

Supporting information

Photothermal theranostics with glutathione depletion and enhanced reactive oxygen species generation for efficient antibacterial treatment

Yuelan Wu^{a,b}, Xiaoxue Liu^{b,c}, Xiaoyu Zhang^{b,c}, Shuping Zhang^{c,d}, Panhong Niu^{d*},

Hua Gao^{a,b,c*}

a. Qingdao University, Qingdao, Shandong 266071, P. R. China

b. Eye Institute of Shandong First Medical University, State Key Laboratory Cultivation Base, Shandong Provincial Key Laboratory of Ophthalmology, Qingdao, Shandong 250071, P. R. China.

c. Medical Science and Technology Innovation Center, Shandong First Medical University & Shandong Academy of Medical Sciences, Jinan, Shandong 250117, P. R. China.

d. Biomedical Sciences College & Shandong Medicinal Biotechnology Centre, Shandong First Medical University & Shandong Academy of Medical Sciences, Jinan, 250117, P. R. China.

* Correspondence:

hgao@sdfmu.edu.cn (Hua Gao) and niupanhong@sdfmu.edu.cn (Panhong Niu)

Table S1. The correlation parameters of photothermal conversion ability of PDA-FDM-23.

Parameter	T_{\max}	T_{suur}	A_{808}	m	c	τ_s	η
Value	55.8	23.3	1.982	0.5g	4.2 J/g	266.364	17.26%

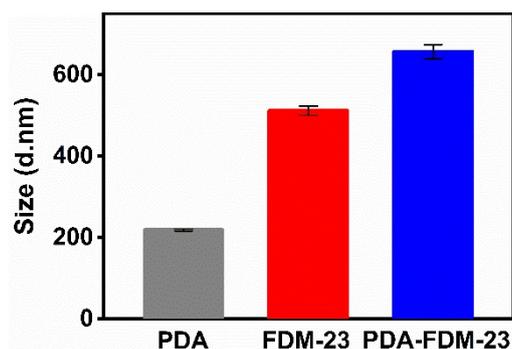


Fig. S1 DLS analysis of PDA, FDM-23, and PDA-FDM-23.

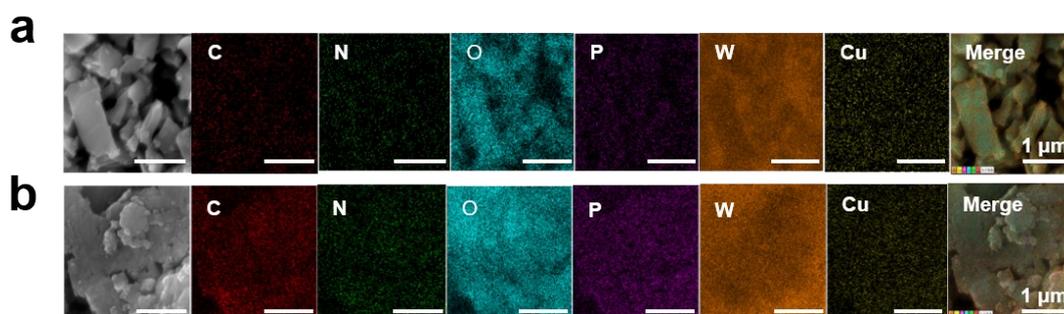


Fig. S2 EDS mapping images of FDM-23 (a) and PDA-FDM-23 (b).

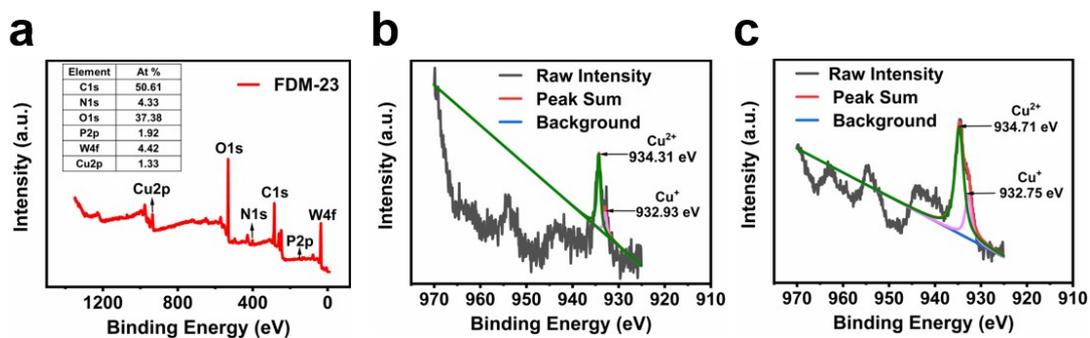


Fig. S3 (a) XPS survey spectra analysis of FDM-23. **(b)** Cu2p core of PDA-FDM-23 level spectra. **(c)** Cu2p core of FDM-23 level spectra.

