

Supporting Information

A colorimetric sensor for the sequential detection of Cu²⁺ and CN⁻ in fully aqueous media: practical performance of Cu²⁺

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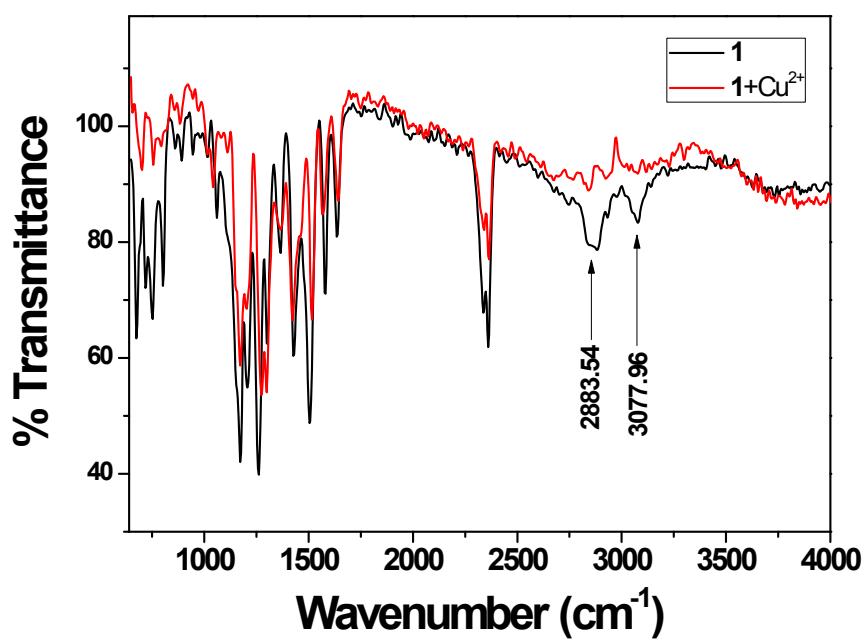


Fig. S1 FT-IR spectra of **1** and Cu²⁺-**2c31** complex.

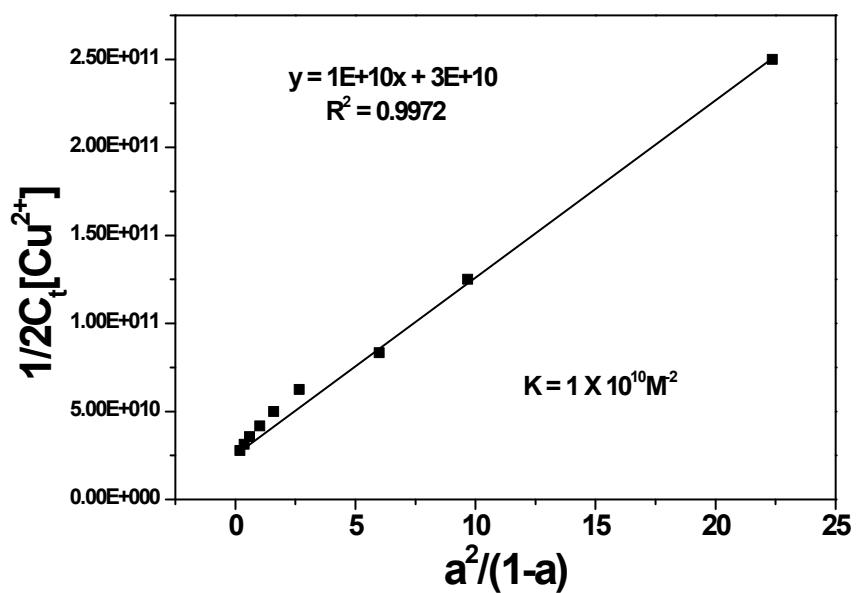


Fig. S2 Li's equation plot (absorbance at 525 nm) of **1**, assuming 2:1 stoichiometry for association between **1** and Cu^{2+} .

