

Supporting Information

Two isomeric and distinguishable H₂S fluorescence probes for monitoring spoilage of eggs as well as visualizing exogenous and endogenous H₂S in living cells

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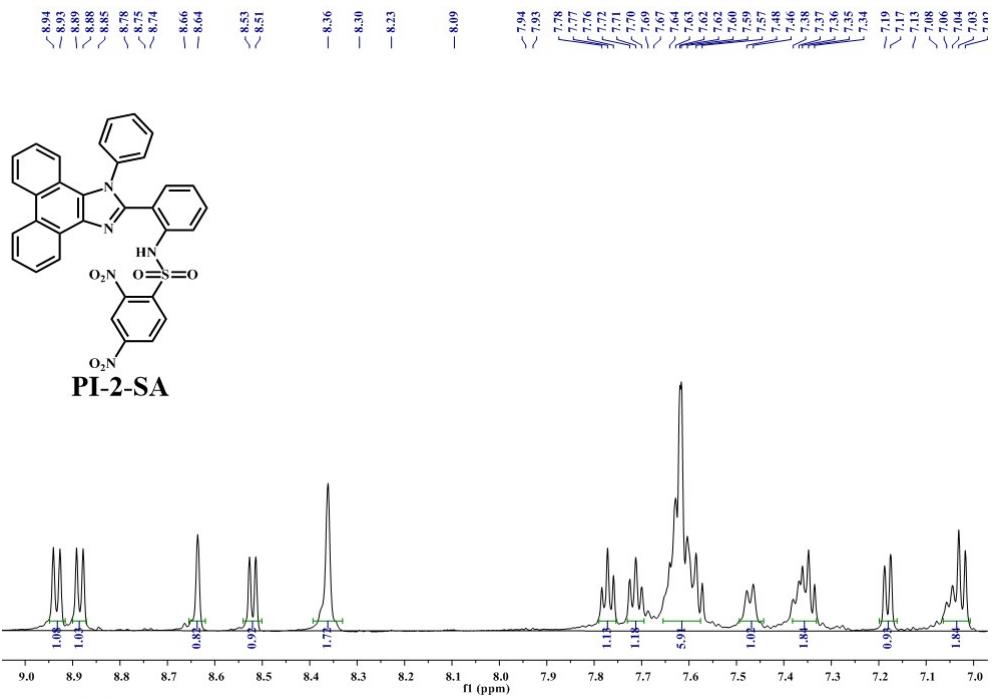


Fig. S1 ^1H NMR spectra of **PI-2-SA** in $\text{DMSO}-d_6$ (600 MHz).

