

Supporting Information

MnCO₃-Au Nanoparticles to Enable Catalytic Tumor Inhibition with Immune Activation

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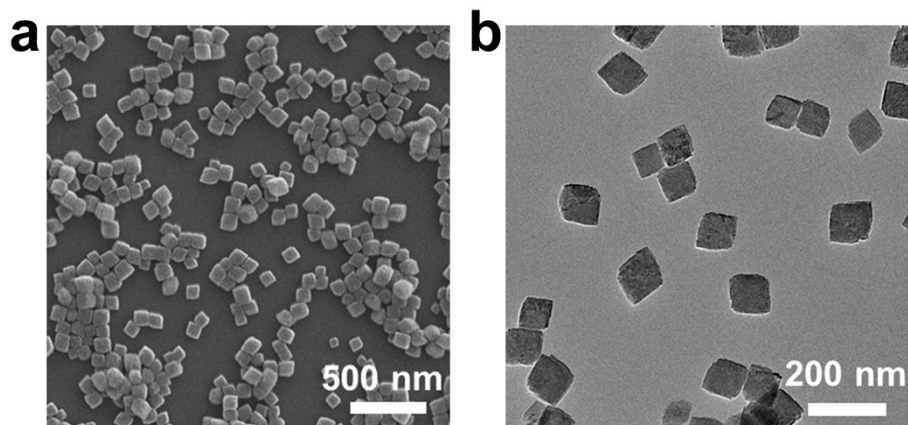


Figure S1. (a) SEM image and (b) TEM image of MnCO_3 nanoparticles.

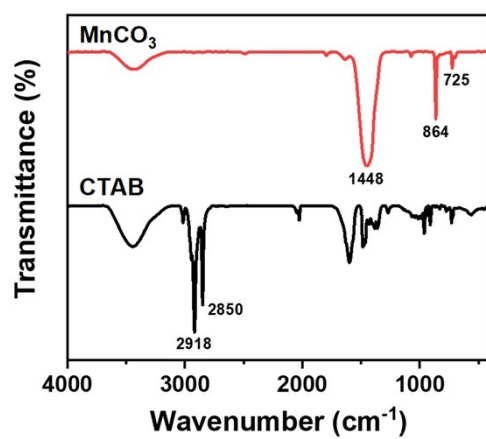


Figure S2. FTIR spectra of CTAB and as-prepared MnCO_3 nanoparticles.

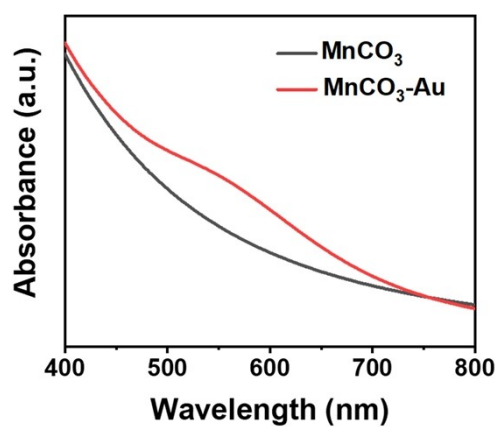


Figure S3. UV-Vis spectra of MnCO_3 and $\text{MnCO}_3\text{-Au}$.

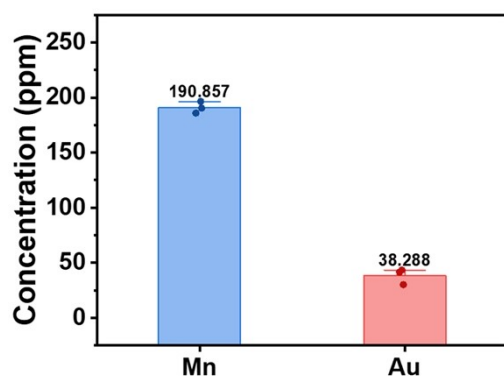


Figure S4. The concentrations of Mn and Au in the digestion solution of MnCO₃-Au nanoparticles determined by ICP-OES.

