




Article

Salt-Templated Nanoarchitectonics of CoSe₂-NC Nanosheets as an Efficient Bifunctional Oxygen Electrocatalyst for Water Splitting

Hong Cao ¹, Hailong Li ^{1,*} , Linhao Liu ², Kangning Xue ¹, Xinkai Niu ¹, Juan Hou ^{1,*} and Long Chen ²

¹ Key Laboratory of Ecophysics, Department of Physics, College of Science, Shihezi University, Shihezi 832003, China; c_hong1002@sina.com (H.C.); x-15937874475@sina.com (K.X.); niuxinkai0424@sina.com (X.N.)

² Key Laboratory for Green Process of Chemical Engineering of Xinjiang Bingtuan, School of Chemistry and Chemical Engineering, Shihezi University, Shihezi 832003, China; thebestleo@sina.cn (L.L.); chenlong2012@sinano.ac.cn (L.C.)

* Correspondence: well09131015@126.com (H.L.); hjuan05@sina.com (J.H.)

Abstract: Recently, the extensive research of efficient bifunctional electrocatalysts (oxygen evolution reaction (OER) and hydrogen evolution reaction (HER)) on water splitting has drawn increasing attention. Herein, a salt-template strategy is prepared to synthesize nitrogen-doped carbon nanosheets encapsulated with dispersed CoSe₂ nanoparticles (CoSe₂-NC NSs), while the thickness of CoSe₂-NC NSs is only about 3.6 nm. Profiting from the ultrathin morphology, large surface area, and promising electrical conductivity, the CoSe₂-NC NSs exhibited excellent electrocatalytic of 10 mA·cm⁻² current density at small overpotentials of 247 mV for OER and 75 mV for HER. Not only does the nitrogen-doped carbon matrix effectively avoid self-aggregation of CoSe₂ nanoparticles, but it also prevents the corrosion of CoSe₂ from electrolytes and shows favorable durability after long-term stability tests. Furthermore, an overall water-splitting system delivers a current density of 10 mA·cm⁻² at a voltage of 1.54 V with resultants being both the cathode and anode catalyst in alkaline solutions. This work provides a new way to synthesize efficient and nonprecious bifunctional electrocatalysts for water splitting.



Citation: Cao, H.; Li, H.; Liu, L.; Xue, K.; Niu, X.; Hou, J.; Chen, L.

Salt-Templated Nanoarchitectonics of CoSe₂-NC Nanosheets as an Efficient Bifunctional Oxygen Electrocatalyst for Water Splitting. *Int. J. Mol. Sci.*

2022, 23, 5239. <https://doi.org/10.3390/ijms23095239>

Academic Editors: Katsuhiko Ariga, Fabien Grasset and Yann Molard

Received: 7 April 2022

Accepted: 2 May 2022

Published: 7 May 2022

Publisher's Note: MDPI stays neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Copyright: © 2022 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<https://creativecommons.org/licenses/by/4.0/>).

Keywords: metal–organic frameworks; nitrogen-doped carbon; CoSe₂; oxygen reduction reaction; hydrogen evolution reaction

1. Introduction

Electrochemical water splitting has been an effective approach to generate sustainable and clean H₂ energy [1–4]. To accelerate the production of hydrogen, numerous endeavors have been attempted to explore advanced and stable electrocatalysts that lower the overpotential of OER and HER [5–7]. Thus far, precious metal-based catalysts (such as Ir-based for OER and Pt-based for HER) are still the most effective oxygen electrocatalysts, but scarcity and high costs have largely hindered commercial applications. Hence, numerous research endeavors have been devoted to finding non-precious metal alternatives, including transition metal phosphates [8], nitrides [9], dichalcogenides [10,11], and borides [12–14], which showed exceptional activity and stability for OER and HER. Among them, cobalt selenide is a promising catalyst candidate. Its intrinsic metallic properties lead it to have higher electrical conductivity and more active edge sites [15–17]. For example, Lan et al. fabricated CoSe₂ spheres via a facile hydrothermal process and the prepared samples exhibited an excellent electrochemical activity toward OER and HER [18].

In order to achieve higher electrochemical catalytic properties and stability for CoSe₂, utilizing a nitrogen-doped carbon material as a supporter to enable CoSe₂ particles is a feasible and widely used method. Metal organic frameworks (MOFs) have shown considerable advantages as precursors, and the well-organized framework consists of inorganic metal ions or clusters and N-containing organic ligands that can be converted in situ to nitrogen-doped carbon through pyrolysis [19–22]. Dong et al. found that pyrolysis and selenization

of in situ grown zeolitic imidazolium framework-67 (ZIF-67) can homogeneously anchor CoSe₂ nanoparticles (CoSe₂/CF) to carbon fiber paper and the obtained CoSe₂/CF shows excellent long-term stability and electrocatalytic properties [23]. However, metal sites in the MOF usually induce shrinkage agglomeration during high temperature calcination, which would disrupt the structure of the MOF and prevent exposure of the active site. Jiao et al. developed the SiO₂ as templates to inhibit the Fe agglomeration during pyrolysis [24]. Notably, compared with the bulk counterparts, two-dimensional (2D) MOF nanosheets are being increasingly studied in electrocatalysis due to their highly exposed active sites [25], large surface area [26], and enhanced conductivity [27]. To date, synthetic methods of 2D MOF nanosheet preparation usually depends on physical exfoliation and chemical vapor deposition (CVD). Tang et al. prepared a series of ultrathin Ni/Co MOF nanosheets with unsaturated coordination metal active sites by a simple ultrasonic method, and demonstrated that ligand-unsaturated metal atoms are the main active centers of electrocatalytic OER. Nevertheless, how to avoid the aggregation and restacking of exfoliated nanosheets is still a challenge. Wurster et al. engineered heterobimetallic catalysts via CVD and the obtained nanosheets exhibited 300 mV overpotential and high turnover frequencies for OER [28]. Considering the low-yield of traditional methods, Huang et al. developed a salt-template confined method to prepare ultrathin ZIF-67 nanosheets [29]. The Co, N co-doped ultra-thin graphene nanosheets exhibited better electrocatalytic performance than commercial Pt/C catalysts. Thus, there is an urgent demand to develop a simple and cost-effective approach for designing 2D MOF derived efficient electrocatalysts with large surface area and fast mass transfer.

