

Electronic Supplementary Information

Enzyme-instructed self-assembly leads to activation of optical properties for selective fluorescence detection and photodynamic ablation of cancer cells

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Figures

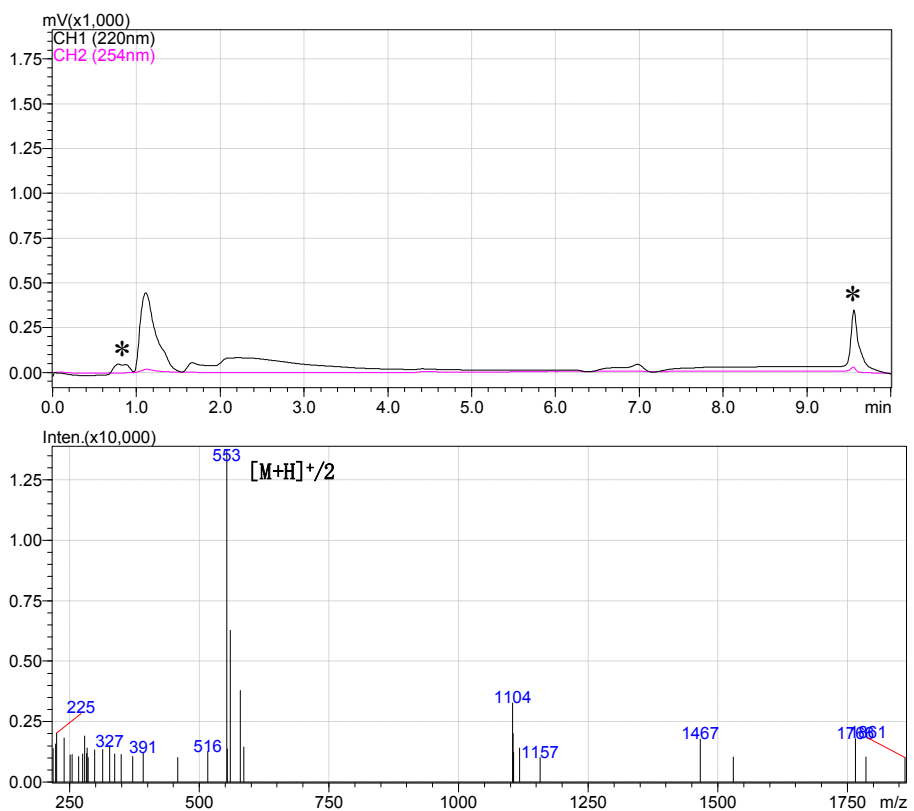


Fig. S1 LC-MS spectra of compound 2 (the stars represent systemic peaks).

