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# Integration detection of mercury(II) and GSH with a fluorescent “on-off-on” switch sensor based on nitrogen, sulfur co-doped carbon dots†

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Using aurine and citric acid as precursors, we have synthesized stable blue-fluorescent nitrogen and sulfur co-doped carbon dots (NS-CDs), with a high quantum yield of up to 68.94% via a thermal lysis method. The fluorescent NS-CDs were employed as a sensitive sensor for the integration detection of Hg<sup>2+</sup> and glutathione (GSH). This was attributed to Hg<sup>2+</sup> effectively quenching the fluorescence of the NS-CDs by static quenching, and then GSH was able to recover the fluorescence owing to the stronger binding between Hg<sup>2+</sup> and the sulfhydryl of GSH. Based on the “on-off-on” tactic, the detection limits of Hg<sup>2+</sup> ions and GSH were 50 nM and 67 nM respectively. The fluorescence sensor was successfully applied to detect Hg<sup>2+</sup> ions and GSH in actual samples (tap water and fetal bovine serum). Furthermore, we have proved that the sensor had good reversibility. Overall, our NS-CDs can serve as effective sensors for environmental and biological analysis in the future.

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## 1. Introduction

As an important zero-dimensional carbon-based nanometer material, carbon dots (CDs) have been studied since 2004 (ref. 1–4) and widely used in biological analysis,<sup>5,6</sup> sensing,<sup>7–9</sup> catalysis,<sup>10,11</sup> bioimaging<sup>12</sup> and photothermal therapy<sup>13–15</sup> and biomedicine.<sup>16</sup> They have various properties, including stable fluorescence photoluminescence,<sup>17,18</sup> low toxicity,<sup>19</sup> excellent biocompatibility,<sup>20</sup> good water solubility<sup>21,22</sup> and low-cost synthesis.<sup>23</sup>

However, the applications of CDs are limited by their low quantum yield (QY) in conventional carbon sources. To increase QY, a hetero atom doping method has been utilized, in which nitrogen (N) and sulfur (S) are the commonest doped elements used to improve the fluorescence performance of CDs and provide targeting properties for special functional groups.<sup>24,25</sup>

In recent years, heavy metals have been considered as hazardous chemicals in the environment. Hg<sup>2+</sup> ions, as one of the most stable inorganic forms of mercury, are known to be highly toxic because of their easy bioaccumulation and inactive sulfhydryl group. Glutathione (GSH) is one of the most ubiquitous low-molecular-weight (LMW) thiol in living cells<sup>26,27</sup> and can cause various physiological dysregulations with an anomalous

concentration. Therefore, GSH<sup>28</sup> has been commonly used as a diagnostic biomarker for multiple diseases. Especially, it is very important to monitor the level of GSH in living cells considering the high GSH concentration in cancer cells.<sup>29,30</sup>

To monitor Hg<sup>2+</sup> ions<sup>31,32</sup> and GSH,<sup>33,34</sup> numerous analytical methods have been developed, including AAS,<sup>35</sup> electrochemical,<sup>36</sup> electrophoresis,<sup>37</sup> HPLC,<sup>38</sup> mass spectrometry,<sup>39</sup> and *etc.*<sup>40</sup> However, most of them are laborious, intricate, and in need of professional operating and expensive instruments. Fluorescence probes providing a better choice of Hg<sup>2+</sup> ions and GSH detection in living cells, tissues, even animals due to the exclusive advantages of high sensitivity,<sup>41,42</sup> simple controls, convenient testing/monitoring and quick response, *etc.* However, most of the conventional fluorescent probes such as small organic fluorescent molecule, and semiconductor quantum dots are relatively toxic and poorly water-soluble, and greatly limit the detection of Hg<sup>2+</sup> ions and GSH in bioassay. Accordingly, CDs as a novel fluorescent probe with excellent biocompatibility has attracted increasing attention.

In this study, we have successfully prepared CDs with citric acid and taurine, with its QY up to 68.94%. With the addition of Hg<sup>2+</sup> ions, the fluorescence of NS-CDs can be quenched efficiently, and the fluorescence was recovered while GSH was added. Moreover, NS-CDs have been tested in actual sample (tap water and fetal bovine serum) to detect Hg<sup>2+</sup> ions and GSH, showing their good selectivity and sensitivity.

## 2. Results and discussion

### 2.1. Characterization of NS-CDs

As shown in Fig. 1a, NS-CDs are well dispersed and have nearly spherical shape. The particle sizes of NS-CDs are mainly

