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3, 7125One-pot synthesis and microstructure analysis of
Fe-doped NiS₂ for efficient oxygen evolution
electrocatalysis†Haofei Zhao,‡ Wenrui Li‡ and Rongming Wang *

The development of low-cost and effective electrocatalysts for water splitting is important for the production and application of green hydrogen as a renewable energy source. In this work, Fe-doped NiS₂ with different doping contents was successfully prepared through a one-pot solvothermal process utilizing a PEGylated deep eutectic solvent. The morphology, microstructure, electronic structure and oxygen evolution reaction (OER) catalytic activity of the synthesized samples are systematically investigated. The sample with an Fe content of ~16% possesses the best electrocatalytic performance with a low overpotential of 257 mV at 10 mA cm⁻², a small Tafel slope of 41 mV dec⁻¹ and a superior stability for 60 h at ~20 mA cm⁻². Stable and homogenous Ni(Fe) oxide porous nanostructures with residual S, including pores of several nanometers, were found after the long-term OER test. X-Ray photoelectron spectroscopy and electron energy loss spectroscopy mapping indicate that the enhanced OER property may be attributed to the local composition variation of Fe at the nanoscale and the synergic effect of Fe and Ni. This study provides insights into the rational design and synthesis of superior transition-metal based water splitting electrocatalysts.

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Introduction

With the imminent threat of global warming and the rapid depletion of fossil fuels, it is urgent to develop green and renewable energy systems. Due to its high energy density, hydrogen is a promising candidate as a carbon-free energy carrier and may play a role in reducing industrial carbon emissions.¹ Water electrolysis using renewable electricity is an attractive strategy to produce hydrogen. However, the oxygen evolution reaction (OER) at the anode is a sluggish kinetic process involving four electrons, thus limiting the overall efficiency. The state-of-the-art electrocatalysts are based on noble metals, and their high cost and insufficient reserves restrict their wide application.² Therefore, it is important to develop low-cost and effective OER catalysts for water splitting.

In the past few decades, earth-abundant-transition-metal based compounds have been widely explored as OER electrocatalysts, including oxides, chalcogenides, nitrides, phosphides *et al.*^{3,4} Recent studies have suggested that transition metal sulfides may

exhibit superior OER performance.^{5–8} Previous studies have revealed that the oxyhydroxides *in situ* formed at the surface of sulfides during the OER catalysis in alkaline media are the real active species.^{9–12} And a systematic study of 3d transition-metal oxyhydroxides reveals that the OER activity of these samples follows a trend of Ni > Co > Fe > Mn.¹³ Thus, nickel sulfides are promising electrocatalysts for water splitting.^{14,15}

Heteroatom doping is an effective strategy to improve the electrocatalytic performance. Yan *et al.* revealed that Fe³⁺ doping in the NiS₂(002) facet lowers the activation energy of H₂ generation of active sites, resulting in a good electrocatalytic hydrogen evolution activity.¹⁶ Han *et al.* reported that confined Fe over doping in Ni₃S₂ induces a synergic effect of size decrease and electronic structure modulation, thus resulting in a 70 mV decrease in the overpotential at 10 mA cm⁻² for the OER.¹⁷ Ding *et al.* suggested that the Fe incorporation in NiS₂ regulates the electronic structure of pyrite NiS₂, increasing its conductivity, and the conductive Fe-doped NiS₂ with an *in situ* electrochemical formed surface layer of oxyhydroxides during the OER exhibits a low overpotential of 260 mV at 10 mA cm⁻².¹⁸ These doped nickel sulfides are usually synthesized through a two-step precursor sulfuration route with a sulfur source under high temperature. Combining the two steps into a one-step synthesis should be cost-effective.

As quasi-ionic liquids, deep eutectic solvents (DES) are promising for applications in nanomaterial synthesis due to

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their unique physicochemical properties, such as low cost, easy preparation, environmental friendliness *etc.*^{19,20} Jiang *et al.* developed a PEGylated DES by simply mixing polyethylene glycol 200 (PEG-200) with thiourea in the molar ratio of 2:1, and demonstrated its prospective application in the one-pot solvothermal synthesis of metal sulfides with tunable composition.²¹ PEG 200 in the solvent could act as a shape-control agent, while thiourea could work as the sulfur source and help to stabilize the M-S bonds in metal sulfides.^{21,22}

In this work, we synthesized a series of Fe-doped NiS₂ with different composition through a one-pot solvothermal process using PEGylated DES, and performed a detailed characterization of the prepared samples. The doped samples exhibit enhanced OER activity compared to pure NiS₂, with the best found in (NiFe)S₂-3 (~16% Fe). (NiFe)S₂-3 needs a low overpotential of 257 mV to deliver a current density of 10 mA cm⁻² and a good stability (> 60 h at ~20 mA cm⁻²) in 1 M KOH.

