## **Supporting Information**

FePt Nanoparticles Embedded in Metal-Organic Framework Nanoparticles for Tumor Imaging and Eradication

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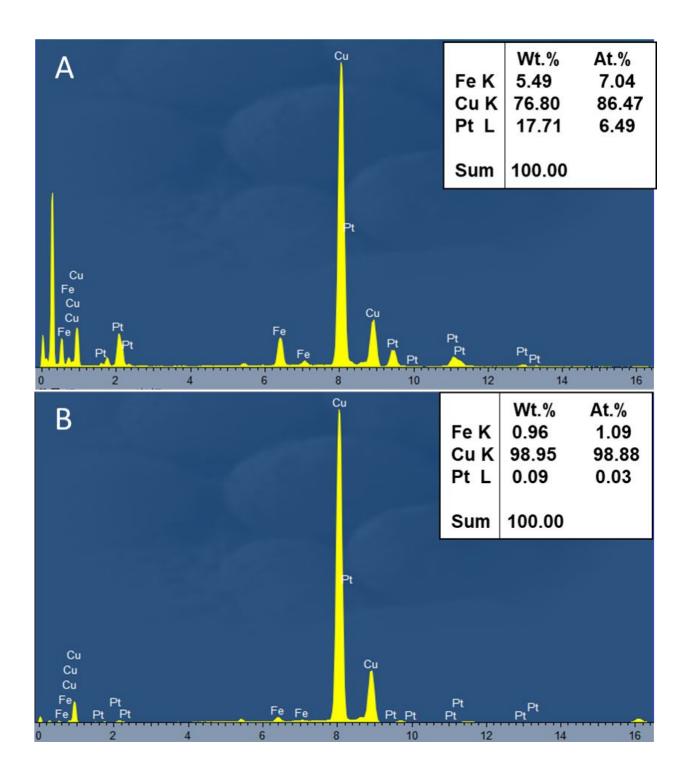


Fig. S1 (A, B) The EDS date of FePt(S) NCs and FePt(R) NCs, respectively.

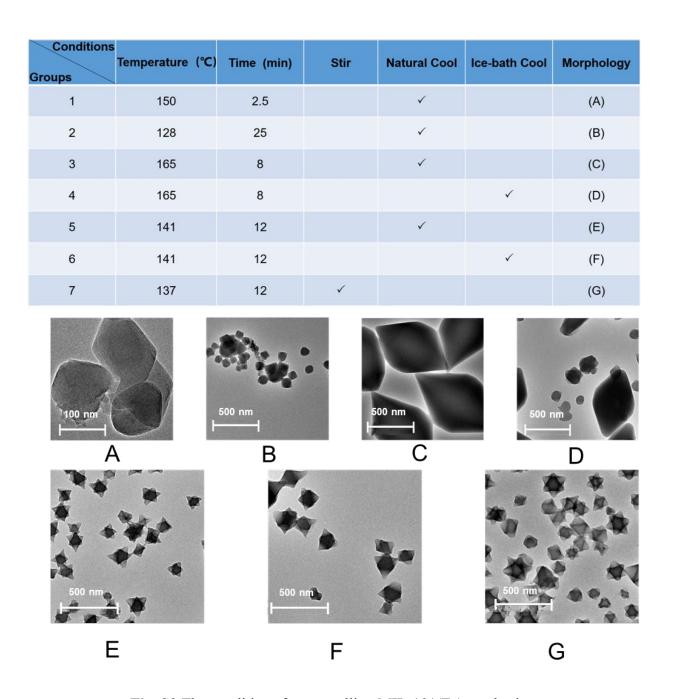


Fig. S2 The conditions for controlling MIL-101(Fe) synthesis.

