content in atomic percentage was employed in these series of characterizations.

exhibits the highest HER activity with overpotentials of 132, 187, and 224 mV at current densities of 10, 50, and 100 mA cm⁻², respectively (Fig. 2b). The decrease in activity as the Se doping content exceeds 9.1% may be resulted from the significantly weakened adsorption of H* at the over-doped Mo-edge Se sites with much positive adsorption free energies (Supplementary Fig. 6). The slopes of linear part of the Tafel plots were used to investigate the HER mechanism over these catalysts. The introduction of Se atoms leads to slight decrease in the Tafel slope of MoS₂ ranging from 75 to 80 mV dec⁻¹, which indicates a Volmer-Heyrovsky mechanism (Supplementary Fig. 7). From the electrochemical impedance spectroscopy (EIS) of the electrode kinetics, the Nyquist plots (Supplementary Fig. 8 and Table 2) reveal a remarkable decrease in the charge-transfer resistance (R_{ct}) from 44.1 Ω of the MoS₂-NF to 11.6 Ω of the Se(9.1)-MoS₂-NF, demonstrating the prominently improved interfacial electron-transfer kinetics over the Se(9.1)-MoS₂-NF. Durability tests show that the HER performance of the Se(9.1)-MoS₂-NF is well maintained for over 30 h at a current density of 10 mA cm⁻², whereas those of the MoS₂-NF and Pt/C (40 wt.%) drop rapidly

(Supplementary Fig. 9). These results demonstrate the positive effect of Se doping in enhancing both the HER activity and stability of the MoS₂.

Properties of Co/Se-codoped MoS₂. Substituting the Mo with Co atoms can effectively activate the adjacent S atoms and improve markedly the HER performance over the MoS₂ basal plane^{10,21,24}. To combine the activating effect of Co-doping with the effect of Se-doping in enriching the edges and boosting the HER activity, we designed a series of Co/Se-codoped MoS₂ nanofoam (Co/Se-MoS₂-NF) to study the synergistic interplay between the dual-elements codoping.

The TEM (Supplementary Fig. 10) and SEM (Supplementary Fig. 11) images show that the Co/Se-MoS₂-NF possesses a similar morphology with that of the Se-MoS₂-NF, containing assemble of small-sized MoS₂ nano-flakes with abundant edges as presented in the HRTEM images (Supplementary Fig. 12). No nanoparticles or large clusters were observed in the TEM and HRTEM images of the Co/Se-MoS₂-NF. The HAADF-STEM and corresponding EDX mappings (Fig. 3a) demonstrate the well-dispersion of Co

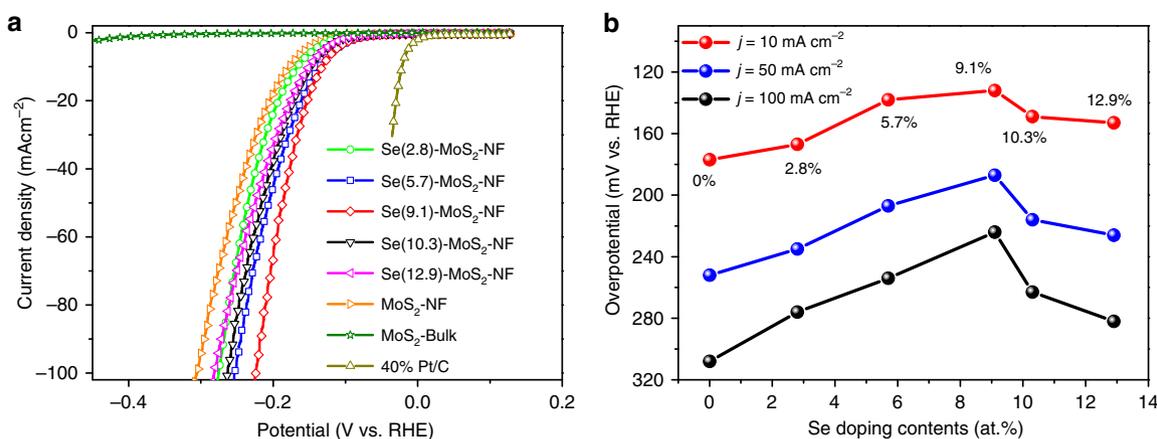


Fig. 2 Electrochemical HER performances of the Se-MoS₂-NF samples. **a** HER polarization curves for the Se-MoS₂-NF with different Se doping contents in comparison with those of bulk MoS₂, MoS₂-NF, and 40% Pt/C. **b** Variation of the overpotentials depending on the Se doping content at current densities of 10, 50, and 100 mA cm⁻², respectively. All catalysts are drop-casted on glassy carbon electrode for HER measurements, which were conducted in a 0.5 M H₂SO₄ electrolyte at 25 °C.

and Se atoms in the structure. This is also confirmed by the XRD analysis showing no other diffraction peaks of Co- or Se-related crystal phases until the Co-doping content reaches 12.2% or more (Supplementary Fig. 13). In addition, the Co K-edge EXAFS characterizations indicate that the Co atoms in the Co/Se-MoS₂-NF are mainly coordinated by S atoms and the Co-S bond in Co/Se-MoS₂-NF is shorter compared with that of the standard CoS sample (Fig. 3g). A Co-Se bond (2.54 Å) is notably longer than a Co-S bond (2.41 Å) according to our DFT calculations but is not observed, which indicates that CoSe phase should not be formed. Besides, the Co valence state in the Co/Se-MoS₂-NF is totally different from those of the Co foil, Co₃O₄, and CoS standard samples and falls in between those of the CoS and Co₃O₄ standard samples (Fig. 3g and Supplementary Fig. 14). Combined with the EXAFS results, it demonstrates that the Co atoms are covalently confined into the MoS₂ lattice rather than being adsorbed on the surface or forming non-MoS₂ hetero-phases such as CoS, CoSe, and Co₃O₄. Atomic-resolution HAADF-STEM imaging of a monolayer Co/Se-MoS₂-NF shows the embedment of Co atoms with a lower contrast at the Mo site and Se atoms being obviously brighter at the S site (Fig. 3b), as confirmed by simulated HAADF-STEM image in Fig. 3c. The Co atom is further identified by using atomic-resolution electron energy loss spectroscopy (EELS), which shows the feature peaks of L₂ and L₃ edges of the Co atom at the site A with lower contrast (Fig. 3c). Distribution of Co and Se atoms being adjacent or separated to each other were observed in the HAADF-STEM image according to the contrast analysis (Fig. 3d), and were confirmed by the simulated HAADF-STEM images and the line intensity profiles (Fig. 3e, f).

