Supporting Information

A ZnO-gated porphyrinic metal-organic framework-based drug delivery system for targeted bimodal cancer therapy

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1. Characterization of porous structure of porMOF



Figure S1. HR-TEM of porMOF



2. pH-dependent release of DOX from the carriers without or with ZnO encapsulation.

Figure S2. Time-dependent fluorescence spectra of DOX released from DOX@porMOF or ZnOgated DOX@porMOF system under different pH conditions. (a) DOX@porMOF at pH 5.0; (b) ZnO-gated DOX@porMOF at pH 5.0; (c) DOX@porMOF at pH 7.4; (d) ZnO-gated DOX@porMOF at pH 7.4. $\lambda_{ex} = 488$ nm.

3. DOX fluorescence measurements in selected cells



c Cell	Fluorescence intensity (a.u.)		ి ²⁰⁰ ⊤	_1	
UCCII	Figure a	Figure b	sity.	a	
1	14.93	24.86	-150 - 11 E		
2	15.26	26.29	. nce	_	
3	15.20	21.25	93 100- S		
4	15.53	26.52	nore		
5	13.10	26.46	9 - 05		
6	15.49	23.72	lati'		
Mean+SD	14.92 ± 0.84	24.85±1.90	∣ ∞ ⁰⊥	without AS1411	with AS1411

Figure S3. DOX fluorescence measurements in selected cells. (a, b) Cell selection from the images of HeLa cells treated with 80 μ g/mL (a) ZnO-gated DOX@porMOF or (b) ZnO-gated DOX@porMOF-AS1411for 4 h. (c) Fluorescence intensities measured from the selected cells. (d) Histogram of the relative average fluorescence intensity.



4. Confocal fluorescence microscopy images of HeLa and NIH3T3 cells

Figure S4. Confocal fluorescence microscopy images of HeLa and NIH3T3 cells after incubating with ZnO-gated DOX@porMOF-AS1411 for different time.