

Supporting Information

Facile one-step synthesis of highly luminescent N-doped carbon dots as efficient fluorescence probe for chromium (VI) detection based on inner filter effect

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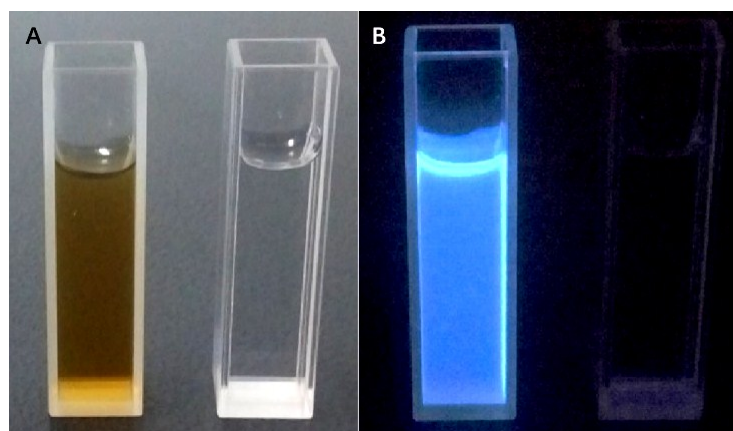


Figure S1 As-prepared reaction mixtures (left) and the starting materials (right) under irradiation with visible (A) and UV light (B).

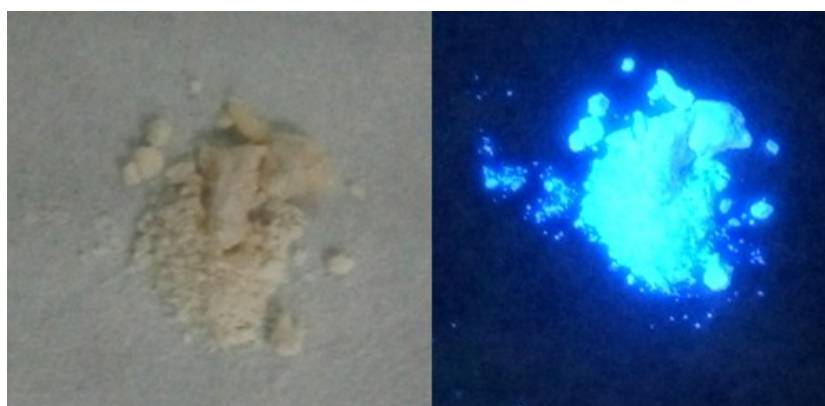


Figure S2. Photographs of the CG-CDs powder under irradiation with visible (left) and UV light (right).

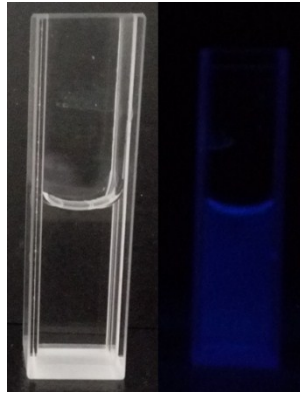


Figure S3. Photograph of the CG-CDs aqueous solution under irradiation with visible (left) and UV light (right)

