

Supporting information

A self-supporting anode with high activity and long-term stability for efficient water-splitting

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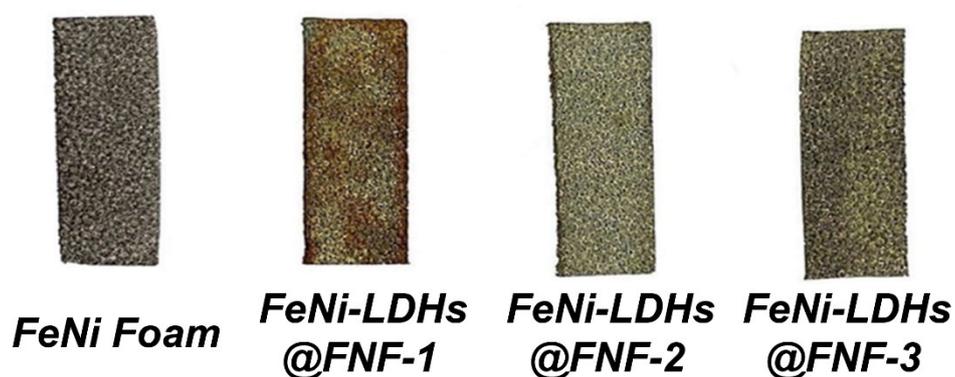


Fig. S1 Appearance of FeNi-LDHs prepared in NiSO₄ solution with varying Ni²⁺ concentration.

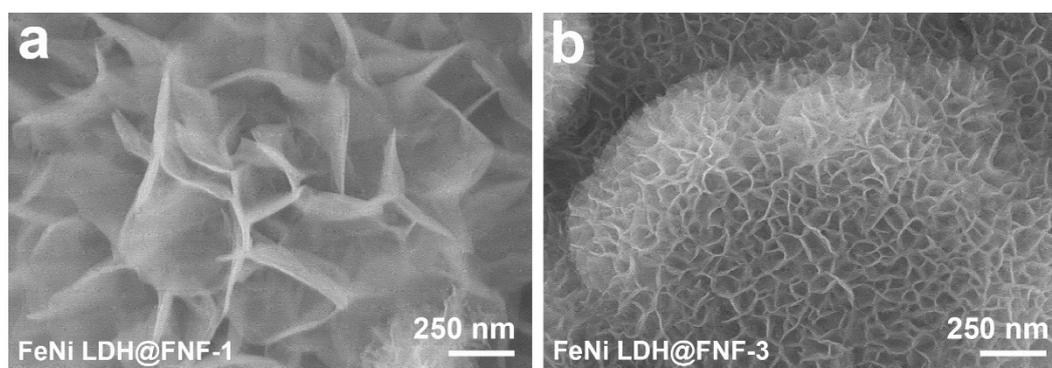


Fig. S2 (a) SEM images of FeNi-LDH@sFNF-1. (b) SEM image of FeNi-LDH@FNF-3.

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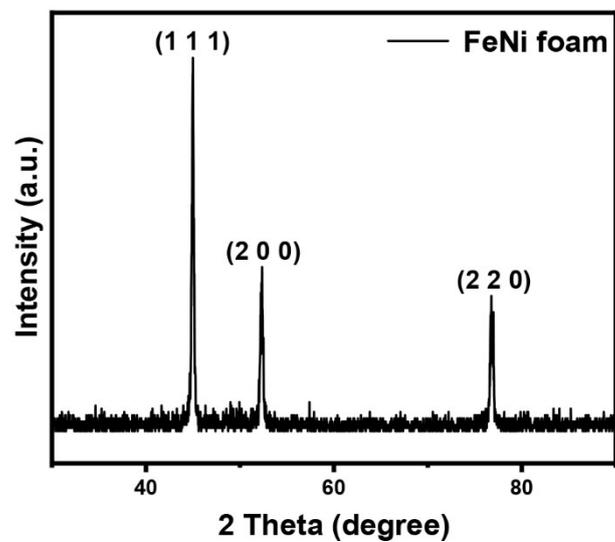


Fig. S3 XRD spectra of FeNi foam.