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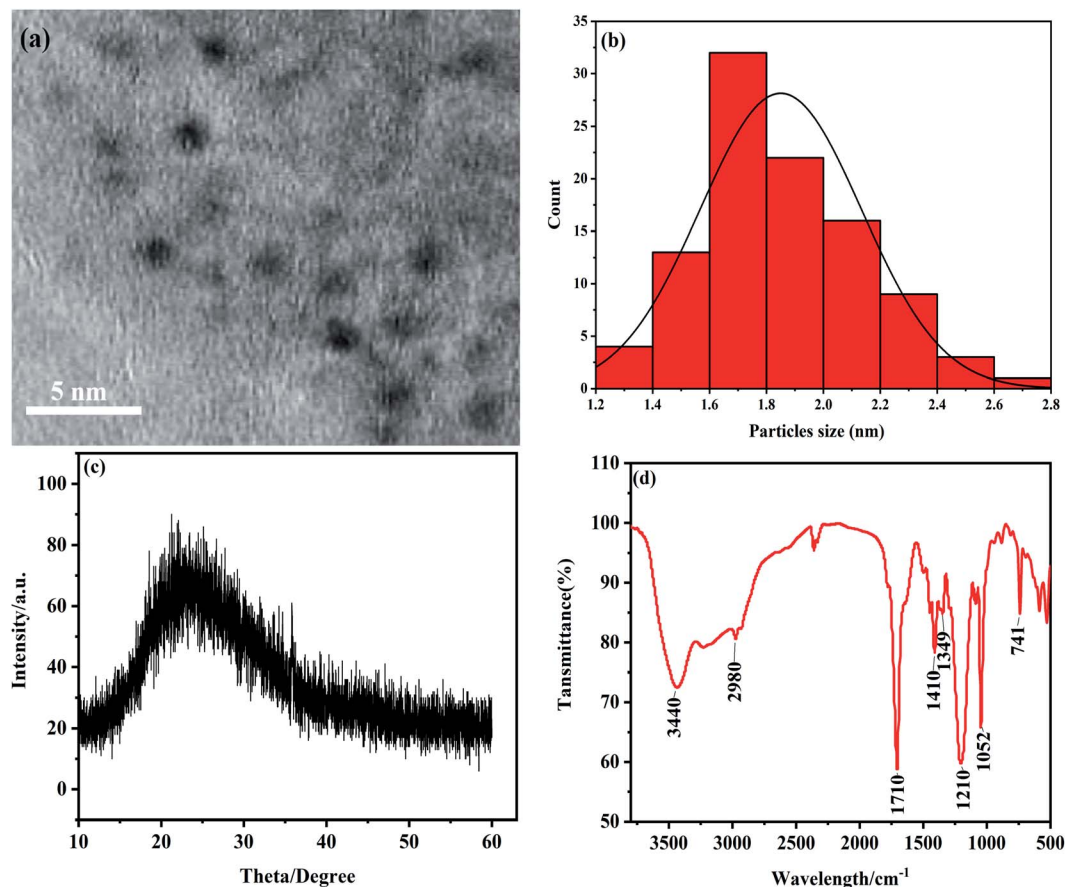


Fig. 1 (a) TEM of the NS-CDs. (b) Histograms showing the particle size distributions of NS-CDs. (c) XRD pattern of the NS-CDs. (d) FTIR spectrum of NS-CDs.

distributed in a narrow range from 1.2 nm to 2.8 nm, with an average particle diameter of 1.7 nm (Fig. 1b). The XRD pattern result indicates a broad peak at  $2\theta = 23^\circ$ , suggesting its attribution to the amorphous structure. Then, we have examined the surface groups on the NS-CDs. The FTIR results shows several strong absorption peaks at  $2500\text{--}3440\text{ cm}^{-1}$ ,  $1710\text{ cm}^{-1}$  and  $1210\text{ cm}^{-1}$ ,  $1349\text{ cm}^{-1}$ ,  $1410\text{ cm}^{-1}$ ,  $1052\text{ cm}^{-1}$ , and  $741\text{ cm}^{-1}$ , which are attributed to O-H groups, C=O and C-O, S=O bonds, C-N bond, O=S=O bonds, and NH bonds, respectively (Fig. 1d). Overall, this suggests NS-CDs have high solubility in water because of their carboxyl, hydroxyl, and amino functional groups on surface.

To further analyze the composed elements of NS-CDs, we have performed XPS analysis, and the full XPS survey spectrum shows the elementary analysis of sulfur (S2p, 169.08 eV), carbon (C1s, 286.08 eV), nitrogen (N1s, 401.08 eV), and oxygen (O1s, 532.08 eV) (Fig. 2a). This indicated NS-CDs includes four elements, C, O, N, and S, with a content of 47.7 wt%, 32.03 wt%, 8.6 wt% and 8.06 wt%, respectively. To reveal the detailed bonds, we have conducted XPS spectrum experiments for each element. C1s spectrum displays four distinct peaks at 288.28 eV (C=O), 286.58 eV (C-S), 285.78 eV (C-O/C-C) and 284.78 eV (C-N) (Fig. 2b). O1s spectrum reveals three O forms, S=O (533.68 eV), C-O (533.18 eV) and C=O (531.68 eV) (Fig. 2c). N1s

spectrum shows a peak at 400.40 eV, which is produced by N-H group (Fig. 2d). S2p spectrum reveals three peaks at 168.78 eV, 167.68 eV and 167.08 eV, which are attributed to O=S=O group, C-S group and S=O group, respectively (Fig. 2e). Hence, we concluded that N and S elements are on the surface of NS-CDs, which is consistent with our FTIR measurements data (Fig. 1d). Both XPS spectra and FTIR spectra suggests the hydrophilicity and stability of NS-CDs due to their functional hydroxyl, amide, amino, and carboxyl/carbonyl groups.