Experimental

Synthesis

The Fe-doped NiS₂ samples were synthesized by a simple solvothermal method. In a typical procedure, a PEGylated DES was firstly prepared by mixing PEG 200 and thiourea in the molar ratio of 2:1.²¹ Then, 0.5 mmol nickel acetate and 0.5 mmol iron(II) acetate were dissolved in 40 ml PEGylated DES and stirred and sonicated for about 1 h to form a transparent homogeneous solution. The solution was subsequently transferred to a 50 ml Teflon-line autoclave, maintained at 200 °C for 10 h, and then cooled down to room temperature naturally. The product was washed with ethanol and water several times. Afterward, the sample was collected by centrifugation (3000 rpm, 3 min) after a low-speed centrifugation (200 rpm, 1 min) to remove large particles (FeS₂ and some aggregations), and dried in an oven at 80 °C for 4 h. This sample was denoted as (NiFe)S₂-3. Following a similar procedure, samples with different Fe doping content were prepared by adding different amounts of acetates while the total amount of Ni and Fe acetates was kept as 1.0 mmol, and labeled as NiS₂ (No Fe doping), (NiFe)S₂-1 (0.2 mmol Fe acetate), (NiFe)S₂-2 (0.4 mmol Fe acetate) and (NiFe)S₂-4 (0.6 mmol Fe acetate).

Characterizations

Powder X-ray diffraction (XRD) patterns were acquired on a Rigaku Ultima IV using Cu K α radiation at a scan rate of 0.02 °C s⁻¹. Transmission electron microscopy (TEM) investigations were performed with a JEOL JEM 2200FS electron microscope. X-Ray photoelectron spectra (XPS) were collected on a PHI5000 VersaProbe III spectrometer with a microfocus and monochromatic Al K α (1486.6 eV) X-ray excitation source. The core-level XPS profiles were fitted using Gaussian-Lorentzian lineshapes with a Shirley-type background.

Electrochemical measurements

Electrochemical tests were performed with a CHI760E electrochemical workstation in a standard three-electrode system in

1 M KOH, using a Pt foil as the counter electrode and a Hg/HgO electrode as the reference electrode. To prepare the working electrode, 3 mg of the prepared samples and 30 μ L 5 wt% Nafion solution were dispersed in 470 μ L ethanol by 30 min sonication to form a homogeneous ink, and then 50 μ L ink was drop-cast on a piece of carbon paper and dried naturally with a catalyst loading of ~0.3 mg cm⁻². The electrolyte was bubbled with O₂ for 30 min before the electrochemical measurements. The prepared working electrodes were activated *via* consecutive cyclic voltammetry (CV) scans for 20 cycles at a scan rate of 10 mV s⁻¹. 100% *iR*-compensated cyclic voltammograms with a scan rate of 2 mV s⁻¹ were used to characterize the OER activity of the samples. The Tafel plots were extracted from the backward sweep of the CV curves to avoid the effect of the redox peaks of Ni. Electrochemical impedance spectroscopy (EIS) was performed in the frequency range from 1 Hz to 100 kHz with an amplitude of 5 mV at the potential of 10 mA cm⁻² for each sample. The electrochemical active surface area (ECSA) was evaluated by carrying out a series of CV measurements in a non-faradaic potential range (0.1 V–0.2 V *vs.* Hg/HgO) at various scan rates (5, 10, 20, 30 and 40 mV s⁻¹). The overpotentials were calculated using the equation: $\eta = E_{\text{Hg/HgO}} + 0.098 \text{ V} + 0.059 \cdot \text{pH} - 1.23 \text{ V}$.

Results and discussion

Structure and morphology

TEM investigations reveal that the as-prepared NiS₂ sample exhibits porous nanostructures (Fig. S1, ESI[†]), while the Ni-Fe sulfides show similar morphology and are composed of irregular nanoparticles of tens of nanometers, as shown in Fig. 1(a–e). Apparent diffraction contrast in the bright-field TEM images indicates the crystal nature of the nanoparticles. Fig. 1(b–e) suggests that the size of the crystallites decreases with the increase of the Fe content, which should have resulted from the decrease of the Ni content in the raw materials and the lattice instability due to Fe over doping.¹⁷

Fig. 1(f) presents typical energy dispersive X-ray spectroscopy (EDX) elemental maps of the nanostructures consisting of several nanocrystals. Homogenous distribution of Ni, Fe and S in the crystallites could be observed in the maps.

The crystal structures of the samples were characterized by XRD. As shown in Fig. 1(g), major peak positions in the XRD patterns matched well with those of NiS₂ (PDF No. 11-99). Minor peaks related to FeS₂ (PDF No. 71-53) can also be distinguished in the spectra of the samples, suggesting the existence of tiny FeS₂ nanoparticles. Detailed investigations indicate that the major diffraction peaks shift to higher angles slightly. The right panel of Fig. 1(g) depicts enlarged (200) peaks of NiS₂. The observed gradually increased positions from 31.58° (NiS₂) to 31.88° ((NiFe)S₂-4) indicate that the lattice parameters of the samples decrease with Fe doping. The cations in disulfides are coordinated by 6 sulfur anions, and the ionic radius of Fe²⁺ (75 pm) is smaller than that of Ni²⁺ (83 pm) under octahedral coordination. The lattice contraction should be