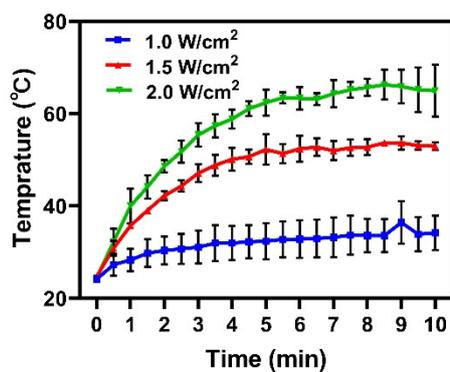


Fig. S4 Heating curves of 100 µg/mL PDA-FDM-23 under laser irradiation with different powers (1.0 W/cm², 1.5 W/cm² and 2.0 W/cm²).

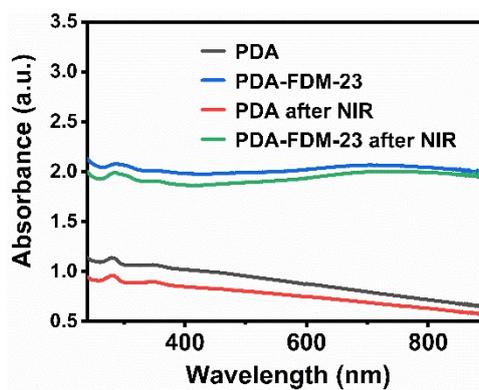


Fig. S5 The UV-vis absorption spectra of PDA-FDM-23 composites before and after laser irradiation (1.5 W/cm², 10 min).

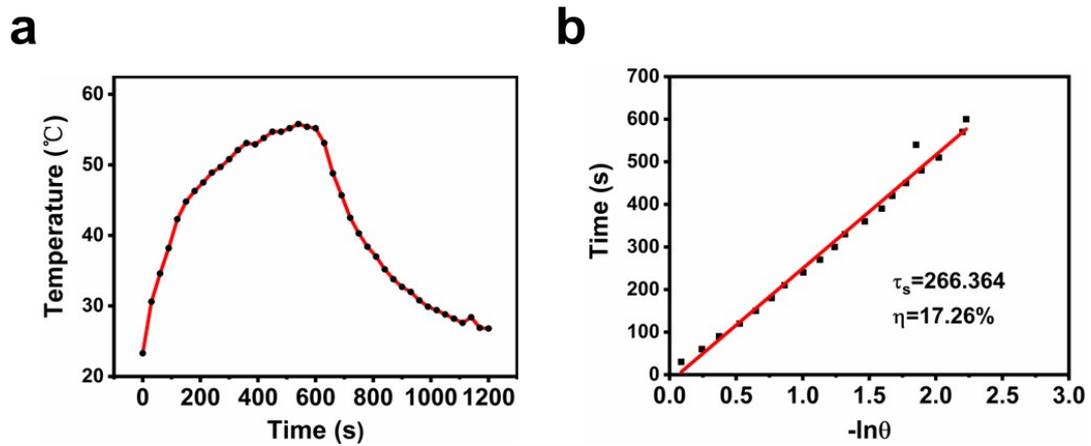


Fig. S6 (a) Heating and cooling curves of PDA-FDM-23 under laser irradiation (808 nm, 1.5 W/cm²). (b) Linear time data versus $-\ln\theta$ obtained from the cooling time.

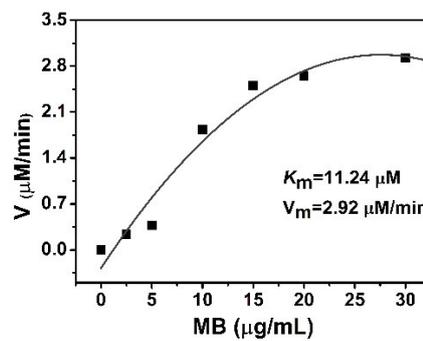


Fig. S7 The Michaelis–Menten curve for PDA-FDM-23.