Compared with pure Se-doping which can reduce the Mo-Mo coordination and increase the edge formation, Co/Se-codoping can further decrease both the Mo-Mo and Mo-S coordination as shown in the Mo K-edge EXAFS spectra (Figs. 1e, 3h). This is also reflected by the increasing peaks of bridging disulfide S₂²⁻ in the S 2p XPS spectra of the Co/Se-MoS₂-NF compared with those of Se-MoS₂-NF and MoS₂-NF (Supplementary Fig. 15), thus demonstrating the effect of the Co/Se-codoping in further promoting the formation of edges. Co/Se-codoping also leads to significant changes to the electronic structures of the S atoms. A notably negative shift in S 2p binding energies of the XPS was observed after the Co/Se codoping (Supplementary Fig. 16). In addition, the S L-edge XANES characterization shows that the intensity of the white line grows obviously higher after the Co/Se

codoping compared with that of single-element Co- or Se-doped MoS₂ with similar morphology (Fig. 3i). Co and Se atoms possess lower electronegativities than those of Mo and S atoms, so that the S atoms adjacent to the Co and Se dopants are more enriched with electrons, which synergistically modifies the electronic structures and hence the reactivity of S atoms^{39,40}.

HER performance of the Co/Se-MoS₂-NF. Considering the optimum Se-doping content of 9.1% in the Se-MoS₂-NF offering the lowest overpotential, we adopted the same weight ratio of precursors for the Se and MoS₂ and prepared a series of Co/Se-MoS₂-NF samples with varied Co content from 4.0% to 14.6% to investigate the synergistic effect of Co/Se-codoping in tuning the HER activity. The onset overpotential of Co(4.0)/Se-MoS₂-NF is significantly decreased compared with those of Se(9.1)-MoS₂-NF and MoS₂-NF (Fig. 4a). As the content of Co increases, the Co(10.4)/Se-MoS₂-NF sample presents the lowest overpotentials of 104, 157, and 188 mV under the current densities of 10, 50, and 100 mA cm⁻², respectively, which is superior to those of Co-doped MoSe₂ and MoS₂ catalysts (Supplementary Fig. 17 and Table 3). Then the activity decreases as the Co content further grows to 12.2% or larger, which may be due to the formation of other crystal phases at high Co content²¹. The Co(10.4)/Se-MoS₂-NF sample also presents a much lower Tafel slope of 67 mV dec⁻¹ than those of 77 mV dec⁻¹ for the Co(10.4)-MoS₂-NF, 75 mV dec⁻¹ for the Se(9.1)-MoS₂-NF and 83 mV dec⁻¹ for the undoped MoS₂-NF, respectively (Fig. 4b). This clearly demonstrates the superior HER kinetics over the Co/Se-codoped MoS₂ via the Volmer-Heyrovsky mechanism, which differs from that over the Pt/C (40 wt.%) electrocatalyst via the Volmer-Tafel mechanism with a Tafel slope of 29 mV dec⁻¹^{41,42}.

The EIS measurements at a constant potential of 150 mV vs. reversible hydrogen electrode (RHE) indicate that the Co(10.4)/Se-MoS₂-NF possesses the lowest R_{ct} value comparing with all other samples (Fig. 4c and Supplementary Table 2). The Tafel slope derived from the EIS reflects purely the charge-transfer kinetics of the catalytic process and is different from that derived from the linear sweep voltammetry (LSV) curves, where the difference stems from the electronic resistances of the catalysts^{43,44}. The much similar values of the EIS and LSV Tafel slopes for the Co(10.4)/Se-MoS₂-NF sample indicates an improved conductivity of the catalyst (Supplementary Fig. 18). The double-layer capacitance (C_{dl}) extracted from the cyclic

