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Supporting Information

Construction of an effective far-red fluorescent and colorimetric platform

containing a merocyanine core for specific and visual detection of

thiophenol in both aqueous medium and living cells

Xin Sun, ^a Mengzhao Wang, ^a Yanan Lu, ^a Zhengliang Lu,^{*,a} Chunhua Fan,^{*,a} Yizhong Lu^{*,b}

^a School of Chemistry and Chemical Engineering, University of Jinan, Jinan 250022, China.

^b School of Materials Science and Engineering, University of Jinan, Jinan 250022, China. Email:

Corresponding author's fax: +86-531-82765475

Corresponding email: zhengliang.lu@yahoo.com; chm_fanch@ujn.edu.cn; mse_luyz@ujn.edu.cn

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Figure S1.¹ H NMR spectrum of FRP-Thio (DMSO-*d*₆, 600 MHz).



Figure S2.¹³ C NMR spectrum of FRP-Thio (DMSO- d_6 , 600 MHz).



Figure S3. HRMS of FRP-Thio (DMSO-*d*₆, 600 MHz).



Figure S4. Time-dependent fluorescence changes of **FRP-Thio** (10 μ M) at 645 nm upon addition of thiophenol (0, 20, 50, 100, and 200 μ M) in DMF aqueous solution (2:8, v/v, PBS buffer 20 mM, pH 7.4).



Figure S5. Color changes of **FRP-Thio** without or with competing analytes and PhSH in DMF-H₂O (2:8, v/v, PBS buffer 20 mM, pH 7.4). a1 and b1) **FRP-Thio**. a2) Addition of various amino acids into a1). a3) Addition of PhSH into a2). b2) Addition of various anions and cations into b1). b3) Addition of PhSH into b2).



Figure S6. pH effect of FRP-Thio toward PhSH in DMF-H₂O (2:8, v/v).



Figure S7. HRMS of FRP-Thio with thiophenol.



Figure S8. MTT assay of SH-SY5Y cells with **FRP-Thio** at different concentration (0-20 μ M) for 24 hours.