## **Supporting Information**

# A Novel Full Solar Light Spectrum Responsive Antimicrobial Agent by WS<sub>2</sub> Quantum Dots for Photocatalytic Wound Healing

## Therapy

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### **EXPERIMENTAL SECTION**

### Materials and chemicals

Commercial WS<sub>2</sub> (99.8 %) was obtained from Alfa Aesar and used without further purification. H<sub>2</sub>SO<sub>4</sub> (95.0 %-98.0 %, analytical reagent), methyl orange (MO), Tris(hydroxymethyl)methyl aminomethane THAM (Tris) and Potassium bromide (KBr) were obtained from Chron Chemical Co. Ltd. (Chengdu, China). Rhodamine B (RhB), 2, 2'-azino-bis (3-ethylbenzothiazoline-6-sulfonic acid) diammonium salt (ABTS), 3,3',5,5'-tetramethylbenzidine (TMB), o-phenylenediamine (OPD), 3, 3'diaminobenzidine (DAB) and 5,5'-dithiobis (2-nitrobenzoic acid) (DTNB) were obtained from Aladdin Company (China, Shanghai). hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>, 30 %) was procured from Jinshan Chemical Reagent Co. Ltd. (Chengdu, China). Reduced glutathione (GSH) was sourced from Nanjing Jiancheng Institute of Biological Engineering. Crystal violet staining solution was purchased from Scientific Phygene (Shanghai, China). Acridine orange and Ethidium bromide (AO-EB) staining kit was acquired from Shanghai Yuanye Bio-Technology Co. Ltd. (China). Sprague Dawley rats. Deionized (DI) water was used in the whole process.



**Fig. S1** TEM of WS<sub>2</sub> QDs. (Scale bar = 50 nm)



**Fig. S2** FT-IR spectra of the  $WS_2$  QDs.



**Fig. S3** Raman spectra of the  $WS_2$  QDs.



Fig. S4 FT-IR spectra of WS<sub>2</sub> QDs.



Fig. S5 UV-vis absorption spectra of MO ( $10 \ \mu g \cdot mL^{-1}$ ) with WS<sub>2</sub> QDs (500  $\ \mu g \cdot mL^{-1}$ ) under UV, Nis and NIR light irradiation at different time (0-180 min).



**Fig. S6** UV-vis absorption spectra of RhB (10  $\mu$ g·mL<sup>-1</sup>) with WS<sub>2</sub> QDs (500  $\mu$ g·mL<sup>-1</sup>) under UV, Nis and NIR light irradiation at different time (0-180 min).



Fig. S7 Peroxidase-like activity of reduced  $WS_2$  QDs is dependent on concentrations (a),  $H_2O_2$  (b), temperature (c), and pH (d).  $WS_2$  QDs show an optimal pH of 4.0-5.0 and optimal temperature around 25-30°C. The insets show the fluorescence spectra of the corresponding reduced  $WS_2$  QDs reaction system.



Fig. S8 Photographs of color changes after GSH treatment at different full spectrum light intervals were determined by Ellman's assay in the absence and presence of WS<sub>2</sub> QDs. The concentration of WS<sub>2</sub> QDs was 50  $\mu$ g·mL<sup>-</sup><sup>1</sup>. GSH without WS<sub>2</sub> QDs as a control showed a significant reduction in color after 60 minutes of light.



Fig. S9 Photographs for the color change after GSH oxidation with different concentrations of  $WS_2$  QDs at different time intervals determined by Ellman's assay.



Fig. S10 Fluorescent staining photograph of *E. coli* treated after exposed (I) Control, (II) UV, (III) Vis, (IV) NIR, (V) Full spectrum, (VI) WS<sub>2</sub> QDs  $(50 \ \mu g \cdot mL^{-1})$ , (VII) WS<sub>2</sub> QDs + UV, (VIII) WS<sub>2</sub> QDs + Vis, (IX) WS<sub>2</sub> QDs + NIR, (X) WS<sub>2</sub> QDs + Full spectrum.



Fig. S11 Fluorescent staining photograph of *S. aureus* treated after exposed (I) Control, (II) UV, (III) Vis, (IV) NIR, (V) Full spectrum, (VI) WS<sub>2</sub> QDs (50  $\mu$ g·mL<sup>-1</sup>), (VII) WS<sub>2</sub> QDs + UV, (VIII) WS<sub>2</sub> QDs + Vis, (IX) WS<sub>2</sub> QDs + NIR, (X) WS<sub>2</sub> QDs + Full spectrum.



Fig. S12 Toxicity experiments with different concentrations of  $WS_2$  QDs for 24 h.