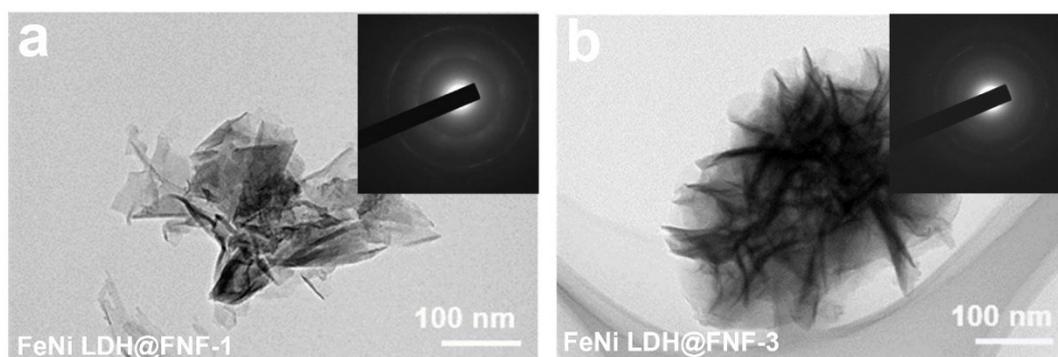


Fig. S4 (a) TEM image of FeNi-LDHs@FNF-1. (b) TEM image of FeNi-LDHs@FNF-3.

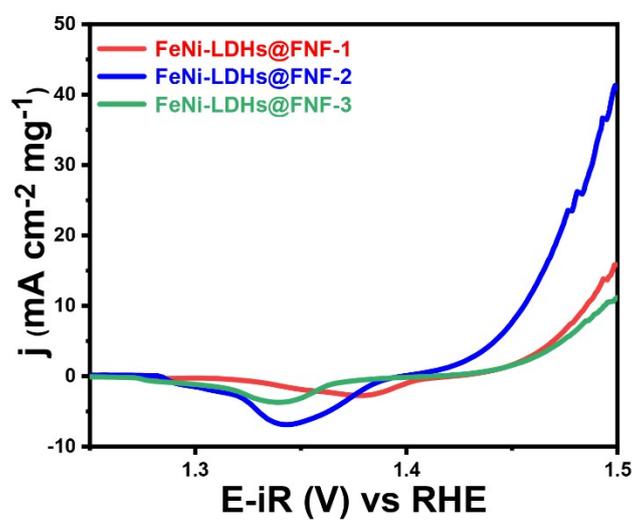


Fig. S5 Polarization curves of FeNi-LDHs normalized with catalyst mass.

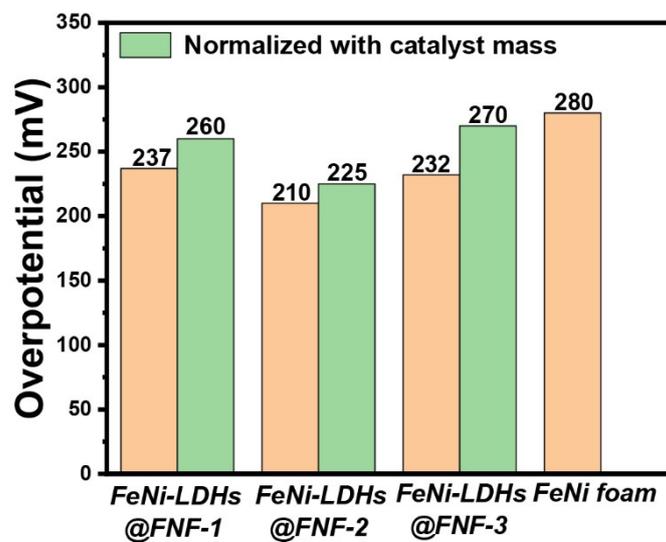


Fig. S6 Overpotential of FeNi-LDHs grown at different concentrations.

