

Supporting information

A ratiometric and colorimetric probe for detecting Hg²⁺ based on naphthalimide-rhodamine and its staining function in cell imaging

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Cell culture and imaging

In light of good cellular permeability of probe **1**, the probe was applied to cell staining. The fluorescence images were taken by an OLYMPUS FV1000 confocal laser scanning microscope. HeLa cells (human cervical cancer cell) were cultured in cell staining (Dulbecco modified Eagle's medium) supplemented with 10% FBS (fetal bovine serum) in an atmosphere of 5% CO₂ and 95% air at 37 °C overnight. Before the cell staining, the cells of DMEM were washed three times with phosphate-buffered saline (PBS). Subsequently, the cells were incubated with probe **1** (20 μM) in the culture medium for 30 min and washed with PBS three times.

Fig. S1. ¹H NMR spectrum of **7** in CD₂Cl₂.

Fig. S2. ¹³C NMR spectrum of probe **7** in CD₂Cl₂.

Fig. S3. The ESI-MS spectrum of **7**.

Fig. S4. ¹H NMR spectrum of **1** in CD₂Cl₂.

Fig. S5. ¹³C NMR spectrum of probe **1** in CD₂Cl₂.

Fig. S6. The ESI-MS spectrum of **1**.

Fig. S7. (A) Nonlinear curve fitting of the fluorescence titration data from 0-21 μM for **1** with Hg²⁺ at 604 nm in CH₃CN-H₂O (7/3, v/v) solution at room temperature; (B) The limit of detection (LOD) of **1** towards Hg²⁺ by fluorescence measured at 604 nm.

Fig. S8. (A) Nonlinear curve fitting of change of absorbance from 0-70 μM for **1** with Hg²⁺ in CH₃CN-H₂O (7/3, v/v) solution at room temperature; (B) The limit of detection (LOD) of change of absorbance of probe **1** towards Hg²⁺ by UV-vis measured.

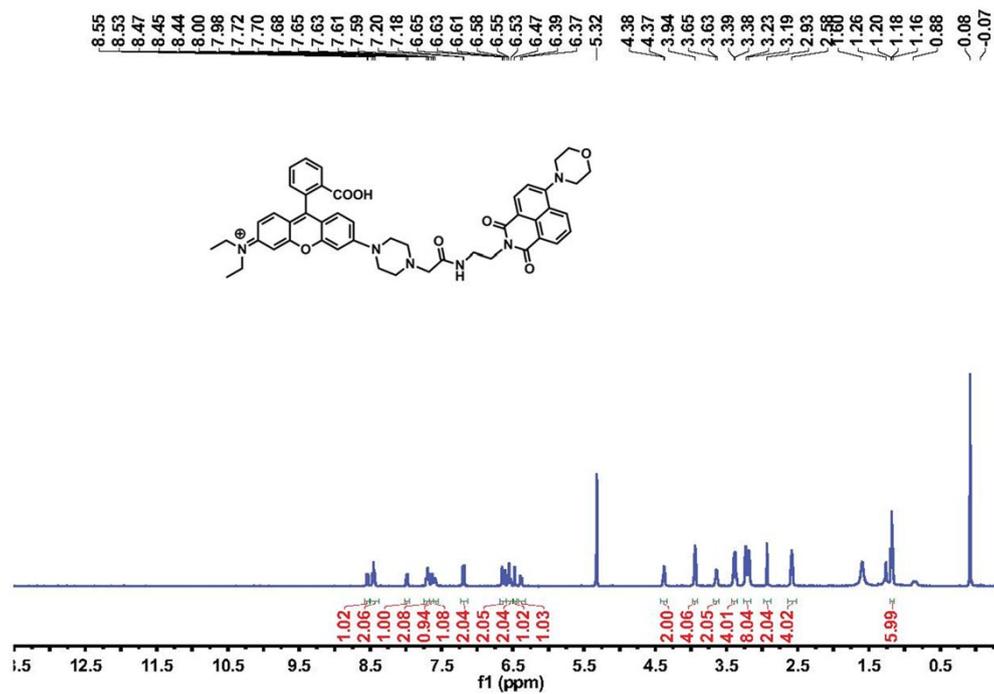


Fig. S1.

