

**Supporting Information**

**Enhancing light harvesting and charge separation of Cu<sub>2</sub>O  
photocathodes with spatially separated noble-metal cocatalysts  
towards highly efficient water splitting**

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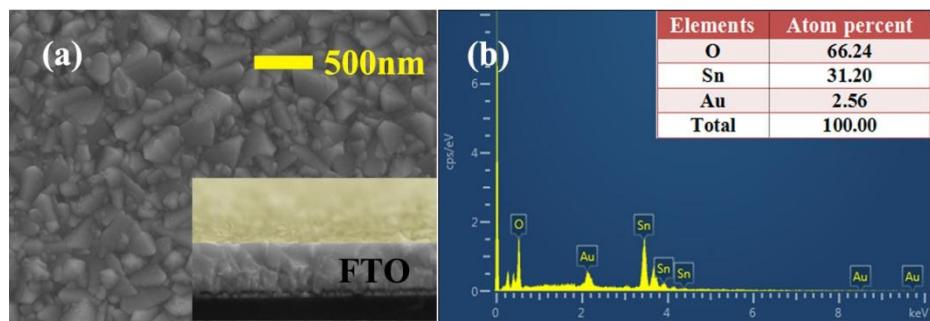


Fig. S1 (a) Top-view and cross-section SEM image and (b) the EDS elemental analysis spectrum of Au

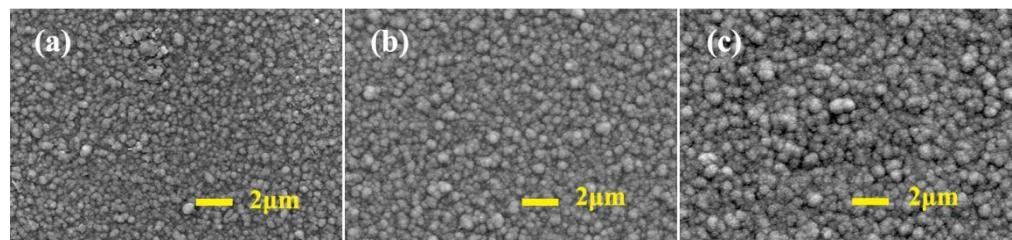


Fig. S2 SEM images of Cu<sub>2</sub>O prepared by varied electrodeposition time: (a) 5min, (b) 15 min, (c) 30 min

Fig. S3 Photocurrent density-voltage curves of bare Cu<sub>2</sub>O with different (a) electrodeposition time and (b) concentration of precursor solution for CuSO<sub>4</sub>

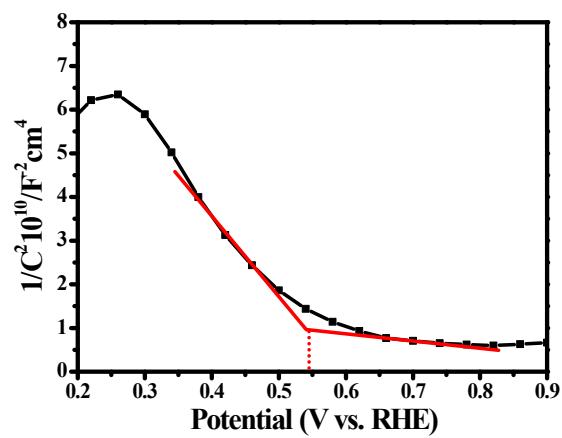


Fig. S4 Mott-Schottky plot of Cu<sub>2</sub>O measured in 0.1 M Na<sub>2</sub>SO<sub>4</sub> solution at 1 kHz

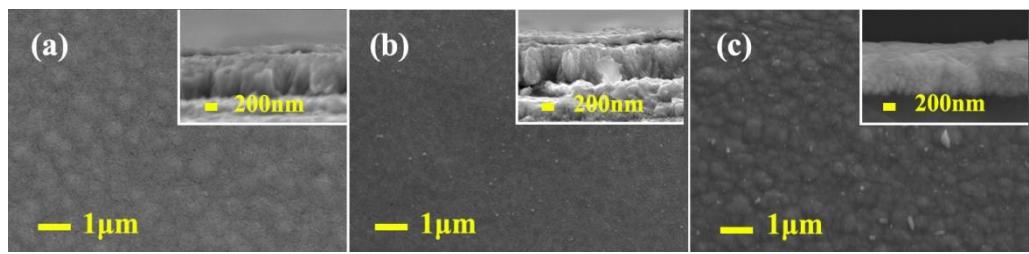


Fig. S5 Top-view and cross-section SEM images of  $\text{Cu}_2\text{O}/\text{Pt}$  photocathodes prepared by different deposition time: (a) 90 s, (b) 120 s, (c) 180 s

