

Supporting Information for
**Green Electrochemical Upcycling of PET Waste via
Bimetallic Catalyst-Driven Ethylene Glycol Selective
Oxidation to Formate**

Contents of the Supporting Information

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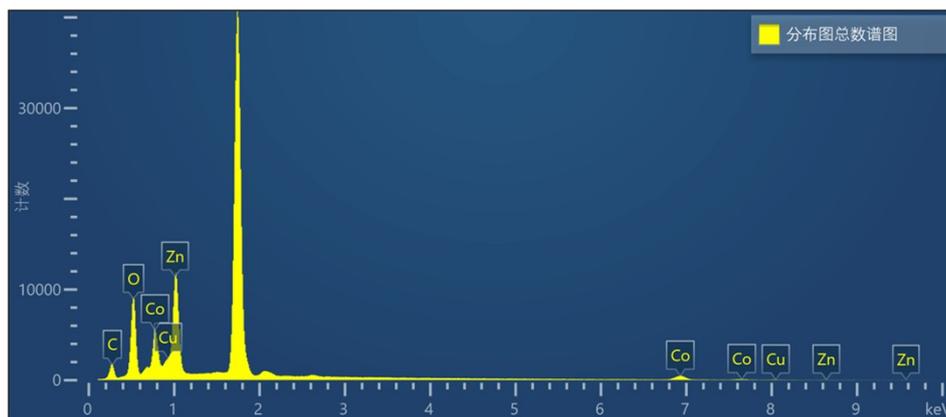


Fig. S1 The EDS analysis of CuO/ZnCuCo₂O₄ catalysts.

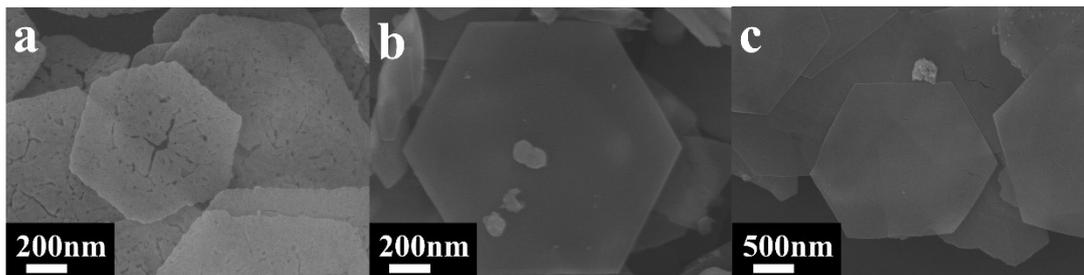


Fig. S2 SEM images of (a) Co_3O_4 , (b) ZnCo_2O_4 , (c) $\text{CuO}/\text{CuCo}_2\text{O}_4$.

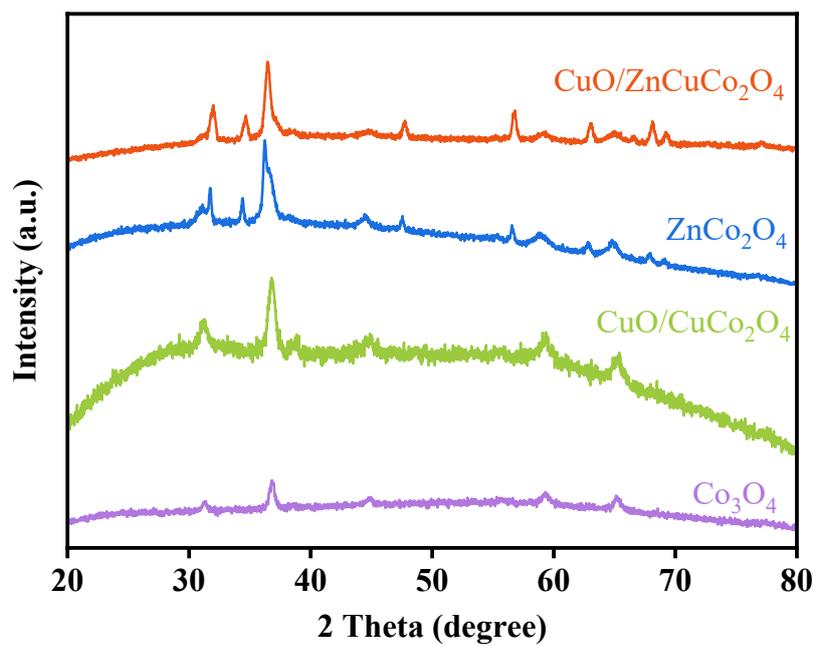


Fig. S3 The XRD patterns of Co_3O_4 , ZnCo_2O_4 , CuCo_2O_4 and $\text{CuO}/\text{ZnCuCo}_2\text{O}_4$ samples.

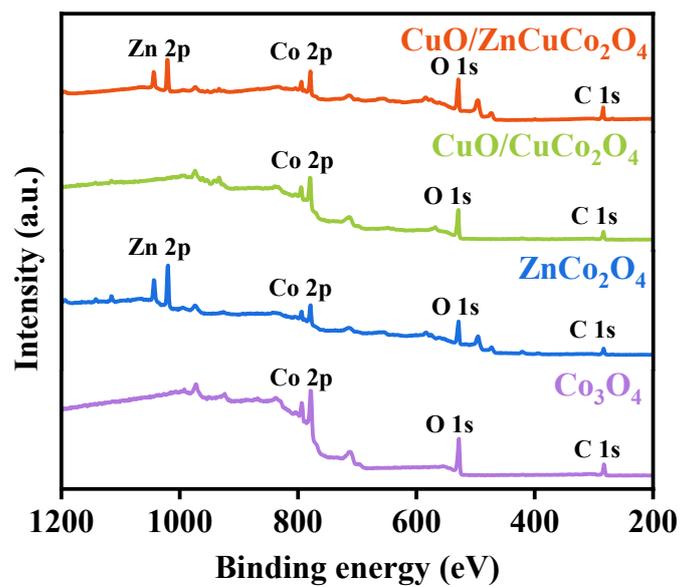


Fig. S4 Survey XPS spectrum of Co₃O₄, ZnCo₂O₄, CuO/CuCo₂O₄ and CuO/ZnCuCo₂O₄.

