

Supporting Information

Trimetallic MOF-Derived CoFeNi/Z-P NC Nanocomposites as Efficient Catalysts for Oxygen Evolution Reaction

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Structural and electronic property characterization

All scanning electron microscope (SEM) images in this article were taken by Zeiss Gemini300, all transmission electron microscope (TEM) images were taken by FEI TF20, X-ray diffraction (XRD) was taken by SmartLab 9KW, X-ray photoelectron spectroscopy (XPS) was taken by Thermo Scientific K- Alpha, the model of the specific surface area analyzer is JW-BK 200.

Electrochemical measurements

Typically, a mixed solvent made up of 250 μL of ethanol (GCS, 99.5%), 20 μL of a 5 weight percent Nafion solution, and 1 mg of Ketjen Black were used to scatter the catalyst. To get a uniform ink, sonicate the mixture in freezing water for 45 minutes. Then, using the drop-casting method, 10 μL of the ink were applied to a polished glassy carbon electrode (GCE) with a diameter of 5 mm, and left to air dry naturally at room temperature. All electrochemical experiments were performed using a conventional three-electrode cell and the AutoLab 302N electrochemical workstation. In this three-electrode system, the working electrode is a modified GCE electrode, the counter electrode is a graphite rod and the reference electrode is a saturated Ag/AgCl electrode.

Utilizing a scan rate of 5 mV s^{-1} , we employed linear sweep voltammetry (LSV) to analyze the polarization curves for OER activity. All figures were calibrated to a reversible hydrogen electrode (RHE) and corrected for IR correction using the formula $E_{\text{RHE}} = E_{\text{Ag/AgCl}} + 0.0592 \text{ pH} + 0.197$. Using the equation: $\eta(\text{V}) = E_{\text{RHE}} - 1.23 \text{ V}$, the overpotential (η) for the hydrogen evolution process (HER) was derived. Electrochemical impedance spectroscopy (EIS) was carried out by applying an AC voltage with a 10 mV amplitude over a frequency range of 1 to 100,000 Hz. Electrical double-layer specific capacitance (C_{dl}) of the materials was determined by cyclic voltammetry (CV) utilizing non-Faradaic overpotentials and sweep rates ranging from 20 to 100 mV s^{-1} .

Figures:

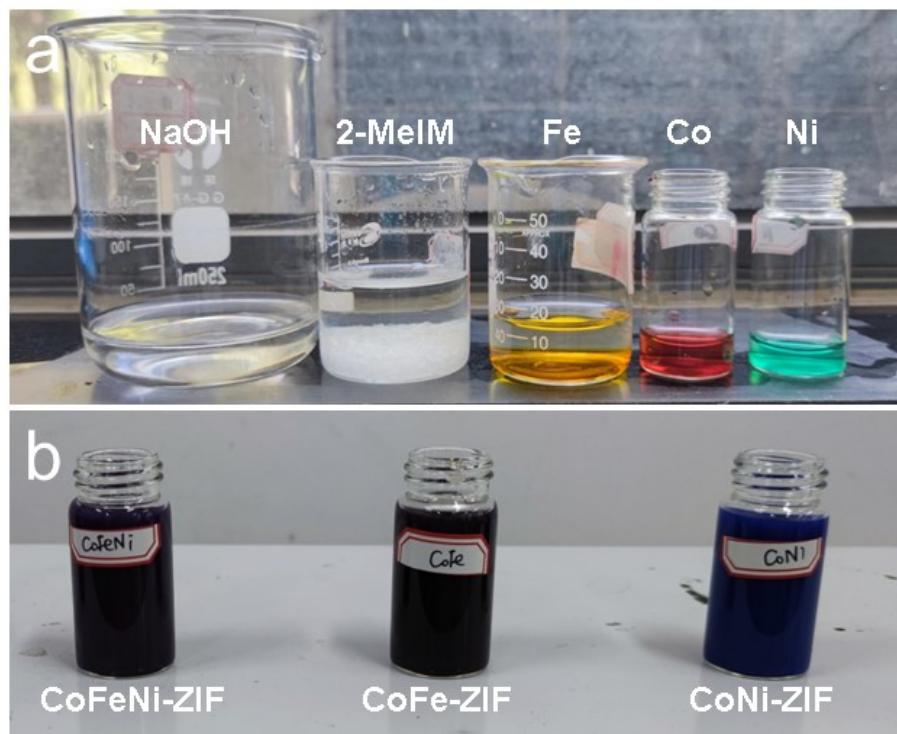


Fig. S1 Optical images of the materials before reaction (a) and after the reaction completed (b)

