Supporting Information

Ferrocene-liposome-PEG: a robust ·OH/lipid peroxides nanoconverter for inducing tumor ferroptosis

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This word file contains: Fig. S1-S14. **Supplementary Figures**



Fig. S1 TEM image of Lp-PEG.



Fig. S2 The stability of Fc-Lp-PEG in PBS (a) and RPMI 1640 (b) at 37 $^\circ C$ 7 days. .



Fig. S3 The encapsulation efficiency (EE) and drug loading efficiency (DLE) of Lp-PEG to Fc at different feeding ratios.



Fig. S4 Detection of the intracellular RhB&Fc-Lp-PEG for 0 h, 1 h, 2h, 4 h, and 6 h by a fluorescence microscope. DAPI was used to stain the nuclei. RhB was used to lable Fc-Lp-PEG.



Fig. S5 The mean fluorescence intensity (MFI) of 4T1 cells with various treatments by H₂DCFDA staining.

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Fig. S6 A) Western blot and B) relative expression (R. e.) analysis of GPX-4 in 4T1 cells treated with Lp-PEG, Fc and Fc-Lp-PEG.



Fig. S7 Optical microscope images of 4T1 cells treated with Fc-Lp-PEG for 0 h, 6 h, 12 h, and 24 h.



Fig. S8 Flow cytometry analysis of 4T1 cells received various treatments *via* fluorescein Annexin V-FITC/PI double labeling.



Fig. S9 Hemolysis percentage of Fc-Lp-PEG at different mass concentrations. The RBCs treated with PBS and H₂O were negative and positive control groups.



Fig. S10 Blood biochemistry analysis of hepatorenal function markers including A) alanine aminotransferase (ALT), B) aspartate aminotransferase (AST), C) alkaline phosphatase (ALP), D) urea nitrogen (BUN), and E) creatinine (CREA). N.S.S. represents no statistical significance.



Fig. S11 The bodyweight variations of 4T1 tumor-bearing mice exposed to various treatments during 14 days.



Fig. S12 The H&E staining images of main organs of 4T1 tumor-bearing mice exposed to various treatments at day 14. The scale bar value was 100 μ m.



Fig. S13 The H&E staining images of main organs of mice exposed to Fc-Lp-PEG at day 30. The scale bar value was 100 μ m.



Fig. S14 The fluorescence emission spectrum of the ZnPC&Fc-Lp-PEG at excitation of 630 nm.