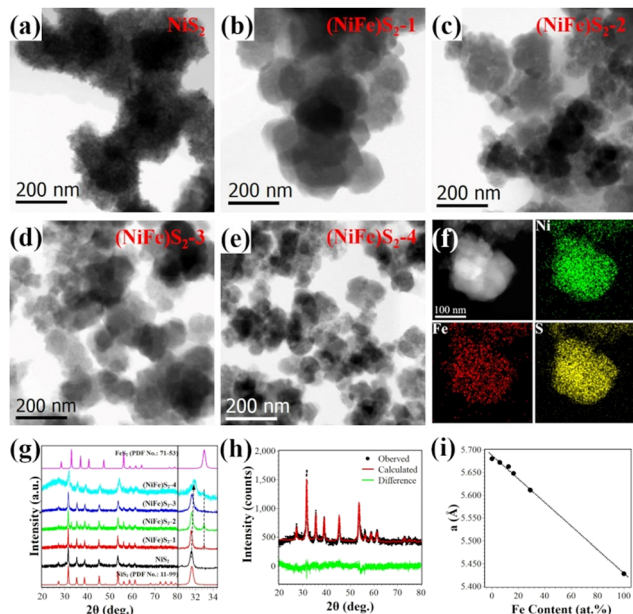


Fig. 1 (a–e) Bright-field TEM images of NiS_2 , $(\text{NiFe})\text{S}_{2-1}$, $(\text{NiFe})\text{S}_{2-2}$, $(\text{NiFe})\text{S}_{2-3}$ and $(\text{NiFe})\text{S}_{2-4}$. (f) High-angle annular dark field image of a nanoparticle composed of several crystallites and corresponding EDX elemental maps of $(\text{NiFe})\text{S}_{2-3}$. (g) XRD patterns (left) and enlarged (200) main peak (right) of the as-prepared samples. (h) Graphic results of the Rietveld refinement fitting of the XRD data of $(\text{NiFe})\text{S}_{2-3}$ using MAUD software. (i) Variation of lattice parameters with Fe content in the prepared samples.

attributed to the replacement of Ni^{2+} by smaller Fe^{2+} . The broadening of the peaks maybe due to the size effect of the nanocrystals, which is consistent with the TEM observations. Quantitative analysis of the XRD data was performed by Rietveld fitting (Fig. 1(h)) using MAUD software.²³ The lattice parameters and average crystal sizes of the samples were retrieved and summarized in Table S1 (ESI†). The Fe contents determined by EDX quantitative analysis are also shown in Table S1 (ESI†). Fig. 1(i) depicts the variation of the lattice parameters with the Fe content. The almost linear relationship between them as previously reported^{18,24} suggests successful doping of Fe in NiS_2 .

High resolution TEM (HRTEM) investigations provide further insights into the nanostructures. Fig. 2(a) shows a representative HRTEM image of $(\text{NiFe})\text{S}_{2-3}$, exhibiting two well-resolved perpendicular lattice fringes with interplanar spacings of 0.40 nm and 0.34 nm, which could be indexed to the (110) and (111) planes of cubic disulfide, respectively. The observed amorphous structure on the surface with 1–3 atomic layers in thickness may have resulted from the residual carbonaceous species or surface oxidation of the sample. The inset shows the fast Fourier transform (FFT) pattern of the marked square in Fig. 2(a). The inhomogeneous contrast in the HRTEM image and the diffusing of the diffraction spots in the FFT pattern suggest lattice variations in the sample. Fig. 2(b) shows the inverse FFT (IFFT) image of the marked area. Contrast with stripes of several nanometers can be distinguished in the IFFT

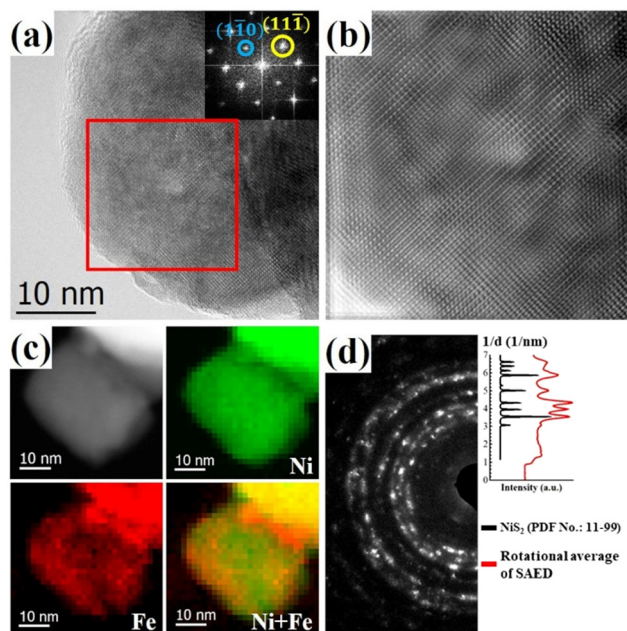


Fig. 2 Representative TEM images of $(\text{NiFe})\text{S}_{2-3}$. (a) High-resolution image and corresponding FFT pattern (inset). (b) Inverse FFT image of the area marked by the red square in (a). (c) EELS elemental maps of a crystallite with a pixel size of 1.6 nm showing the local distributions of Ni and Fe. (d) SAED pattern (left) and the comparison of the peak positions between its rotational average and the reference pattern of NiS_2 (right).

image, which may come from the local variation of the lattice parameters with different doping of Fe at the nanometer scale. The local distributions of Ni and Fe at the nanometer scale were obtained with electron energy loss spectroscopy (EELS) mapping in scanning TEM mode with a nanometer-size electron beam. As shown in Fig. 2(c), Fe distributes throughout the nanocrystal inhomogeneously at the nanometer scale and is rich in the surfaces/interfaces. The composition variation could induce strains and electronic structure modulation, which may benefit the electrocatalytic performance.¹⁷ The selected-area electron diffraction (SAED) pattern of $(\text{NiFe})\text{S}_{2-3}$ in Fig. 2(d) exhibits a pronounced ring structure, which can be indexed with NiS_2 , indicating the polycrystalline nature of the Fe-doped NiS_2 .

