Electronic supplementary information (ESI)

A Colorimetric and Ratiometric Fluorescent Probe for Copper(II) with Large Red Shift and Its Ratiometric Imaging in Living Cells

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Fig. S1 Time-dependent fluorescent behavior of probe 1 with different equiv. of Cu^{2+} in H₂O-CH₃CN HEPES buffer (pH 7.4, 25 mM) (50%, V/V).



Fig. S2 Fluorescence spectra of the probe 1 (10 μ M, $\lambda_{ex} = 376$ nm) upon the addition of various metal ions (15 μ M) in HEPES buffered (pH 7.4, 25 mM) H₂O-Ethanol (50%, V/V).



Fig. S3 Fluorescence spectra of the probe 1 (10 μ M, $\lambda_{ex} = 375$ nm) upon the addition of various metal ions (15 μ M) in HEPES buffereded (pH 7.4, 25 mM) H₂O-DMSO (50%, V/V).



Fig. S4 Fluorescence spectra of the probe 1 (10 μ M, $\lambda_{ex} = 371$ nm) upon the addition of various metal ions (15 μ M) in HEPES buffered (pH 7.4, 25 mM) H₂O-CH₃CN (70%, V/V).



Fig. S5 Fluorescence spectra of the probe 1 (10 μ M, $\lambda_{ex} = 376$ nm) upon the addition of various metal ions (15 μ M) in HEPES buffered (pH 7.4, 25 mM) H₂O-CH₃CN (90%, V/V).



Fig.S6 The ratio of fluorescence intensity of the probe 1 at 554 nm and 411nm in the presence of absence of Cu^{2+} at different pH conditions.



Fig. S7 ¹H NMR spectrum of probe 1 in CDCl₃



Fig. S8 Mass (ESI) spectrum of probe 1 (M+1).



Fig. S9 Mass (ESI) spectrum of the reaction products of probe 1 with Cu²⁺.