

Revised Electronic Supplementary Information

Construction of Ratiometric Phosphorescent Assay with Long-Lived Carbon Quantum Dots and Inorganic nanoparticles for its application in environmental and biological system

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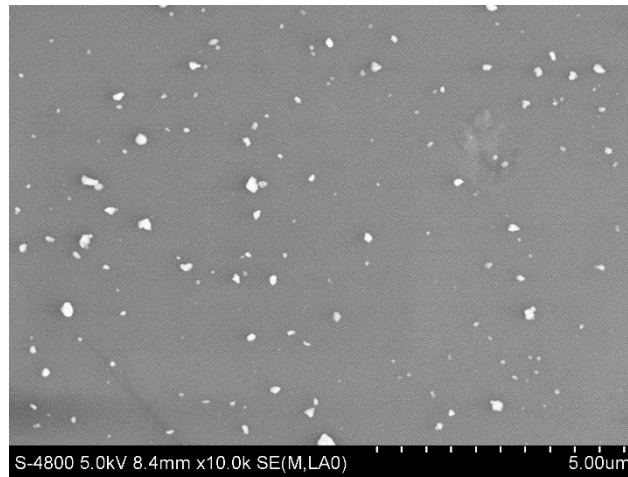


Fig. S1 The SEM image of $\text{CaTiO}_3:\text{Pr}^{3+}@\text{SiO}_2$.

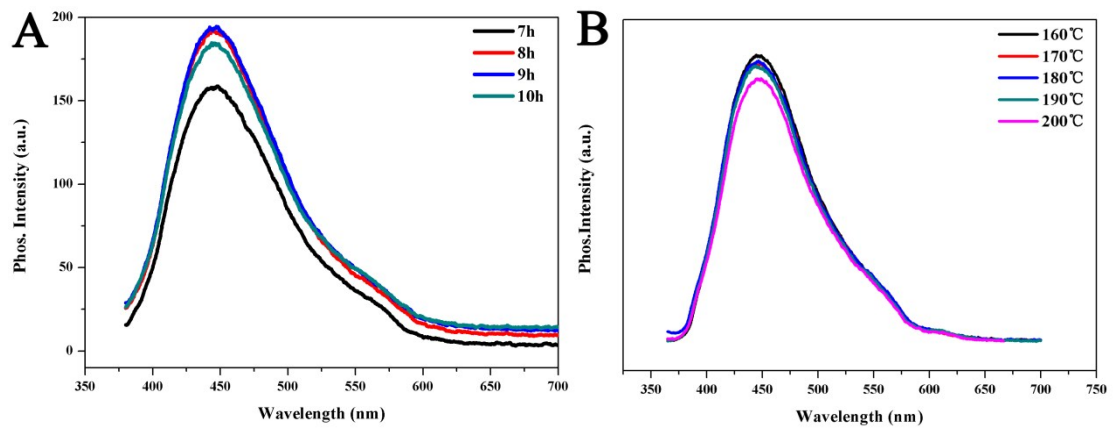


Fig.S2 The optimization of reaction time (A) and reaction temperature (B) of CDs.

