Supporting Information

Fluorescent Imaging and Precise Suppression of Bacterial Infection in Chronic Wounds by Porphyrin-Based Metal-Organic Framework Nanorod

Lihua Zhang ^{a, b, c}, Minzhi Ouyang ^d, Yufei Zhang ^c, Huanxiang Wang ^c, Ziyun Huang ^c, Libei He ^c, Yanli Lei ^c, Zhen Zou *^c, Feng Feng *^{a, b}, Ronghua Yang *^{c, e}

^aSchool of Chemistry and Material Science, Shanxi Normal University, Linfen 041004, China

^bCollege of Chemistry and Environmental Engineering, Shanxi Datong University, Datong 037009, China

^cHunan Provincial Key Laboratory of Cytochemistry, School of Chemistry and Food Engineering, Changsha University of Science and Technology, Changsha 410004, China

^dDepartment ultrasound diagnosis, The Second Xiangya Hospital, Central South University, Changsha, 410011, P. R. China

^eCollege of Chemistry and Chemical Engineering, Hunan Normal University, Changsha 410081, China

*Corresponding authors email: kobe474779970@126.com; Yangrh@pku.edu.cn

Lihua Zhang, Minzhi Ouyang and Yufei Zhang contributed equally to this work.

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Figure S1 The FTIR of TCPP and Zn-TCPP nanorods.



Figure S2 (A) Size distribution (B) photographs of Zn-TCPP in various aqueous media incubated

for 30 h at 37°C.



Figure S3 MTT assay of (A) MCF-7 and (B) L929 cells treated with Zn-TCPP nanorods of different concentrations for 24 h. The error bars were the standard deviations from five

independent measures.



Figure S4 Hemolytic tests of Zn-TCPP with different concentrations. Ultrapure water and PBS

were also used for positive and negative control.



Figure S5 Fluorescence images of MCF-7 cells of incubation with different pH. (A) pH=6, (B)

pH=7, (C) pH=8.



Figure S6 (A) Fluorescence images of in vivo model with extend chronic wound, the circle area was inoculated with 50 μ L *S. aureus* of 1×10⁶ CFU mL⁻¹. (B) The measurement results of pH glass electrodes on different areas of the chronic wound.



Figure S7 (A) Fluorescence spectrum and (B) signal-to-back ratio trend of SOSG fluorescent probe treatment with Zn-TCPP nanorods under illumination of near-infrared light ($\lambda ex = 660$ nm).



Figure S8 CLSM of SOSG fluorescent probe treatment with Zn-TCPP nanorods under

illumination of near-infrared light ($\lambda ex = 660$ nm).



Figure S9 Local photodynamic inactivation of Staphylococcus aureus. The red dotted line range is

the illumination range.



Figure S10 H&E and Masson images of the normal tissues on the first day and eighth day.