

Supplementary Material

Dual-strategy of interface and reconstruction engineering to boost efficient alkaline water and seawater oxidation

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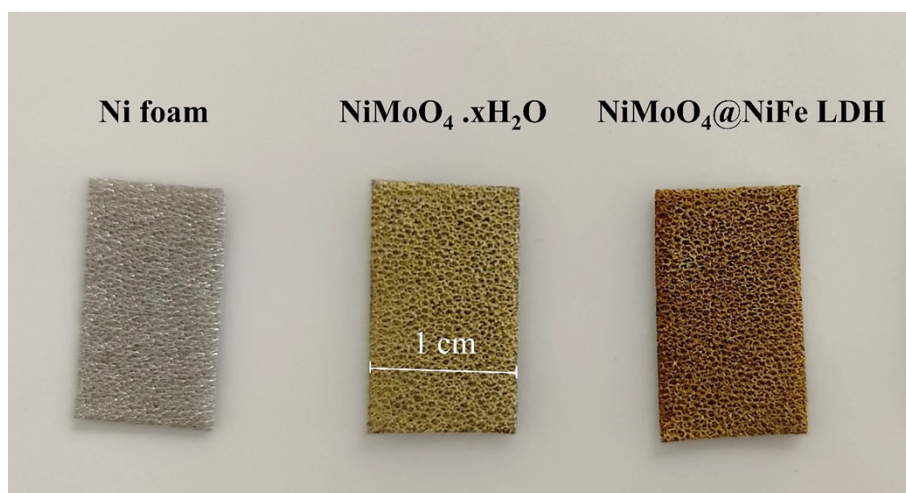


Fig. S1. Optical photographs of Ni foam, NiMoO₄.xH₂O and NiMoO₄@NiFe LDH.

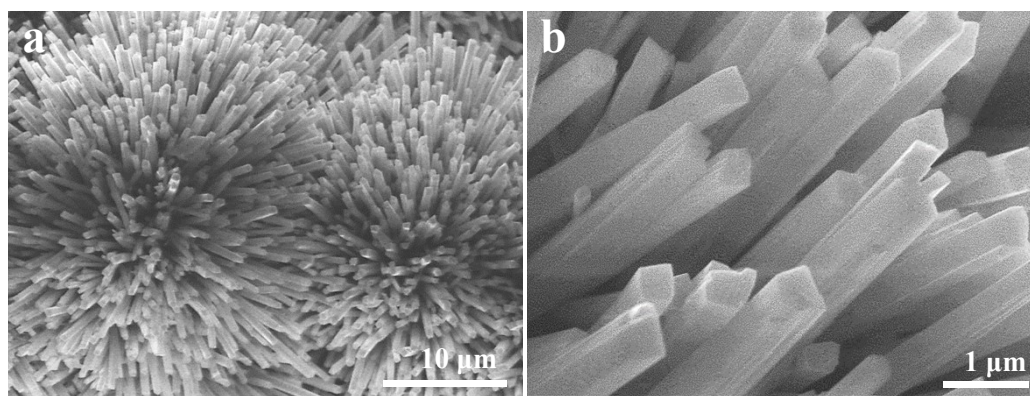


Fig. S2. SEM images at low (a) and high (b) magnifications for NiMoO₄.xH₂O.

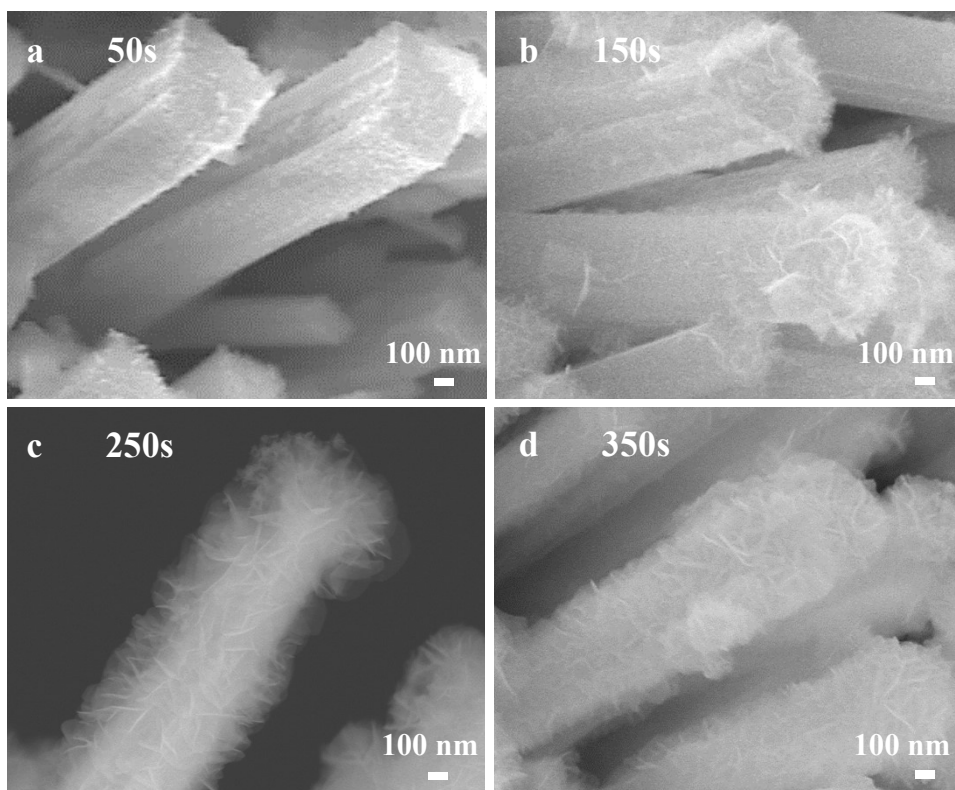


Fig. S3. SEM images of NiMoO₄@NiFe LDH with different electrodeposition time of NiFe LDH.

