

Supporting Information

Improving the oxygen evolution performance of iron–manganese oxyhydroxides by Cr doping

Jiayang Yu, Tianmi Tang, Jingqi Guan* and Yupeng Guo*

Department of Physical Chemistry, College of Chemistry, Jilin University,
Changchun 130021, PR China. *E-mail: guanjq@jlu.edu.cn (J.Q. Guan),
guoyupeng@jlu.edu.cn (Y.P. Guo)

Synthesis of FeMnCr_{0.6}OOH/NF

FeMnCr_{0.6}OOH/NF was synthesized using hydrothermal synthesis method. 0.2703 g FeCl₃·6H₂O, 0.1979 g MnCl₂·4H₂O, 0.1599 g CrCl₃, 0.24 g NH₂CONH₂ and 0.075 g PVP were dissolved in 15 mL H₂O via ultrasonic treatment. Subsequently, the solution was transferred into a 20 mL Teflon autoclave and a piece of clear NF was added, which was heated at 100 °C for 12 hours. Finally, the resulted NF was washed three times with ethanol and distilled water and dried at 60 °C for 12 hours, which was nominated as FeMnCr_{0.6}OOH/NF. For comparison, FeMnCr_nOOH/NF (n = 0.4, 0.6, 0.8 and 1.0), FeOOH/NF, MnOOH/NF, CrOOH/NF, FeMnOOH/NF, MnCr_{0.6}OOH/NF, FeCr_{0.6}OOH/NF, FeMnNiOOH/NF, FeMnCoOOH/NF, FeMnCuOOH/NF, FeMnZnOOH/NF, and FeMnScOOH/NF samples were

synthesized by a similar method. FeMnCr_{0.6}O_x/NF was prepared by calcining FeMnCr_{0.6}OOH/NF in air at 500 °C for 2 h.

Characterization

Powder X-ray diffraction (XRD) patterns were collected by a Rigaku D/MAX-2600 scanning. Transmission electron microscope (TEM) images were observed by a FEI Tecnai F20. High-resolution TEM (HRTEM) images were recorded on a JEM-2100 transmission electron microscope at 200 kV. Scanning electron microscopy (SEM) images were recorded on a ZEISS GeminiSEM 300 field emission scanning electron microscope. X-ray photoelectron spectroscopy (XPS) was recorded on a Thermo Fisher Scientific Nexsa G2.

Electrochemical measurements

All electrochemical measurements were performed on a CHI760E electrochemical working station using a typical three-electrode configuration in 1 M KOH aqueous solution. Linear sweep voltammetry (LSV) polarization curves were acquired at a scan rate of 5 mV·s⁻¹. Electrochemical impedance spectroscopy (EIS) measurements were performed at open-circuit potential in the frequency range from 100 kHz to 0.1 Hz with an a.c. perturbation of 5 mV. All potentials measured were calibrated to RHE using the following equation: $E \text{ (versus RHE)} = E \text{ (versus SCE)} + 0.241 \text{ V} + 0.0591 \text{ pH}$.

The Faradaic efficiency was calculated using the equations:

$$FE = (V/V_m) e N_A Z/Q$$

where V is the volume of gas (L), V_m is the standard molar volume (22.4 L mol⁻¹), e is the electron charge, N_A is the Avogadro number (6.02×10^{23}), Z is the number of electrons needed to form O₂ molecular (for O₂, $Z = 4$), and Q is the amount of power consumed during electrolysis (C)

The TOF value was calculated using the equations:

$$TOF = J_A A/(4 \cdot n \cdot F)$$

where J_A is the current density at 95% iR-corrected overpotential = 300 mV, A is the area of the working electrode (cm²), n is the amount of substance at the active site (mol), F is the Faraday constant.