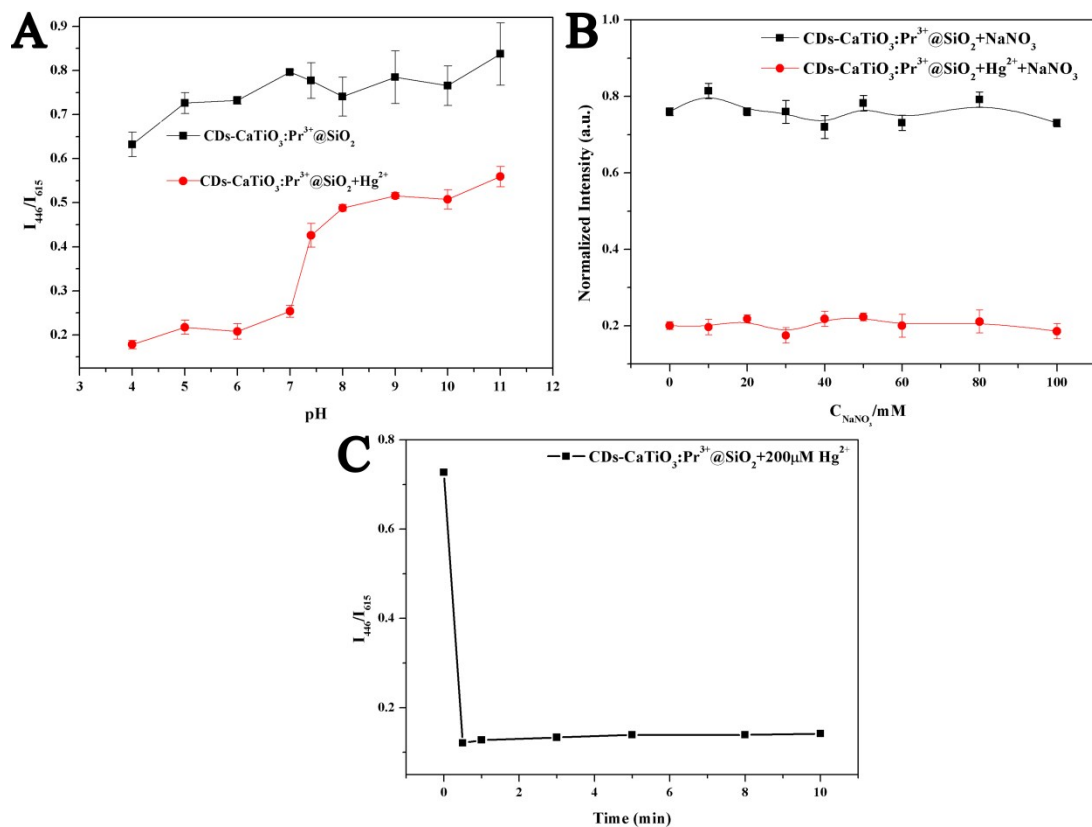


Fig.S3 Effect of pH on the ratio phosphorescence intensity (I_{446}/I_{615}) of CDs-CaTiO₃:Pr³⁺@SiO₂ with HEPES buffer solution (10 mM, pH 7.0) in the absence (black line) and presence of Hg²⁺ (200 μM) (red line) (A) and the ratio phosphorescence intensity (I_{446}/I_{615}) in the absence and presence of Hg²⁺(200 μM) under different ionic strengths (NaNO₃) (B); The ratio phosphorescence intensity (I_{446}/I_{615}) of CDs-CaTiO₃:Pr³⁺@SiO₂ in the presence of Hg²⁺ (200 μM) versus incubation time (C).

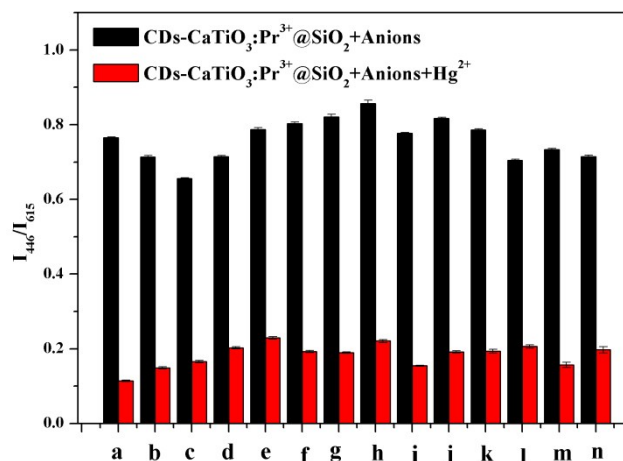


Fig.S4 Phosphorescence responses of CDs-CaTiO₃:Pr³⁺@SiO₂ upon the addition of different anions (200 μM) (a) Blank (b) F⁻ (c) Cl⁻ (d) Br⁻ (e) I⁻ (f) C₂O₄²⁻ (g) CO₃²⁻ (h) HPO₄²⁻ (i) H₂PO₄⁻ (j) PO₄³⁻ (k) NO₃⁻ (l) S²⁻ (m) SO₃²⁻ (n) PPA in absence and presence of Hg²⁺ (200 μM) (λ_{ex} =345 nm), where I_{446} and I_{615} refer to the phosphorescence intensity of CDs and CaTiO₃:Pr³⁺@SiO₂, respectively.