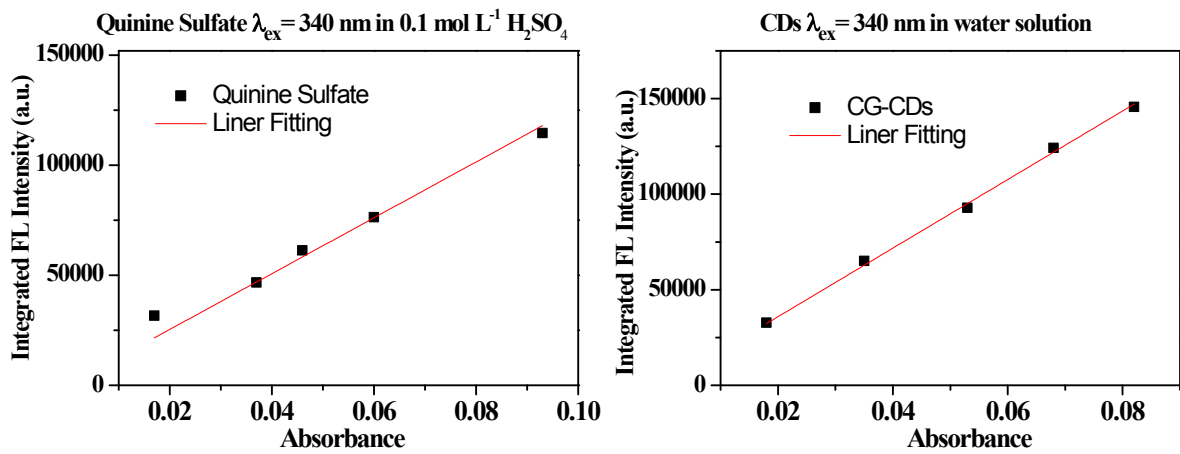


Figure S4. Plots of integrated fluorescence (FL) intensity of quinine sulfate (referenced dye) and CG-CDs synthesized from citric acid and glycine as a function of optical absorbance at 340 nm.

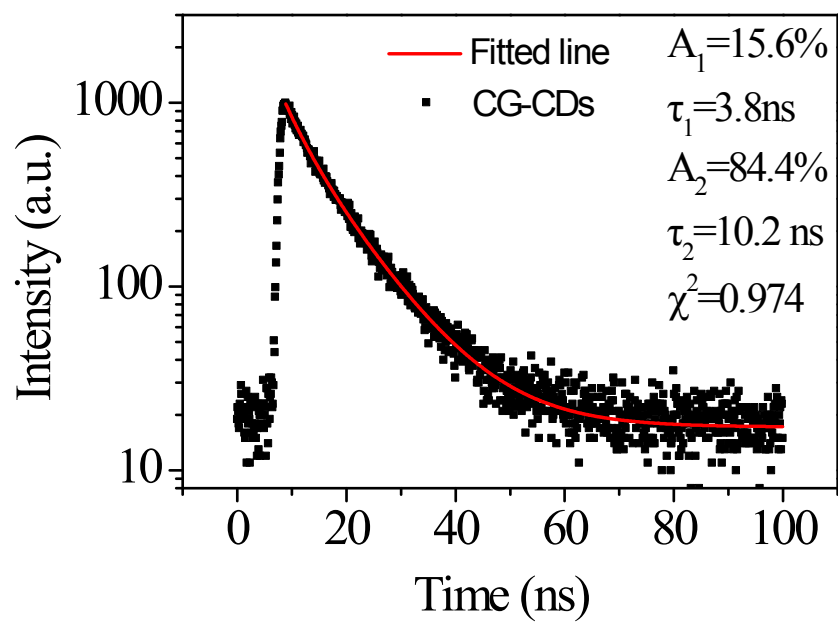


Figure S5 Fluorescence decay spectra and fitting curves of CG-CDs.

