## **Electronic Supplementary Information**

## Selectively electrolyzing CO<sub>2</sub> to ethylene by Cu-Cu<sub>2</sub>O/rGO catalyst derived from copper hydroxide nanostrands/graphene oxide

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- 1. Figures S1-S13.
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**Fig. S1.** SEM images of (a) Copper hydroxide nanostrands (CHNs); (b) S9 CNHS/GO before electrochemical reduction; (c) S11 CNHS/GO before electrochemical reduction.



Fig. S2. SEM images of S10 Cu-Cu<sub>2</sub>O/rGO in large scale.



Fig. S3. XRD patterns of S9-S11 synthesized by in situ electrochemical reduction.



**Fig. S4.** The in-situ Raman spectra of the CHNs/GO electrochemical reduction process in the wavenumber range from 200 to 700 cm<sup>-1</sup>.



**Fig. S5.** (a) The activity measurement of S10 by potential scanning from -0.6 to -2.0 V vs. SHE; (b) the chronoamperometry curves of S10 at different potentials vs. SHE.



Fig. S6. FE of H<sub>2</sub> for CO<sub>2</sub> ERR from S1 to S11 samples at -1.3 V vs. SHE.



Fig. S7. NMR spectrum recorded from the liquid products after  $CO_2$  ERR by using S10 catalyst at -1.3 V vs. SHE.



Fig. S8. NMR spectrum recorded from the liquid products after  $CO_2$  ERR by using S10 catalyst at -1.4 V vs. SHE.



Fig. S9. NMR spectrum recorded from the liquid products after  $CO_2$  ERR by using S10 catalyst at -1.5 V vs. SHE.



Fig. S10. NMR spectrum recorded from the liquid products after  $CO_2$  ERR by using S10 catalyst at -1.6 V vs. SHE.



Fig. S11. NMR spectrum recorded from the liquid products after  $CO_2$  ERR by using S10 catalyst at -1.7 V vs. SHE.



Fig. S12. NMR spectrum recorded from the liquid products after  $CO_2$  ERR by using S10 catalyst at -1.8 V vs. SHE.



Fig. S13. SEM image of S10 after catalytic  $CO_2$  ERR for 50 hours at -1.3 V vs. SHE.

Sample	CHNs溶液 (ml)	GO (2mg/ml)
S1	5	1
S2	10	1
S3	15	1
S4	20	1
S5	30	1
S6	40	1
<b>S</b> 7	50	1
S8	70	1
S9	100	1
S10	125	1
S11	150	1

**Table S1.** The CHNs and GO volume ratio for different samples.

vs. SHE	C <sub>2</sub> H <sub>4</sub>	$C_2H_6$	H <sub>2</sub>	СО
-1.3 V	55.4%	37.6%	6.7%	0.3%
-1.4 V	68.2%	10.2%	20.5%	1.1%
-1.5 V	43.2%	2.9%	51.2%	2.7%
-1.6 V	17.8%	1.2%	77.5%	3.7%
-1.7 V	4.6%	0.7%	91%	3.7%
-1.8 V	1.8%	0	95.9%	2.3%

**Table S2.** The FE of the gas products from  $CO_2$  ERR at different potential by suing S10 as catalyst.

Electrolyzer	Catalyst	Electrolyte	Potential	Current density	Stability	C <sub>2</sub> H <sub>4</sub> (FE	C <sub>2</sub> total (FE	Ref.
			(V)	$(mA/cm^2)$	(h)	%)	%)	
Flow cell	OH <sup>-1</sup> modified 25 nm Cu	7 M KOH	-0.55 vs RHE	275	150	70	83%	2
	NPs/graphite/carbon black/PTFE							
H cell	Tens nm Cu-CuO <sub>x</sub> /carbon black	0.1 M KHCO <sub>3</sub>	-1.3 vs RHE	20	9	53	74	4
Flow cell	CuO-Cu <sub>2</sub> O/carbon black	1 M KHCO <sub>3</sub>	-1.6 vs RHE	160	12	46	-	7
H cell	250-300 nm Cu-CuOx-I/copper foil	0.1 M KHCO <sub>3</sub>	-0.9 vs RHE	31.2	22	47	80	9
H cell	Cu <sub>x</sub> @Cu <sub>2</sub> O nano covex/carbon paper	0.1 M KHCO <sub>3</sub>	-1.2 vs RHE	14.8	12	59.3	90.5	11
Flow cell	Cu-Cu <sup>+</sup> on CuSiO <sub>3</sub> /carbon black	0.1 M KHCO <sub>3</sub>	-1.1 vs RHE	20.2	6	51.8	70	12
Flow cell	C <sub>60</sub> -Cu-Cu <sup>+</sup> composite	1.0 M KOH	-1.4 vs RHE	366	12	~35	61	14
Flow cell	Cu/Cu <sub>2</sub> O NPs/Carbon nanotubes	2 M KOH	-	800	3	45	79	17
Flow cell	OH <sup>-</sup> modified Cu NPs/carbon black	0.1 M KHCO <sub>3</sub>	-3.4 cell	316	35	55.6	-	22
			voltage					
Flow cell	NH <sub>3</sub> Cl modified Cu/Cu <sub>2</sub> O/carbon black	0.1 M KHCO <sub>3</sub>	-	200	50	~10	~75	24
Flow cell	Poly ionic liquid/Cu-Cu <sub>2</sub> O hybrids	1 M KOH	-0.85 vs RHE	304	10	~44	76.1	35
Flow cell	Cu-Cu <sub>2</sub> O/rGO	1 M KHCO <sub>3</sub>	-1.3 vs SHE	16	50	55.4	93	This work
			-1.4 vs SHE	26	50	68.2	78.4	

**Table S3.** The Faraday efficiency of  $C_2H_4$  and total  $C_2$  through the  $CO_2$  ERR catalyzed by  $Cu^0$ - $Cu^+$  based catalyst reported recently and this work.