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A multifunctional biodegradable brush polymer-drug conjugate for paclitaxel/gemcitabine co-delivery and tumor imaging

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Supporting Figures

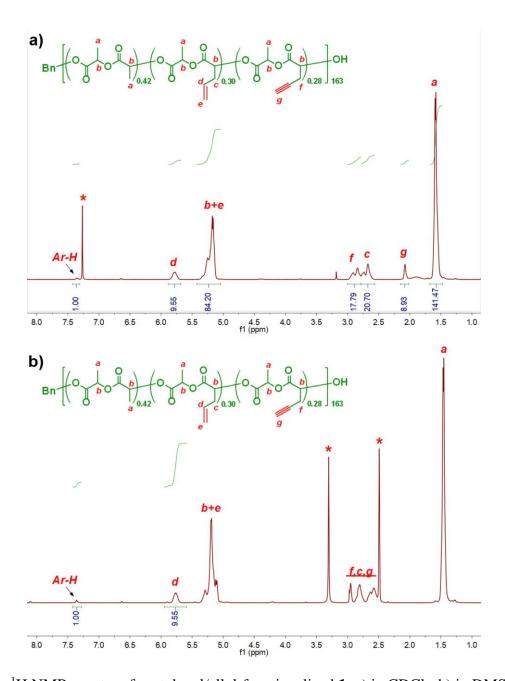


Fig. S1 ¹H NMR spectra of acetylenyl/allyl-functionalized **1**: a) in CDCl₃, b) in DMSO-*d*₆.

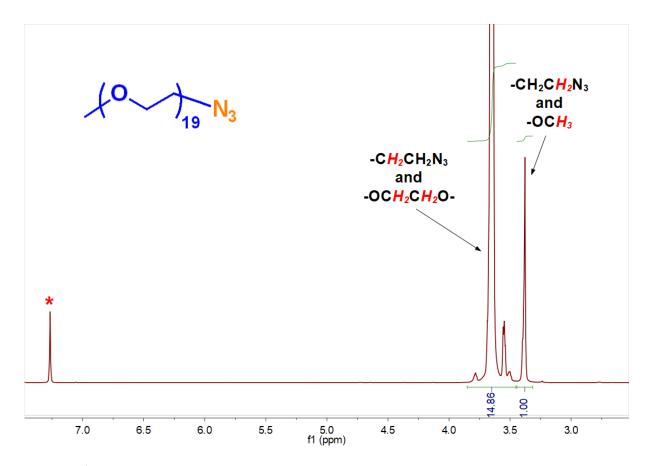


Fig. S2 $\,^{1}$ H NMR spectra of CH₃O-PEG-N₃ in CDCl₃.

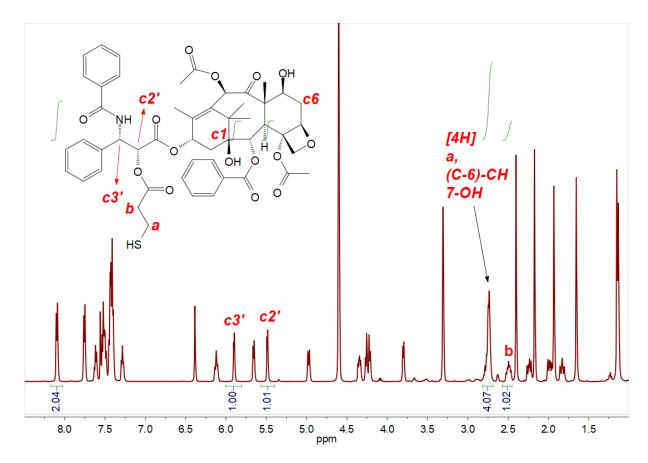


Fig. S3 $\,^{1}$ H NMR spectrum of PTX-SH in CDCl₃/CD₃OD (v/v, 1:1).

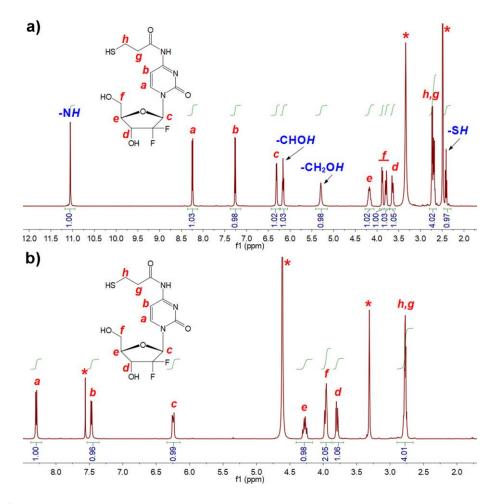


Fig. S4 1 H NMR spectra of GEM-SH in a) DMSO- d_6 , b) CDCl₃/CD₃OD (v/v, 1:1).