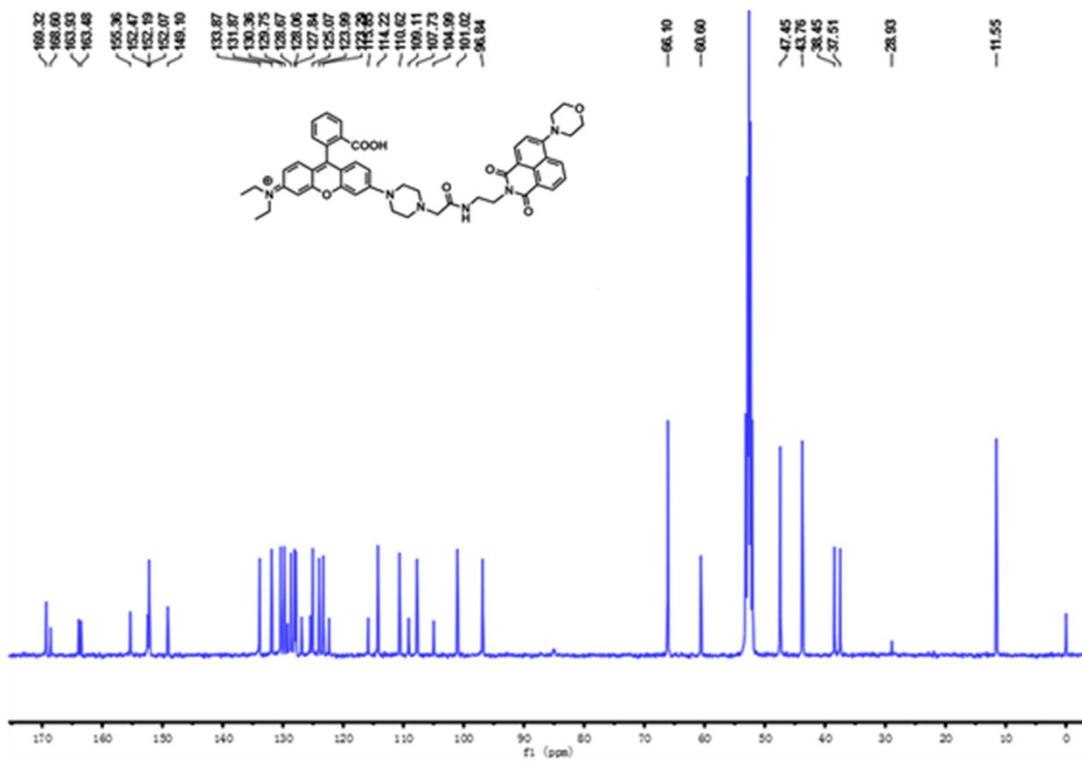


Fig. S2.