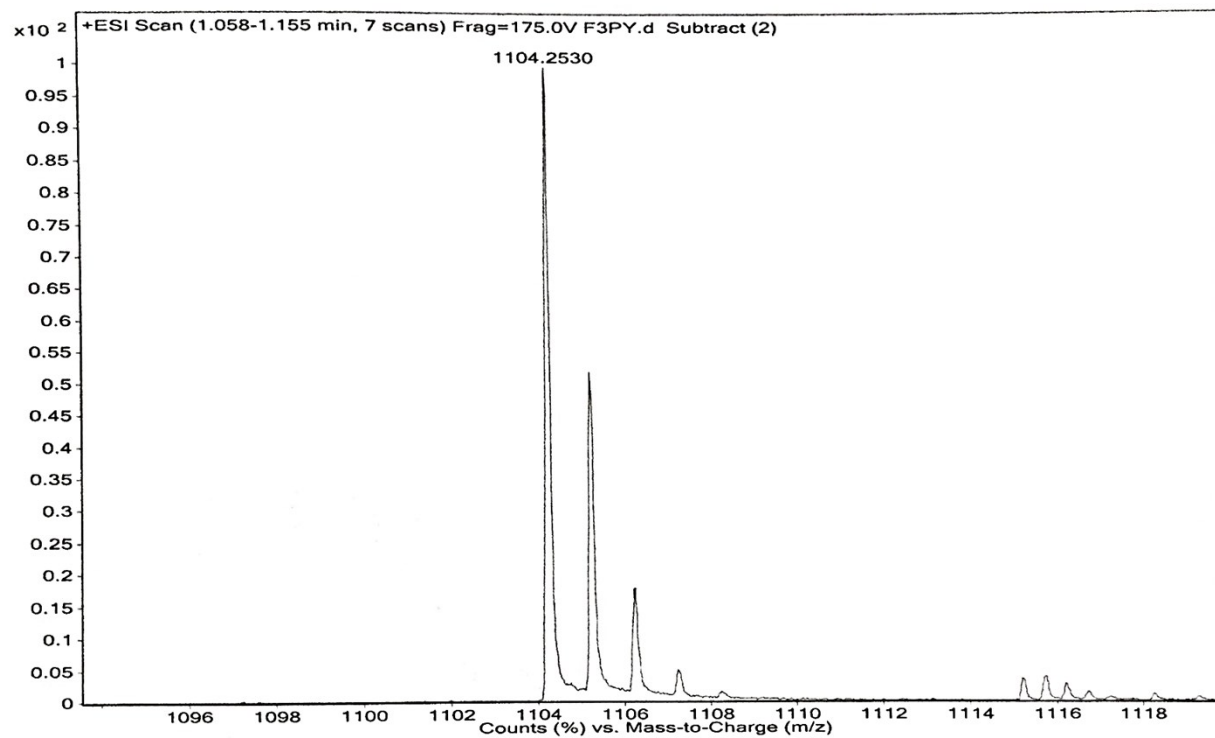


Fig. S3 HR-MS spectrum of compound **2**.

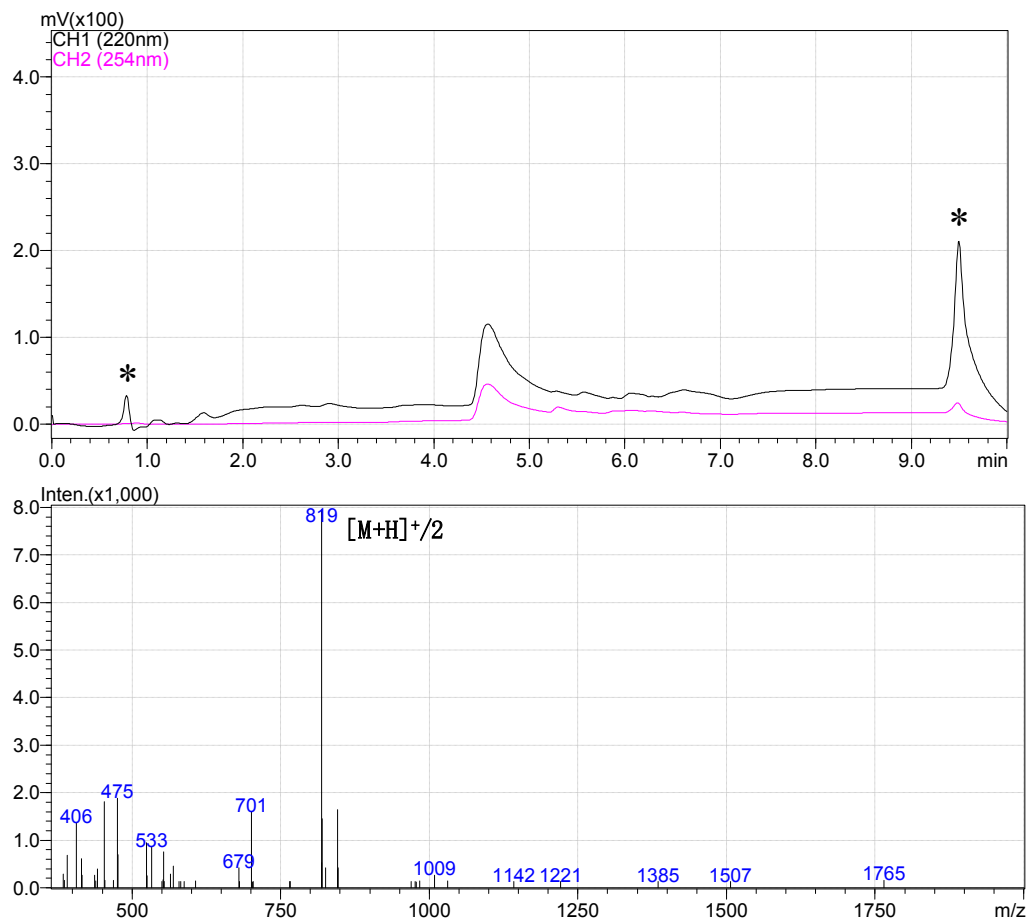


Fig. S4 LC spectra of TPE-Py-FpYGpYGpY (the stars represent systemic peaks).