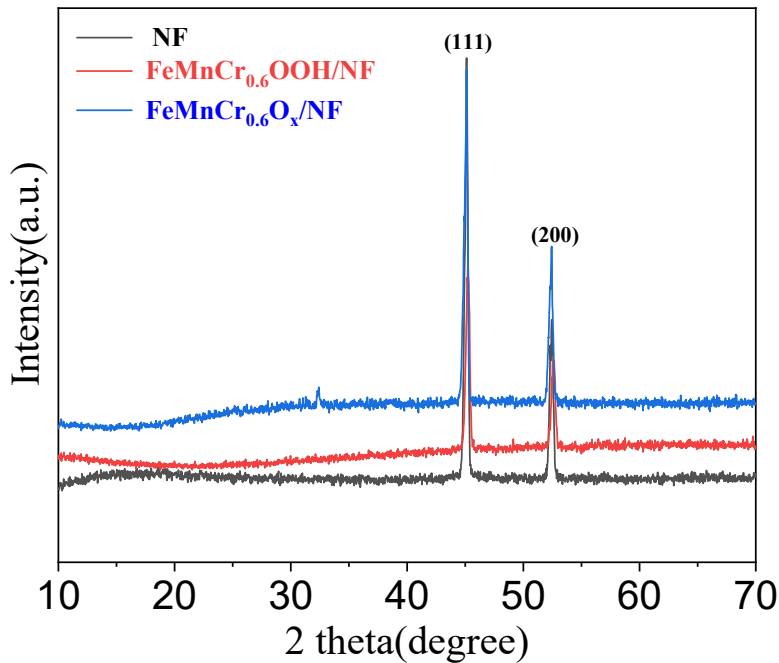


Fig. S1. XRD patterns of NF, FeMnCr_{0.6}OOH/NF and FeMnCr_{0.6}O_x/NF.

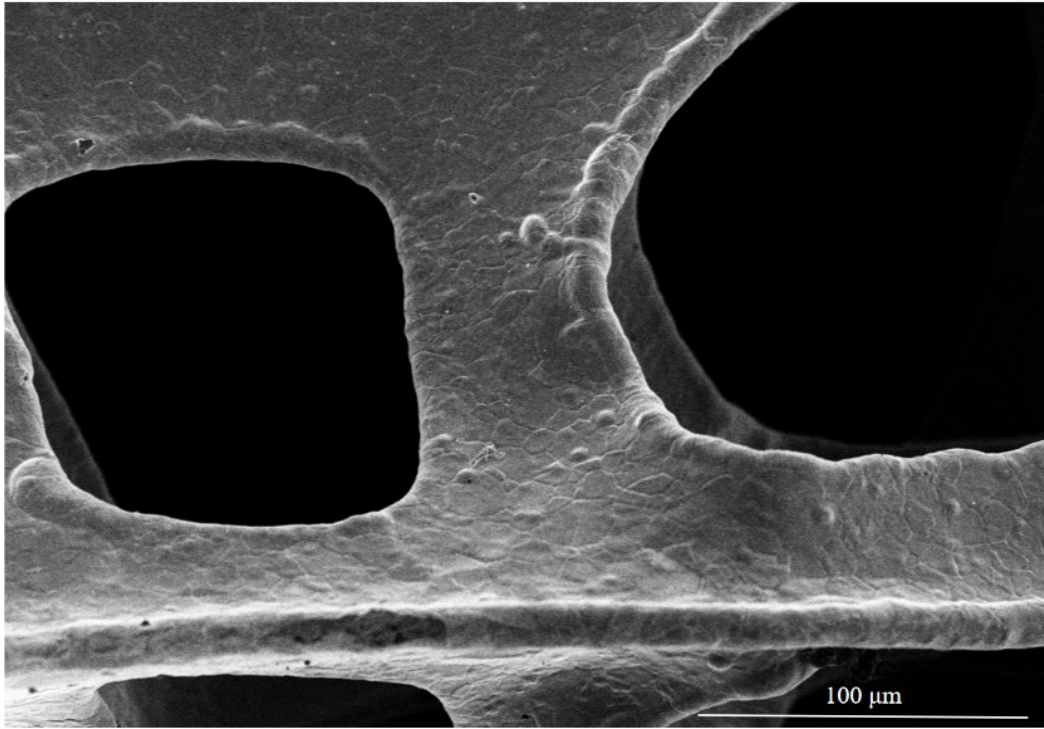


Fig. S2. SEM image of NF.