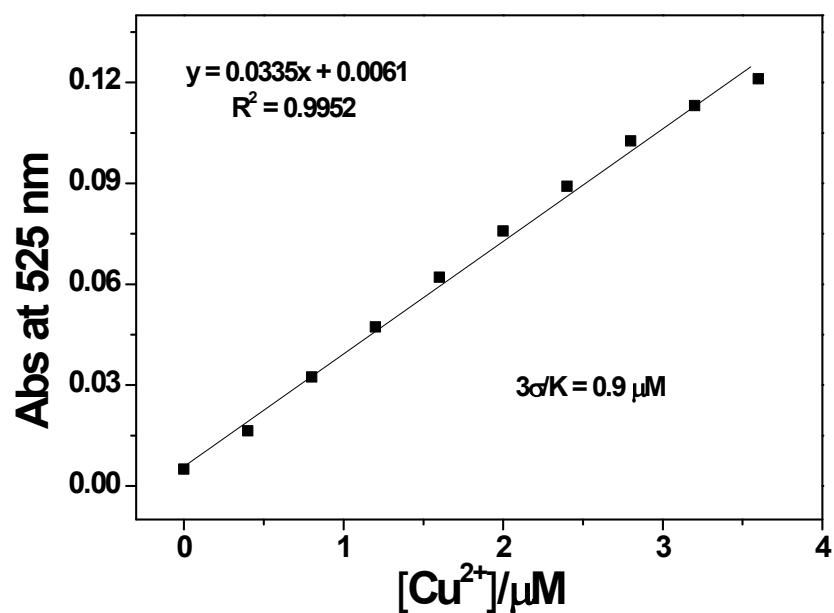


Fig. S3 Determination of the detection limit based on change in the ratio (absorbance at 525 nm) of **1** (10 μM) with Cu^{2+} .

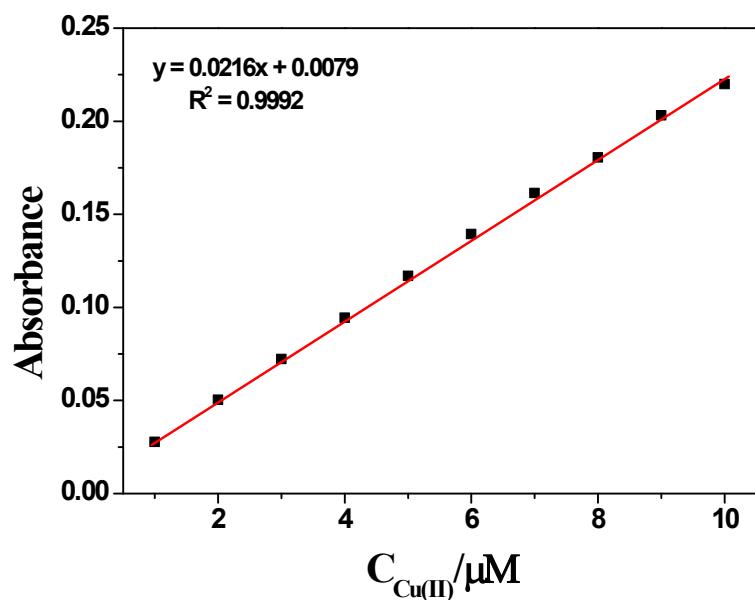


Fig. S4 Absorption intensity (at 525 nm) of **1** as a function of Cu^{2+} concentration. $[1] = 30 \mu\text{mol/L}$ and $[\text{Cu}^{2+}] = 1.00\text{-}10.00 \mu\text{mol/L}$ in 10 mM bis-tris buffer-DMSO solution (8:2, pH 7.0).