Here, CoSe₂-NC nanosheet electrocatalysts were prepared using NaCl as a template. Inorganic salt has excellent chemical stability and a smooth surface, which is suitable as a template to build 2D structures. During pyrolysis, the outer layer of ZIF-67 served as the nitrogen source and carbon source for the in situ synthesis of nitrogen-doped carbon. The resultant compounds showed a higher electrochemically active surface area (ECSA) and stability. It also displays excellent OER and HER activity. In addition, the water-splitting cell, based on CoSe₂-NC bifunctional catalysts, shows good electrochemical performance, demonstrating that the catalysts with 2D MOF-derived nanosheet structures have great potential for practical applications.

2. Results and Discussion

2.1. Characterization of CoSe₂-NC NSs

Commercial NaCl powder was selected as template for the one-step synthesis of 2D CoSe₂ nanosheets. A schematic diagram of the material fabrication process is shown in Figure 1. First, NaCl powder was mixed with cobalt (II) nitrate and 2-methylimidazole precursors. After vigorous grinding, the organic ligands of imidazole coordinate to Co²⁺ at room temperature (Figure S1). It should be noted that, in this approach, the excess of NaCl is used to avoid the aggregation of Co during selenization. The EDS images of the NaCl@MOF (Figure S2) show the uniform growth of ZIF-67 on the NaCl surface. SEM image of the 2D MOF which removed the salt template (Figure S3) evidenced that the salt template successfully synthesized ZIF-67 nanosheets. The diffraction peaks of NaCl@MOF and NaCl@CoSe₂ (Figure S4) both match the NaCl crystallinity, and no other peaks were displayed, indicating that the formed ZIF-67 layer was relatively thin. Afterwards, the ZIF-67 shell and Se powder were calcined at 750 °C in a N₂-protected tube furnace and converted to CoSe₂-NC.

To further expand the applicability of the synthetic method, amorphous nanosheets with a thickness of about 3.6 nm were effectively corroborated by XRD, Raman, and AFM. As shown in Figure 2, the nanosheets showed no obvious diffraction peaks, indicating that CoSe₂-NC NSs are amorphous materials [30]. The graphite (002) peak in CoSe₂-NC NSs implied graphitization. Compared with the 3D counterpart, the diffraction peaks of CoSe₂-NC NPs were a perfect match with the simulated patterns (JCPDS, No. 09-0234), suggesting high crystallinity. As is known, the activity and number of exposed active

sites directly affect the activity of electrocatalysts. Compared to its crystalline counterpart, the non-crystalline structure possesses more unsaturated coordination sites and effective active sites. Raman spectroscopy (Figure 2b) showed two peaks. The peak at 1355 cm^{-1} was due to the disordered sp^3 carbon (D-band) and the peak at 1580 cm^{-1} indicated the existence of graphite sp^2 carbon (G-band) [31]. Typically, the G-band corresponds to the lattice characteristics of graphite, while the D-band corresponds to the vibrational modes of carbon atoms at the edges of graphene [32,33]. The degree of carbon disorder is usually estimated by the value of I_D/I_G [34,35]. The calculated ratios of I_D/I_G were 1.63 for $\text{CoSe}_2\text{-NC NSs}$ and 1.26 for $\text{CoSe}_2\text{-NC NPs}$. The results showed that the salt template prepared nanosheets with a higher I_D/I_G than the 3D structure, which indicated that the nanosheets had abundant defects and were considered catalytically active sites [36,37]. The AFM images (Figure 2c) were also used to evaluate the thickness of $\text{CoSe}_2\text{-NC NSs}$. As revealed in the AFM images, Figure 2d suggests that $\text{CoSe}_2\text{-NC NSs}$ exhibited ultrathin nanosheets with a thickness of 3.6 nm. Ultrathin nanosheets can expose abundant catalytic active sites during the OER and HER multiphase reaction interface.

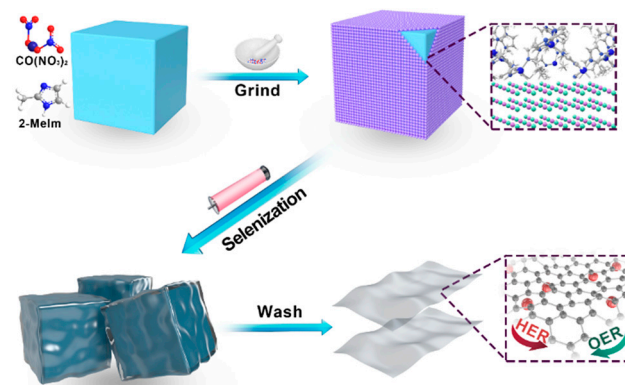


Figure 1. Schematic illustration of the overall synthetic procedure of $\text{CoSe}_2\text{-NC NSs}$.

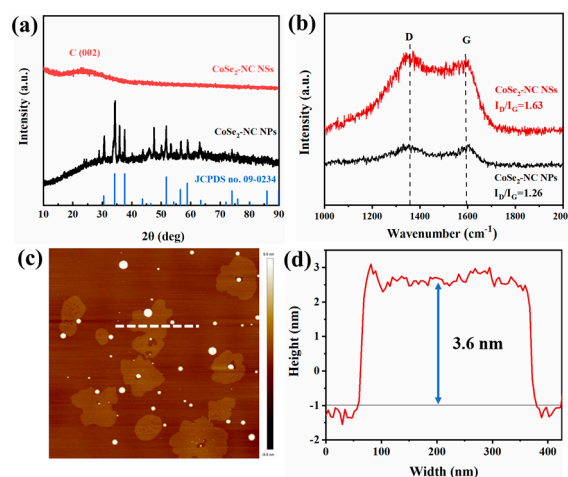


Figure 2. (a) XRD pattern of $\text{CoSe}_2\text{-NC NSs}$ and $\text{CoSe}_2\text{-NC NPs}$, (b) Raman spectra of $\text{CoSe}_2\text{-NC NSs}$ and $\text{CoSe}_2\text{-NC NPs}$, (c,d) AFM image of $\text{CoSe}_2\text{-NC NSs}$.

To obtain more details of the structure, the morphologies of the $\text{CoSe}_2\text{-NC NSs}$ were observed using TEM. Figure 3a demonstrates that $\text{CoSe}_2\text{-NC}$ was in a sheet-like morphology. According to Figure S6, the dodecahedral shapes of the $\text{CoSe}_2\text{-NC NPs}$ could be well preserved, with dimensions of around 500 nm. Figure 3b shows that the CoSe_2 particles were densely interconnected with the graphene layers. HRTEM showed that the interplanar distance of the lattice fringes was 0.258 nm, corresponding to the (111) plane of CoSe_2 , and 0.35 nm, corresponding to the (002) plane of graphite [38]. Moreover, the elemental mapping images (Figure 3d–h) verified that abundant C and N distributed throughout the

entire sample. The results indicated the CoSe₂ particles were encapsulated in nitrogen-doped carbon layers, which can induce greater stability during the corrosion of electrolytes.

