Supporting Information for

Self-Healing Dynamic Hydrogel Microparticles with Structural Color

for Wound Management

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Supplementary Figures



Fig. S1 H-NMR spectrum of DEX-BA



Fig. S2 H-NMR spectrum of DEX-CA



Fig. S3 Images showing that histidine promotes gelation of the dynamic hydrogel. When the pictures were taken, the tube was changed from the upright state to the inclined state. The dashed lines represent the outline of the sample



Fig. S4 FT IR spectra of DEX-BA, DEX-CA and dynamic hydrogel



Fig. S5 Temperature sensitivity of dynamic hydrogels with different concentrations



Fig. S6 SEM image of DEX-CA/BA dynamic hydrogel. Scale bar is 10 µm



Fig. S7 Real-time rheology test of DEX-CA/BA dynamic hydrogel



Fig. S8 Temperature rising curve of the photothermal hydrogel under different NIR power irradiation



Fig. S9 a-b Images of gelation process of DEX-CA/BA with or without eumenitin and VEGF. **c** Rheology test of the DEX-CA/BA without drugs. **d** Rheology test of the DEX-CA/BA with drugs



Fig. S10 Drug release curve of Rhodamine B labeled-eumenitin and FITC-BSA with or without NIR irradiation



Fig. S11 a-b Images and the estimated hemolysis rate of different treatments including control group, NIR irradiation, and CMPs plus NIR irradiation



Fig. S12 Fluorescent images of NIH 3T3 cells treated with different strategies. Scale bar represents $200 \,\mu$ m.



Fig. S13 a-c Fluorescent images of NIH 3T3 cells treated with different strategies including control group, NIR irradiation, and CMPs plus NIR irradiation. **d** Analysis of relative activity of NIR 3T3 cells in different groups. Scale bars are 50 µm



Fig. S14 Bright field and fluorescent images of CMPs co-culturing with NIH 3T3 cells



Fig. S15 Data analysis of tuber number in different groups



Fig. S16 Live/dead fluorescent images of bacteria with different treatments, including control group and NIR irradiation. Scale bars are 100 μm



Fig. S17 a CMPs applied on the wound. **b-c** Images of CMPs-treated wound and local magnification of structural color microspheres. Scale bars are 5 mm in (**b**) and 500 μ m in (**c**)







Fig. S19 H&E staining images of main organs from rats in different groups. Scale bars are 500 μm



Fig. S20 Blood routine and biochemisity analysis of rats in different groups