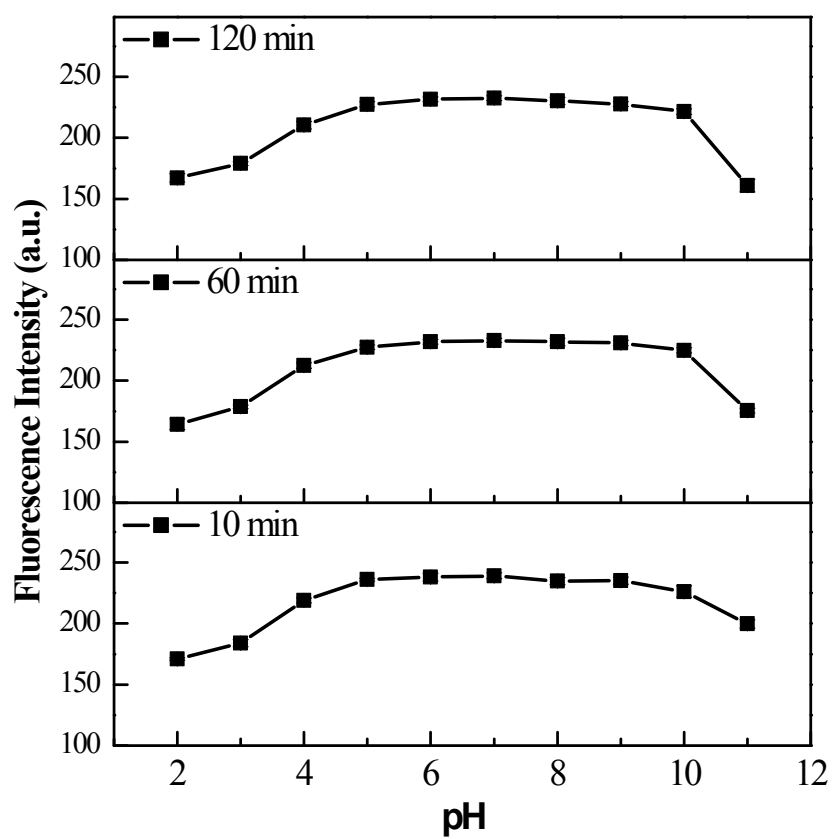


Figure S6. Effects of pH value on the fluorescence intensity of CG-CDs within 120 min.

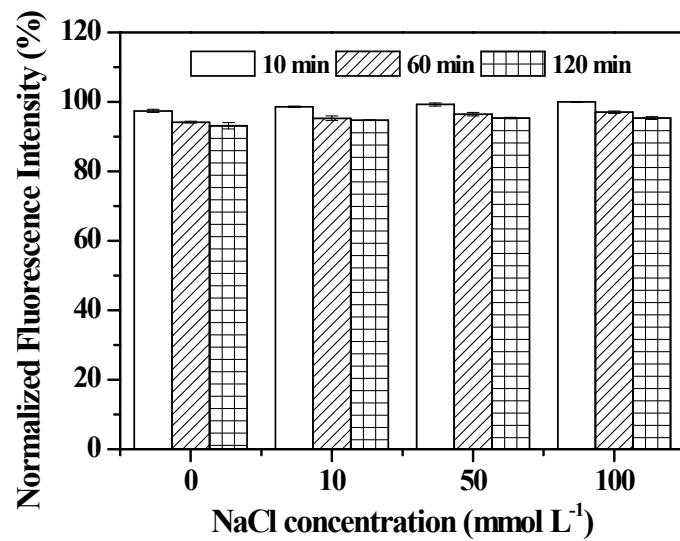


Figure S7. Effects of ionic strength of NaCl on the fluorescence intensity of CG-CDs within 120 min.

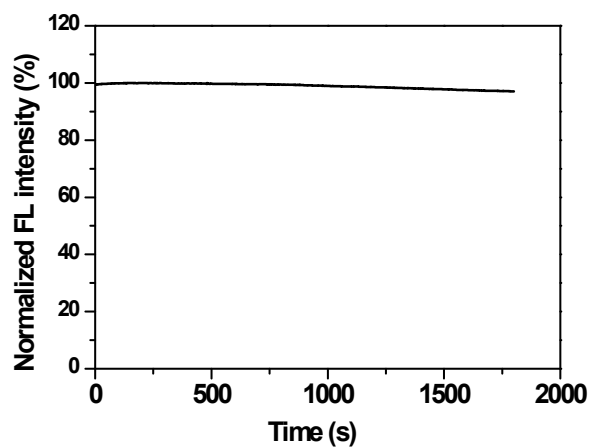


Figure S8. Photostability of fluorescence intensity for CG-CDs under continuous excitation at 340 nm for 30 min.