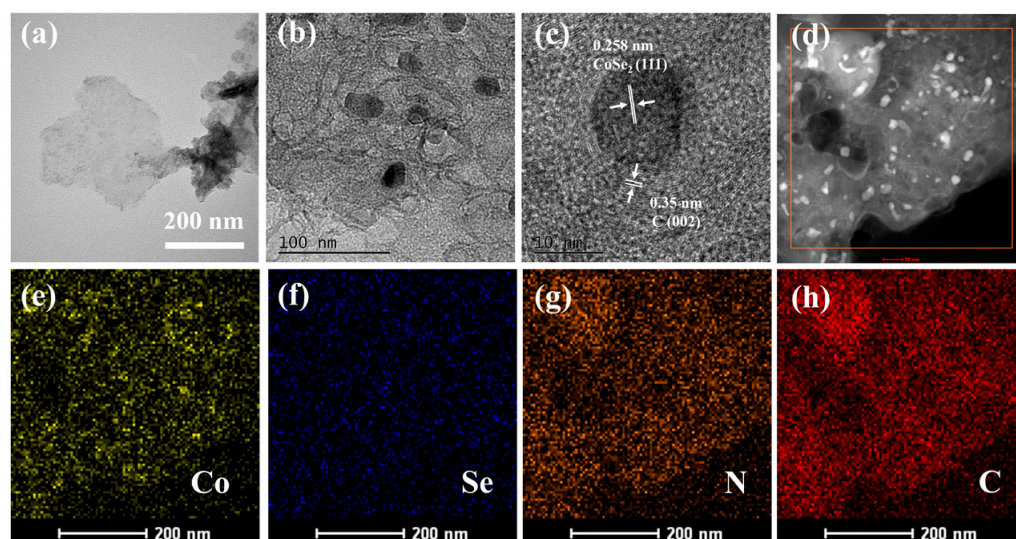


Figure 3. (a,b) TEM images of the CoSe₂-NC NSs, (c) HRTEM image of the CoSe₂-NC NSs, (d–h) EDS elemental mapping images of CoSe₂-NC NSs.

2.2. Electronic States of CoSe₂-NC NSs

The elemental compositions of CoSe₂-NC NSs were determined by XPS analysis. The XPS spectrum of Co 2p (Figure 4a) can be divided into Co 2p_{3/2} and Co 2p_{1/2}, which were located at 780.7 and 796.6 eV, and the corresponding satellite peaks were at 786.5 eV and 803.2 eV, respectively. These results indicated the presence of Co²⁺ [39]. The measured binding energy of 778.1 eV, relative to the reported binding energy of metallic Co, indicated that the Co on the catalyst surface was oxidized by Se elements [40]. The measured binding energy of 778.1 eV, relative to the reported binding energy of metallic Co, indicated that Co on the catalyst surface was oxidized by Se elements [41]. Pyridinic N and pyrrolic N can both coordinate with Co, so the peaks at 781.8 and 797.8 eV could be assigned to Co-N structures. In the Se spectral region (Figure S10), the two main characteristic peaks of CoSe₂-NC NSs were located at 54.9 and 55.8 eV, which correspond to the Se 3d_{5/2} and 3d_{3/2} orbitals of Se^{2−}, respectively. In addition, the peak located at 60.1 eV indicated the presence of Se-O bonds on the surface. These relative peaks were from Co²⁺ coordinated to Se ions. The peaks of the C 1s spectrum (Figure 4b) at 284.3 eV, which could be assigned to sp² hybridized carbons, and the peaks at 285.4 and 286.9 eV, due to the N-sp² C and N-sp³, demonstrated the successful doping of N into carbon [42]. Notably, the C 1s spectrum of CoSe₂-NC NPs (Figure 4e) could be only deconvoluted into sp² C and N-sp² C. The presence of the sp³ carbon atoms could disrupt the long-range order of the carbon network and were considered to be defective sites in the sp² carbon matrix. The analysis results in Figure 4c show that the characteristic peaks were located at 398.4, 399.6, and 400.7 eV, corresponding to pyridine N, pyrrole N, and graphite N, further confirming the formation of nitrogen-doped graphitic carbon [43]. Figure 4f shows that the pyrrole N peak of cobalt selenide is prominent, indicating that the three-dimensional structure of CoSe₂-NC NPs has more pyrrole N in the annealing process. However, the pyrrole N species are nitrogen atoms in a five-membered C-N heterocyclic structure, which are unstable due to their special structure. These results indicated that the CoSe₂ particles were successfully encapsulated into N-doped carbon (NC) matrix. For increasing active sites, doping the carbon matrix with nitrogen heteroatoms is useful. In addition, the nitrogen formed a strong bond with the internal atoms, which resulted in a high stability of the composite [44].

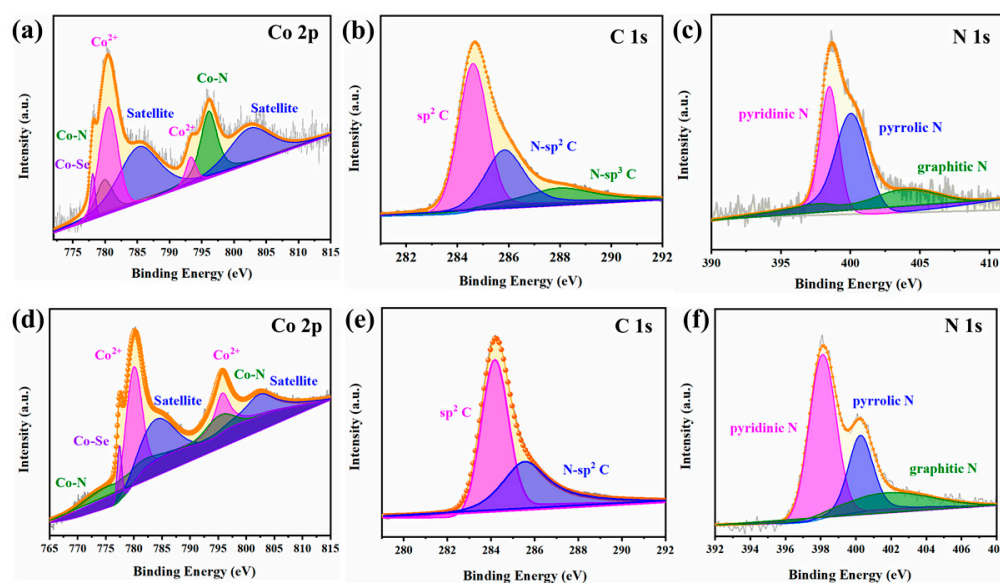


Figure 4. (a–c) High-resolution XPS spectra of Co 2p, C 1s and N 1s, for the CoSe₂-NC NSs, (d–f) High-resolution XPS spectra of Co 2p, C 1s, and N 1s for the CoSe₂-NC NPs.