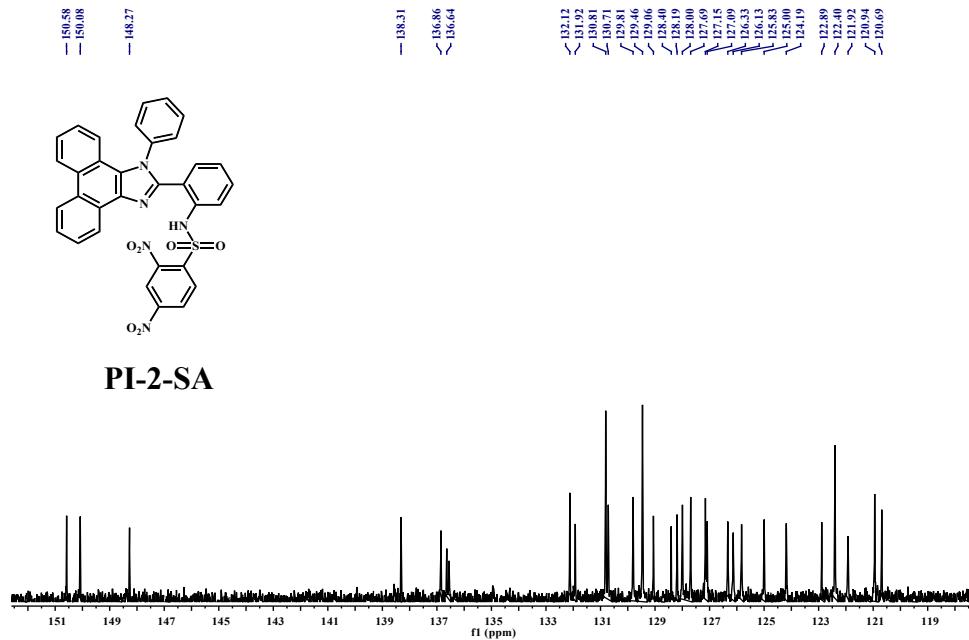


Fig. S2 ^{13}C NMR spectra of **PI-2-SA** in $\text{DMSO}-d_6$ (150 MHz).

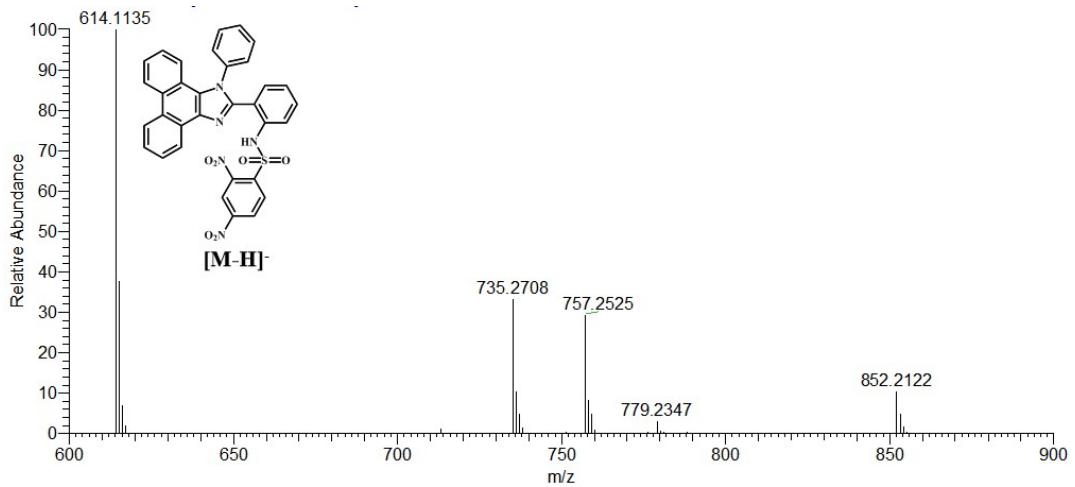


Fig. S3 ESI-MS spectrum of PI-2-SA.

