Carbon dot decorated Co₃O₄ nanozymes responsive to NIR-II window

for mild photothermal-enhanced nanocatalytic therapy

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Figure S1. Raman spectrum of CD nanozymes.



Figure S2. The survey XPS (a), high-resolution C 1s (b), N 1s (c), and O 1s (d) spectra of CDs.



Figure S3. The hydrodynamic diameter of Co_3O_4 (a), CDs (b), and CD@ Co_3O_4 (c).



Figure S4. Element mapping images of CD@Co₃O₄.



Fig. S5. Photographs of $CD@Co_3O_4$ PBS solution, saline solution, and FBS solution stored for different periods of time (0, 1d, and 7 d).



Figure S6. The survey XPS (a), high-resolution Co 2p (b), and O 1s (c) spectra of Co₃O₄.



Figure S7. The NIR-II thermographic images of CD, Co₃O₄, and CD@Co₃O₄.



Figure S8. (a) High-resolution Co 2p XPS spectrum of CD@Co₃O₄. (b, c) High-resolution Co 2p XPS

spectrum of Co_3O_4 (b) and $CD@Co_3O_4$ (c) incubated with H_2O_2 .



Figure S9. Absorption spectra of the oxidized TMB catalyzed by Co₃O₄ at pH 6.5.



Figure S10. Absorption spectra of the oxidized TMB catalyzed by $CD@Co_3O_4$ (a) or Co_3O_4 (b) at pH 7.4.



Figure S11. O_2 generation upon the addition of CD@Co₃O₄ and H₂O₂ (0.40 mM) at varied pH (4.5, 6.5, or 7.4).



Figure S12. The GSH depletion activity evaluation of Co_3O_4 (a) and $CD@Co_3O_4$ (b).



Figure S13. (a) Confocal images of Hela cells treated with ICG labeled $CD@Co_3O_4$. (b) Cellular uptake of ICG labeled $CD@Co_3O_4$ determined by flow cytometry.



Figure S14. (a) Cell viability of and Hela cells treated with Co_3O_4 at varied concentrations (0-300 µg/mL). (b) Cell viability of and Hela cells treated with Co_3O_4 at varied concentrations (0-300 µg/mL) with the addition of H_2O_2 (50 µM). (c) Cell viability of and Hela cells treated with Co_3O_4 at varied concentrations (0-300 µg/mL) with the addition of H_2O_2 (50 µM) with the addition of H_2O_2 (50 µM) under the mild NIR-II laser treatments.



Figure S15. (a) Cell viability of and Hela cells treated with CDs at varied concentrations (0-300

 μ g/mL). (b) Cell viability of and Hela cells treated with CDs at varied concentrations (0-300 μ g/mL) with the addition of H₂O₂ (50 μ M). (c) Cell viability of and Hela cells treated with CDs at varied concentrations (0-300 μ g/mL) with the addition of H₂O₂ (50 μ M) under the mild NIR-II laser treatments.



Fig. S16. NIR fluorescence spectrum of ICG and ICG@CD@Co₃O₄.



Figure S17. Time-dependent NIR fluorescence intensity of ICG-labeled $CD@Co_3O_4$ in the major organs and tumor tissues based on the fluorescence imaging results.



Figure S18. The body weight curves of the treated mice after different treatments (n=5).



Figure S19. H&E-stained images obtained from the major organs (heart, liver, spleen, lung, and kidney) of mice in different treatment groups.



Figure S20. (a-b) Biochemical blood analysis (a) and hematological index (b) of the mice that were sacrificed at 18 days after different treatments. The terms of biochemical blood analysis include ALB, ALT, AST, TP, UREA, TBIL, CREA, and GLOB. The terms of hematological index include PLT, MCV, MCHC, MCH, HCT, Hb, WBC, and RBC.