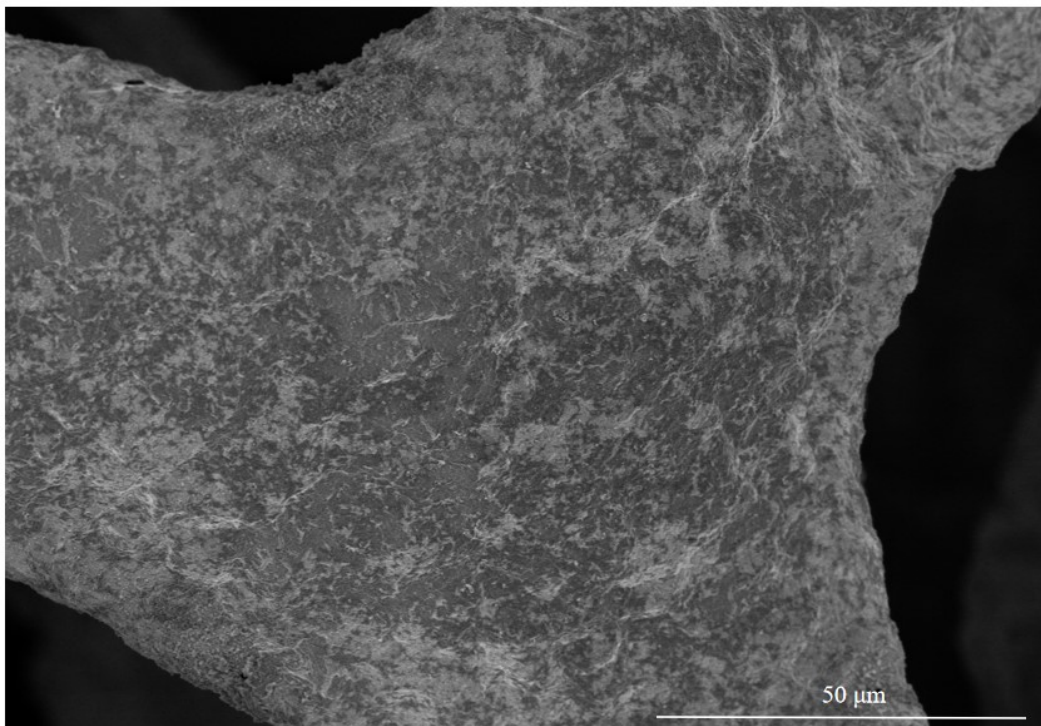


Fig. S3. SEM image of FeMnCr_{0.6}OOH/NF.

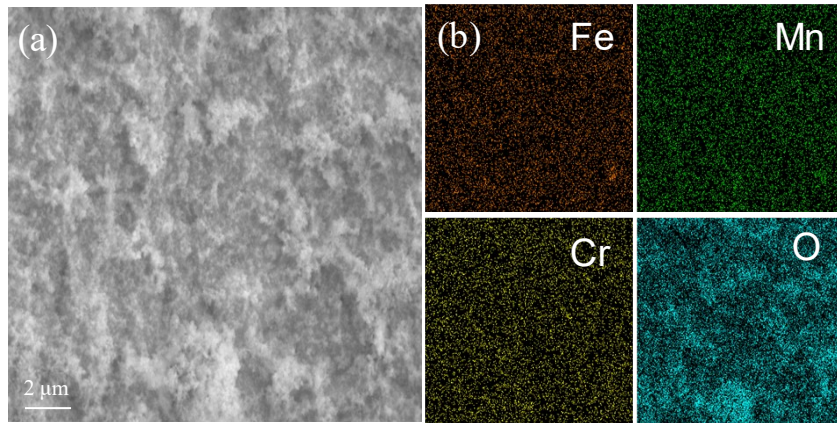


Fig. S4. (a) SEM image and (b) element mapping of FeMnCr_{0.6}OOH/NF.