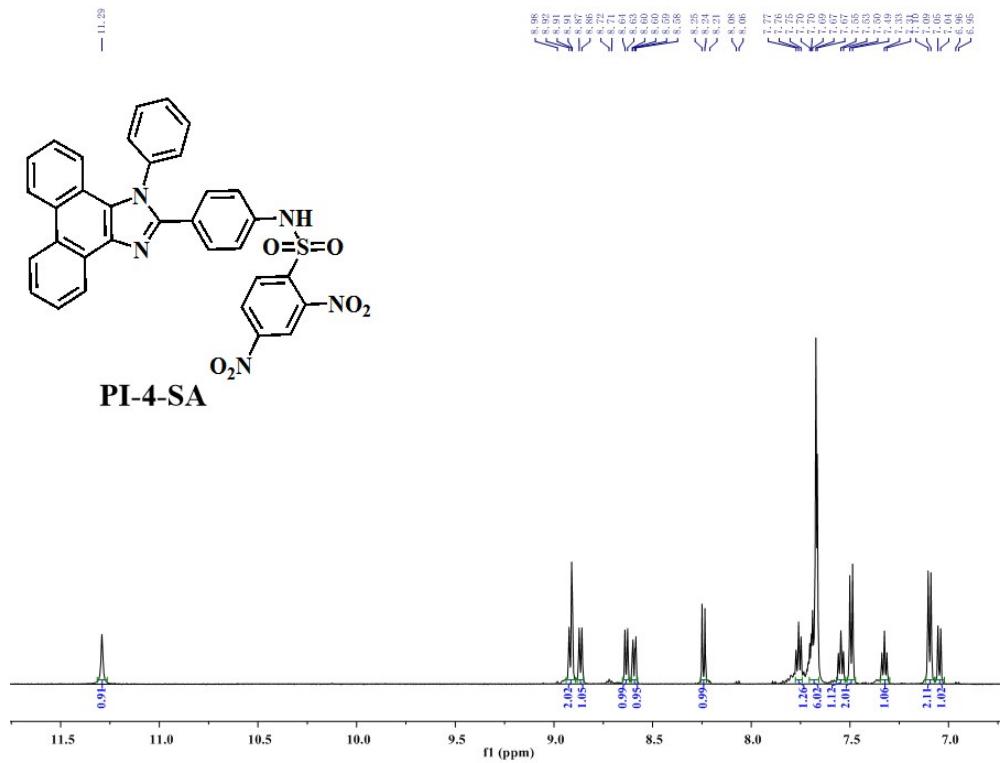


Fig. S4 ^1H NMR spectra of PI-4-SA in $\text{DMSO}-d_6$ (600 MHz).

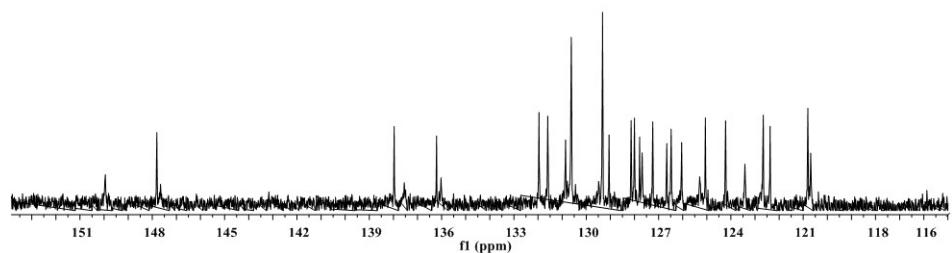
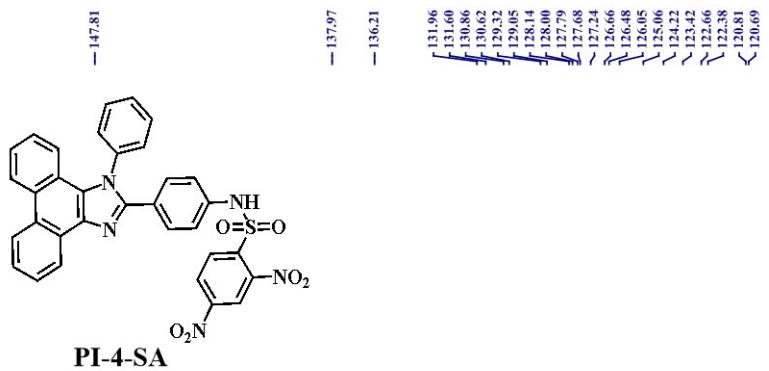


Fig. S5 ^{13}C NMR spectra of PI-4-SA in $\text{DMSO}-d_6$ (150 MHz).

