

Supporting information

Anodic fabrication of nanostructured Cu_xS and CuNiS_x thin films and their hydrogen evolution activities in acidic electrolytes

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Figure S1

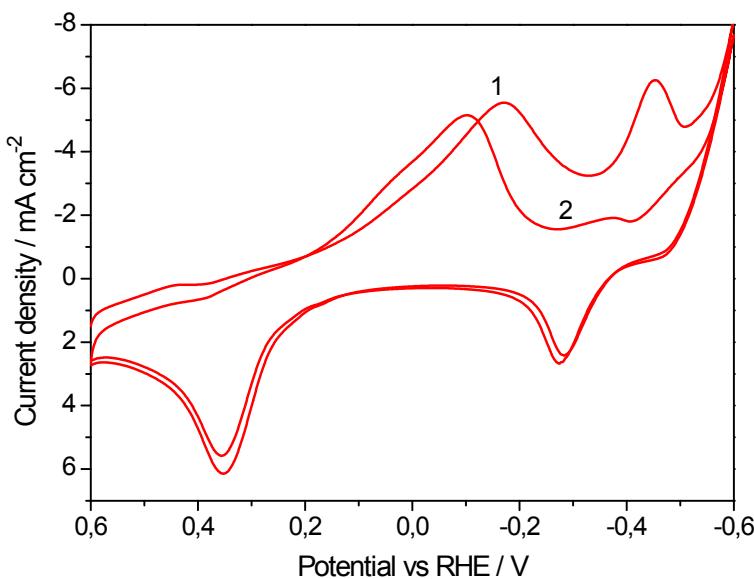


Figure S1 : Repeated cyclic voltammograms of CuxS in 0.5 M Na₂SO₄. 1 – Represent the first cycle, 2 indicates the lower reduction current density for the second cathodic peak.