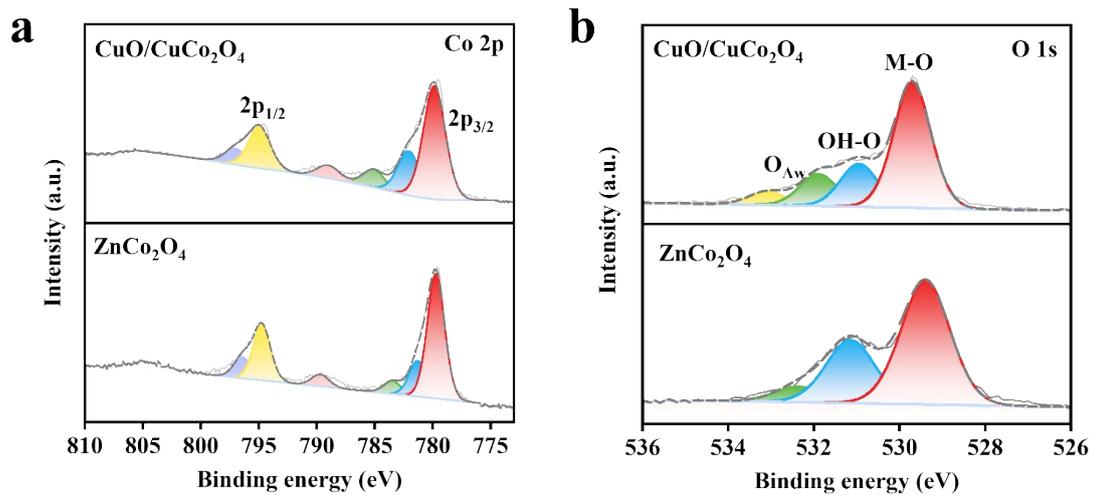


Fig. S5 High-resolution (a) Co 2p spectra, (b) O 1s XPS spectra of ZnCo₂O₄ and CuO/CuCo₂O₄.

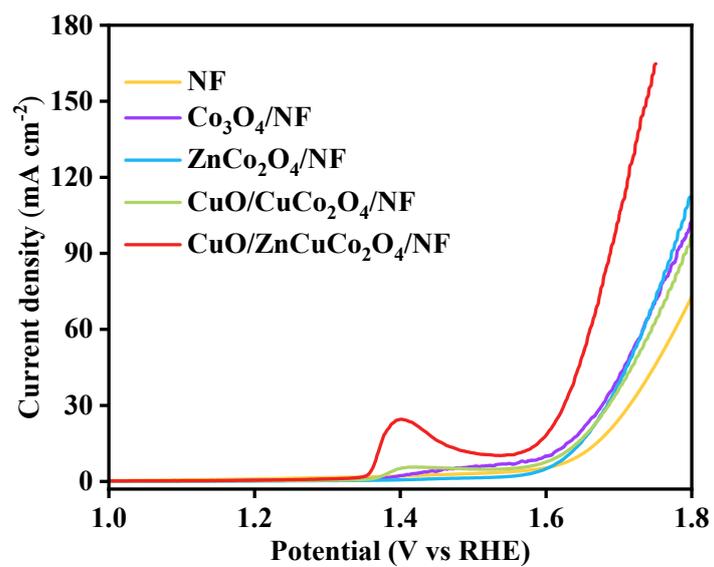


Fig. S6 LSV plots of CuO/ZnCuCo₂O₄/NF and comparison samples in 1 M KOH for OER.

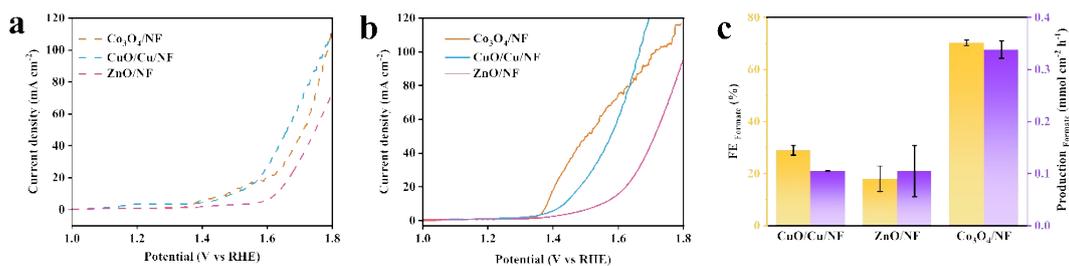


Fig. S7 (a) LSV plots with Co_3O_4 and single-metal catalysts added without EG, (b) with EG. (c) FE and production of converting EG to formate under the condition of 1.6 V vs RHE for Co_3O_4 and its comparative catalyst.

