

Supplementary Information

Construction of GSH-triggered cationic fluoropolymer as two-in-one nanoplatform for combined chemo/gene therapy

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Table S1. Relative transfection efficiency of PSSF (according to PEI 25k) at various mass ratios (with reference to Figure 4C).

	w/w=10	w/w=15	w/w=20	w/w=25
HeLa	1.58	5.35	9.93	14.11
U2OS	5.51	3.91	3.32	2.67

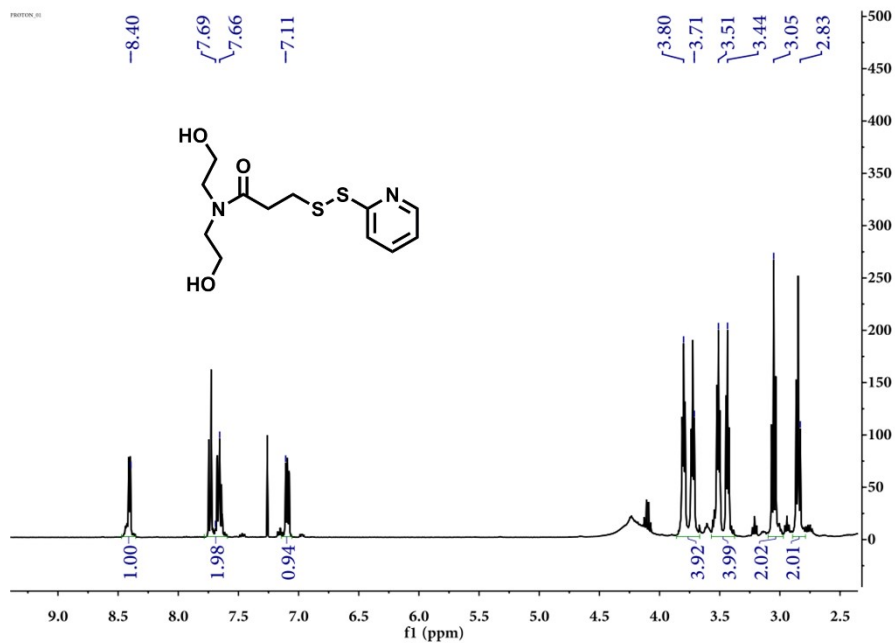


Figure S1. ¹H NMR spectrum of compound 2 in CDCl₃.

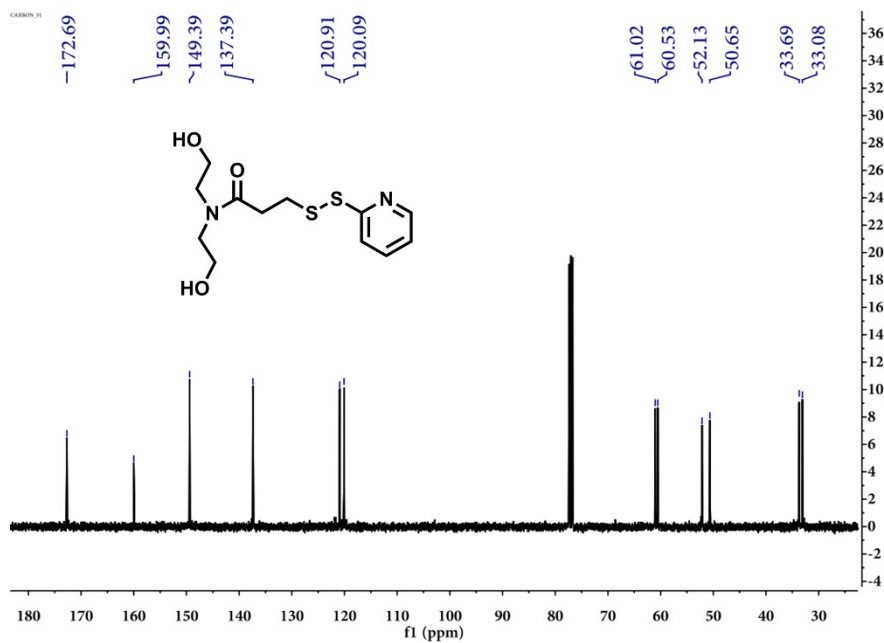


Figure S2. ^{13}C NMR spectrum of compound **2** in CDCl_3 .