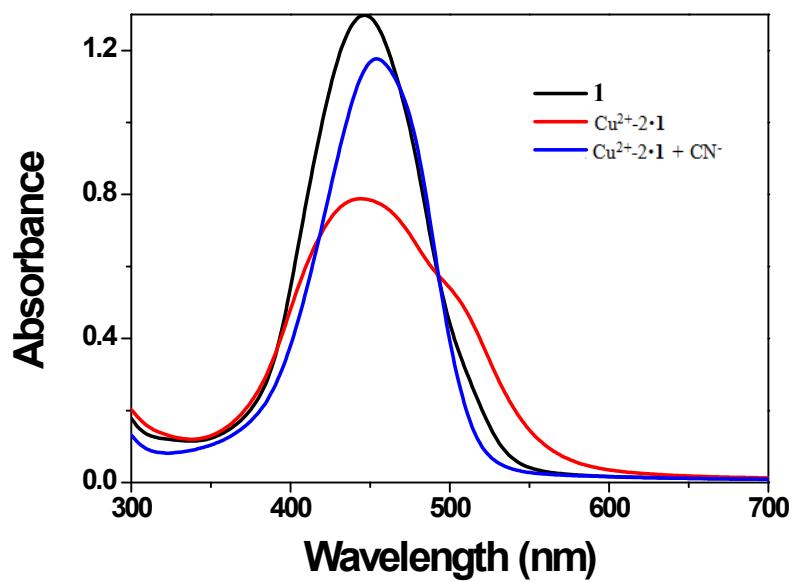


Fig. S5 UV-vis spectra of **1** (30 μM), $\text{Cu}^{2+}\text{-}2\bullet\text{1}$ (Cu^{2+} = 15 μM), and $\text{Cu}^{2+}\text{-}2\bullet\text{1} + \text{CN}^-$ (200 equiv) in bis-tris buffer solution.

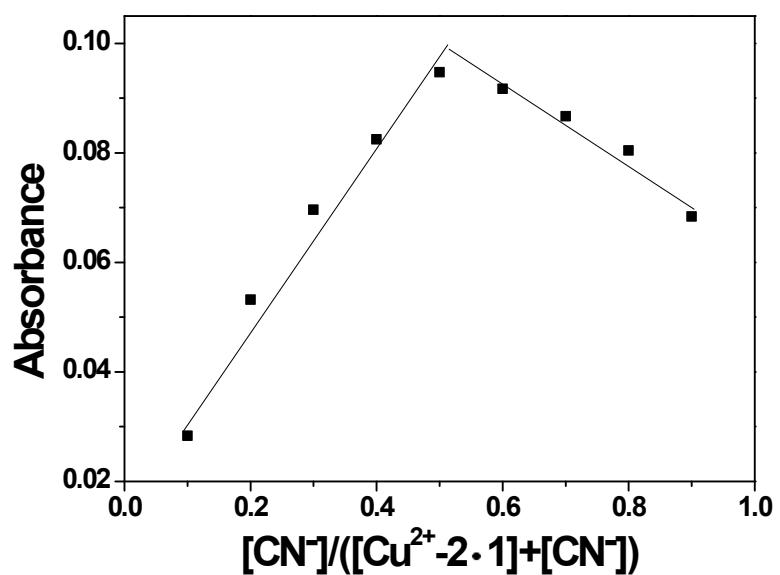


Fig. S6 Job plot of $\text{Cu}^{2+}\text{-2}\text{-1}$ complex and CN^- , where the intensity at 454 nm was plotted against the mole fraction of CN^- . The total concentrations of CN^- with $\text{Cu}^{2+}\text{-2}\text{-1}$ complex were 100 μM .