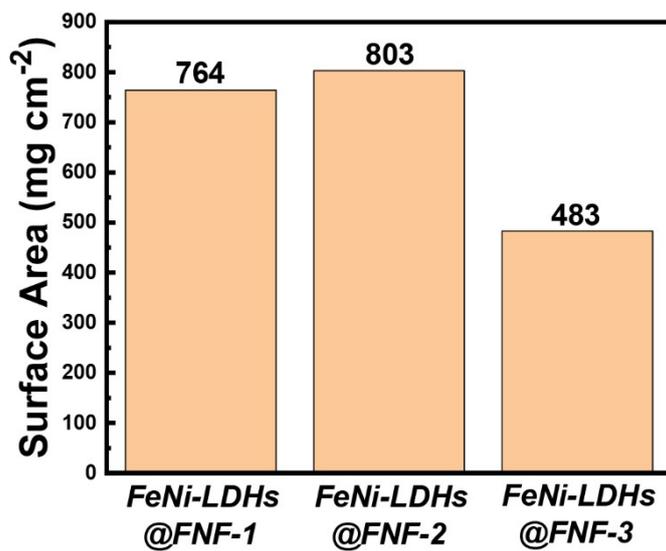


Fig. S7 BET surface area of FeNi-LDHs grown at different concentrations.