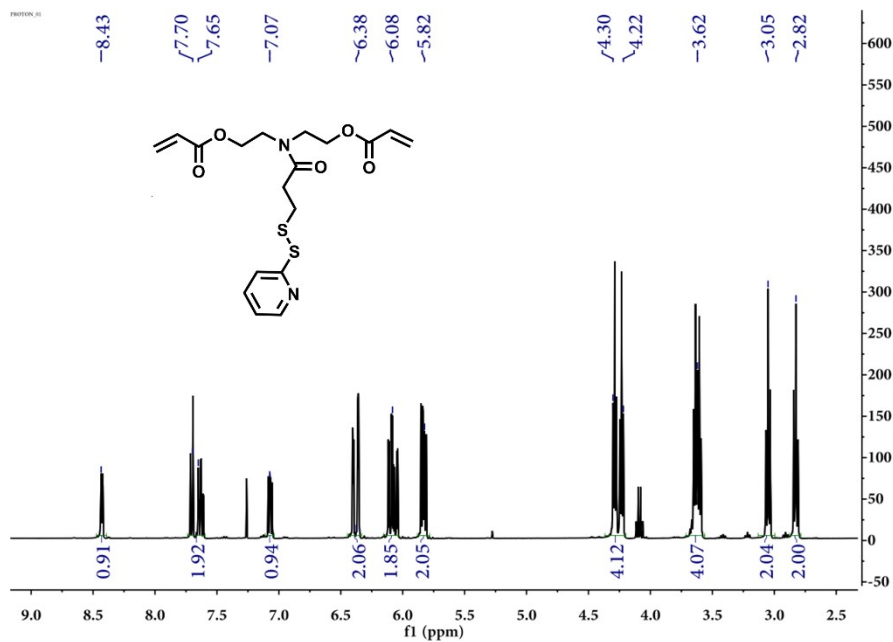


Figure S3. ^1H NMR spectrum of compound **SS** in CDCl_3 .

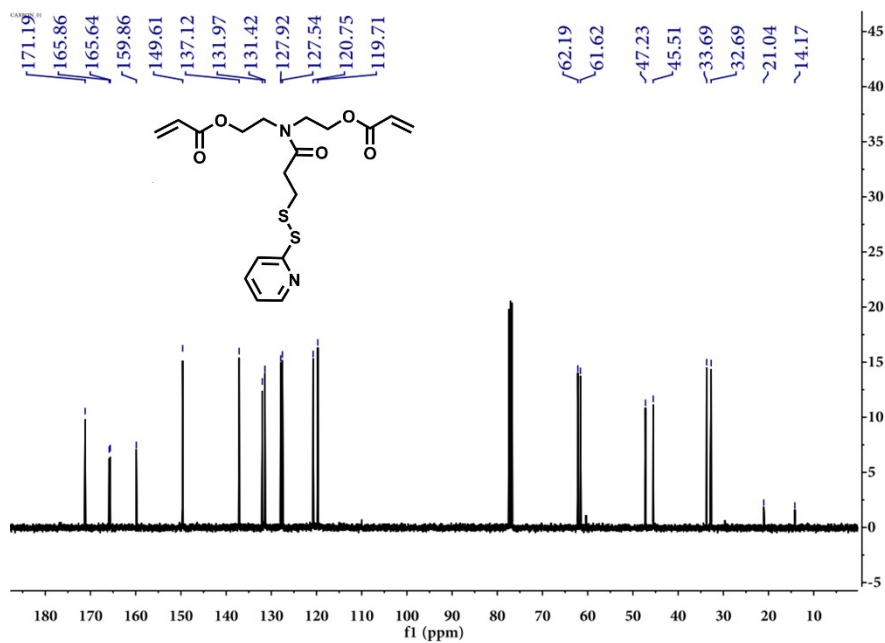


Figure S4. ¹³C NMR spectrum of compound SS in CDCl₃.