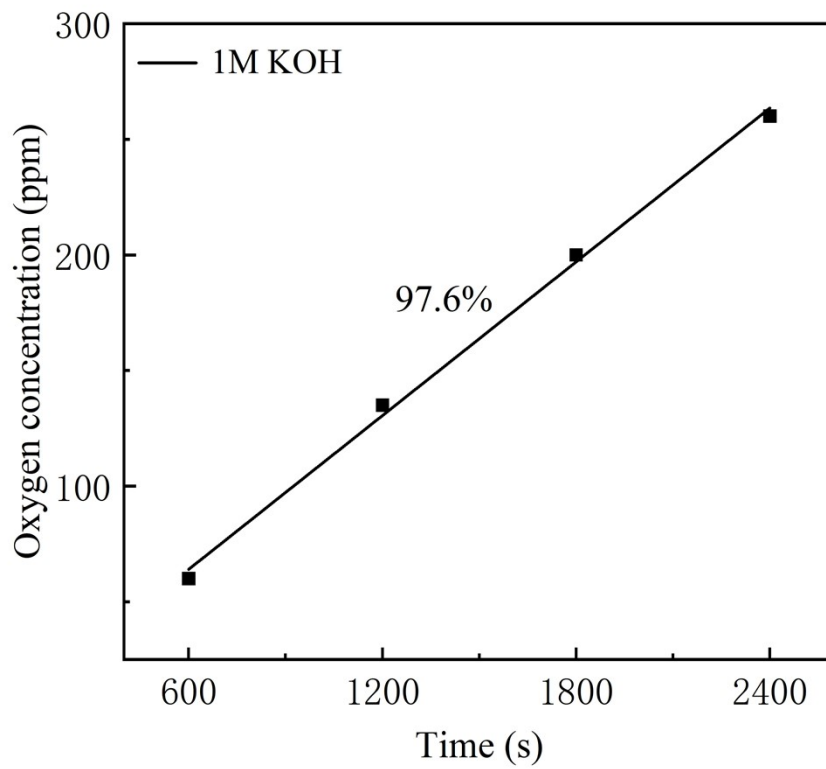


Fig. S5. Faraday efficiency of FeMnCr_{0.6}OOH/NF in 1M KOH at 10 mA cm⁻².

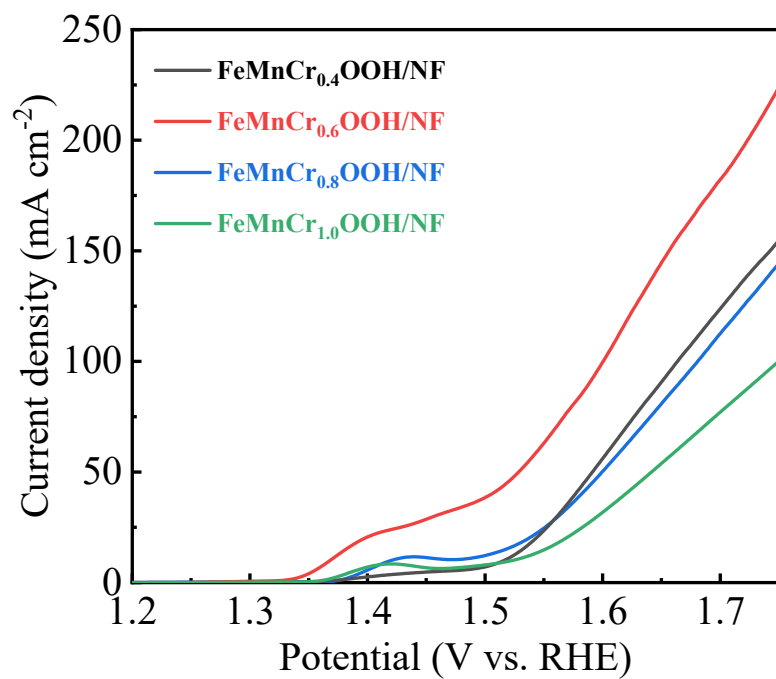


Fig. S6. (a) OER polarization curves of FeMnCr_nOOH/NF ($n = 0.4-1.0$) at a scan rate of 5 mV s^{-1} in 1M KOH.