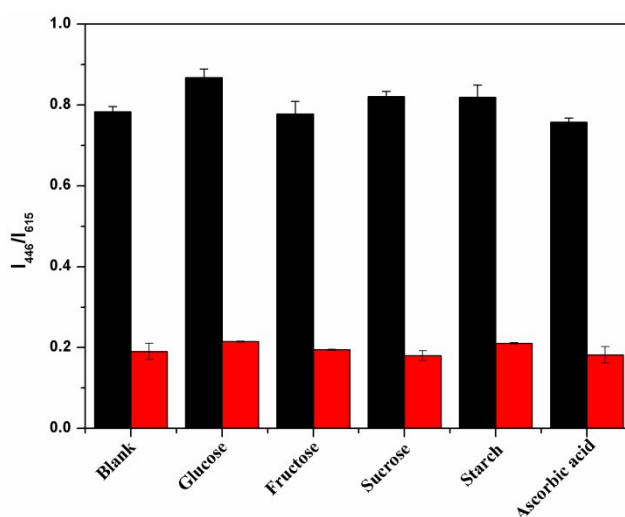


Fig.S5 Phosphorescence responses of CDs-CaTiO₃:Pr³⁺@SiO₂ upon the addition of glucose, fructose, sucrose, starch and ascorbic acid (200 μM) in the absence and presence of Hg²⁺ (200 μM).

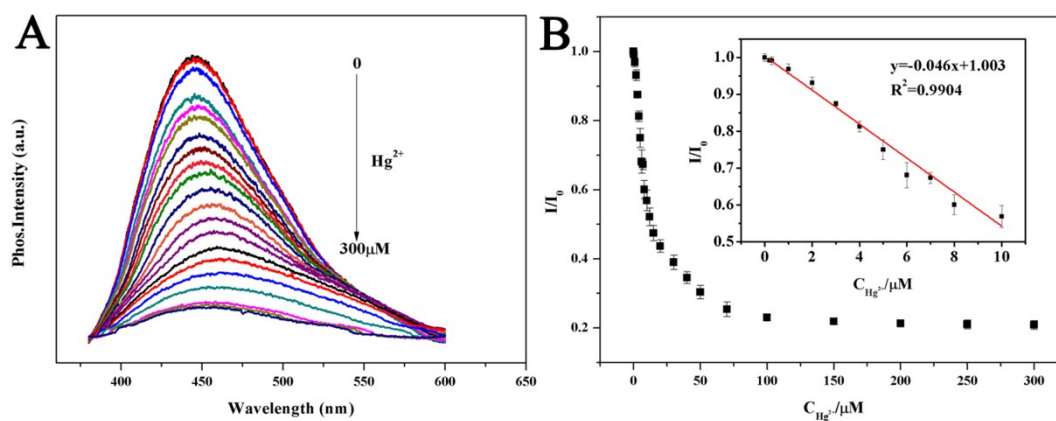


Fig.S6 The phosphorescence spectra of CDs in the presence of different Hg^{2+} concentrations from 0 to 300 μM in HEPES buffer solution (10 mM, pH 7.0) (A); Plot of I/I_0 versus the concentration of Hg^{2+} from 0 to 300 μM (where I_0 refers to the phosphorescence intensity of CDs and I is the phosphorescence intensity of CDs with the addition of various Hg^{2+} concentrations) (B).

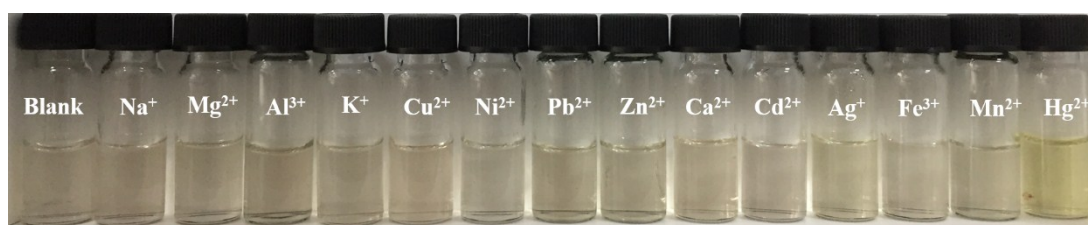


Fig. S7 Images of CDs were taken with addition of various metal ions under sunlight.

