

Supporting information

Highly stable and water dispersible polymer-coated CsPbBr₃ nanocrystals for Cu-ion detection in water

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Experimental section:

Characterization methods: UV-visible absorption spectra of NCs were recorded with a Shimadzu UV-2700 Spectrophotometer. Steady-state photoluminescence (PL) spectra were collected with Ocean Insight MAYA 2000 Pro high sensitivity Spectrometer using a 370 nm UV excitation source. X-Ray diffraction (XRD) analysis was carried out with the Bruker D8 diffractometer using Cu-K_α ($\lambda=1.54 \text{ \AA}$) as incident radiation. Transmission electron microscopy (TEM) and high-resolution transmission electron microscopy (HRTEM) imaging were recorded by Jeol-JEM-2010 microscope operated at 200 kV to determine the shape and size of the NCs. The Fourier Transform Infrared (FTIR) analysis was carried out using JASCO FT/IR-6600 Infrared Spectrometer to confirm the silica and polymers encapsulation around the NCs.

Ion detection test: To probe the Cu²⁺ ions, the PL intensity of the samples was recorded after the subsequent addition of different metal ions. Firstly, aqueous solutions containing metal ions at same concentration (0.1 mM) including Ni²⁺, Al³⁺, In³⁺, Co²⁺, Fe²⁺, Fe³⁺, Zn²⁺ and Cu²⁺ were prepared. The NC aqueous solution was prepared by dispersing directly dispersing powder PbN-4 sample in DI water (concentration~ 5 mg/mL). Next, 100 μL of PbN-4 NC aqueous solution was taken in 7 glass vials and 100 μL of each of the aqueous solutions containing the metal ions were added and the PL intensity of the NC solution was observed.

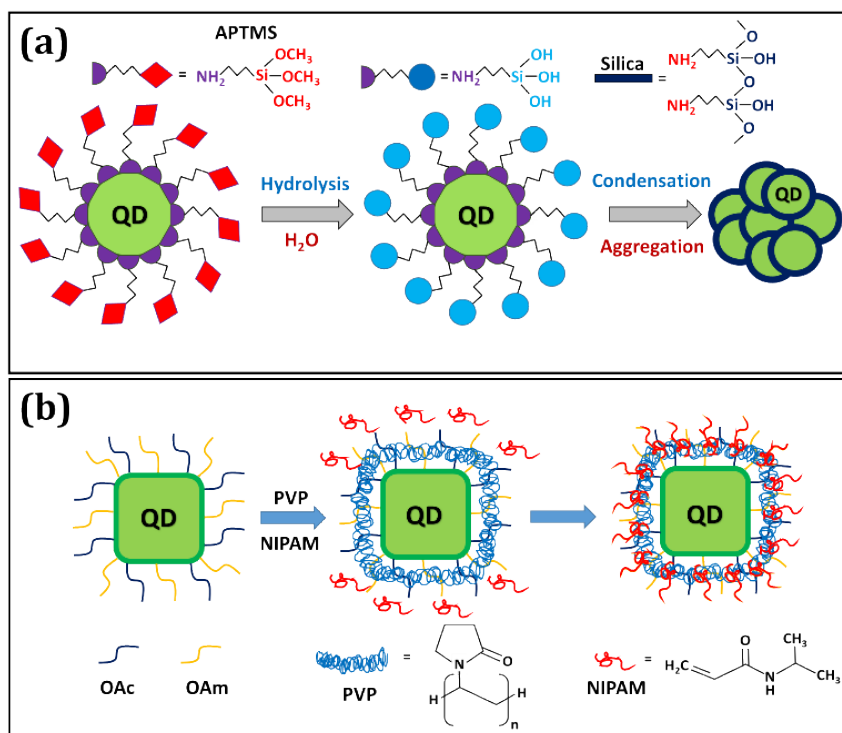


Figure S1. (a) Schematic representation of hydrolysis and condensation processes of APTMS for silica shelling around the perovskite NCs, and (b) Schematic representation of polymer coating around PbN NCs.