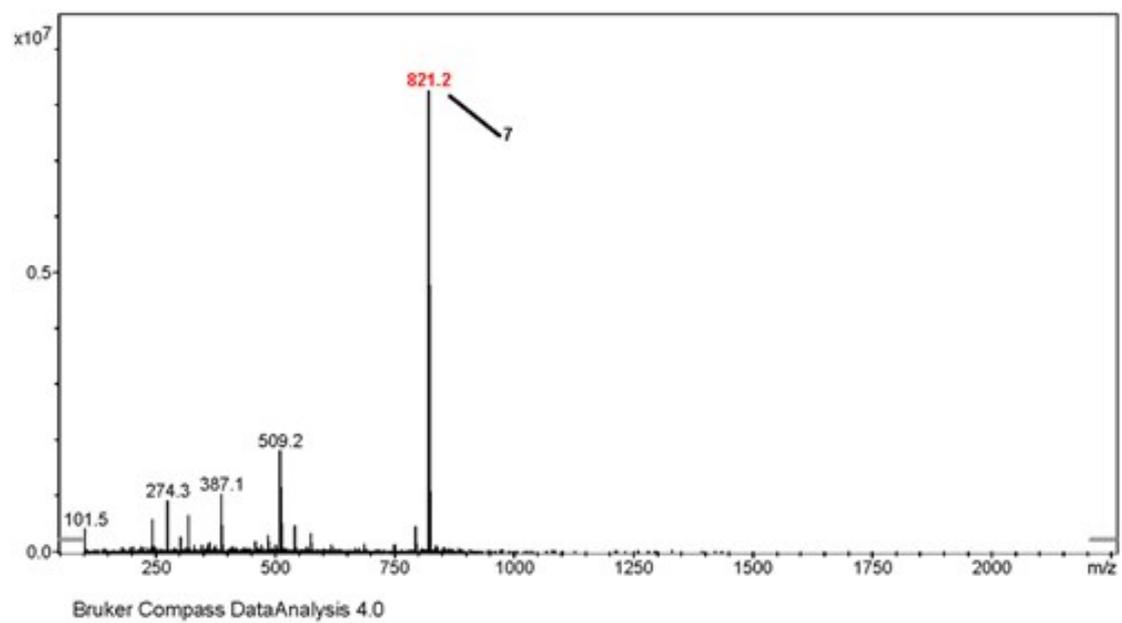


Fig. S3

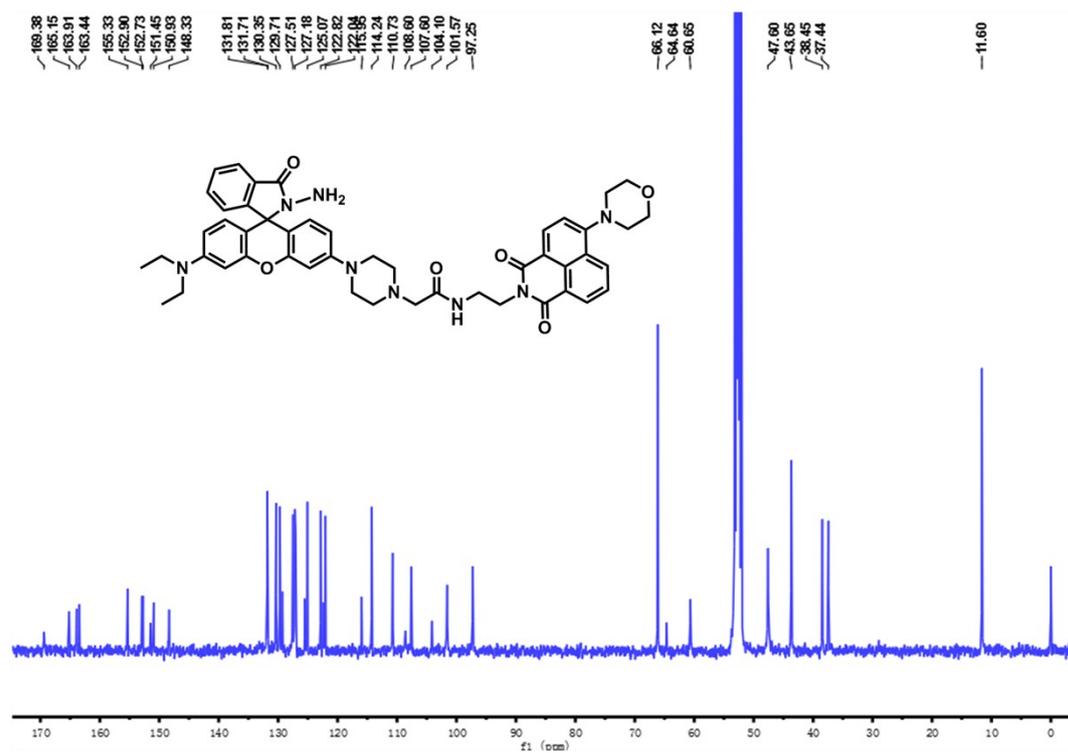
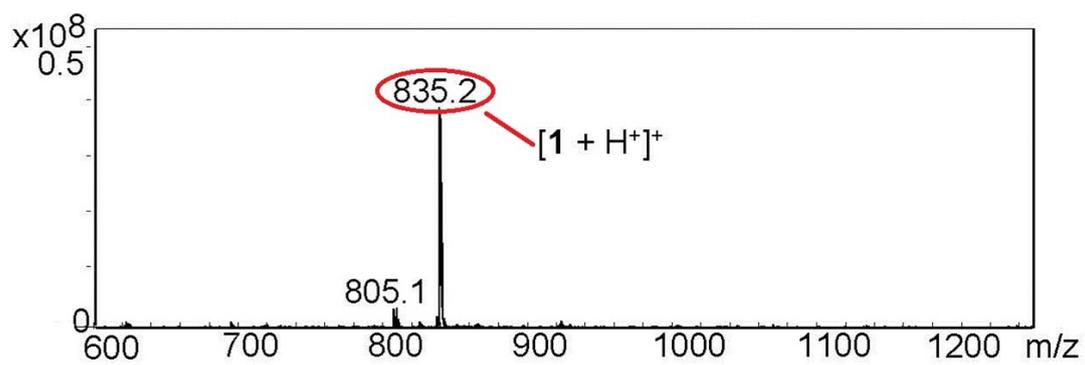


