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Supporting Information

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Inhibiting Solid Tumor Growth In Vivo by Non-Tumor-Penetrating Nanomedicine

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Table S1. Characterizations of the PPD and DOX-PPD micelles.

Entry	R_h (nm)[a] at 25 °C	Zeta potential[b] (mV)	DLC of DMXAA[c] (wt%)	DLC of DOX[d] (wt%)
PPD	18.7 ± 3.4	-4.61 ± 0.51	11.8	--
DOX-PPD	31.5 ± 9.4	-2.97 ±1.04	11.1	5.4

a. Measured by DLS; b. Estimated at pH 7.4 at 25 °C, a mean ± STD of 6 measurements; c. Determined by UV-Vis at 343 nm; d. Determined by UV-Vis at 480 nm.

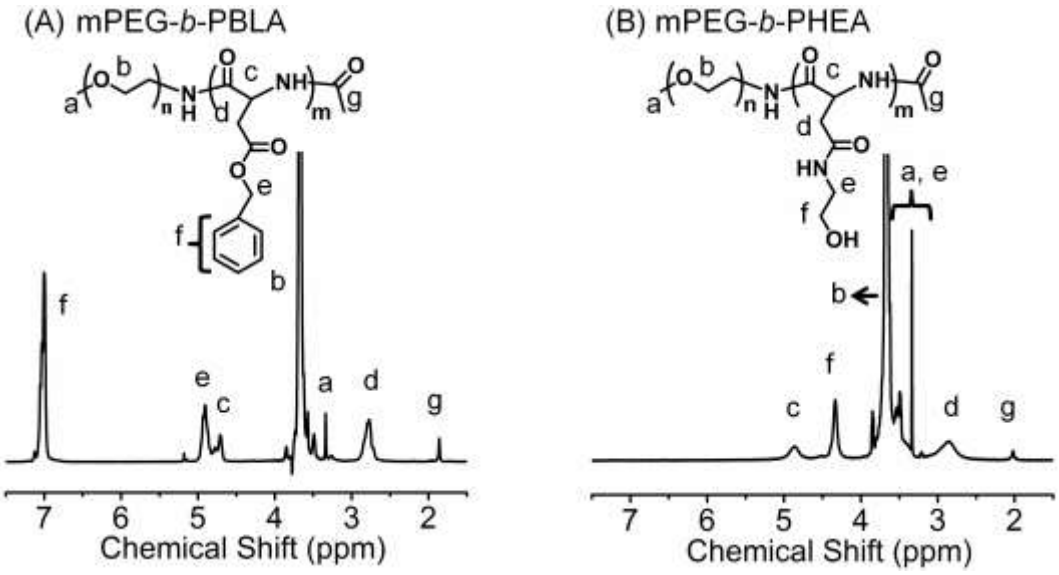

Figure S1. ^{1}H NMR spectra of mPEG-*b*-PBLA (A), mPEG-*b*-PHEA (B) in CF_3COOD.

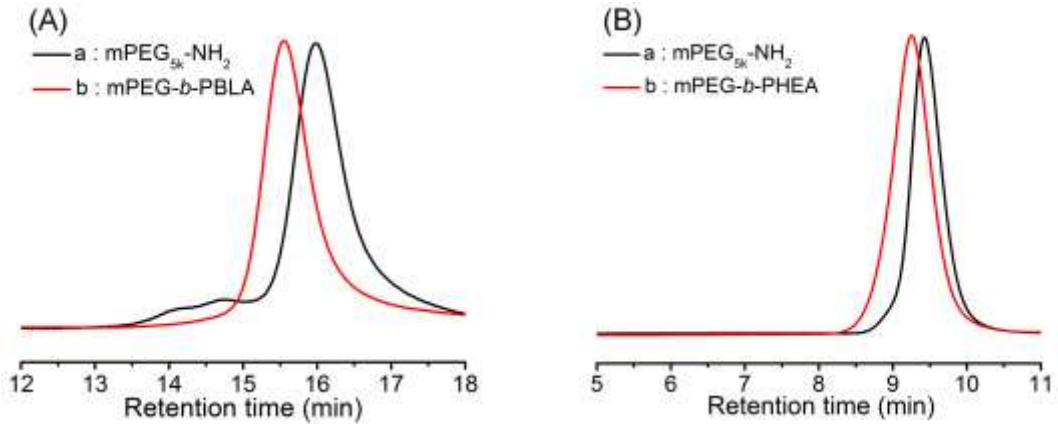

Figure S2. (A), GPC traces for $mPEG_{5k}$-NH_2 (a) and mPEG-*b*-PBLA (b) using DMF as eluent. (B), GPC traces for $mPEG_{5k}$-NH_2 (a) and mPEG-*b*-PHEA (b) using acetate buffer solution (0.1 M, pH 2.8) as eluent.