XPS analysis

The surface chemical states of Ni, Fe and S in $(\text{NiFe})\text{S}_{2-3}$ were investigated with XPS. The survey spectrum clearly proves the presence of Ni, Fe and S (Fig. 3(a)), while the C and O should be due to the surface adsorbed contaminants and/or oxidation. The $\text{Ni } 2p_{3/2}$ region exhibits a strong peak at 853.4 eV along with a satellite peak at 860.3 eV, corresponding to Ni^{2+} in disulfides,^{25–27} as shown in Fig. 3(b). The shoulder peak at 855.7 eV might be due to the presence of Ni^{3+} .^{11,28,29} The Fe $2p_{3/2}$ spectrum overlaps with two broad Ni LMM Auger peaks.^{30–32} The peak positions, widths, area ratio *etc.* of the two Auger peaks are obtained by fitting the XPS spectrum of NiS_2 ranging from 701 eV and 717 eV (Fig. S2, ESI†), and used in



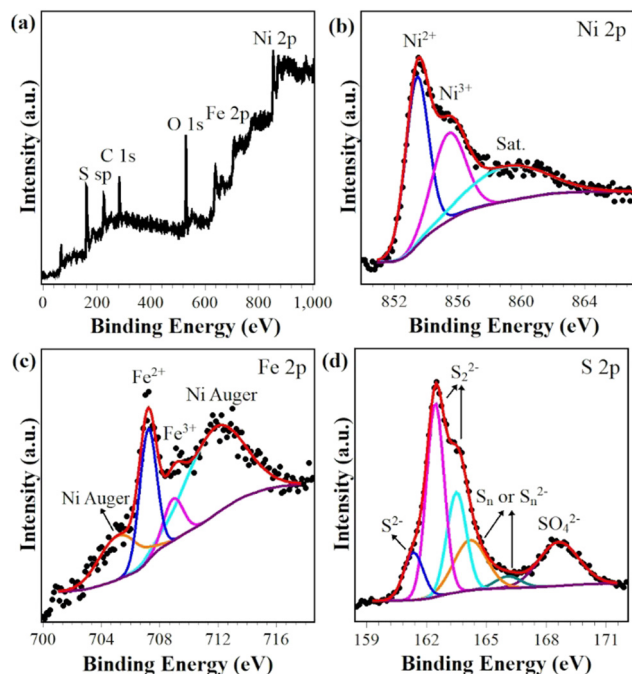


Fig. 3 XPS survey spectrum (a) and core-level spectra with deconvolution (b) Ni 2p, (c) Fe 2p and (d) S 2p of (NiFe) S_2 -3.

the fitting of the Fe 2p $_{3/2}$ spectrum. The fitted narrow peaks at 707.3 eV and 709.2 eV could be attributed to Fe $^{2+}$ in disulfides and Fe $^{3+}$, respectively.^{11,26,33} The S 2p profile could be deconvoluted to 6 peaks, as demonstrated in Fig. 3(d). The peaks at 162.5 eV and 163.5 eV unequivocally originate from 2p spin-orbital splitting of S $_2^{2-}$,^{33–35} and the peaks at 164.2 eV and 166.1 eV are ascribed to polysulfide S $_n$ or S $_n^{2-}$ often presented at the disulfide surfaces.^{33,36} The fitting peaks at 161.4 eV and 168.6 eV could be assigned to S $_2^{2-}$ and SO $_4^{2-}$ species, respectively.^{26,29,37} The XPS spectra of other samples exhibit similar features (Fig. S3, ESI †) and could be well fitted with the same components, the fitting parameters are listed in Table S2 (ESI †) (Ni 2p $_{3/2}$) and Table S3 (ESI †) (S 2p). Thus, these analysis results further confirm the successful doping of Fe in NiS $_2$.

Electrochemical oxygen evolution activity

The OER activities of Fe-doped NiS $_2$ were investigated in O $_2$ -saturated 1.0 M KOH using a standard three-electrode system. Blank CFP was also tested under the same conditions, and exhibited negligible activity. The 100% iR -compensated cyclic voltammograms with a scan rate of 2 mV s $^{-1}$ were shown in Fig. 4(a). The undoped sample (NiS $_2$) needs a high overpotential (η_{10}) of 411 mV at 10 mA cm $^{-2}$. With Fe doping, the η_{10} values are significantly reduced, and (NiFe) S_2 -3 exhibits the best activity with a low overpotential of 257 mV, which is comparable or even superior to that of other reports, such as Fe $_{0.1}$ Ni $_{0.9}$ S $_2$ (260 mV),¹⁸ V $_{0.1}$ Ni $_{0.9}$ S $_2$ (290 mV),³⁴ (NiFe) S_2 (230 mV),¹¹ Fe–NiS $_2$ /C (256 mV),³⁸ (NiFe) S_2 @Graphene (320 mV),²⁶ (NiCo) S_2 (270 mV),²⁷ Fe $_{0.2}$ Co $_{0.8}$ S $_2$ (290 mV)³⁹ *etc.* A detailed comparison with some reported transition-metal based OER catalysts (sulfides, phosphides, hydroxides *etc.*)

