Electronic Supplementary Information

Cu²⁺ integrated carbon dots as efficient bioprobe for selective sensing of guanine nucleobase

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Scheme S1. Synthetic scheme of CuCD (molar ratio between EDTA-2Na, $2H_2O$ and $CuCl_2$ · $2H_2O$ was 1:1).



Scheme S2. Synthetic scheme of CuCD (molar ratio between EDTA-2Na, $2H_2O$ and $CuCl_2$ · $2H_2O$ was 2:1).



Scheme S3. Synthetic scheme of CuCD (molar ratio between EDTA-2Na, $2H_2O$ and $CuCl_2$ · $2H_2O$ was 1:2).



Fig. S1 FEGTEM images of CuCD.



Fig. S2 DLS study of CuCD.



Fig. S3 Deconvolution spectra of (a) C 1s (b) O 1s and (c) N 1s of CuCD.



Fig. S4 (a) XRD, and (b) EDX analysis of CuCD. Sample was crusted on glass grid for the EDX experiment.



Fig. S5 Tauc's plot of CuCD.



Fig. S6 Stern-Volmer plot of guanine (1-10 nM) in aqueous solution of CuCD (250 µg/mL).



Fig. S7 Structures of (a) adenine, (b) guanine, (c) cytosine and (d) thymine nucleobases.



Fig. S8 (a) Structure of guanine and (b) proposed binding model of CuCDs-guanine complex.



Fig. S9 Interference studies between guanine (20 nM) and samples of (a) CuCD (250 μ g/mL) and adenine (20 nM) mixture, (b) CuCD (250 μ g/mL) and cytosine (20 nM) mixture and (c) CuCD (250 μ g/mL) and thymine (20 nM) mixture.



Fig. S10 Selectivity of CuCD (250 μ g/mL) to guanine over other metal ions and biomolecules (500 nM). (a) Fluorescence intensity plot (b) relative intensity of CuCD in presence of different metal ions and biomolecules. The error bars represent the standard deviations.



Fig. S11 FEGTEM images of (a) CuCD, (b) CuCD-guanine complex and (c) CuCD, (d) CuCD-guanine complex at higher resolution.



Fig. S12 Suspension stability index of CuCD solution where $[CuCD] = 500 \ \mu g/mL$ (a) CuCD with respect to FBS concentration (0-75%) in DMEM media, (b) CuCD with respect to number of days in 10% FBS in DMEM media and (c) photostability of CuCD solution under UV (wavelength 365 nm, power 12 W) light irradiation up to 200 min.



Fig. S13 Cell viability experiment of CuCD where $[CuCD] = 25-200 \ \mu g/mL$ in NIH3T3, B16F10 and MCF-7 cells after 12 h of incubation. The standard deviation was in the range of 1-3% in triplicate experiments.



Fig. S14 Bright field and fluorescence microscopic images of cells after 12 h incubation with CuCD where $[CuCD] = 200 \ \mu g/mL$ in case of (a,b) treated NIH3T3 cells with guanine (500 nM) and (c) corresponding mean fluorescence intensity for NIH3T3.