

Supporting Information

Enzyme-triggered on-demand release of H₂O₂-self-supplying

CuO₂@Fe₃O₄ Nanoagent for enhanced chemodynamic

antimicrobial therapy and wound healing

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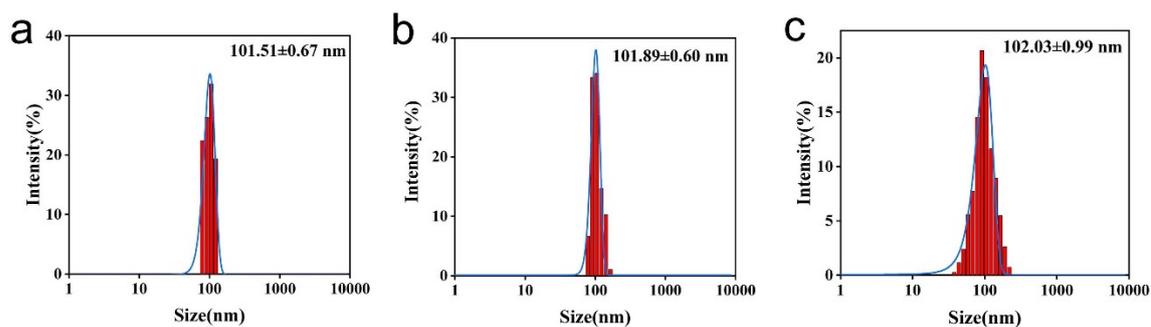
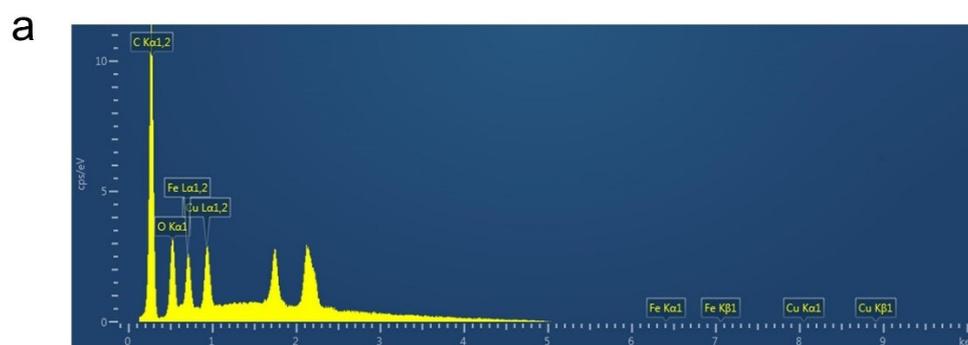


Figure S1. DLS size distribution of Fe_3O_4 (a), $\text{CP@Fe}_3\text{O}_4$ (b) and $\text{HA-CP@Fe}_3\text{O}_4$ (c).



b

Element Line	Weight %	Weight % Sigma
C	45.0	0.5
Fe	29.7	0.6
Cu	17.0	0.4
O	8.2	0.2

Figure S2. Applying EDS for elemental analysis. A) Qualitative analysis: The peaks in the EDS spectrum represent C, Fe, Cu and O elements in $\text{HA-CP@Fe}_3\text{O}_4$. B) Quantitative analysis: Quantitative analysis is to obtain the content of C, Fe, Cu and O in $\text{HA-CP@Fe}_3\text{O}_4$ by X-ray intensity.

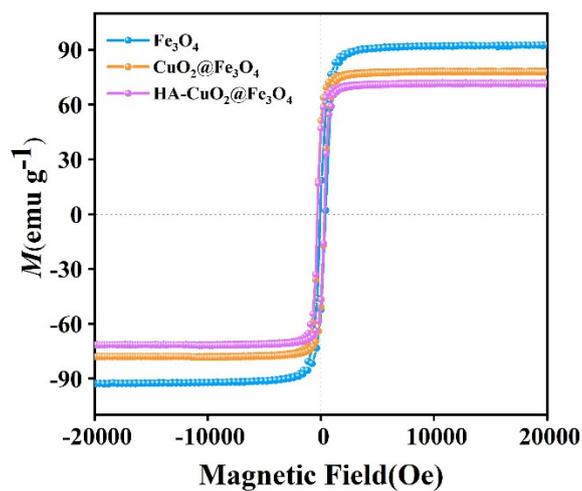


Figure S3. Room-temperature magnetic hysteresis loops of Fe_3O_4 , $\text{CP@Fe}_3\text{O}_4$ and $\text{HA-CP@Fe}_3\text{O}_4$.

