Electronic Supplementary Information

Self-Assembly of Paramagnetic Amphiphilic Copolymers for Synergistic Therapy

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Fig. S1. Synthesis of PCL-*b*-PIEtMn.



Fig. S2. ¹H NMR spectra (400MHz, CDCl₃) of PCL, PCL-MDFC, PCL-*b*-PDMAEMA

and PCL-*b*-PIEt.



Fig. S3. Zeta potential of NPs with different mass ratio of PCL-*b*-PEG and PCL-*b*-PIEtMn.



Fig. S4. Stability of paramagnetic NPs in DPBS and culture medium.



 DLC_{DOX} = (weight of loaded drug / weight of polymer) = 5.3% DLC_{IR780} = (weight of loaded drug / weight of polymer) = 2.0%

Fig. S5. Standard curve and drug loading contents (DLC) of (a) DOX and (b) IR-780.



Fig. S6. CLSM images and fluorescence intensity analysis of MCF-7 cells after incubation with free (a) DOX, (b) DOX&IR-780@NPs L-, and (c) DOX&IR-780@NPs L+, respectively.



Fig. S7. Cross-section CLSM images of MCF-7 cells after incubation with DOX&IR-

780@NPs L+. (a) Nucleus, (b) cell membrane, (c) DOX, and (d) overlay.



Fig. S8. Cytotoxicity of paramagnetic NPs. MCF-7 cells were incubated with paramagnetic NPs for 48 h before the ATPlite assay.



Fig. S9. Biodistribution of IR-780 in 4T1 tumor-bearing mice after intravenous injection of free IR-780 and DOX&IR-780@NPs at different time points.



Fig. S10. Qualitative analysis of IR-780 intensities in different organs and tumors after injection of free IR-780 and DOX&IR-780@NPs at different time points. (a) 12 h, (b) 48 h.



Fig. S11. H&E staining images of major organs after treatments. Scale bar is 100 µm.