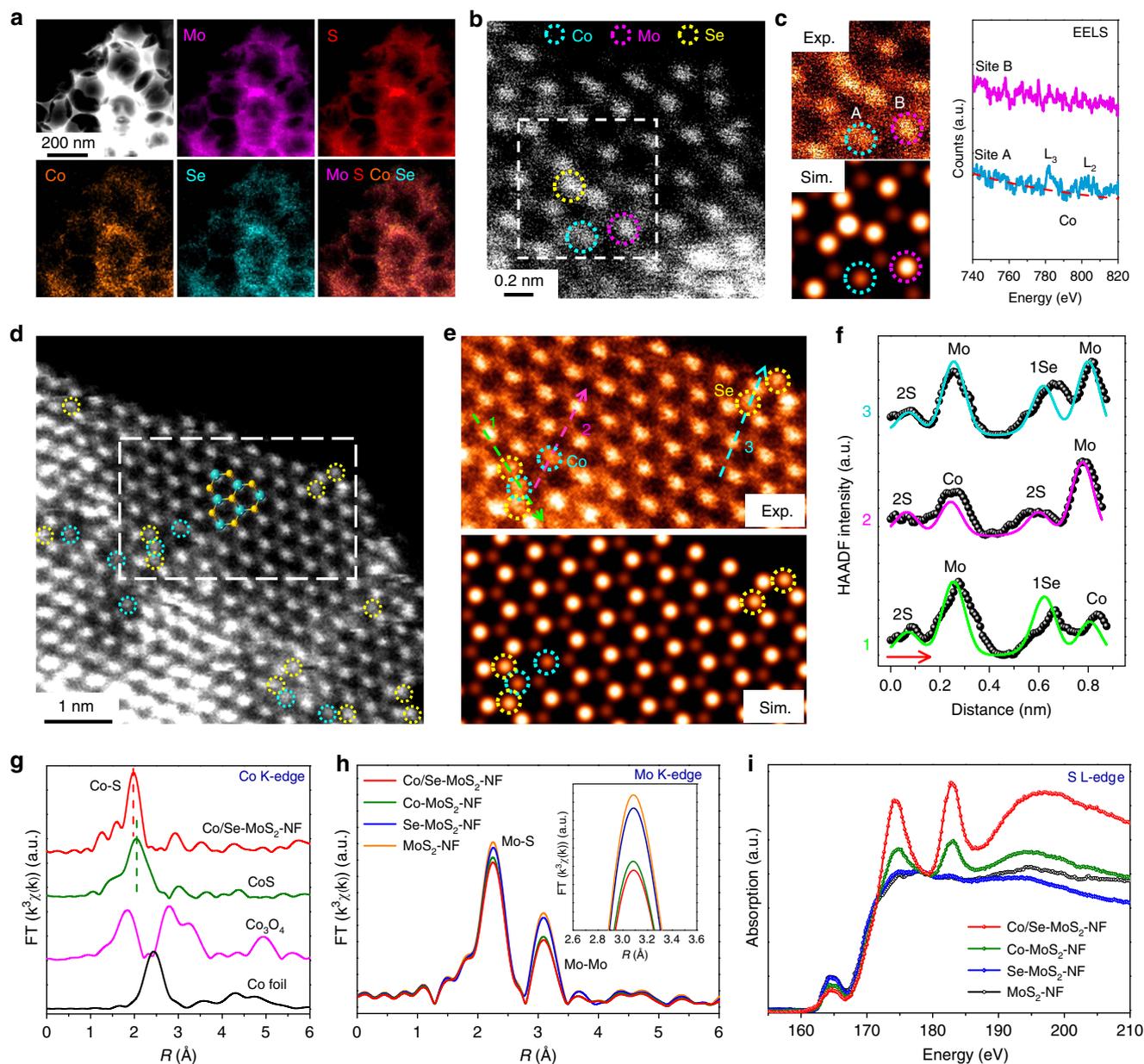


Fig. 3 Structural and electronic properties of the Co/Se-doped MoS₂ nanofoam (Co/Se-MoS₂-NF). **a** HAADF-STEM image and corresponding EDX mappings of the Co/Se-MoS₂-NF. **b** Atomic-resolution HAADF-STEM image of the Co/Se-MoS₂-NF. **c** Enlarged and simulated images of the white dashed line region in **b** and the EELS spectra obtained at the site A and B via line scan. **d** Low pass filtered large area atomic-resolution HAADF-STEM image of the Co/Se-MoS₂-NF. **e** Enlarged and simulated images of the white dashed line in **d**. **f** The measured intensity profile along three lines labeled in **e**. **g** Co K-edge k^3 -weighted EXAFS spectra of Co/Se-MoS₂-NF sample in comparison with CoS, Co₃O₄, and Co foil, respectively. Mo K-edge k^3 -weighted EXAFS spectra (**h**) and S L-edge total electron yield X-ray absorption spectra (**i**) of Co/Se-MoS₂-NF sample compared with Co-MoS₂-NF, Se-MoS₂-NF, and MoS₂-NF. The Co/Se-MoS₂-NF sample with 10.4% Co-doping content in atomic percentage was employed in these series of characterizations.

voltammetry (CV) measurements was used to estimate the electrochemically active surface areas. The Co(10.4)/Se-MoS₂-NF shows a higher C_{dl} of 45.7 mF cm⁻² compared with those of 26.3 mF cm⁻² for the Co(10.4)-MoS₂-NF, 30.7 mF cm⁻² for the Se(9.1)-MoS₂-NF, and 15.8 mF cm⁻² for the MoS₂-NF, respectively (Supplementary Figs. 19 and 20). This demonstrates that the Co(10.4)/Se-MoS₂-NF possesses a higher catalytically active surface area and exposes more active sites⁴².

More interestingly, the Co(10.4)/Se-MoS₂-NF exhibits both superior HER activities and stability at practically high current densities. At 208 mA cm⁻², the overpotential becomes lower than that of the 40 wt.% Pt/C catalyst (Fig. 4d and Supplementary Fig. 21). At a higher large current density of 1000 mA cm⁻², the

overpotential is only 382 mV and significantly lower than that of 671 mV over the Pt/C catalyst. The faradic efficiency for H₂ production in a wide range of working potential is well maintained at ~100%, which indicate that there is almost no other reduction product (Supplementary Fig. 22). To the best of our knowledge, such HER activity of the catalyst also surpasses those of all previously reported heteroatom-doped MoS₂ catalysts (Supplementary Table 3). More importantly, the activity can be well maintained for more than 360 h without decay at the large current density of 1000 mA cm⁻², while the activity of Pt/C catalyst will drop quickly within 50 h at the same test condition (Fig. 4e). The XRD (Supplementary Fig. 23) and SEM image (Supplementary Fig. 24) of the Co(10.4)/Se-MoS₂-NF show no

