## **Electronic Supplementary Information**

## Modulating Interleaved ZnO Assembly with CuO Nanoleaves for Multifunctional Performance: Perdurable CO<sub>2</sub> Gas Sensor and Visible Light Catalyst

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**Figure S1.a** Particle size distribution calculated from FE-SEM image (N-150 particles) given in Fig. 1h illustrating hierarchical ZnO spheres synthesized at 500°C for 2 h.



Figure S1.b X-ray diffractograms of Zinc hydroxide carbonate (ZHC) precursor synthesized at 120°C for 6 h, ZnO calcined at 300°C (Z-3), 400°C (Z-4) and 500°C (Z-5) for 2 h.



Figure S2 Nitrogen adsorption-desorption isotherms ZnO microspheres (Z-4). Inset pore size

distribution curve.



Figure S3 Representative stacked XPS spectrum of (a) pure ZnO, (b) pure CuO, (c) CZ-1:1 and

(d) 1 wt%Ag-CZ-1:1.



Figure S4 XPS core level spectra of 1 wt.%Ag-CZ-1:1,where (a) Ag 3d and (b) O 1s.



**Figure S5** FE-SEM and TEM micrographs of **(a-b)** pure ZnO obtained at 400°C for 2 h, **(c-d)** pure CuO obtained at 50°C for 12 h and **(e-f)** CuO/ZnO composite in 1:8 mole ratio (CZ-1:8) obtained at 50°C for 12 h. The inset in each case represents the corresponding SAED pattern. Pink arrows and yellow circles represent CuO nanoleaves and ZnO spheres respectively.



**Figure S6 (a)** Scanning electron micrograph reveals the actual area of CuO/ZnO composite used for elemental quantification, **(b)** layered EDS micrograph, elemental maps demonstrate uniform presence of **(c)** zinc (Zn), **(d)** copper (Cu), **(e)** oxygen (O), **(f)** silver (Ag) and **(g)** energy dispersive X-ray spectroscopy analysis of CuO/ZnO (CZ-1:1) decorated with 5 wt.%Ag.



Figure S7 (a) Sensor response characteristics of CZ-1:8 composite as a function of operating temperature towards different  $CO_2$  gas concentrations and (b) response/recovery curves for composites towards 1000 ppm  $CO_2$  gas at 320°C.



**Figure S8.a** X-ray diffractograms of 1 wt.%Ag-CZ-1:8 composite where **(a)** before and **(b)** after 40 days of continuous sensing performance towards 1000 ppm CO<sub>2</sub> gas balanced in dry air at 320°C.



Figure S8.b XPS core level spectrum of Ag after 40 days of continuous sensing performance

towards 1000 ppm CO<sub>2</sub> gas balanced in dry air at 320°C.



**Figure S9** UV-Vis absorbance spectra depicting the change in concentration of MB under visible irradiation as a function of time for various photocatalyst.



**Figure S10** UV-Vis absorbance spectra depicting the change in concentration of MB under visible irradiation as a function of time for various photocatalyst.



Figure S11.a Chemical stability testing of the CZ-1:8 interleaved assembly under visible light irradiation for four consecutive cycles.



Figure S11.b Rate of photocatalytic degradation of MB under visible light irradiation in presence of CZ-1:8 for four consecutive cycles.



Figure S12 X-ray diffractograms where (a) as-synthesized CZ-1:8 composite, (b) after 40 days of  $CO_2$  gas sensing performance and (c) after 4 consecutive cycles of visible light induced MB degradation.



Figure S13 TEM micrograph of CZ-1:8 composite after 40 days of continuous CO<sub>2</sub> gas sensing test.



Figure S14 TEM micrograph of CZ-1:8 composite after 4 consecutive photocatalytic cycles of MB degradation studies.