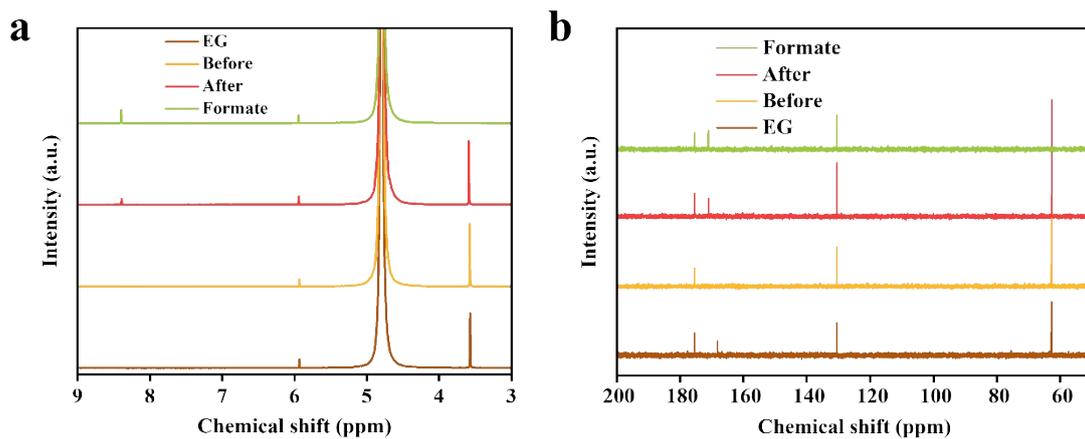


Fig. S8 Products before and after electro-reforming on $\text{CuO}/\text{ZnCuCo}_2\text{O}_4/\text{NF}$ electrode (a) ^1H NMR spectra, (b) ^{13}C NMR spectra.

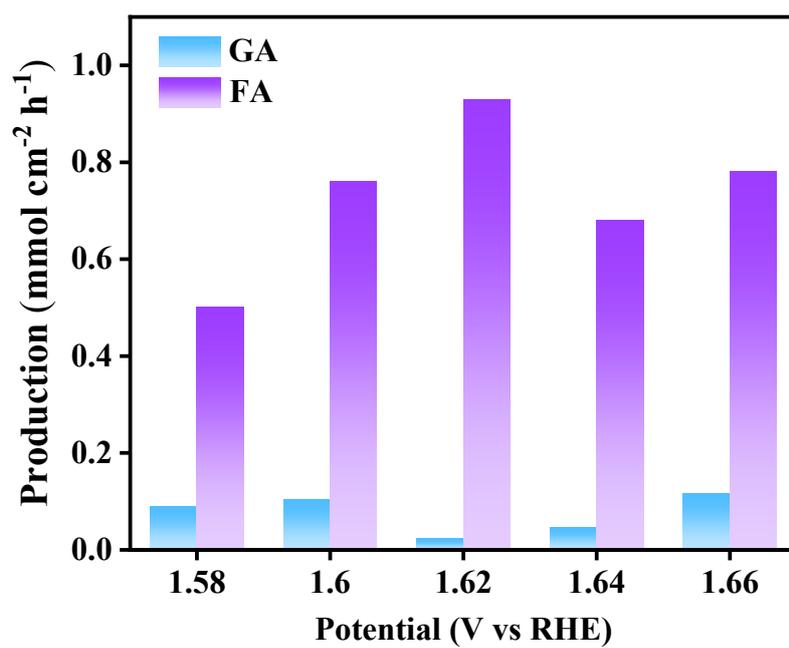


Fig. S9 The production of CuO/ZnCuCo₂O₄/NF under different potentials.

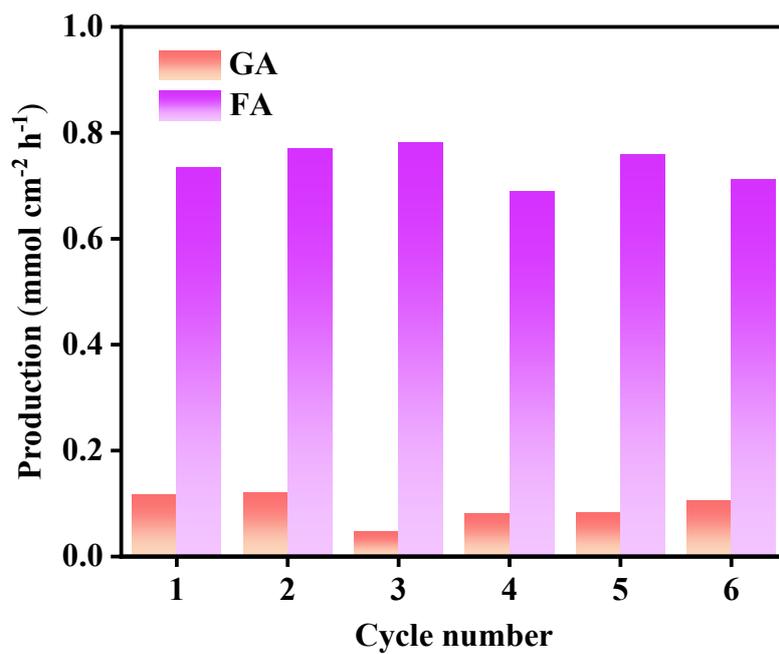


Fig. S10 The yields of CuO/ZnCuCo₂O₄/NF with consecutive six cycles at 1.62 V vs RHE.

