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Supporting Information

Chemodynamic/photothermal synergistic therapy based on Ce-

doped Cu-Al layered double hydroxide

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Scheme S1. Schematic illustration of the bottom-up method of synthesizing CuAlCe-LDH.

Table S1.	The feed	ratio and	the actual	ratio of LDI	H determined b	ov ICP.
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Sample	Feed ratio	Actual ratio	References
Cu Al-LDH	2:1	2.47 : 1	this work
Cu Al Ce-LDH	2:0.5:0.5	2.32 : 0.57 : 0.43	this work
Cu Al Ce-LDH	2:0.67:0.33	2.21:0.65:0.34	this work
Cu Al Ce-LDH	2:0.75:0.25	2.37 : 0.75 : 0.28	this work
Mg Al Ce-LDH	3:0.8:0.2	0.73 : 0.24 : 0.014	1, 2
Ni Fe Ce-LDH		doping 5% Ce	3

Table S2. The K_M and V_{max} values of different Fenton catalysts.

Sample	<i>K_M</i> (mM)	<i>V_{max}</i> (M⋅s ⁻¹)	References
ICG/CuAlCe-LDH	1.57	4.88×10 ⁻⁶	this work
FeAl-LDH	0.16	1.47×10 ⁻⁶	4
PEG/Fe-LDHs	0.09	1.76×10 ⁻⁶	4
Fe ₃ O ₄ NPs	26.08	6.17×10 ⁻⁸	5
Mn-NS	26.40	7.04×10 ⁻⁸	6
Fe ₃ O ₄ @PPy@GOD NCs	4.94	1.13×10 ⁻⁸	7



Fig. S1 XRD pattern of CuAl-LDH (2:1) nanosheets after restocking.



Fig. S2 H_2O_2 reacts with CuAl-LDH and CuAlCe-LDH to oxidate TMB at pH=6.5, and the absorbance (650 nm) of reactants is determined *via* UV spectrum.



Fig. S3 TEM image of CAC-LDH nanosheets with corresponding EDX mapping images for Cu, Al, Ce and O, respectively.



Fig. S4 The hydrodynamic size of CAC-LDH.



Fig. S5 Zeta potential of ICG and CAC-LDH.



Fig. S6 The UV-vis-NIR of ICG aqueous solution before and after adsorption with CAC-LDH.



Fig. S7 FTIR spectra of CAC-LDH, ICG and ICG/CAC-LDH, respectively.



Fig. S8 (A) TEM and (B) AFM of ICG/CAC-LDH.



Fig. S9 Size distribution of ICG/CAC-LDH in water, PBS, and culture medium (DMEM).



Fig. S10 UV-vis-NIR spectra of ICG, CAC-LDH and ICG/CAC-LDH, respectively.



Fig. S11 Release profiles of copper (A) and cerium (B) from ICG/CAC-LDHs under various conditions. Error bars represented for standard deviation, n = 3.



Fig. S12 Cu (I) detected by the selective sequestering agent neocuproine.



Fig. S13 FL spectra of terephthalate (TA) oxidized by \cdot OH generated from the reactions between ICG/CAC-LDH and H₂O₂: (A) without 808 nm laser irradiation; (B) with 808 nm laser irradiation.



Fig. S14 TEM images of CAC-LDH after different treatments for various periods of time.



Fig. S15 Mass extinction coefficient of ICG (A) and ICG/CAC-LDH (B) at 808 nm. Normalized absorbance intensity at λ = 808 nm divided by the characteristic length of cell (A/L) at varying concentrations.



Fig. S16 Photostability tests of ICG and ICG/CAC-LDH for three cycles.



Fig. S17 Normalized absorbance of ICG and ICG/CAC-LDH at 808 nm in solutions at different pH values with H_2O_2 (0.1 mM).



Fig. S18 Cytotoxicity tests with different concentrations of CuAlCe-LDH and H_2O_2 in different pH conditions.



Fig. S19 Relative viabilities of HepG2 cells after incubated with ICG and ICG/CAC-LDH at various concentrations (quantified by ICG: 0, 5, 10, 15, 20, 25 μ g·mL⁻¹) at pH 6.5 with 808 nm laser irradiation.



Fig. S20 GSH content of HepG2 cells treated with different concentrations of CAC-LDH (0–50 μ g·mL⁻¹) for 24 h.



Fig. S21 ROS levels of DCFH-DA stained HepG2 cells with different treatments.



Fig. S22 Linear relationship between PA signal and ICG/CAC-LDH concentration under the conditions of (A) GSH (1 mM) and (B) H_2O_2 (0.1 mM). Linear relationship between T_1 -MR signal and Cu(II) concentration under the conditions of (C) GSH (1 mM) and (D) H_2O_2 (0.1 mM).

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