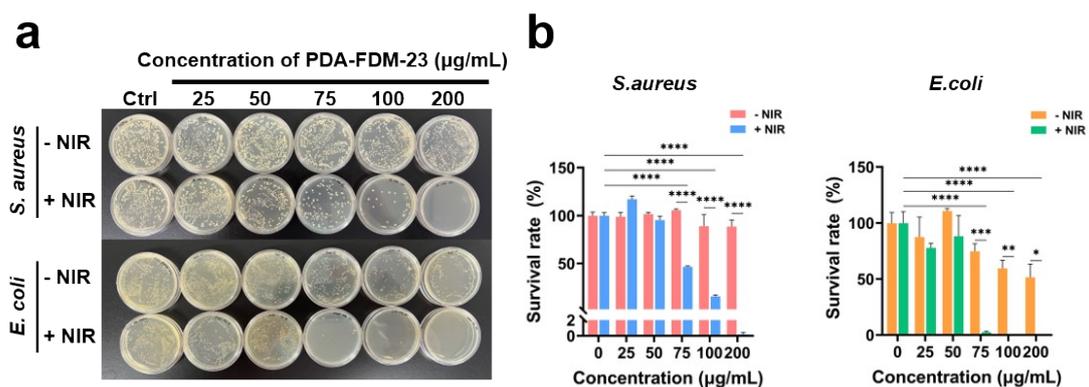


Fig. S8 (a) Plate photographs of *S. aureus* and *E. coli* treated with different concentrations of PDA-FDM-23 in the presence and absence of light (1.5 W/cm², 20 minutes). (b) Relative bacterial survival rates of *S. aureus* and *E. coli* based on (a). Data

were represented as means \pm SEM (n = 3). ns: no significant difference, **** P < 0.0001, *** P < 0.001, ** P < 0.01.

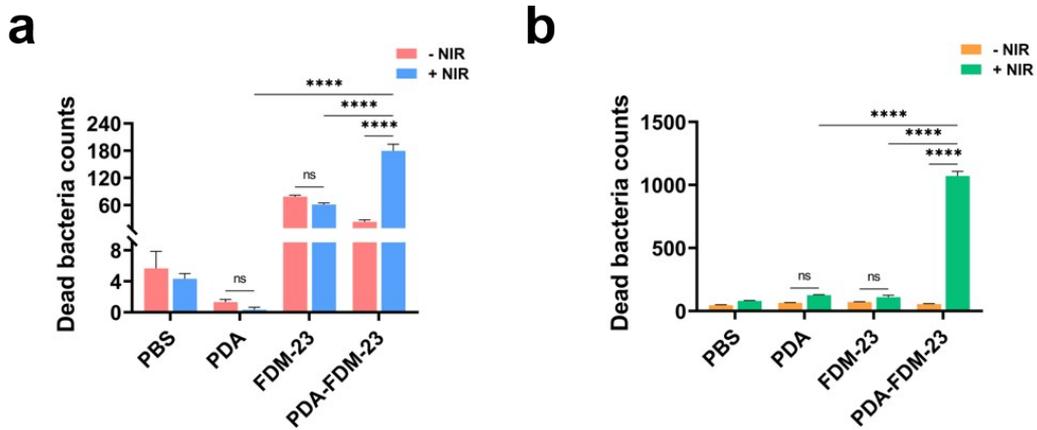


Fig. S9 The dead bacteria count of PI-labeled *S. aureus* (a) and *E. coli* (b). Data were represented as means \pm SEM (n = 3). ns: no significant difference. **** P < 0.0001.

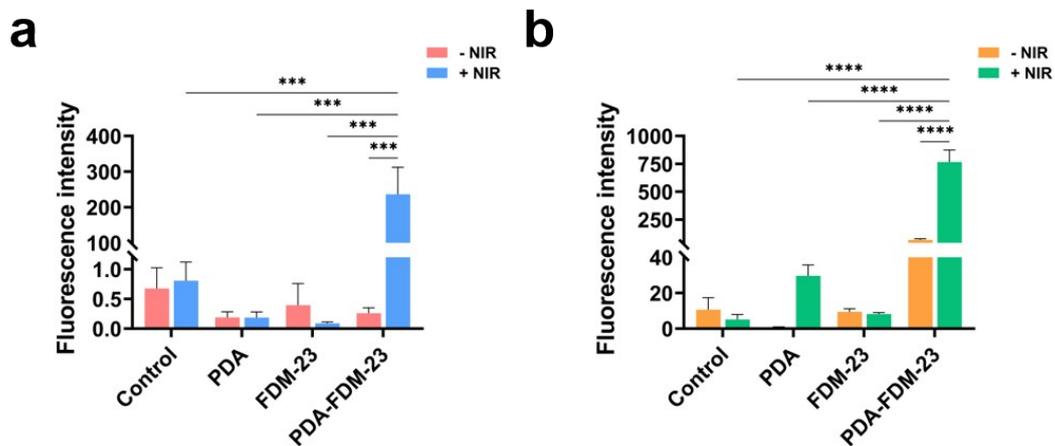


Fig. S10 Relative quantification of ROS fluorescence intensity after incubation with *S. aureus* (a) and *E. coli* (b) in each group. Data were represented as means \pm SEM (n = 3). *** P < 0.001, **** P < 0.0001.