Electronic supplementary information

A Michael addition reaction-based fluorescent probe for malononitrile detection and its applications in aqueous solution, living cells and zebrafish

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Fig. S5. Time-dependent fluorescence spectral changes of probe Hcy-DCV (10 μ M) upon the addition of malononitrile (100 μ M) in PBS buffer (pH=7.4, 10 mM) at 25°C ($\lambda_{ex} = 300$ nm, $\lambda_{em} = 460$ nm).



Fig. S6. Color changes of **Hcy-DCV** solutions (10 μ M) upon the addition of different interfering analysts and malononitrile (200 μ M) under natural light and UV radiation. (1, K⁺; 2, Cu²⁺; 3, Mg²⁺; 4, Ca²⁺; 5, Fe³⁺; 6, Cl⁻; 7, NO₂⁻; 8, SO₄²⁻; 9, CO₃²⁻; 10, L-Phe; 11, L-Ile; 12, L-Lys; 13, L-Pro; 14, L-Arg; 15, Acetoacetate; 16, Acetylacetone; 17, Dimethyl sulfone; 18, Dimethyl sulfoxide; 19, Malonic acid; 20, Diethyl malonate; 21, Nitromethane; 22, Methyl cyanoacetate; 23, Acetonitrile; 24, *p* - Nitrophenylacetonitrile; 25, Hydrazine; 26, CNCH₂CN).



Fig. S7. Partial ¹H NMR spectra of probe **Hcy-DCV** in the absence (a) and presence (b) of malononitrile in DMSO- d_6 (400 MHz).



Fig. S8. HR-MS of the reaction product of probe Hcy-DCV and malononitrile.



Fig. S9. Cell viability of H1975 cells after the treatment by different concentrations of Hcy-DCV (0-32 μ M).