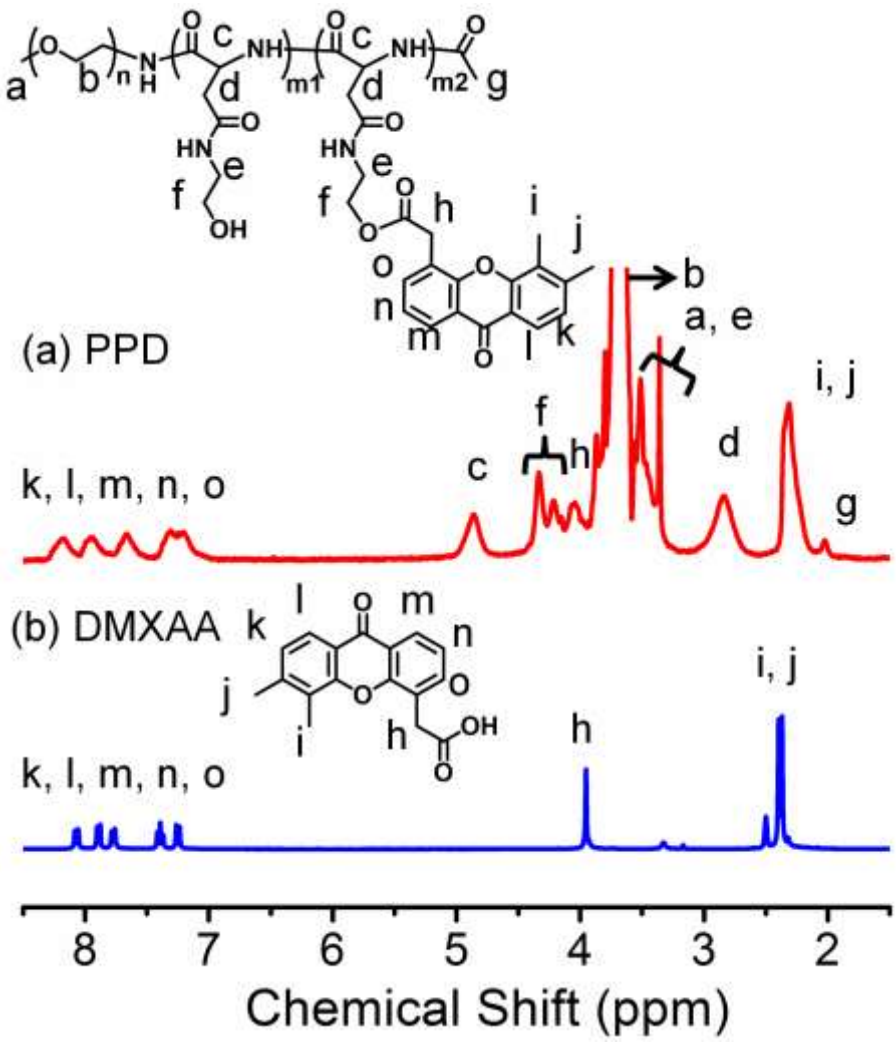

Figure S3. ^{1}H NMR spectra of PPD (a) and DMXAA (b) in CF_3COOD.

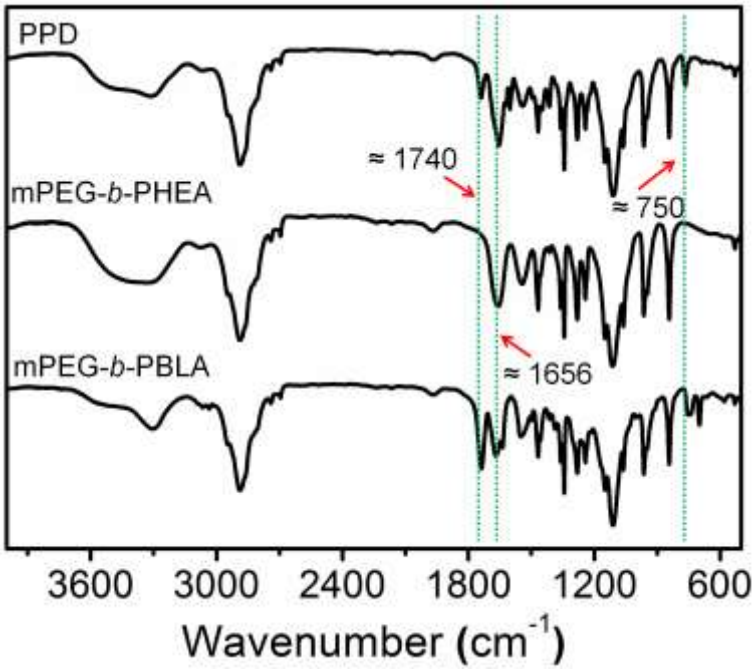

Figure S4. FT-IR spectra of mPEG-*b*-PBLA, mPEG-*b*-PHEA and PPD.