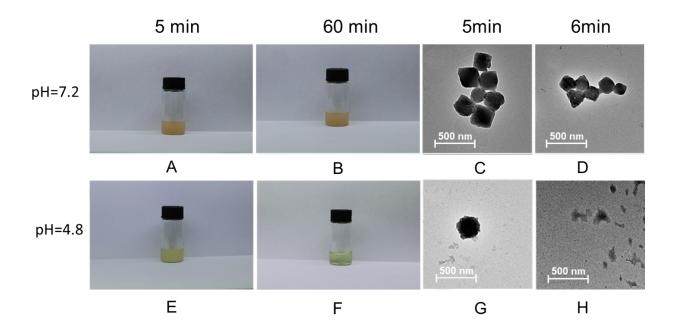
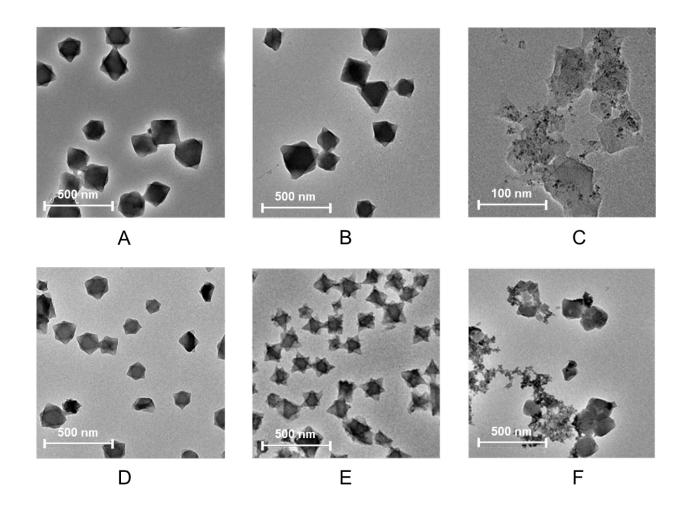
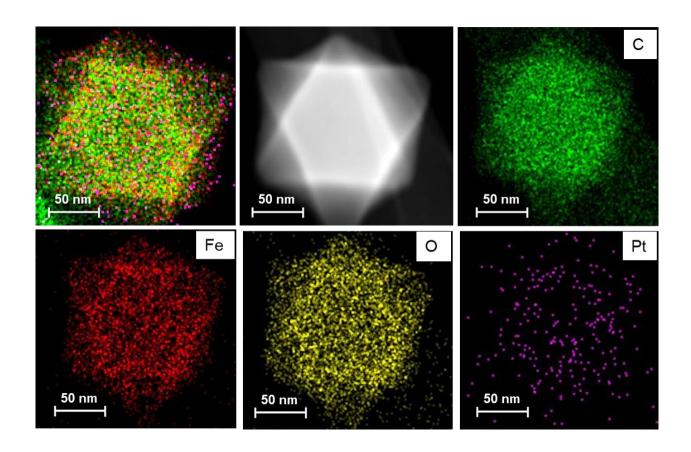


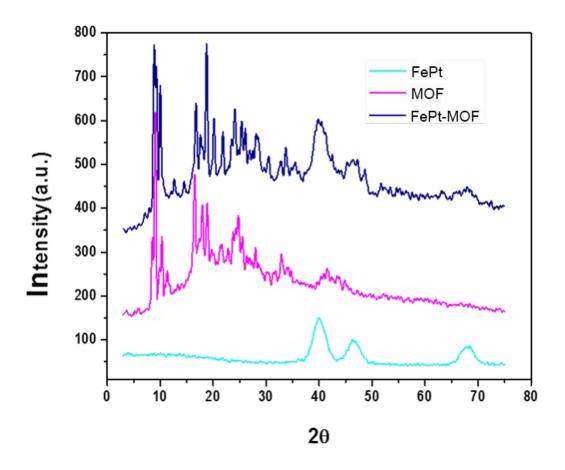
Fig. S3 The stability of the MIL-101(Fe) under the conditions of pH 7.2 and pH 4.8, respectively.