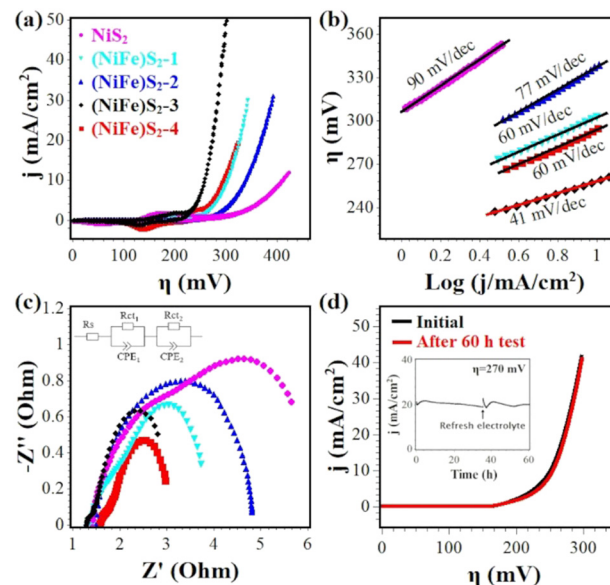


Fig. 4 (a) Cyclic voltammograms of NiS $_2$, (NiFe) S_2 -1, (NiFe) S_2 -2, (NiFe) S_2 -3 and (NiFe) S_2 -4 with 100% iR correction at a scan rate of 2 mV s $^{-1}$ in 1 M KOH. (b) Tafel plots extracted from the backward sweep of the corresponding voltammograms. (c) Nyquist plots of the EIS data in the frequency range of 100 kHz to 1 Hz; the inset is the equivalent circuit model used to fit the data. (d) Linear sweep voltammograms of (NiFe) S_2 -3 before and after 60 h test at ~ 20 mA cm $^{-2}$; the inset is the measured i - t curve.

can be found in Table S5 (ESI †). The Fe content of (NiFe) S_2 -3 is $\sim 16\%$, close to the optimal Fe content of Fe-doped Ni based OER electrocatalysts in previous reports.^{25,26,40,41} For better evaluation of the electrocatalytic performance, the Tafel plots were derived from the backward sweep of the voltammograms.⁴² As depicted in Fig. 4(b), (NiFe) S_2 -3 also shows the smallest Tafel slope of 41 mV dec $^{-1}$, indicating a fast reaction kinetics. For all samples that have similar ECSA (Fig. S4, ESI †), it is reasonable to conclude that (NiFe) S_2 -3 should possess a high intrinsic OER activity.

EIS measurements were carried out to characterize the charge-transfer resistance during the OER catalysis. The Nyquist plots of the EIS data are shown in Fig. 4(c) and fitted with a simplified Randles equivalent circuit (inset), where R_s is the solution resistance (~ 1.5 Ω), R_{ct1} and R_{ct2} represent the internal resistance of the electrode and the interfacial resistance between the electrode and the electrolyte, respectively.⁴³ As listed in Table S4 (ESI †), (NiFe) S_2 -3 displays a low total resistance ($R_{ct1} + R_{ct2}$) of ~ 2.3 Ω , suggesting a fast charge transfer during the OER process.

Fig. 4(d) shows the stability test results of (NiFe) S_2 -3. After being tested for 60 h at ~ 20 mA cm $^{-2}$, the activity loss is negligible and the overpotential to achieve 10 mA cm $^{-2}$ increased slightly.

Sample after the OER durability test

As shown in Fig. 5(a), (NiFe) S_2 -3 transformed to porous nanostructures after a long-time durability test (~ 20 mA cm $^{-2}$, 60 h), and holes of several nanometers presented in the tested sample, which may help to expose reaction sites and provide