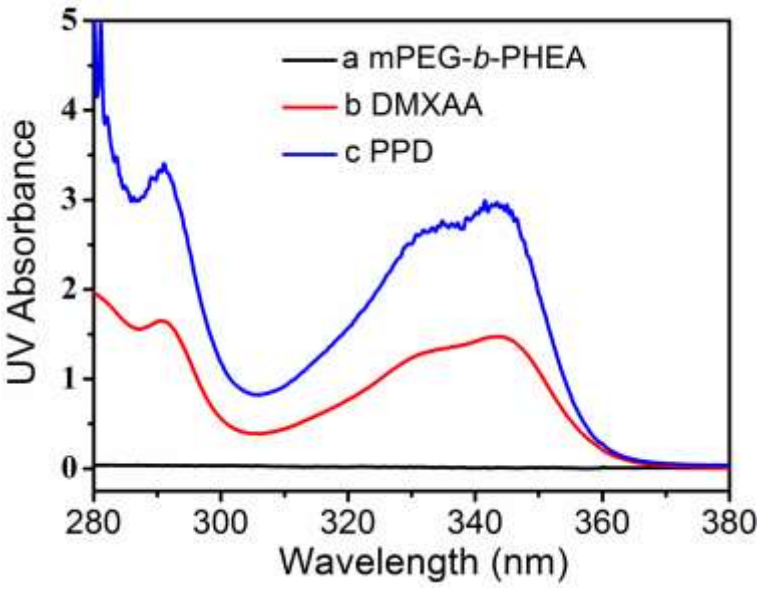

Figure S5. The UV-Vis measurements of mPEG-*b*-PHEA (1.0 mg mL^{-1}), DMXAA (0.0625 mg mL^{-1}) and PPD (1.0 mg mL^{-1}) using DMF as a solvent.

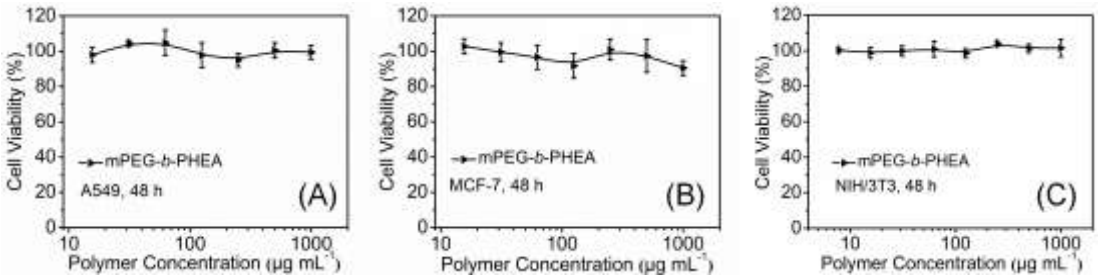

Figure S6. *In vitro* cytotoxicities of mPEG-*b*-PHEA to A549, MCF-7 and NIH/3T3 cells (A, B and C).