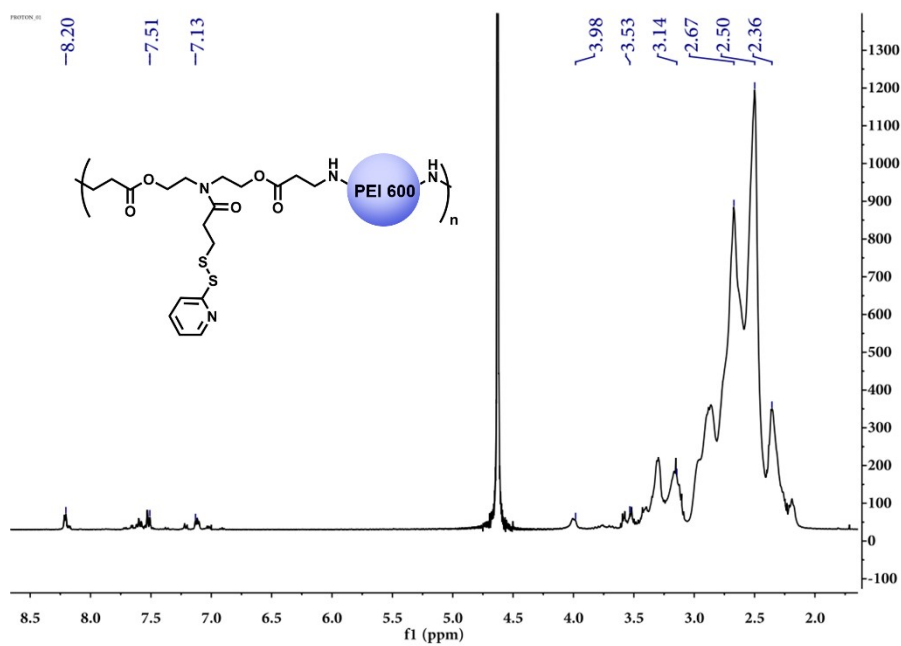


Figure S5. ¹H NMR spectrum of PSS in D₂O.

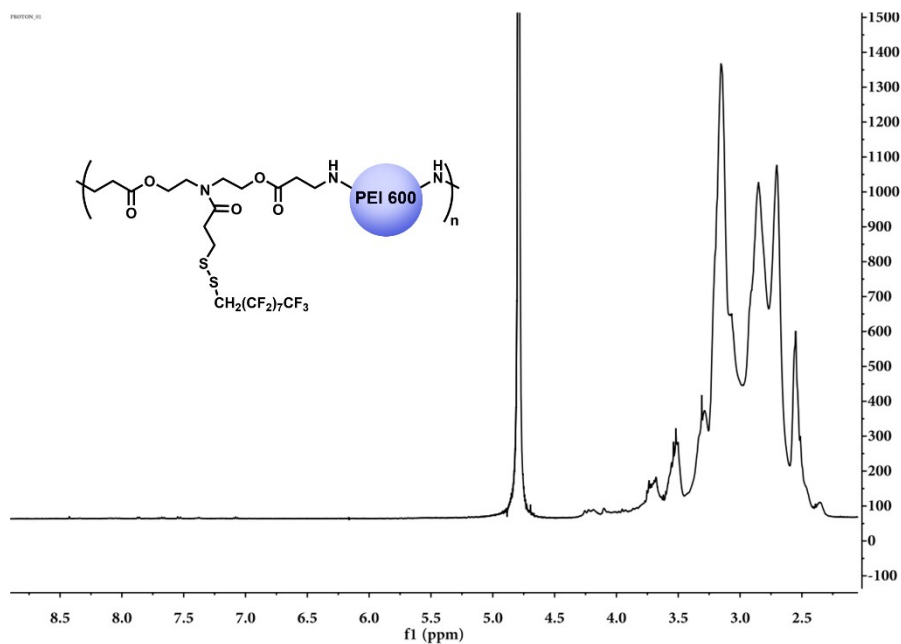


Figure S6. ¹H NMR spectrum of PSSF in D₂O.