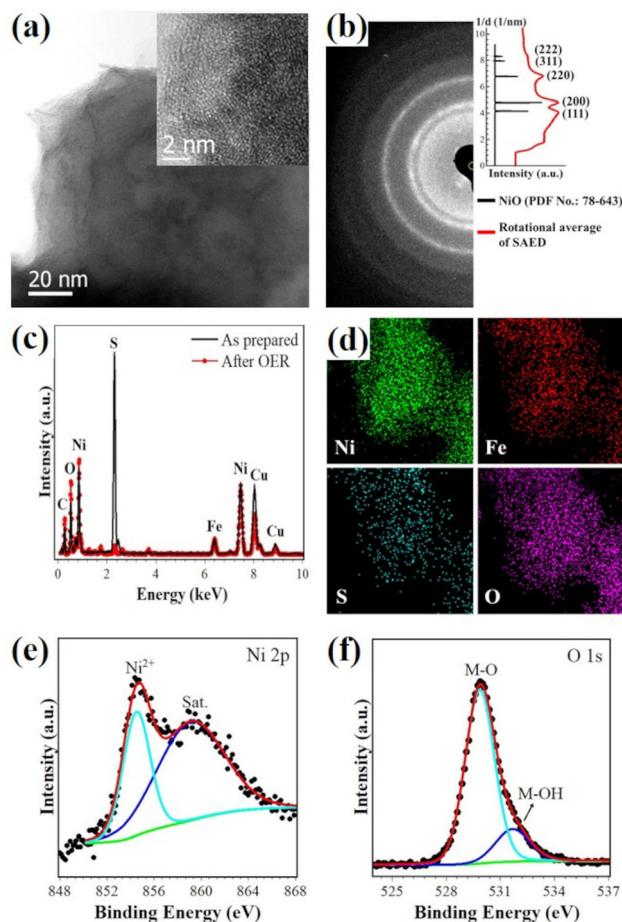


Fig. 5 Morphology, composition and surface chemical states of (NiFe) S_{2-3} after the durability test ($\sim 20 \text{ mA cm}^{-2}$, 60 h). (a) Bright-field and high-resolution TEM images (inset). (b) SAED pattern (left panel) and indexing (right panel), (c) comparison of the EDX spectra before and after the OER test, (d) EDX elemental maps of Ni, Fe, S and O, and the (e) Ni 2p and (f) O 1s XPS profiles.

pathways of mass transport, thus contributing to the superior electroactivity.^{10,44} The inset in Fig. 5(a) shows an HRTEM image, suggesting that the nanostructures are composed of many poorly-crystallized nanocrystals. Compared with the EDX spectrum of the as-prepared sample (Fig. 5(c)), the intensity of the S peak in that of the tested sample is significantly reduced, while the O peak increased, indicating the desulfurization and oxidation of the sample in the strongly oxidative environment during OER catalysis. Elemental maps of Ni, Fe, S, and O suggest that these elements are distributed uniformly (Fig. 5(d)). The polycrystalline rings in the SAED pattern could be well indexed with NiO (PDF No.: 78-643). The Ni 2p_{3/2} XPS profile is well fitted with a peak at 854.5 eV along with a satellite peak at 859.1 eV (Fig. 5(e)), which could be attributed to Ni²⁺ in NiO.^{28,45} The O 1s spectrum is composed of two components with the binding energies of 529.9 eV and 531.7 eV (Fig. 5(f)), which could be assigned to metal-O and metal-OH bands, respectively.^{45–47} These characterization results suggest the formation of nanoporous Ni(Fe) oxides after the stability test.

It has been shown that sulfides will be oxidized (at least on the surface) to oxyhydroxides in the strong oxidative environment during OER catalysis in an alkaline solution, which should be the real active species.^{9,11,18,32,48} Based on the reports on the redox behavior of Ni-Fe oxyhydroxides,^{49–51} it is reasonable to conclude that Ni(Fe) oxides result from the electrochemical reduction of *in situ* formed oxyhydroxides with the gradual decrease of the applied potential, and would be oxidized to oxyhydroxides when the potential increases. It should be noted that the sample is not fully desulfurized even after 60 h electrolysis at $\sim 20 \text{ mA cm}^{-2}$, and the residual S may play a role in the high OER activity.²⁵

Previous studies suggest that the stable electrocatalytic performance of sulfides during OER catalysis may be ascribed to the *in situ* electrochemical formation of a stable core-shell heterostructure consisting of an active oxyhydroxide layer and conductive disulfide core,^{11,12,18} stable oxyhydroxides,^{10,25,52} etc. We investigated the composition change of (NiFe) S_{2-3} during the durability test with TEM EDX. As shown in Fig. S5 (ESI[†]), the significant change in the EDX spectra is the reduction of the S peak, indicating the desulfurization of the prepared sample. After being tested for 1 h at $\sim 20 \text{ mA cm}^{-2}$, the S peak is still relatively strong, and is significantly reduced after being tested for 5 h, which is close to that of the sample being tested for 60 h. This result indicates the gradual transformation of (NiFe) S_{2-3} during the long-term electrocatalysis. Considering the negligible activity loss after a 60 h test (Fig. 4(d)), it is reasonable to conclude that the long-term stable catalytic performance may be due to the *in situ* electrochemically-formed stable oxyhydroxides.

Conclusions

In summary, we have successfully synthesized and systematically investigated Fe doped NiS₂ via a one-pot solvothermal process in PEGylated DES. When used as OER catalysts, the Fe-doped samples exhibit enhanced activity compared to the undoped one. The sample containing $\sim 16\%$ Fe ((NiFe) S_{2-3}) shows the best electrocatalytic activity, and only needs a low overpotential of 257 mV to achieve a current density of 10 mA cm^{-2} and displays a small Tafel slope of 41 mV dec^{-1} . Furthermore, (NiFe) S_{2-3} maintains its activity even after a stability test of 60 h at $\sim 20 \text{ mA cm}^{-2}$. The post-OER (NiFe) S_{2-3} is found to be nanoporous Ni(Fe) oxide with residual S, which should be *in situ* electrochemically transformed to active oxyhydroxides during the OER catalysis. Detailed investigations reveal the presence of local lattice distortions and nanoscale variation of Fe doping content, which may contribute to the superior performance of (NiFe) S_{2-3} . This work may contribute to the development of a low-cost and effective non-noble electrocatalyst for water splitting.

Conflicts of interest

There are no conflicts to declare.