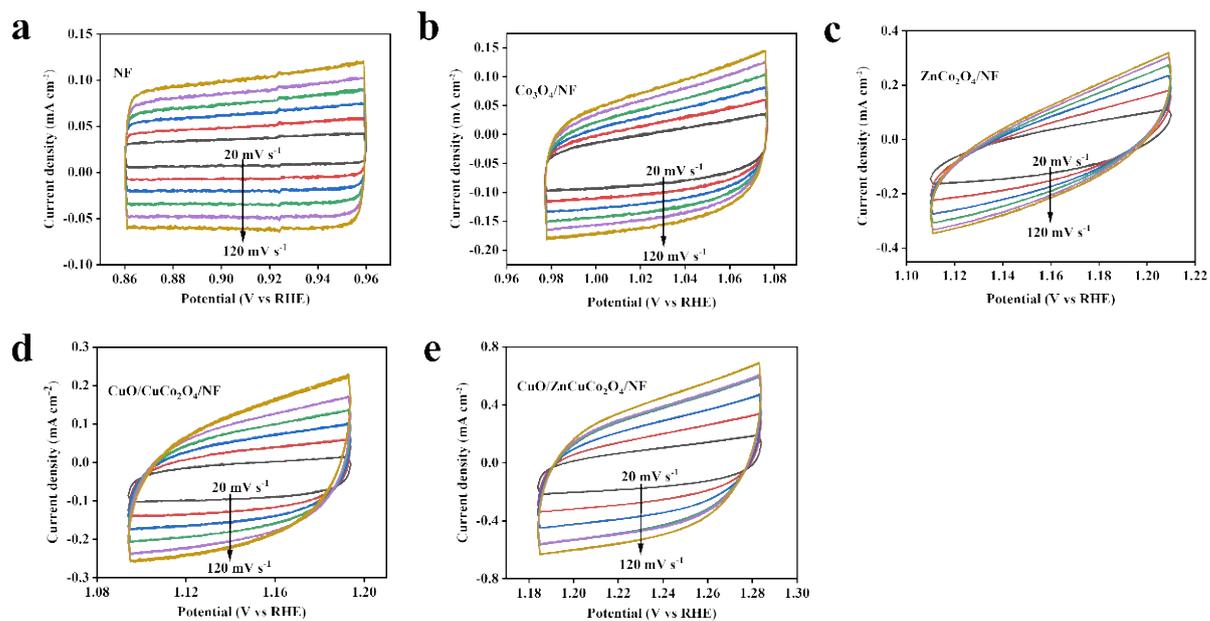


Fig. S11 The CV curves of bare NF (a), Co₃O₄/NF (b), ZnCo₂O₄/NF (c), CuO/CuCo₂O₄/NF (d) and CuO/ZnCuCo₂O₄/NF (e) samples.

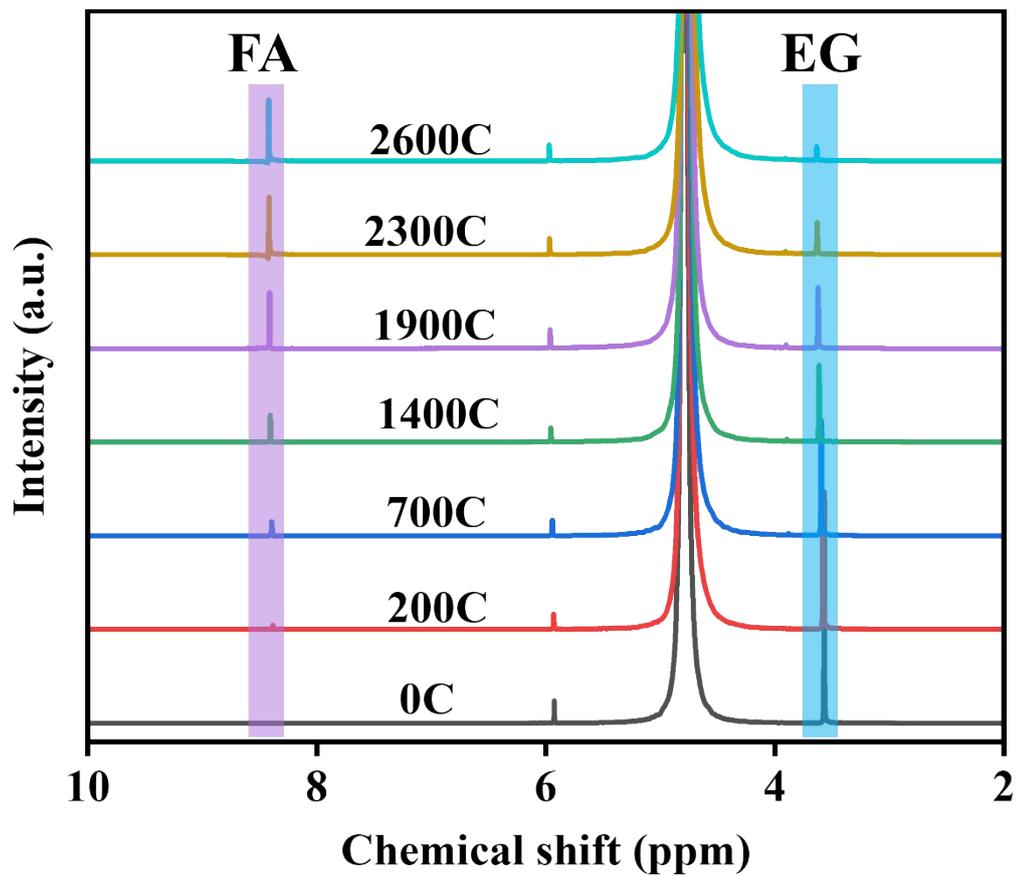


Fig. S12 ^1H NMR measurements of EG oxidation on $\text{CuO}/\text{ZnCuCo}_2\text{O}_4/\text{NF}$ electrode.