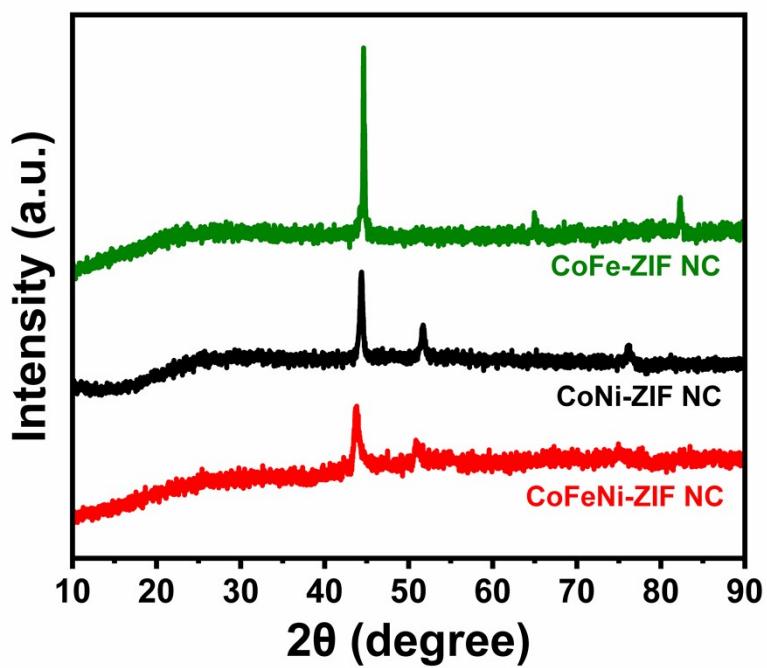


Fig. S2 The XRD patterns of CoFe-ZIF NC, CoNi-ZIF NC and CoFeNi-ZIF NC

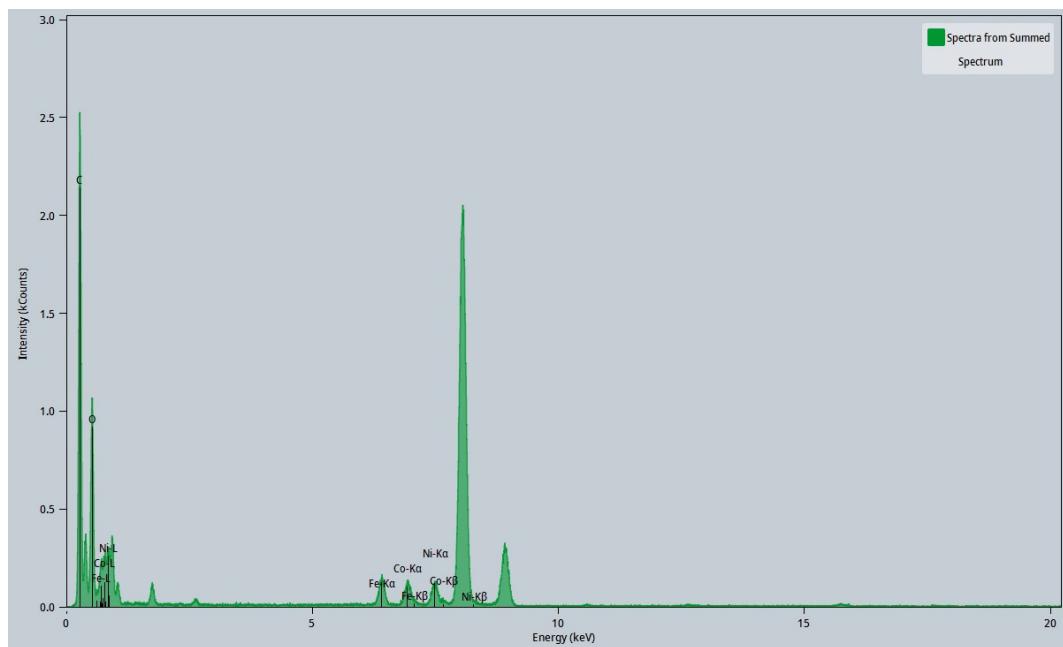