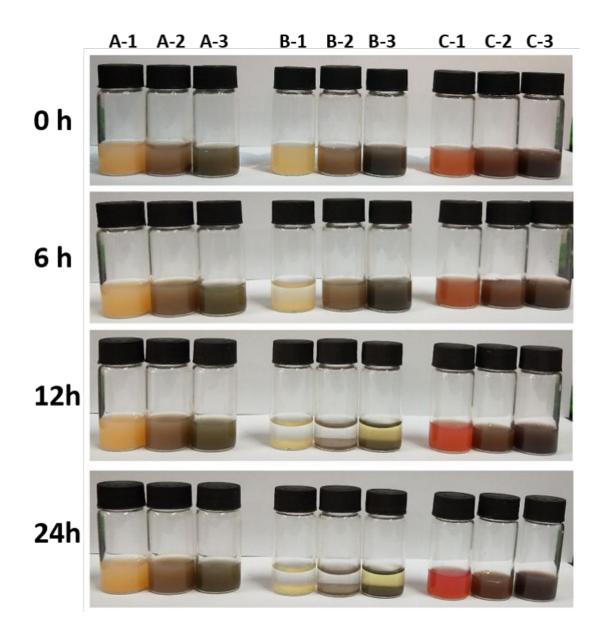
**Fig. S4** The loading capacity of the MIL-101(Fe) to FePt(R/S). (A, B, C) FePt(S) (20 mg/ 30 mg/ 40 mg), FeCl<sub>3</sub>·6H<sub>2</sub>O (0.23 g, 1.384 mol) and PTA (0.402 g, 1.384 mol) were dissolved in 60 mL DMF, heated at 141°C. (D, E, F) FePt(R) (2 mg/ 4 mg/ 6 mg), FeCl<sub>3</sub>·6H<sub>2</sub>O (0.23 g, 1.384 mol) and PTA (0.402 g, 1.384 mol) were dissolved in 60 mL DMF, heated at 141°C.



**Fig. S5** HAADF STEM image of FePt(R)-MOF nanocomposites. Corresponding elemental mapping of FePt(R)-MOF nanocomposites.



**Fig. S6** XRD patterns of FePt, MOF and FePt-MOF NCs.



**Fig. S7** The dispersion of the as-prepared MOF (1), FePt(R)-MOF-tLyp (2) and FePt(S)-MOF-tLyp-1 (3) were dissolved in  $H_2O$  (A), PBS (B) and DMEM medium (C) for 0, 6, 12 and 24 h, respectively.

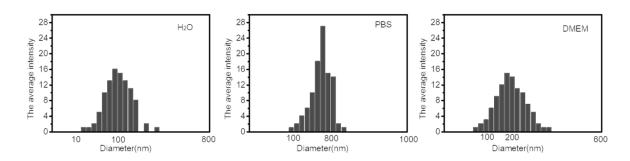


Fig. S8 The size of MOF after 24 h in H<sub>2</sub>O, PBS and DMEM medium, respectively.

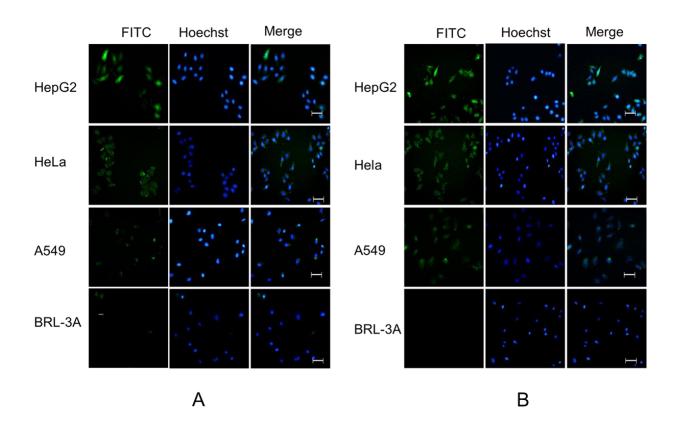
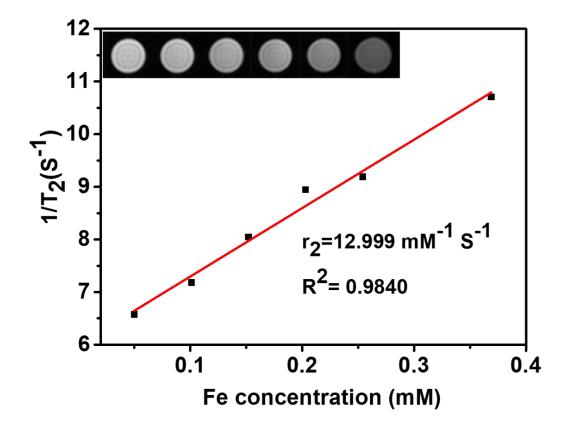


