

Supporting Information

For

Nickel foam and stainless steel mesh as electrocatalysts for hydrogen evolution reaction, oxygen evolution reaction and overall water splitting in alkaline media

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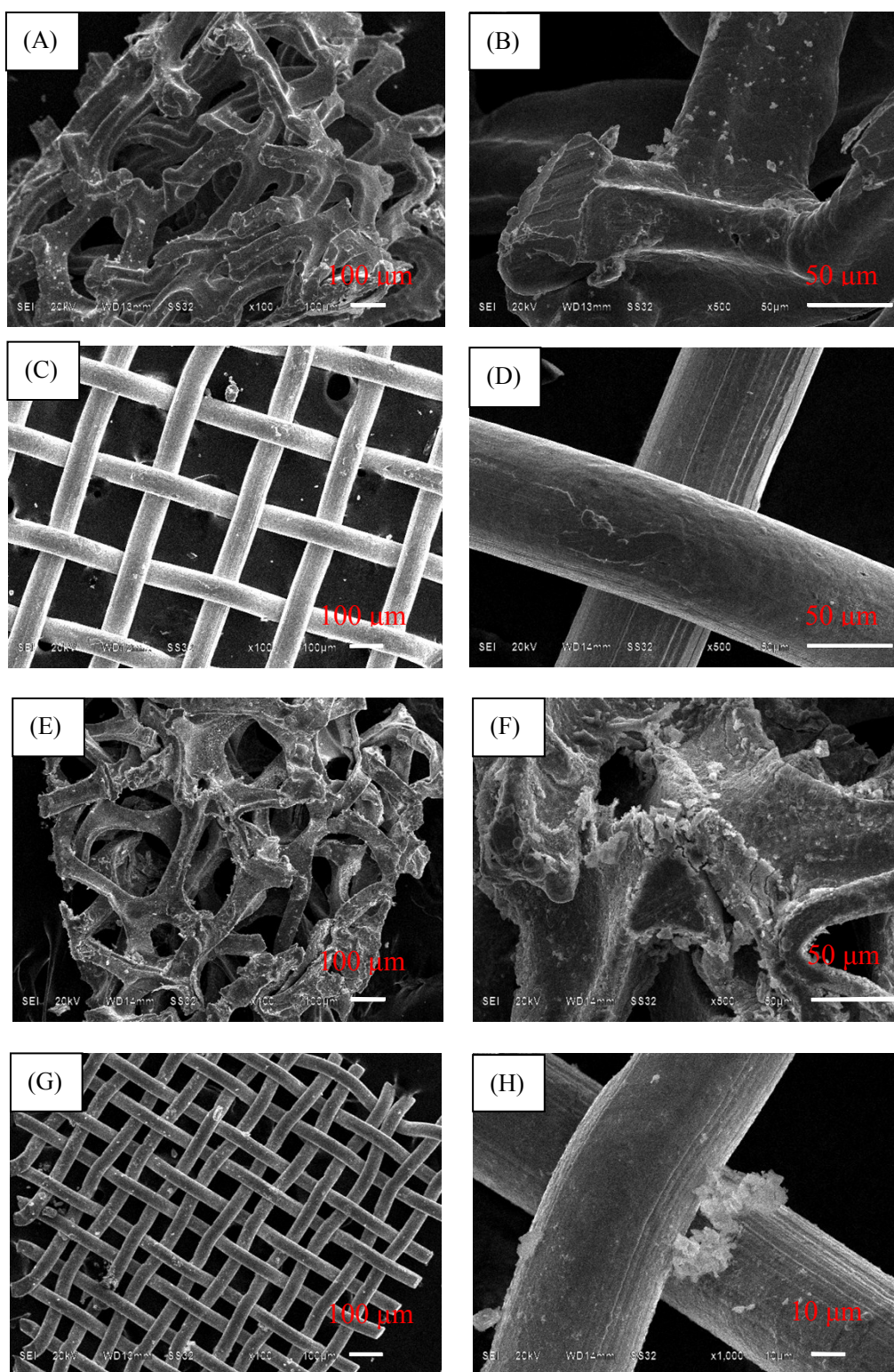


Fig. S1. The SEM images of Cu foam (A, B) and Ni mesh (C, D) before use. The SEM images of Ni foam (E, F) and SS mesh (G, H) after the stability test.

