## Graphene Oxide Composite for Selective Recognition, Capturing, Photothermal Killing of Bacteria Over Mammalian Cells

## Gang Ma<sup>1</sup>, Junjie Qi<sup>1,\*</sup>, Qifan Cui<sup>2</sup>, Xueying Bao<sup>2</sup>, Dong Gao<sup>2</sup> and Chengfen Xing<sup>2,\*</sup>

- <sup>1</sup> National-Local Joint Engineering Laboratory for Energy Conservation in Chemical Process Integration and Resources Utilization, School of Chemical Engineering and Technology, Hebei University of Technology, Tianjin 300131, P.R. China; magang0629@163.com (G.M.); qijunjie@hebut.edu.cn (J.Q.)
- <sup>2</sup> Key Laboratory of Hebei Province for Molecular Biophysics, Institute of Biophysics, Hebei University of Technology, Tianjin 300401, P.R. China; cuiqifan1228@163.com (Q.C.); XueyingBao2017@163.com (X.B.); gaodong@iccas.ac.cn (D.G.); xingc@hebut.edu.cn (C.X.)
- \* Correspondence: xingc@hebut.edu.cn; Tel/Fax: 86-22-60435642 (C.X.); qijunjie@hebut.edu.cn (J.Q.)



Figure S1. Standard curve of GO-PEG-NH2 in PBS.



**Figure S2.** (a) AFM image of GO deposited on mica substrate. (b) The height profile of the AFM image. (c) Hydrodynamic diameter of GO measured by DLS. (d) Hydrodynamic diameter of GO-PEG-NH<sub>2</sub> measured by DLS. (e) Photograph of GO-PEG-NH<sub>2</sub> dispersed in different culture media for 24 h. (f) SEM image of GO-PEG-NH<sub>2</sub>.



Figure S3. CLSM images of (a, b) E. coli, (c, d) CCRF-CEM.



**Figure S4.** Gel imaging of *E. coli* and *S. aureus* colonies. (**a**, **e**) Plate photographs of the *E. coli* and *S. aureus* LB agar plate without GO-PEG-NH<sup>2</sup> under dark. (**b**, **f**) Plate photographs for *E. coli* and *S. aureus* LB agar plate supplemented with GO-PEG-NH<sup>2</sup> (50 µg/mL) under dark. (**c**, **g**) Plate photographs for *E. coli* and *S. aureus* LB agar plate upon 808 nm laser irradiation (1.5 W/cm<sup>2</sup> for 5 min). (**d**, **h**) Plate photographs for *E. coli* and *S. aureus* LB agar plate supplemented with GO-PEG-NH<sup>2</sup> upon 808 nm laser irradiation.

**Table S1.** Evaluation of *E. coli* colonies by plate counting method (The power density of the 808 nm laser is 1.5 W/cm<sup>2</sup>, the irradiation time is 5 min, and the concentration of GO-PEG-NH<sub>2</sub> is 50 µg/mL).

<b>Experimental Condition</b>	Number of Colonies
Control/Non-Laser	$734 \pm 1$
Control/Laser	$676 \pm 23$
GO-PEG-NH <sub>2</sub> /Non-Laser	$611 \pm 15$
GO-PEG-NH <sub>2</sub> /Laser	9 ± 7

**Table S2.** Evaluation of *S. aureus* colonies by plate counting method (The power density of the 808 nm laser is 1.5 W/cm<sup>2</sup>, the irradiation time is 5 min, and the concentration of GO-PEG-NH<sub>2</sub> is 50  $\mu$ g/mL).

Experimental Condition	Number of Colonies
Control/Non-Laser	776 ± 31
Control/Laser	$767 \pm 34$
GO-PEG-NH2/Non-Laser	$714 \pm 22$
GO-PEG-NH <sub>2</sub> /Laser	$6 \pm 5$

**Table S3.** Evaluation of *S. aureus* and *E. coli* colonies by plate counting method (The power density of the 808 nm laser is  $1.5 \text{ W/cm}^2$ , and the irradiation time is 5 min).

Concentration (µg/mL)	Number of Colonies (E. coli)	Number of Colonies (S.
		aureus)
Control	734 ± 1	776 ± 31
10	$258 \pm 46$	$279 \pm 47$
30	$69 \pm 30$	$66 \pm 35$
50	$9\pm7$	$6 \pm 5$
70	$3 \pm 2$	$1 \pm 1$