## 2.2. Optical properties of NS-CDs

Next, we determined the optical properties of NS-CDs. NS-CDs display an absorption peak at 332 nm because of the absorption of connected carbonyl/amine group<sup>43</sup> with its maximum excitation wavelength and emission at 345 nm and 443 nm, respectively (Fig. 3a). The aqueous solution of NS-CDs displays colorless under sunlight but emits blue fluorescence while under UV lamp (Fig. 3a inset). The CIE color coordinate of NS-CDs is (0.1545, 0.1551), further indicating NS-CDs show blue fluorescence with UV radiation (Fig. 3c). With the excitation wavelength of NS-CDs ranging from 325 nm to 385 nm, their emission peak shifts from 439 nm to 446 nm, showing redshift (Fig. 3b). This suggests that N and S doping elements contribute to a surface condition and uniform size distribution of NS-

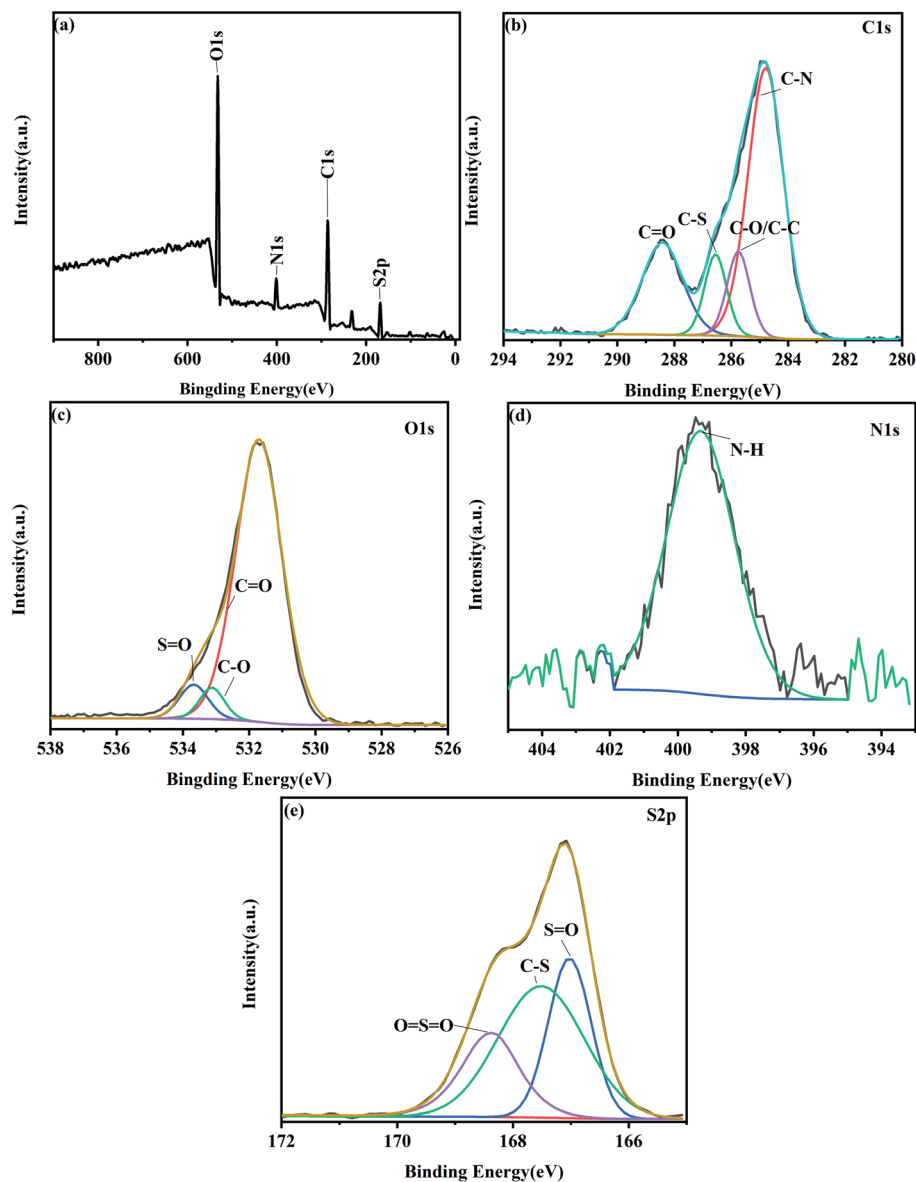


Fig. 2 (a) XPS survey of the NS-CDs. (b) XPS spectrum of C1s. (c) XPS spectrum of O1s. (d) XPS spectrum of N1s. (e) XPS spectrum of S2p.

CDs.<sup>44</sup> 2D-emission plots show that a photoluminescence center could be observed at the emission wavelength of 443 nm and excitation wavelength of 345 nm (Fig. 3d). In addition, the fluorescence QY of NS-CDs was calculated to be 68.94%.

Furthermore, ionic strength, illumination, and pH value were examined to assess the fluorescence stability of NS-CDs. With increasing the concentration of sodium chloride (from 0 to 4 M), there was no obviously change on the fluorescence intensity of NS-CDs, suggesting their excellent tolerance in ion solutions (Fig. S1†). In addition, the fluorescence intensity of NS-CDs remains unaltered under the UV light irradiation for at least 90 min, showing their stability under UV light (Fig. S2†). The fluorescence intensity of NS-CDs gradually increases when pH value increases from 1 to 7, and decreases when pH value raised from 7 to 13, with the maximum intensity at pH value of 7 (Fig. S3†). In general, we have confirmed that NS-CDs have

a stable and excellent optical performance even under extreme environmental conditions.

