

Fabrication of Poly Dopamine@poly (Lactic Acid-Co-Glycolic Acid) Nanohybrids for Cancer Therapy via a Triple Collaboration Strategy

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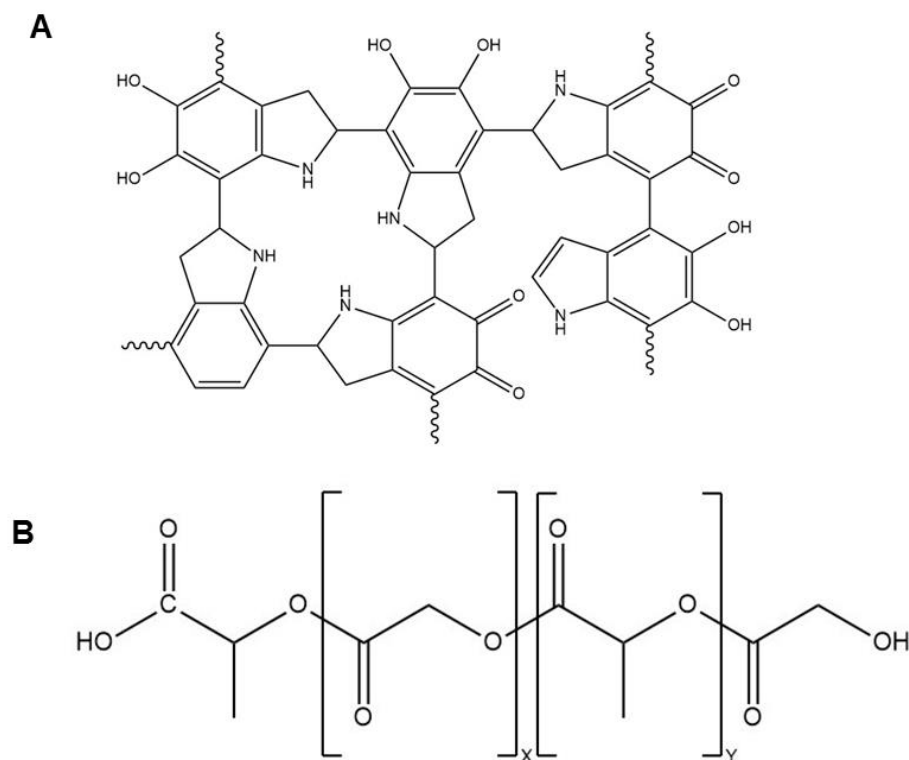


Figure S1. The structure PDA (A) and PLGA (B).

Table S1. Influence of the mass of DOX on encapsulation efficiency (EE) and loading content (LC) of PDA@PLGA/DC NPs.

PLGA (mg)	DOX (μ g)	EE (%)	LC (%)
10	100	0	0
10	200	0	0
10	300	33	0.99
10	400	40	1.6
10	500	16.5	0.83

Table S2. Influence of the mass of CA4 on encapsulation efficiency (EE) and loading content (LC) of PDA@PLGA/DC NPs.

PLGA (mg)	CA4 (μ g)	EE (%)	LC (%)
10	100	41.8	0.42
10	200	23.4	0.47
10	300	0	0
10	400	0	0
10	500	0	0

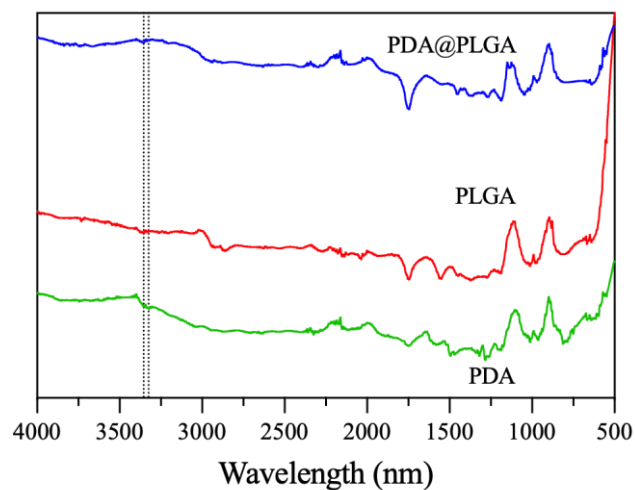


Figure S2. FT-IR spectra of PDA@PLGA NPs , PLGA NPs and PDA NPs.