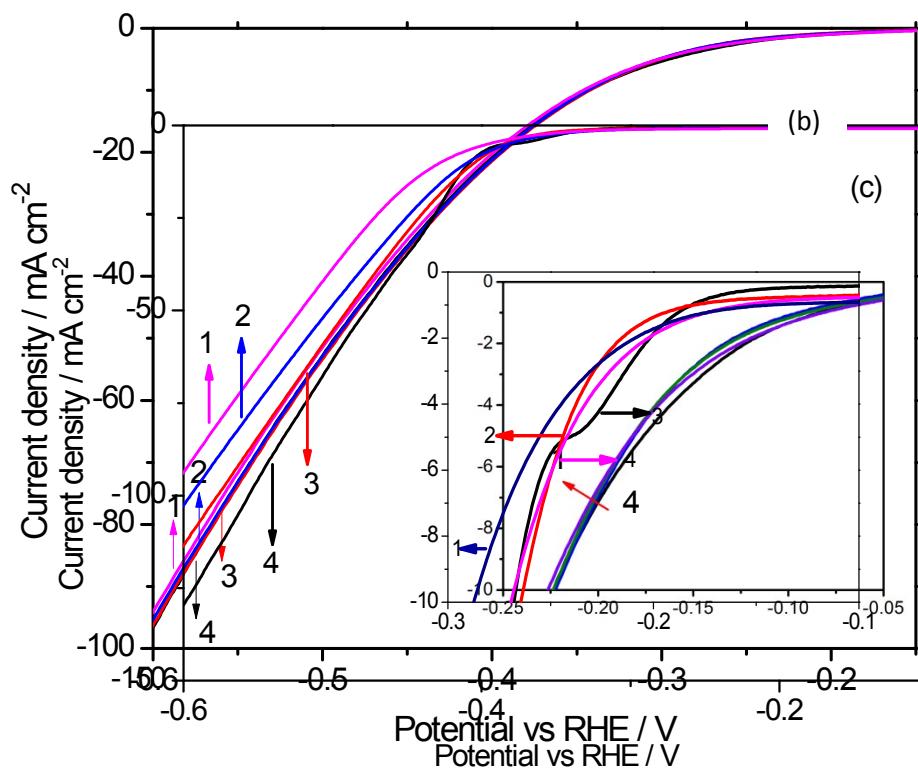
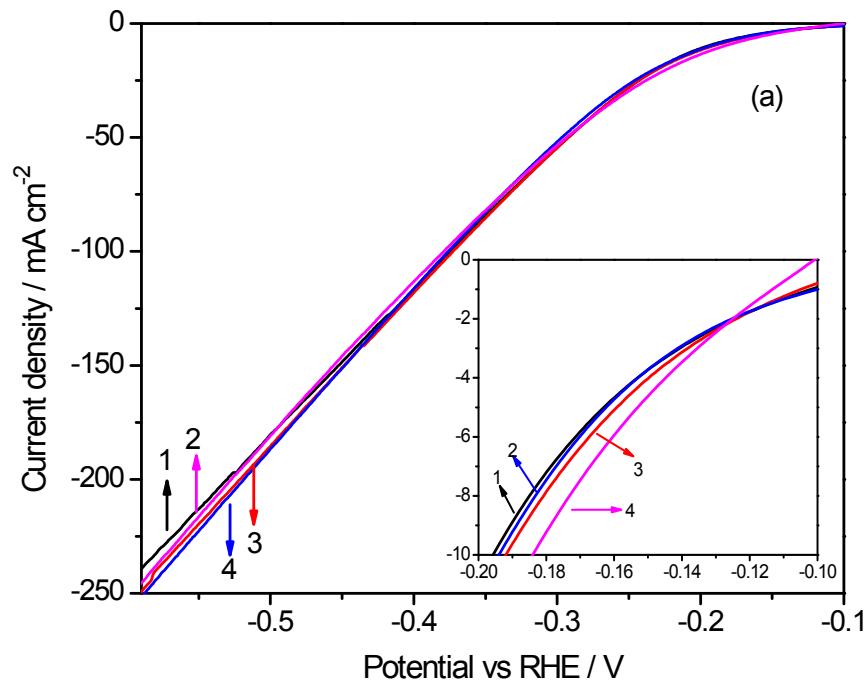
