Highly efficient oxygen evolution and stable water splitting by coupling NiFe LDH with metal phosphides

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ABSTRACT It is a great challenge to develop highly active oxygen evolution reaction (OER) electrocatalysts with superior durability. In this study, a NiFe layered double hydroxide-decorated phosphide (NiFe LDH@CoP/NiP₃) was constructed to display satisfactory OER activity and good stability for water splitting in alkaline media. At an overpotential of 300 mV, NiFe LDH@CoP/NiP₃ achieved a current density of 82 mA cm⁻² for the OER, which was 9.1 and 2.3 times that of CoP/NiP₃ and NiFe LDH, respectively. Moreover, the reconstruction behavior, during which oxyhydroxides formed, was studied by a combination of X-ray photoelectron spectroscopy, Raman spectroscopy, and scanning electron microscopy. A synergistic effect between NiFe LDH and CoP/NiP₃ was also observed for the hydrogen evolution reaction. Furthermore, when NiFe LDH@CoP/NiP₃ acted as both the cathode and anode for overall water splitting, a high current density of 100 mA cm⁻² was maintained for more than 275 h. In addition, under Xe light irradiation, a solar-to-hydrogen efficiency of 9.89% was achieved for solar-driven water splitting. This work presents the coupling of different active compositions, and can provide a reference for designing bifunctional electrocatalysts.

Keywords: bifunctional electrocatalysis, oxygen evolution reaction, phosphides, layered double hydroxides, water splitting

INTRODUCTION

Efficient electrocatalytic overall water splitting is expected to alleviate the increasingly severe energy shortage and

environmental crisis [1–4]. However, the slow kinetics of the oxygen evolution reaction (OER) and poor durability of catalysts under alkaline conditions [5–8]. Although RuO₂, IrO₂ and others have been utilized, there is an urgent necessity to develop earth-abundant OER catalysts with high activity, good stability, and low cost [9–12]. In previous studies, a series of OER catalysts in alkaline solutions have been developed, including metal oxides [13,14], hydroxides [15], selenides [16], sulfides [17,18], carbides [19], phosphides [20,21], and others [22–24]. Although phosphides generally show good hydrogen evolution reaction (HER) activity because of their similar catalytic mechanism to that of hydrogenase [25], their OER activities are unsatisfactory. Additionally, during the OER, phosphides are usually converted to oxyhydroxides, which are considered to be real active sites [26]. The investigation of reconstruction behavior requires more attention [27].

In contrast, layered double hydroxides (LDHs) are regarded as excellent OER catalyst candidates [28,29]. They have large specific surface areas and porous structures, which are beneficial for increasing the number of active sites and promoting mass transfer. However, because of their weak hydrogen adsorption, LDHs exhibit relatively poor HER activity, thus hindering their potential as bifunctional OER/HER catalysts. Synergistic effects play important roles in boosting bifunctional performance, such as in the OER/HER [30]. Yang *et al.* [31] decorated LDH with CoNiSe₂ to adjust H₂O chemisorption and

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produce reactive H intermediates, thus leading to efficient overall water splitting. Liang *et al.* [32] electrodeposited amorphous NiFe hydroxides onto a NiFeP surface to construct a hybrid structure. The resulting electronic interactions boosted water oxidation by lowering the adsorption energy of H₂O. Thus, it could be proposed that metal phosphides and NiFe LDH may compensate each other's weaknesses when combined, achieving bifunctional OER/HER activity [33,34].

In this study, we coupled NiFe LDH with metal phosphides to boost the OER activity and stability of water splitting in alkaline media. For the OER, at an overpotential of 300 mV, NiFe LDH@CoP/NiP₃ achieved a current density of 82 mA cm⁻², which was 9.1 and 2.3 times that of CoP/NiP₃ and NiFe LDH, respectively. The reconstruction behavior during the OER was also studied. As both the cathode and anode in an electrolytic cell, NiFe LDH@CoP/NiP₃, demonstrated robust stability for 275 h at a high current density of 100 mA cm⁻². Using the simulated solar illumination, water splitting driven by solar cells achieved a solar-to-hydrogen (STH) efficiency of 9.89%.

EXPERIMENTAL SECTION

Materials

Fe(NO₃)₃·9H₂O, Ni(NO₃)₂·6H₂O, CO(NH₂)₂, NaH₂PO₂·H₂O and NH₄F were purchased from Sinopharm Chemical Reagent Co, Ltd. (SCRC). Co(NO₃)₂·6H₂O and C₃H₄N₂ were purchased from Shanghai Macklin Biochemical Co, Ltd. All the reagents were of analytical grade and used directly without further purification.