Results and discussion:

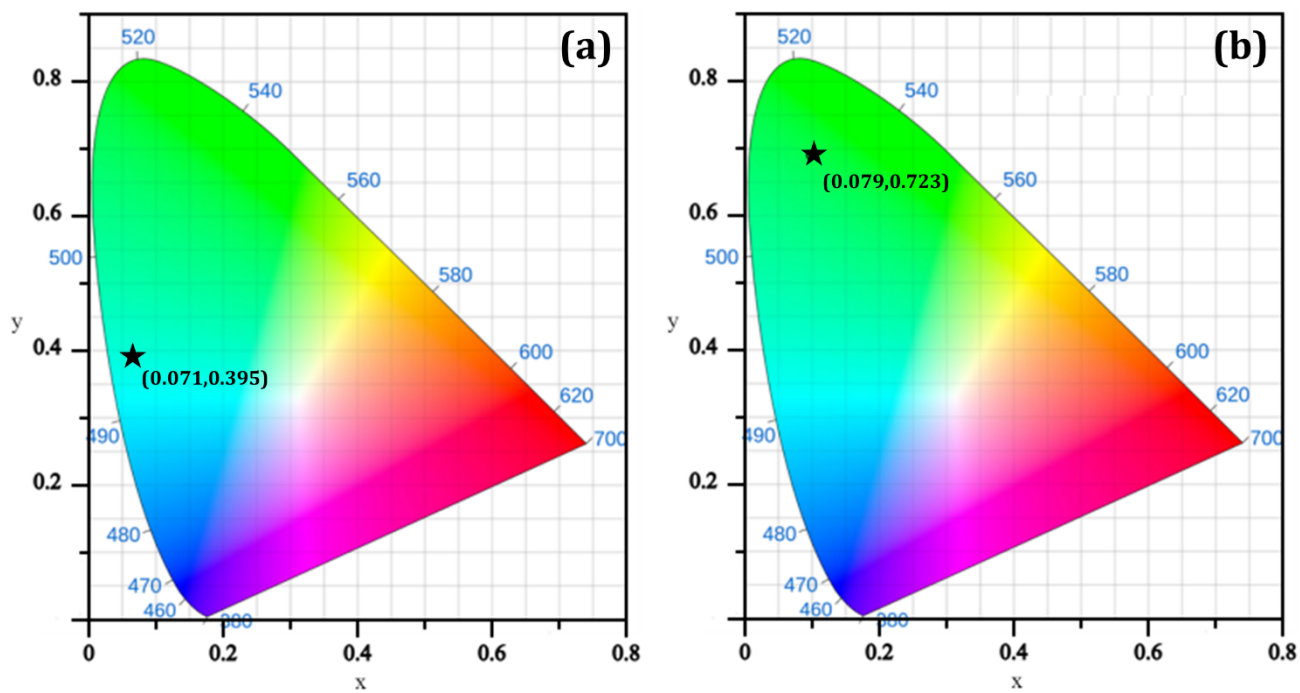


Figure S3. CIE color coordinates of (a) CPB@SiO₂ and (b) PbN-3 NCs.