Acknowledgements

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Notes and references

- J. Zhu, L. Hu, P. Zhao, L. Y. S. Lee and K. Y. Wong, *Chem. Rev.*, 2020, **120**, 851–918.
- J. Song, C. Wei, Z. F. Huang, C. Liu, L. Zeng, X. Wang and Z. J. Xu, *Chem. Soc. Rev.*, 2020, **49**, 2196–2214.
- J. Mohammed-Ibrahim, *J. Power Sources*, 2020, **448**, 227375.
- Z. P. Wu, X. F. Lu, S. Q. Zang and X. W. Lou, *Adv. Funct. Mater.*, 2020, **30**, 1–20.
- S. Shit, S. Chhetri, W. Jang, N. C. Murmu, H. Koo, P. Samanta and T. Kuila, *ACS Appl. Mater. Interfaces*, 2018, **10**, 27712–27722.
- Y. Wu, X. Liu, D. Han, X. Song, L. Shi, Y. Song, S. Niu, Y. Xie, J. Cai, S. Wu, J. Kang, J. Zhou, Z. Chen, X. Zheng, X. Xiao and G. Wang, *Nat. Commun.*, 2018, **9**, 1425.
- H. Li, S. Chen, Y. Zhang, Q. Zhang, Q. Zhang, X. Jia, L. Gu, X. Sun, L. Song and X. Wang, *Nat. Commun.*, 2018, **9**, 2452.
- X. Feng, Q. Jiao, H. Cui, M. Yin, Q. Li, Y. Zhao, H. Li, W. Zhou and C. Feng, *ACS Appl. Mater. Interfaces*, 2018, **10**, 29521–29531.
- S. Jin, *ACS Energy Lett.*, 2017, **2**, 1937–1938.
- X. Zou, Y. Wu, Y. Liu, D. Liu, W. Li, L. Gu, H. Liu, P. Wang, L. Sun and Y. Zhang, *Chem*, 2018, **4**, 1139–1152.
- M. Zhou, Q. Weng, X. Zhang, X. Wang, Y. Xue, X. Zeng, Y. Bando and D. Golberg, *J. Mater. Chem. A*, 2017, **5**, 4335–4342.
- Y. Liu, S. Ju, C. Wang, M. Li, W. Zhu, D. Chen, A. Yuan and G. Zhu, *Ind. Eng. Chem. Res.*, 2019, **58**, 18976–18985.
- R. Subbaraman, D. Tripkovic, K. C. Chang, D. Strmcnik, A. P. Paulikas, P. Hirunsit, M. Chan, J. Greeley, V. Stamenkovic and N. M. Markovic, *Nat. Mater.*, 2012, **11**, 550–557.
- P. Luo, H. Zhang, L. Liu, Y. Zhang, J. Deng, C. Xu, N. Hu and Y. Wang, *ACS Appl. Mater. Interfaces*, 2017, **9**, 2500–2508.
- M. Yao, B. Sun, L. He, N. Wang, W. Hu and S. Komarneni, *ACS Sustainable Chem. Eng.*, 2019, **7**, 5430–5439.
- J. Yan, H. Wu, P. Li, H. Chen, R. Jiang and S. Liu, *J. Mater. Chem. A*, 2017, **5**, 10173–10181.
- C. Han, W. Li, C. Shu, H. Guo, H. Liu, S. Dou and J. Wang, *ACS Appl. Energy Mater.*, 2019, **2**, 5363–5372.
- X. Ding, W. Li, H. Kuang, M. Qu, M. Cui, C. Zhao, D. C. Qi, F. E. Oropeza and K. H. L. Zhang, *Nanoscale*, 2019, **11**, 23217–23225.
- F. M. Perna, P. Vitale and V. Capriati, *Curr. Opin. Green Sustainable Chem.*, 2020, **21**, 27–33.
- B. B. Hansen, S. Spittle, B. Chen, D. Poe, Y. Zhang, J. M. Klein, A. Horton, L. Adhikari, T. Zelovich, B. W. Doherty, B. Gurkan, E. J. Maginn, A. Ragauskas, M. Dadmun, T. A. Zawodzinski, G. A. Baker, M. E. Tuckerman, R. F. Savinell and J. R. Sangoro, *Chem. Rev.*, 2021, **121**, 1232–1285.
- J. Jiang, C. Yan, X. Zhao, H. Luo, Z. Xue and T. Mu, *Green Chem.*, 2017, **19**, 3023–3031.
- C. Lamiel, V. H. Nguyen, M. Baynosa, D. C. Huynh and J. J. Shim, *J. Electroanal. Chem.*, 2016, **771**, 106–113.
- L. Lutterotti, S. Matthies, H. R. Wenk, A. S. Schultz and J. W. Richardson, *J. Appl. Phys.*, 1997, **81**, 594–600.
- L. Wang, T. Y. Chen, C. L. Chien, J. G. Checkelsky, J. C. Eckert, E. D. Dahlberg, K. Umemoto, R. M. Wentzcovitch and C. Leighton, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2006, **73**, 144402.
- T. Wang, G. Nam, Y. Jin, X. Wang, P. Ren, M. G. Kim, J. Liang, X. Wen, H. Jang, J. Han, Y. Huang, Q. Li and J. Cho, *Adv. Mater.*, 2018, **30**, 1–9.
- C. Liu, H. Ma, M. Yuan, Z. Yu, J. Li, K. Shi, Z. Liang, Y. Yang, T. Zhu, G. Sun, H. Li and S. Ma, *Electrochim. Acta*, 2018, **286**, 195–204.
- J. Zhang, X. Bai, T. Wang, W. Xiao, P. Xi, J. Wang, D. Gao and J. Wang, *Nano-Micro Lett.*, 2019, **11**, 2.
- P. Dubey, N. Kaurav, R. S. Devan, G. S. Okram and Y. K. Kuo, *RSC Adv.*, 2018, **8**, 5882–5890.
- P. Xu, J. Li, J. Luo, L. Wei, D. Zhang, D. Zhou, W. Xu and D. Yuan, *Sci. Rep.*, 2018, **8**, 1–9.
- G. Zhang, Y. S. Feng, W. T. Lu, D. He, C. Y. Wang, Y. K. Li, X. Y. Wang and F. F. Cao, *ACS Catal.*, 2018, **8**, 5431–5441.
- D. L. Legrand, G. M. Bancroft and H. W. Nesbitt, *Int. J. Miner. Process.*, 1997, **51**, 217–228.
- W. Li, H. Zhao, H. Li and R. Wang, *Nanoscale Adv.*, 2022, **4**, 1220–1226.
- M. V. Morales-Gallardo, A. M. Ayala, M. Pal, M. A. Cortes Jacome, J. A. Toledo Antonio and N. R. Mathews, *Chem. Phys. Lett.*, 2016, **660**, 93–98.
- H. Liu, Q. He, H. Jiang, Y. Lin, Y. Zhang, M. Habib, S. Chen and L. Song, *ACS Nano*, 2017, **11**, 11574–11583.
- L. Gao, C. Guo, X. Liu, X. Ma, M. Zhao, X. Kuang, H. Yang, X. Zhu, X. Sun and Q. Wei, *New J. Chem.*, 2020, **44**, 1711–1718.
- V. Toniazio, C. Mustin, J. M. Portal, B. Humbert, R. Benoit and R. Erre, *Appl. Surf. Sci.*, 1999, **143**, 229–237.
- X. Han, X. Wu, C. Zhong, Y. Deng, N. Zhao and W. Hu, *Nano Energy*, 2017, **31**, 541–550.
- L. Xie, D. Zhao, J. Dai, Z. Wu and L. Li, *Catalysts*, 2019, **9**, 458.
- Y. Zhou, M. Luo, Z. Zhang, W. Li, X. Shen, W. Xia, M. Zhou and X. Zeng, *Appl. Surf. Sci.*, 2018, **448**, 9–15.
- J. Landon, E. Demeter, N. Inoğlu, C. Keturakis, I. E. Wachs, R. Vasić, A. I. Frenkel and J. R. Kitchin, *ACS Catal.*, 2012, **2**, 1793–1801.
- N. Cheng, Q. Liu, A. M. Asiri, W. Xing and X. Sun, *J. Mater. Chem. A*, 2015, **3**, 23207–23212.
- S. Anantharaj, S. R. Ede, K. Karthick, S. Sam Sankar, K. Sangeetha, P. E. Karthik and S. Kundu, *Energy Environ. Sci.*, 2018, **11**, 744–771.
- J. Jiang, F. Sun, S. Zhou, W. Hu, H. Zhang, J. Dong, Z. Jiang, J. Zhao, J. Li, W. Yan and M. Wang, *Nat. Commun.*, 2018, **9**, 2885.
- X. Y. Yu and X. W. David Lou, *Adv. Energy Mater.*, 2018, **8**, 1–37.