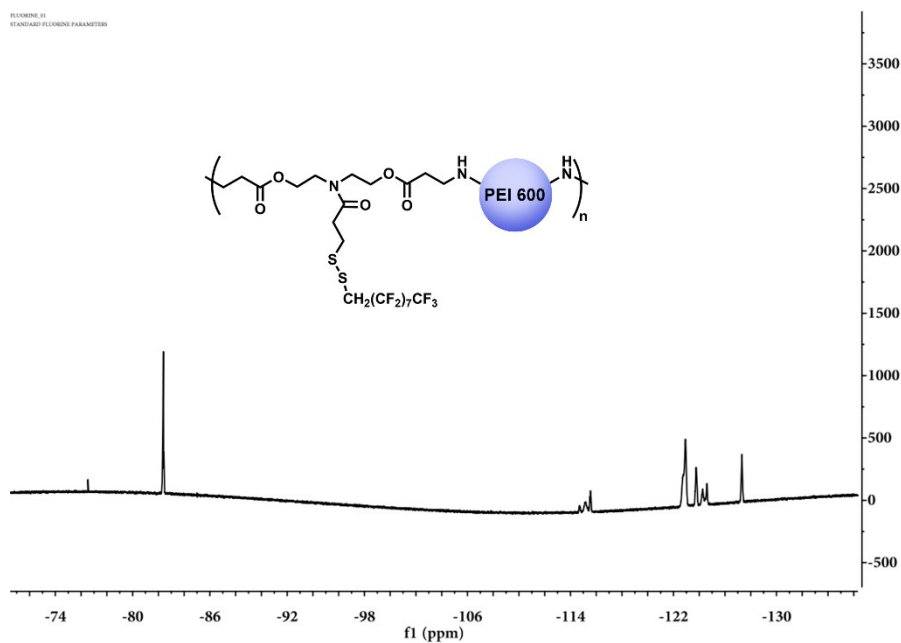


Figure S7. ¹⁹F NMR spectrum of PSSF in MeOD.