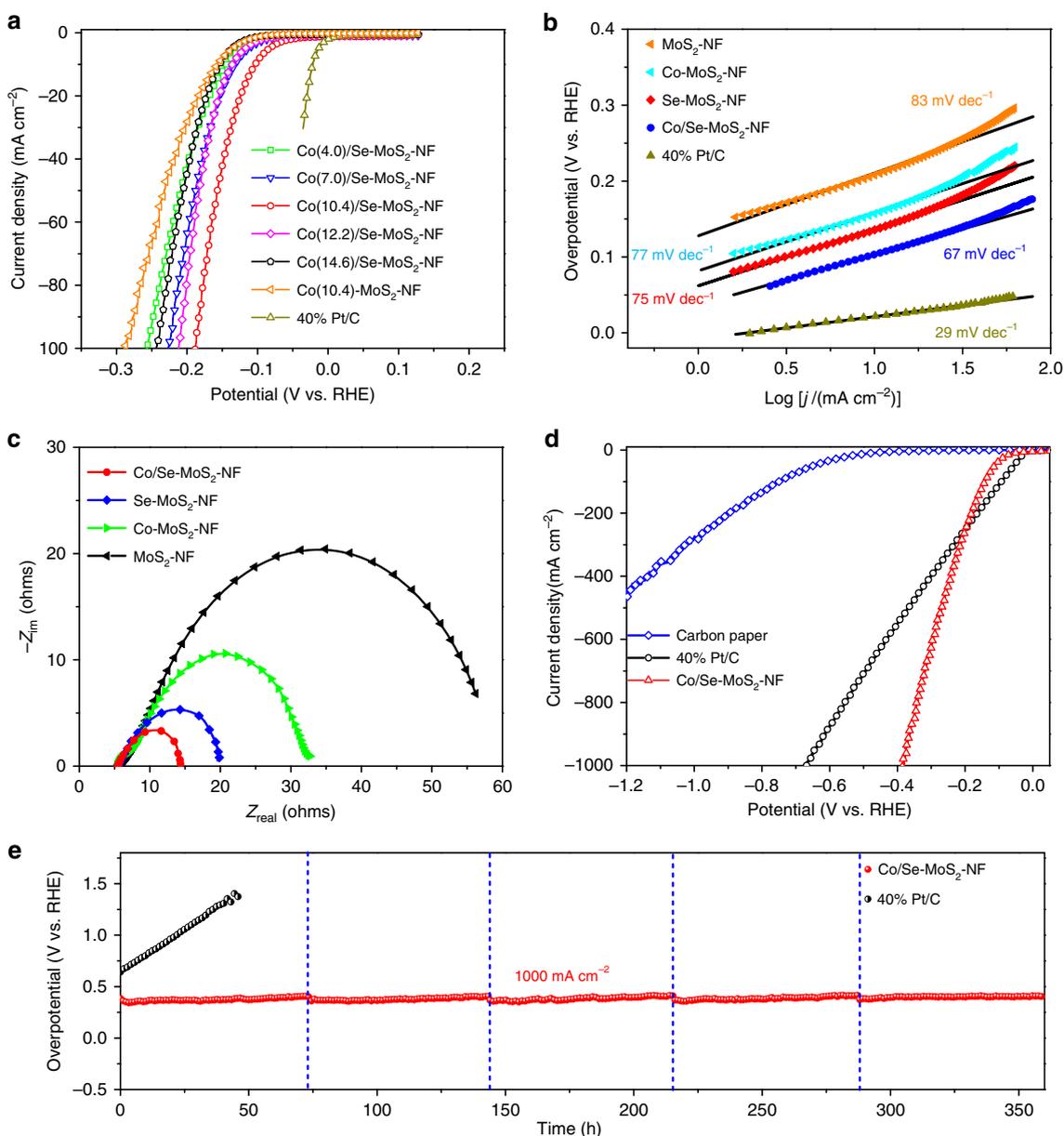


Fig. 4 Synergistic effect of Co/Se-codoping on the HER performance of MoS₂ nanofoam. **a** HER polarization curves for Co/Se-MoS₂-NF with different Co-doping contents in comparison with Co-MoS₂-NF and 40% Pt/C. All samples are drop-casted on glassy carbon electrode for measurements. The Tafel plots (**b**) and the electrochemical impedance spectroscopy (EIS) Nyquist plots (**c**) of Co/Se-MoS₂-NF sample compared with Co-MoS₂-NF, Se-MoS₂-NF, MoS₂-NF, and commercial 40% Pt/C. **d** HER polarization curves for Co/Se-MoS₂-NF under large current densities in comparison with 40% Pt/C and carbon paper. Co/Se-MoS₂-NF and 40% Pt/C are drop-casted on carbon paper for measurements. **e** The chronopotentiometric measurements of long-term stability for Co/Se-MoS₂-NF and 40% Pt/C at 1000 mA cm⁻². The vertical dotted lines in **e** mark the time span (every 72 h) of replacing the electrolyte in the 360 h chronoamperometry test. All the HER measurements were conducted in a 0.5 M H₂SO₄ electrolyte at 25 °C.

change in the crystallographic and morphology structure after the 360 h of durability test. The inductively coupled plasma optical emission spectrometry (ICP-OES) measurement shows that the content of Pt in the sample after the durability test is below the detection limit, thus indicating that Pt is not likely deposited on the Co(10.4)/Se-MoS₂-NF sample. These results demonstrate that the dual-element Co/Se-codoping can synergistically enhance both the HER activity and stability of MoS₂, even under industrial-level large current densities.

Understanding the synergistic effect of Co/Se-codoping. DFT calculations were performed to study the synergistic effect of dual-element Co/Se-codoping in improving the HER performance of

MoS₂ (Fig. 5). Co-doping significantly enhances the adsorption of hydrogen on the basal plane S site with adsorption free energy of H* (ΔG_{H^*}) decreasing from 2.01 to -0.32 eV (Fig. 5a). However, it also inevitably strengthens the adsorption of hydrogen on the Mo-edge S site with ΔG_{H^*} of -0.47 eV, which leads to over-binding of H* and is unfavorable for the HER at the Co-doped Mo-edge sites. Introducing Se into the Co-doped MoS₂ compensates the over-strong adsorption of H* on both the Co-doped in-plane and edge sites and thereby moderately weakens their adsorption activity for H*. At the Co/Se-codoped in-plane, Mo-edge, and S-edge sites, the ΔG_{H^*} are increased to -0.21, -0.14, and 0.10 eV, respectively, from those of -0.32, -0.46, and -0.10 eV at the corresponding Co-doped sites (Fig. 5a, b). This indicates an improved HER activity at the Co/Se-codoped in-plane

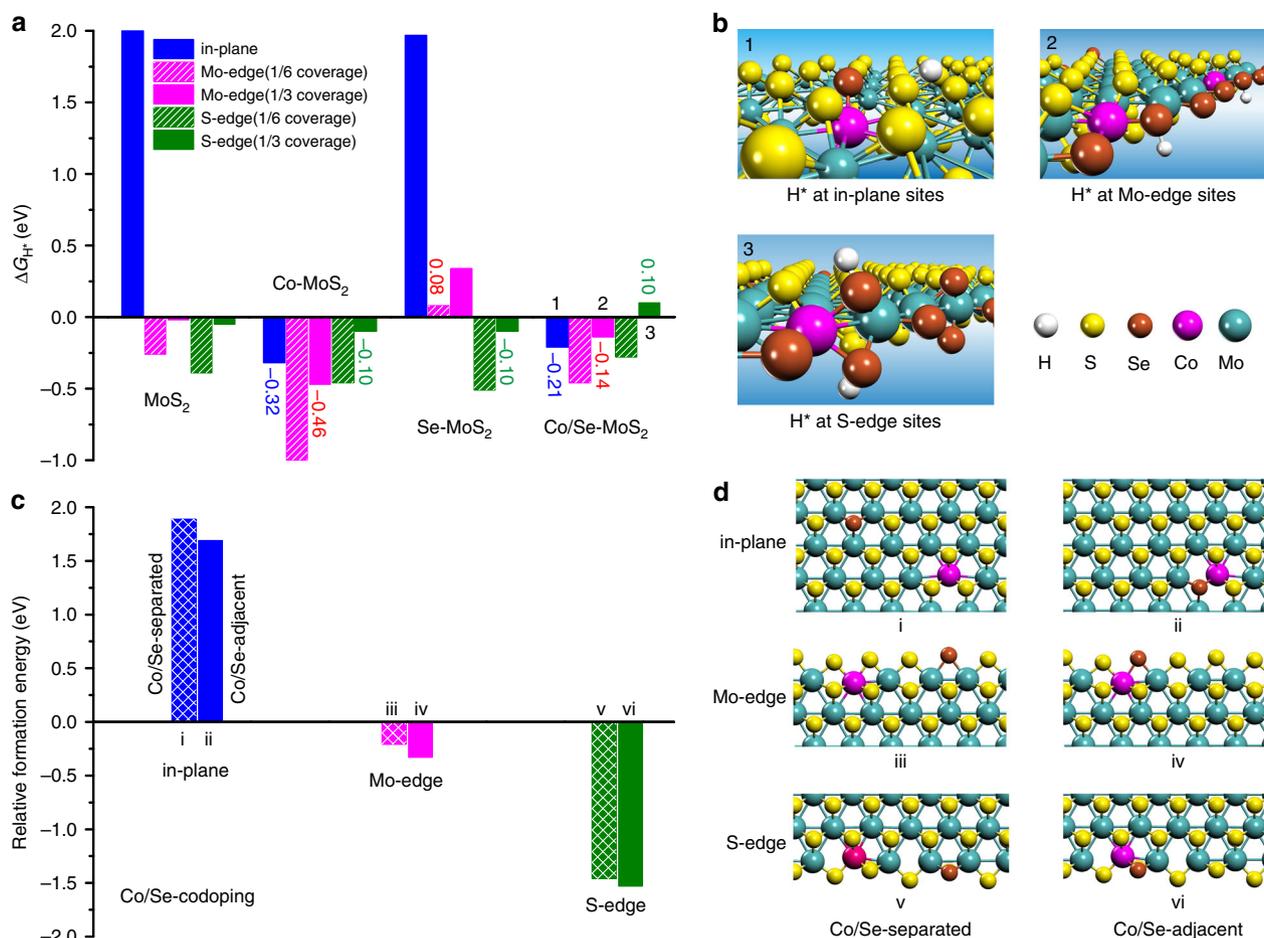


Fig. 5 Theoretical studies of the Co-doping, Se-doping, and Co/Se-codoping effect in tuning the HER performance of MoS₂. **a** The adsorption free energies of H* (ΔG_{H^*}) at the in-plane, Mo-edge, and S-edge sites of pure, Co-doped, Se-doped, and Co/Se-codoped MoS₂, denoted as MoS₂, Co-MoS₂, Se-MoS₂, Co/Se-MoS₂, respectively. The hydrogen coverage at the Mo-edge is defined according to the terminating S or Se monomer sites, while that at the S-edge is defined according to the terminating S or Se dimer sites which are the active sites for the HER. **b** Adsorption structures of H* at the in-plane, Mo-edge and S-edge sites of Co/Se-MoS₂. **c** Relative formation energies of the Co/Se-codoped structures with different doping configurations of Co and Se being separated (Co/Se-separated, grid bars) or adjacent (Co/Se-adjacent, solid bars) to each other. **d** Structures of Co/Se-codoped basal plane, Mo-edge, and S-edge with Co/Se-separated and Co/Se-adjacent configurations (Supplementary Fig. 26).