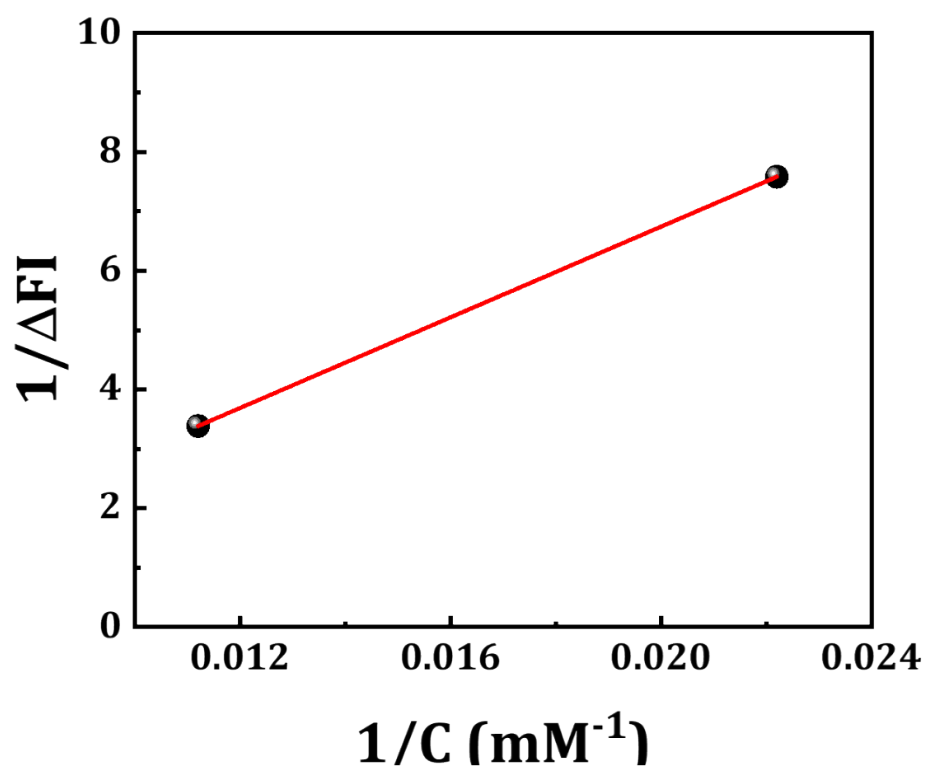


Figure S4. Double-reciprocal plot of $[C]$ of NIPAM versus the change in the fluorescence intensity (ΔFI) of PbN NCs.

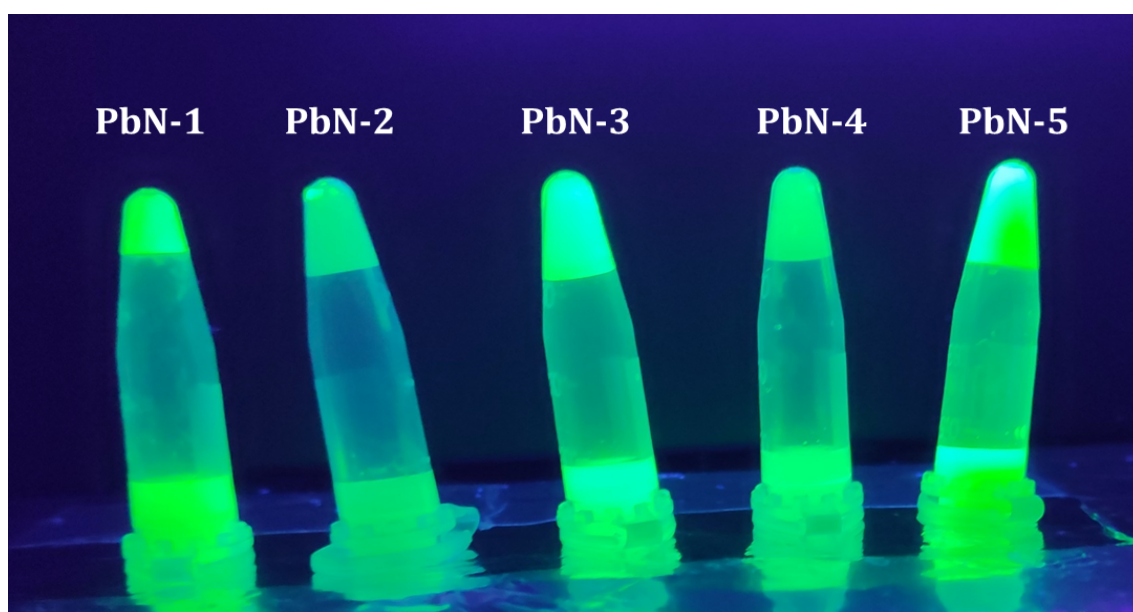
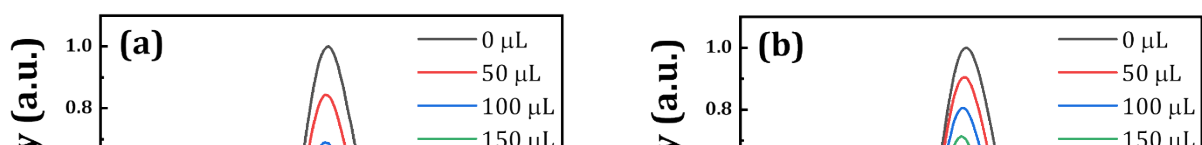