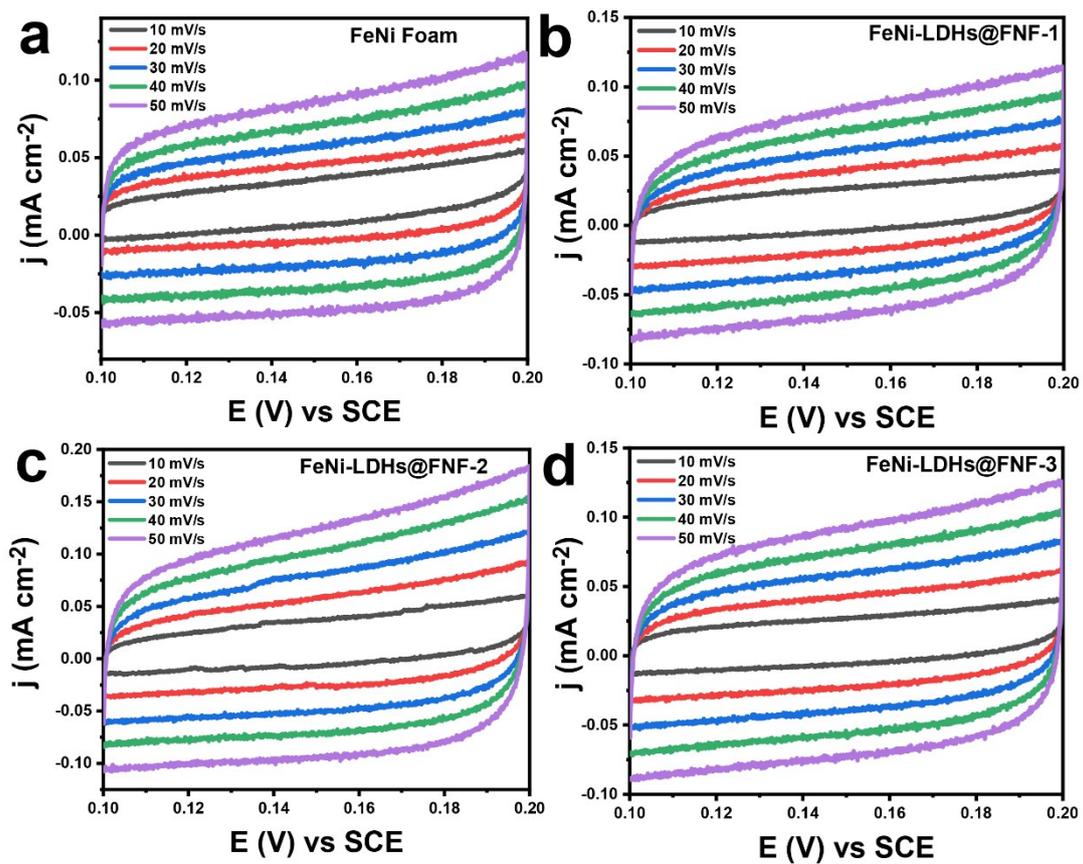


Fig. S8 (a, b, c, d) ECSA curves of FeNi-LDHs grown on FeNi foam under different NiSO_4 concentrations.

Table S1 Process comparison of different OER catalysts (* is the catalyst developed in this article)

Catalyst	Preparation	Chemicals
FeNi-LDHs-rGO	<ol style="list-style-type: none"> 1. Synthesis of FeNi LDH by hydrothermal method. 2. Decarbonisation of FeNi LDH. 3. Synthesis of GO from natural graphite by a modified Hummers method. 4. Embedding graphene oxide into the FeNi LDH layer by anion exchange. 5. Reduction of FeNi-GO LDH to FeNi-rGO LDH. 	ferrous chloride, nickel chloride, urea aqueous, trisodium trate, hydrochloric acid, sodium chloride, natural graphite, hydrazine hydrate, ammonium
NiFe-LDHs/CNT	<ol style="list-style-type: none"> 1. Oxidization of carbon nanotubes. 2. Synthesis of NiFe LDH on multi-walled carbon nanotubes by solvothermal method. 	carbon nanotubes sulfuric acid sodium nitrate potassium permanganate hydrogen peroxide N,N-dimethylformamide nickel acetate ferrous nitrate
N-NiFe-LDHs	Synthesis of N-NiFe LDH on nickel foam in the chemical bath.	nickel acetate hexamethylene tetramine

		iron nitrate ethanol
n-NiFe-LDHs/NGF	<p>1. Preparation of magnesium oxide template by surfactant-assisted hydrothermal reaction.</p> <p>2. Synthesis of NGF via CVD method with the as-obtained MgO as templates.</p> <p>3. Synthesis of n-NiFe LDH/NGF via urea-assisted precipitation with NGF as the substrate.</p>	magnesium oxide polyethylene glycol methane/ammonia hydrochloric acid N-methylpyrrolidone nickel nitrate ferric nitrate urea
NiCo-LDHs/Ni NiMn-LDHs/Ni NiCoMn-LDHs/Ni	Synthesis of NiCo LDH, NiMn LDH and NiCoMn LDH on nickel foam by hydrothermal method.	nickel acetate cobalt acetate manganese acetate sodium carbonate sodium hydroxide urea
*FeNi-LDHs/FeNi	Synthesis of FeNi-LDHs on FeNi foam at room temperature.	nickel sulfate

Some calculation formulas involved in the article:

The overpotential is calculated by the following equation:

$$E \text{ (vs. RHE)} = E \text{ (vs. SCE)} + 1.0672 \text{ V}$$

$$\eta_{OER} = E \text{ (vs. RHE)} - 1.23 \text{ V}$$

The ECSA is calculated by the following equation:

$$ECSA = Cdl/Cs(40 \mu F cm^{-2})$$

The electric quantity of Ni reduction is calculated by the following equation:

$$Q = \frac{\int j \cdot t dt}{m(Ni)}$$

$\int j \cdot t dt$ is obtained by the integration of the j-t curve (Fig.S4). $m(Ni)$ is obtained from the catalyst mass (By measuring the quality difference of samples before and after chemical corrosion. FeNi-LDHs@FNF-1: 2 mg, FeNi-LDHs@FNF-2: 2 mg, FeNi-LDHs@FNF-3: 3 mg) and XPS element content.