Support Information

Linear and High-Molecular-Weight Poly-porphyrins for

Efficient Photodynamic Therapy

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



Figure S2. (A) ¹H NMR spectrum of compound 2 in CDCl₃. (B) Mass spectrum of compound 2. Obsered $[M+H]^+ = 889.6$.



Fig

ure S3. (A) ¹H NMR spectrum of **compound 4b** in CDCl₃. (B) ¹H NMR spectrum of **compound 4** in CDCl₃. (C) ¹³C NMR spectrum of **compound 4**.



Figure S4. HRMS spectrum of compound 4. Observed $[M+Na]^+ = .323.1115$ and Observed $[M+H]^+ = .301.1293$.



Figure S5. (A) ¹H NMR spectrum of **compound 3** in CDCl₃. (B) MALDI-TOF-MS of **compound 3**. Obsered $[M+H]^+ = 952.4$.



Figure S6. ¹H NMR spectrum of **pP** in CDCl₃.



Figure S7. ¹H NMR spectrum of **pP-PEG** in *d*-DMSO.



Figure S8. ¹H NMR spectrum of AZ-amine in *d*-DMSO.



Figure S9. ¹H NMR spectrum of **pP-PEG-AZ** in *d*-DMSO.





Figure S10. GPC curves of pP, pP-PEG and pP-PEG-AZ.



Figure S11. Size distribution of pP-PEG-AZ NPs formed by two methods.



Figure S12 Particle sizes of pP-PEG-AZ NPs for diffient time incubation.



Figure S13. SEM image of pP-PEG-AZ NPs (\overline{A} and \overline{B}) and pP-PEG NPs (\overline{C} and \overline{D}).Bar = 500 nm.



Figure S14. (A) (A) UV-vis spectra of the monomer 2, pP (no Zn), and pP-PEG-AZ NPs. (B) UV-vis absorbance spectra of monomer 2 (No Zn) and monomer A (Zn coordinated). (C) Fluorescence emission spectrum of the monomer 2 and pP (λ_e = 420 nm). Solvent: DMSO. Concentration: porphyrin moiety of 7.5*10⁻⁶M.



Figure S15 Intracellular internalization level of the monomer, pP, pP-PEG NPs, and pP-PEG-AZ NPs in MCF-7 cells. Incubation time: 12 h. n = 3.



Figure S16. H&E staining images of normal tissues of the mice after different treatments.