Figure S5. Photographic image of PbN NCs dispersed in water under a UV lamp.



	0 μL	50 μL	100 μL	150 μL	200 μL	250 μL	300 μL	350 μL	400 μL	450 μL	500 μL	550 μL	600 μL	650 μL	700 μL	750 μL	800 μL
CPB@SiO ₂																	
PbN-1																	
PbN-2																	
PbN-3																	
PbN-4																	

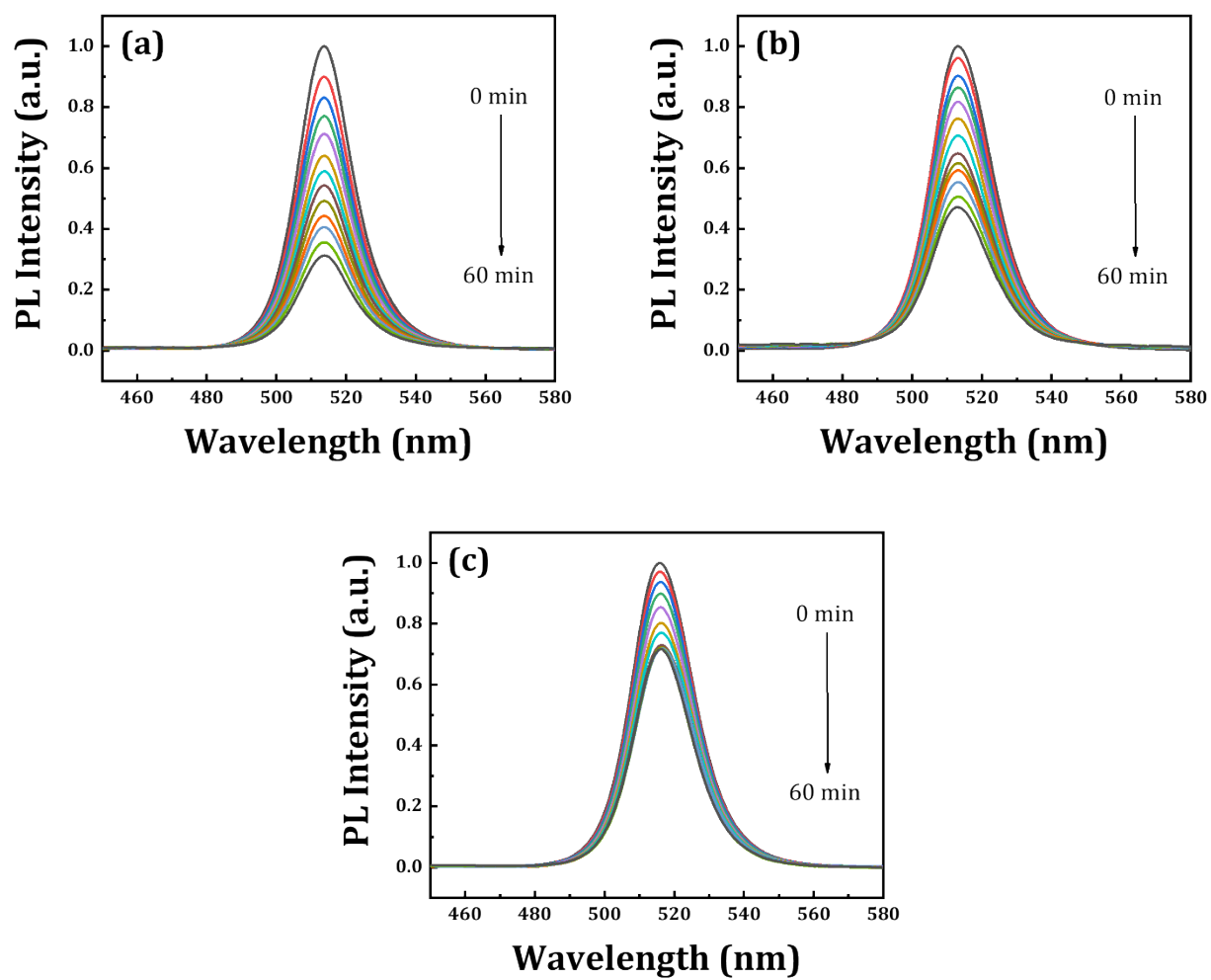


Figure S8. Water stability of (a) PbN-2, (b) PbN-3 and (c) PbN-5 as a function of time.










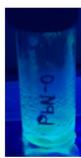











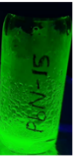
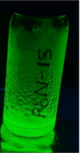

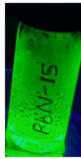



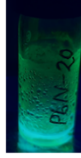

	0 min	15 min	30 min	45 min	60 min
CPB@SiO ₂					
PbN-1					
PbN-2					
PbN-3					
PbN-4					
PbN-5					

Figure S9. Water stability test 2: Photographs of samples at a time interval of 15 mins after addition of DI water under a UV Lamp.