and Mo-edge sites with reduced overpotentials. In comparison, pure Se-doping only notably weakens the H* adsorption at the Mo-edge but still lead to a good HER activity at the Mo-edge and S-edge sites with ΔG_{H^*} of 0.08 eV and -0.10 eV, respectively (Fig. 5a). Thus, the Co/Se-codoping brings a compromise between the enhancement effect of Co-doping and weakening effect of Se-doping on the hydrogen adsorption at the codoped sites, leading to more favorable hydrogen adsorption activities for the HER. In addition, Se-doping also prominently stabilizes the Co-doped basal plane and edges by forming Co-Se bonds (Fig. 5c, d), where the formation energies of Co/Se-codoped basal plane, Mo-edge, and S-edge can be reduced by 0.21, 0.12, and 0.07 eV per Co-Se bond formed, respectively, compared with those with Co and Se separated from each other. These results demonstrate a synergy between the activating effect of inner-layer Co-doping and the stabilization effect of surface Se-doping in the Co/Se-codoped MoS₂, which significantly improves the HER performance by simultaneously enriching active sites and optimizing their hydrogen adsorption activities.

Discussion

In summary, via co-confining Se in the surface and Co in the inner layer of the MoS₂ combined with fabricating a nanofoam

architecture, we achieve both ultrahigh activity and stability for the acidic HER. The activating effect of the inner-layer Co-doping combined with the stabilizing effect of the surface Se-doping enables the formation of abundant active sites in both the basal plane and the edges with optimized hydrogen adsorption activity at the codoped sites. The Co/Se-MoS₂-NF presents the highest HER activity among the previously reported heteroatom-doped MoS₂ catalysts under industrial-level large current densities. At a high current density of 1000 mA cm⁻², the Co/Se-MoS₂-NF exhibits a much lower overpotential of 382 mV than that of 671 mV over the 40 wt.% Pt/C catalyst, and the high activity can be maintained with long-term stability of more than 360 h without decay. Such a strategy of engineering the HER activity of MoS₂ via co-confining multielements offers a promising and feasible route of developing high-performance and low-cost MoS₂ catalysts for large-scale production of clean energy hydrogen.

Methods

Synthesis of SiO₂ template. The uniform SiO₂ nanosphere template was prepared by adding 9.6 g tetraethyl orthosilicate into 200 mL of ethanol solution under stirring for 10 min, followed by adding 8 mL of deionized water and 8 mL of ammonia. Then, the mixture was continuously stirred under room temperature until all the solvent was evaporated and was dried at 60 °C to obtain SiO₂ template.

Synthesis of Se-doped MoS₂ nanofoam (Se-MoS₂-NF). The Se-MoS₂-NF was synthesized through a one-pot chemical synthesis method. First, 0.4 g (NH₄)₆Mo₇O₂₄·4H₂O, specified amount of selenium powder and 1.6 g SiO₂ nanosphere powder were dispersed in 25 mL deionized water, followed by continuously stirring under room temperature until all the solvent was evaporated. Then, the gained solid was dried under 80 °C for 12 h. After that, the gained powder and 10 mL CS₂ were both transferred into 40 mL stainless steel autoclave in the atmosphere of Ar and maintained at 400 °C for 4 h. The final product was treated with HF (aq.) under room temperature overnight, followed by washing several times with water and absolute ethanol and drying at 80 °C. For comparison, the MoS₂ nanofoam (MoS₂-NF) was synthesized via the same method as Se-MoS₂-NF without adding selenium powder. The MoS₂-FL was synthesized by using 0.9 g (NH₄)₆Mo₇O₂₄·4H₂O dissolved in 20 mL deionized water and 10 mL CS₂, and undergoing the same process as the synthesis of the MoS₂-NF without using SiO₂ template. All the doping contents in final samples were measured by ICP-OES. We have synthesized a series of Se-MoS₂-NF samples with Se doping contents of 2.8, 5.7, 9.1, 10.3, and 12.9% in atomic percentage by the above methods.

Synthesis of Co/Se-codoped MoS₂ nanofoam (Co/Se-MoS₂-NF). For the synthesis of Co/Se-MoS₂-NF samples, typically, 0.4 g (NH₄)₆Mo₇O₂₄·4H₂O, 1.6 g SiO₂ nanospheres powder, 30 mL deionized water, specified amount of Co (NO₃)₂·6H₂O and selenium powder were mixed homogeneously, followed by continuously stirring under room temperature until the solvent was evaporated and drying at 80 °C. Then, the gained powder and 10 mL CS₂ were transferred into 40 mL stainless steel autoclave and followed by the same process as the synthesis of the Se-MoS₂-NF. For comparison, the Co-doped MoS₂ nanofoam (Co-MoS₂-NF) was synthesized by the same chemical method as the Co/Se-MoS₂-NF without adding selenium powder. All the doping contents in final samples were measured by ICP-OES.

Materials characterization. TEM, HAADF-STEM, HRTEM, and EDX mappings measurements were carried out on the Phillips Analytical FEI Tecnai20 electron microscope operated at an accelerating voltage of 200 kV. SEM measurements were carried out on Hitachi S5500 operated at 30 kV. Atomic-resolution HAADF-STEM images and EELS mapping were performed on a spherical aberration-corrected Nion U-HERMES100. The experiments were conducted at 60 kV accelerating voltage to reduce radiation damage and also with relatively low electron dose. EELS spectra processing was performed using the open-source software HYPERSPY (formerly EELS LAB) to remove X-ray strikes⁴⁵. Atomic-resolution HAADF-STEM image simulation was performed in the QSTEM software. XRD patterns were recorded on Rigaku Ultima IV diffractometer. Cu Kα radiation ($\lambda = 0.15406$ nm) at 40 kV and 30 mA was used as the X-ray source. XPS measurements were carried out on a Thermo ESCALAB 250Xi spectroscopy using Al Kα x-rays as the excitation source. Raman measurements were performed on a home-modified Invia confocal Raman microscope (Renishaw, UK) with Leica DM2500 microscopes, which was operated with a 633 nm excitation laser at a power of 26.7 μW. XANES and EXAFS were measured at the BL14W1 beamline of the Shanghai Synchrotron Radiation Facility (SSRF) recorded in a transmission mode. The S L-edge XANES spectra were collected in total electron yield mode at the beamline BL10 in Hefei National Synchrotron Radiation Laboratory (NSRL). ICP-OES test was conducted on Shimadzu ICPS-8100.