2.3. Electrochemical Performance of CoSe₂-NC NSs

The performance of the OER catalysts were tested in 1 M KOH electrolyte. Typically, the potential of the OER at a current density of 10 mA·cm^{−2} is defined as E_{j=10}. The overpotential is subtracted from the E_{j=10} value of the catalyst by 1.23 V. Figure 5a shows the LSV curves for the OER electrocatalytic performance of CoSe₂-NC NSs, CoSe₂-NC NPs, Co-NC, and IrC using a three-electrode configuration. It is a remarkable that the CoSe₂-NC NSs catalyst exhibited an advanced electrocatalytic activity with an overpotential of 246.7 mV, which is evidently better than that of other catalysts, and is even better than IrC (overpotential = 322.8 mV. The Tafel slope of CoSe₂-NC NSs (72.66 mV·dec^{−1}), CoSe₂-NC NPs (91.97 mV·dec^{−1}), Co-NC (78.03 mV·dec^{−1}), and IrC (90.67 mV·dec^{−1}) are displayed in Figure 5b, and suggests that the morphology of the nanosheet directly increases OER activity. Figure 5c shows the overpotential and Tafel slope data of CoSe₂-NC NSs, CoSe₂-NC NPs, Co-NC, and IrC, indicating that CoSe₂-NC NSs have a far greater OER activity than the other samples. The synergistic effect of CoSe₂ and nitrogen-doped carbon contributes to the outstanding OER activity of CoSe₂-NC NSs. Meanwhile, the stability of the catalyst in electrolyte is a key parameter. As shown in Figure 5d, CoSe₂-NC NSs maintained a stable voltage at current density of 10 mA·cm^{−2} after 132 h of chronopotentiometry testing. The outstanding stability mainly originates from the carbon layers protecting the CoSe₂ nanoparticles.

To assess the HER electrocatalytic activity, CoSe₂-NC NSs, CoSe₂-NC NPs, Co-NC, and PtC were explored in 1 M KOH electrolyte using a three-electrode cell. The CoSe₂-NC NSs catalyst exhibited favorable electrocatalytic properties (Figure 6a). The overpotential of CoSe₂-NC NSs was only 75.6 mV, which is much less than CoSe₂-NC NPs (121.1 mV) and Co-NC (133.6 mV). The Tafel slope of 114.4 mV·dec^{−1} was measured for CoSe₂-NC NSs, and was lower than CoSe₂-NC NPs and Co-NC. As shown in Figure 6d, the ECSA of CoSe₂-NC NSs (13.43 mF·cm^{−2}) was much higher than CoSe₂-NC NPs (10.48 mF·cm^{−2}), Co-NC (12.08 mF·cm^{−2}), and IrC (7.25 mF·cm^{−2}). The results indicate that there were more active sites in CoSe₂-NC NSs and that these abundant active sites were derived from the low-dimensional nanosheets, which have a large specific surface area (Figure S10). To further explore the HER kinetics, EIS measurements (Figure 6e) were taken in 1.0 M KOH. The smaller semicircle diameter of CoSe₂-NC NSs (1.21 Ω) demonstrates a smaller charge-transfer resistance. This reveals that the electrochemical impedance of the CoSe₂-NC NSs is much lower, which can effectively accelerate the charge transfer between the electrocatalyst and electrolyte interface.

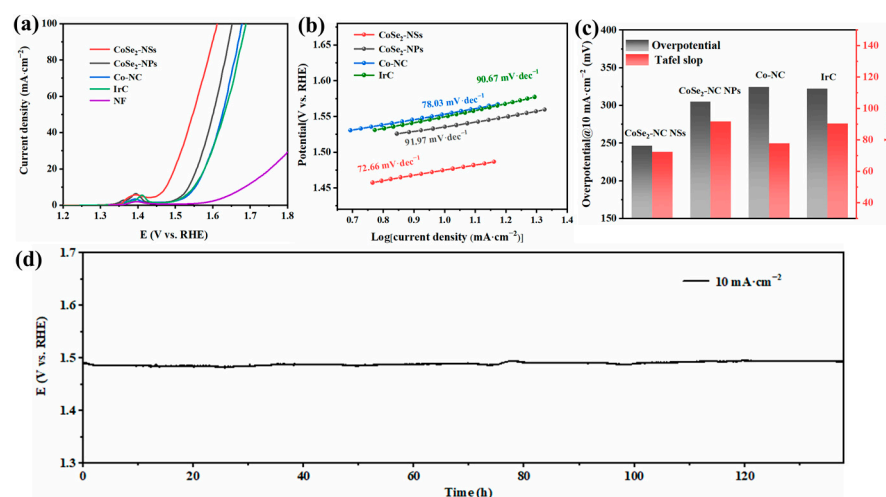


Figure 5. (a) LSV curves of CoSe₂-NC NSs, CoSe₂-NC NPs, Co-NC, IrC and NF, (b) Tafel slopes of CoSe₂-NC NSs, CoSe₂-NC NPs, Co-NC and IrC, (c) Tafel slopes and overpotential for OER of CoSe₂-NC NSs, CoSe₂-NC NPs, Co-NC and IrC, (d) stability of CoSe₂-NC NSs in chronopotentiometry curve.