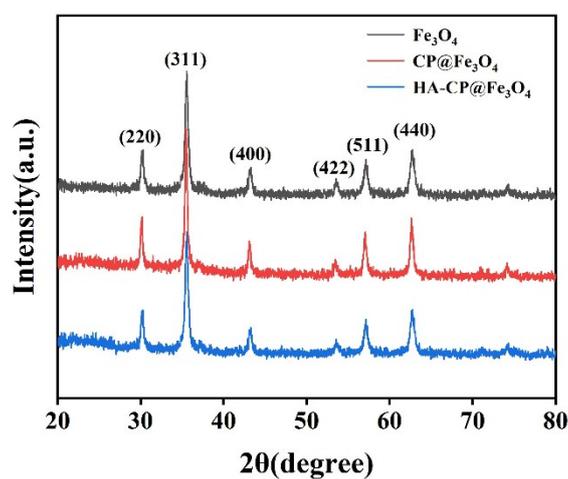


Figure S4. XRD patterns of Fe_3O_4 (black line) $\text{CP@Fe}_3\text{O}_4$ (red line) and $\text{HA-CP@Fe}_3\text{O}_4$ (blue line).

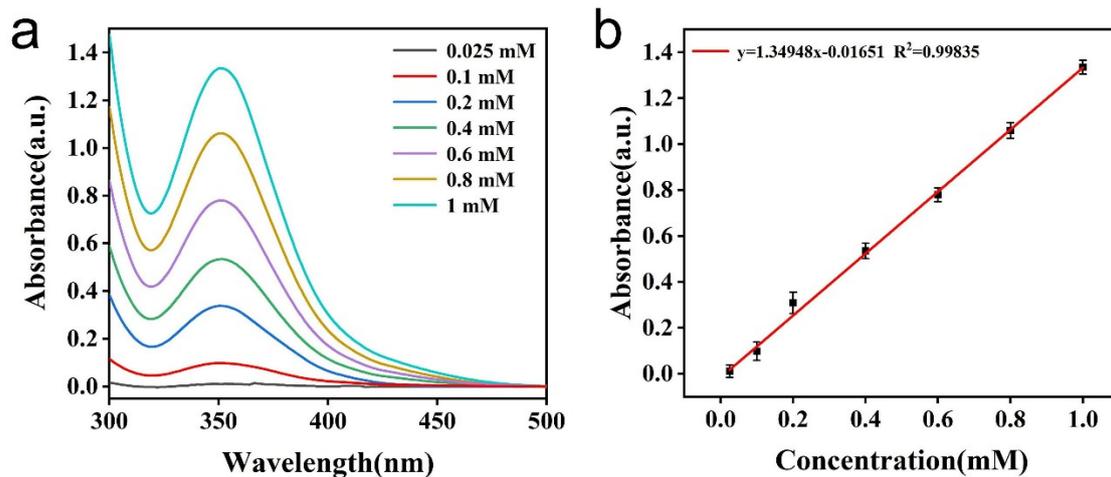


Figure S5. (a) The concentration of H₂O₂ was quantified by the absorbance of the product from the reaction of KI with H₂O₂. (b) Taking a fixed KI concentration as the reference, the absorbance was measured under different concentrations of H₂O₂ to obtain a calibration curve.

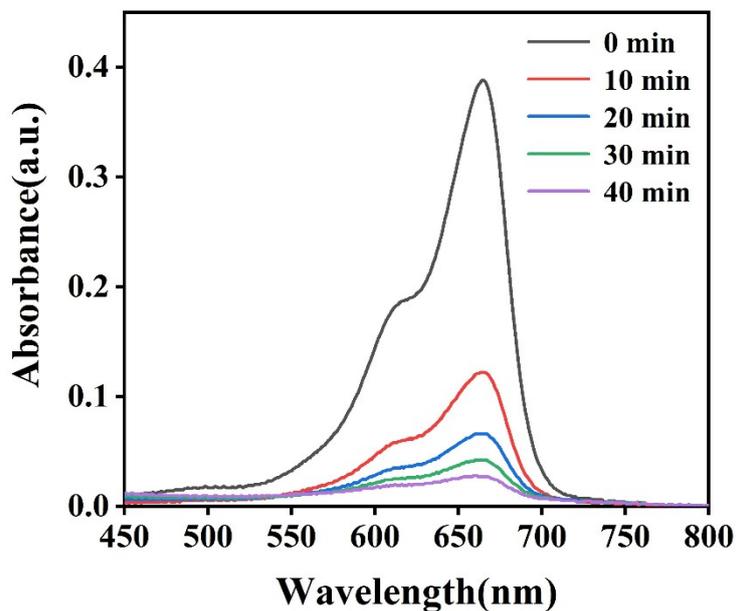


Figure S6. Absorption spectra of MB solution with HA-CP@Fe₃O₄ at increasing time intervals.