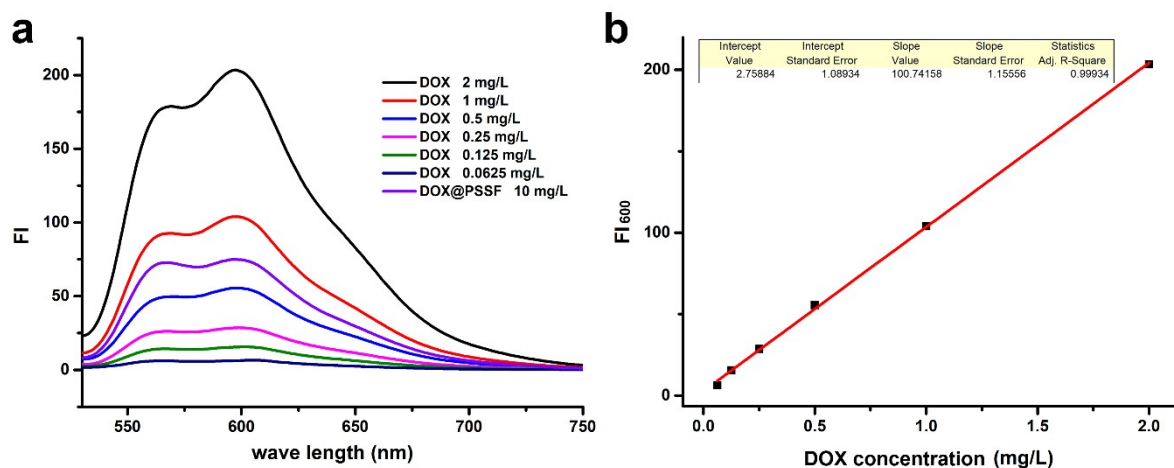


Figure S8. (a) The fluorescence spectrum of DOX and DOX@PSSF at different concentrations. (b) Standard curve of FI₆₀₀-DOX concentration.

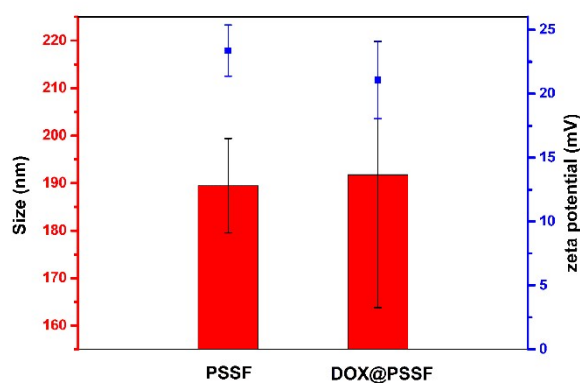


Figure S9. Particle size (columns) and zeta potential (dots) of PSSF and DOX@PSSF.

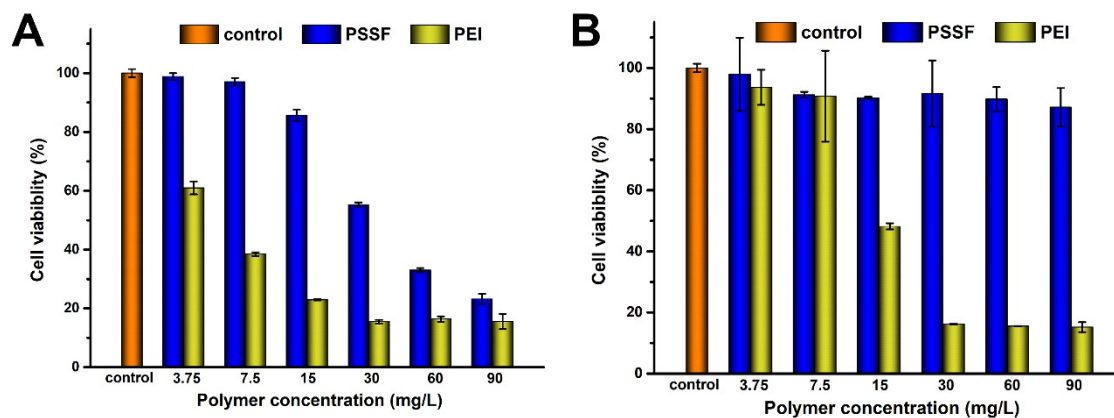


Figure S10. Cell viability after treatment with different PSSF concentrations for 48 h in 7702(A) and DC(B) cells

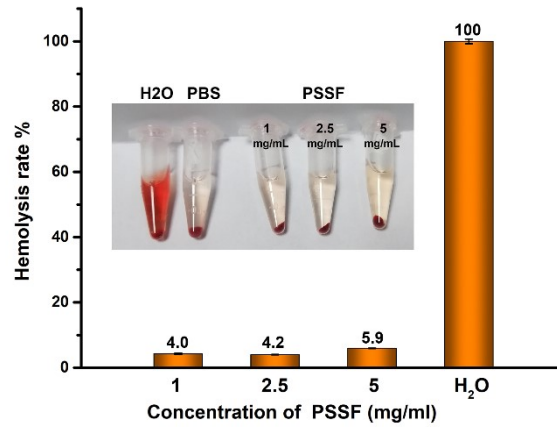


Figure S11. Hemolysis rate of **PSSF** at different concentrations after incubation with red blood for 2 h. Data represent mean \pm SD (n=3).