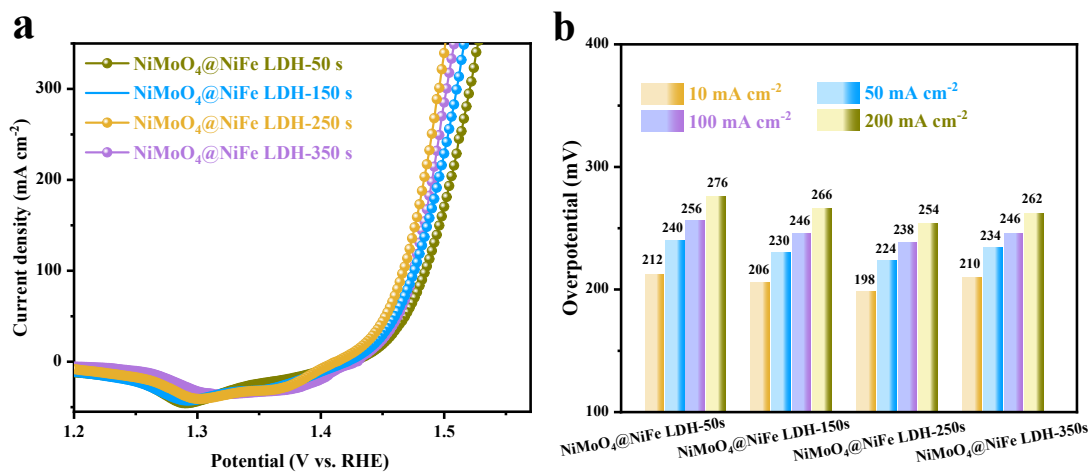


Fig. S4. OER polarization curves of NiMoO₄@NiFe LDH with different electrodeposition time of NiFe LDH tested in 1 M KOH electrolyte. (b) Comparison of the overpotentials required for these catalysts to attain current densities of 10, 50, 100, and 200 mA cm⁻² for OER.

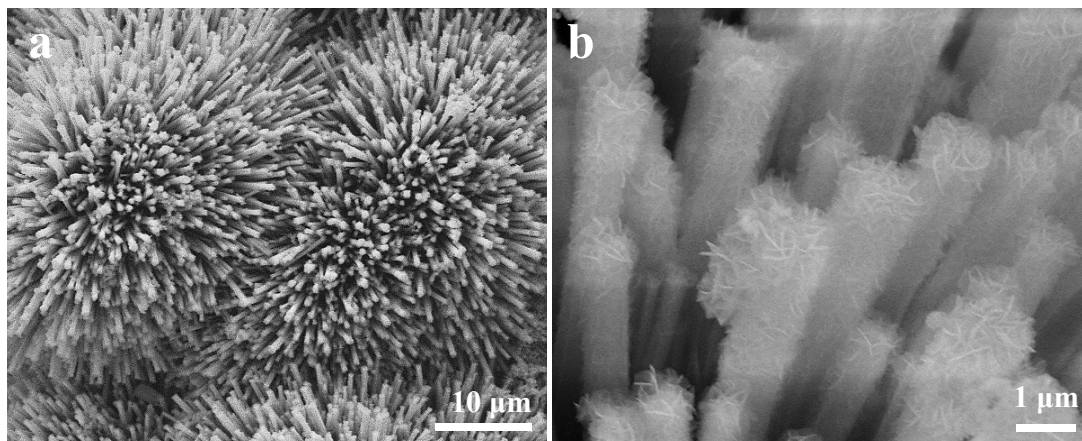


Fig. S5. SEM images at low (a) and high (b) magnifications for NiMoO₄@NiFe LDH-250 s.

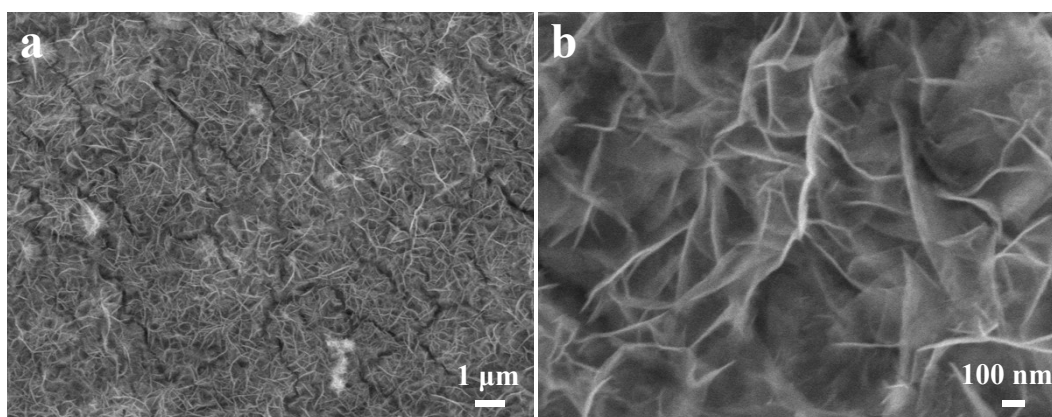


Fig. S6. SEM images at low (a) and high (b) magnifications for NiFe LDH/NF.