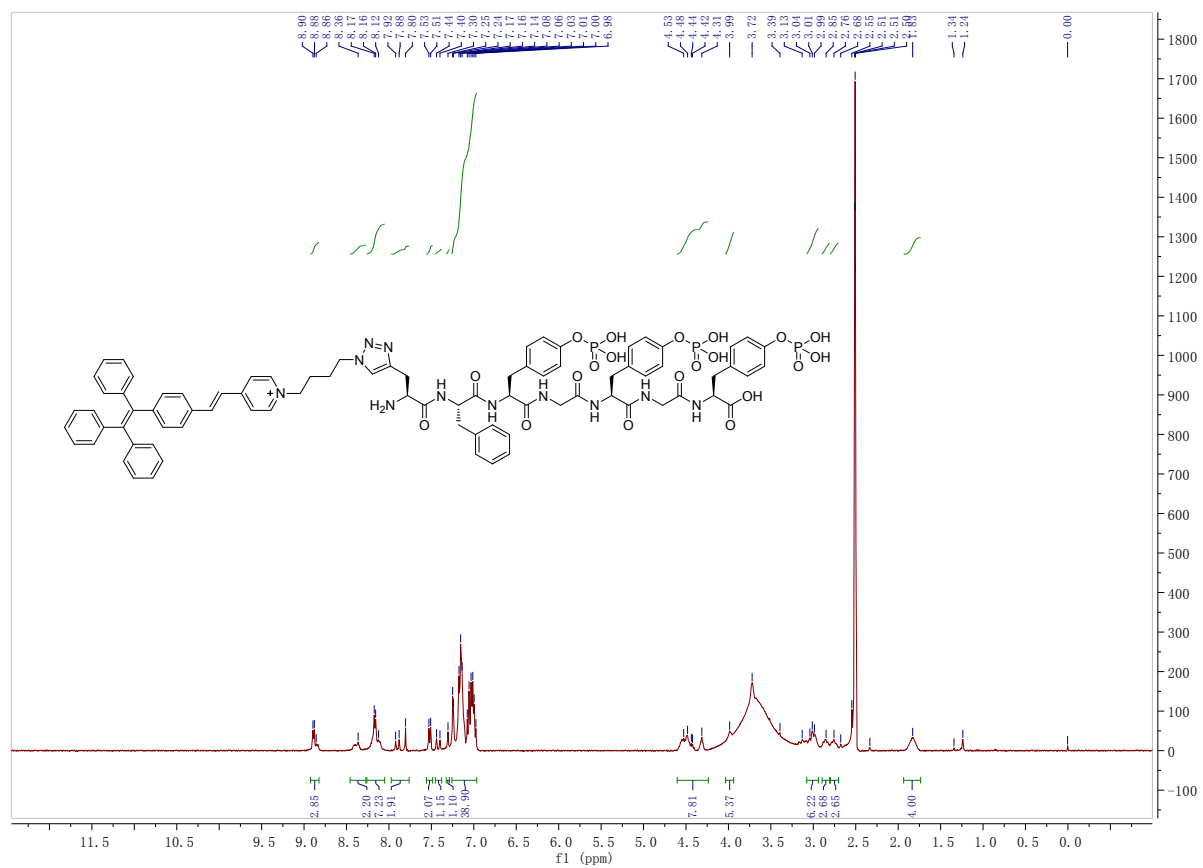


Fig. S5 ¹H NMR spectrum of TPE-Py-FpYGpYGpY in DMSO-*d*₆.

