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Green synthesis of fluorescent carbon quantum dots for the detection of mercury (II) and

glutathione

Daraksha Bano, Vijay Kumar, Vikas Kumar Singh, Syed Hadi Hasan*

Nano-Material Research Laboratory, Department of Chemistry, Indian Institute of Technology

(BHU), Varanasi -221005, U.P., India.

*Corresponding author's details:

E-mail; shhasan.apc@itbhu.ac.in, vijuevs@gmail.com

Phone No.: +91-542-6702861

Mobile No.: +91 9839089919

Calculations:

Quantum yield measurement

The quantum yield (QY) is measured by using a quinine sulfate (QS) as a standard having QY equal to 0.54 in 0.1 M H₂SO₄.

$$QY_{x} = QY_{s} \cdot \frac{I_{x}}{I_{s}} \frac{A_{s}}{A_{x}} \cdot \frac{\eta_{x}^{2}}{\eta_{s}^{2}}$$
(1)

Where 'QY' stands for the FL quantum yield, 'I' used for the integrated FL emission intensity at the excitation of 360 nm, 'A' denotes the optical density, ' η ' is the refractive index of a given solvent (for the distilled water $\eta_x/\eta_s = 1$), 'x' for the synthesized CQDs, and 's' used for the standard used.

Stern-Volmer quenching constant

The quenching efficiency of the synthesized CQDs was calculated by the following Stern-Volmer (SV) quenching equation:

$$F_0/F = 1 + K_{SV}[Hg^{2+}]$$
⁽²⁾

Where 'F' used for the FL emission intensity at various concentrations of absorber Hg^{2+} , F_0 being the absorber at $[Hg^{2+}] = 0$, and K_{SV} is the SV quenching constant



Figure S1 (a) Photostability of CQDs, showing the fluorescence of CQDs remain almost same even after incubating 5 months at 4 $^{\circ}$ C, (b) stability under the high ionic strength after the addition of different concentration of NaCl (0, 10, 20, 30, 40, 50 mM).



Figure S2 represents the optimization of pH ranges from 3 to 13, showing that prepared CQDs is independent of the pH used.



Figure S3 The kinetic stability of CQDs- Hg^{2+} system, indicating 5 min time is optimum to complete the quenching mechanism.



Figure S4 Fluorescence decay curve for the Hg^{2+} detection analysis.



Figure S5 Interference study under various conditions $[Hg^{2+}] = 0.05 \text{ mM}$, $[Al^{3+}] = [Pb^{2+}] = [Ca^{2+}]$, $[Mg^{2+}] = 5 \text{ mM}$, $[Zn^{2+}] = [Cd^{2+}] = [Ni^{2+}] = [Fe^{2+}] = 10 \text{ mM}$, $[Cu^{2+}] = [Fe^{3+}] = 0.01 \text{ mM}$.



Figure S6 The relationship between the FL emission intensity variation and the Hg^{2+} concentration.



Figure S7 The fluorescence response of CQDs/Hg²⁺ solution towards different essential amino acids of concentration 40 μ M where F and F₀ are fluorescence intensities of CQDs/Hg²⁺/amino acid and CQDs respectively.