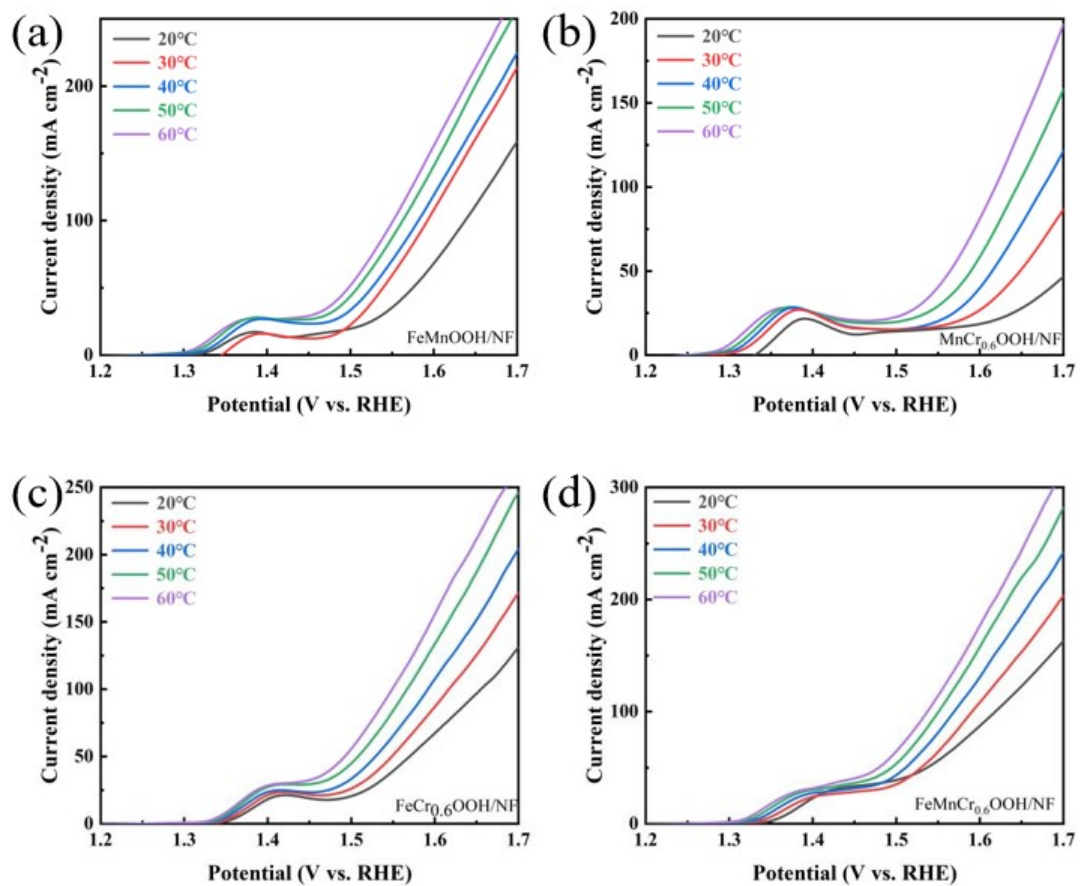


Fig. S7. OER polarization curves of FeMnOOH/NF, MnCr_{0.6}OOH/NF, FeCr_{0.6}OOH/NF and FeMnCr_{0.6}OOH/NF in 1 M KOH with scan rate 5 mV s⁻¹ at 20°C, 30°C, 40°C, 50°C, and 60°C.