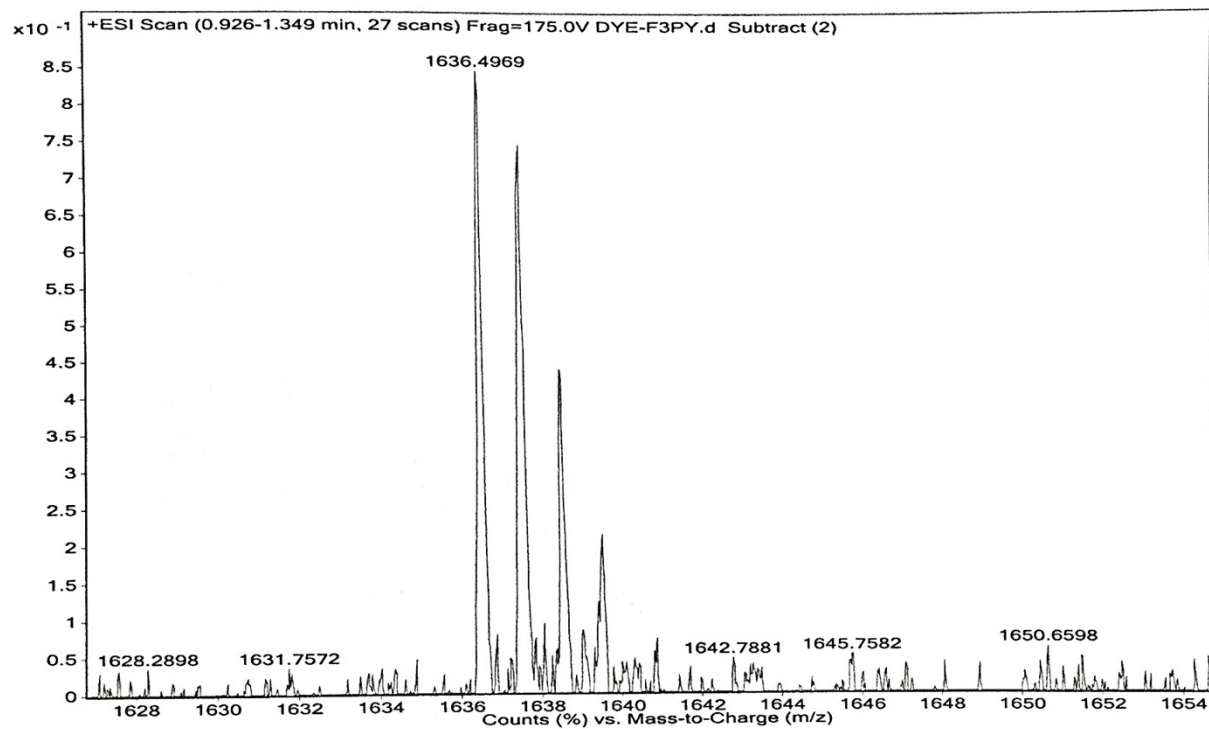


Fig. S6 HR-MS spectrum of TPE-Py-FpYGpYGpY.

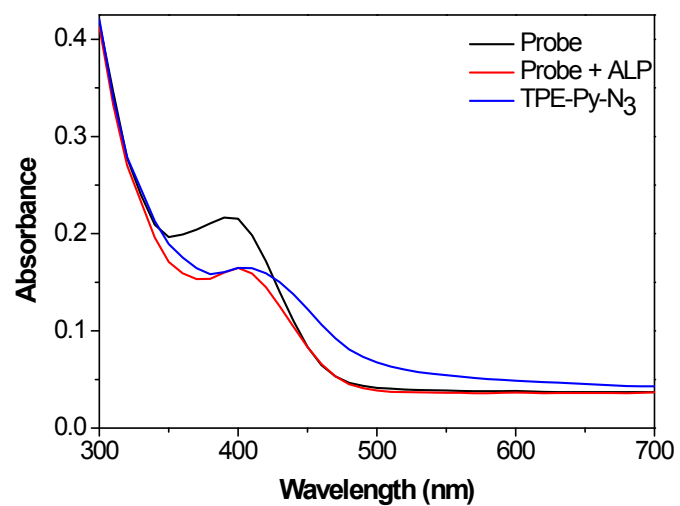


Fig. S7 UV-vis absorption spectra of different samples as indicated in PBS buffer.