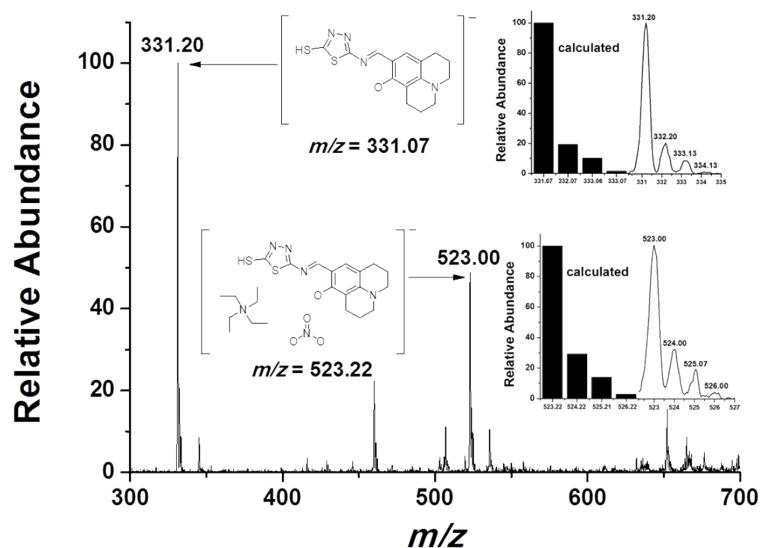


Fig. S7 Negative-ion electrospray ionization mass spectrum of Cu²⁺-2>1 (0.1 mM) upon addition of CN⁻ (1 equiv).

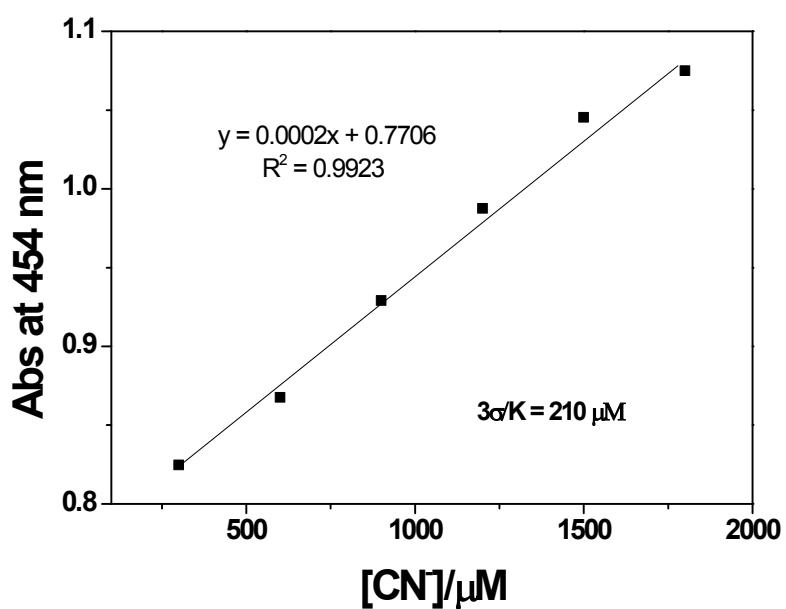


Fig. S8 Determination of the detection limit based on change in the ratio (absorbance at 454 nm) of Cu²⁺-2 ω 1 (15 μM) with CN⁻.

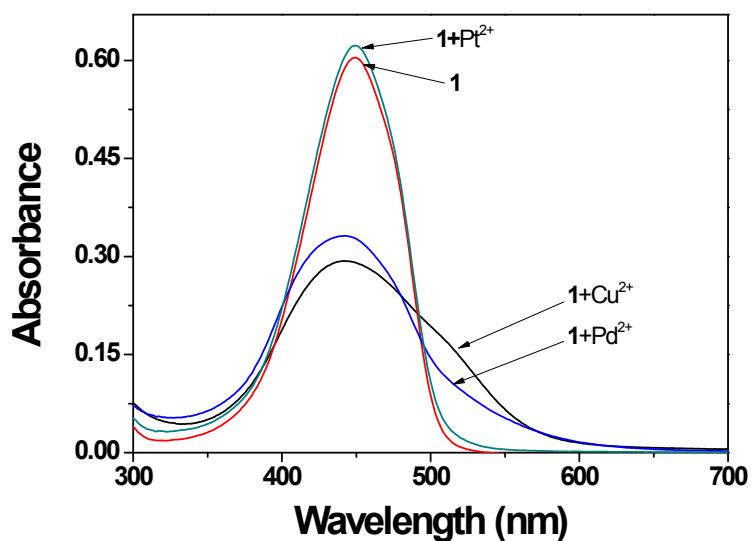


Fig. S9 UV-vis spectra of **1** (10 μ M), **1+Cu²⁺**, **1+Pd²⁺** and **1+Pt²⁺** (0.5 equiv) in bis-tris buffer solution.