Fig. S9 The fluorescence image of FePt(S)-MOF-tLyp-FITC NCs (A) and FePt(R)-MOF-tLyp-FITC NCs (B) , respectively. (scale bars:  $200~\mu m$ )



**Fig. S10** The MR imaging  $(1/T_2)$  of FePt-MOF NCs treated with 4T1 cells. Relaxivity values were obtained from the slopes of linear fits of experimental data. (Inset:  $T_2$ -Weighted MR images of different concentrations of FePt-MOF nanocomposites)

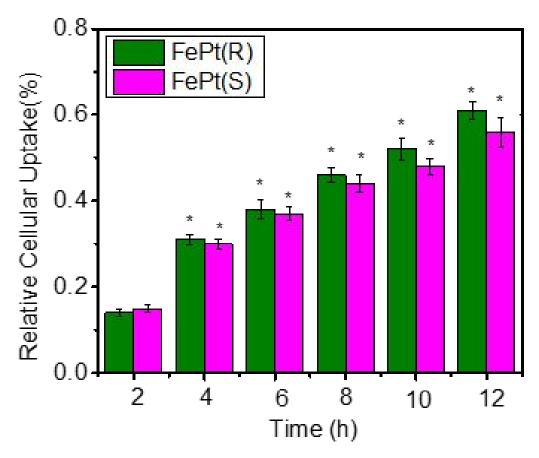
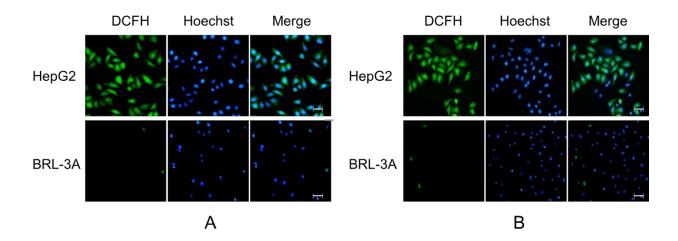
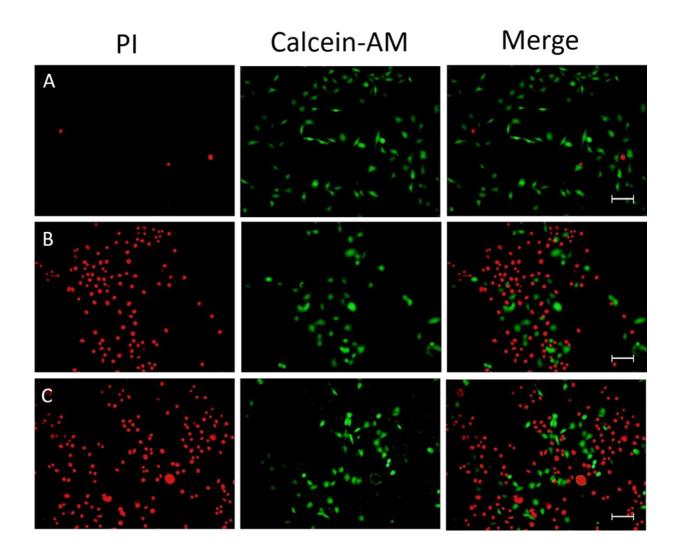