Fig. S3 The EDS composition of CoFeNi-ZIF

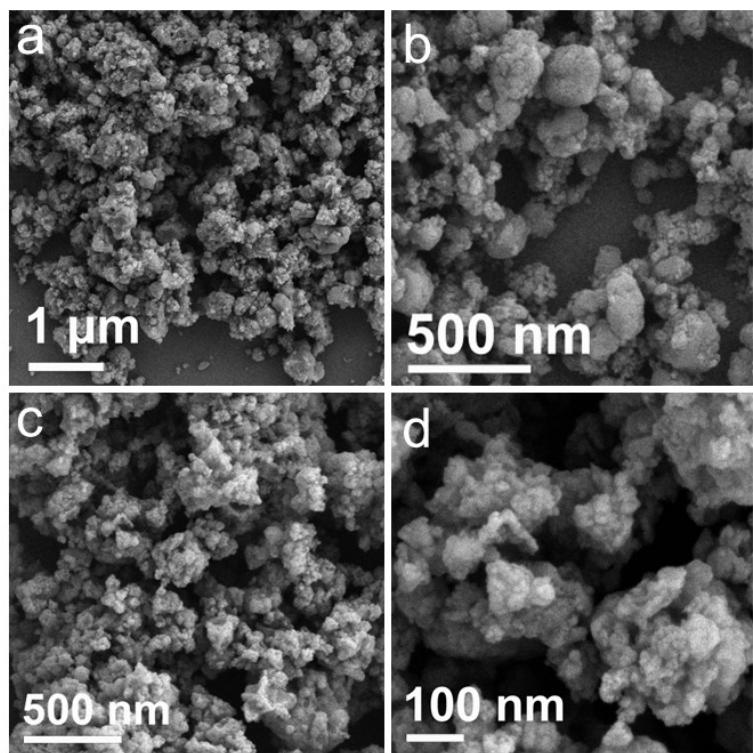


Fig. S4 (a-b) SEM images of CoFeNi-ZIF. (c-d) SEM images of CoFeNi-Z.