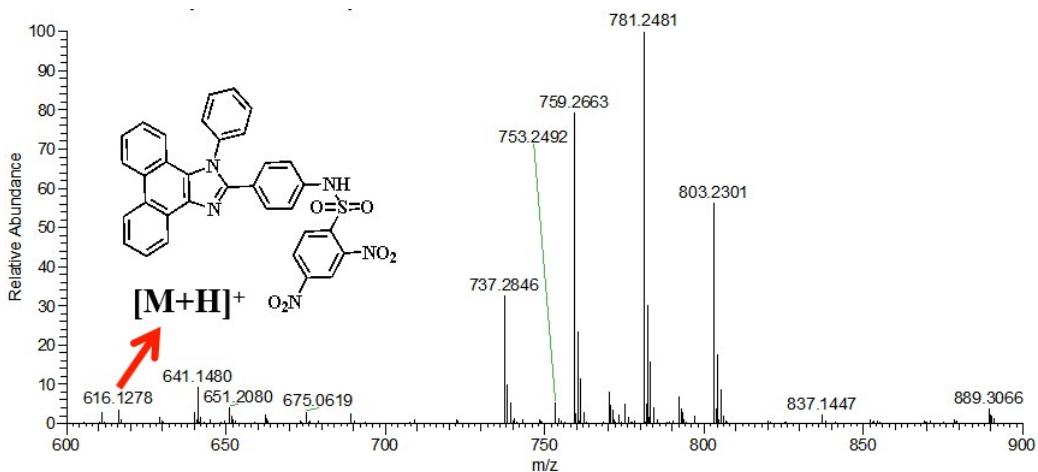


Fig. S6 ESI-MS spectrum of PI-4-SA.

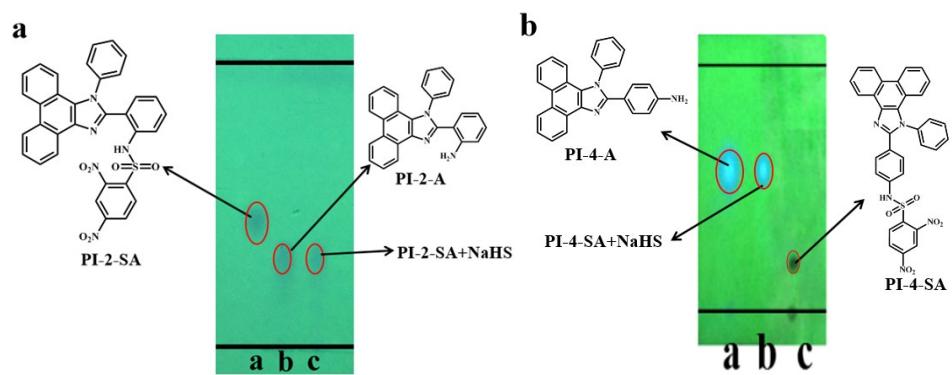


Fig. S7 Thin layer chromatography (TLC) analysis of the reaction of **PI-2-SA** (10 μ M) with NaHS (50 μ M) (a) and **PI-4-SA** with NaHS (0.25 mM) (b) in DMF/H₂O (v/v=1/1, Hepes 10 μ M, pH =7.4) solution.

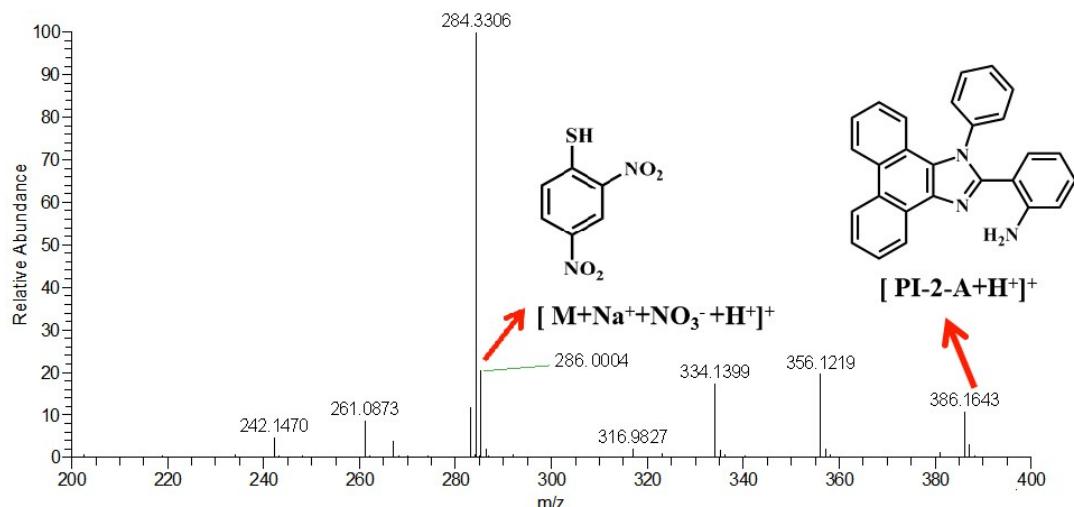


Fig. S8 ESI-MS spectrum for the reaction products of **PI-2-SA** with NaHS.