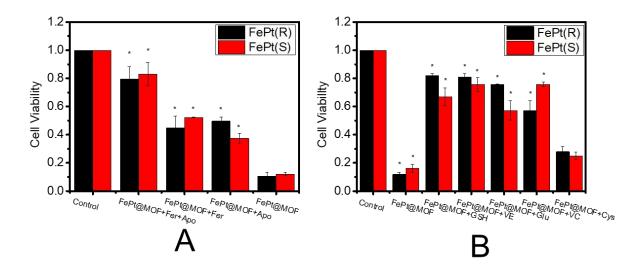
Fig. S11 The relative cellular uptake of FePt(S)-MOF-tLyp NCs and FePt(R)-MOF-tLyp NCs toward 4T1 and L02 cells under different incubation times at the Fe concentration of 30  $\mu$ g/mL. (\*p < 0.05, t-test)



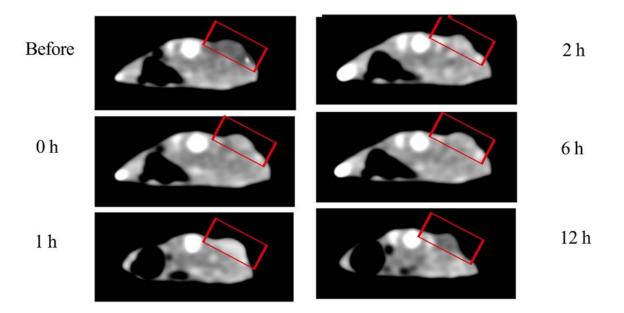
**Fig. S12** Fluorescence microscopy images of DCFH-DA labeled HepG2 and BRL-3A cells incubated with FePt(S)-MOF-tLyp-1 NCs (A) and FePt(R)-MOF-tLyp-1 NCs (B) for 8h at the Fe concentration of 25 mg mL<sup>-1</sup>, respectively. (scale bars:  $200 \, \mu m$ )



**Fig. S13** Fluorescent images of 4T1 cells co-stained by Calcein-AM (green fluorescence, live cells) and PI (red fluorescence, dead cells) of different treatments (A: Control, B: FePt(S)-MOF-tLyp-1 NCs, C: FePt(R)-MOF-tLyp-1 NCs. (scale bars: 200 μm)



**Fig. S14** (A) The corresponding survival rates of FePt-MOF-tLyp-1 NCs , FePt-MOF-tLyp-1 NCs + Lip-1 + Apo, FePt-MOF-tLyp-1 NCs + Lip-1 and FePt-MOF-tLyp-1 NCs + Apo loaded 4T1 cells. (B) Viabilities of 4T1 cells processed with FePt-MOF-tLyp-1 NCs, FePt-MOF-tLyp-1 NCs +VE, FePt-MOF-tLyp-1 NCs +VC, FePt-MOF-tLyp-1 NCs +Glu, FePt-MOF-tLyp-1 NCs +GSH and FePt-MOF-tLyp-1 NCs +Cys. (\*p < 0.05, t-test)



**Fig. S15** In vivo CT imaging (axial plane) of a 4T1 tumor-bearing mouse at different time intervals after intravenous injection with FePt-MOF-tLyp-1 NCs. Tumor tissue was indicated with red pane.

