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Supplementary information

Berberine mediated fluorescent gold nanoclusters in biomimetic erythrocyte ghosts as a nanocarrier for enhanced photodynamic treatment

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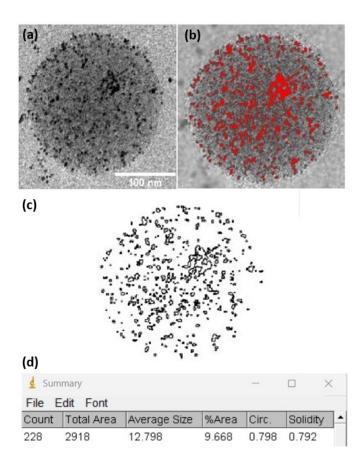


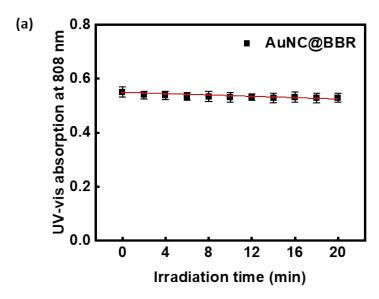
Figure S1 Characterization of AuNC@BBR nanocomposite: (a) HR-TEM image of AuNC@BBR complex. Scale bar represents 100 nm; Quantification with ImageJ (b) thresholding, (c) edge detection, and (d) summary of the quantification.

Journal Name ARTICLE

 ${\it Table S1 A quantification summary for multiple AuNC@BBR complex using ImageJ Software}$

Slice of AuNC@BBR TEM image	Average Size	%Area	Mean	Circ.	Solidity
1	17.574	15.116	65.194	0.726	0.753
2	16.752	16.371	36.346	0.711	0.746
3	19.222	14.074	52.772	0.725	0.756
4	10.762	10.098	47.723	0.846	0.798
5	16.875	17.352	52.96	0.849	0.845
6	18.946	18.523	76.592	0.867	0.883
7	15.527	10.651	79.338	0.891	0.911

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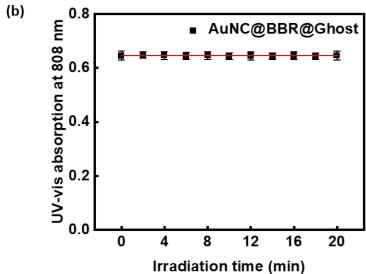


Figure S2 Photostability of (a) AuNC@BBR; and (b) AuNC@BBR@Ghost under 808 nm laser (0.75 W/cm²) irradiation within 20 min by UV-visible absorbance spectra

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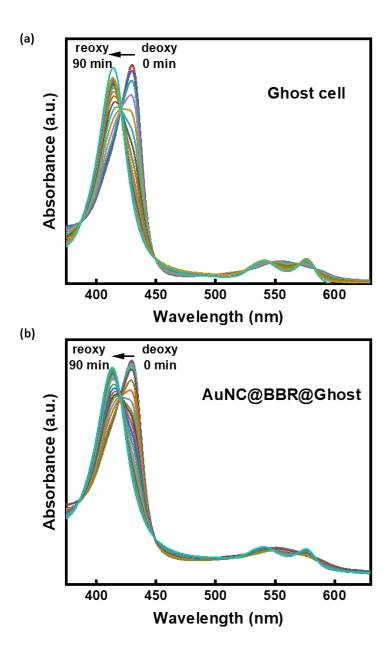


Figure S3 Time-dependent UV-Vis absorbance spectra of (a) Ghost; and (b) AuNC@BBR@Ghost in the deoxygenated and reoxygenated conditions.

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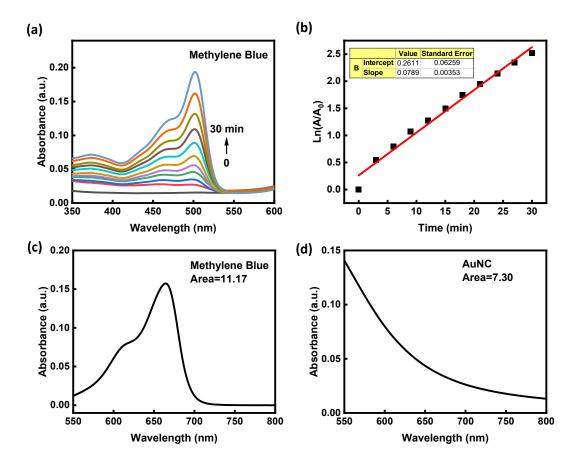


Figure S4 (a) Time-dependent absorbance of DCF at 502 nm caused by methylene blue (MB) under 808 nm laser irradiation; (b) The rate constants for DCF absorbance in the presence of MB; (c & d) The absorption peak area of MB and AuNC respectively.

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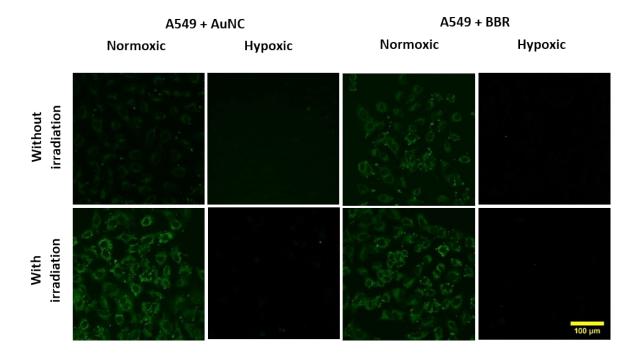


Figure S5 Identification of reactive oxygen species generation in A549 cell line using oxidized DCFDA, observed by fluorescence microscopy (Excitation: 488 nm; Emission: 520-550 nm; Objective: 20x) Scale bar represents 50 μ m.

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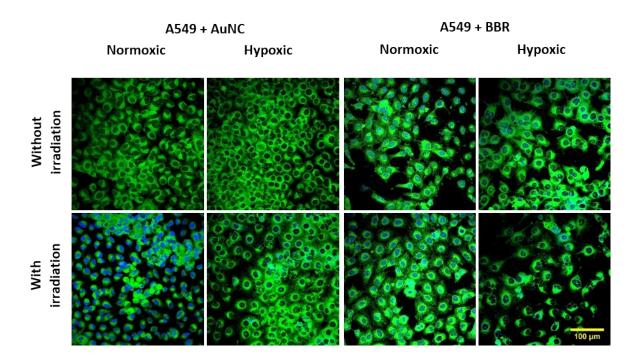


Figure S6 Discrimination of live and dead A549 normoxia and hypoxia cell line using violet fluorescent and DiOC5 staining, observed by confocal fluorescence microscopy (Objective: 20x) Scale bar represents $100 \mu m$.