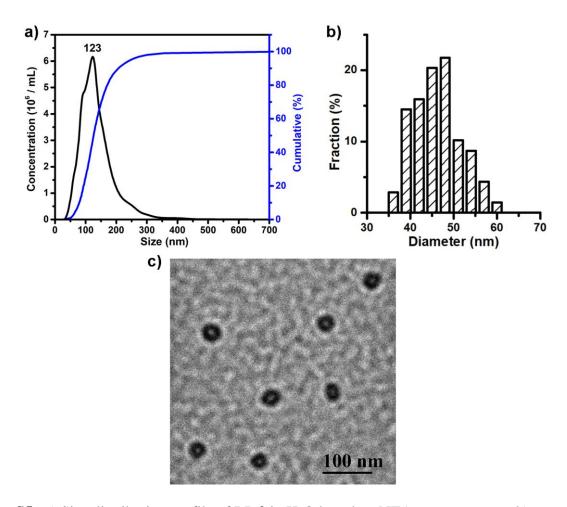


Fig. S5 a) Size distribution profile of BP **2** in H₂O based on NTA measurements. b) Histogram of diameter distribution of BP **2** according to the TEM images. c) TEM images of BP **2**. TEM samples were stained by RuO₄.

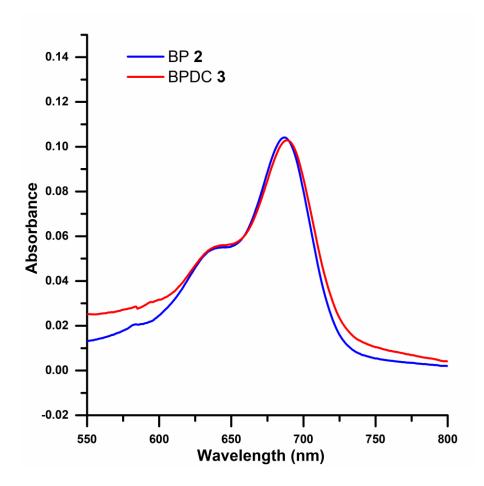


Fig. S6 Absorbance spectra of BP 2 and BPDC 3.

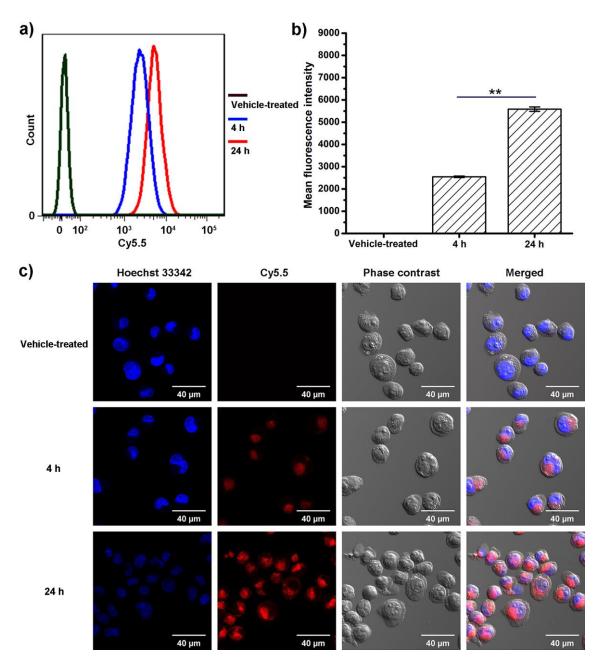


Fig. S7 a) Flow cytometry data showing cellular uptake of BP **2** in MIA PaCa-2 cells at 4 and 24 h post treatment compared with untreated cells. b) Mean fluorescence intensity of Cy5.5 obtained from flow cytometry at 4 and 24 h post treatment. c) Confocal microscopy images of MIA PaCa-2 cells at 4 and 24 h post treatment with BP **2**, compared with untreated cells. Cell nuclei were counterstained with Hoechst 33342. Data were reported as mean \pm standard deviation (SD) of three independent experiments. One-way ANOVA was used to assess the statistical significance of the data. **p < .01 was considered as statistical highly significant.