

High sensitive sensing of hydroquinone and catechol based on β -cyclodextrin modified carbon dots

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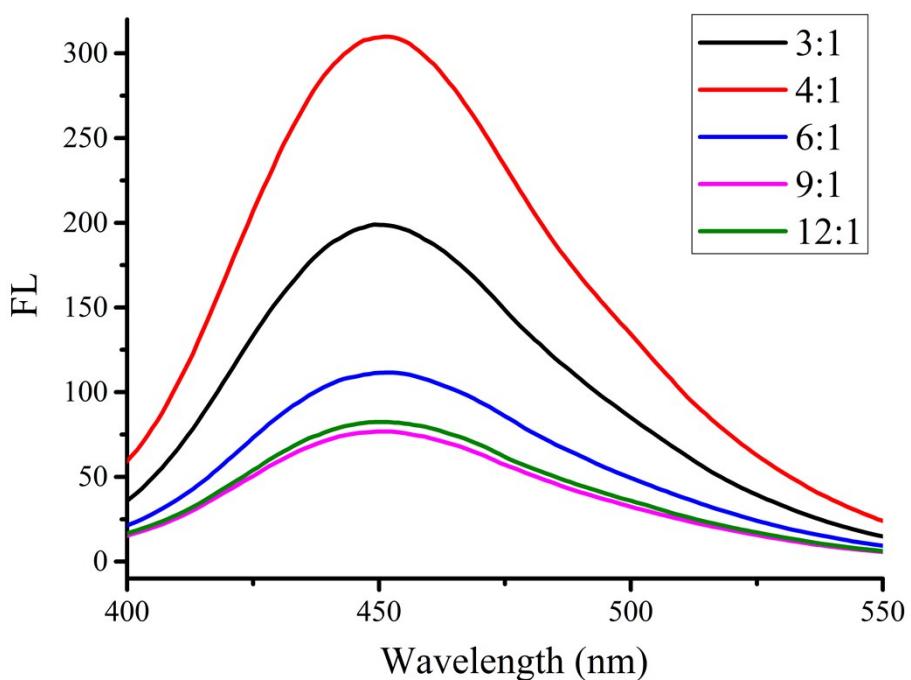


Fig. S1. Fluorescence spectra of C-dot and amono-6-OTs- β -CD mixed with the different ratio ([mono-6-OTs- β -CD]/[C-dot]v/v: (black) 3 : 1, (red) 4 : 1, (blue) 6 : 1, (pink) 9 : 1, (green) 12 : 1.

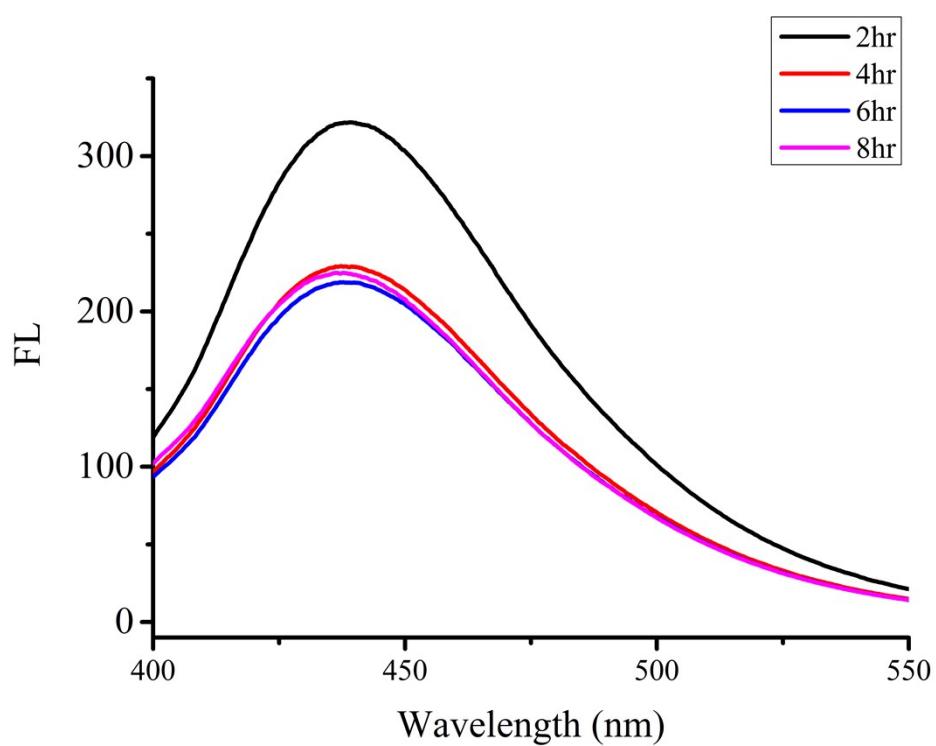


Fig. S2. Fluorescence spectra of the synthesized C-dot@ β -CD at variety incubate time.