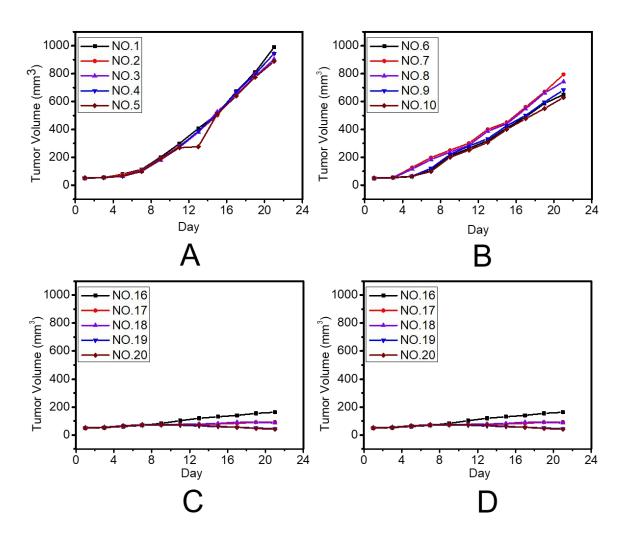
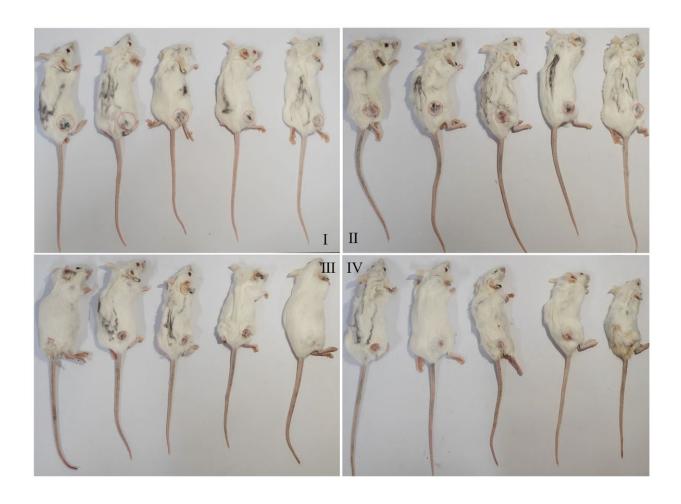


Fig. S16 The body weight of every mouse during different treatments.



**Fig. S17** The images of mouse for different groups. ( I : PBS; II : MOF-tLyp-1; III : FePt(R)-MOF-tLyp-1; IV: FePt(S)-MOF-tLyp-1).

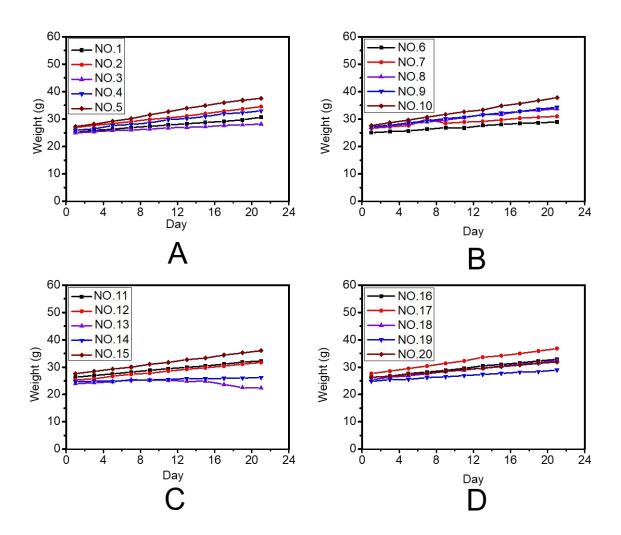
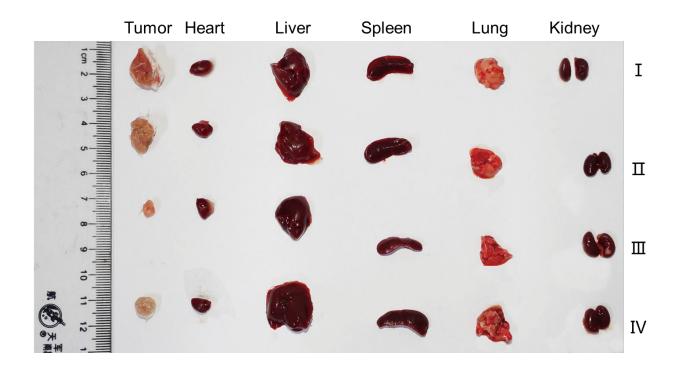


Fig. S18 The volume of tumors in tumor-bearing mice of every mouse during different treatments.



**Fig. S19** The photographs of major organs and tumors after various treatments. ( I : PBS; II : MOF-tLyp-1; III: FePt(R)-MOF-tLyp-1; IV: FePt(S)-MOF-tLyp-1)