### 2.3. Detection of Hg<sup>2+</sup> and GSH

To evaluate their selectivity of specific metal ions, we investigated the fluorescence response of NS-CDs to common metal ions. Compared with other ions, Hg<sup>2+</sup> ions have a stronger interaction with NS-CDs, with outstanding fluorescence quenching (Fig. 4a).<sup>45,46</sup> When possible interference ions coexists, the addition of coexisted ions exert no effect on the fluorescence quenching caused by the Hg<sup>2+</sup> ions, further indicating a high selectivity of NS-CDs for Hg<sup>2+</sup> ions detection even in the presence of possible interference ions (Fig. 4b).

The fluorescence spectra were recorded with adding Hg<sup>2+</sup> ions of various concentrations into the NS-CDs solution. The fluorescence intensity drops quickly when the concentration of

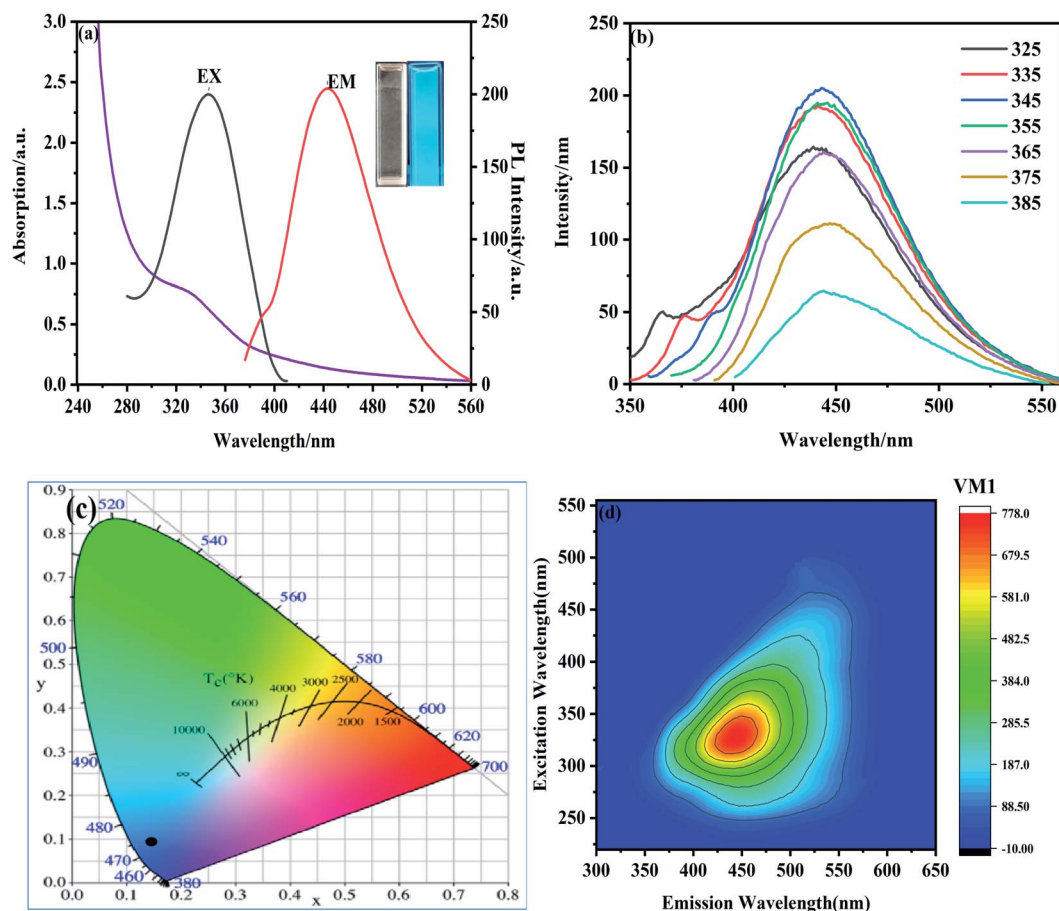


Fig. 3 (a) UV-vis absorption spectrum, the excitation of the NS-CD, the emission of the NS-CDs. Inset: photographs of the NS-CDs under white light (left) and UV light at 365 nm (right). (b) Fluorescence spectra of NS-CDs under different excitation wavelengths. (c) CIE diagram of NS-CDs. (d) Excitation-emission map of NS-CDs.