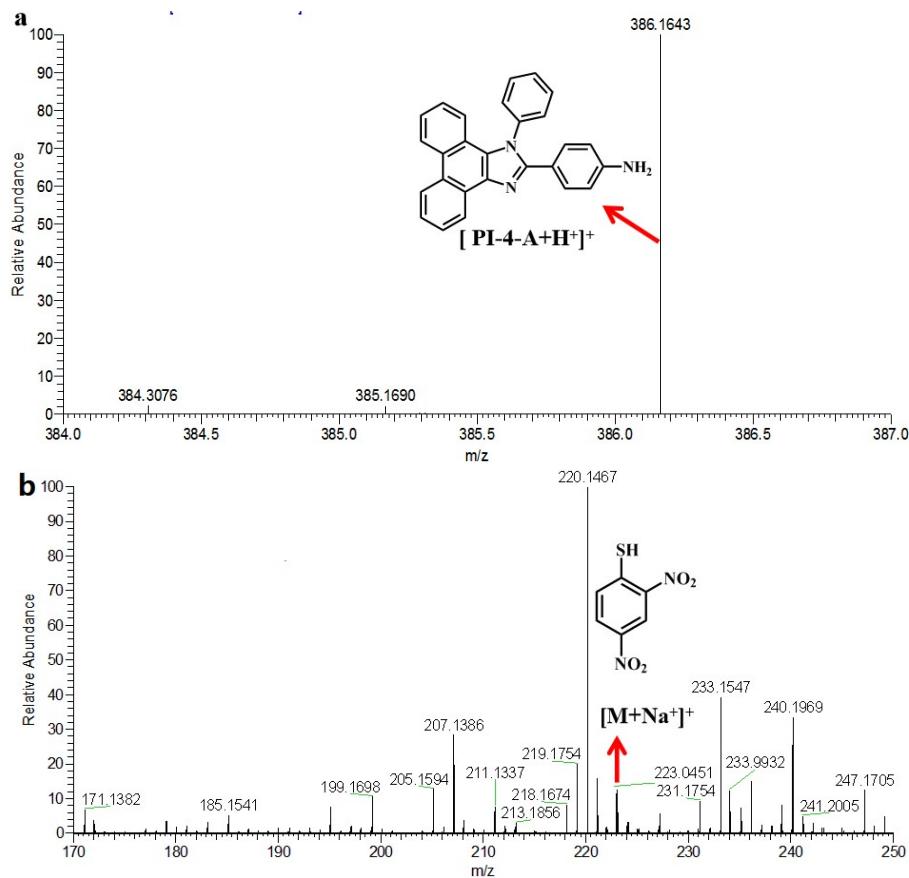


Fig. S9 ESI-MS spectrum for the reaction products of PI-4-SA with NaHS (a for PI-4-A and b for 2,4-dinitro

phenthiol).

