Supplementary Information (ESI) for Dalton Transactions, 2009/12/10 Supporting Information

For

Colorimetric Sensing of Cu(II): Cu(II) Induced Deprotonation of an Amide Responsible for Color Changes

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Figure S1. Fluorescence spectra of **1** (100 μ M) in a methanol-H₂O solution (v/v = 4/1, 20 mM Hepes buffer, pH 7.0) in the presence of different metal ions (100 μ M).



Figure S2. Job's plot of **2**-Ni²⁺ (top) and **2**-Co²⁺ (bottom) complexes, where the absorbance at 483 nm (**2**-Ni²⁺) or the fluorescence at 520 nm (**2**-Co²⁺) was plotted against the mole fraction of Ni²⁺ or Co²⁺ at a constant total concentration of 1.0×10^{-4} M in a methanol-H₂O solution (v/v = 4/1, 20 mM Hepes buffer, pH 7.0).



Figure S3. Absorption spectra of 1 (100 μ M) at different pH values in a methanol-water solution (v/v = 4:1, 20 mM buffer).



Figure S4. pH titration of Cu(II) binding with chemosensor **2** (10^4 M) in a methanol-H₂O solution (v/v = 4/1, 20 mM buffer). Both concentrations of Cu(II) was 10^4 M.



Figure S5. pH titration of Ni(II) (top) or Co(II) (bottom) binding with chemosensor **2** (10^{-4} M) in a methanol-H₂O solution (v/v = 4/1, 20 mM buffer). Both concentrations of Ni(II) (top) or Co²⁺ were 10^{-4} M.



Figure S6. IR spectra of 1 and its mixture with Cu(II) in methanol.