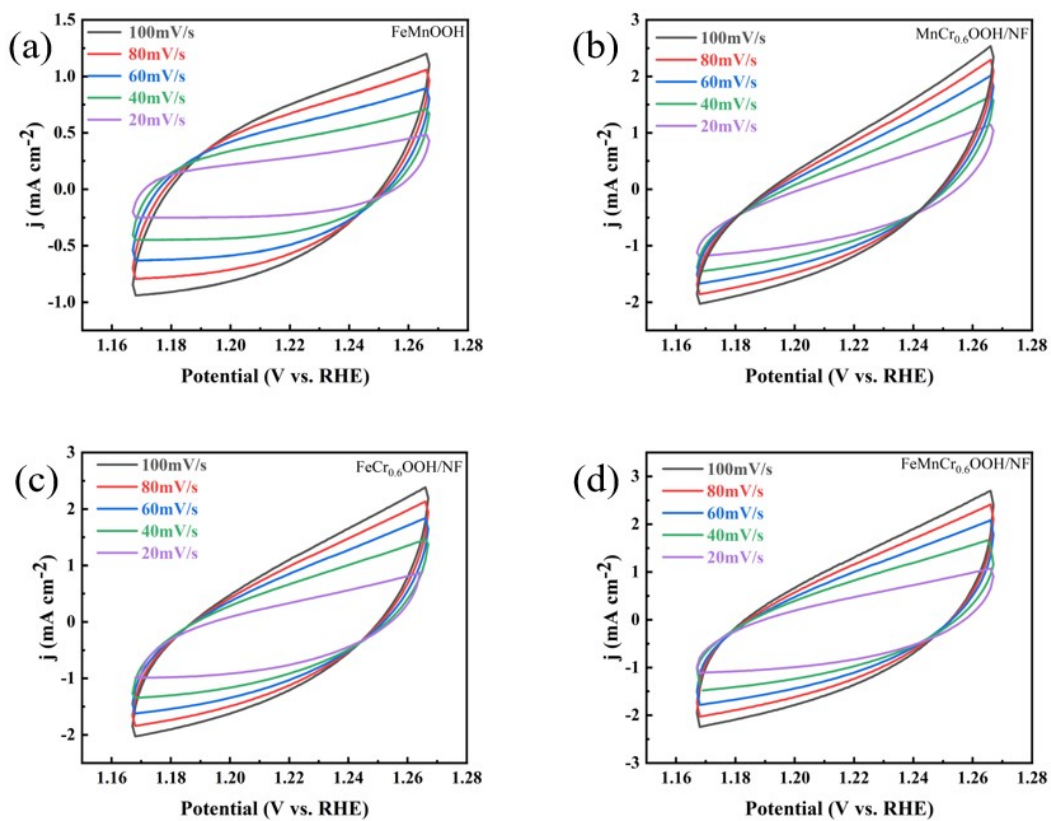


Fig. S8. CVs of (a) FeMnOOH/NF, (b) MnCr_{0.6}OOH/NF, (c) FeCr_{0.6}OOH/NF, and (d) FeMnCr_{0.6}OOH/NF measured in a non-Faradaic region at different scan rate in 1 M KOH.

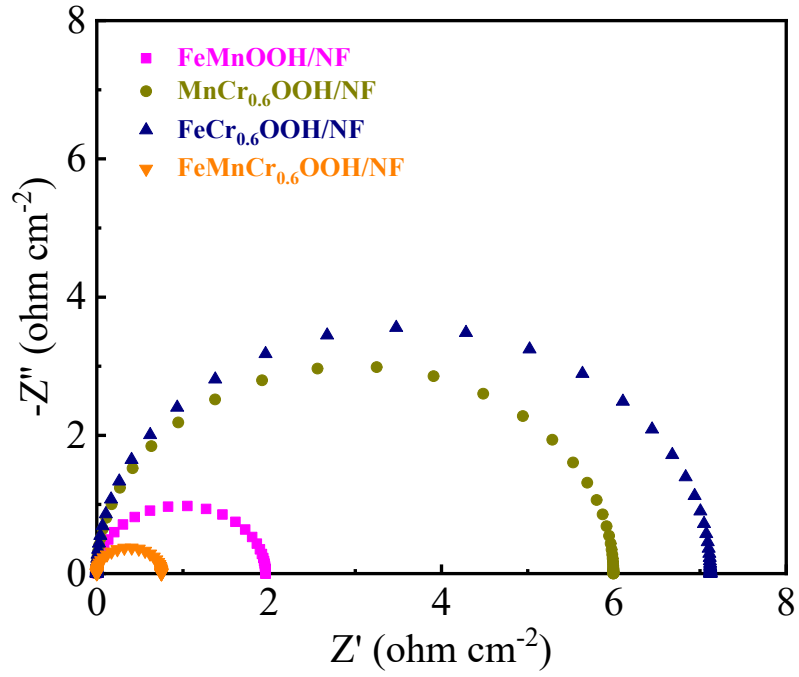


Fig. S9. Nyquist plots of the EIS test for the FeMnOOH/NF, MnCr_{0.6}OOH/NF, FeCr_{0.6}OOH/NF, and FeMnCr_{0.6}OOH/NF.