Electrochemical measurements. Electrochemical measurements were performed in a three-electrode electrochemical cell equipped with a gas flow controlling system on electrochemical workstations (CHI 760E and CHI 680 C). We employed the glassy carbon rotating disk electrode with a diameter of 5 mm covered by a thin catalyst film as working electrode to perform HER measurements at low-level current densities of 10, 50, and 100 mA cm⁻². Typically, 4 mg catalyst and 2 mg carbon black (Vulcan XC-72) was suspended in 1 mL ethanol with 20 μL Nafion solution (5 wt.%, Du Pont) to form a homogeneous ink assisted by ultrasound. Then 25 μL of the ink was spread onto the surface of glassy carbon by a micropipette and dried under room temperature. The final loading amount of catalysts on work electrode is 0.5 mg cm⁻². Graphite rod and Ag/AgCl (saturated KCl-filled) were used as counter electrode and reference electrode, respectively. HER tests were conducted in an Ar-saturated 0.5 M H₂SO₄ electrolyte at 25 °C under 1600 rpm. The LSV polarization curve was measured at 5 mV/s in the potential range of -0.1 to 1.0 V (vs. Ag/AgCl). Before measurements, the sample were repeatedly swept from -0.4 to 0.3 V (vs. Ag/AgCl) in the electrolyte until steady voltammogram curve was obtained. The long-term stability tests were performed by using chronopotentiometric measurement on the carbon fiber paper (CFP) with a catalyst loading of 0.5 mg cm⁻². EIS was evaluated at -0.15 V (vs. RHE) between 100 kHz and 0.1 Hz with a 5 mV AC potential perturbation. Tafel analysis from EIS measurements was performed at various working potential from 100 kHz to 0.1 Hz with a 15 mV AC potential perturbation. To determine the double-layer capacitance (C_{dl}), the CV were performed at various scan rates (20, 40, 60 mV s⁻¹, etc.) between 0.1 V and 0.2 V (vs. RHE).

We loaded the catalyst on carbon paper to measure the HER activity and stability at large current densities up to 1000 mA cm⁻². The Tafel slopes of the Pt/C catalyst loaded on carbon paper and glassy carbon electrode are almost the same,

which indicating that their HER kinetics are identical (Supplementary Fig. 25). The Fig. 4d, e show polarization curves and chronopotentiometric measurements under large current densities, respectively, which were obtained by using a three-electrode H-type electrochemical cell equipped with a gas flow controlling system. For the preparation of the working electrode, 4 mg catalyst and 2 mg carbon black (Vulcan XC-72) were suspended in 0.5 mL ethanol with 25 μL Nafion solution (5 wt.%, Du Pont) to form a homogeneous ink assisted by ultrasound. Then, the ink was dropped on CFPs (1 cm × 1 cm) and dried under room temperature. For the characterization of HER polarization curves, an Ag/AgCl (saturated KCl-filled) electrode and graphite rod were used as reference and counter electrode, respectively. For the characterization of chronopotentiometric test, Pt mesh was used as the counter electrode owing to the instability of graphite rod under large current densities. In order to avoid effectively the possible Pt contamination during measurements, cathode and anode compartments were separated with Nafion 117 membrane in H-type electrochemical cell. To weaken the influence of reactant concentration for catalyst activity, the electrolyte was replaced every 72 h.

All potentials were 85% iR corrected and referenced to the RHE by the equation $E(\text{RHE}) = E(\text{Ag/AgCl, saturated KCl-filled}) - iR + 0.227$. The iR correction was performed according to a previously reported method^{46,47}. The R for iR correction was determined by impedance measurements.

Calculation of the faradaic efficiency. The controlled potential electrolysis was carried out to determine the faradaic efficiency of H₂ using a three-electrode H-type electrochemical cell. During the controlled potential electrolysis, the electrolyte in both anodic and cathodic compartments were stirred slightly to enhance the diffusion of reactant. Quantification of the H₂ were performed using the gas chromatograph (Shimadzu GC 2014) with a thermal conductivity detector. The faradaic efficiency of H₂ are calculated from GC chromatogram peak areas at a given potential as follow:

$$i_{\text{H}_2} = v_{\text{H}_2} \times V \times \frac{2Fp_0}{RT}, \quad (1)$$

$$FE_{\text{H}_2} = \frac{i_{\text{H}_2}}{i_{\text{total}}} \times 100\%, \quad (2)$$

where i_{H_2} is partial current density for H₂, i_{total} is total current density, v_{H_2} is volume concentration of H₂ based on the calibration of the GC, V is gas volume flow rate, and F is the Faraday's constant 96,485 C mol⁻¹. $p_0 = 101.325$ kPa, $R = 8.314$ J mol⁻¹ K⁻¹, and $T = 298.15$ K.

DFT calculations. DFT calculations were performed using the Vienna Ab-initio Simulation Package⁴⁸⁻⁵¹. The projector augmented-wave pseudopotential method with Perdew-Burke-Ernzerhof exchange-correlation functional and a plane-wave cutoff energy of 400 eV was adopted⁵²⁻⁵⁵. The standard version of PAW pseudopotentials without using semi-core states were used for all elements. Van der Waals correction was calculated using the Zero-damping DFT-D3 method of Grimme^{56,57}. A tri-layer model of MoS₂ was built in a 6 × 6 supercell to simulate the in-plane doping of Co and Se. A nanoribbon model of MoS₂ with six repeated units along the belt direction was built to simulated the Mo-edge and S-edge. The Mo edge was saturated with S monomers and the S edge was saturated with a combination of S monomers and dimers^{58,59}. The vacuum thicknesses were set larger than 15 Å between the layers or the ribbons. A Monkhorst-Pack k-point sampling of 1 × 1 × 1 was selected for all models⁶⁰. In structural optimizations, the residual forces between atoms were converged below 0.02 eV/Å. Dipole correction was applied to decouple the interactions between the slabs or nanoribbons.

The free energy of (H⁺ + e⁻) was calculated as that of ½ H₂. The free energy of gas molecules was calculated as $E_{\text{total}} + ZPE + \int_0^T C_p dT - TS$, where E_{total} is DFT calculated total energy, ZPE is the zero-point energy, T is temperature, $\int_0^T C_p dT$ is the integrated heat capacity from 0 K to T , and S is the entropy. For the ZPE + $\int_0^T C_p dT - TS$ parts, we adopt the experimental data from the NIST database (Supplementary Table 4). The free energies of adsorbed species were calculated as $E_{\text{total}} + ZPE + \int_0^T C_{v(\text{vib})} dT - TS_{\text{vib}}$, where the $\int_0^T C_{v(\text{vib})} dT$ and S_{vib} are the integrated heat capacity from 0 K to T and the entropy parts from non-imaginary vibrations based on harmonic oscillation approximation.