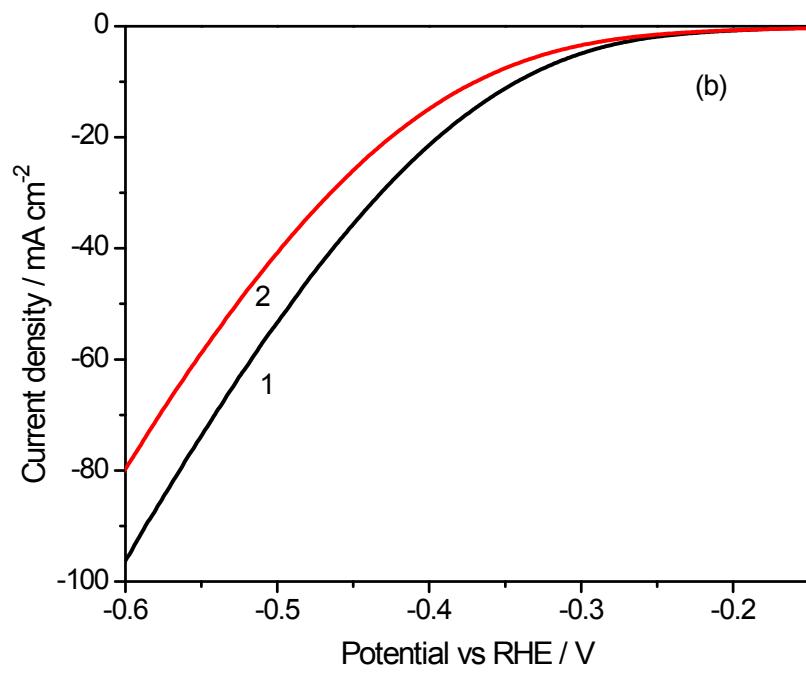
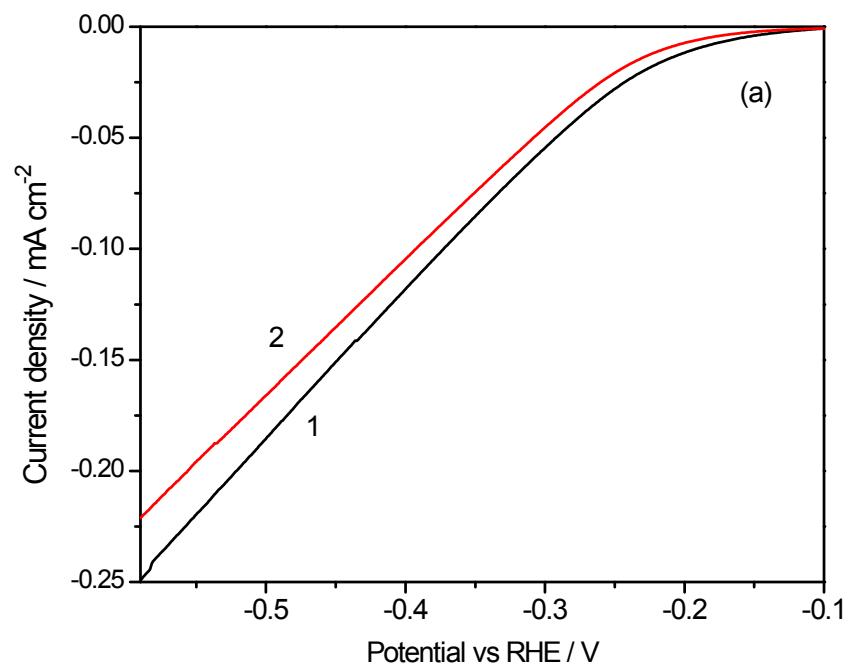


