## **Supporting Information**

## Visual determination of ferric ions in aqueous solution based on a high selectivity and sensitivity ratiometric fluorescent nanosensor

Haolan Wu<sup>a</sup>, Lian Yang<sup>b</sup>, Linfeng Chen<sup>c</sup>, Fang Xiang<sup>d</sup> and Hongyu Gao<sup>f,\*</sup>

<sup>a</sup> School of Water Resources and Environment, China University of geosciences (Beijing), Beijing 100083, China.

<sup>b</sup> Research Center for Sewage Recycling Engineering of Beijing, Beijing 100124, China

<sup>c</sup> Faculty of Material Science and Chemistry, China University of Geosciences, Wuhan, 430074China

<sup>d</sup> School of Foreign Language, China University of Geosciences, Wuhan 430074, China

<sup>*f*</sup> Institute for resources and environmental engineering, Shanxi University, Taiyuan 030000, China. E-mail: <u>sailor0611@163.com</u>



Scheme.S1. The synthetic route of the RhB

<sup>1</sup>H NMR (400MHz, CDCl<sub>3</sub>) δ (ppm): 7.97 (d, 1H), 7.45 (m, 4H), 7.11 (d, 1H), 6.48 (m, 4H), 6.20 (m, 2H), 3.91 (m, 6H), 3.31(m, 4H), 2.31 (s, 6H), 1.15 (t, 6H).

<sup>13</sup>C NMR (400MHz, d<sub>6</sub>-DMSO) δ (ppm): 165.46, 156.82, 152.11, 150.14, 140.95, 134.45, 129.08, 127.16, 122.02, 120.71, 118.51, 61.32, 49.79, 15.97, 13.74.

Elemental analysis: calcd for C<sub>34</sub>H<sub>34</sub>N<sub>4</sub>O<sub>5</sub>: C, 70.57; H, 5.92; N, 9.68. Found: C,



Fig. S1. The <sup>1</sup>H NMR spectrum of RhB in CDCl<sub>3</sub>



Fig. S2 The  ${}^{13}$ C NMR spectrum of RhB in  $d_6$ -DMSO



Fig. S3. The TEM image of (a) CdTe QDs and (b) CdTe@SiO<sub>2</sub> QDs.



**Fig. S4.** (a) The stability of the RhB-CdTe@SiO<sub>2</sub> QDs sensor in acetonitrile/water mixtures (1:4 v/v, 1/1; pH=7.0). (B) The stability of the RhB-CdTe@SiO<sub>2</sub> QDs sensor in the presence of 4.5  $\mu$ M of Fe<sup>3+</sup>.

Rhodamine-based fluorescence sensor for Fe <sup>3+</sup>	LOD	linear ranges(mol/L <sup>-1</sup> )
An "off-on" rhodamine-based fluorescence probe 1	14 nM	6.0×10 <sup>-8</sup> -7.2×10 <sup>-6</sup>
Ferric ion chemosensor based on rhodamine derivative <sup>2</sup>	0.32 µM	0-2.0×10 <sup>-5</sup>
Rhodamine-based "off-on" chemosensors for ferric ion <sup>3</sup>	5.0 µM	$5.0 \times 10^{-6}$ - $2.0 \times 10^{-5}$
Rhodamine hydroxamate as fluorescent chemosensor <sup>4</sup>	1.0 µM	$1.0 \times 10^{-6}$ - $2.0 \times 10^{-5}$
Rhodamine-based thiacalix[4]arene fluorescent sensor <sup>5</sup>	35.0 nM	$5.0 \times 10^{-6}$ - $6.0 \times 10^{-5}$
This Work	20.5 nM	0-3.5×10 <sup>-6</sup>

Table S1 The comparison of the LOD and linear range for the detection of  $Fe^{3+}$  by other rhodamine-based fluorescence sensor.

## References

- 1. J. Mao, Q. He and W. Liu, Anal. Bioanal. Chem., 2010, 80, 2093.
- 2. H. Liu, X. Wan, L. Gu, T. Liu and Y. Yao, *Tetrahedron*, 2014, 70, 7527-7533.
- Z. Yang, M. She, B. Yin, J. Cui, Y. Zhang, W. Sun, J. Li and Z. Shi, *J. Org. Chem.*, 2012, 77, 1143.
- 4. K. S. Moon, Y. K. Yang, S. Ji and J. Tae, *Tetrahedron Lett.*, 2010, 51, 3290-3293.
- 5. X. Y. Zheng, W. J. Zhang, L. Mu, X. Zeng, S. F. Xue, Z. Tao and T. Yamatob, *J. Incl. Phenom. Macro.*, 2010, 68, 139-146.