Electronic Supplementary Information (ESI)

A fluorescent probe with aggregation-induced emission characteristics for distinguishing homocysteine over cysteine and glutathione

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Fig. S1¹ H NMR spectrum of TPE-Py.



Fig. S2 ¹³C NMR spectrum of TPE-Py.



Fig. S3 High resolution mass spectrum of TPE-Py.



Fig. S4 High resolution mass spectrum of TPE-Py-Hcy.







Fig. S6¹³C NMR spectrum of TPEPh-Py.



Fig. S7 High resolution mass spectrum of TPEPh-Py.



Fig. S8 UV-vis spectra of TPE-Py and TPEPh-Py (3 $\mu M)$ in acetonitrile.



Fig. S9 (A) PL spectra of TPEPh-Py (3 μ M) in acetonitrile and acetonitrile–water mixtures with different water fractions (f_w). (B) Plots of emission intensity versus the composition of the aqueous mixtures of TPEPh-Py. Inset: photographs of TPEPh-Py in acetonitrile–water mixtures with f_w values of 0 and 90 vol%.



Fig. S10 UV-vis spectra of TPE-Py (3 μ M) in acetonitrile/phosphate buffer (20 : 80 v/v, 20 mM, pH 8) with different concentration of (A) Hcy and (B) Cys (0-60 mM).



Fig. S11 PL spectra of TPE-Py (3 μ M) in acetonitrile/phosphate buffer (20 : 80 v/v, 20 mM, pH 8) with different concentration of GSH (0-60 mM).



Fig. S12 PL spectra of TPEPh-Py (3 μ M) in acetonitrile/phosphate buffer (20 : 80 v/v, 20 mM, pH 8) with different concentration of Hcy (0-60 mM).

Fig. S13 Normalized fluorescence responses of TPE-Py (3 μ M) to changing Hcy concentrations (1.5, 3.0, 6.0, 12.0, 18.0 μ M) in acetonitrile/phosphate buffer (20 : 80 v/v, 20 mM, pH 8). Detection limit = 3.46 × 10⁻⁷ M. F = I_{455}/I_{550} .

Fig. S14 PL spectra of TPE-Py (3 μ M) in acetonitrile/phosphate buffer (20 : 80 v/v, 20 mM, pH 8) with different amino acids (60 mM).

Fig. S15 Emission ratio I_{455}/I_{550} of TPE-Py in the absence or presence of 60 mM typical typical thiol compounds and non-thiol nucleophiles.