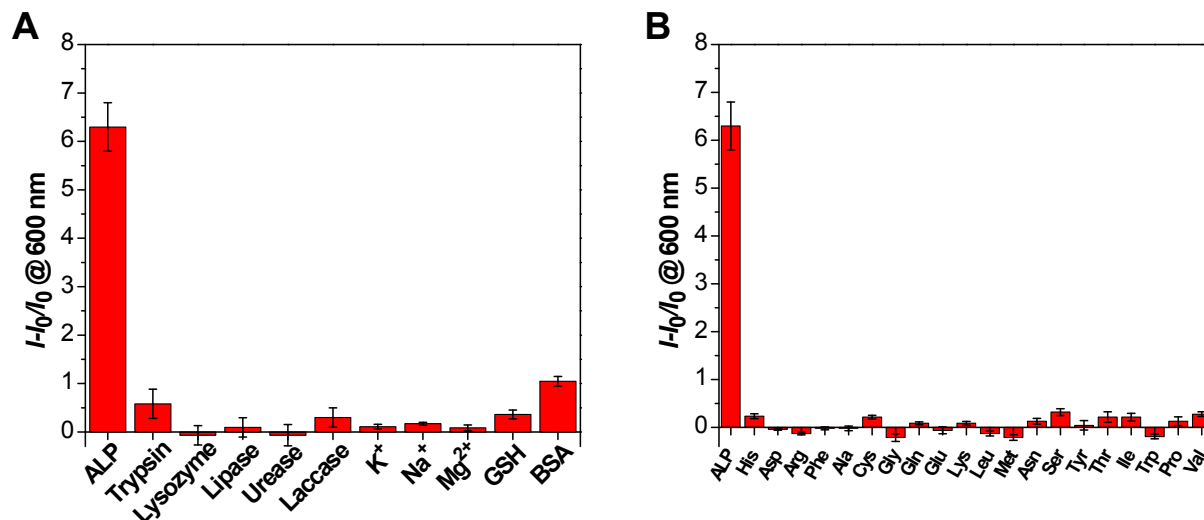


Fig. S8 Plot of $(I-I_0)/I_0$ versus (A) different enzymes, important ions and biomolecules as well as (B) various amino acids. I and I_0 are the PL intensities at 600 nm of TPE-Py-FpYGpYGpY (10 μ M) with and without treatment at 37 °C for 30 min. 1 U mL⁻¹ for ALP, lysozyme, lipase, and urease; 100 μ g mL⁻¹ for trypsin and laccase; 1 mM for K^+ , Na^+ , Mg^{2+} , GSH, and amino acids.