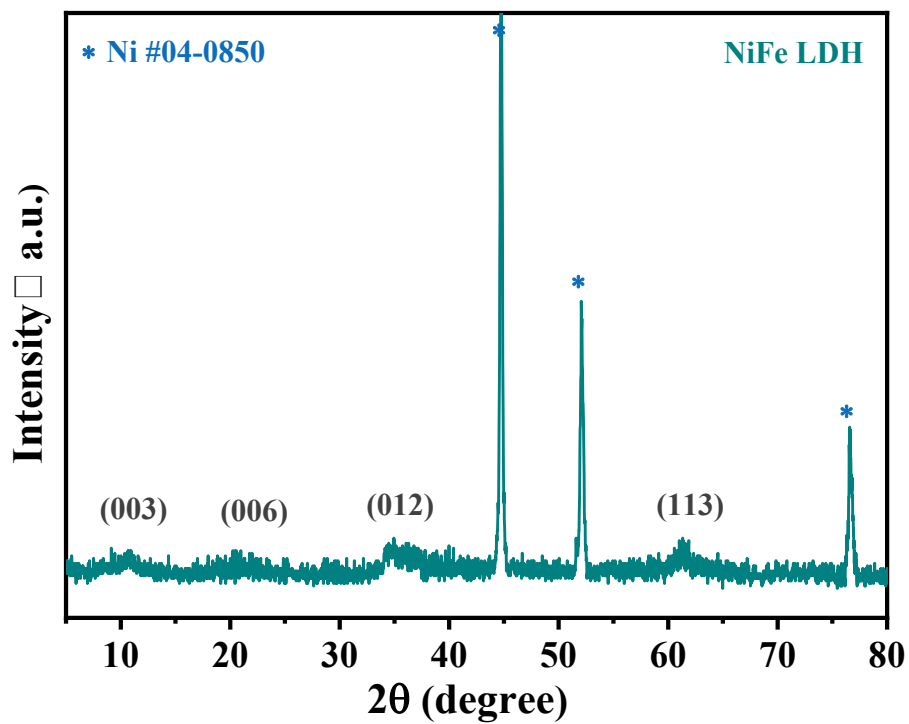


Fig. S7. XRD patterns of NiFe LDH.

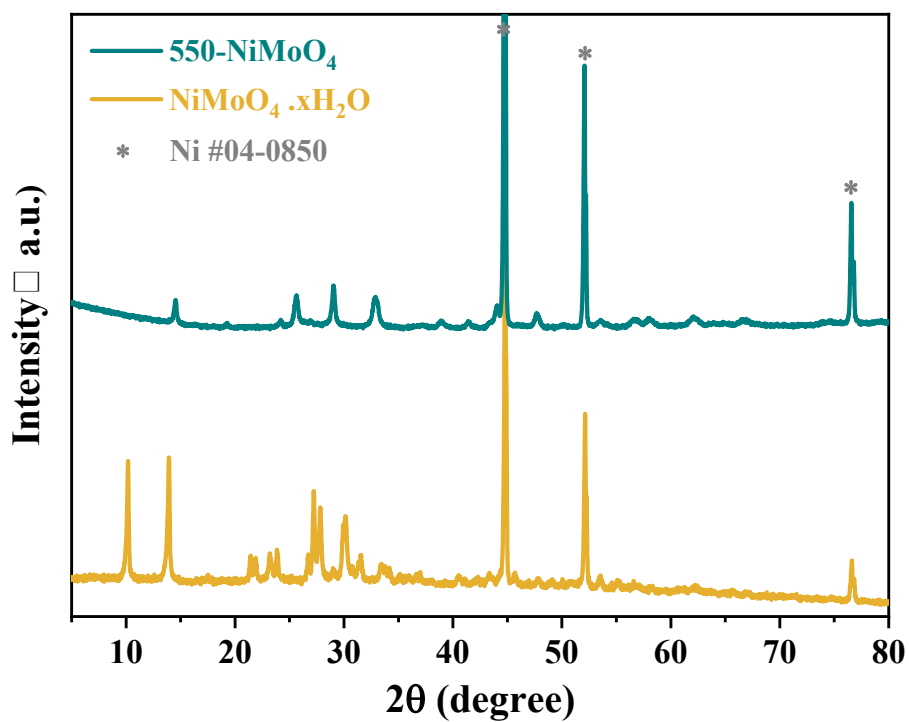


Fig. S8. XRD patterns of annealed NiMoO₄ and NiMoO₄.xH₂O.

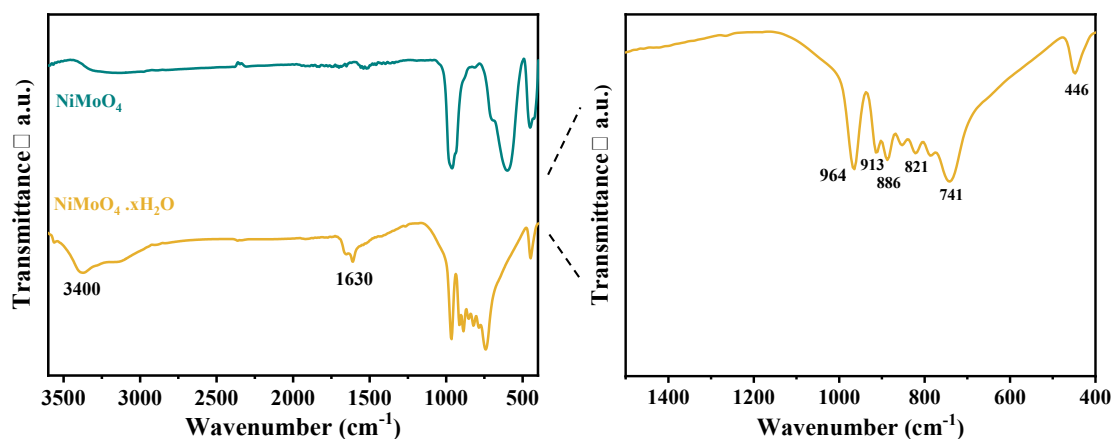


Fig. S9. FTIR spectra of annealed NiMoO₄ and NiMoO₄·xH₂O with the magnified spectra.