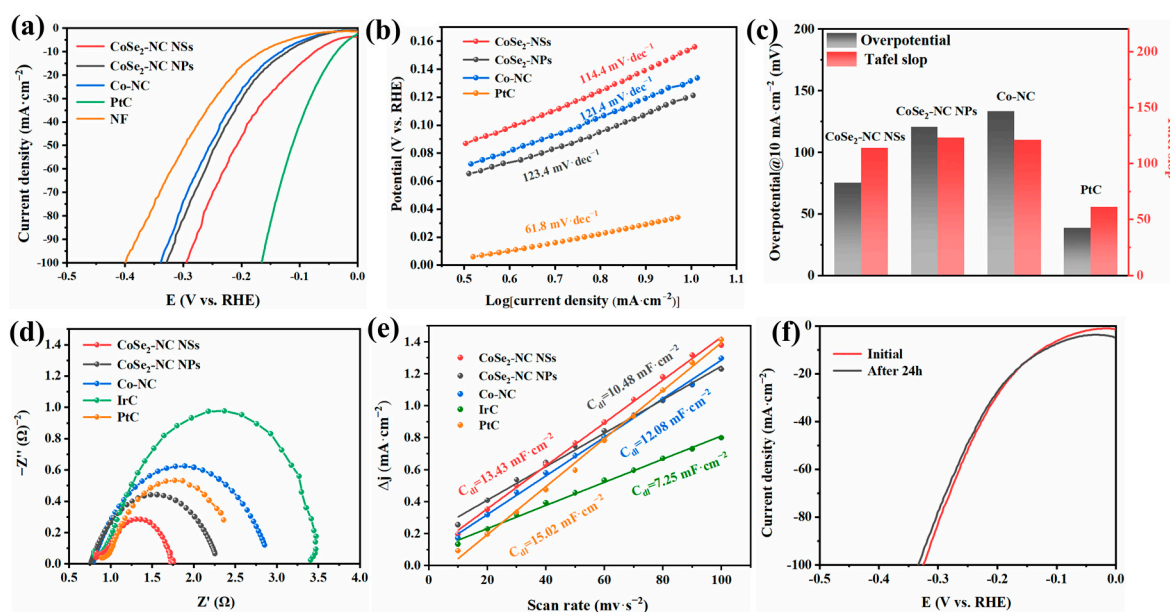


Figure 6. (a) LSV curves of CoSe₂-NC NSs, CoSe₂-NC NPs, Co-NC, PtC and NF. (b) Tafel slopes of CoSe₂-NC NSs, CoSe₂-NC NPs, Co-NC and PtC. (c) Tafel slopes and overpotential for HER of CoSe₂-NC NSs, CoSe₂-NC NPs, Co-NC and PtC. (d) C_{dl} curves of CoSe₂-NC NSs, CoSe₂-NC NPs, Co-NC, PtC and IrC. (e) EIS test of CoSe₂-NC NSs, CoSe₂-NC NPs, Co-NC, PtC and IrC. (f) LSV curves initial and after 24 h of continuous CV.

In view of the high performance of the prepared CoSe₂-NC NS electrodes for both OER and HER, a two-electrode cell was set up in 1.0 M KOH using CoSe₂-NC NSs as both the cathode and anode. Figure 7a shows the LSV curves of the CoSe₂-NC NSs, which exhibited excellent overall water splitting activity. For the LSV measurement, the cell voltage of the CoSe₂-NC NSs-based water splitting cell at 10 mA·cm⁻² was only 1.54 V, even below that of the PtC || IrC (1.65 V). After 12 h of constant current testing, the catalyst voltage showed no obvious change, which indicated that the catalyst has good electrochemical activity and stability. The salt template promotes the formation of ultrathin nanosheets. The strong bonding of CoSe₂ and the carbon layer ensures the immobilization of the active component, which is beneficial for improving the durability of the electrochemical process for overall water splitting.

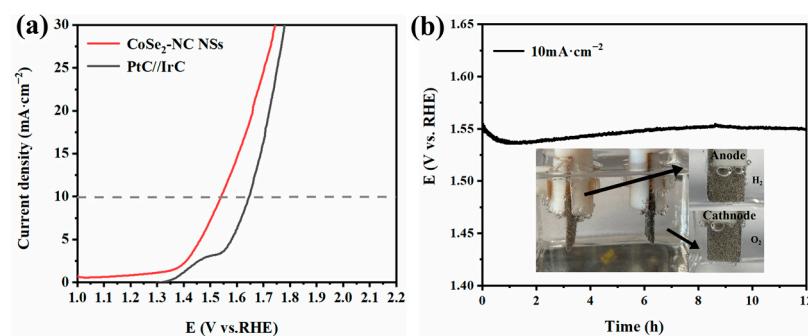


Figure 7. (a) LSV curves of the CoSe₂-NC NSs and PtC | IrC with a two-electrode system in alkaline electrolyte. (b) Stability of CoSe₂-NC NSs in 1 M KOH at a current density of 10 mA·cm⁻² over 12 h.

3. Materials and Methods

3.1. Materials and Reagents

Co(NO₃)₂·6H₂O (99%), 2-methylimidazole (2-MeIm, 99%), and selenium powder (99.999%) were supplied by Sigma Aldrich (Missouri, USA). Methanol (99%), ethanol (99%), NaCl and KOH were bought from Chemical Reagent (Guangzhou, China). Nafion solution (5%) was purchased from Hesen (Shanghai, China). The ultrapure water (18 MΩ) used in the experiments was prepared using Hhitech equipment (Shanghai, China). Commercial catalysts (Pt/C, 20 wt%, Ir/C, 5 wt%) for comparison were bought from Macklin (Shanghai, China). All chemicals in the experiment were used directly without further purification.

3.2. Synthesis of Co-NC

A total of 1.97 g 2-methylimidazole was dissolved in a mixed solvent with 20 mL of methanol and 20 mL of ethanol. Meanwhile, 0.87 g Co(NO₃)₂·6H₂O was dissolved in another mixed solvent with 20 mL of methanol and 20 mL of ethanol. Then, the above two solutions were mixed under continuous stirring for 1 min and the final solution was kept at room temperature for 24 h. The resultant purple ZIF-67 precipitate was collected using centrifugation and washed several times with ethanol and ultrapure water and then dried in an oven at 60 °C for 12 h.

3.3. Synthesis of CoSe₂-NC NPs

The ZIF-67 particles and 0.1 g Se powder were dispersed in ceramic boats and the temperature in the furnace was raised to 750 °C at a rate of 2 °C·min⁻¹. After that, the furnace was naturally cooled to room temperature. During the pyrolysis, the furnace was under a N₂ atmosphere. To remove the free metal ions, the prepared black powder product was stirred in 0.5 M hydrochloric acid for 12 h. The samples were collected by centrifugation and washed repeatedly with deionized water and then dried at 80 °C.

3.4. Synthesis of CoSe₂-NC NSs

First, 0.363 g Co(NO₃)₂·6H₂O, 0.411 g 2-methylimidazole and 4 g NaCl salt were mixed and ground in a mortar. After that, ZIF-67 coated on NaCl nanocrystals surface (denoted as NaCl@ZIF-67) was obtained. Then, the as-obtained product and 0.1 g Se powder were selenized under a N₂ atmosphere at 750 °C for 2 h, 2 °C·min⁻¹. After being cooled, the powders were washed with 0.5 M HCl solution and deionized water to remove the NaCl templates and impurities.