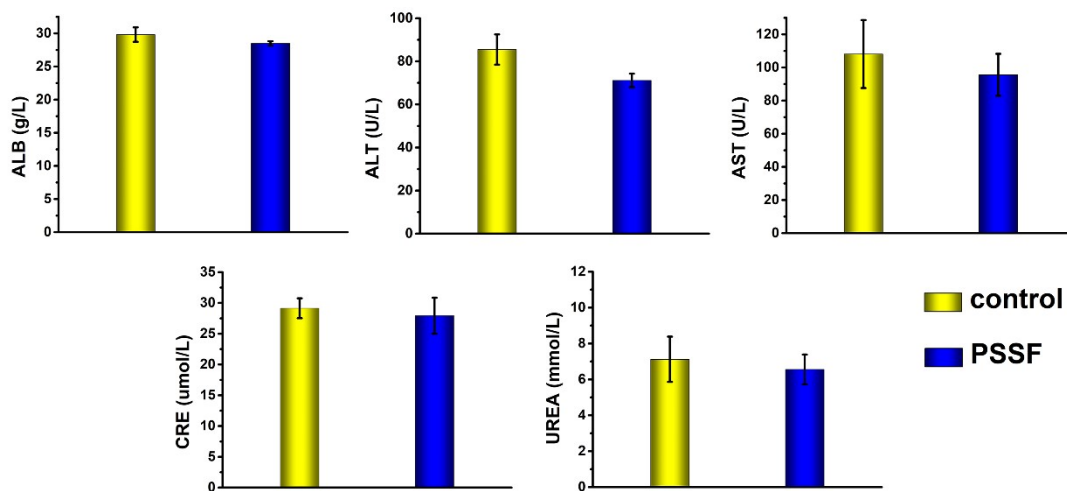


Figure S12. The blood biochemical indexes (ALB, ALT, AST, CRE, UREA) of mice injected with **PSSF** (50 mg/kg, once every other day, 3 times in total).