Preparation of the catalysts

Synthesis of CoP/NiP₃

Firstly, Co(NO₃)₂·6H₂O (1 mmol), NH₄F (5 mmol) and CO(NH₂)₂ (5 mmol) were mixed and dissolved in 30 mL deionized (DI) water. Before hydrothermal reaction, Ni foams were successively cleaned with 3 mol L⁻¹ HCl, ethanol, and pure water under ultrasonication for 15 min to remove surface oxides and impurity. The mixture and a piece of Ni foam (2 × 1 cm²) were shifted to Teflon-lined stainless-steel autoclave and warmed up to 120°C and maintained for 8 h. The precursor was washed with DI water and ethanol for three times and dried in vacuum at 60°C. Secondly, C₃H₄N₂ (0.30 g) and the as-prepared precursor were put on the upper and downstream of quartz boat, respectively. Then the obtained sample was calcined to 280°C (2°C min⁻¹) and maintained for 2 h in

N₂. Next, phosphorization was realized at 260°C under the condition of the second step except using NaH₂PO₂·H₂O.

Synthesis of NiFe LDH@CoP/NiP₃

The synthesis of NiFe LDH was referred to literature [34]. Fe(NO₃)₃·9H₂O (0.25 mmol), Ni(NO₃)₂·6H₂O (0.75 mmol) and CO(NH₂)₂ (1.25 mmol) were mixed and dissolved in 30 mL DI water. The mixture and the above CoP/NiP₃ (2 × 1 cm²) were shifted to a Teflon-lined stainless-steel autoclave and warmed up to 120°C and maintained for 10 h. The sample was washed with DI water and ethanol for three times and dried in vacuum at 60°C.

Synthesis of NiFe LDH

NiFe LDH was synthesized by the similar process as NiFe LDH@CoP/NiP₃. Fe(NO₃)₃·9H₂O (0.25 mmol), Ni(NO₃)₂·6H₂O (0.75 mmol) and CO(NH₂)₂ (1.25 mmol) were mixed and dissolved in 30 mL DI water. Then, the mixture and a piece of treated Ni foam (2 × 1 cm²) instead of CoP/NiP₃ were shifted to a Teflon-lined stainless-steel autoclave and warmed up to 120°C and maintained for 10 h. The sample was washed with DI water and ethanol for three times and dried in vacuum at 60°C.

Characterizations

The crystalline structures were tested by X-ray diffraction (XRD) on D/max 2550 VB with 40 kV and Cu Kα radiation (λ = 1.5418 Å). The morphology of the samples was characterized by a field emission scanning electron microscope (FE-SEM, Quanta 250 FEG) and transmission electron microscope (FE-TEM, FEI Tecnai G2 F20). X-ray photoelectron spectroscopy (XPS) measurements were performed on Thermo Fisher Scientific K-Alpha with a monochromatic Al-Kα X-ray source. All XPS spectra were calibrated by shifting the detected adventitious C 1s peak to 284.4 eV. Raman spectra were measured by the Horiba Jobin Yvon HR Evolution with a 532-nm excitation laser.

Electrochemical test

OER and HER tests were performed in a three-electrode system of an electrochemical station (CHI 660E) at room temperature. A graphite rod and a saturated calomel electrode (SCE) were used as the counter electrode and the reference electrode, respectively. The prepared NiFe LDH@CoP/NiP₃ was directly used as the work electrode without further treatment. The reference electrode is converted into a potential relative to the reversible hydrogen electrode (RHE): $E(\text{RHE}) = E(\text{SCE}) + 0.2415 +$

0.0591 $\times$ pH. The water splitting was conducted in 1 mol L⁻¹ KOH through a two-electrode system which employed NiFe LDH@CoP/NiP₃ as both cathode and anode. All the linear sweep voltammetry (LSV) of HER, OER and water splitting were tested at the sweep rate of 2 mV s⁻¹ with 90% internal resistance (IR) compensation.

For the solar-driven water splitting, the STH efficiency can be defined as the following equation:

$$\text{STH (\%)} = \text{Chemical energy output} / \text{Solar energy input} = j \times A \times E_f \times \text{FE}_{\text{H}_2} / (P_s \times A),$$

where j represents the operating current density; A represents the effective illuminated area; E_f is the standard thermodynamic potential difference between HER and OER (1.23 V). FE_{H_2} is the Faradic efficiency of HER which is regarded as 100%; P_s is the power of simulated solar illumination (100 mW cm⁻²).