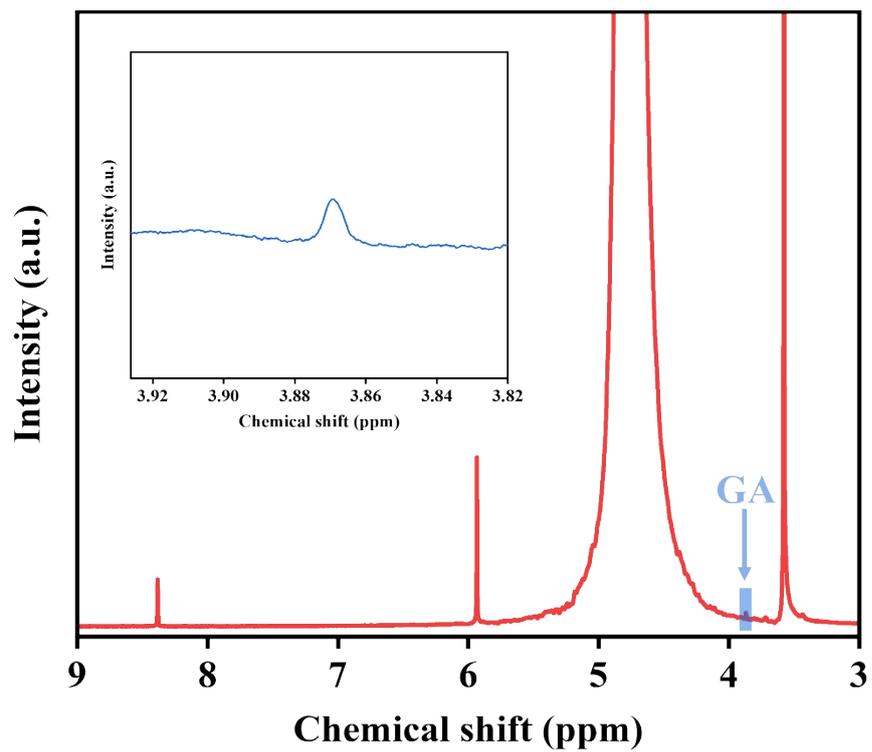


Fig. S13 ^1H NMR of $\text{CuO}/\text{ZnCuCo}_2\text{O}_4/\text{NF}$ at an input charge of 200C.

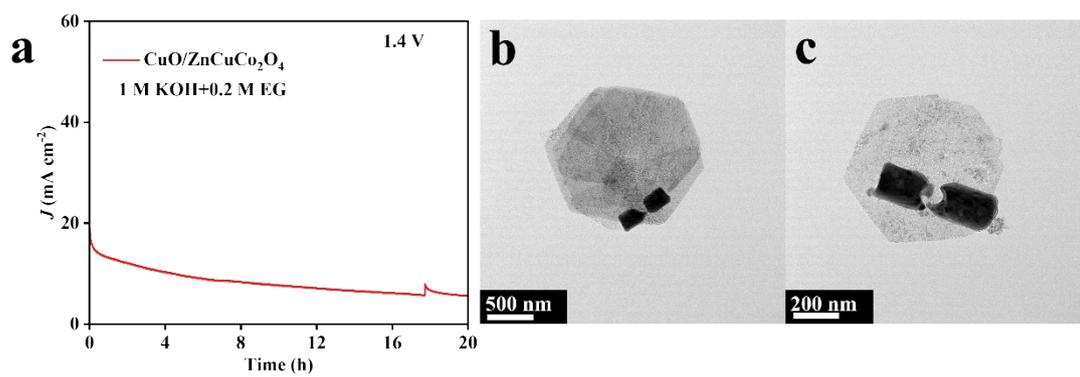


Fig. S14 (a) Chronoamperometric measurement of CuO/ZnCuCo₂O₄/NF for EGOR. (b) (c) TEM images of CuO/ZnCuCo₂O₄ catalyst at different magnifications after 20 hours.

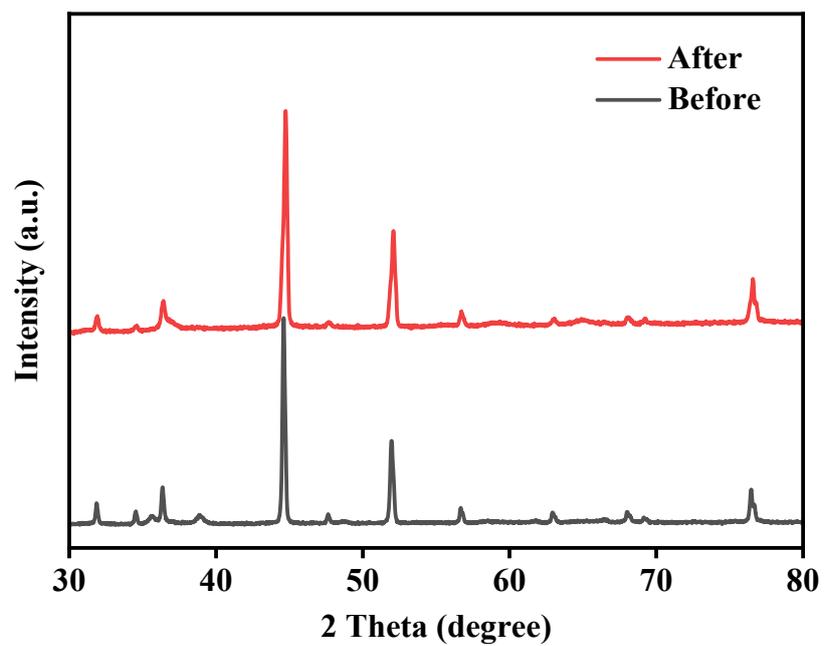


Fig. S15 XRD patterns of CuO/ZnCuCo₂O₄/NF electrode before and after reaction.