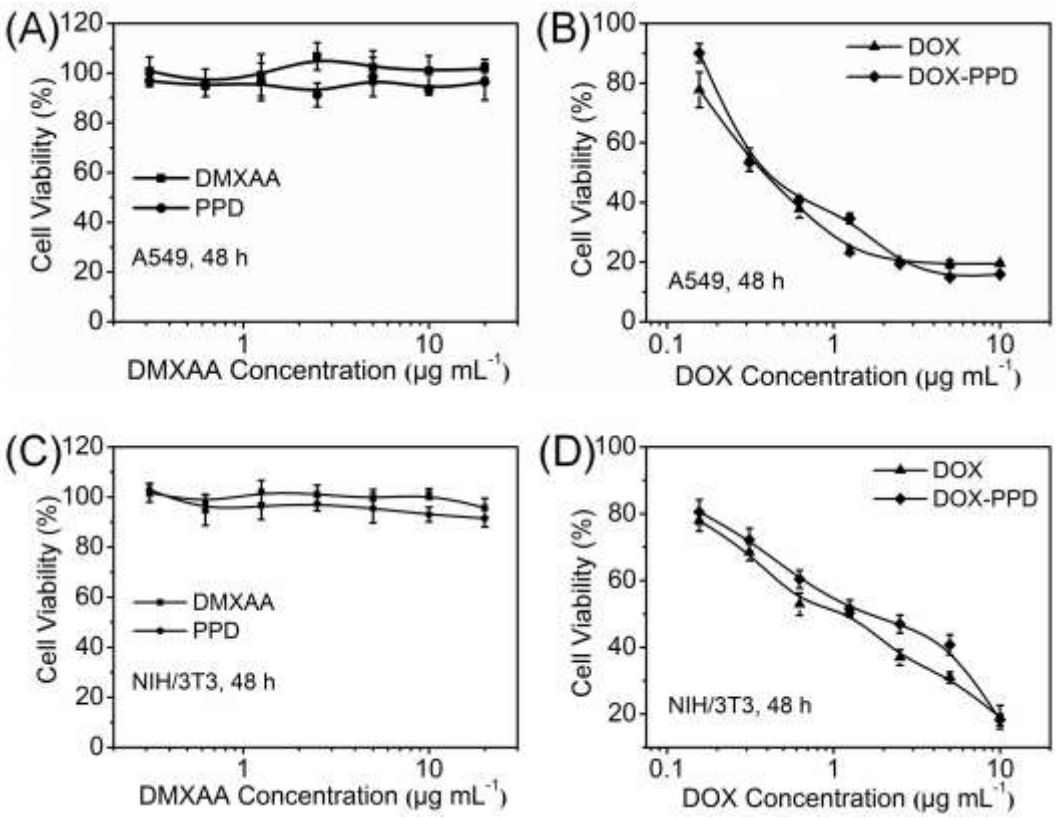

Figure S7. *In vitro* cytotoxicity of free DMXAA, PPD, free DOX and DOX-PPD to A549 and NIH/3T3 cells.

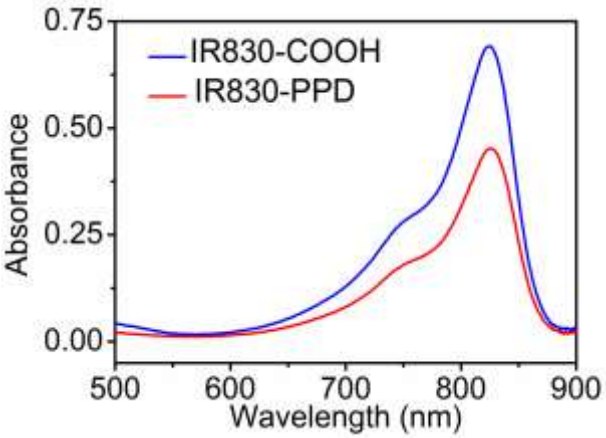

Figure S8. UV/Vis spectra of IR830-COOH and IR830-PPD in DMF (concentrations: IR830-COOH, 0.005 mg mL^{-1}; IR830-PPD, 0.05 mg mL^{-1}).

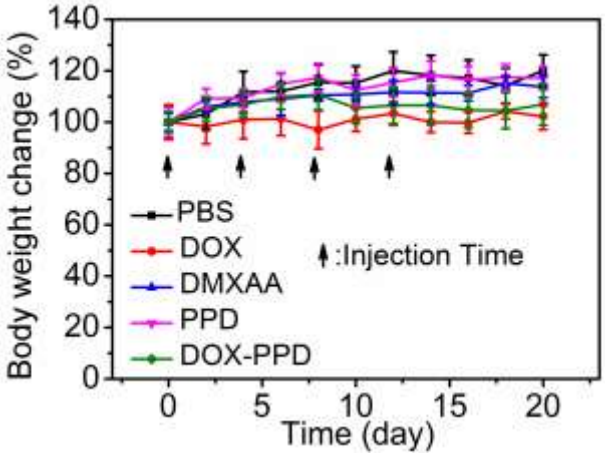

Figure S9. Body weight change of MCF-7 tumor bearing nude mice during different treatments: PBS, free DOX (5 mg kg^{-1}), DMXAA (10 mg kg^{-1}), PPD (10 mg DMXAA kg^{-1}) and DOX-PPD (5 mg DOX kg^{-1} and 10 DMXAA kg^{-1}). The data are shown as mean ± SD (n = 6).