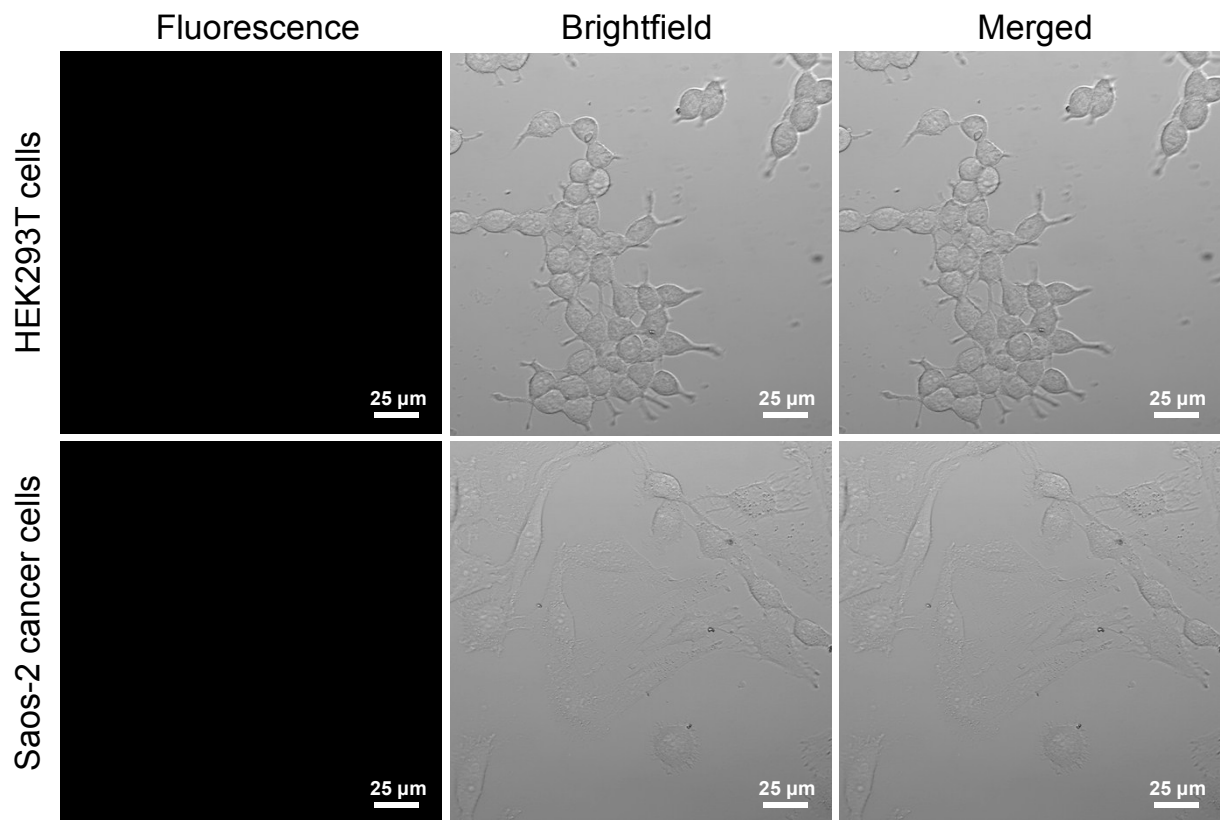


Fig. S9 CLSM images of live HEK293T cells and Saos-2 cancer cells without any probe incubation.