Figure S3 : LSVs of different electrodes before and after chronoamperometric stability tests a) CuNiS_x, (b) CuNi, (c) Cu_xS



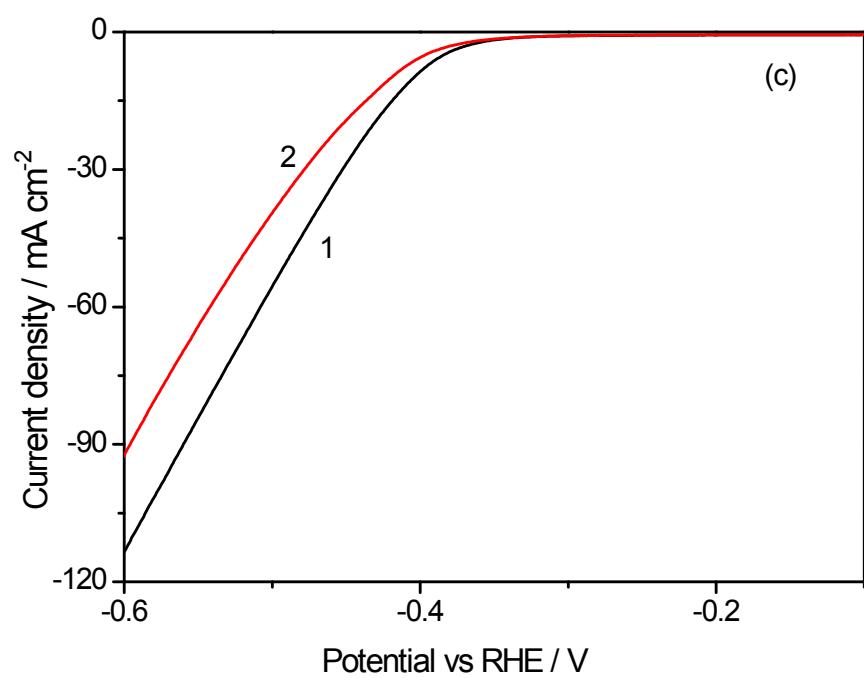
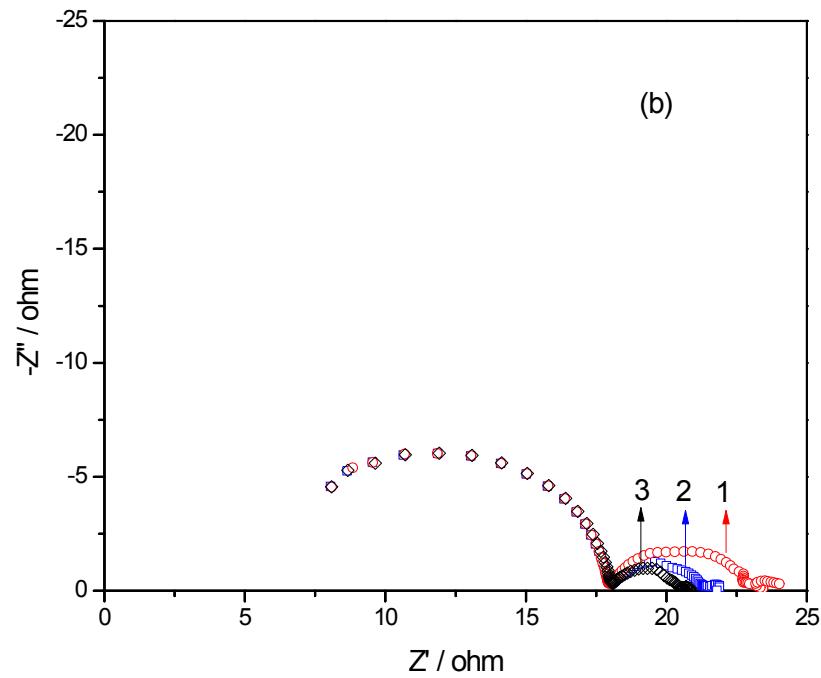
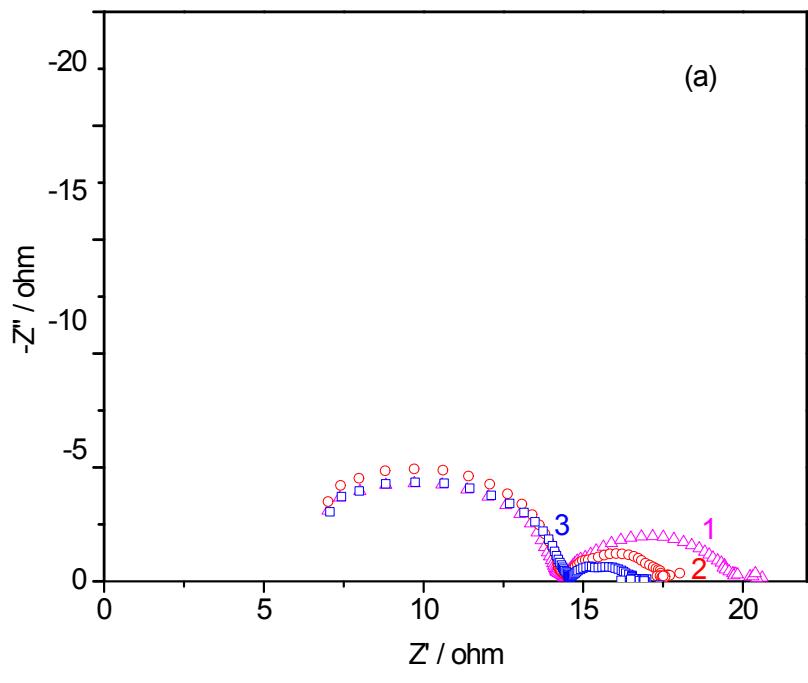


Figure S4:Potential dependant Nyquist plots of (a) CuNiS_x, (b) CuNi, (c) Cu_xS in 0.5 M H₂SO₄. 1 – at -0.2 V , 2 at -0.3 V and 3 at -0.4 V vs RHE



