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## **Electronic Supplementary Information (ESI)**

for

## A Novel Colorimetric and Near-infrared Fluorescent Probe for Hydrogen Peroxide Imaging in Vitro and in Vivo

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Fig. S1 Absorption spectra of probe DCM-B1 (5  $\mu$ M) before (red line) and after reacting with H<sub>2</sub>O<sub>2</sub> (100  $\mu$ M, blue line).



Fig. S2 Emission spectra of probe DCM-B1 in the presence of different equivalents of  $H_2O_2$  (0, 0.5, 1.0, 2.0, 4.0, 6.0, 8.0, 10.0, 15.0, 20.0 eq, 30 min) excited at 560 nm.



Fig. S3 A linear correlation between emission intensities and concentrations of  $H_2O_2$ .



**Fig. S4** Fluorescence intensity of 5  $\mu$ M DCM-B1 to the testing species in PBS buffer solution (20 mM, 50% DMSO, pH 7.4) at 700 nm excited at 560 nm. Bars represent fluorescence intensity during 0, 10, 30 and 60 min after addition of various compounds excited at 560 nm.



**Fig. S5** cytotoxicity of DCM-B2 probe. Cell viability of MCF-7 cells incubated with different concentration DCM-B2 probe.



Fig. S7 <sup>13</sup>C NMR spectrum of compound DCM-OH



Fig. S8 <sup>1</sup>H NMR spectrum of compound DCM-B1



Fig. S9 <sup>13</sup>C NMR spectrum of compound DCM-B1



Fig. S10 IR spectrum of compound DCM-B1



Fig. S11 <sup>1</sup>H NMR spectrum of compound DCM-B2



Fig. S12 <sup>13</sup>C NMR spectrum of compound DCM-B2



Fig. S13 IR spectrum of compound DCM-B2



Fig. S14 Mass spectrum of compound DCM-OH



Fig. S15 Mass spectrum of the solution of compound DCM-B2 after treated with  $\rm H_2O_2$