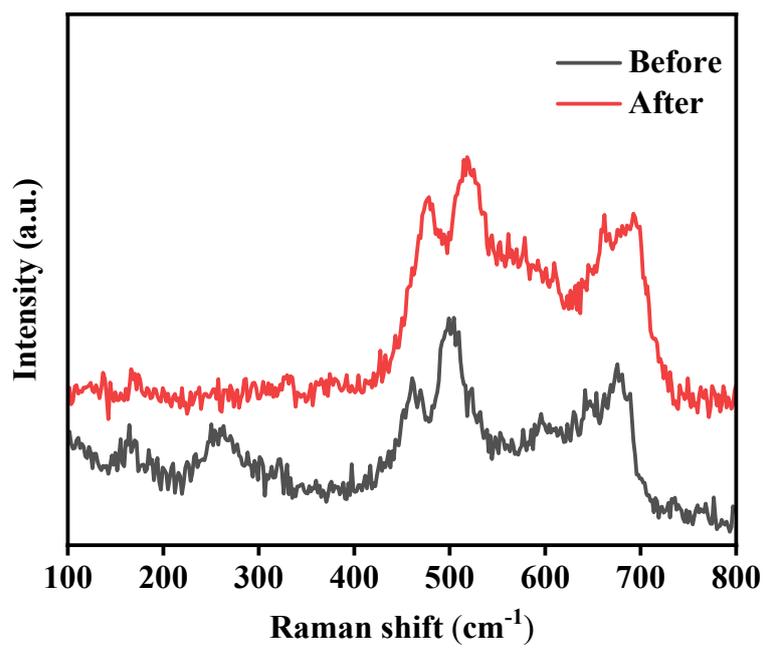


Fig. S16 Raman patterns of CuO/ZnCuCo₂O₄/NF electrode before and after electrocatalysis.

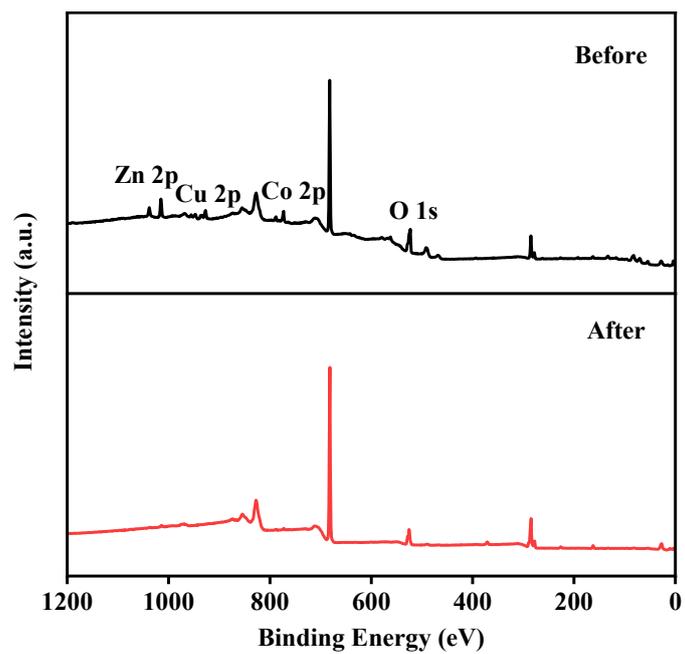


Fig. S17 Survey XPS spectra of CuO/ZnCuCo₂O₄/NF electrode before and after electrolysis.

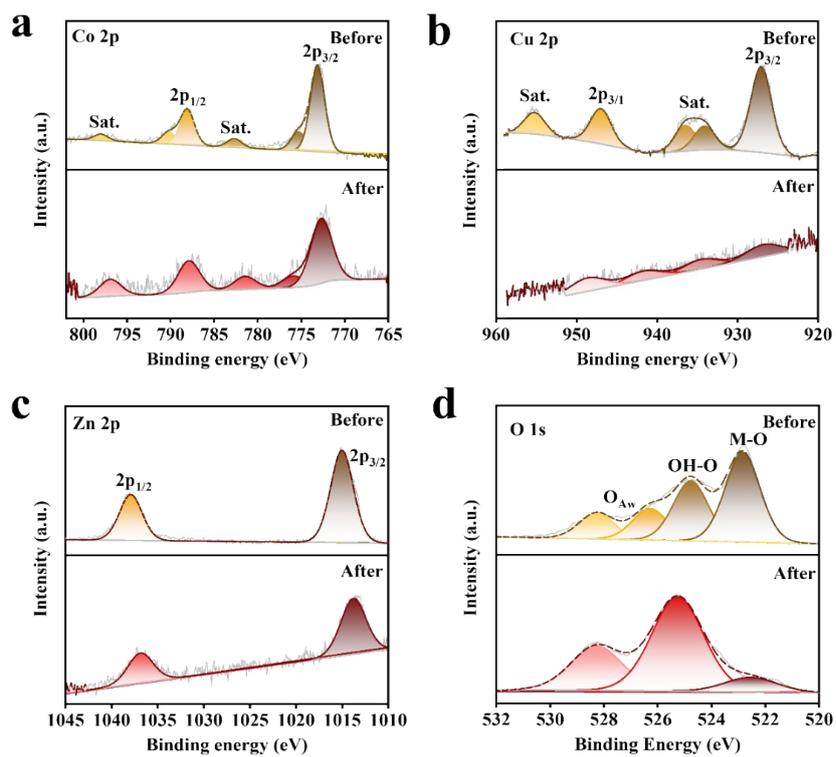


Fig. S18 XPS spectra of CuO/ZnCuCo₂O₄/NF electrode before and after electrolysis.