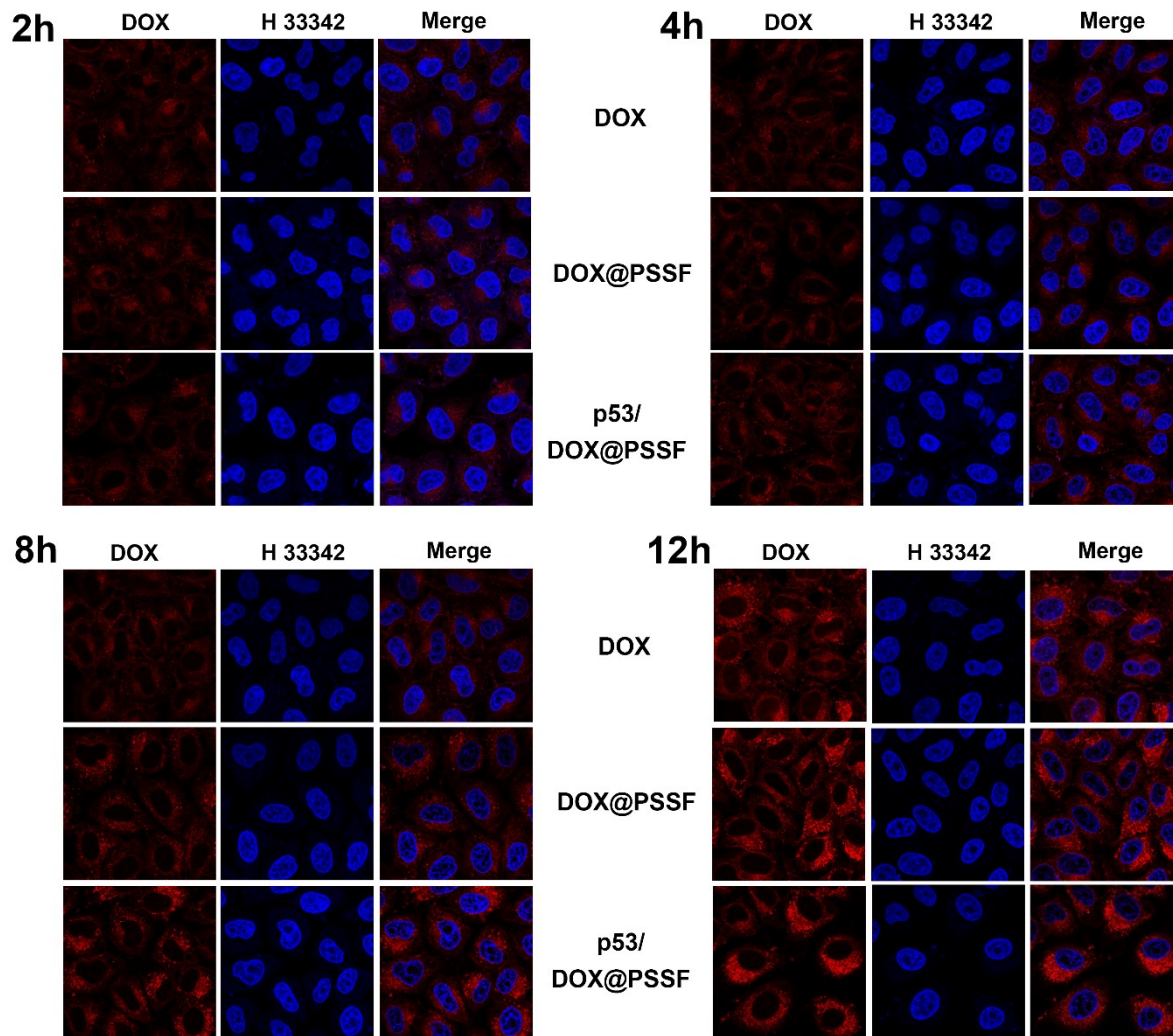


Figure S13. Cellular uptake of DOX at 2, 4, 8 and 12 h under different treatments (DOX: 2 $\mu\text{g/mL}$, p53: 1.85 $\mu\text{g/mL}$).

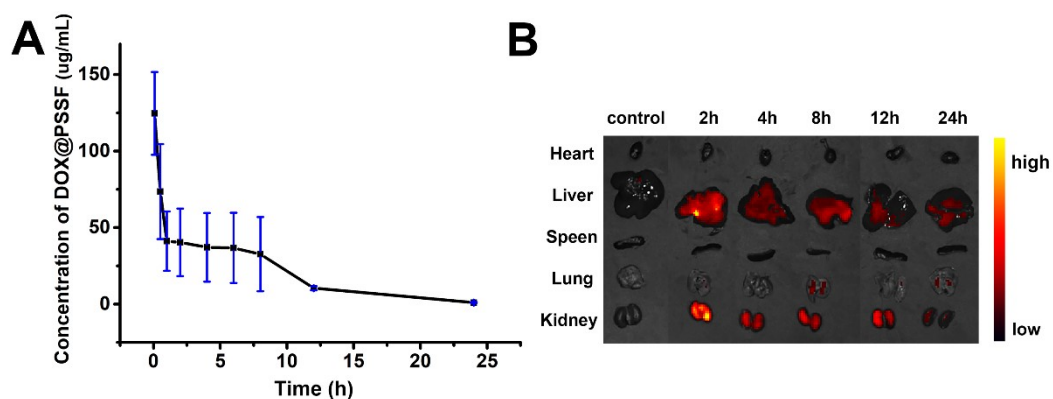


Figure S14. After mice tail vein injection, the blood retention (A) and bio-distribution of DOX@PSSF in major organs (B) at various time points.