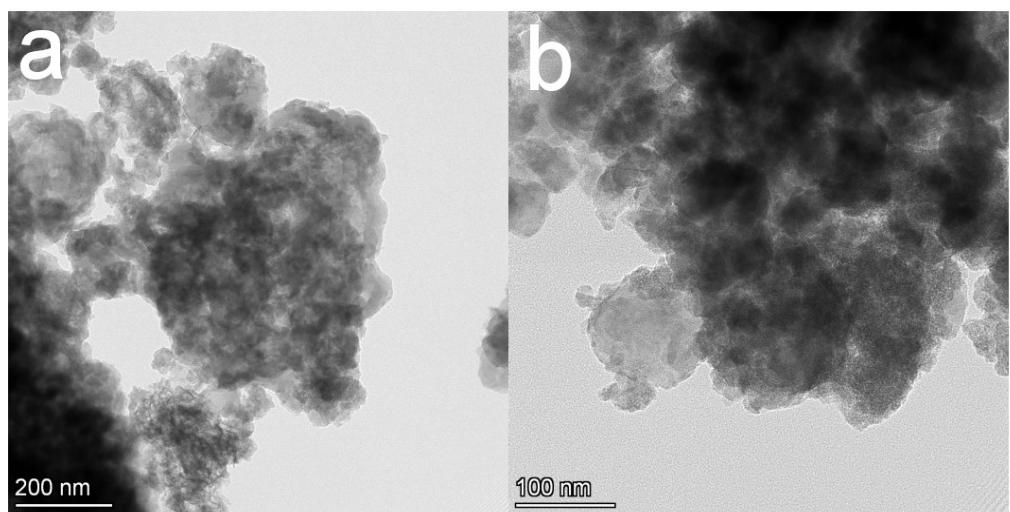


Fig. S5 The TEM images of CoFeNi-Z at 200 nm and 100 nm

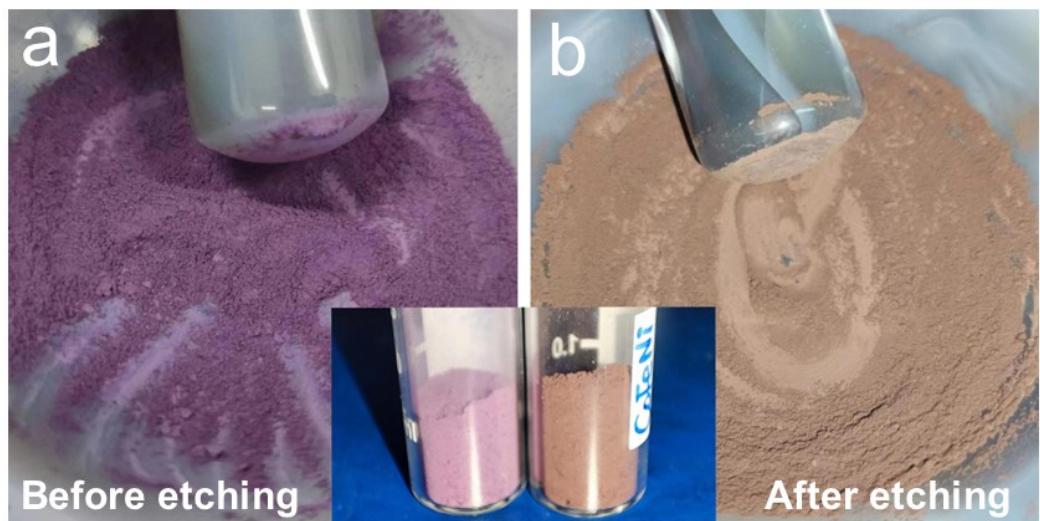


Fig. S6 Optical images of solid samples of CoFeNi-ZIF (a) and CoFeNi-Z (b)