The formation energies were calculated by using the free energies of H₂, H₂S, pristine MoS₂, and MoS₂, and Co bulk as the references. The chemical potential of each element is calculated as:

$$\mu(\text{H}) = 1/2G(\text{H}_2), \quad (3)$$

$$\mu(\text{S}) = G(\text{H}_2\text{S}) - G(\text{H}_2), \quad (4)$$

$$\mu(\text{Mo}) = G(\text{MoS}_2) - 2\mu(\text{S}), \quad (5)$$

$$\mu(\text{Se}) = 1/2[G(\text{MoSe}_2) - \mu(\text{Mo})], \quad (6)$$

$$\mu(\text{Co}) = 1/2G(\text{Co} - \text{bulk})^*. \quad (7)$$

*The supercell of hcp Co bulk contains two Co atoms.

Then the formation energy (E_{form}) of a structure $\text{Mo}_x\text{Co}_y\text{S}_z\text{Se}_m$ is given by:

$$E_{\text{form}} = E_{\text{total}}(\text{Mo}_x\text{Co}_y\text{S}_z\text{Se}_m) - x\mu(\text{Mo}) - y\mu(\text{Co}) - z\mu(\text{S}) - m\mu(\text{Se}). \quad (8)$$

The relative formation energy of a Co-doped, Se-doped, or Co/Se-codoped MoS_2 edge is calculated by using its formation energy minus that of the corresponding undoped pure MoS_2 edge. This is to decouple the doping effect on the formation energy from that of creating undoped pure MoS_2 edges.

Data availability

The data that support the findings of this study are available from the corresponding authors on request.

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References

- Dresselhaus, M. S. & Thomas, I. L. Alternative energy technologies. *Nature* **414**, 332–337 (2001).
- Turner, J. A. Sustainable hydrogen production. *Science* **305**, 972 (2004).
- Stamenkovic, V. R., Strmcnik, D., Lopes, P. P. & Markovic, N. M. Energy and fuels from electrochemical interfaces. *Nat. Mater.* **16**, 57–69 (2017).
- Zou, X. & Zhang, Y. Noble metal-free hydrogen evolution catalysts for water splitting. *Chem. Soc. Rev.* **44**, 5148–5180 (2015).
- Chen, Z. et al. A review on non-precious metal electrocatalysts for PEM fuel cells. *Energy Environ. Sci.* **4**, 3167–3192 (2011).
- Norskov, J. K. & Christensen, C. H. Toward efficient hydrogen production at surfaces. *Science* **312**, 1322 (2006).
- Hinnemann, B. et al. Biomimetic hydrogen evolution: MoS_2 nanoparticles as catalyst for hydrogen evolution. *J. Am. Chem. Soc.* **127**, 5308–5309 (2005).
- Jaramillo, T. F. et al. Identification of active edge sites for electrochemical H_2 evolution from MoS_2 nanocatalysts. *Science* **317**, 100 (2007).
- Deng, J. et al. Highly active and durable non-precious-metal catalysts encapsulated in carbon nanotubes for hydrogen evolution reaction. *Energy Environ. Sci.* **7**, 1919–1923 (2014).
- Deng, J. et al. Triggering the electrocatalytic hydrogen evolution activity of the inert two-dimensional MoS_2 surface via single-atom metal doping. *Energy Environ. Sci.* **8**, 1594–1601 (2015).
- Deng, D. et al. Catalysis with two-dimensional materials and their heterostructures. *Nat. Nanotechnol.* **11**, 218–230 (2016).
- Mao, J., Wang, Y., Zheng, Z. & Deng, D. The rise of two-dimensional MoS_2 for catalysis. *Front. Phys.* **13**, 138118 (2018).
- Wang, Y. et al. Catalysis with two-dimensional materials confining single atoms: concept, design, and applications. *Chem. Rev.* **119**, 1806–1854 (2019).
- Qi, K. et al. Single-atom cobalt array bound to distorted 1T MoS_2 with ensemble effect for hydrogen evolution catalysis. *Nat. Commun.* **10**, 5231 (2019).
- Kibsgaard, J., Chen, Z., Reinecke, B. N. & Jaramillo, T. F. Engineering the surface structure of MoS_2 to preferentially expose active edge sites for electrocatalysis. *Nat. Mater.* **11**, 963–969 (2012).
- Xie, J. et al. Defect-rich MoS_2 ultrathin nanosheets with additional active edge sites for enhanced electrocatalytic hydrogen evolution. *Adv. Mater.* **25**, 5807–5813 (2013).
- Deng, J. et al. High-performance hydrogen evolution electrocatalysis by layer-controlled MoS_2 nanosheets. *Rsc. Adv.* **4**, 34733–34738 (2014).
- Li, H. et al. Activating and optimizing MoS_2 basal planes for hydrogen evolution through the formation of strained sulphur vacancies. *Nat. Mater.* **15**, 364–364 (2016).
- Albu-Yaron, A. et al. MoS_2 hybrid nanostructures: from octahedral to quasi-spherical shells within individual nanoparticles. *Angew. Chem. Int. Ed.* **50**, 1810–1814 (2011).
- Yun, Q. et al. Three-dimensional architectures constructed from transition-metal dichalcogenide nanomaterials for electrochemical energy storage and conversion. *Angew. Chem. Int. Ed.* **57**, 626–646 (2018).
- Deng, J. et al. Multiscale structural and electronic control of molybdenum disulfide foam for highly efficient hydrogen production. *Nat. Commun.* **8**, 14430 (2017).
- Shi, Y. et al. Energy level engineering of MoS_2 by transition-metal doping for accelerating hydrogen evolution reaction. *J. Am. Chem. Soc.* **139**, 15479–15485 (2017).
- Xiong, Q. et al. Cobalt covalent doping in MoS_2 to induce bifunctionality of overall water splitting. *Adv. Mater.* **30**, 1801450 (2018).
- Lau, T. H. M. et al. Transition metal atom doping of the basal plane of MoS_2 monolayer nanosheets for electrochemical hydrogen evolution. *Chem. Sci.* **9**, 4769–4776 (2018).
- Wu, W. et al. Activation of MoS_2 basal planes for hydrogen evolution by Zinc. *Angew. Chem. Int. Ed.* **58**, 2029–2033 (2019).
- Meng, X. et al. Three-dimensionally hierarchical MoS_2 /graphene architecture for high-performance hydrogen evolution reaction. *Nano Energy* **61**, 611–616 (2019).
- Li, H. et al. Growth of alloy $\text{MoS}_2\text{Se}_{2(1-x)}$ nanosheets with fully tunable chemical compositions and optical properties. *J. Am. Chem. Soc.* **136**, 3756–3759 (2014).
- Mann, J. et al. 2-dimensional transition metal dchalcogenides with tunable direct band gaps: $\text{MoS}_{2(1-x)}\text{Se}_{2x}$ monolayers. *Adv. Mater.* **26**, 1399–1404 (2014).
- Zhang, X. et al. Phonon and raman scattering of two-dimensional transition metal dichalcogenides from monolayer, multilayer to bulk material. *Chem. Soc. Rev.* **44**, 2757–2785 (2015).
- Konkena, B. et al. MoSSe @reduced graphene oxide nanocomposite heterostructures as efficient and stable electrocatalysts for the hydrogen evolution reaction. *Nano Energy* **29**, 46–53 (2016).
- Allred, A. L. Electronegativity values from thermochemical data. *J. Inorg. Nucl. Chem.* **17**, 215–221 (1961).
- Li, K. & Xue, D. Estimation of electronegativity values of elements in different valence states. *J. Phys. Chem. A* **110**, 11332–11337 (2006).
- Diener, A., Neumann, T., Kramar, U. & Schild, D. Structure of selenium incorporated in pyrite and mackinawite as determined by XAFS analyses. *J. Contam. Hydrol.* **133**, 30–39 (2012).
- Gao, M.-R. et al. An efficient molybdenum disulfide/cobalt diselenide hybrid catalyst for electrochemical hydrogen generation. *Nat. Commun.* **6**, 5982 (2015).
- Hu, J. et al. Engineering stepped edge surface structures of MoS_2 sheet stacks to accelerate the hydrogen evolution reaction. *Energy Environ. Sci.* **10**, 593–603 (2017).
- Xu, Q. et al. Unsaturated sulfur edge engineering of strongly coupled MoS_2 nanosheet-carbon macroporous hybrid catalyst for enhanced hydrogen generation. *Adv. Energy Mater.* **9**, 1802553 (2019).
- Vrubel, H., Merki, D. & Hu, X. Hydrogen evolution catalyzed by MoS_3 and MoS_2 particles. *Energy Environ. Sci.* **5**, 6136–6144 (2012).
- Ravel, B. & Newville, M. Data analysis for X-ray absorption spectroscopy using IFEFFIT. *J. Synchrotron Radiat.* **12**, 537–541 (2005).
- Costello, M. T. & Kasrai, M. Study of surface films of overbased sulfonates and sulfurized olefins by X-ray absorption near edge structure (XANES) spectroscopy. *Tribol. Lett.* **24**, 163–169 (2006).
- Li, D. et al. Polarized X-ray absorption spectra and electronic structure of molybdenite (2H-MoS_2). *Phys. Chem. Miner.* **22**, 123–128 (1995).
- Conway, B. E. & Tilak, B. V. Interfacial processes involving electrocatalytic evolution and oxidation of H_2 , and the role of chemisorbed H. *Electrochim. Acta* **47**, 3571–3594 (2002).
- Merki, D. et al. Fe, Co, and Ni ions promote the catalytic activity of amorphous molybdenum sulfide films for hydrogen evolution. *Chem. Sci.* **3**, 2515–2525 (2012).
- Vrubel, H., Moehl, T., Grätzel, M. & Hu, X. Revealing and accelerating slow electron transport in amorphous molybdenum sulphide particles for hydrogen evolution reaction. *Chem. Commun.* **49**, 8985–8987 (2013).
- Meiron, O. E. et al. Improved catalytic activity of $\text{Mo}_{1-x}\text{W}_x\text{Se}_2$ alloy nanoflowers promotes efficient hydrogen evolution reaction in both acidic and alkaline aqueous solutions. *Nanoscale* **9**, 13998–14005 (2017).
- De la Peña, F. et al. Mapping titanium and tin oxide phases using EELS: an application of independent component analysis. *Ultramicroscopy* **111**, 169–176 (2011).
- Lukowski, M. A. et al. Enhanced hydrogen evolution catalysis from chemically exfoliated metallic MoS_2 nanosheets. *J. Am. Chem. Soc.* **135**, 10274–10277 (2013).
- Sheng, W., Gasteiger, H. A. & Shao-Horn, Y. Hydrogen oxidation and evolution reaction kinetics on platinum: acid vs alkaline electrolytes. *J. Electrochem. Soc.* **157**, B1529–B1536 (2010).
- Kresse, G. & Hafner, J. Ab initio molecular dynamics for liquid metals. *Phys. Rev. B* **47**, 558–561 (1993).
- Kresse, G. & Furthmüller, J. Efficient iterative schemes for ab initio total-energy calculations using a plane-wave basis set. *Phys. Rev. B* **54**, 11169–11186 (1996).
- Kresse, G. & Hafner, J. Ab initio molecular-dynamics simulation of the liquid-metal-amorphous-semiconductor transition in germanium. *Phys. Rev. B* **49**, 14251–14269 (1994).
- Kresse, G. & Furthmüller, J. Efficiency of ab-initio total energy calculations for metals and semiconductors using a plane-wave basis set. *Comp. Mater. Sci.* **6**, 15–50 (1996).
- Kresse, G. & Joubert, D. From ultrasoft pseudopotentials to the projector augmented-wave method. *Phys. Rev. B* **59**, 1758–1775 (1999).
- Perdew, J. P., Burke, K. & Ernzerhof, M. Generalized gradient approximation made simple. *Phys. Rev. Lett.* **78**, 1396–1396 (1997).