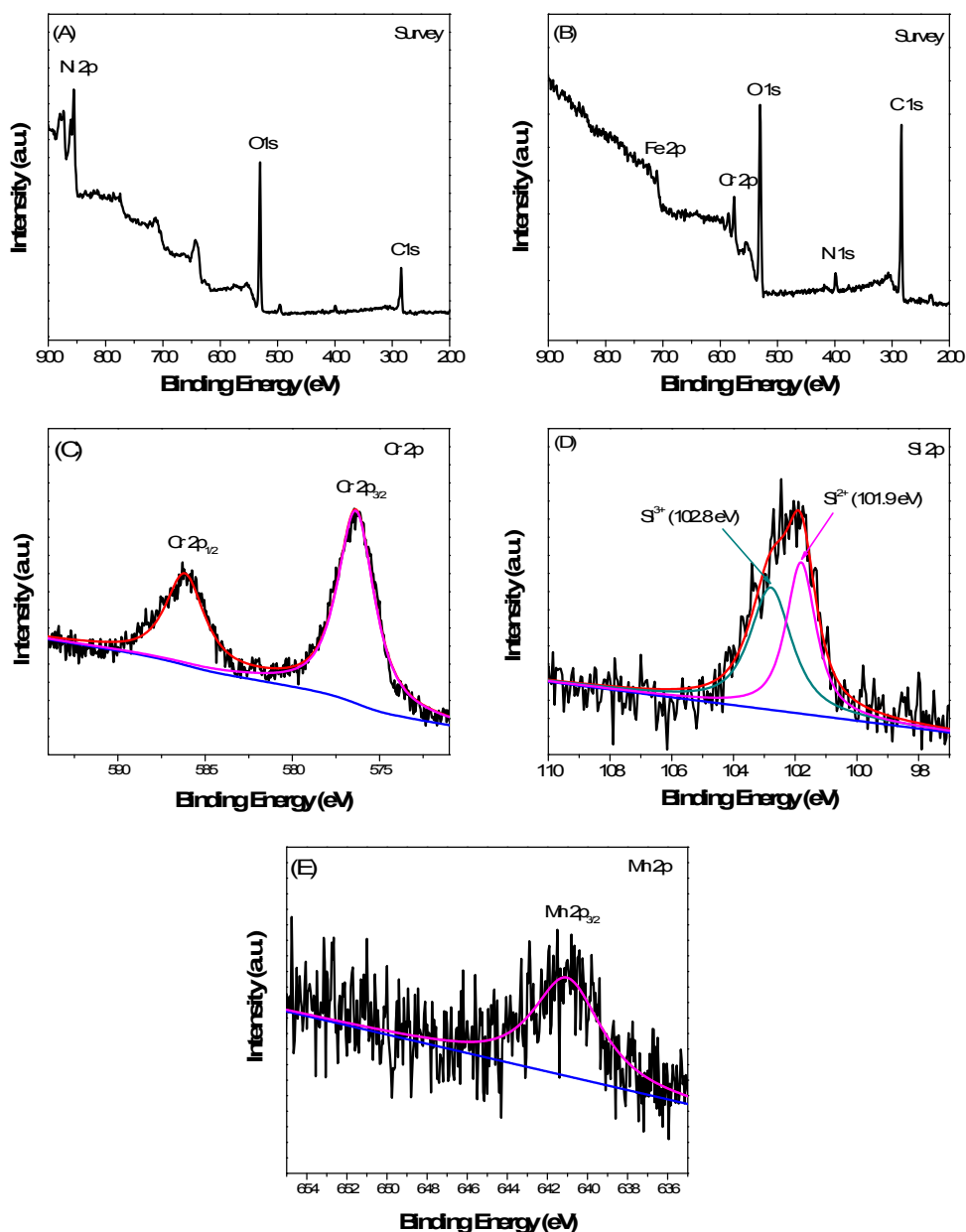


Fig. S2. The Survey XPS spectra of Ni foam (A) and SS mesh (B) before use; The high resolution XPS spectra of (C) Cr 2p and (D) Si 2p (E) Mn 2p of SS mesh before use.

In the Cr 2p region, the peaks at 586.3 and 576.4 eV correspond to the Cr 2p of Cr³⁺ [1, 2]. The high-resolution Si 2p XPS spectra of SS mesh are shown in Fig. S2D, 102.8 eV was characteristics of Si³⁺, while the peaks at 101.9 eV was characteristics of Si²⁺ [3]. The spin-orbit components of the Mn 2p_{3/2} photoemission were located at 641.2 eV, which corresponds to MnO and Mn₂O₃ [4, 5].