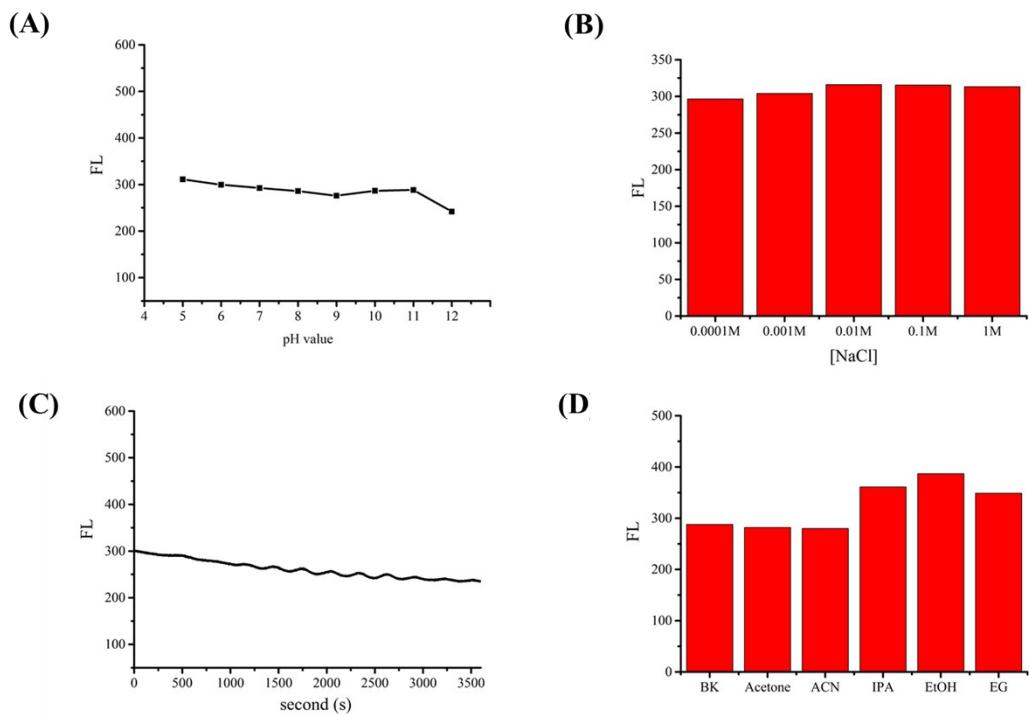


Fig. S3. (A) The stability of C-dot@ β -CD at pH ranges from 5 to 12, (B) Stability of C-dot@ β -CD in different concentrations of NaCl ranging from 0.1 mM to 1M, (C) Photostability of C-dot@ β -CD, (D) Influence of different solvents (80% of total volume) on the fluorescence properties.

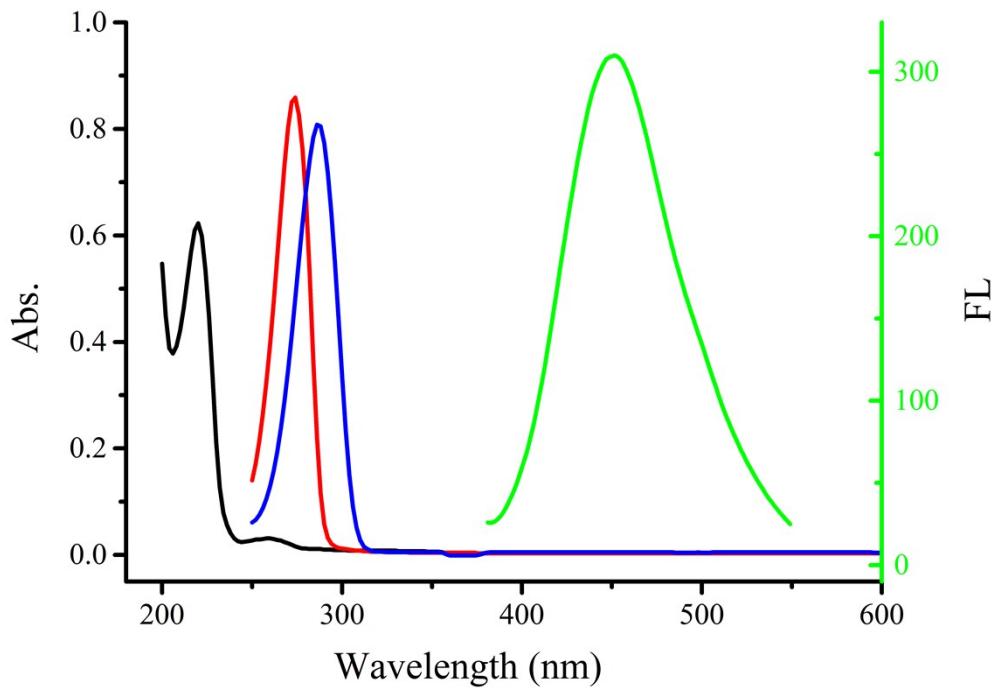


Fig. S4. UV–vis absorption of C-dot@ β -CD(black), CC (red), HQ(blue) and C-dot@ β -CD fluorescent emission(green).