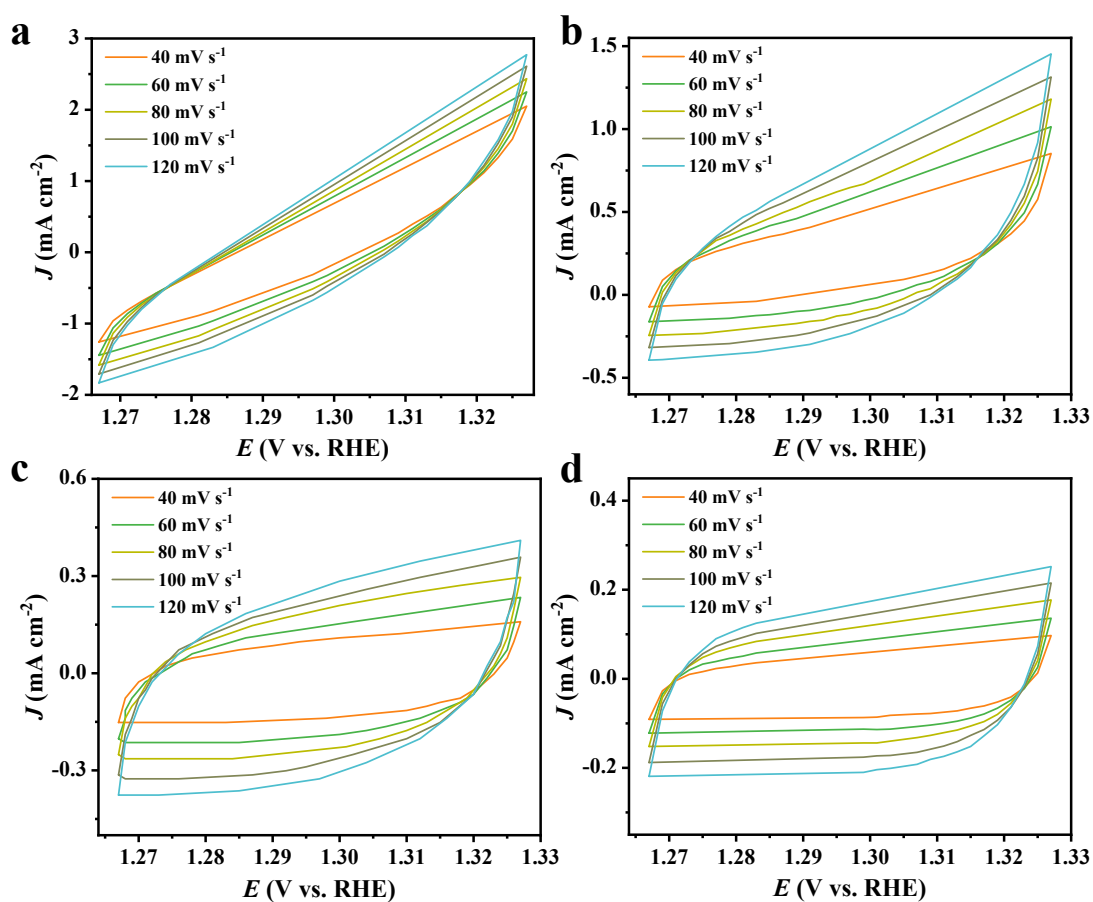


Fig. S10. Typical cyclic voltammograms at different scan rates. (a) NiMoO₄@NiFe LDH, (b) NiMoO₄·xH₂O, (c) NiFe LDH and (d) NF with scan rates ranging from 40 mV/s to 120 mV/s with an interval point of 20 mV/s.

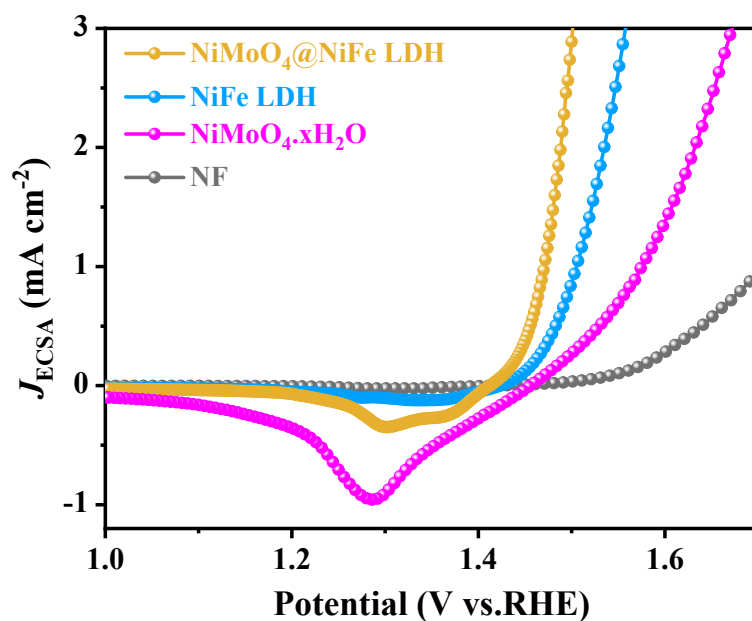


Fig. S11. ECSA normalized polarization curves for NiMoO₄@NiFe LDH, NiFe LDH, NiMoO₄.xH₂O and NF.