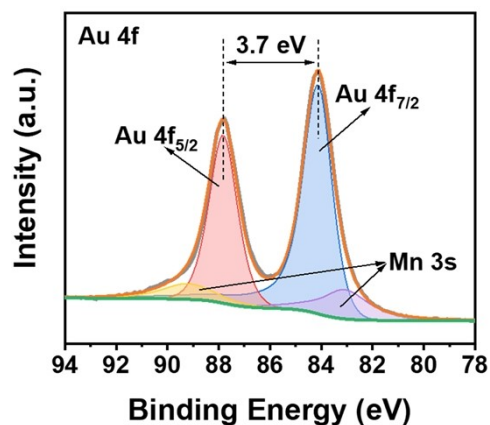


Figure S5. High-resolution Au 4f XPS spectrum of MnCO₃-Au.

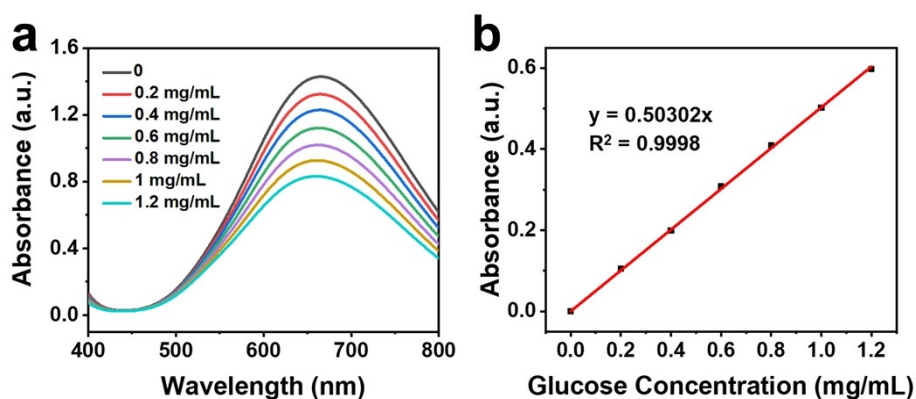


Figure S6. (a) UV-Vis absorption spectra of Fehling's reagent reacted with different concentrations of glucose. (b) The standard curve of absorbance difference at 663 nm versus glucose concentration.

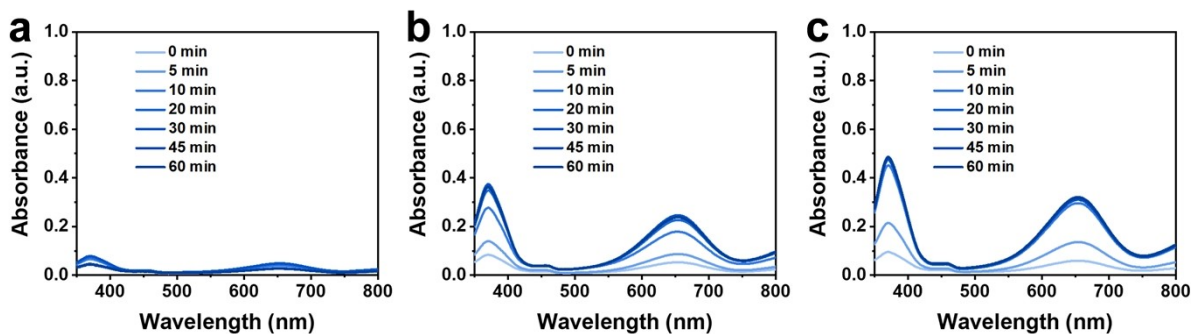


Figure S7. UV-Vis absorption spectra of TMB oxidized by generated H_2O_2 in (a) 0 mg/mL, (b) 0.5 mg/mL, and (c) 1 mg/mL glucose solutions at different time points with the addition of 500 $\mu\text{g/mL}$ $\text{MnCO}_3\text{-Au}$.