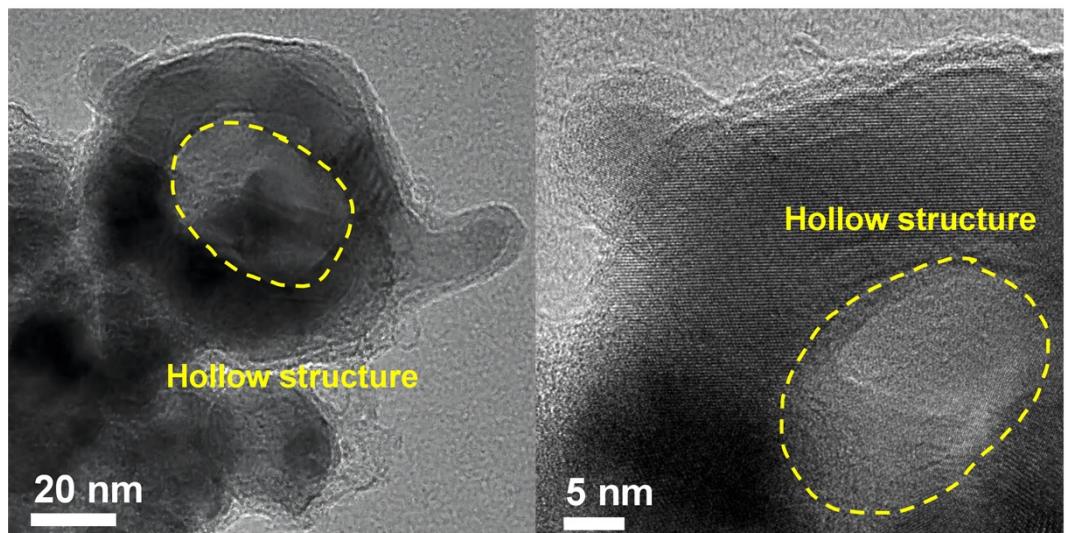


Fig. S7 HRTEM images of CoFeNi/Z-P NC

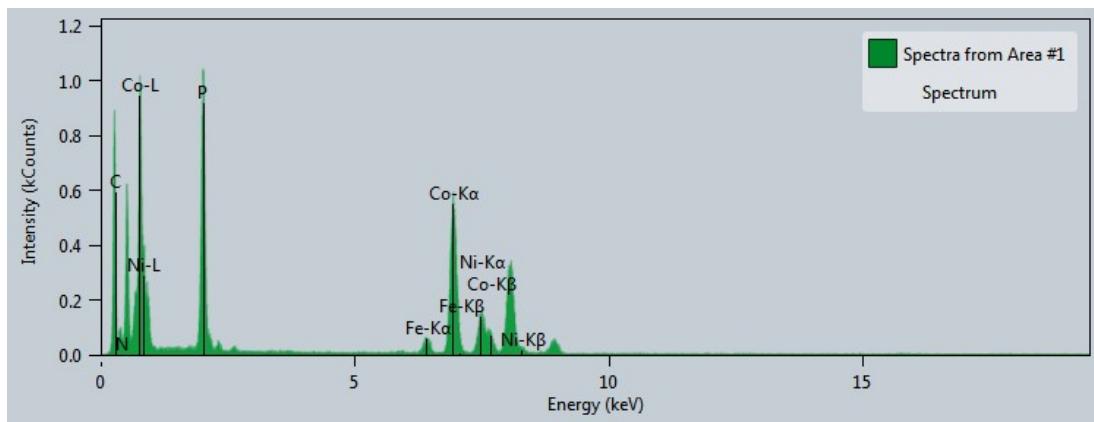


Fig. S8 The EDS composition of CoFeNi/Z-P NC

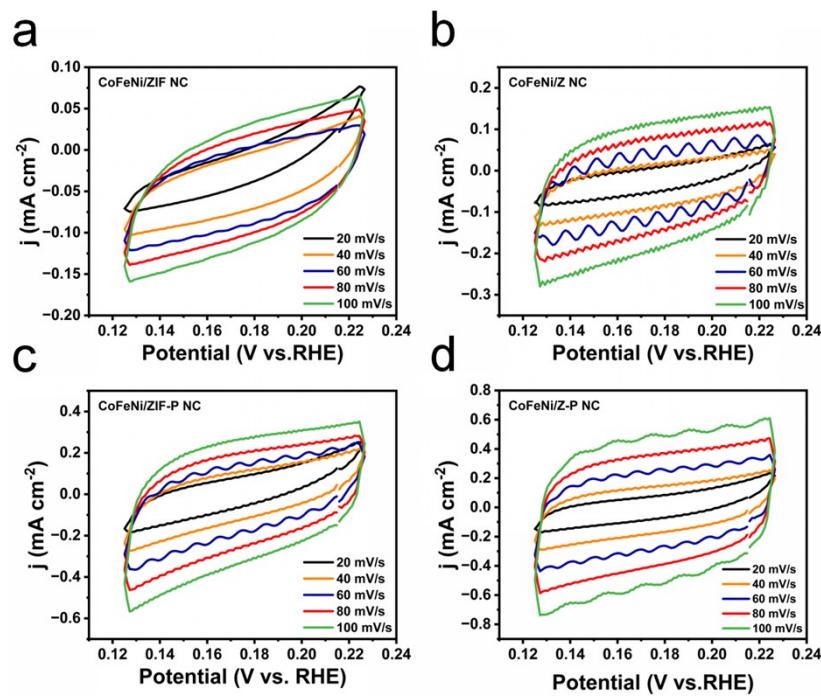


Fig. S9 Cyclic voltammograms of CoFeNi/ZIF NC, CoFeNi/Z NC CoFeNi/ZIF-P NC and CoFeNi/Z P NC within the range of 0.13 to 0.23 V vs RHE with scan rate from 20 to 100 mV s^{-1} in 1 M KOH.