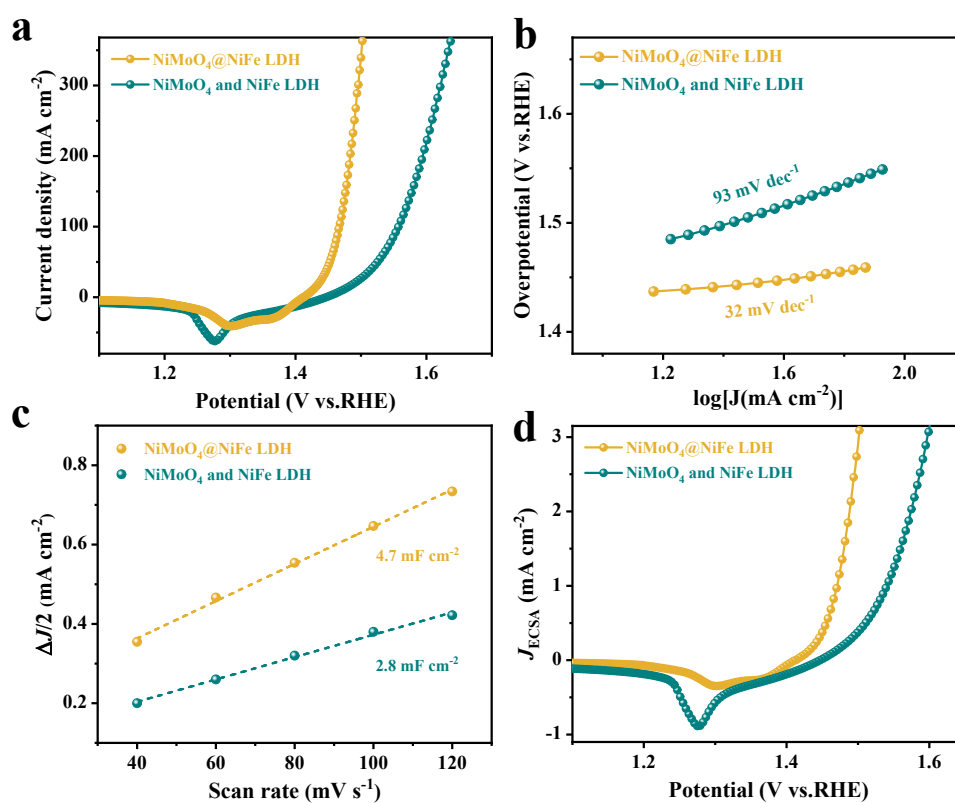


Fig. S12. (a) OER Polarization curves, (b) corresponding Tafel plots, (c) electrical double layer capacitance (C_{dl}) and (d) ECSA normalized polarization curves of core-shell NiMoO₄@NiFe LDH catalysts and physically mixed composite.

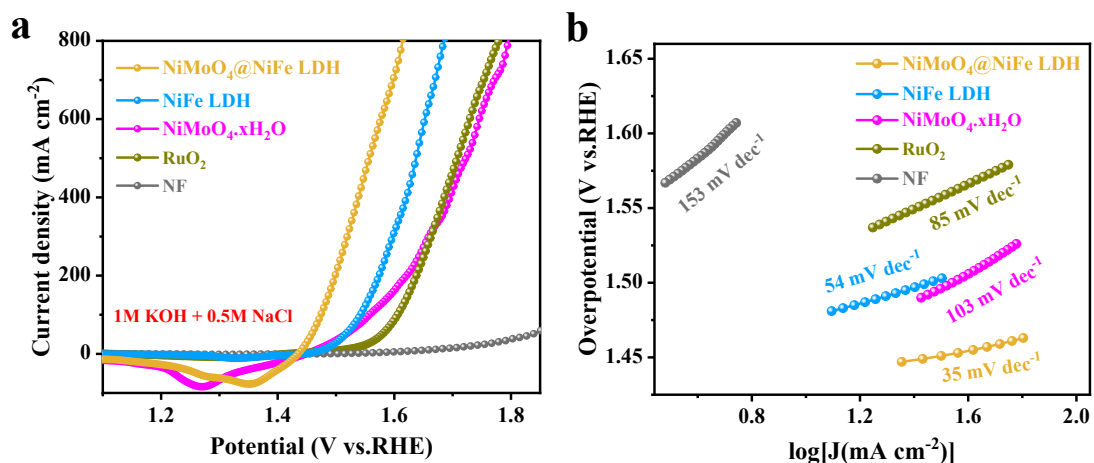


Fig. S13. (a) OER Polarization curves and (b) corresponding Tafel plots of the studied catalysts in 1M KOH + 0.5 M NaCl electrolyte.



Fig. S14. The digital photograph of the results for ClO⁻ formation in 1 M KOH seawater electrolyte after OER stability testing of NiMoO₄@NiFe LDH at 100 mA cm⁻² for 80 h. The formation of hypochlorite (ClO⁻) was examined via the potassium iodide starch paper.