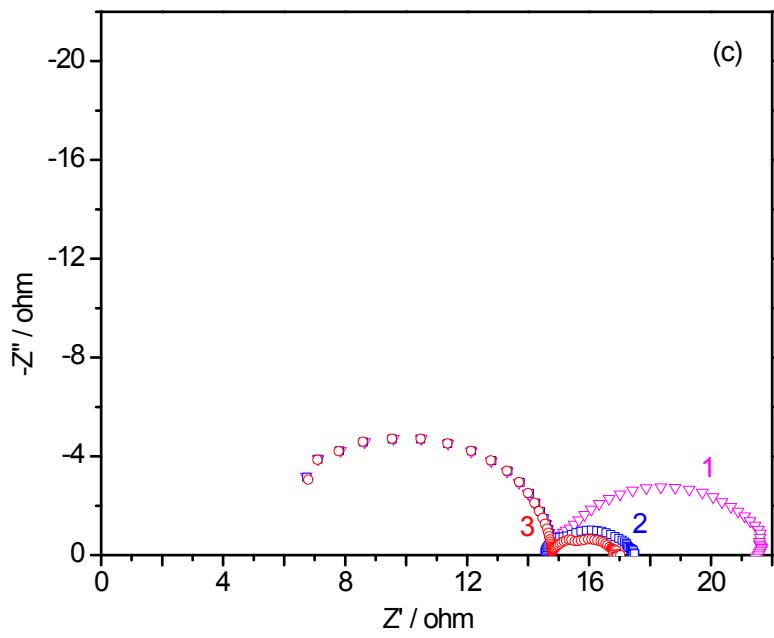
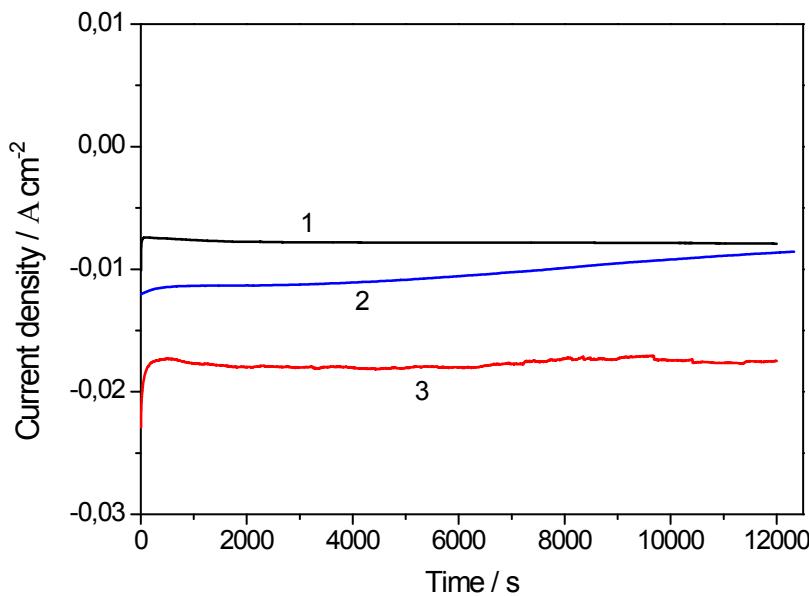


Figure S5: Constant potential stability test of (1) Cu_xS, (2) CuNi, (3) CuNiS_x performed in 0.5M H₂SO₄ using carbon rod as counter electrode.



At higher oxygen evolution rates, the surface of carbon rod undergo formation of loose carbon particles and come into the electrolyte solution. However the current density behaviour are similar to the Pt counter.

Table S1: Comparison of overpotential requirement and Tafel slope values for different electrocatalyst and the present thin film electrode materials.

Electrocatalyss	Cathodic overpotential for 10 mA cm ⁻² in 0.5 M H ₂ SO ₄ / mv	Tafel Slope / mV/dec
Pt/C	30	30
MoS ₂	230	101
MoS ₂ /RGO	150	40
MoS ₂ -CoSe ₂	68	36
CoSe ₂	130	48
NiSe ₂	230	57
NiS ₂	240	42
CoS ₂	230	44
FeS ₂	270	62
*Cu _x S	270	102
*CuNiS _x	213	98
*CuNi	195	128
Cu - bare	465	158

‘present work, direct deposition by electrochemical method, without any conductive binders

References

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