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Figure S1.TEM images of CuCoMnO_x-200 (a), CuCoMnO_x-300 (b), CuCoMnO_x-400 (c)and CuCoMnO_x-500 (d).



Figure S2. CO conversion (a,c) and temperature (b,d) curves of $CoMnO_x$ -T and $CuCoMnO_x$ -T in CO-PROX at 250 mW/cm².



Figure S3. CO conversion and O_2 selectivity of CuCoMnO_x-300 for photothermal CO-PROX.



Figure S4. CO conversion of thermal catalysis over $CoMnO_x$ -T (a) and $CuCoMnO_x$ -T (b).



Figure S5. comparation of CO conversion of the $CoMnO_x$ and $CuCoMnO_x$ catalysts for thermal and photothermal CO-PROX.



Figure S6. CO conversion and temperature curve of $CuCoMnO_x$ -300 in CO-PROX at 250mW/cm² illuminated under different optical composition.



Figure S7. CO conversions of pure Co_3O_4 and Mn_2O_3 , binery $CoMnO_x$ and ternary $CuCoMnO_x$ calcined at 300 °C (a) and CO conversions of $CuCoMnO_x$ -300 with different copper amount in photothermal CO-PROX at 250mW/cm².



Figure S8. CO conversion of $CuCoMnO_x$ -300 in CO-PROX at different illumination power densities (a) and at different weight hourly space velocity (b) at 250 mW/cm² (b).



Figure S9. PAE values of $CuCoMnO_x$ -300 catalyst under different illumination densities.



Figure S10. UV-Vis-IR absorption spectrums of (a) CoMnO_x-T, (b) CuCoMnO_x-T.



Figure S11. XPS spectra of C 1s (A) and Mn 3s (B) of $CoMnO_x$ -300 (a), CuCoMnO_x-200 (b), CuCoMnO_x-300 (c) , CuCoMnO_x-400 (d) and CuCoMnO_x-500 (e).



Figure S12. DRIFTS spectra (1900-1200 cm⁻¹) of $CoMnO_x$ -300 (a) and CuCoMnO_x-300 (b) under the reaction stream at different temperature.



Figure S13. DRIFTS spectra (3800-2200 cm⁻¹) of CoMnO_x-300 (a) and CuCoMnO_x-300 (b).