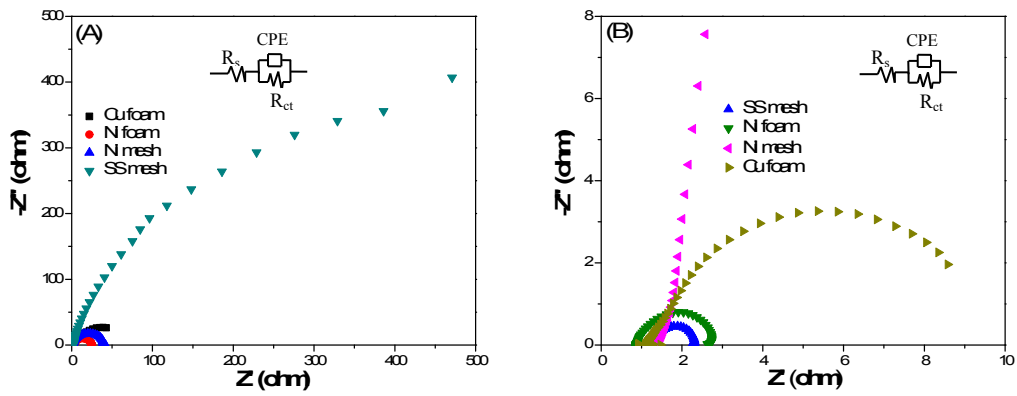


Fig. S3. Nyquist plots of EIS of samples with an amplitude of 5 mV (A) from 10^5 Hz to 0.1 Hz (HER); (B) from 10 kHz to 0.01 Hz (OER).

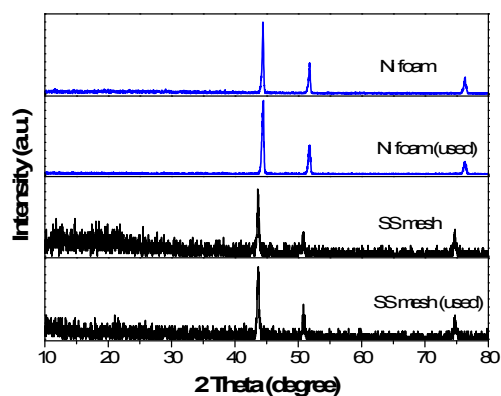


Fig. S4. Comparison of the XRD patterns of Ni foam and SS mesh before and after the stability test.

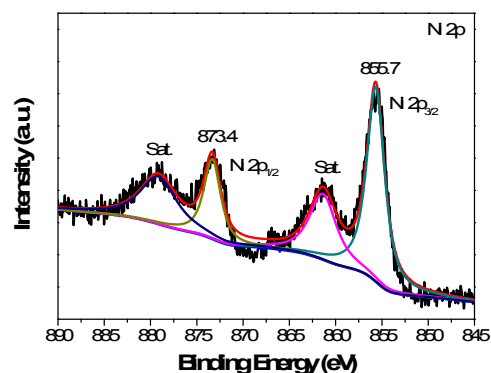


Fig. S5. The XPS spectra of Ni 2p of Ni foam after the stability test.

The XPS results of the Ni foam after the HER stability test are shown in the Fig. S5. The peaks at binding energies of 873.4 and 855.7 eV can be assigned to Ni 2p_{1/2} and Ni 2p_{3/2} of NiO, respectively [6]. The satellite peak at around 879.2 eV and 861.4 eV are two shake-up type peaks of nickel at the high binding energy side of the Ni 2p_{1/2} and Ni 2p_{3/2} edge [7]. Comparing the XPS data of the Ni 2p before (Fig. 3A) and after the stability test (Fig. S5), it can be seen that the peak at 852.1 eV disappeared after the electrocatalysis test, which shows the metallic nickel has been completely oxidized to NiO after the electrocatalysis test in 1 M KOH solution.