54. Blöchl, P. E. Projector augmented-wave method. *Phys. Rev. B* **50**, 17953–17979 (1994).
55. Perdew, J. P., Burke, K. & Ernzerhof, M. Generalized gradient approximation made simple. *Phys. Rev. Lett.* **77**, 3865–3868 (1996).
56. Grimme, S., Antony, J., Ehrlich, S. & Krieg, H. A consistent and accurate ab initio parametrization of density functional dispersion correction (DFT-D) for the 94 elements H–Pu. *J. Chem. Phys.* **132**, 154104 (2010).
57. Grimme, S., Ehrlich, S. & Goerigk, L. Effect of the damping function in dispersion corrected density functional theory. *J. Comput. Chem.* **32**, 1456–1465 (2011).
58. Lauritsen, J. V. et al. Atomic-scale insight into structure and morphology changes of MoS₂ nanoclusters in hydrotreating catalysts. *J. Catal.* **221**, 510–522 (2004).
59. Raybaud, P. et al. Ab initio study of the H₂–H₂S/MoS₂ gas–solid interface: the nature of the catalytically active sites. *J. Catal.* **189**, 129–146 (2000).
60. Monkhorst, H. J. & Pack, J. D. Special points for Brillouin-zone integrations. *Phys. Rev. B* **13**, 5188–5192 (1976).

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Author contributions

D.D. conceived the project and designed the experiments. Z.Z. performed the materials synthesis, materials characterization, and electrochemical measurements. L.Y. conducted the DFT calculations. M.G., X.C., W.Z., and C.M. performed the atomic-resolution HAADF-STEM measurements. L.W. and J.Z. assisted with the S L-edge XANES measurements. X.M., J.H., Y.T., and J.M. assisted with partial materials characterization. S.W.

conducted the Raman measurements. Z.T. gave the valuable discussions and suggestions. Z.Z., L.Y., and D.D. co-wrote the paper.

Competing interests

The authors declare no competing interests.

Additional information

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