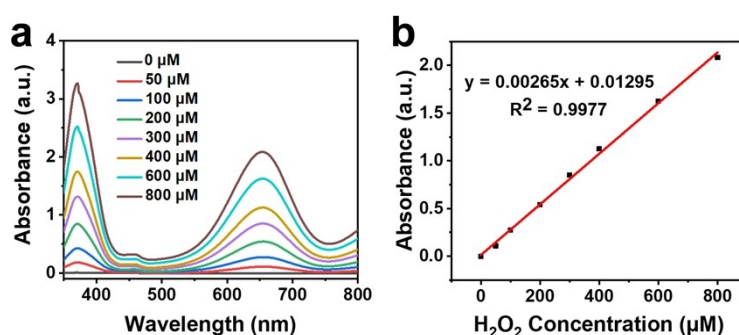


Figure S8. (a) UV-Vis absorption spectra of TMB oxidized by various concentrations of H_2O_2 . (b) The standard curve of absorbance at 652 nm versus H_2O_2 concentration.

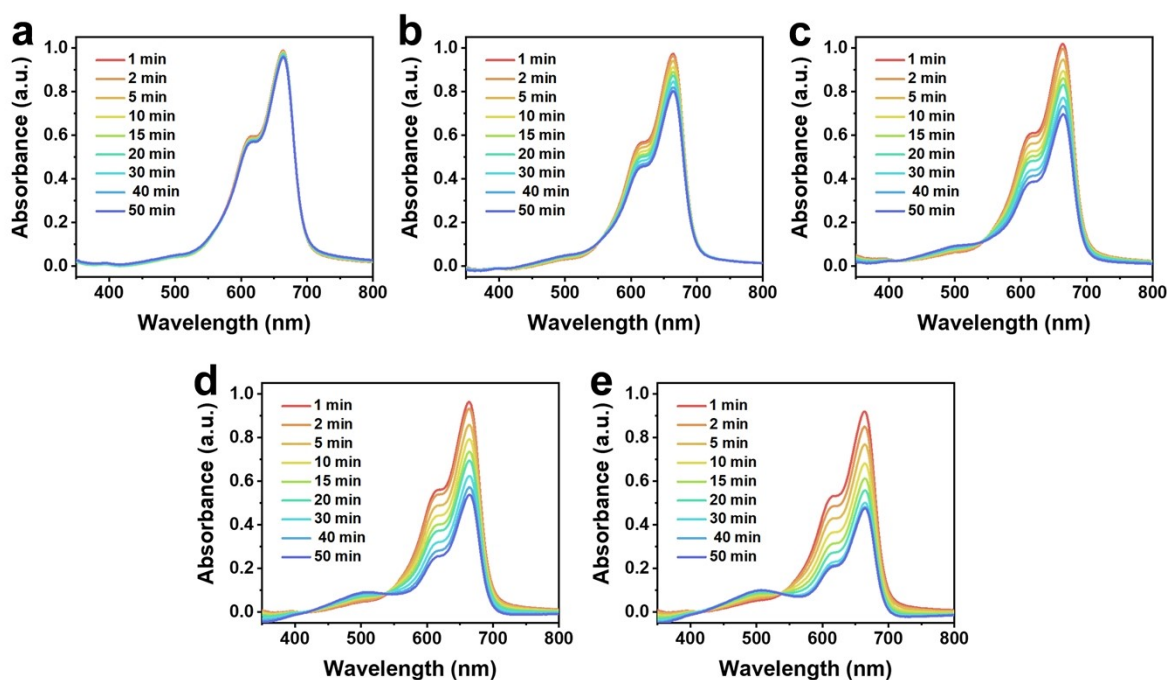


Figure S9. UV-Vis absorption spectra of MB solutions containing 500 $\mu\text{g/mL}$ $\text{MnCO}_3\text{-Au}$ and (a) 0 mg/mL, (b) 0.1 mg/mL, (c) 0.3 mg/mL, (d) 0.5 mg/mL, (e) 1 mg/mL glucose.