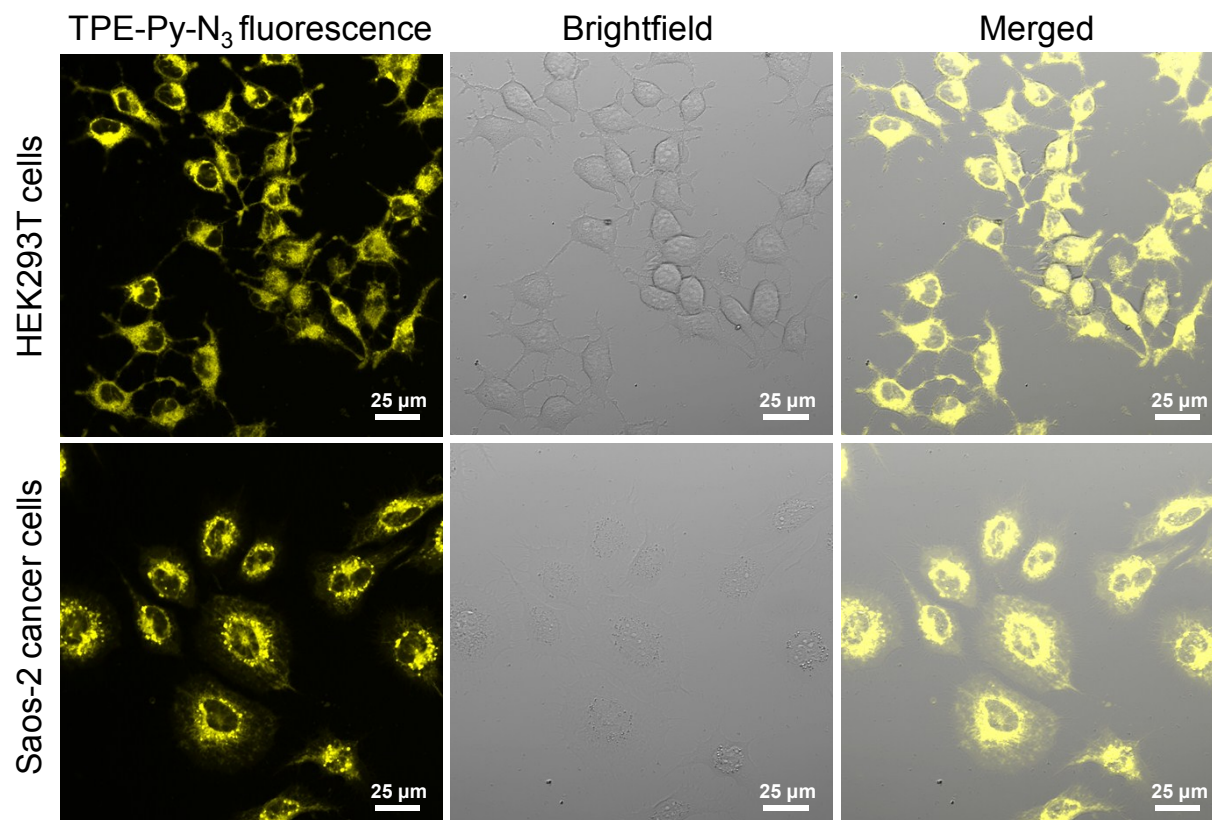


Fig. S10 CLSM images of live HEK293T cells and Saos-2 cancer cells incubated with TPE-Py-N₃ (5 μM) at 37 °C for 2 h.

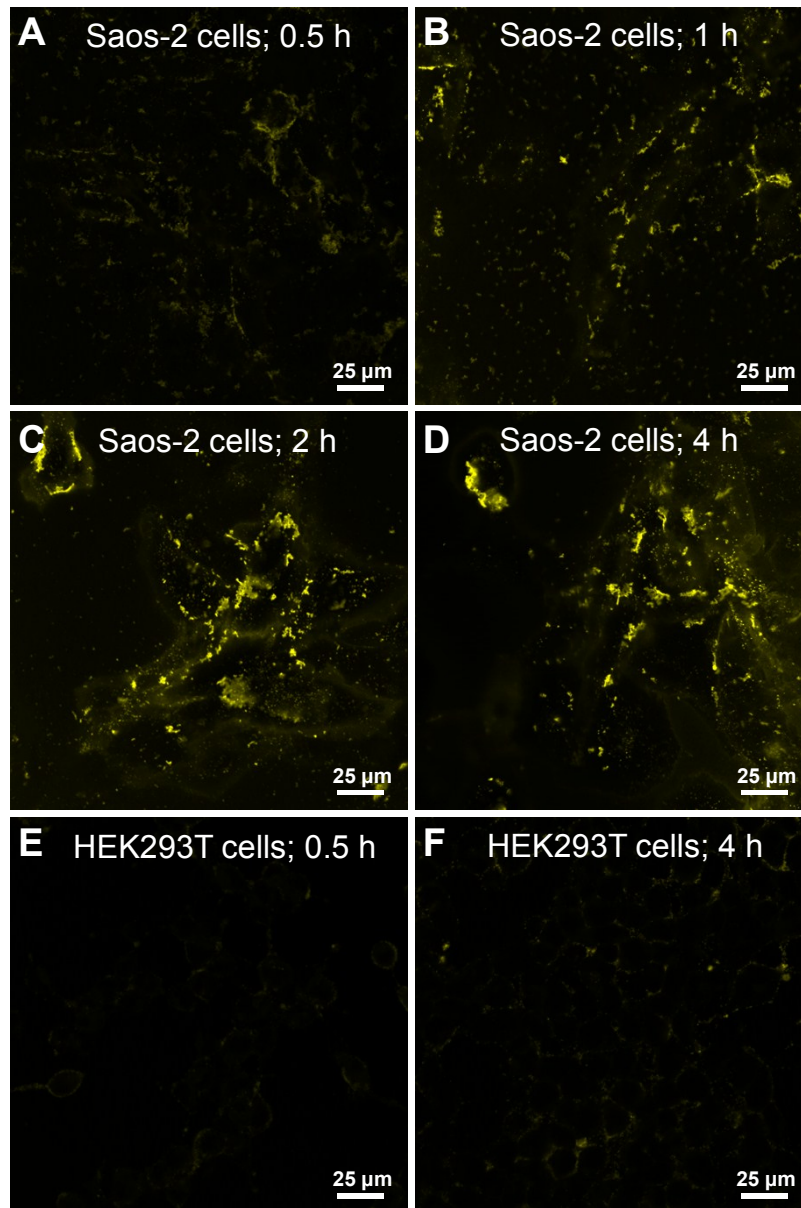


Fig. S11 CLSM images of Saos-2 cancer cells incubated with TPE-Py-FpYGpYGpY (10 μM) at 37 °C for (A) 0.5 h, (B) 1 h, (C) 2 h, and (D) 4 h. CLSM images of HEK293T cells incubated with TPE-Py-FpYGpYGpY (10 μM) at 37 °C for (E) 0.5 h and (F) 4 h.

