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Supporting Information

Polymeric Nanoparticles Controlled by On-chip Self-Assembly Enhance Cancer Treatment Effectiveness

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Scheme S1. Schematic illustration of (a) Formation of NPs via SMR chip. pPBA and DOX solution were pumped into two inlets, and the synthesized NPs were pumped out via outlet. Scale bar = 10 mm. (b) chemical interaction for NP formation and dissociation depending on pH.



Figure S1. (a) Synthesis scheme of pPBA and NPs. Diol of DOX can interact with PBA of pPBA. (b) 1 H NMR spectra of pPBA.



Figure S2. NP formation at various ratios via (a) BM method and (b) MV method with Re 250, and then the NPs were analyzed with DLS (n = 3).



Figure S3. (a) Schematic illustration of PBA fluorescence quenching. (b) Quenching of PBA fluorescence of pPBA, BMPs and MVPs at Re 250 and (c) Re-dependency of MVP2 and MVP4. (d) CFD results for mixing efficiency of SMR at each height depending on Re.



Figure S4. (a) Cytotoxicity of DOX and the NPs evaluated by MTT assays after 48 h incubation. (b) Cytotoxicity of pPBA at each cell line after 48 h incubation. Negligible cytotoxicity of pPBA was observed.



Figure S5. (a) Confocal fluorescence images of Daoy cell showing cellular uptake of NPs for 4 h. Blue, red, and green images represent nucleus, DOX, and pPBA labelled with

fluorescence dye, respectively. Scale bar is 50 μ m. (b) Relative DOX and pPBA mean fluorescence intensities of (a). Each MFI was normalized maximum intensity.