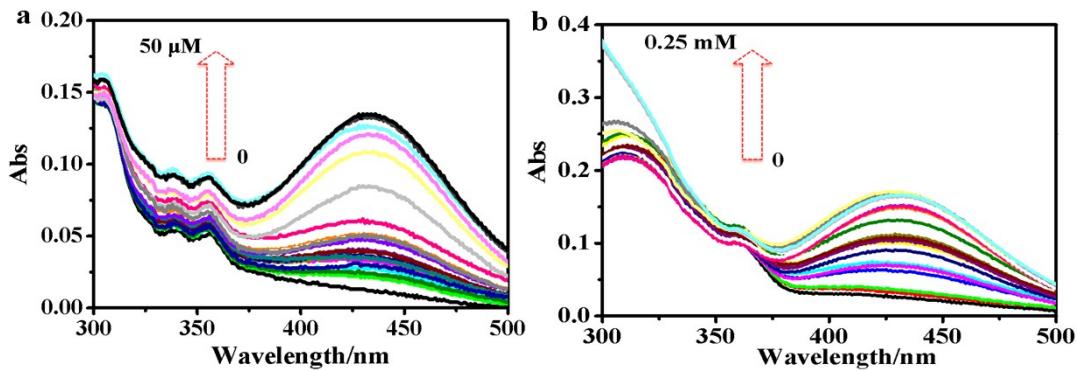


Fig. S10 UV-vis spectra of PI-2-SA (10 μ M) (a) and PI-4-SA (10 μ M) (b) with various amount of NaHS in DMF/H₂O

(v/v=1/1, Hepes 10 μ M, pH =7.4).

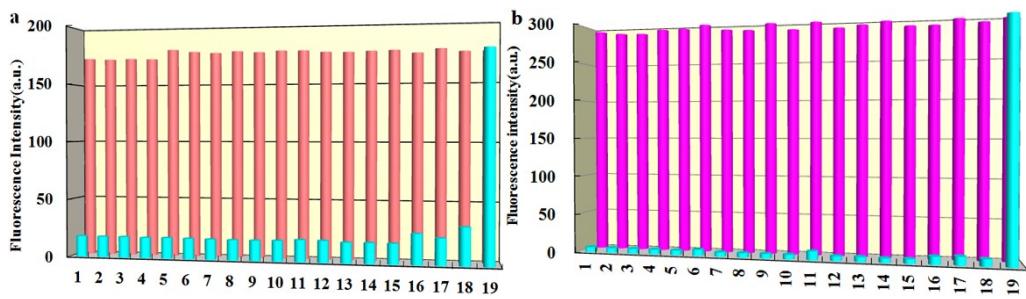


Fig. S11 Fluorescence intensities of **PI-2-SA** (10 μ M, λ_{ex} =310 nm) (a) and **PI-4-SA** (10 μ M, λ_{ex} =329 nm) (b) upon

the addition of various relevant matters (1.blank; 2.Br⁻; 3.Cl⁻; 4.F⁻; 5.BrO³⁻; 6.I⁻; 7.SO₃²⁻; 8.HSO₃⁻; 9.SCN⁻; 10.NO₃⁻; 11.H₂PO₄⁻; 12.CO₃²⁻; 13.HCO₃⁻; 14.AcO⁻; 15.GSH; 16.L-Cys; 17.Hcy; 18.S²⁻; 19.HS⁻) in the presence of NaHS with mole ratio of 1 : 2 between NaHS and analyte.

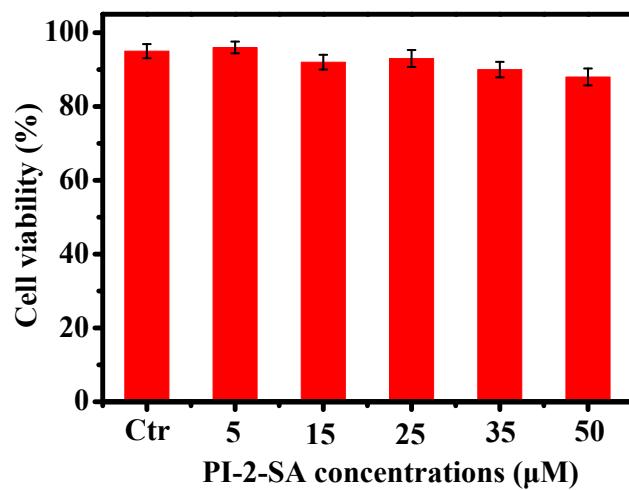


Fig. S12 MTT assay of HeLa cells in the presence of different concentrations of **PI-2-SA**.