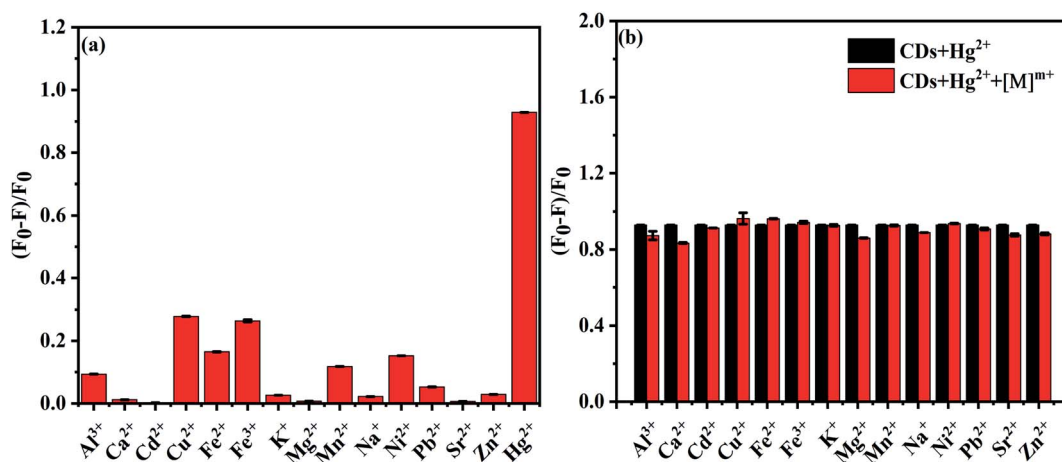


Fig. 4 (a) Selectivity of the NS-CDs to different metal ions (30  $\mu\text{M}$ ). (b) Bar diagram of fluorescence intensity of NS-CDs to  $\text{Hg}^{2+}$  (30  $\mu\text{M}$ ) with or without interfering metal ions (30  $\mu\text{M}$ ).

$\text{Hg}^{2+}$  increases from 0  $\mu\text{M}$  to 100  $\mu\text{M}$  (Fig. 5a). The relationship between the relative fluorescence intensity ( $(F_0 - F)/F_0$ ) of NS-CDs and the concentration of  $\text{Hg}^{2+}$  ions is shown in Fig. 5b. When the concentrations of  $\text{Hg}^{2+}$  ions range from 0–10  $\mu\text{M}$ ,

there is a strong linear correlation, with the relative coefficient ( $R^2$ ) of 0.9932. The fitted linear equation is  $(F_0 - F)/F_0 = 0.0776c(\text{Hg}^{2+}) - 0.4405$ , where  $F$  and  $F_0$  are the fluorescence intensities of NS-CDs with or without adding  $\text{Hg}^{2+}$  ions, respectively.

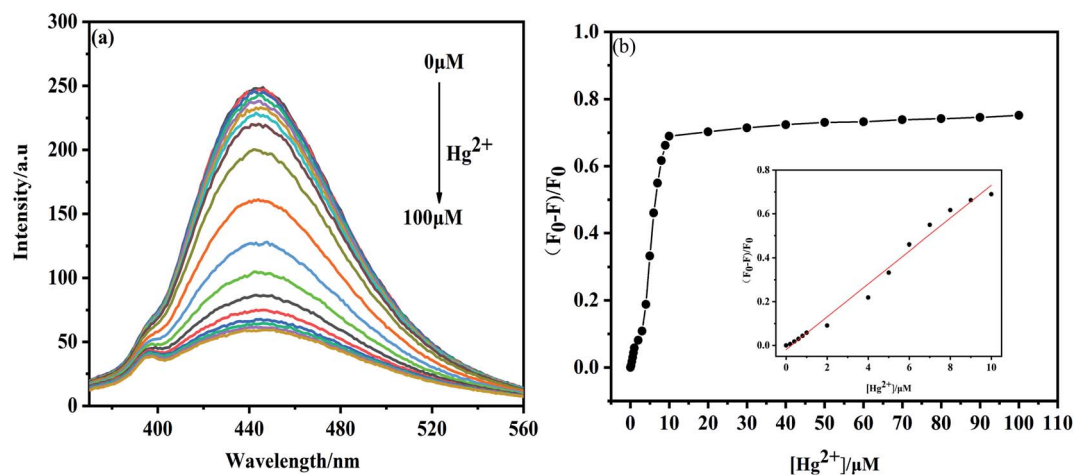


Fig. 5 (a) Fluorescence spectra of NS-CDs after addition of different concentrations of  $\text{Hg}^{2+}$ . (b) Relation of the  $(F_0 - F)/F_0$  and  $\text{Hg}^{2+}$  concentration (0  $\mu\text{M}$  to 100  $\mu\text{M}$ ). Insert: linear relationship of the  $(F_0 - F)/F_0$  and  $\text{Hg}^{2+}$  concentration (0  $\mu\text{M}$  to 10  $\mu\text{M}$ ).

Table 1 Comparison of different probes for the sensing of  $\text{Hg}^{2+}$

Precursor	Probes	Linear range ( $\mu\text{M}$ )	LOD (nM)	References
Cucumber juice	N/S/P-CD	1–70	180	47
Folic acid	N-CQDs	0–75	230	48
CA and APTES	Si/N-CDs	0.5–30	140	49
Citric acid, ethylenediamine, and methyl blue	NSCDs	0.7–15	540	50
Citric acid and 2,6-diaminopyridine	N-CQDs	0.2–1.2	85	51
Citric acid and polyacrylamide	CDs-AgNPs	0.01–2.5	3.6	52
Gadenia fruits	N/S-CDs	0.2–20	320	32
L-Cysteine and citric acid	S,N-CDs/AgNPs	$1.5 \times 10^{-3}$ to 2.0	0.51	53
Citric acid and urea	N-CDs@NaFZ	0.1–1.5	$5.5 \times 10^{-3}$	54
Citric acid and taurine	NS-CDs	0–100	50	This work

According to  $3\delta/k$ , we have determined the detection limit of  $\text{Hg}^{2+}$  ions at 50 nM. Table 1 shows the linear range and LOD of  $\text{Hg}^{2+}$  are compared with previously literature values.