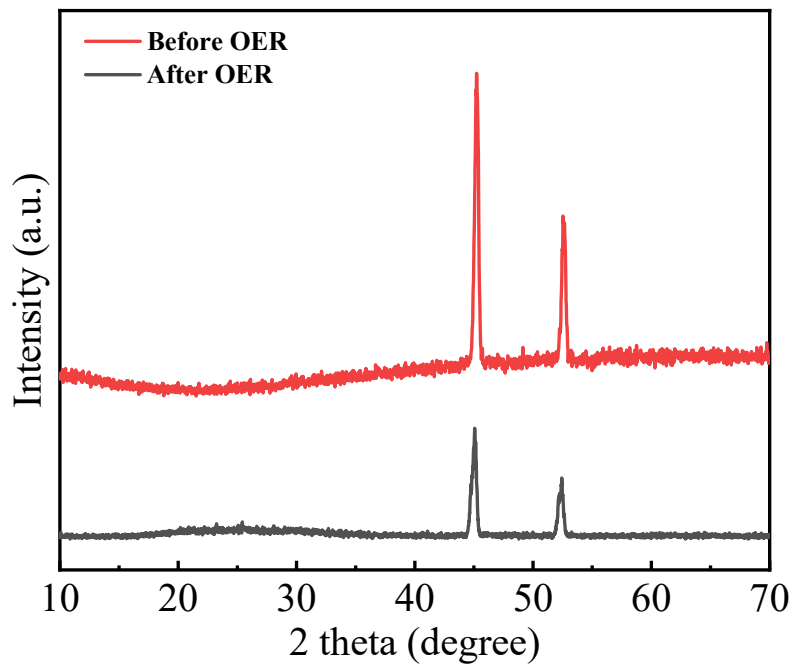


Fig. S10. XRD patterns of FeMnCr_{0.6}OOH/NF before and after OER.

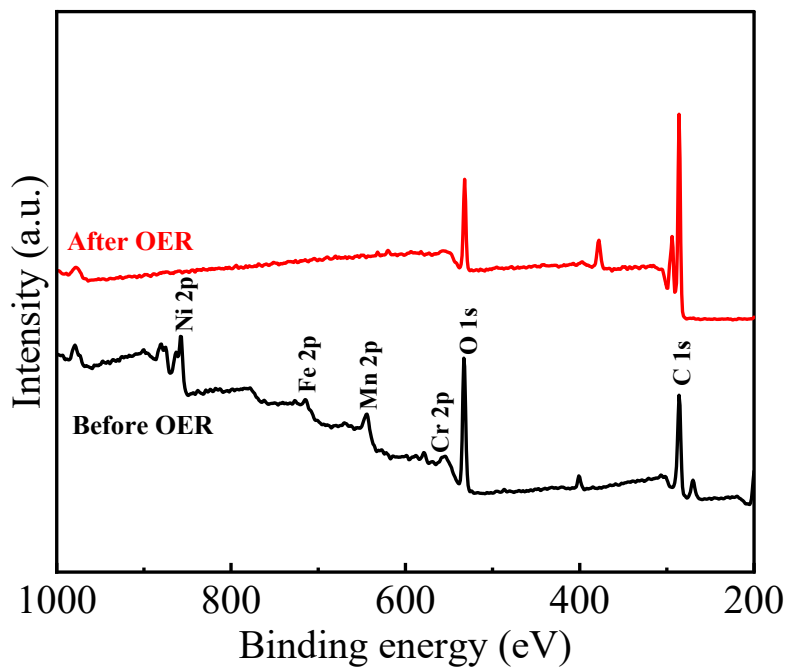


Fig. S11. XPS spectra of FeMnCr_{0.6}OOH/NF before and after OER.