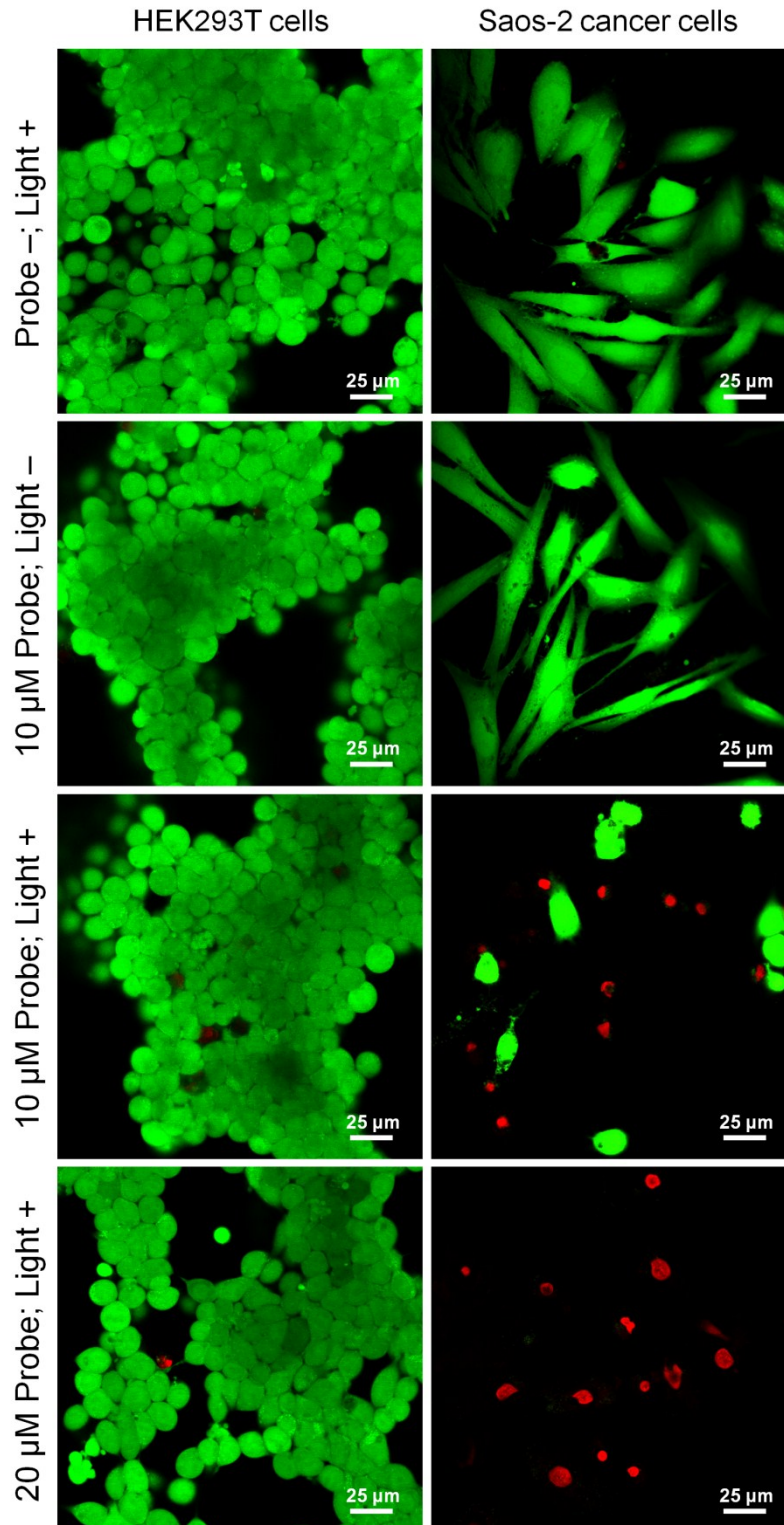


Fig. S12 Live/dead staining with CLSM imaging verifies the PDT efficacy of the probe on HEK293T cells and Saos-2 cancer cells.