When we examined the selectivity of the NS-CDs- $\text{Hg}^{2+}$  system to different amino acids, other small biomolecules cannot or little alter the fluorescence, except for GSH (Fig. 6a). The quenched fluorescence is only recovered when GSH is

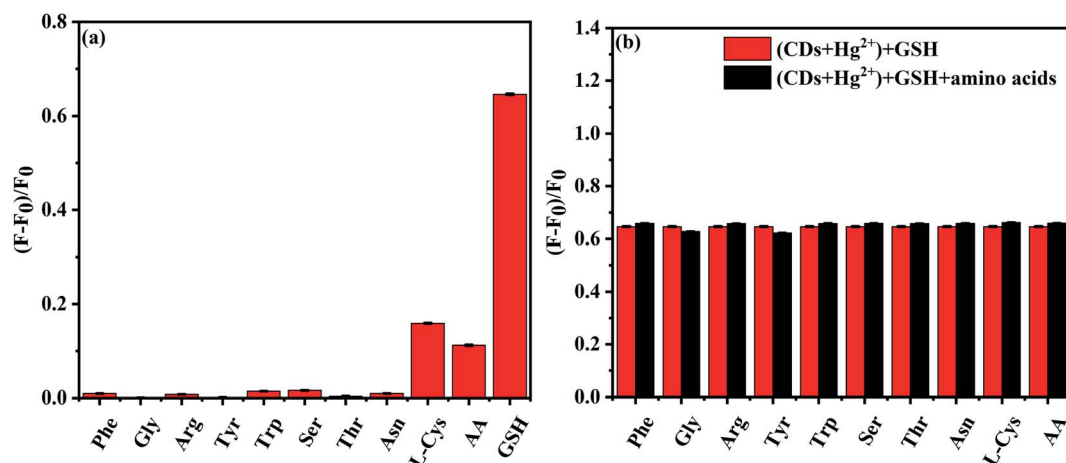


Fig. 6 (a) Selectivity of the NS-CDs- $\text{Hg}^{2+}$  system to different biomolecules (30  $\mu\text{M}$ ). (b) Bar diagram of intensity of NS-CDs- $\text{Hg}^{2+}$  system to GSH (30  $\mu\text{M}$ ) with or without interfering biomolecules (30  $\mu\text{M}$ ).



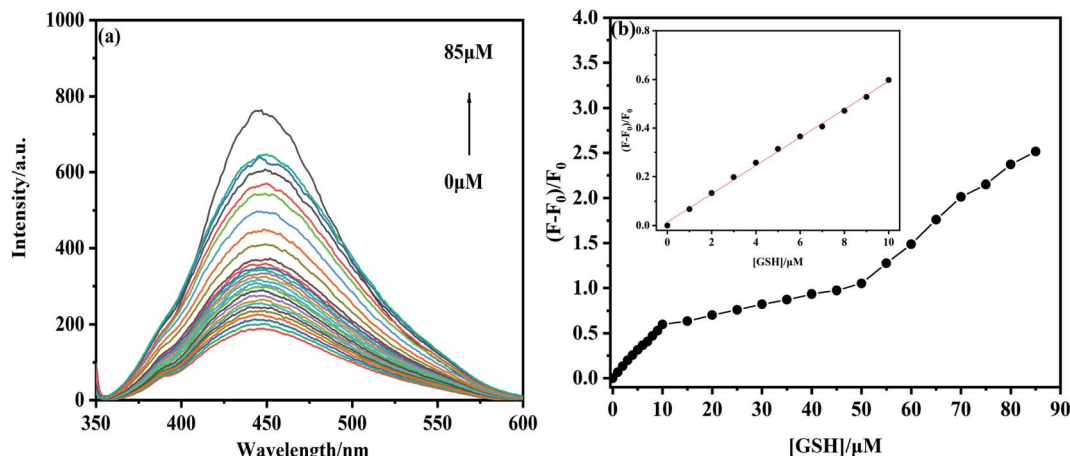


Fig. 7 (a) Fluorescence spectra of NS-CDs-Hg<sup>2+</sup> after addition of different concentrations of GSH. (b) Relation of the  $(F - F_0)/F_0$  and GSH concentration (0  $\mu\text{M}$  to 85  $\mu\text{M}$ ). Inset: linear relation of the  $(F - F_0)/F_0$  and GSH concentration (0  $\mu\text{M}$  to 10  $\mu\text{M}$ ).