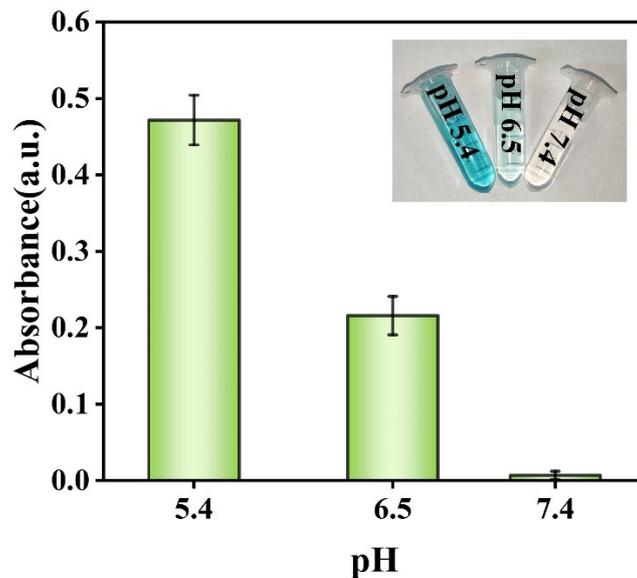


Figure S7. Fenton activity of HA-CP@Fe₃O₄ (200 µg/mL) at different pH after 60 min reaction.

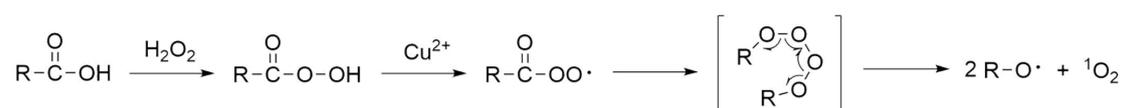


Figure S8. Russell reaction mechanism.

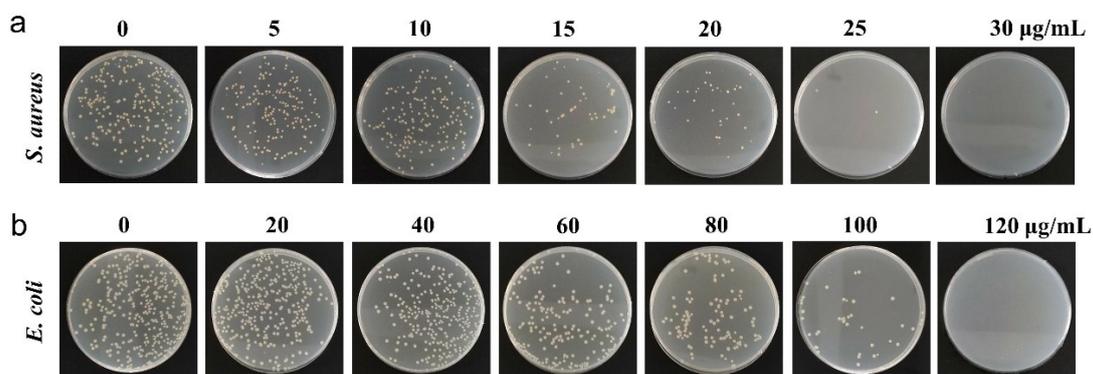


Figure S9. (a) Images of the LB agar plates of *S. aureus* treated with different concentrations of HA-CP@Fe₃O₄. (b) Images of the LB agar plates of *E. coli* treated with different concentrations of CP@Fe₃O₄.

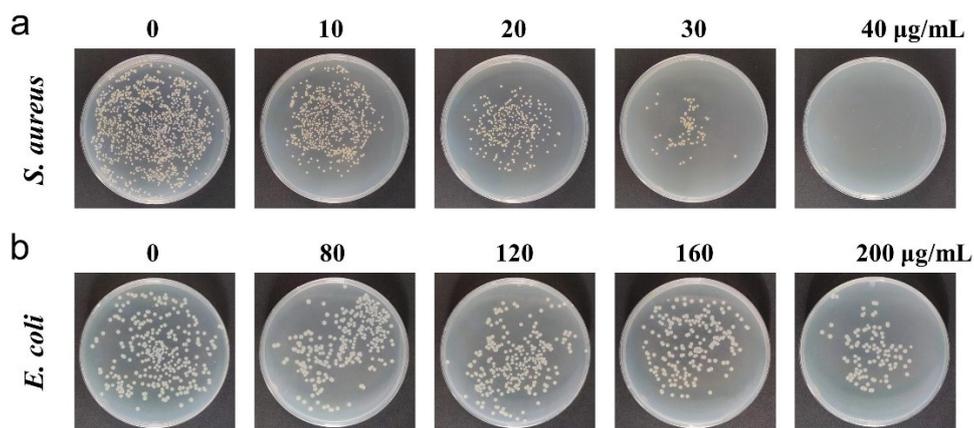


Figure S10. (a) Images of the LB agar plates of *S. aureus* treated with different concentrations of CP@Fe₃O₄. (b) Images of the LB agar plates of *E. coli* treated with different concentrations of HA-CP@Fe₃O₄.