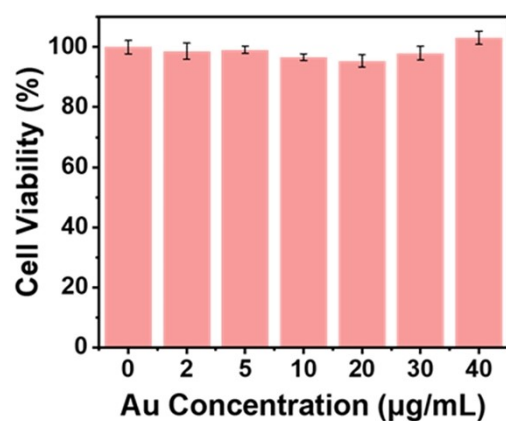


Figure S10. 4T1 cell viabilities after incubation with different concentrations of AuNPs for 24 h.

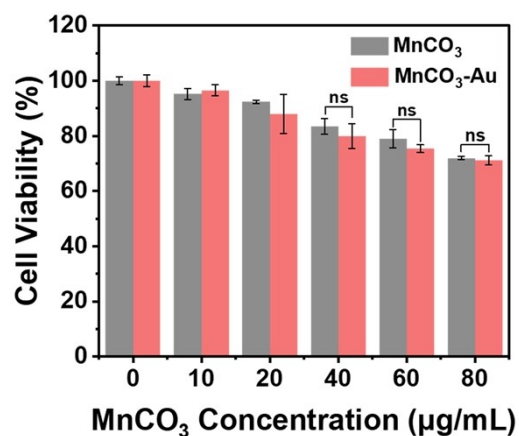


Figure S11. 4T1 cell viabilities after incubation with MnCO₃ and MnCO₃-Au in glucose-free RPMI 1640 medium.

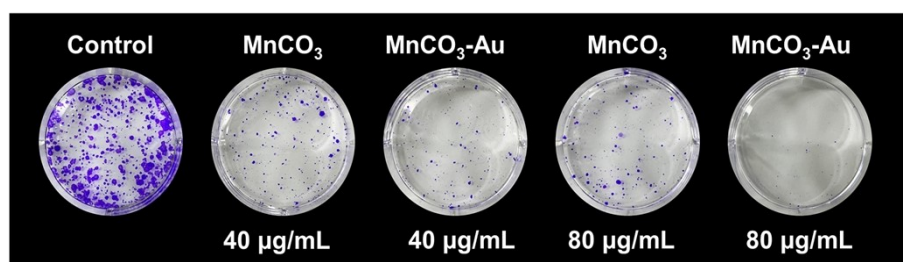


Figure S12. Colony formation of 4T1 cells after different treatments.

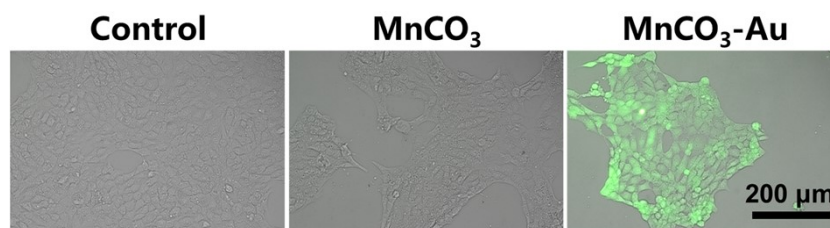


Figure S13. The bright-field images merged with fluorescence images of DCFH-DA stained 4T1 cells after being incubated with PBS, MnCO_3 , and $\text{MnCO}_3\text{-Au}$.

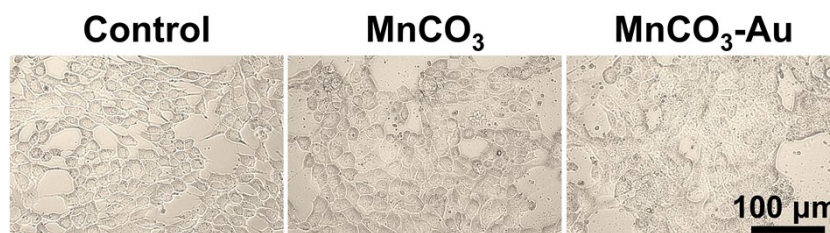


Figure S14. The bright-field images of JC-1 stained 4T1 cells after being treated with PBS, MnCO_3 , and $\text{MnCO}_3\text{-Au}$.