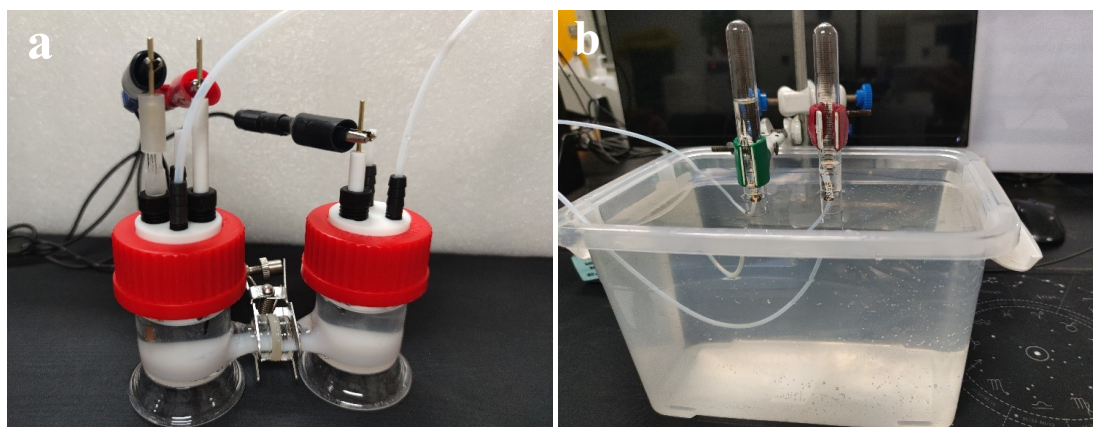


Fig. S15. Photograph of the (a) H type cell and (b) gas collection device.

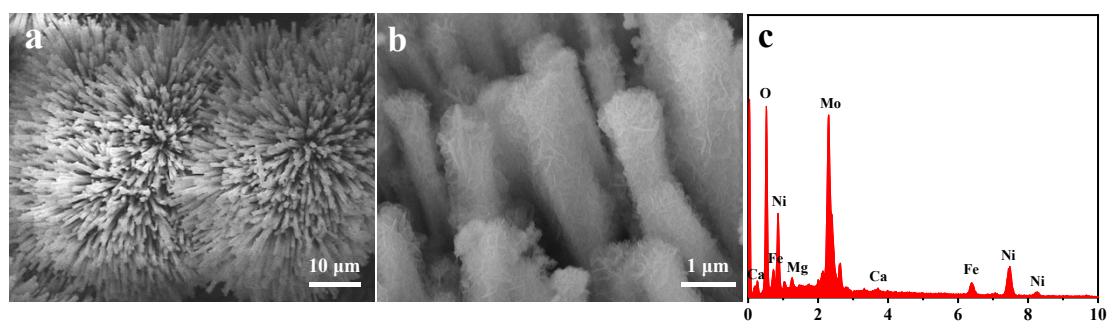


Fig. S16. (a-b) SEM images and (c) EDS spectrum of $\text{NiMoO}_4@\text{NiFe}$ LDH catalyst after immersion in natural seawater for 15 days.

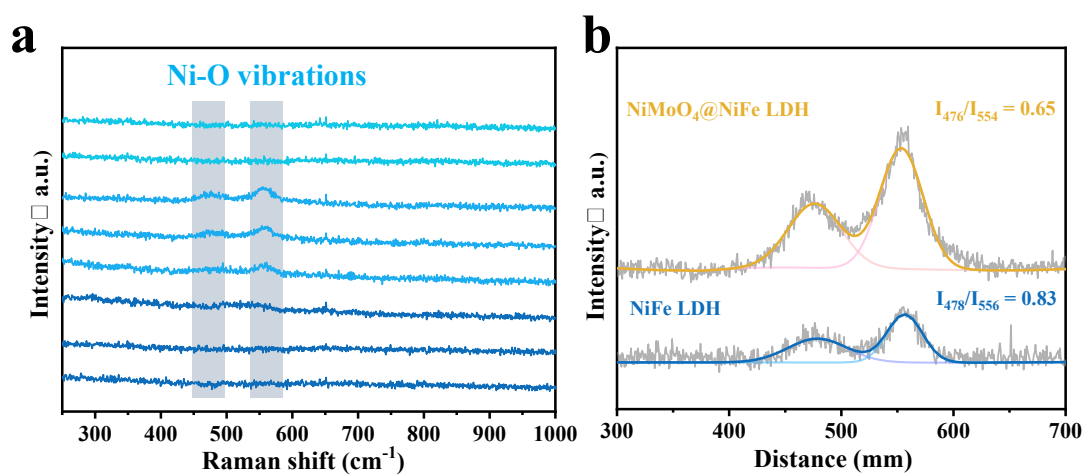


Fig. S17. (a) In-situ Raman spectra of NiFe LDH in 1 M KOH during the OER process. (b) Raman bands of $\text{NiMoO}_4@\text{NiFe}$ LDH and NiFe LDH measured at 1.6 V versus RHE.