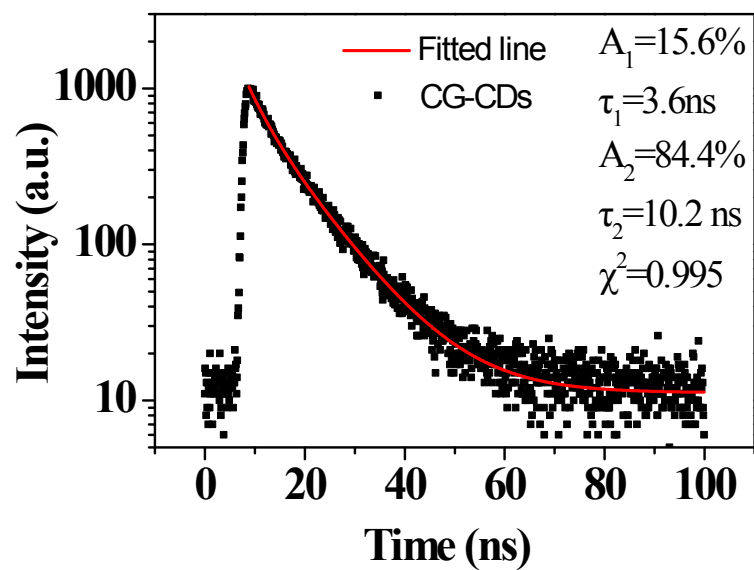


Figure S9. Fluorescence decay and fitting curves of CG-CDs in the presence of Cr (VI) (200 μM).

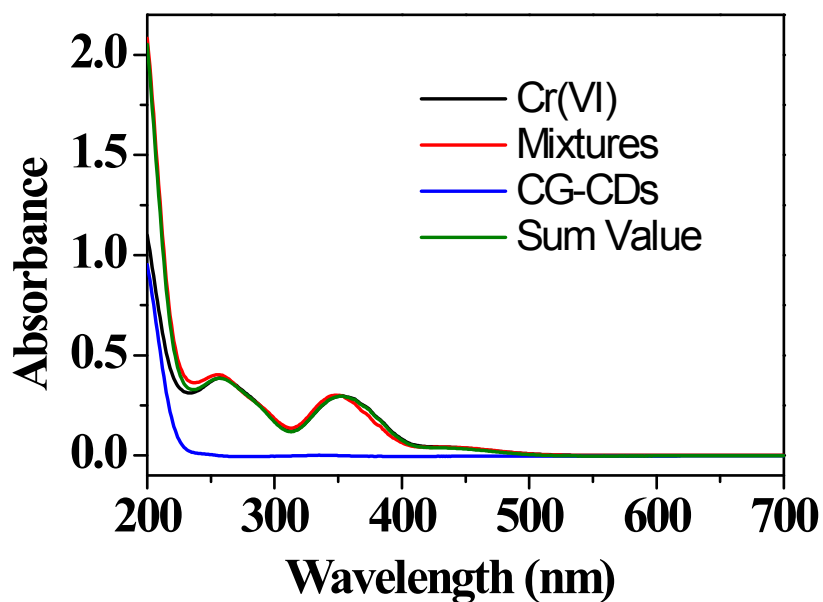


Figure S10. The UV-vis absorption spectra of CG-CDs, Cr(VI), CG-CDs- Cr(VI) mixtures and the sum value of absorbance of CG-CDs and Cr(VI).