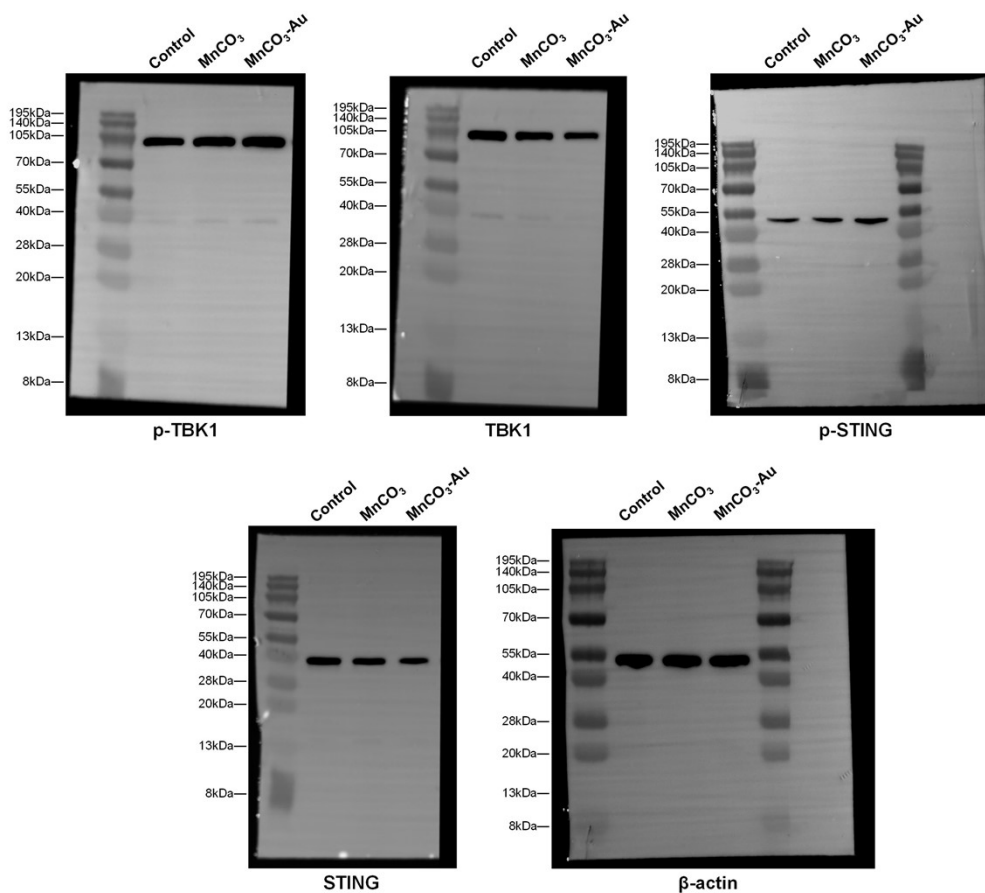


Figure S15. Raw data associated with Fig. 4d. Lanes were not cropped or rearranged when presenting the results in Fig. 4d.

