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Electronic Supplementary Information

Glutathione- and Light-Controlled Generation of Singlet Oxygen for Triggering Drug

Release in Mesoporous Silica Nanoparticles

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Fig. S1 TEM images of (a,c) **Pc@MSN** and (b,d) **Pc/Dox@MSN**. Scale bars denote 200 nm (a,b) or 50 nm (c,d).



Fig. S2 Hydrodynamic diameter (D_h) distribution of **Pc@MSN** (black) and **Pc/Dox@MSN** (red) in water as determined by DLS.



Fig. S3 Change in fluorescence spectrum ($\lambda_{ex} = 610 \text{ nm}$) of **Pc/Dox@MSN** ([ZnPc] = 0.5 μ M) upon incubation with (a) 5 mM or (b) 5 μ M of GSH in PBS with 0.5% Tween 80 monitored over 30 min.



Fig. S4 (a) Fluorescence spectra ($\lambda_{ex} = 485 \text{ nm}$) of Pc/Dox@MSN (0.331 mg MSN mL⁻¹) at different irradiation time after being incubated with 5 mM of GSH in DMF for 30 min. (b) Fluorescence spectra of the supernatant of the centrifuged samples after various irradiation time ($\lambda > 610 \text{ nm}$, fluence rate = 40 mW cm⁻²).



Fig. S5 Changes in fluorescence spectra ($\lambda_{ex} = 485 \text{ nm}$) of the dialysates of **Pc/Dox@MSN** in PBS with 0.5% Tween 80 after incubation with (a-c) 5 mM or (d,e) 5 μ M of GSH for 30 min (a,c,d) with or (b,e) without light irradiation ($\lambda > 610 \text{ nm}$, fluence rate = 40 mW cm⁻²) for 20 s, and then under ambient conditions for a period of 24 h. The control with (c) pre-incubation of





Fig. S6 Cytotoxic effects of Pc@MSN and Pc/Dox@MSN on HepG2 cells after being incubated for (a) 24 h, (b) 12 h, or (c, d) 6 h in the absence or presence of light for (a - c) 10 s or (d) 30 s ($\lambda > 610$ nm, fluence rate = 40 mW cm⁻²). Data are expressed as the mean ± standard deviation, each performed in quadruplicate.









Fig. S8 (a) 1 H and (b) $^{13}C{^{1}H}$ NMR spectra of 4 in CDCl₃.





Fig. S9 (a) ¹H and (b) ¹³C{¹H} NMR spectra of 5 in CD_2Cl_2 with a trace amount of CD_3OD .