Decoration of PdAg dual-metallic alloy nanoparticles on Z-scheme α-Fe₂O₃/CdS for manipulable products via photocatalytic reduction of carbon dioxide

Shuhui Yang¹, Xi Ke¹, Kang Wang¹, Menglong Zhang^{1*} and Dongxiang Luo^{23*}

¹Institute of Semiconductors, South China Normal University, Guangzhou, 510006, P.R. China ²School of Chemistry and Chemical Engineering/Institute of Clean Energy and Materials/Guangzhou Key Laboratory for Clean Energy and Materials/Huangpu Hydrogen Innovation Center, Guangzhou University, Guangzhou 510006, PR China

³School of Materials and Energy, Guangdong University of Technology, 510006, P.R. China



Figure S1. SEM image of α -Fe₂O₃ nanorod.



Figure S2. a) PXRD patterns of α -Fe₂O₃/CdS/PdAg 1:2; b) TEM image of pristine PdAg NPs and illustration of lattice fringe, statistical distribution of PdAg NPs size; c) TEM image of ternary composites α -Fe₂O₃/CdS/PdAg scraped from a slide; d) EDS spectrum of pristine PdAg NPs, in which red for Pd peak and green for Ag peak. The PXRD patterns and lattice fringes are corresponding to JCPDS 46-1088 (FTO), 33-0664 (α -Fe₂O₃), and 41-1049 (CdS).



Figure S3. UV-vis spectrum of series photocatalysts.



Figure S4. XPS spectrum of Pd 3d and Ag 3d in PdAg and α -Fe₂O₃/CdS/PdAg.



Figure S5. TRPL spectra of α -Fe₂O₃/CdS and α -Fe₂O₃/CdS/PdAg (2:1).



Figure S6. a) Linear sweep voltammetry and b) chronoamperometry at 0.3 V vs RHE of α -Fe₂O₃, CdS and α -Fe₂O₃/CdS.



Figure S7. Relative photocurrent change at -0.3 V vs RHE of α-Fe₂O₃/CdS/PdAg 1:2 in 10 min.



Figure S8. PXRD patterns of α -Fe₂O₃/CdS/PdAg 1:2 before and after electrochemical reaction.