Supporting Information

A multifunctional nanoplatform for cancer chemo-photothermal synergistic therapy and overcoming multidrug resistance

Yunmei Peng,^{a,b,1} Junpeng Nie,^{a,b,1} Wei Cheng,^{b,1} Gan Liu,^c Dunwan Zhu,^{d,*} Linhua Zhang,^d Chaoyu Liang,^b Lin Mei,^c Laiqiang Huang,^{a,b,*} and Xiaowei Zeng^{a,b,c,*}

^a School of Life Sciences, Tsinghua University, Beijing 100084, China

^b Division of Life and Health Sciences, Graduate School at Shenzhen, Tsinghua University, Shenzhen 518055, China

^c School of Pharmaceutical Sciences (Shenzhen), Sun Yat-sen University, Guangzhou 510275, China

^d Institute of Biomedical Engineering, Chinese Academy of Medical Sciences & Peking Union Medical College, Tianjin Key Laboratory of Biomedical Materials, Tianjin 300192, China

¹ These authors contributed equally to this work.

* Corresponding authors.

E-mail: zeng.xiaowei@sz.tsinghua.edu.cn or xwzeng@163.com (X. Zeng)

E-mail: huanglq@sz.tsinghua.edu.cn (L. Huang)

E-mail: zhudunwan@bme.pumc.edu.cn (D. Zhu)



Scheme S1. The oxidative self-polymerization mechanism of dopamine and the conjugation mechanism between H_2N -TPGS and PDA coating.



Figure S1. DLS size distribution of DTX-loaded PLGA NPs (A), PLGA NPs@PDA (B), PLGA NPs@PDA-PEG (C) and PLGA NPs@PDA-TPGS (D).



Figure S2. XPS spectra analysis of PLGA NPs, PLGA NPs@PDA, PLGA NPs@PDA-PEG and PLGA NPs@PDA-TPGS.



Figure S3. *In vitro* release profiles of the DTX-loaded PLGA NPs, DTX-loaded PLGA NPs@PDA and DTX-loaded PLGA NPs@PDA-TPGS. (A) without NIR; (B) with NIR.



Figure S4. DTX concentration-time profile following intravenous administration of Taxotere®, DTX-loaded PLGA NPs@PDA, DTX-loaded PLGA NPs@PDA-PEG and DTX-loaded PLGA NPs@PDA-TPGS in SD rats at the DTX dose of 10 mg/kg (n = 5).

DMEM			
Samples(n=3)	Size (nm)	PDI	ZP(mV)

Table S1. The dispersivity of DTX-loaded PLGA NPs@PDA-TPGS in PBS and

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DTX-loaded PLGA NPs @PDA-TPGS (PBS)	181.2 ± 2.8	0.129	3.2±0.3
DTX-loaded PLGA NPs @PDA-TPGS (DMEM)	179.5 ± 5.7	0.113	3.5±0.3

Table S2. IC₅₀ values of Taxotere[®], DTX-loaded PLGA NPs@PAD-PEG, DTX-loaded PLGA NPs@PAD-TPGS and DTX-loaded PLGA NPs@PAD-TPGS+NIR on MCF-7 cells after 24 h and 48 h incubation

	IC ₅₀ (µg/ml)			
Incubation		DTX-loaded	DTX-loaded	DTX-loaded PLGA
time (h)	Taxotere®	PLGA NPs @	PLGA NPs	NPs@PAD-
		PAD-PEG	@PAD-TPGS	TPGS+NIR
24	19.21±1,53	21.61±1.37	23.00±1.45	10.77±1.87
48	10.96±0.78	4.64±0.69	2.16±0.73	1.41±0.45

Table S3. IC₅₀ values of Taxotere®, DTX-loaded PLGA NPs@PAD-PEG, DTX-loaded PLGA NPs@PAD-TPGS and DTX-loaded PLGA NPs@PAD-TPGS+NIR on MCF-7/ADR cells after 24 h and 48 h incubation

		IC ₅₀ (μg/ml)			
Incubation		DTX-loaded	DTX-loaded	DTX-loaded PLGA	
time (h)	Taxotere®	PLGA NPs @	PLGA NPs	NPs@PAD-	
		PAD-PEG	@PAD-TPGS	TPGS+NIR	
24	17.06±1.829.	22.45±1,92	5.65±1,37	1.45±1.09	
48	73±0.66	4.25±0.54	1.59±0.61	0.67±0.39	