3.5. Material Characterization

X-ray diffraction (XRD) was carried out using an X' Pert PRO with a Cu Ka radiation diffractometer ($k = 1.5418 \text{ \AA}$). Raman spectra were detected using a Horiba JobinYvon, LabRAM HR800. Atomic force microscopy (AFM) images were determined using a Bruker Dimension ICON. The morphology and structures of the catalysts were measured by scanning electron microscopy (SEM, ZEISS Sigma 300) and an energy dispersive spectrom-

eter (EDS). Transmission electron microscopy (TEM) was performed on a Tecnai G2 F30. The Brunauer–Emmet–Teller (BET) surface areas were measured at 77 K using a NOVA 2000 (Quantachrome, Boynton Beach, FL, USA). X-ray photoelectron spectroscopy (XPS) analyses were performed on a Thermo Scientific K-Alpha with Al Ka radiation.

3.6. Electrochemical Performance

Prior to catalyst loading, nickel foam was acid washed to remove oxide impurities from the surface. The 1 cm × 1 cm nickel foam was immersed in 1 M HCl solution and sonicated for 15 min, and then sonicated in deionized water and anhydrous ethanol solution for 15 min. A homogeneous suspension of the catalyst was formed by dispersing 2 mg of catalyst in a mixture of 250 µL of deionized water, 700 µL of anhydrous ethanol, and 50 µL of Nafion with sonication for 1 h. Then, the suspension (100 µL) was dripped onto the pre-polished nickel foam and dried in a vacuum at 60 °C. The mass loading of the active materials in this paper was, on average, 0.2 mg·cm^{−2}.

Electrocatalytic OER and HER measurements were tested at room temperature in a standard three-electrode setup, which was carried out on a CHI 760E electrochemical workstation (Chenhua, Shanghai, China). Nickel foam with electrocatalyst, graphite rod, and Ag/AgCl electrode filled with saturated KCl were selected as the working electrode, counter electrode, and reference electrode, respectively. The electrolyte used was KOH solution at 1.0 M (pH 13.6). To ensure the O₂/H₂O equilibrium at 1.23 V vs. RHE, all electrochemical experiments were performed in the O₂ saturated condition. Linear sweep voltammetry (LSV) and cyclic voltammetry (CV) curves were measured at a scan rate of 5 mV·s^{−1}. Tafel slopes were calculated according to Tafel equation: $\eta = a + b \log(j)$. CV was measured in the potential window of a non-Faraday process at different scan rates from 10 to 100 mV·s^{−1}. The slope C_{dl} was obtained by fitting the current density versus scan rate as a linear relationship. To further investigate electrocatalytic kinetics, electrochemical impedance spectroscopy (EIS) measurements were carried out in the frequency range 10 kHz to 10 mHz. The stability of the catalyst was determined by chronopotentiometry measurements at $j = 10 \text{ mA} \cdot \text{cm}^{-2}$.

4. Conclusions

In summary, CoSe₂ nanoparticles embedded into nitrogen-doped carbon nanosheets were successfully synthesized using a salt-template strategy. The ultrathin nanosheets formed by the salt could effectively avoid self-aggregation of the CoSe₂ particles, while the 2D structure could promote more efficient electron transfer between reactants and catalysts. In comparison with CoSe₂-NC NPs and bare Co-NC NPs, the CoSe₂-NC NSs exhibited remarkable OER and HER properties. In addition, the robust structure can maintain excellent stability during the reaction. As a result, this work provides a new strategy for the design of CoSe₂-based bifunctional electrocatalysts with excellent catalytic performance and long-term stability.

Supplementary Materials: The following supporting information can be downloaded at: <https://www.mdpi.com/article/10.3390/ijms23095239/s1>.

Author Contributions: Writing, review and editing, H.C., methodology, H.L., formal analysis, L.L. and X.N., data curation, K.X., visualization and supervision, J.H., funding acquisition, L.C. All authors have read and agreed to the published version of the manuscript.

Funding: This work was supported by the National Natural Science Foundation of China (No. 51962032) and the Autonomous Region Graduate Student Innovation Program (XJ2021G141).

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: Not applicable.

Conflicts of Interest: The authors declare no conflict of interest.