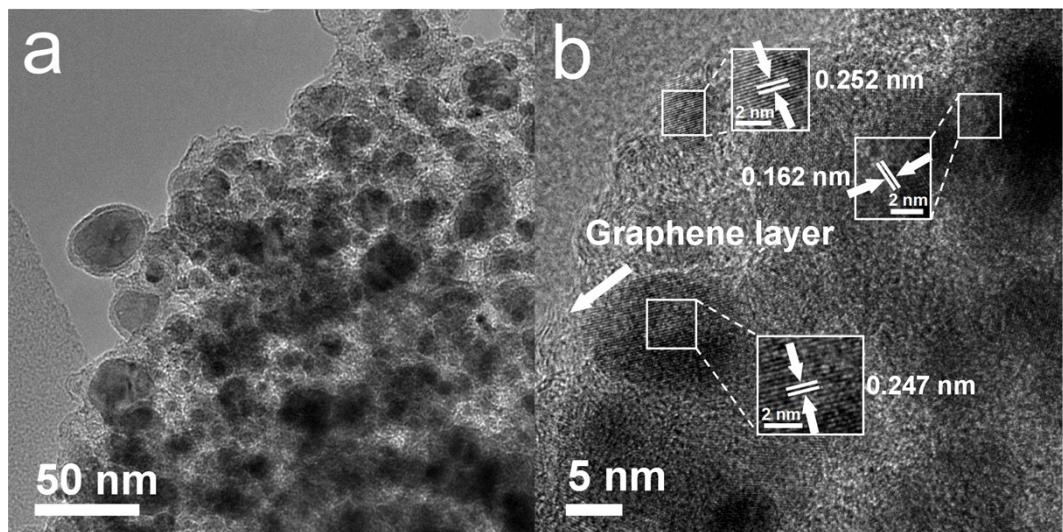


Fig. S10 TEM image and HRTEM image (a and b) of CoFeNi/Z-P NC after stability test

Table S1. Chemical compositions of CoFeNi/Z-P NC by XPS measurement

Sample	C (atom %)	N (atom %)	Co (atom %)	Fe (atom %)	Ni (atom %)	P (atom %)
CoFeNi/Z-P NC	89.88	2.12	1.77	1.63	0.91	3.68

Table S2. Comparison of the OER electrocatalytic performance for different catalysts

Catalyst	Overpotential (10 mA cm ⁻²)	Tafel slope (mV dec ⁻¹)	Ref.
CoFeNi/Z-P NC	244	66.2	This work
Fe-Co-N-PC-900	430	93	1
(Co, Ni)Se ₂ @NiFe LDH	277	114	2
Fe-NiCo-S	247	111	3
NiCoP/NC PHCs	297	51	4
Fe ₁ Co ₁ -P/C	360	58.4	5
ZIF-FeCo-F-300	250	51.2	6
Fe ₁ Co ₃ -NC-1100	349	99.93	7
CoFeP@C	336	82.5	8
A _{2.7} B-MoF-FeCo _{1.6}	288	39	9
Co ₃ O ₄ /Co-Fe oxide DSNBs	297	61	10

Table S3. Comparison of RuO₂ parameter appearing in the literature

Catalyst s	Electrolyte	Overpotential		Ref.
		at 10 mA cm ⁻²	Tafel slope (vs. RHE)	
RuO ₂	1 M KOH	344 mV	97.4 mV dec ⁻¹	This Work
RuO ₂	1 M KOH	328 mV	130 mV dec ⁻¹	3
RuO ₂	1 M KOH	378 mV	65.5 mV dec ⁻¹	12
RuO ₂	1 M KOH	287 mV	108.6 mV dec ⁻¹	13
RuO ₂	1 M KOH	315 mV	135.7 mV dec ⁻¹	14
RuO ₂	1 M KOH	327 mV	356.44 mV dec ⁻¹	15
RuO ₂	1 M KOH	-	141 mV dec ⁻¹	8
RuO ₂	1 M KOH	332 mV	133.8 mV dec ⁻¹	17
RuO ₂	1 M KOH	264 mV	192 mV dec ⁻¹	18
RuO ₂	1 M KOH	298 mV	124 mV dec ⁻¹	19
RuO ₂	0.1MKOH	380 mV	95 mV dec ⁻¹	20

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