Fig. S5



Bruker Compass DataAnalysis 4.0

Fig. S6

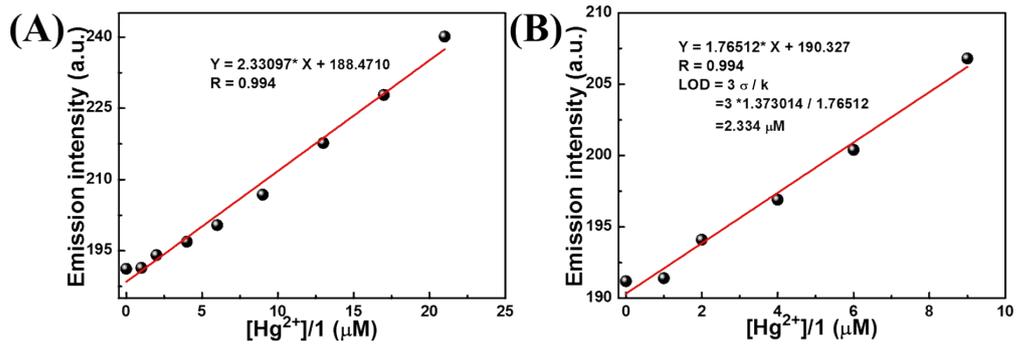


Fig. S7

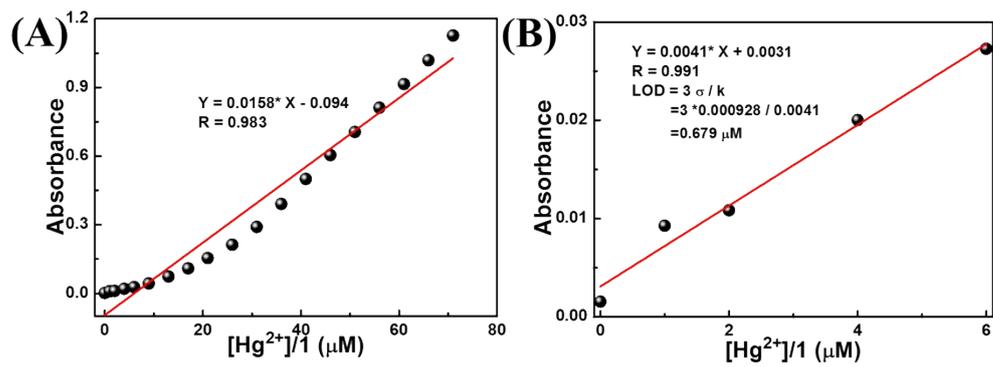


Fig. S8