References

- Wang, Z.; Li, C.; Domen, K. Recent developments in heterogeneous photocatalysts for solar-driven overall water splitting. *Chem. Soc. Rev.* **2019**, *48*, 2109–2125. [\[CrossRef\]](#) [\[PubMed\]](#)
- Li, H.; Chen, L.; Jin, P.; Li, Y.; Pang, J.; Hou, J.; Peng, S.; Wang, G.; Shi, Y. NiCo₂S₄ microspheres grown on N, S co-doped reduced graphene oxide as an efficient bifunctional electrocatalyst for overall water splitting in alkaline and neutral pH. *Nano Res.* **2022**, *15*, 950–958. [\[CrossRef\]](#)
- Pang, J.X.; Fu, H.H.; Kong, W.W.; Jiang, R.; Ye, J.H.; Zhao, Z.Y.; Hou, J.; Sun, K.S.; Zheng, Y.; Chen, L. Design of NiCo₂O₄ nanoparticles decorated N, S co-doped reduced graphene oxide composites for electrochemical simultaneous detection of trace multiple heavy metal ions and hydrogen evolution reaction. *Chem. Eng. J.* **2022**, *433*, 133854. [\[CrossRef\]](#)
- Fan, C.; Zhai, X.; Chen, L.; Peng, S.; Jiang, R.; Yu, J.; Li, Y.; Zhang, Y.; Kong, W.; Ge, G.; et al. Synthesis and electrocatalytic mechanism of ultrafine MFe₂O₄ (M: Co, Ni, and Zn) nanocrystallites: M/Fe synergistic effects on the electrochemical detection of Cu(II) and hydrogen evolution reaction performances. *J. Mater. Chem. A* **2021**, *9*, 22277–22290. [\[CrossRef\]](#)
- Ye, J.H.; Zhai, X.W.; Chen, L.; Guo, W.; Gu, T.T.; Shi, Y.L.; Hou, J.; Han, F.; Liu, Y.; Fan, C.C.; et al. Oxygen vacancies enriched nickel cobalt based nanoflower cathodes: Mechanism and application of the enhanced energy storage. *J. Energy Chem.* **2021**, *62*, 252–261. [\[CrossRef\]](#)
- You, B.; Sun, Y. Innovative Strategies for Electrocatalytic Water Splitting. *Acc. Chem. Res.* **2018**, *51*, 1571–1580. [\[CrossRef\]](#)
- Wang, D.; Chang, Y.-X.; Li, Y.-R.; Zhang, S.L.; Xu, S.L. Well-dispersed NiCoS₂ nanoparticles/rGO composite with a large specific surface area as an oxygen evolution reaction electrocatalyst. *Rare Metals* **2021**, *40*, 3156–3165. [\[CrossRef\]](#)
- He, P.; Yu, X.Y.; Lou, X.W. Carbon-Incorporated Nickel-Cobalt Mixed Metal Phosphide Nanoboxes with Enhanced Electrocatalytic Activity for Oxygen Evolution. *Angew. Chem.* **2017**, *56*, 3897–3900. [\[CrossRef\]](#)
- Huang, T.; Chen, Y.; Lee, J.-M. Two-Dimensional Cobalt/N-Doped Carbon Hybrid Structure Derived from Metal–Organic Frameworks as Efficient Electrocatalysts for Hydrogen Evolution. *ACS Sustain. Chem. Eng.* **2017**, *5*, 5646–5650. [\[CrossRef\]](#)
- Sheng, H.; Janes, A.N.; Ross, R.D.; Kaiman, D.; Huang, J.; Song, B.; Schmidt, J.R.; Jin, S. Stable and selective electrosynthesis of hydrogen peroxide and the electro-Fenton process on CoSe₂ polymorph catalysts. *Energy Environ. Sci.* **2020**, *13*, 4189–4203. [\[CrossRef\]](#)
- Kong, D.; Wang, H.; Lu, Z.; Cui, Y. CoSe₂ nanoparticles grown on carbon fiber paper: An efficient and stable electrocatalyst for hydrogen evolution reaction. *J. Am. Chem. Soc.* **2014**, *136*, 4897–4900. [\[CrossRef\]](#) [\[PubMed\]](#)
- Gupta, S.; Patel, M.K.; Miotello, A.; Nainesh, P. Metal Boride-Based Catalysts for Electrochemical Water-Splitting: A Review. *Adv. Funct. Mater.* **2019**, *30*, 1906481–1906509. [\[CrossRef\]](#)
- Guo, F.; Wu, Y.; Chen, H.; Liu, Y.; Yang, L.; Ai, X.; Zou, X. High-performance oxygen evolution electrocatalysis by boronized metal sheets with self-functionalized surfaces. *Energy Environ. Sci.* **2019**, *12*, 684–692. [\[CrossRef\]](#)
- Masa, J.; Weide, P.; Peeters, D.; Sinev, L.; Xia, W.; Sun, Z.; Somsen, C.; Muhler, M.; Schuhmann, W. Amorphous Cobalt Boride (Co₂B) as a Highly Efficient Nonprecious Catalyst for Electrochemical Water Splitting: Oxygen and Hydrogen Evolution. *Adv. Energy Mater.* **2016**, *6*, 1502313–1502323. [\[CrossRef\]](#)
- Zhang, L.; Lu, C.; Ye, F.; Pang, R.; Liu, Y.; Wu, Z.; Shao, Z.; Sun, Z.; Hu, L. Selenic Acid Etching Assisted Vacancy Engineering for Designing Highly Active Electrocatalysts toward the Oxygen Evolution Reaction. *Adv. Mater.* **2021**, *33*, e2007523. [\[CrossRef\]](#) [\[PubMed\]](#)
- Li, K.; Zhang, J.; Wu, R.; Yu, Y.; Zhang, B. Anchoring CoO Domains on CoSe₂ Nanobelts as Bifunctional Electrocatalysts for Overall Water Splitting in Neutral Media. *Adv. Sci.* **2016**, *3*, 1500426. [\[CrossRef\]](#)
- Zhong, W.; Wang, Z.; Gao, N.; Huang, L.; Lin, Z.; Liu, Y.; Meng, F.; Deng, J.; Jin, S.; Zhang, Q.; et al. Coupled Vacancy Pairs in Ni-Doped CoSe for Improved Electrocatalytic Hydrogen Production Through Topochemical Deintercalation. *Angew. Chem.* **2020**, *59*, 22743–22748. [\[CrossRef\]](#) [\[PubMed\]](#)
- Lan, K.; Li, J.; Zhu, Y.; Gong, L.; Li, F.; Jiang, P.; Niu, F.; Li, R. Morphology engineering of CoSe₂ as efficient electrocatalyst for water splitting. *J. Colloid Interface Sci.* **2019**, *539*, 646–653. [\[CrossRef\]](#)
- Duan, J.; Chen, S.; Zhao, C. Ultrathin metal-organic framework array for efficient electrocatalytic water splitting. *Nat. Commun.* **2017**, *8*, 15341. [\[CrossRef\]](#)
- Xu, Y.; Li, B.; Zheng, S.; Wu, P.; Zhan, J.; Xue, H.; Xu, Q.; Pang, H. Ultrathin two-dimensional cobalt–organic framework nanosheets for high-performance electrocatalytic oxygen evolution. *J. Mater. Chem. A* **2018**, *6*, 22070–22076. [\[CrossRef\]](#)
- Li, C.; Zhao, D.-H.; Long, H.L.; Li, M. Recent advances in carbonized non-noble metal–organic frameworks for electrochemical catalyst of oxygen reduction reaction. *Rare Met.* **2021**, *40*, 2657–2689. [\[CrossRef\]](#)
- Zhao, S.; Wang, Y.; Dong, J.; He, C.T.; Yin, H.; An, P.; Zhao, K.; Zhang, X.; Gao, C.; Zhang, L.; et al. Ultrathin metal–organic framework nanosheets for electrocatalytic oxygen evolution. *Nat. Energy* **2016**, *1*, 16184. [\[CrossRef\]](#)
- Sun, C.; Dong, Q.; Yang, J.; Dai, Z.; Lin, J.; Chen, P.; Huang, W.; Dong, X. Metal–organic framework derived CoSe₂ nanoparticles anchored on carbon fibers as bifunctional electrocatalysts for efficient overall water splitting. *Nano Res.* **2016**, *9*, 2234–2243. [\[CrossRef\]](#)
- Jiao, L.; Zhang, R.; Wan, G.; Yang, W.; Wan, X.; Zhou, H.; Shui, J.; Yu, S.H.; Jiang, H.L. Nanocasting SiO₂ into metal-organic frameworks imparts dual protection to high-loading Fe single-atom electrocatalysts. *Nat. Commun.* **2020**, *11*, 2831. [\[CrossRef\]](#) [\[PubMed\]](#)