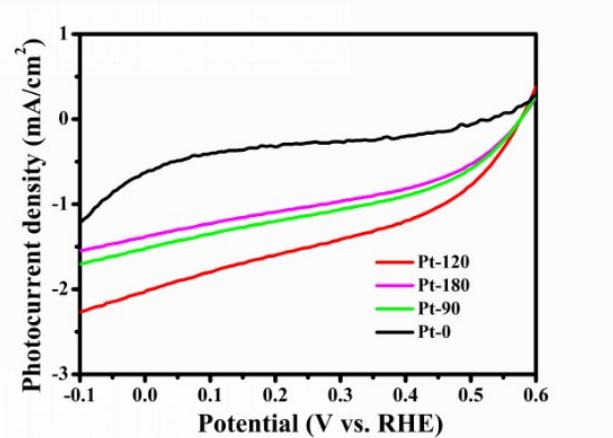


Fig. S6 Photocurrent density-voltage curves of  $\text{Cu}_2\text{O}/\text{Pt}-x$  photocathodes in 5 mM  $\text{H}_2\text{PtCl}_6$  aqueous solutions for various times. Pt-0: reference  $\text{Cu}_2\text{O}$  NGs, Pt-90: 90 s; Pt-120: 120 s; Pt-180: 180 s

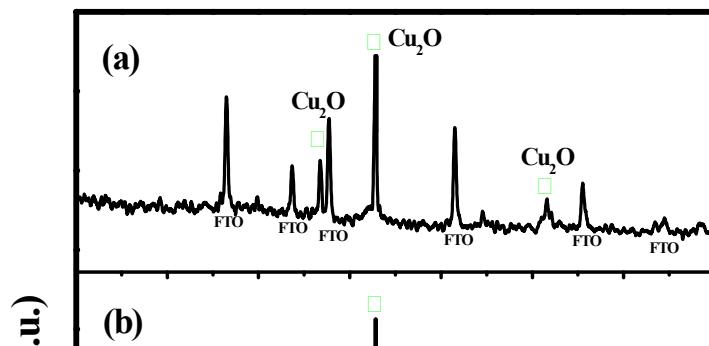


Fig. S7 XRD (a) before and (b) after PEC stability measurement for the pristine Cu<sub>2</sub>O and (c) after PEC stability measurement for Au/Cu<sub>2</sub>O/Pt composite

Fig. S8 Cyclic voltammetry of (a) pristine Cu<sub>2</sub>O and (b) Au/Cu<sub>2</sub>O/Pt composite

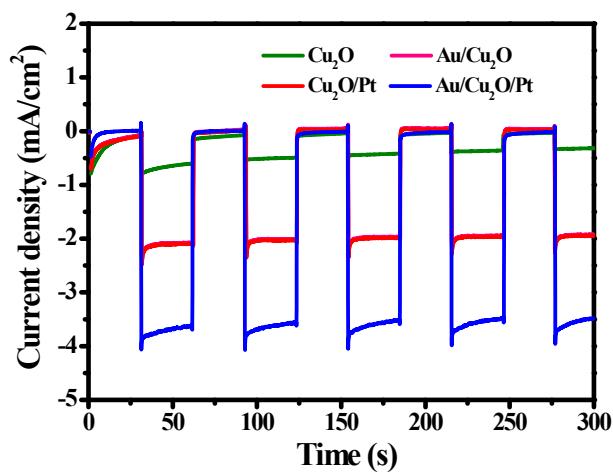


Fig. S9 Photocurrent density-time characteristics in 0.1 M Na<sub>2</sub>SO<sub>4</sub> solution for pristine Cu<sub>2</sub>O, Cu<sub>2</sub>O/Pt, Au/Cu<sub>2</sub>O and Au/Cu<sub>2</sub>O/Pt electrodes at 0 V vs. RHE

Table S1 EIS fitting results of  $R_{ct}$  for Cu<sub>2</sub>O NGs, Au/Cu<sub>2</sub>O, Cu<sub>2</sub>O/Pt and Au/Cu<sub>2</sub>O/Pt photocathodes as shown in Fig. 9

Sample	Cu <sub>2</sub> O	Au/Cu <sub>2</sub> O	Cu <sub>2</sub> O/Pt	Au/Cu <sub>2</sub> O/Pt
$R_{ct}$ ( $\Omega$ )	1546	1318	453	250