RESULTS AND DISCUSSION

To realize the composite of CoP/NiP₃ and NiFe LDH, CoP/NiP₃ was synthesized, followed by the growth of NiFe LDH (Fig. S1). The specific quantities of raw materials are stated in Table S1. XRD patterns were recorded to obtain crystallographic information (Fig. 1a). Table S2 summarizes the peaks for NiFe LDH@CoP/NiP₃ and their attributions. The peaks located at $2\theta = 45.1^\circ$, 52.3° and 76.8° correspond to the (111), (200), and (220) crystallographic planes of metallic Ni (PDF#03-1051), respectively. The peaks at $2\theta = 21.4^\circ$ and 39.4° indicate the presence of nickel hydroxides (PDF#73-1520). The peak at $2\theta = 36.1^\circ$ corresponds to the (111) crystallographic plane of CoP (PDF#29-0497). The peaks at 16.0° and 36.3° correspond to the (200) and (013) crystallographic planes of NiP₃ (PDF#73-1242), respectively. XPS was then conducted to determine the surface composition and chemical valence (Figs S2–S7). As shown in Fig. 1b, NiFe LDH@CoP/NiP₃ presents peaks corresponding to Fe, Co,

Ni, C, P and O. The coexistence of Fe and P illustrates that NiFe LDH@CoP/NiP₃ was successfully obtained. The element contents are stated in Table S3. We noted that the Fe content was only 0.1 wt%, which may suggest the low NiFe LDH content. Due to the low contents of Fe and Ni in the raw materials, the relative content of NiFe LDH was low; this explains the lack of recognizable characteristic peaks associated with NiFe LDH in the XRD patterns. Additionally, a weak peak at 26.6° , attributed to amorphous carbon, was observed, which resulted from small quantities of CO(NH₂)₂ and C₃H₄N₂ in the raw materials [35,36].

In the SEM images (Fig. 2a and Fig. S8), NiFe LDH@CoP/NiP₃ shows an arborized structure (a typical characteristic of metal phosphides) on Ni foam. This structure could facilitate intimate contact between the catalyst and electrolyte to boost the OER kinetics [37]. The energy dispersive spectra (EDS) of NiFe LDH@CoP/NiP₃ also supported the presence of Fe, Co, Ni, C, P, etc. (Fig. S9), in accordance with the XPS spectra. Furthermore, the uniform distribution of these elements was confirmed by the TEM elemental mapping images (Fig. 2b, c). Fig. 2d presents the morphology of NiFe LDH@CoP/NiP₃. The phosphides were decorated with thin LDH nanosheets. The high-resolution TEM images (Fig. 2e, f) show the presence of CoP and NiP₃ particles. The clear crystal planes with d -spacing of 0.248 and 0.208 nm correspond to the (111) plane of CoP and the (321) plane of NiP₃, respectively.

The OER performance was evaluated in 1 mol L⁻¹ KOH solution (Fig. 3a). As shown, NiFe LDH@CoP/NiP₃ only required an overpotential of 239 mV to drive 10 mA cm⁻², outperforming its isolated NiFe LDH and CoP/NiP₃ counterparts [38]. At a large current density of 100 mA cm⁻², NiFe LDH@CoP/NiP₃ also displayed a significant overpotential advantage (310 mV) compared

Figure 1 (a) XRD patterns and (b) XPS survey of various samples.

Figure 2 (a) SEM images of NiFe LDH@CoP/NiP₃, (b) TEM and (c) corresponding elemental mapping images. (d) TEM and (e, f) high-resolution TEM images of NiFe LDH@CoP/NiP₃.

Figure 3 (a) OER polarization curves in 1.0 mol L⁻¹ KOH. (b) Overpotentials at 100 mA cm⁻² (left) and current densities at $\eta = 300 \text{ mV}$ (right) of different samples. (c) Tafel plots of different samples. (d) The overpotentials of recently reported electrocatalysts in 1.0 mol L⁻¹ KOH.

with NiFe LDH and CoP/NiP₃. In particular, NiFe LDH@CoP/NiP₃ only required an overpotential of 300 mV to reach a current density of 82 mA cm⁻², which is 164.0, 9.1 and 2.3 times that of Ni foam, CoP/NiP₃ and NiFe LDH, respectively (Fig. 3b). The Tafel slope value of NiFe LDH@CoP/NiP₃ was 94 mV dec⁻¹ (Fig. 3c), which indicates faster kinetics than the other samples [39].

Based on the above result, the enhanced OER activity should be attributed to the synergy of CoP/NiP₃ and NiFe LDH [40]. Compared with both LDH catalysts and phosphide catalysts reported recently, NiFe LDH@CoP/NiP₃ exhibited a relatively favorable performance at both 10 and 100 mA cm⁻² (Fig. 3d and Tables S4 and S5). The NiFe LDH@CoP/NiP₃ demonstrated good stability for

more than 70 h at an overpotential of 300 mV (Fig. S10).