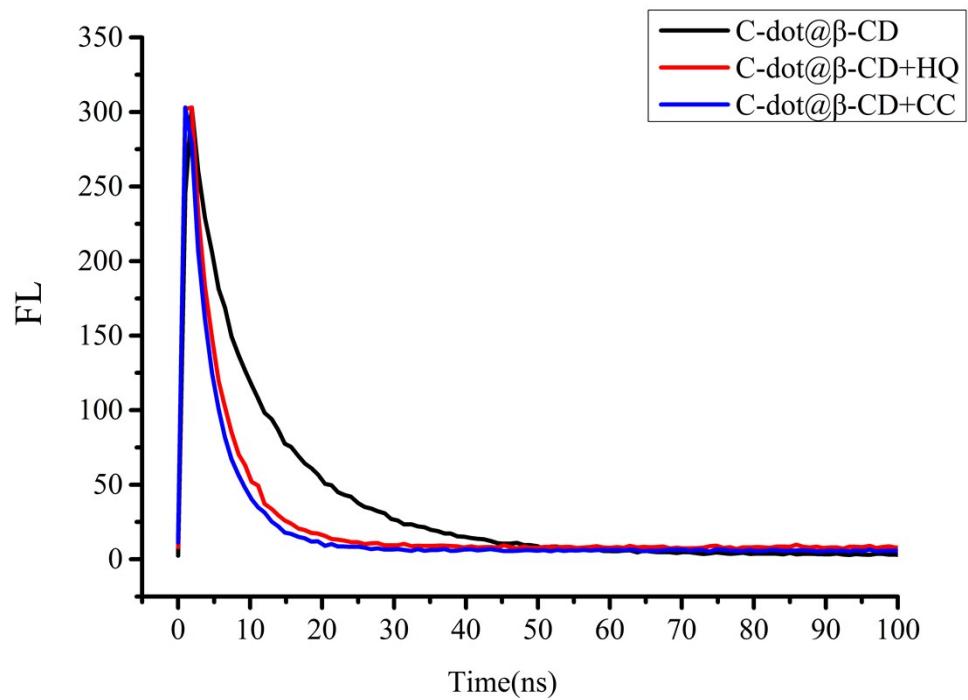


Fig. S5. Time-resolved decay of C-dot@ β -CD (black) with HQ (red), with CC(blue).

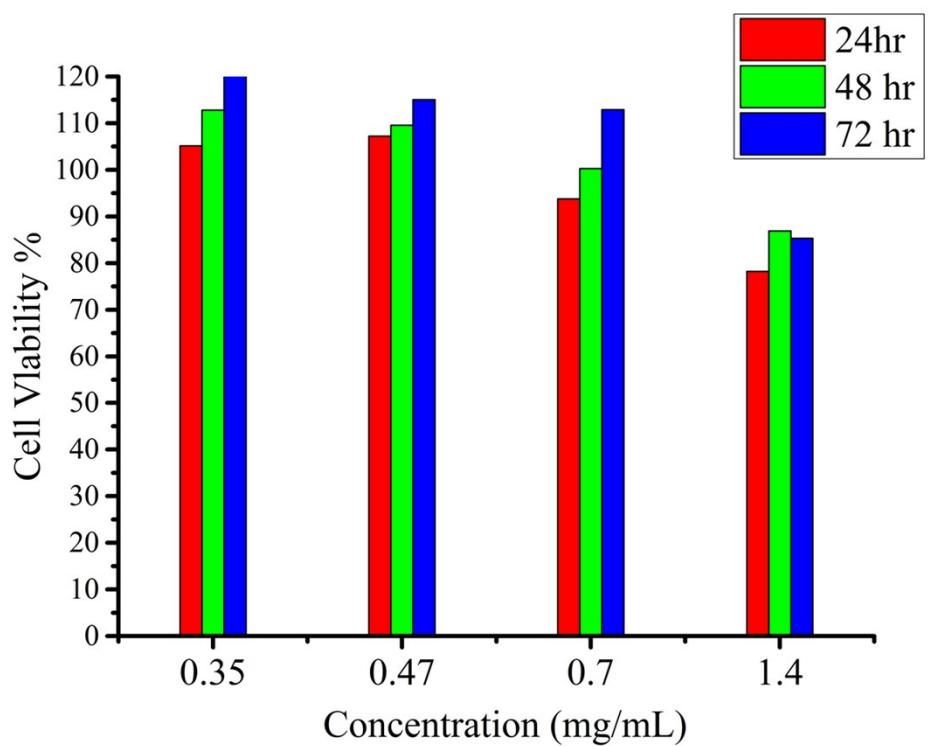


Fig. S6. Cell viability assay of human HeLa cells against C-dot@ β -CD at arranged concentrations from 0.35-1.4 mg/mL.

TableS1. Time-resolved decay of C-dots in the absence and presence of catechol and hydroquinone

Sample	τ_1 (ns)	Area (%)	τ_2 (ns)	Area (%)	Average τ (ns)
C-dot@ β -CD	13.65	64.08	3.38	35.92	9.96
C-dot@ β -CD + CC	5.11	62.80	1.73	37.20	3.85
C-dot@ β -CD + HQ	5.63	63.68	1.96	36.32	4.30