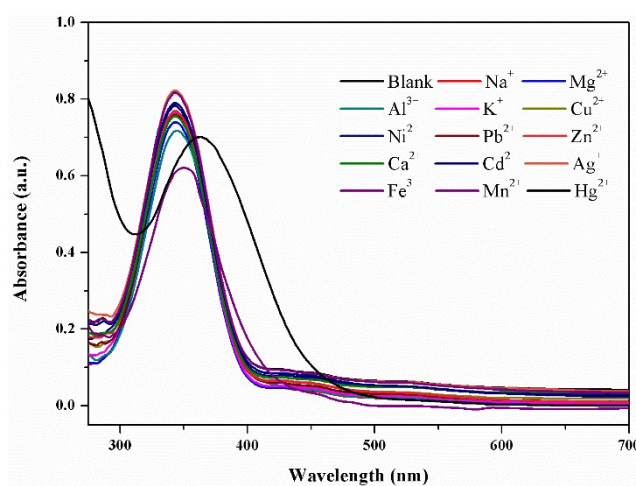


Fig. S8 The UV-Vis absorption spectra of CDs in the presence of different metal ions.

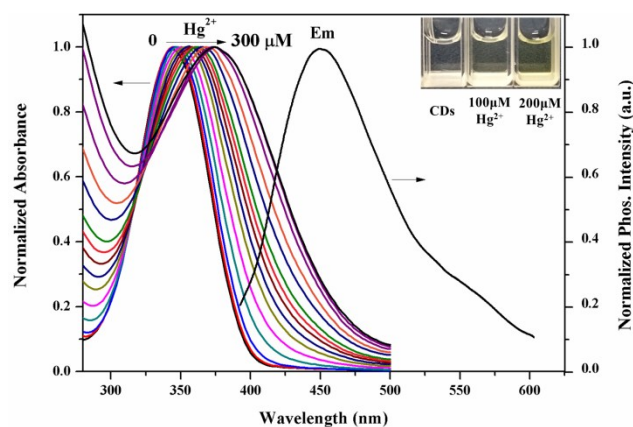


Fig.S9 The absorption spectra of CDs in the presence of various Hg^{2+} concentrations and the emission spectrum of CDs under the excitation wavelength of 345 nm.

Table S1. Comparison of different analysis materials for the detection of Hg^{2+} in water samples

Analysis Material	Analytes	Linear range (μM)	Detection limit (nM)	Real Sample	Detection methods	Reference
$\text{Eu}^{3+}/\text{CDs@MOF-253}$	Hg^{2+}	0.065–150	0.065	River sample Fountain water	Ratiometric fluorescence	1
$\text{GO-PPV@MSN-NH}_2\text{@SRh6G}$	Hg^{2+}	0.060-0.18	71	Tap water —	Fluorescence	2
Carbon nanoparticles -Rhodamine B	Hg^{2+}	0–6	42	—	Ratiometric fluorescence	3
Dual-emission Carbon dot	Hg^{2+}	0-40	9.0	serum samples River water	Ratiometric fluorescence	4
CDs- Rhodamine B	Hg^{2+}	0.5-10	25	—	Ratiometric fluorescence	5
BSA-Ag NCs	Hg^{2+}	0.05-25	48.7	—	Ratiometric fluorescence	6
Pyrene-functionalized magnetic nanoparticles	Hg^{2+}	4.0 - 40	50	Tap water Pond water Human Serum	Fluorescence	7
CDs-Ag NCs	Hg^{2+}	0-0.5	28	Lake water Tap water Mineral water	Ratiometric fluorescence	8
CDs-CdSe@ZnS QDs	Hg^{2+}	0.2-2.0	100	Tap water Lake water	Ratiometric fluorescence	9
CDs- $\text{CaTiO}_3\text{:Pr}^{3+}$ @ SiO_2	Hg^{2+}	0.07-10	9.65	Tap water Pond water Human Serum	Ratiometric phosphorescence	This work

References

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