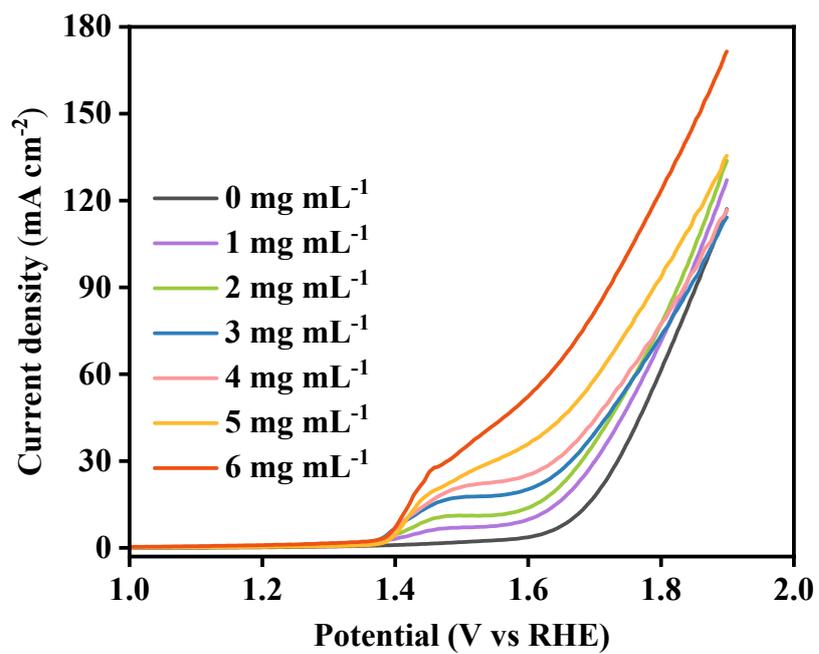


Fig. S19 LSV curves of CuO/ZnCuCo₂O₄/NF in 1 M KOH with different concentrations of PET.

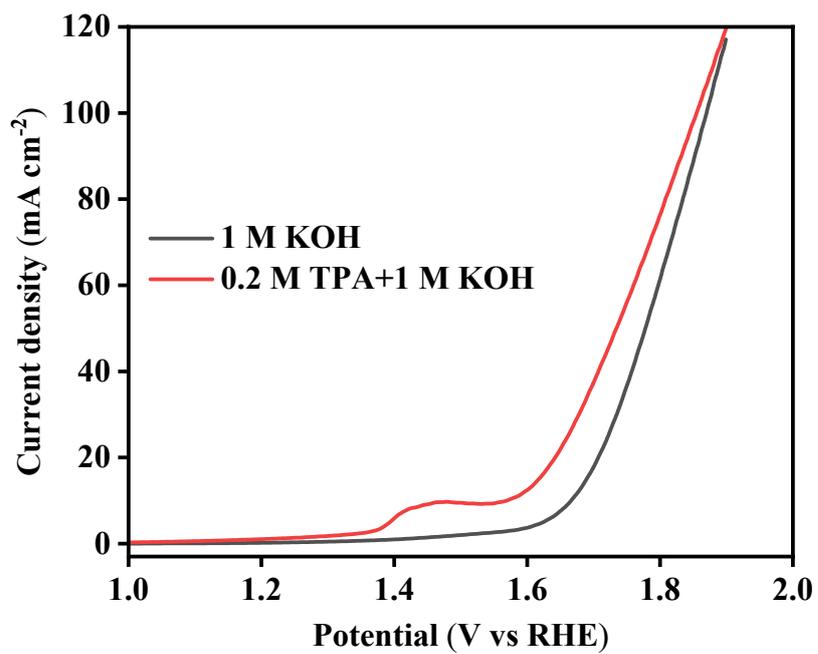


Fig. S20 LSV plots for CuO/ZnCuCo₂O₄/NF electrode in 1 M KOH with and without TPA addition.

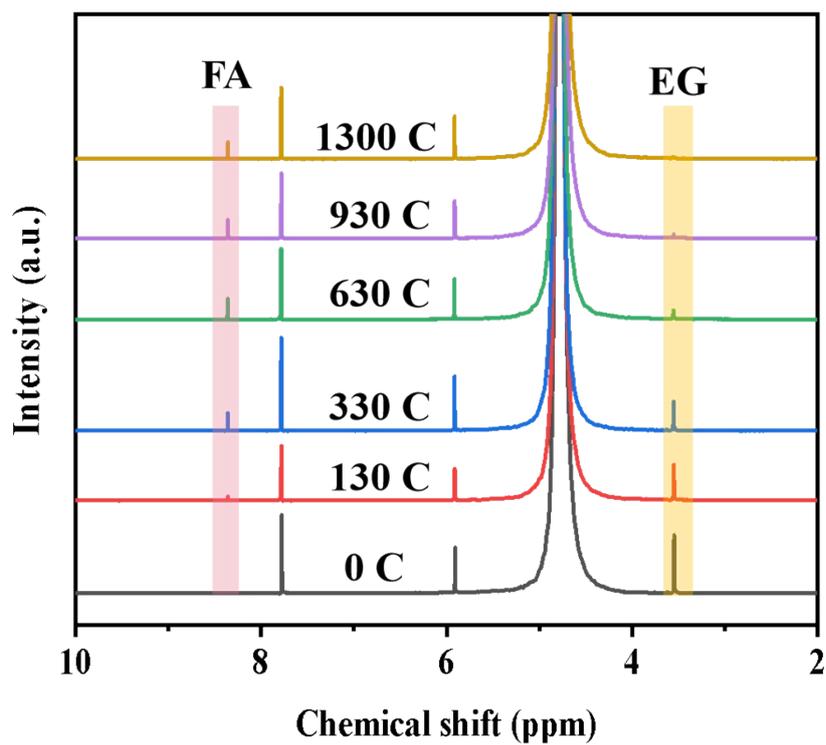


Fig. S21 ^1H NMR measurements of PET hydrolysate oxidation on $\text{CuO}/\text{ZnCuCo}_2\text{O}_4/\text{NF}$ electrode.