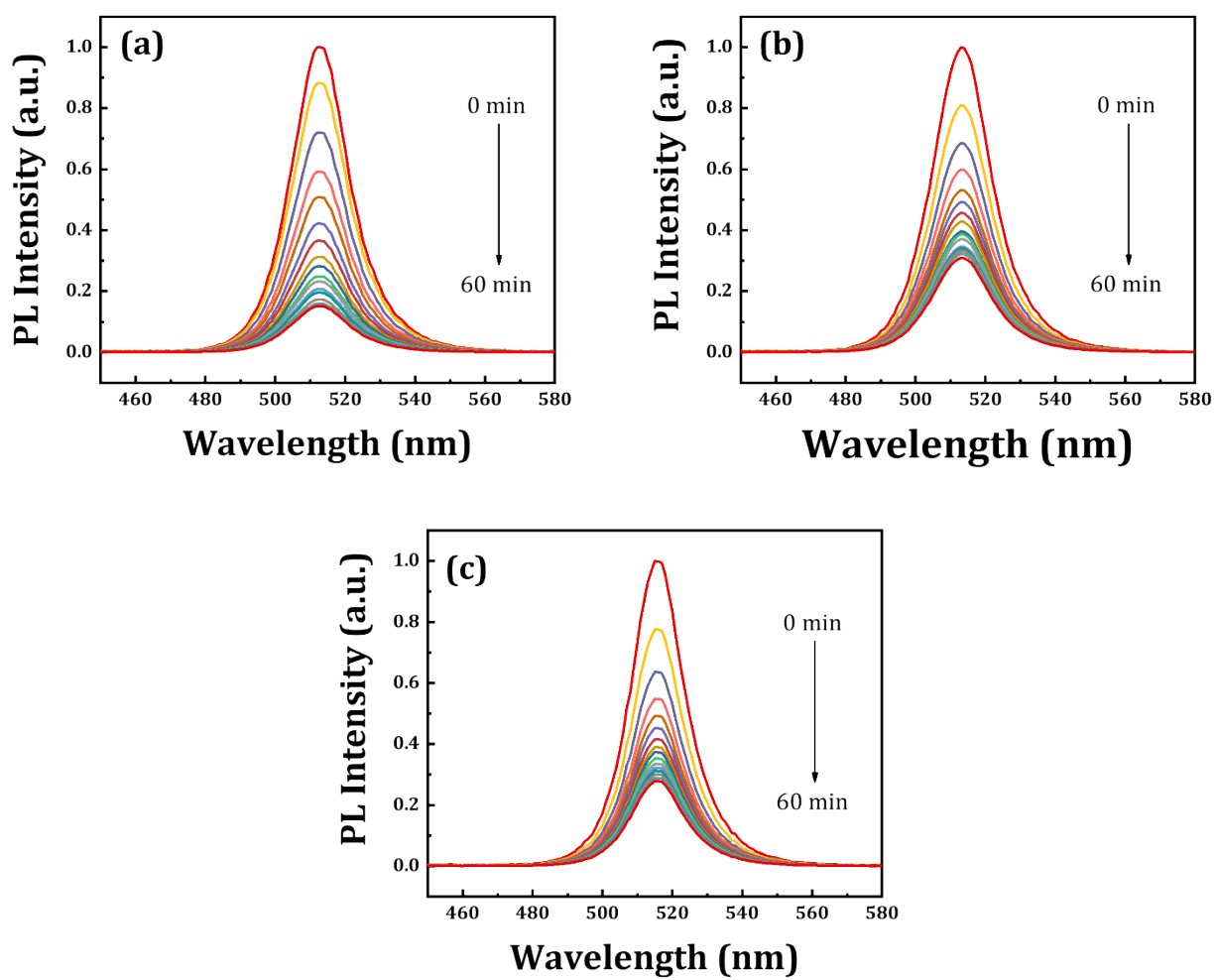


Figure S10. Heat stability of (a) PbN-2, (b) PbN-3 and (c) PbN-5 at 60°C as a function of time.