It has been reported that phosphides exhibit self-reconstruction phenomena during the OER process [41–43]. The reconstructed catalysts, with oxyhydroxides on the surface, inherit the advantages of nanostructures, which are stable and act as the true active species [40]. To investigate the reconstruction of NiFe LDH@CoP/NiP₃ during OER, XPS and Raman spectra were recorded before and after OER. In Fig. 4a, the Co 2p_{3/2} peaks at 774.8, 780.3 and 796.4 eV correspond to the Co⁰, oxidized Co²⁺/Co³⁺, and satellite peak, respectively [44]. The weakened Co⁰ peak of NiFe LDH@CoP/NiP₃ after OER indicates the improved oxidation state of the Co species. As shown in Fig. 4b, the binding energies of Ni 2p_{3/2} (854.9 eV) and Ni 2p_{1/2} (872.7 eV) shift to 855.3 and 873.0 eV, respectively, demonstrating the improved oxidation state after OER [45]. In the O 1s spectra of NiFe LDH@CoP/NiP₃ (Fig. 4c), the peaks at 530.4 and 531.5 eV are attributed to M–O bonds (M representing metal) and oxygen defects in LDH, respectively. The peak at 532.5 eV corresponds to the adsorbed O resulting from hydroxyl groups (such as adsorbed H₂O) [46]. In the Raman spectra of NiFe LDH@CoP/NiP₃, the peaks located at 188, 321 and 534 cm^{−1} correspond to O–M–O, O–H and M–O–M vibrations in LDH, respectively (Fig. 4d and Table S6) [47–49]. After the OER, the peaks

at 188 and 321 cm^{−1} were not present because of the conversion of hydroxides into oxyhydroxides. Under anodic potentials, the hydrophilic, conductive and chemically stable M–OOH may be formed. This could serve as the real active composition to boost OER activity [50–53]. The crystal structure and surface morphology of NiFe LDH@CoP/NiP₃ after OER were also investigated. After the OER process, the phosphide peak intensity was reduced, which may be attributed to the formation of amorphous oxyhydroxides. In the SEM images (Fig. S11a, b), the initial arborized structure was seen to have transformed into a finer flower-like structure after the OER. This change in surface morphology also confirmed the reconstruction of the catalyst, consistent with the afore-described XPS and Raman results. After reconstruction of the phosphides, the M–OOH active site was introduced, and the transformed flower-like structure exposes more active sites. These factors significantly improve the catalyst activity.

Subsequently, the HER activity of NiFe LDH@CoP/NiP₃ was evaluated, which demonstrated clear superiority over those of CoP/NiP₃ and NiFe LDH (Fig. 5a). The overpotentials of NiFe LDH@CoP/NiP₃ at different current densities are listed in Fig. S12. To reach a current density of 50 mA cm^{−2}, the required overpotential was only 206 mV (Fig. 5b left). For NiFe LDH@CoP/NiP₃, an

Figure 4 (a–c) The corresponding high-resolution XPS spectra of Co 2p, Ni 2p and O 1s. (d) Raman spectra of NiFe LDH@CoP/NiP₃.

Figure 5 (a) HER polarization curves in 1.0 mol L⁻¹ KOH. (b) Overpotentials at 50 mA cm⁻² (left) and current densities at $\eta = 300$ mV (right) of various samples. (c) Tafel plots of different samples.

Figure 6 (a) Overall water splitting polarization curves in 1.0 mol L⁻¹ KOH. (b) The required cell voltage of electrocatalysts at 10 mA cm⁻² in recent years. (c) Stability test of NiFe LDH@CoP/NiP₃. Insets: photograph evolving from the electrodes at the beginning of stability test and after 275 h. (d) J-V curve of a solar cell and the polarization curves of NiFe LDH@CoP/NiP₃ for water splitting in a two-electrode system.

overpotential of 300 mV resulted in a current density of 315 mA cm⁻², which is 9.0, 5.3 and 3.4 times that of Ni foam, CoP/NiP₃, and NiFe LDH, respectively (Fig. 5b right). Fig. 5c shows that the Tafel slope of NiFe LDH@CoP/NiP₃ is also the smallest (74 mV dec⁻¹), indicating the Volmer-Heyrovsky mechanism [54].