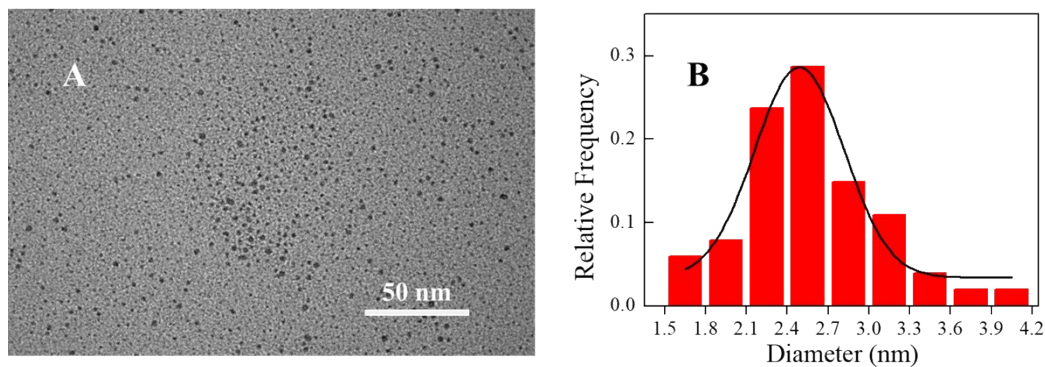


Figure S11. TEM image (A) and size distribution histogram (B) of CG-CDs in present of chromium (VI) ($100 \mu\text{mol L}^{-1}$), the average size was 2.6 nm, which is similar to the pure CG-CDs.

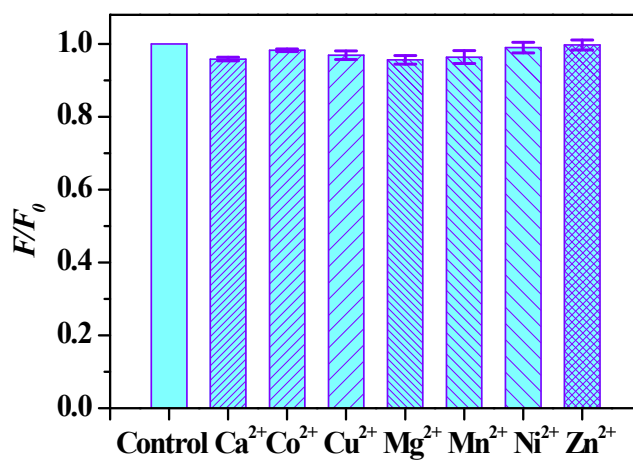


Figure S12. Fluorescence intensity response of CG-CDs in the presence of 200 μM solution of various metal ions. The blank represents the fluorescence response of the solution of CG-CDs without adding any metal ion. The concentration of each kind of ion was 100 μM .

Table S1. Parameters for QY calculation

| | Quinine Sulfate | | | | | CDs | | | | |
|---------------|----------------------|-------|-------|-------|--------|----------------------|-------|-------|--------|--------|
| Abs | 0.017 | 0.037 | 0.046 | 0.06 | 0.093 | 0.018 | 0.035 | 0.053 | 0.068 | 0.082 |
| Integrated FL | 31662 | 46722 | 61250 | 76330 | 114662 | 32756 | 65095 | 92882 | 124230 | 145645 |
| lope | 1.27×10 ⁶ | | | | | 1.79×10 ⁶ | | | | |
| QY | 55% | | | | | 78% | | | | |

FL: fluorescence; **QY:** quantum yield.

Table S2. The comparison of the determination of Cr(VI)

| Methods | Time consumption | Linear detection range | Detection limit | Reference |
|---|------------------|--|---|-----------|
| Fluorescence method with carbon dot | 1 min | 0.01-50 μM | / | [1] |
| Electrochemical detection based on gold nanoparticles | / | 0.1-105 μM | 0.03 μM | [2] |
| Colorimetric method with gold nanoparticle | 30 min | 0.1-20 μM | 0.088 μM | [3] |
| Fluorescence method with AIQx modified SBA-15 | / | 0.32-5.8 μM | 7.7 nM | [4] |
| Colorimetric method with Ce(VI) and 1,5-DPC modified Paper microfluidic devices | 10 min | 0.23-3.75 μg (4.42-7.2 μmol) | 0.12 μg (2.31 μmol) | [5] |
| Colorimetric method with silver nanoparticle | / | 1 nM-1 mM | 1 nM | [6] |
| Fluorescence method with graphitic carbon nitride nanosheets | 10 min | 0.6-300 μM | 0.15 μM | [7] |
| boron and nitrogen co-doped carbon dots | 1 min | 1.39-260 μM | 0.28 μM | [8] |
| nitrogen and sulfur co-doped carbon dots | 10 min | 2-160 μM | 1.72 μM | [9] |

References

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