- 45 R. A. Patil, C. P. Chang, R. S. Devan, Y. Liou and Y. R. Ma, *ACS Appl. Mater. Interfaces*, 2016, **8**, 9872–9880.
- 46 H. Ali-Löytty, M. W. Louie, M. R. Singh, L. Li, H. G. Sanchez Casalongue, H. Ogasawara, E. J. Crumlin, Z. Liu, A. T. Bell, A. Nilsson and D. Friebe, *J. Phys. Chem. C*, 2016, **120**, 2247–2253.
- 47 Q. Liu, J. Huang, Y. Zhao, L. Cao, K. Li, N. Zhang, D. Yang, L. Feng and L. Feng, *Nanoscale*, 2019, **11**, 8855–8863.
- 48 R. He, X. Huang and L. Feng, *Energy Fuels*, 2022, **36**, 6675–6694.
- 49 A. G. Marrani, V. Novelli, S. Sheehan, D. P. Dowling and D. Dini, *ACS Appl. Mater. Interfaces*, 2014, **6**, 143–152.
- 50 M. Gorlin, J. F. De Araujo, H. Schmies, D. Bernsmeier, S. Dresch, M. Gliech, Z. Jusys, P. Chernev, R. Kraehnert, H. Dau and P. Strasser, *J. Am. Chem. Soc.*, 2017, **139**, 2070–2082.
- 51 D. Di Girolamo, M. Piccinni, F. Matteocci, A. G. Marrani, R. Zannoni and D. Dini, *Electrochim. Acta*, 2019, **319**, 175–184.
- 52 D. Y. Chung, P. P. Lopes, P. Farinazzo Bergamo Dias Martins, H. He, T. Kawaguchi, P. Zapol, H. You, D. Tripkovic, D. Strmcnik, Y. Zhu, S. Seifert, S. Lee, V. R. Stamenkovic and N. M. Markovic, *Nat. Energy*, 2020, **5**, 222–230.