Table 2 Comparison of different fluorescence sensors to GSH

Precursor	Probes	Linear range ( $\mu\text{M}$ )	LOD (nM)	References
Boric acid and ethylenediamine	B,N-CDs/MnO <sub>2</sub>	0–90	320	55
O-Phenylenediamine and citric acid	CQDs	1–80	300	56
Eggshell membrane ashes	C-Dots	$5 \times 10^2$ to $8 \times 10^4$	480	57
Citric acid monohydrate and L-cysteine	S-CDs/AuNPs	0.01–5.0	3.6	58
Citric acid and taurine	NS-CDs-Hg <sup>2+</sup>	0–85	67	This work

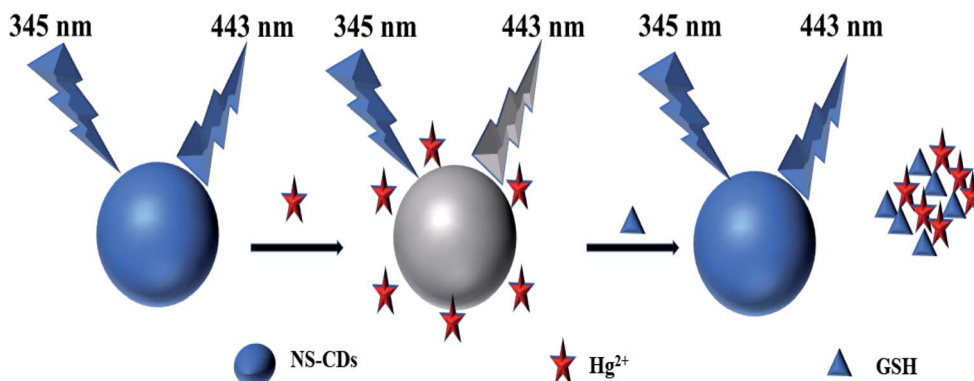
added into the NS-CDs-Hg<sup>2+</sup> system. Consequently, the NS-CDs-Hg<sup>2+</sup> system could be served as a turn-on sensor to detect GSH specifically. When other biomolecules were mixed with GSH, the effect of coexistent biomolecules can be neglected, suggesting the specificity of the NS-CDs-Hg<sup>2+</sup> system to GSH detection (Fig. 6b).

With GSH concentration increased, the fluorescence intensity of NS-CDs-Hg<sup>2+</sup> is gradually returned (Fig. 7a). Then, we calculated the relation between the relative fluorescence intensity  $((F - F_0)/F_0)$  of NS-CDs-Hg<sup>2+</sup> and the concentration of GSH. The relative intensity of NS-CDs-Hg<sup>2+</sup> shows a strong linear correlation to the concentration of GSH in the range of 0–10  $\mu\text{M}$ , with the related coefficient ( $R^2$ ) of 0.9932 (Fig. 7b). The

linear fitting equation is  $(F - F_0)/F_0 = 0.05792c(\text{GSH}) + 0.01393$ , where  $F$  and  $F_0$  are the fluorescence intensities of NS-CDs-Hg<sup>2+</sup> with or without adding GSH, respectively. The detection limit of GSH is calculated at 67 nM. The NS-CDs-Hg<sup>2+</sup> as the fluorescence sensor to test GSH is also compared with the other works in Table 2. In conclusion, these results demonstrate the NS-CDs can detect Hg<sup>2+</sup> and GSH at “on-off-on” mode in a sensitive and specific manner.

#### 2.4. Mechanism of the “on-off-on” mode

In this work, the NS-CDs displays a fascinating “on-off-on” experimental phenomena with the gradual addition of Hg<sup>2+</sup> and GSH. Scheme 1 illustrates a proposed sensing mechanism of



Scheme 1 Schematic diagram of Hg<sup>2+</sup> and GSH determinations.

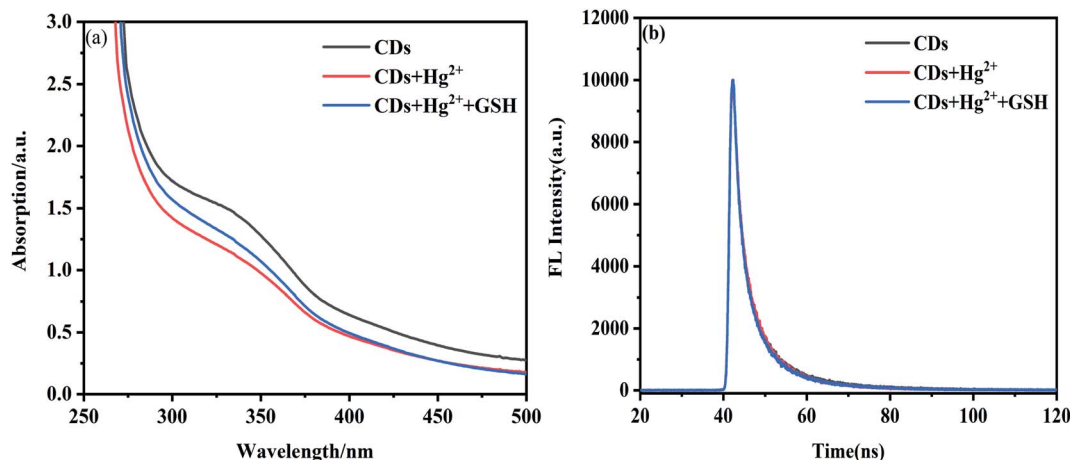


Fig. 8 (a) UV-vis spectra of NS-CDs,  $\text{Hg}^{2+}$ , and GSH system. (b) Fluorescence decay curves of the NS-CDs, NS-CDs- $\text{Hg}^{2+}$ , and NS-CDs- $\text{Hg}^{2+}$  + GSH.