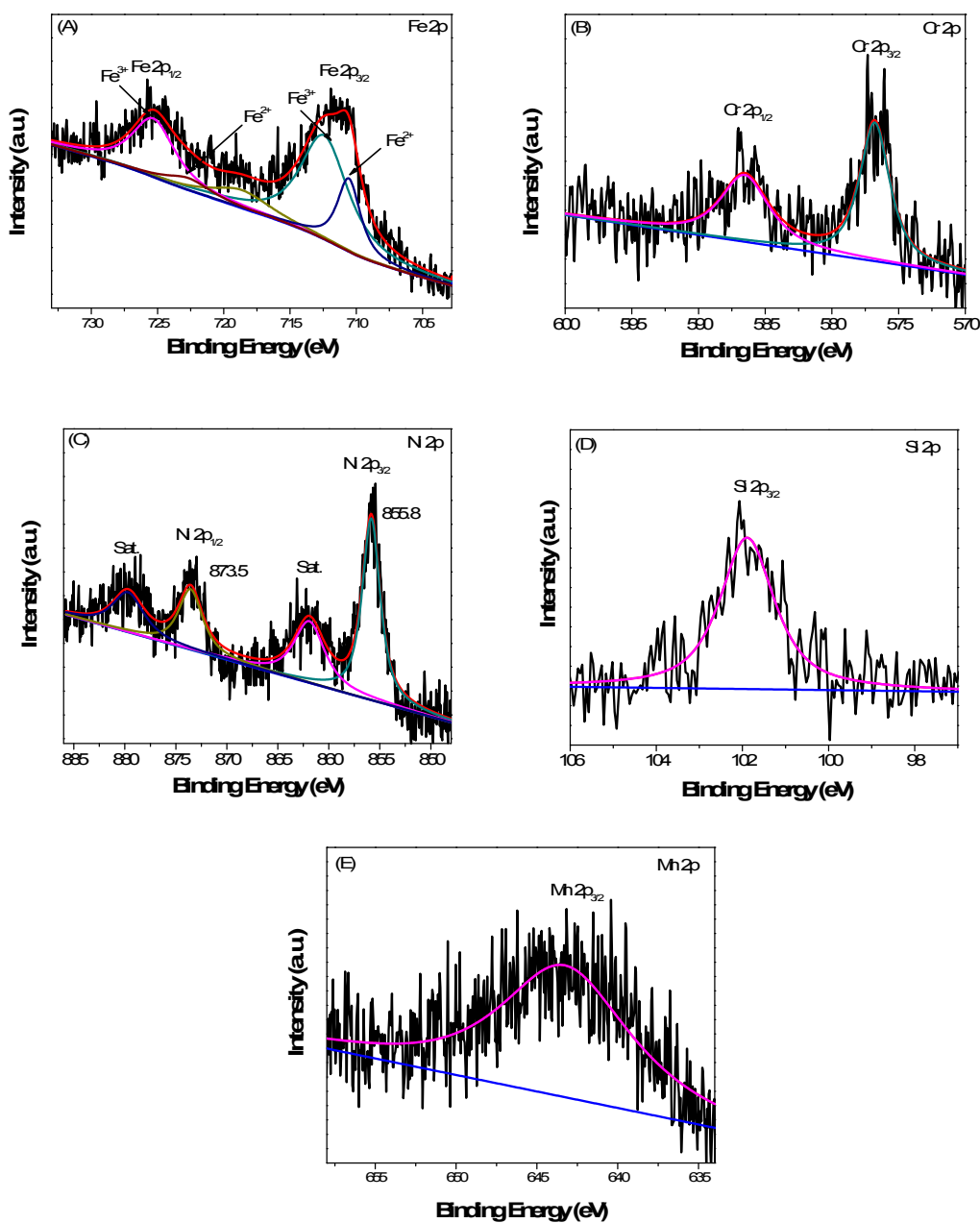


Fig. S6. The XPS spectra of Fe 2p (A), Cr 2p (B), Ni 2p (C), Si 2p (D) and Mn 2p (E) of SS mesh after the stability test.

The XPS results of the SS mesh after the OER stability test are shown in the Fig. S6. Fig. S6A shows the high resolution Fe 2p XPS of SS mesh after the stability test. The XPS spectrum of Fe 2p displays four peaks at 710.6, 712.5, 722.7 and 725.5 eV, which are attributed to the binding energy of Fe 2p_{3/2} and Fe 2p_{1/2}, respectively. The peaks at 710.6 and 722.7 eV can be attributed to FeO [8], whereas the peaks at 712.5 and 725.5 eV can be assigned to Fe₂O₃ [8, 9]. Comparing the XPS data of the Fe 2p before and after the stability