Considering the above results, NiFe LDH@CoP/NiP₃ shows potential as a bifunctional OER/HER electrocatalyst. We then explored the overall water splitting performance (Fig. S13). Using NiFe LDH@CoP/NiP₃ as both the cathode and anode, a cell voltage of 1.64 V was

required to reach a current density of 10 mA cm⁻² (Fig. 6a), which is comparable to most of the reported results summarized in Fig. 6b, based on Tables S7 and S8. Notably, the constant current test at 100 mA cm⁻² lasted for 275 h with negligible activity loss (Fig. 6c). As seen in the inset in Fig. 6c, H₂ bubbles at the cathode and the O₂ bubbles at the anode were continuously produced during the entire electrolysis process, further illustrating the excellent stability of the NiFe LDH@CoP/NiP₃ electrode. The high durability of this water splitting system undoubtedly makes it promising for application. We

speculated that, as a self-supporting electrode, the Ni foam achieved close contact with the catalyst layer, which reduced the charge transfer resistance and enhanced the durability under large current densities [55]. Additionally, the three-dimensional porous structure increased the area of exposed active sites, hence boosting the catalytic activity [56]. Then, we designed a simple solar-driven water splitting system, which comprised a commercial Si solar plane (16 cm^2) and electrodes. Fig. S14a, b show the electrolyzed water test, which used NiFe LDH@CoP/NiP₃ as both the cathode and anode in 1.0 mol L^{-1} KOH solution under Xe light irradiation as simulated sunlight. H₂ and O₂ were spontaneously produced under the simulated solar irradiation. From the current-voltage (*J-V*) curve of the solar cell and the polarization curve of the water splitting system, the operating current density at 1.59 V was calculated to be $\sim 8\text{ mA cm}^{-2}$, which corresponded to an STH efficiency of 9.89% (Fig. 6d) [57,58]. Thus, NiFe LDH@CoP/NiP₃ displayed potential to be used in solar-driven water splitting.

CONCLUSIONS

In summary, NiFe LDH@CoP/NiP₃ was constructed as a bifunctional OER/HER catalyst, which demonstrated improved performance over the sum of its isolated components. Under alkaline condition, NiFe LDH@CoP/NiP₃ achieved an OER current density of 82 mA cm^{-2} (at an overpotential of 300 mV), which was 9.1 times that of CoP/NiP₃. Moreover, the reconstruction behavior during the OER was studied. In particular, for overall water splitting, the sample displayed excellent durability for 275 h at 100 mA cm^{-2} . Under Xe light irradiation, the water splitting STH efficiency was 9.89%. A detailed study of the synergistic mechanism between CoP/NiP₃ and NiFe LDH is currently under way. This work presents the coupling of different active compositions and their synergistic catalytic performance, and can provide a reference for designing bifunctional electrocatalysts.

Received 25 August 2020; accepted 17 November 2020;
published online 2 February 2021

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Acknowledgements This work was financially supported by Hunan Provincial Science and Technology Plan Project (2017TP1001 and 2020JJ4710), the National Key R&D Program of China (2018YFB0704100) and the State Key Laboratory Fund.

Author contributions Song C and Liu Y performed the experiments

and sample preparation; Song C and Wang Y wrote the paper with support from Lei Y; Tang S was responsible for the data processing; Li W and Zeng J contributed to the theoretical analysis; Li Q, Chen L and Peng H provided constructive suggestions. All authors contributed to the general discussion.

Conflict of interest The authors declare no conflict of interest.

Supplementary information Supporting data are available in the online version of the paper.

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层状镍铁双氢氧化物与金属磷化物耦合实现高效析氧和稳定的全解水

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摘要 研制具有优良稳定性的高活性析氧反应(OER)电催化剂是一个巨大的挑战. 这项工作构建了层状镍铁双氢氧化物修饰的磷化物(NiFe LDH@CoP/NiP₃), 其在碱性介质中呈现出了令人满意的OER活性和良好的全解水稳定性. 在300 mV的过电位下, NiFe LDH@CoP/NiP₃的电流密度为82 mA cm⁻², 分别是CoP/NiP₃和NiFe LDH的9.1倍和2.3倍. 通过X射线光电子能谱、拉曼和扫描电镜表征, 研究了OER过程中的重构行为. 氢析出性能测试也论证了NiFe LDH与CoP/NiP₃之间的协同效应. 此外, NiFe LDH@CoP/NiP₃同时作为阴极和阳极进行全解水时, 可以维持100 mA cm⁻²的高电流密度超过275小时. 此外, 在氙灯的照射下, 太阳能驱动的水分解可以实现9.89%的光产氢效率. 这项工作实现了不同活性成分间的耦合, 为双功能电催化剂的设计提供了参考.