Supplementary Materials

Microwave-assisted synthesis of nitrogen doped carbon dots using prickly pear as the carbon source and its application as highly selective sensor for Cr(VI) and as patterning agent

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Sample name	Integrated	Absorbance	Refractive	Quantum yield
	emission	(A) at 360 nm	index of	(Q) at 360 nm
	intensity (I)		solvent (η)	
Quinine	542308	0.132	1.33	0.54 (known
sulphate				standard value)
N-CDs	80758	0.0752	1.33	0.148

 Table S1. Quantum yield calculation of N-CDs.

Sample	τ_1/ns (%)	$\tau_2/ns~(\%)$	$\tau_{\rm ave}/ns$
N-CDs	1.46 (47)	8.10 (53)	4.979
N-CDs + Cr(VI)	1.40 (44)	7.56 (56)	4.849

Table S2. Fluorescence lifetime data with bi-exponential fit of the fluorescence decay curvesof N-CDs and NCDs+Cr(VI).

Sample No.	Spiked (µM)	Intensity (I0)(a.u)	Intensity of Sample (I) (a.u)	I0/I (a.u)	Calculated Concentration (µM)
DW-1	15	3.9745	2.89	1.373	15.1
DW-2	30	3.9745	2.23	1.786	31.8
TW-1	25	3.9745	2.45	1.625	25.8
TW-2	35	3.9745	2.05	1.936	37.8

 Table S3. Different parameters for analysis of real sample using Stern-Volmer equation.



Fig. S1. Raman spectrum of N-CDs.



Fig. S2. Fluorescence decay curve of N-CDs recorded at 360 nm excitation.



Fig. S3. Bar diagram showing change in fluorescence intensity of N-CDs at different pH ranging from 2 to 12. ($\lambda ex = 360 \text{ nm}$, $\lambda emi = 450 \text{ nm}$). (Red marking with metal ion).



Fig. S4. Fluorescence spectra of N-CDs with 0.5 to 5.0 % of NaCl is shown. The intensity of the fluorescence band as a function of concentration of NaCl is shown in the inset.



Fig. S5. Plot of the change in fluorescence intensity of N-CDs in presence of different concentrations of Cr(VI) as a function of time (0 to 60 min).



Fig. S6. Fluorescence spectral change of N-CDs in presence of various metal ions ($\lambda_{ex} = 360$ nm).



Fig. S7. Fluorescence decay curve of N-CDs in presence of Cr(VI).