test, it can be seen that the peak at 706.4 eV disappeared after the stability test, which shows the metal iron after the stability test has been completely oxidized. In addition, the XPS peak of Fe^{2+} after the stability test decreased, and the peak area of Fe^{3+} was relatively increased, which shows the part of the ferrous iron is converted into ferric iron, and Fe elements was existed as a mixture of FeO and Fe_2O_3 after the stability test. Fig. S6B shows the high resolution Cr 2p XPS spectrum of SS mesh after the stability test, the peaks at binding energies of 586.6 and 576.7 eV can be assigned to Cr 2p of Cr^{3+} [1, 2]. Comparing the XPS data of the Cr 2p before and after the stability test, it can be seen that the peak of Cr element slightly weakens after the stability test. Fig. S6C shows the high resolution Ni 2p XPS of SS mesh after the stability test, the peaks at binding energies of 873.5 and 855.8 eV can be assigned to Ni $2p_{1/2}$ and Ni $2p_{3/2}$ of NiO , respectively [6]. The satellite peak at around 879.7 eV and 861.9 eV are two shake-up type peaks of nickel at the high binding energy side of the Ni $2p_{1/2}$ and Ni $2p_{3/2}$ edge [7], which shows that Ni elements of SS mesh was existed as NiO after the stability test. Fig. S6D shows the high resolution Si 2p XPS spectrum of SS mesh after the stability test, the peaks at 101.9 eV were characteristics of Si^{2+} [3], which indicates that trivalent Si is completely converted into divalent Si after the stability test. Fig. S6E shows the high resolution Mn 2p XPS spectrum of SS mesh after the stability test, the peaks at binding energies of 643.3 eV can be assigned to MnO_2 [2], which indicates that trivalent Mn is converted into tetravalent Mn after the electrocatalysis test in 1 M KOH solution.

Table S1. The catalytic activity data of different catalysts for HER .

Catalyst	η / (V vs RHE) ($j = 10 \text{ mA cm}^{-2}$)	Ref
δ -MnO ₂	80 mV	[10]
Fe/N-HCS-0.5-800	170 mV	[11]
NOPHC ₁₀ -900	290 mV	[12]
NiCoP	118 mV	[13]
Ni ₃ S ₂	200 mV	[14]
W(S _{0.48} Se _{0.52}) ₂	260 mV	[15]
Ni _{0.9} Fe _{0.1} /NC	219 mV	[16]
NiCo ₂ S ₄	210 mV	[17]
Ni ₃ S ₂	223 mV	[18]
Co ₄ Mo ₂ @NC	218 mV	[19]
Mo(S _{0.49} Se _{0.51}) ₂	271.3 mV	[20]
Ni foam	217 mV	This work

Table S2. The catalytic activity data of different catalysts for OER.

Catalyst	η / (V vs RHE) ($j = 10 \text{ mA cm}^{-2}$)	Ref
Co ₉ S ₈	434 mV	[21]
MnO ₂ -CoP ₃	288 mV	[22]
CuO-Co-0.2	394 mV	[23]
Ni ₃ S ₂	187 mV	[24]
Ni ₃ Se ₂	290 mV	[25]
NiCo-LDH	271 mV	[26]
N-NiFe	230 mV	[27]
Ni _{2.53} Ir NCs	302 mV	[28]
CoS	297 mV	[29]
CoP	330 mV	[30]
SS mesh	277 mV	This work

Table S3. Water splitting cell voltage of different electrocatalysts.

Catalyst	E (j = 10 mA cm ⁻²)	Ref
NiFe/NiCo ₂ O ₄ /NF NiFe/NiCo ₂ O ₄ /NF	1.67 V	[31]
Ni ₅ P ₄ Ni ₅ P ₄	1.7 V	[32]
Er-doped CoP Er-doped CoP	1.58 V	[33]
Ni _{0.2} Co _{0.8} Se Ni _{0.2} Co _{0.8} Se	1.592 V	[34]
CoS-RGO CoS-RGO	1.77 V	[35]
NESSP NESS	1.74 V	[36]
SS scrubber SS scrubber	1.98 V	[37]
Co ₁ -Fe ₁ -B-P Co ₁ -Fe ₁ -B-P	1.68 V	[38]
ZIF-8-C6 ZIF-8-C4	1.82 V	[39]
Co(OH) ₂ @NCNTs@NF Co(OH) ₂ @NCNTs@NF	1.72 V	[40]
Ni foam Stainless steel mesh	1.74 V	This work

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