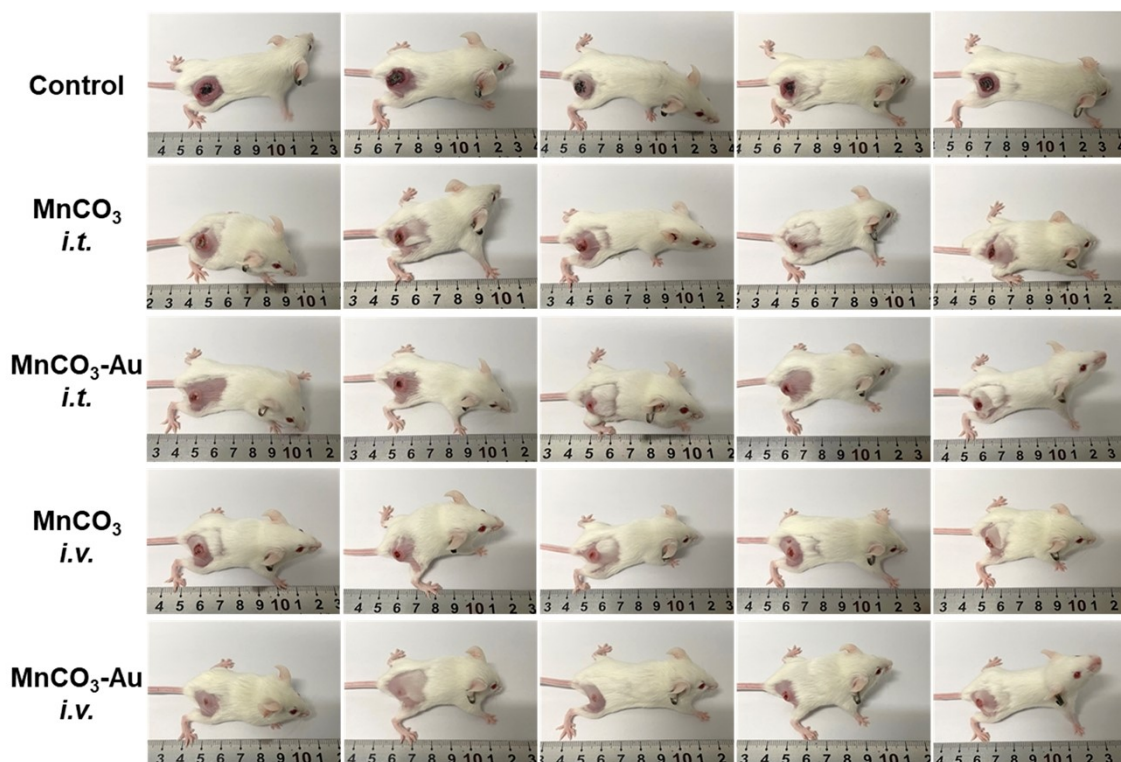


Figure S16. Photographs of tumor-bearing mice on day 14 after different treatments.

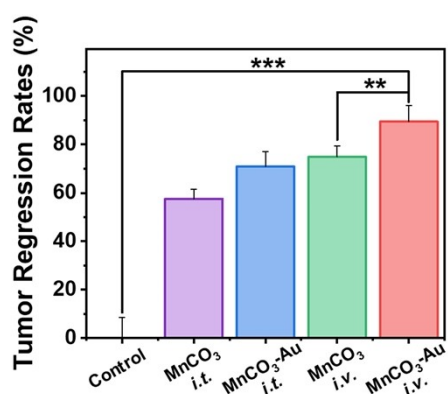


Figure S17. Tumor regression rates on day 14 after different treatments.

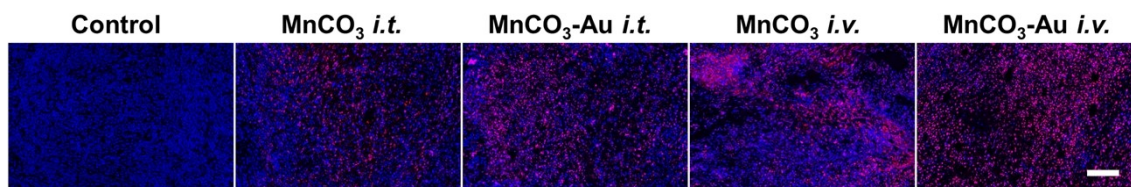


Figure S18. TUNEL immunofluorescence staining images of the tumors collected on day 14 after different treatments (scale bar: 100 μ m).

