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Supplementary data for

A rhodol-enone dye platform with dual reaction triggers for specific detection of Cys

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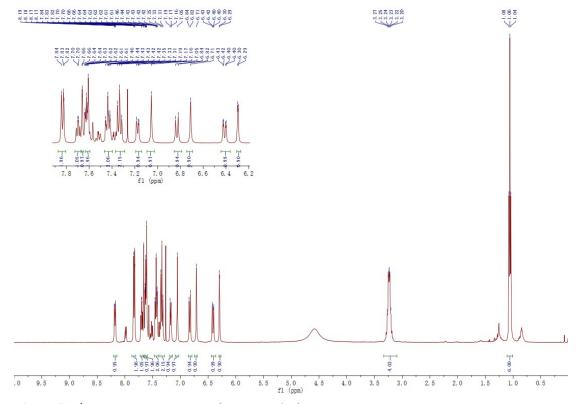


Figure S1. ^1H NMR spectroscopy of compound of BL.

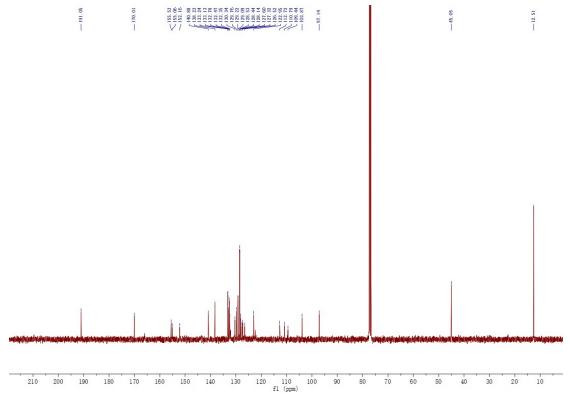


Figure S2. ¹³C NMR spectroscopy of compound of BL.

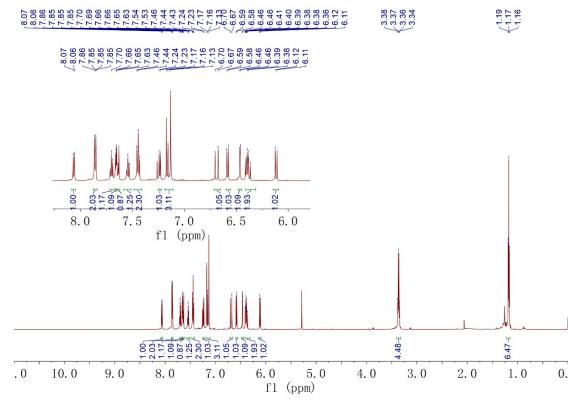


Figure S3. ¹H NMR spectroscopy of compound of probe **BL-C**.

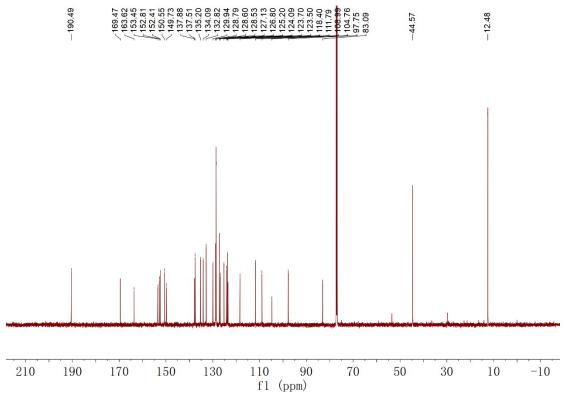
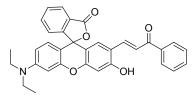


Figure S4. ¹³C NMR spectroscopy of compound of probe BL-C.



Exact Mass: 517.18892

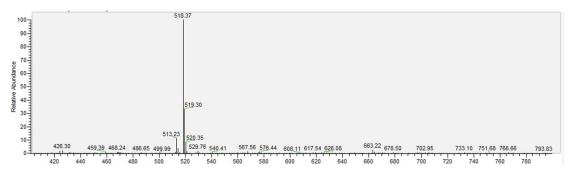


Figure S5. MS spectroscopy of **BL** (M+H)⁺.

Exact Mass: 571.19949

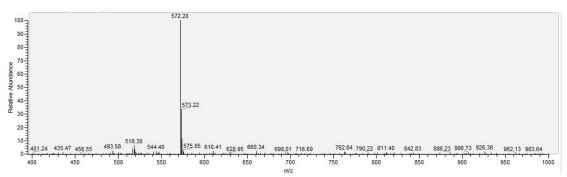


Figure S6. MS spectroscopy of probe **BL-C** (M+H)⁺.