25. Zheng, Y.R.; Gao, M.R.; Gao, Q.; Li, H.H.; Xu, J.; Wu, Z.Y.; Yu, S.H. An efficient CeO₂/CoSe₂ Nanobelt composite for electrochemical water oxidation. *Small* **2015**, *11*, 182–188. [[CrossRef](#)]
26. Liu, X.; Liu, Y.; Fan, L.Z. MOF-derived CoSe₂ microspheres with hollow interiors as high-performance electrocatalysts for the enhanced oxygen evolution reaction. *J. Mater. Chem. A* **2017**, *5*, 15310–15314. [[CrossRef](#)]
27. Zhang, Y.; Zhang, C.; Guo, Y.; Liu, D.; Yu, Y.; Zhang, B. Selenium vacancy-rich CoSe₂ ultrathin nanomeshes with abundant active sites for electrocatalytic oxygen evolution. *J. Mater. Chem. A* **2019**, *7*, 2536–2540. [[CrossRef](#)]
28. Wurster, B.; Grumelli, D.; Hotger, D.; Gutzler, R.; Kern, K. Driving the Oxygen Evolution Reaction by Nonlinear Cooperativity in Bimetallic Coordination Catalysts. *J. Am. Chem. Soc.* **2016**, *138*, 3623–3626. [[CrossRef](#)]
29. Huang, L.; Zhang, X.; Han, Y.; Wang, Q.; Fang, Y.; Dong, S. In situ synthesis of ultrathin metal–organic framework nanosheets: A new method for 2D metal-based nanoporous carbon electrocatalysts. *J. Mater. Chem. A* **2017**, *5*, 18610–18617. [[CrossRef](#)]
30. Liu, C.; Wang, J.; Wan, J.; Cheng, Y.; Huang, R.; Zhang, C.; Hu, W.; Wei, G.; Yu, C. Amorphous Metal–Organic Framework-Dominated Nanocomposites with Both Compositional and Structural Heterogeneity for Oxygen Evolution. *Angew. Chem.* **2020**, *59*, 3630–3637. [[CrossRef](#)]
31. Chang, S.H.; Danilovic, N.; Chang, K.C.; Subbaraman, R.; Paulikas, A.P.; Fong, D.D.; Highland, M.J.; Baldo, P.M.; Stamenkovic, V.R.; Freeland, J.W.; et al. Functional links between stability and reactivity of strontium ruthenate single crystals during oxygen evolution. *Nat. Commun.* **2014**, *5*, 4191. [[CrossRef](#)] [[PubMed](#)]
32. Ding, J.; Bu, L.; Guo, S.; Zhao, Z.; Zhu, E.; Huang, Y.; Huang, X. Morphology and Phase Controlled Construction of Pt–Ni Nanostructures for Efficient Electrocatalysis. *Nano Lett.* **2016**, *16*, 2762–2767. [[CrossRef](#)] [[PubMed](#)]
33. Wang, T.; Yang, C.; Liu, Y.; Yang, M.; Li, X.; He, Y.; Li, H.; Chen, H.; Lin, Z. Dual-Shelled Multidoped Hollow Carbon Nanocages with Hierarchical Porosity for High-Performance Oxygen Reduction Reaction in Both Alkaline and Acidic Media. *Nano Lett.* **2020**, *20*, 5639–5645. [[CrossRef](#)] [[PubMed](#)]
34. Gui, Y.; Liu, X.; Dou, Y.; Zhang, L.; Al-Mamun, M.; Jiang, L.; Yin, H.; He, C.-T.; Zhao, H. Manipulating the assembled structure of atomically thin CoSe₂ nanomaterials for enhanced water oxidation catalysis. *Nano Energy* **2019**, *57*, 371–378. [[CrossRef](#)]
35. Aboagye, A.; Elbohy, H.; Kelkar, A.D.; Qiao, Q.; Zai, J.; Qian, X.; Zhang, L. Electrospun carbon nanofibers with surface-attached platinum nanoparticles as cost-effective and efficient counter electrode for dye-sensitized solar cells. *Nano Energy* **2015**, *11*, 550–556. [[CrossRef](#)]
36. Zhang, H.; Liu, X.; He, G.; Zhang, X.; Bao, S.; Hu, W. Bioinspired synthesis of nitrogen/sulfur co-doped graphene as an efficient electrocatalyst for oxygen reduction reaction. *J. Power Sources* **2015**, *279*, 252–258. [[CrossRef](#)]
37. Zhao, D.; Zhang, S.; Yin, G.; Du, C.; Wang, Z.; Wei, J. Effect of Se in Co-based selenides towards oxygen reduction electrocatalytic activity. *J. Power Sources* **2012**, *206*, 103–107. [[CrossRef](#)]
38. Wu, X.; Han, S.; He, D.; Yu, C.; Lei, C.; Liu, W.; Zheng, G.; Zhang, X.; Lei, L. Metal Organic Framework Derived Fe-Doped CoSe₂ Incorporated in Nitrogen-Doped Carbon Hybrid for Efficient Hydrogen Evolution. *ACS Sustain. Chem. Eng.* **2018**, *6*, 8672–8678. [[CrossRef](#)]
39. Cai, T.; Zhao, L.; Hu, H.; Li, T.; Li, X.; Guo, S.; Li, Y.; Xue, Q.; Xing, W.; Yan, Z.; et al. Stable CoSe₂/carbon nanodice@reduced graphene oxide composites for high-performance rechargeable aluminum-ion batteries. *Energy Environ. Sci.* **2018**, *11*, 2341–2347. [[CrossRef](#)]
40. Gupta, S.; Patel, N.; Miotello, A.; Kothari, D. Cobalt-Boride: An efficient and robust electrocatalyst for Hydrogen Evolution Reaction. *J. Power Sources* **2015**, *279*, 620–625. [[CrossRef](#)]
41. Kim, J.; Kim, H.; Kim, S.-K.; Ahn, S.-H. Electrodeposited amorphous Co–P–B ternary catalyst for hydrogen evolution reaction. *J. Mater. Chem. A* **2018**, *6*, 6282–6288. [[CrossRef](#)]
42. Zhou, W.; Lu, J.; Zhou, K.; Yang, L.; Ke, Y.; Tang, Z.; Chen, S. CoSe₂ nanoparticles embedded defective carbon nanotubes derived from MOFs as efficient electrocatalyst for hydrogen evolution reaction. *Nano Energy* **2016**, *28*, 143–150. [[CrossRef](#)]
43. Liu, H.; Jin, M.; Zhan, D.; Wang, J.; Cai, X.; Qiu, Y.; Lai, L. Stacking faults triggered strain engineering of ZIF-67 derived Ni–Co bimetal phosphide for enhanced overall water splitting. *Appl. Catal. B Environ.* **2020**, *272*, 118951. [[CrossRef](#)]
44. Xia, B.Y.; Yan, Y.; Li, N.; Wu, H.B.; Lou, X.W.; Wang, X. A metal–organic framework-derived bifunctional oxygen electrocatalyst. *Nat. Energy* **2016**, *1*, 15006. [[CrossRef](#)]