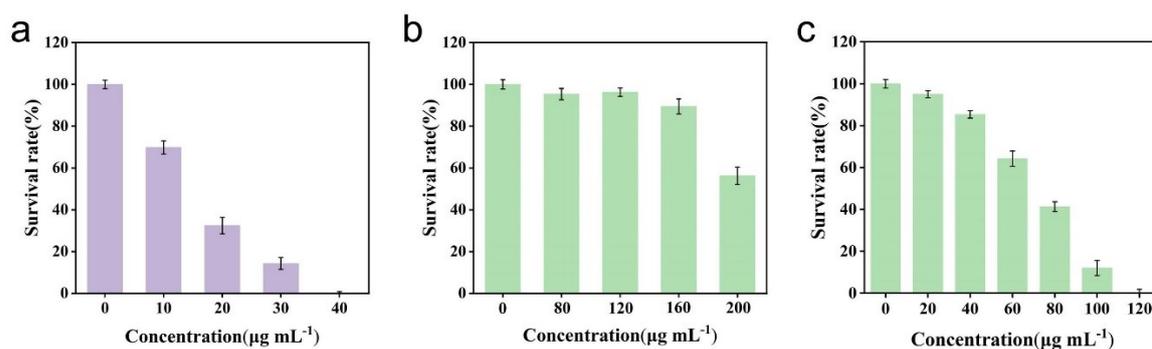


Figure S11. (a) Statistical analysis of *S. aureus* treated with different concentrations of CP@Fe₃O₄. (b) Statistical analysis of *E. coli* treated with different concentrations of HA-CP@Fe₃O₄. (c) Statistical analysis of *E. coli* treated with different concentrations of CP@Fe₃O₄.

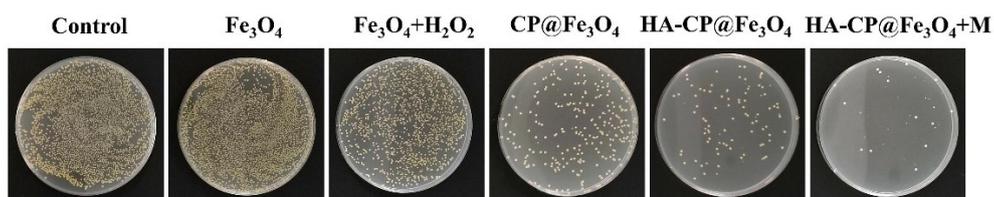


Figure S12. Bacteria isolated from wound tissue after different treatments were cultured on agar plates.

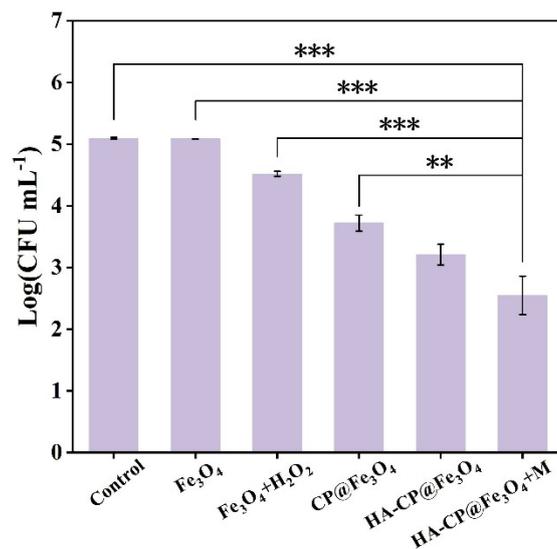


Figure S13. Bacterial viability from the infected wound tissues after treatment with different groups.

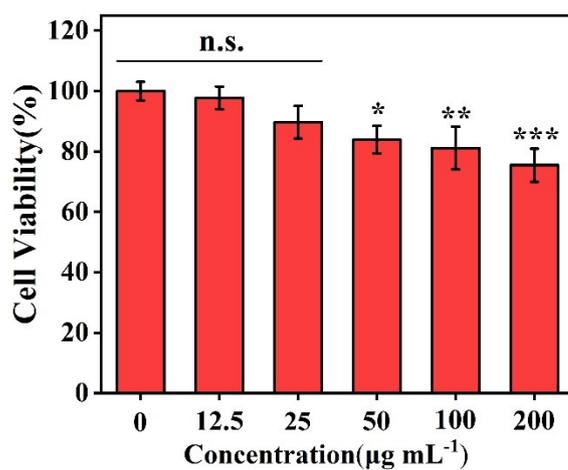


Figure S14. The survival rate of L929 cells after treatment with different concentrations of CP@Fe₃O₄ for 24 h.