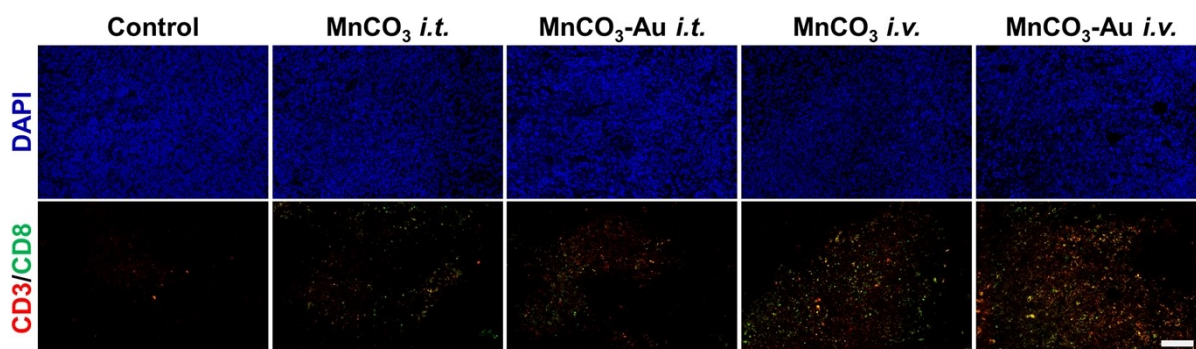


Figure S19. Immunofluorescence staining images of CD8⁺ T cells in the tumors collected on day 14 after different treatments. CD3 is stained in red fluorescence and CD8 is stained in green fluorescence (scale bar: 100 μ m).

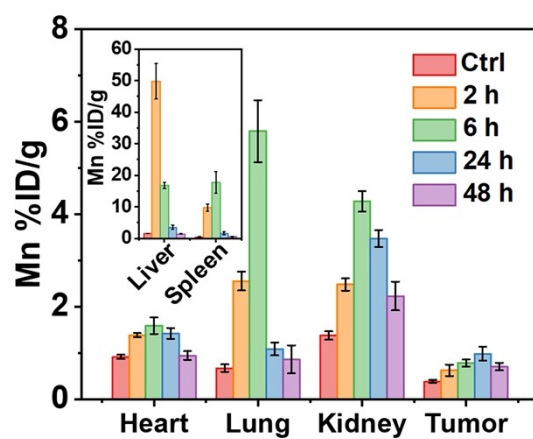


Figure S20. Biodistribution of Mn at various time points after intravenous injection of MnCO₃-Au ($n = 3$, mean \pm SD).

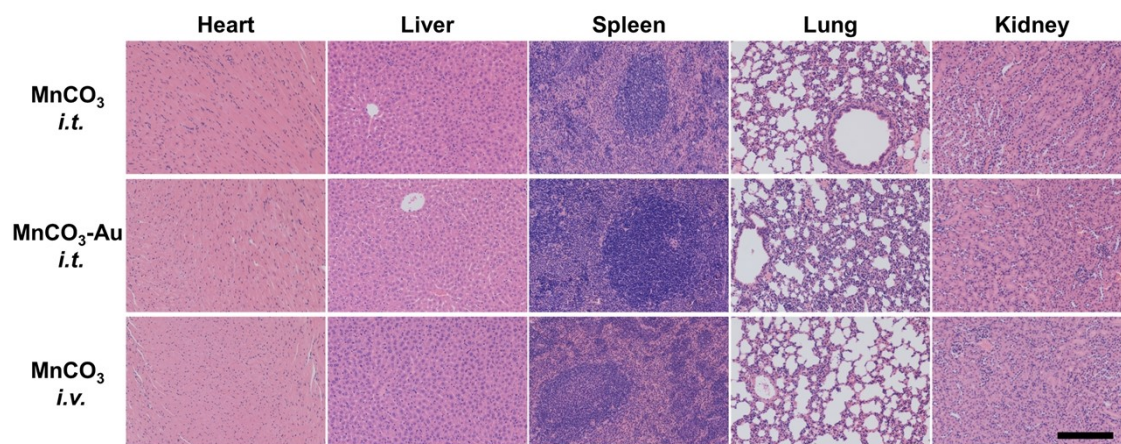


Figure S21. H&E staining images of major organs collected from mice after different treatments (scale bar: 200 μ m).