## Supplementary

## White Light Powered Antimicrobial Nanoagents for Triple Photothermal, Chemodynamic and Photodynamic Based Sterilization

Hua Tian<sup>a,c,‡</sup>, Houjuan Zhu<sup>b,‡,\*</sup>, Yuling Xue<sup>a</sup>, Maonan Wang<sup>d</sup>, Kuoran Xing<sup>a</sup>, Zibiao

Li<sup>b</sup>, Xian Jun Loh<sup>b</sup>, Enyi Ye<sup>b</sup>, Xianguang Ding<sup>e</sup>, Bang Lin Li<sup>f</sup>, Xueqiong Yin<sup>c,\*</sup>, David

Tai Leong<sup>a,</sup>\*

<sup>a</sup>Department of Chemical and Biomolecular Engineering, National University of Singapore, Singapore 117585, Singapore.

<sup>b</sup>Institute of Materials Research and Engineering (IMRE), Agency for Science, Technology and Research (A\*STAR), 2 Fusionopolis Way, Innovis #08-03, Singapore 138634, Republic of Singapore.

<sup>c</sup>Hainan Provincial Fine Chemical Engineering Research Centre, Hainan University, Haikou, Hainan, 570228, P.R. China.

<sup>d</sup>Department of Pathology, Xiangya Hospital, School of Basic Medical Sciences, Central South University, Changsha, China.

<sup>e</sup>Key Laboratory for Organic Electronics and Information Displays & Jiangsu Key Laboratory for Biosensors, Nanjing University of Posts & Telecommunications, Nanjing, 210023 China.

<sup>f</sup>School of Chemistry and Chemical Engineering, Southwest University, Chongqing 400715, P. R. China.

\*Corresponding author. E-mail: <u>cheltwd@nus.edu.sg;</u> <u>zhu\_houjuan@imre.a-</u> <u>star.edu.sg;</u> <u>yxq88@hotmail.com</u>

‡Equal contributions



Fig. S1 Nanodevice characterization. TEM images of (a)  $CoS_x$  QDs, (b) CDs, (c) GCDCO, (d) Particle size of  $CoS_x$  QDs, (e) XRD pattern of CDs.



**Fig. S2** Photothermal evaluation of  $CoS_x$  QDs irradiated by sunlight. (a) Photothermal heating curves of  $CoS_x$  QDs solutions with various concentrations irradiated by simulated sunlight. (b) Infrared thermal images of  $CoS_x$  QDs solutions at the indicated concentrations and irradiated for different durations. (c) Photothermal stability of  $CoS_x$  QDs over 5 on/off cycles of simulated sunlight irradiation (200-2500 nm, 2335)



**Fig. S3** (a)  $A/A_0$  of different DAB reaction systems at different time without  $H_2O_2$ . (b) Degradation of MB of different reaction systems without  $H_2O_2$  at different time.



**Fig. S4** ROS production from GCDCO. (a) Time-dependent fluorescence changes without cold-light irradiation (400-800 nm, 91 mW/m<sup>2</sup>). (b) The relative intensity of different reaction systems with cold-light irradiation at 10 min.



Fig. S5. Cell viability of HCE cells (a) and LO2 cells (c) incubated with various GCDCO at different concentration (0-53.33  $\mu$ g/mL) for 6 h. (b) Cell viability of HCE cells treated with GOx, CDs, CoS<sub>x</sub> QDs and GCDCO at concentrations of 0.83, 3.33 and 13.33  $\mu$ g/mL. (d) Cell viability of LO2 cells treated with GOx, CDs, CoS<sub>x</sub> QDs and GCDCO at concentrations of 0.83, 3.33 and 13.33  $\mu$ g/mL in absence of glucose.



**Fig. S6.** Minimum inhibitory concentration (MIC) for different concentrations of GCDCO nanoagents for 12 h.



**Fig. S7**. Antibacterial characterization of *E. coli in vitro*. (a) Photographs of survived bacterial colonies of *E. coli* without sunlight and with sunlight treatment in the absence and presence of glucose. Corresponding antibacterial activity of GOx (5  $\mu$ g/mL), CDs (5  $\mu$ g/mL), CoS<sub>x</sub> QDs (10  $\mu$ g/mL) and GCDCO (10  $\mu$ g/mL) without sunlight and with sunlight treatment in the absence (b) and presence of glucose (c) measured by OD values of bacteria dispersion. Corresponding antibacterial activity of GOx, CDs, CoS<sub>x</sub> QDs and GCDCO without sunlight and with sunlight treatment in the absence (d) and presence of glucose (e) measured by bacteria counting.



**Fig. S8.** Antibacterial characterization of *E. coli in vitro*. (a) Photographs of survived bacterial colonies of *E. coli* without sunlight and with sunlight treatment in the absence and presence of glucose. Corresponding antibacterial activity of GOx (2.5  $\mu$ g/mL), CDs (2.5  $\mu$ g/mL), CoS<sub>x</sub> QDs (5  $\mu$ g/mL) and GCDCO (5  $\mu$ g/mL) without sunlight and with sunlight treatment in the absence (b) and presence of glucose (c) measured by OD values of bacteria dispersion. Corresponding antibacterial activity of GOx, CDs, CoS<sub>x</sub> QDs and GCDCO without sunlight and with sunlight treatment in the absence (d) and presence of glucose (e) measured by bacteria counting.

Commite	Atomic conc. (%)		
Sample –	S 2p	Co 2p	
GCDCO	1.38	0.47	
CoS <sub>x</sub> QDs	0.81	0.39	

Table S1	Elemental	ana	lysis
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