NS-CDs. On the one hand, the fluorescence of NS-CDs can be quenched by  $\text{Hg}^{2+}$  because the combination between  $\text{Hg}^{2+}$  and S-containing groups can form the  $\text{Hg}^{2+}$ -S bonds.<sup>59,60</sup> The structure change of NS-CDs- $\text{Hg}^{2+}$  is demonstrated by FTIR spectrometry, compared to the FTIR spectrum of NS-CDs, the characteristic peak of S=O group ( $1349\text{ cm}^{-1}$ ) is disappeared. The result further illustrate the formation of  $\text{Hg}^{2+}$ -S bonds (Fig. S4†). On the other hand, since the sulfhydryl of GSH has a stronger binding interaction with  $\text{Hg}^{2+}$ , it can take away  $\text{Hg}^{2+}$  from the surface of NS-CDs and then recover the fluorescence of NS-CDs.

The fluorescence sensing procedure was demonstrated by fluorescence lifetime and UV-vis experiments. In Fig. 8a, the absorption peak of NS-CDs (345 nm) is weakened after the addition of  $\text{Hg}^{2+}$ , which proves  $\text{Hg}^{2+}$  compounds with NS-CDs. However, the absorption peaks intensity is strengthened with the addition of GSH, proving that  $\text{Hg}^{2+}$  is removed from the surface of NS-CDs.<sup>61</sup>

The measured FL decay curves exhibit a double-exponential function (Fig. 8b). The average excitation lifetime of NS-CDs is

9.89 ns, containing long lifetime of 11.39 ns (50.15%) and short lifetime of 2.25 ns (49.85%). With the addition of  $\text{Hg}^{2+}$  and GSH, there is no change in the average fluorescence lifetime of NS-CDs, indicating the fluorescence quenching mechanism is static quenching.<sup>62,63</sup>

### 2.5. Detection of $\text{Hg}^{2+}$ and GSH in actual samples

To investigate the feasibility and sensitivity of the method, the N-CDs was applied to determine  $\text{Hg}^{2+}$  in tap water. The tap water samples obtained from the laboratory and was not any pretreatment. The recoveries of  $\text{Hg}^{2+}$  are 94.0–105.2 (RSD < 3.4%) in Table S1.† Tap water contains some other elements, we further study the interferences experiment of ions commonly present in tap water (Fig. S5†), the result shows the addition of metal ions did not interfere with the fluorescence intensity of NS-CDs/tap water system, further illustrating the metals does not affect the test results of actual samples.

Future, practical applications of N-CDs- $\text{Hg}^{2+}$  sensor for GSH determination in fetal bovine serum sample were executed. The measurement of GSH in fetal bovine serum is  $0.35\text{ }\mu\text{M}$ , and the recoveries of GSH are 98.4–102.5% (RSD < 2.4%) (Table S2†). All the results indicate that the N-CDs sensor have high precision and accuracy for determining  $\text{Hg}^{2+}$  and GSH in the practical detection.

Reversibility (also known as fatigue resistance) is one of the important indicators for evaluating “on-off-on” fluorescent sensors. In order to study the reversibility of NS-CDs for  $\text{Hg}^{2+}$  and GSH detection,  $\text{Hg}^{2+}$  and GSH were added alternately to the NS-CDs solution (Fig. 9). The results reflects the fluorescent intensity shows an alternating change, and retains good fatigue resistance after 10 cycles, indicating that the fluorescent sensor has good reversibility.

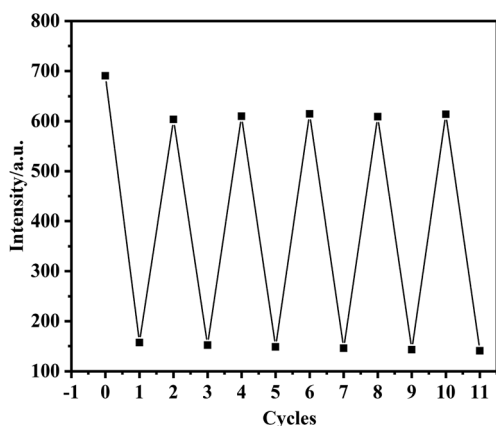


Fig. 9 The reversibility of the NS-CDs by alternately adding  $\text{Hg}^{2+}$  and GSH.

## 3. Conclusions

In summary, NS-CDs based fluorescent “on-off-on” was prepared for detecting  $\text{Hg}^{2+}$  and GSH. The blue emission NS-

CDs showed excellent optical performance, good fluorescent stability, with the fluorescence quantum yield up to 68.94%. The fluorescence of NS-CDs solution was quenched (namely turn-off) when  $\text{Hg}^{2+}$  ions were added and subsequently recovered (turn-on) when GSH was added. These results illustrate that the NS-CDs can be used as fluorescence sensor to detect  $\text{Hg}^{2+}$  ions and GSH through fluorescent “on-off-on” detection mode, with the detection limits as low as 50 nM and 67 nM, respectively. Furthermore, the fluorescent sensor had good reversibility. Consequently, the NS-CDs offered an effective platform for biological analysis and environmental analysis.

## Conflicts of interest

The researcher claims no conflicts of interests.

## Acknowledgements

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