Exact Mass: 638.20867

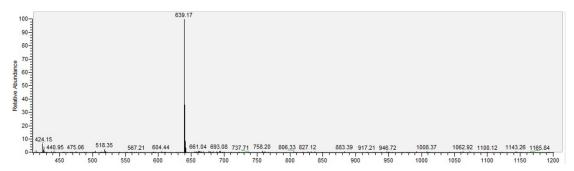


Figure S7. MS spectroscopy of probe BL-C after the treatment with Cys (M+H)⁺.

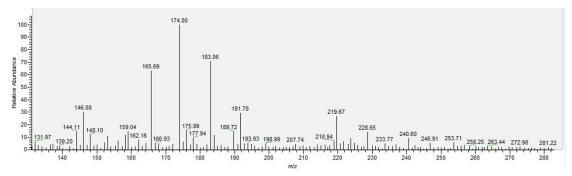


Figure S8. MS spectroscopy of the lactam produced by the reaction of probe **BL-C** with Cys (M-H)⁻.

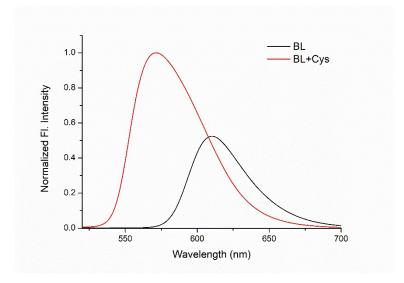


Figure S9. Fluorescence spectra of **BL** (10 μ M) in the absence and presence of Cys (100 μ M).

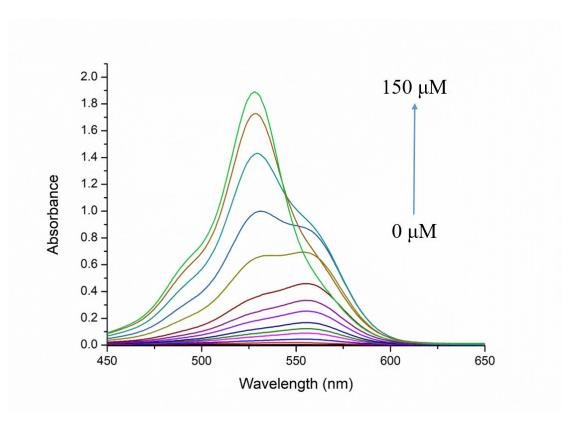


Figure 10. UV/Vis absorption responses of probe **BL-C** (10 μ M) toward different concentrations of Cys (0-150 μ M).

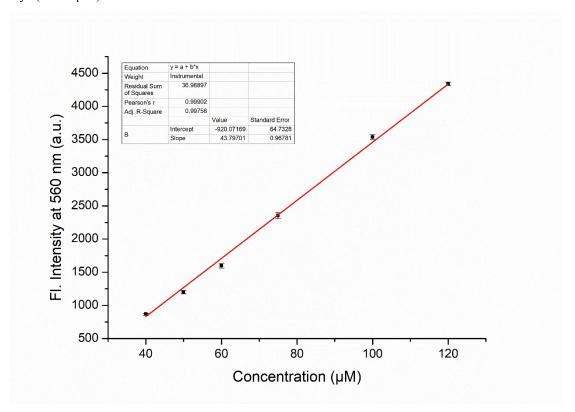


Figure S11. Fluorescence intensity at 560 nm of probe BL-C versus Cys concentration (40-120 μ M) with λ_{ex} =480 nm. Error bars, SD (n = 3).

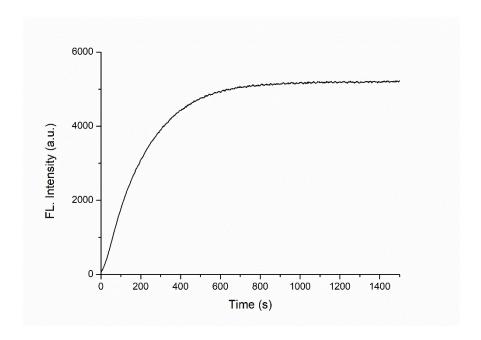


Figure S12. The time-dependent changes of the fluorescence intensity at 560 nm in the presence of 150 μM Cys