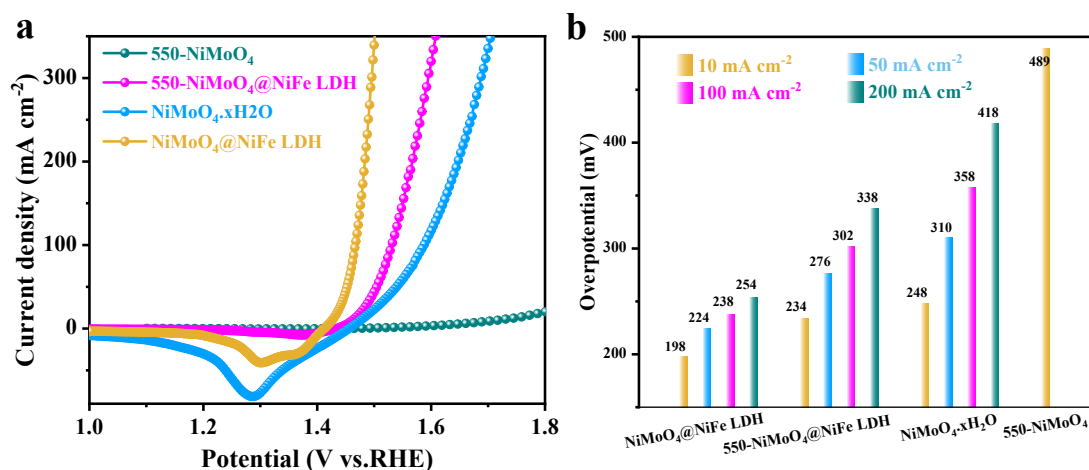


Fig. S18. OER polarization curves of NiMoO₄·xH₂O, NiMoO₄@NiFe LDH, annealed NiMoO₄ and annealed NiMoO₄@NiFe LDH in 1 M KOH electrolyte. (b) Comparison of the overpotentials required for these catalysts to attain current densities of 10, 50, 100, and 200 mA cm⁻² for OER.

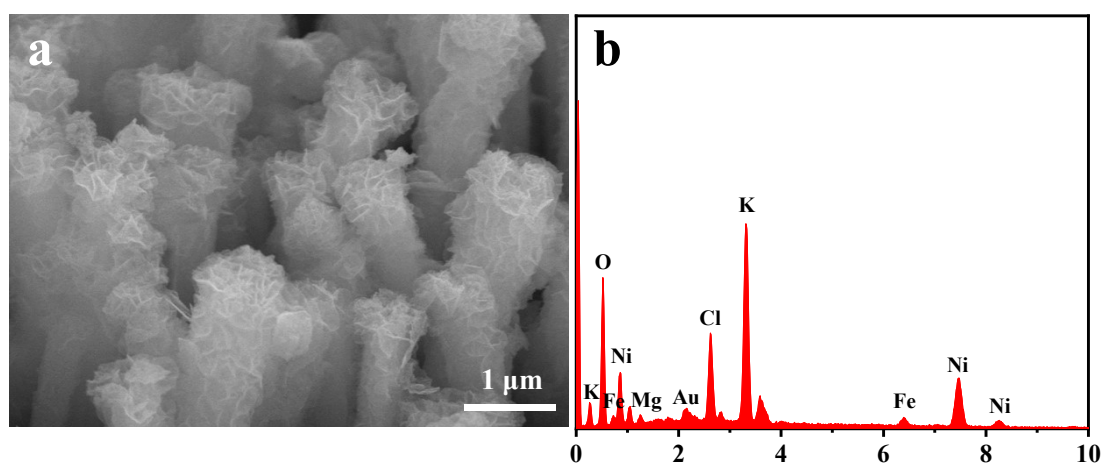


Fig. S19. (a) SEM images and (b) EDS spectrum of NiMoO₄@NiFe LDH catalyst after OER test in 1 M KOH seawater.

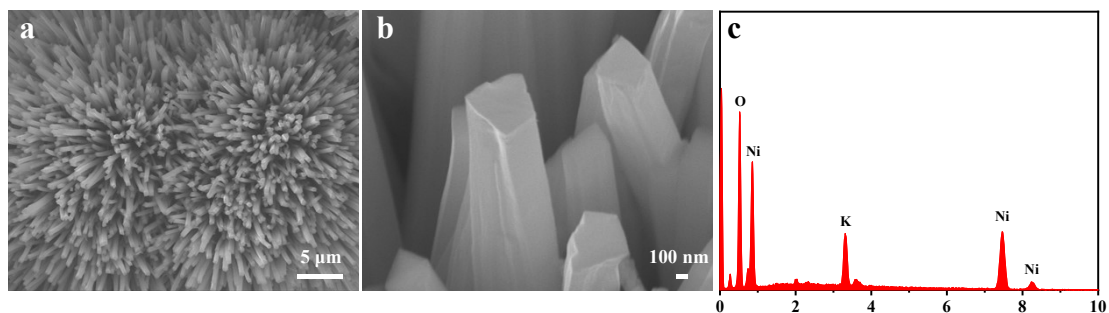


Fig. S20. (a-b) SEM images and (c) EDS spectrum of $\text{NiMoO}_4 \cdot x\text{H}_2\text{O}$ catalyst after OER test in 1 M KOH.

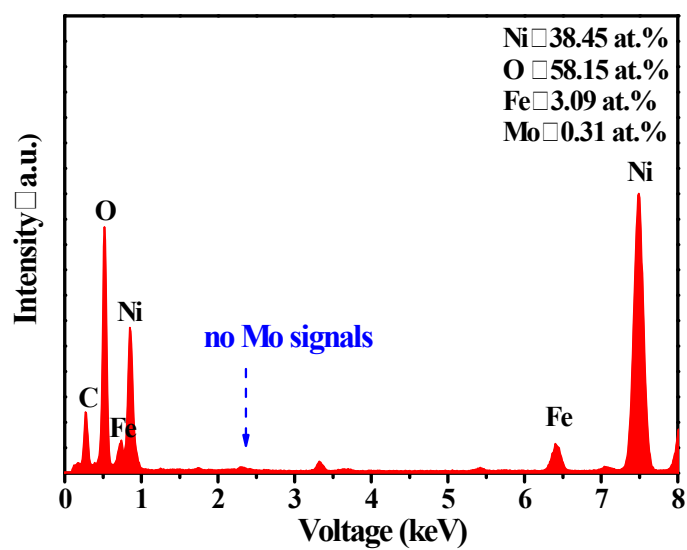


Fig. S21. EDS map sum spectrum of $\text{NiMoO}_4 @ \text{NiFe}$ LDH catalyst after OER test in 1 M KOH.