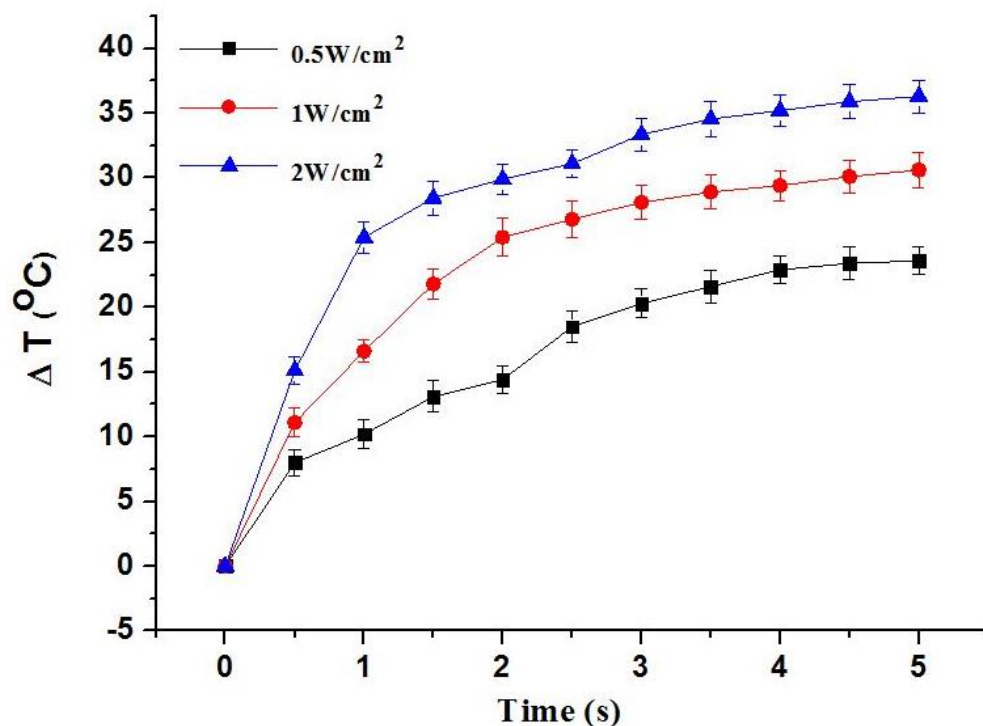


Figure S3. Temperature variation curves of PDA@PLGA/DC NPs at various power intensities with the same concentration at 150 $\mu\text{g/mL}$.

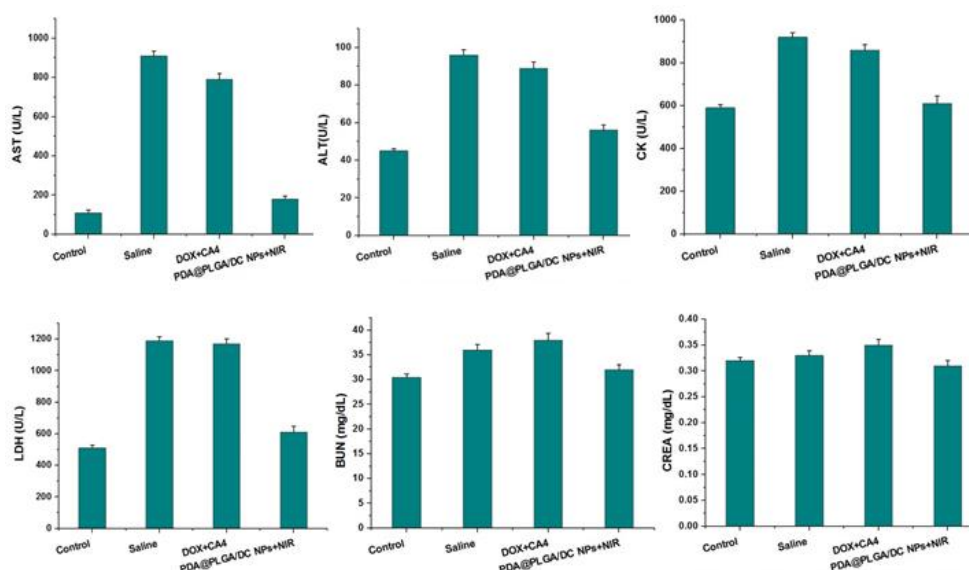


Figure S4. Serum biochemical examination of drug-loaded nanoparticles treated mice by saline, free DOX+CA4 and PDA@PLGA/DC NPs+NIR. Normal BALB/c mice served as the control group. Data were presented as mean \pm SD (n=5). Hepatic function indicators: AST, aspartate transaminase, ALT, alanine aminotransferase; Cardiac function indicators: CK, creatine kinase, LDH, lactate dehydrogenase; Renal function indicators: BUN, blood urea nitrogen, CREA, creatinine.