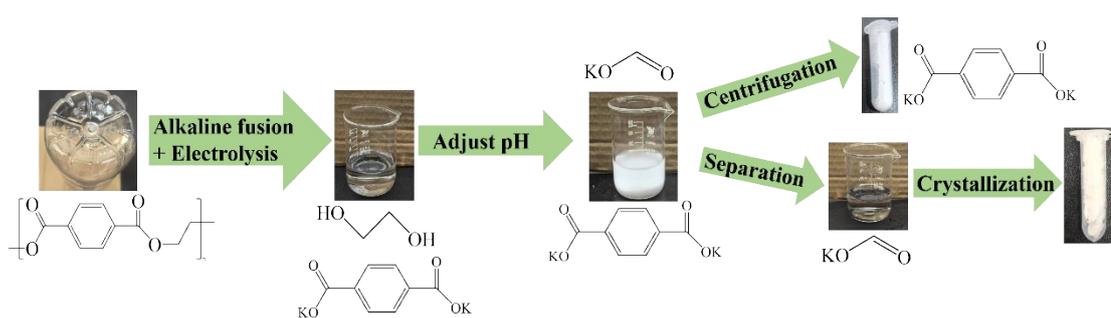


Fig. S22 Schematic illustration of the process of PET electro-reforming and product separation.

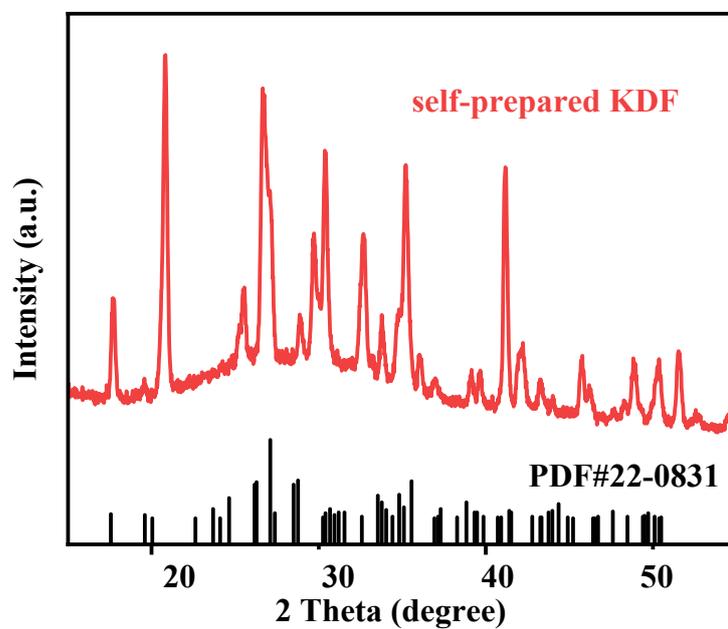


Fig. S23 The XRD pattern of KDF prepared from PET electrolyte.

Table S1. Comparison of state-of-the-art catalysts for ethylene glycol and PET hydrolysate.

Catalysts	Electrolyte	Stability	Overpotential	Current density (mA cm ⁻²)	FE (%)	Formate yield (mmol cm ⁻² h ⁻¹)	Ref.
CuO/ZnCu Co₂O₄/NF	1 M KOH + 0.2 M EG	20 H (1.4 vs RHE)	271 mV@100 mA cm ⁻²	100 (1.42 vs RHE)	95.5	0.78	This work
NiFe- LDH/NF	0.5 M EG + 1 M KOH	24 H (1.4 vs RHE)	122 mV@100 mA cm ⁻²	100 (1.47 vs RHE)	94.3	-	[1]
Ni_{0.9}Mn_{0.1} LDH	1 M KOH + 1 M EG	12 H	180 mV@10 mA cm ⁻²	100 (1.37 vs RHE)	85.8	-	[2]
CoFe- LDH/NF- 10	1 M KOH + 0.1 M EG	-	240 mV@10 mA cm ⁻²	10 (~1.44 vs RHE)	49.7	-	[3]

NiCu_{60s}/NF	1 M KOH + 0.3 M EG	24 H (1.42 V vs RHE)	307 mV@100 mA cm ⁻²	50 (~1.42 vs RHE)	95.8	0.70	[4]
R-NiCo/NF	1.0 M KOH + 0.1 M EG	10 H (1.45 V vs RHE)	264 mV@50 mA cm ⁻²	50 (1.43 vs RHE)	98.2	0.63	[5]
Cu(F)@Cu O@ Ni(OH)₂	1.0 M KOH + 0.1 M EG	24 H (1.4 V vs RHE)	200 mV@50 mA cm ⁻²	50 (1.36 vs RHE)	-	-	[6]
NiCo₂O₄	1.0 M KOH + 0.3 M EG	18 H (1.5 V vs RHE)	244 mV@150 mA cm ⁻²	100 (1.40 V vs RHE)	90.4	-	[7]

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