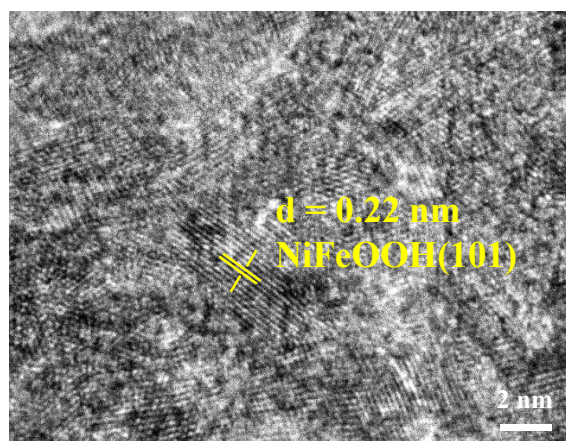


Fig. S22. The high-magnification TEM image of NiMoO₄@NiFe LDH catalyst after OER test in 1 M KOH.

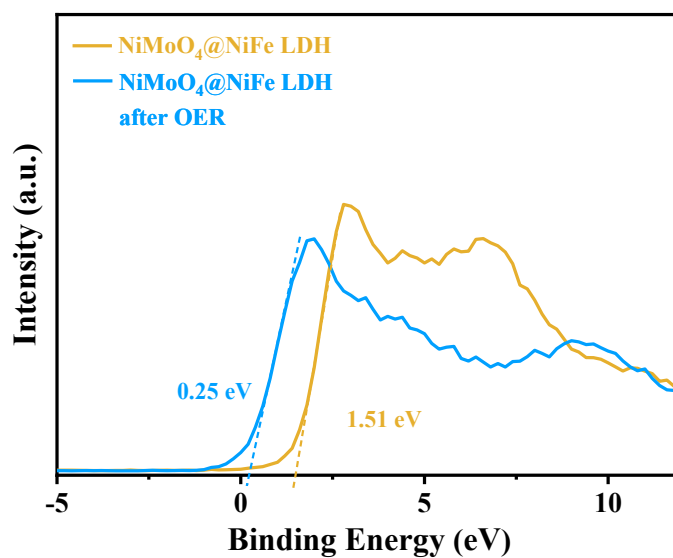


Fig. S23. Valence band spectra of NiMoO₄@NiFe LDH before and after OER catalysis in 1 M KOH.

Table S1. Comparison of OER performance of NiMoO₄@NiFe LDH with some previously reported NiFe-based catalysts in 1.0 M KOH solution.

Catalysts	Electrolyte	η (mV) at 10 mA cm ⁻²	Tafel slope (mV/dec)	Reference
NiMoO₄@NiFe LDH	1.0 M KOH	198	32	<i>This work</i>
Fe-doped β -Ni(OH) ₂	1.0 M KOH	219	53	1
hcp-NiFe@NC	1.0 M KOH	226	41	2
Fe _{0.052} Ni-POMo	1.0 M KOH	255.3 \pm 1.6	43.8	3
NiFe-based SURMOFs	0.1 M KOH	300	44.3	4
NiFe-LDH	1.0 M KOH	275	56.7	5
Ni Fe MOF	0.1 M KOH	240	34	6
NiOOH/FeOOH NBs	1.0 M KOH	246	41	7
Ni _{0.8} Co _{0.1} Fe _{0.1} OxHy	1.0 M KOH	239	45.4	8
Ni ^{II} Fe ^{III} @NC	1.0 M KOH	360	81	9
NiFe@N doped carbon	0.1 M KOH	350	56	10
Fe/(Ni)OOH	1.0 M KOH	290	32	11
Ni _{2/3} Fe _{1/3} LDH	1.0 M KOH	310	76	12
Fe-S-NiMoO ₄ /MoO ₃	1.0 M KOH	212	41	13
Ni (Fe)OxHy	1.0 M KOH	218 \pm 5	31 \pm 4	14
NiCoFe-P- NP@NiCoFe-PBA	1.0 M KOH	223	78	15

NiFe/NiFe-OH	1.0 M KOH	222	41	16
MnCo-CH@NiFe-OH	1.0 M KOH	186	49	17
a-NiFe-OH/NiFeP/NF	1.0 M KOH	199	39	18

Table S2. Comparison of OER performance of NiMoO₄@NiFe LDH with some previously reported catalysts for seawater oxidation.

Catalysts	Electrolyte	η (mV) at 100 mA cm ⁻²	η (mV) at 500 mA cm ⁻²	Referenc e
NiMoO₄@NiFe LDH	1 M KOH seawater	251	349	<i>This work</i>
Ni ₃ S ₂ /Co ₃ S ₄	1 M KOH seawater	360	440	19
Ni ₃ FeN@C	1 M KOH seawater	283	351	20
MoS ₂ -(FeNi) ₉ S ₈	1 M KOH seawater	256	329	21
Gd-Mn ₃ O ₄ @ CuO- Cu(OH) ₂	1 M KOH seawater	-	400	22
Ir ₁ /Ni _{1.6} Mn _{1.4} O ₄	0.5 M KOH seawater	330	-	23
B-Co ₂ Fe LDH	1 M KOH seawater	310	376	24
NiFe LDH	1 M KOH seawater	247	296	25
CoPx@FeOOH	1 M KOH seawater	283	337	26

S-(Ni,Fe)OOH	1 M KOH seawater	300	398	27
Ni ₃ S ₂ /Fe-NiPx	1 M KOH seawater	290	336	28

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