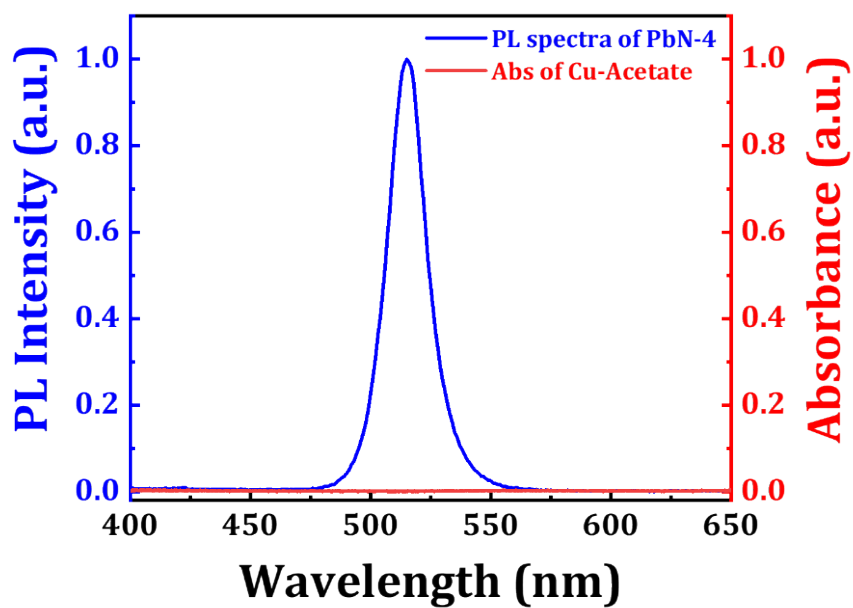


Figure S11. Absorption and PL spectra of Cu-acetate and PbN-4 NCs as shown in legends.

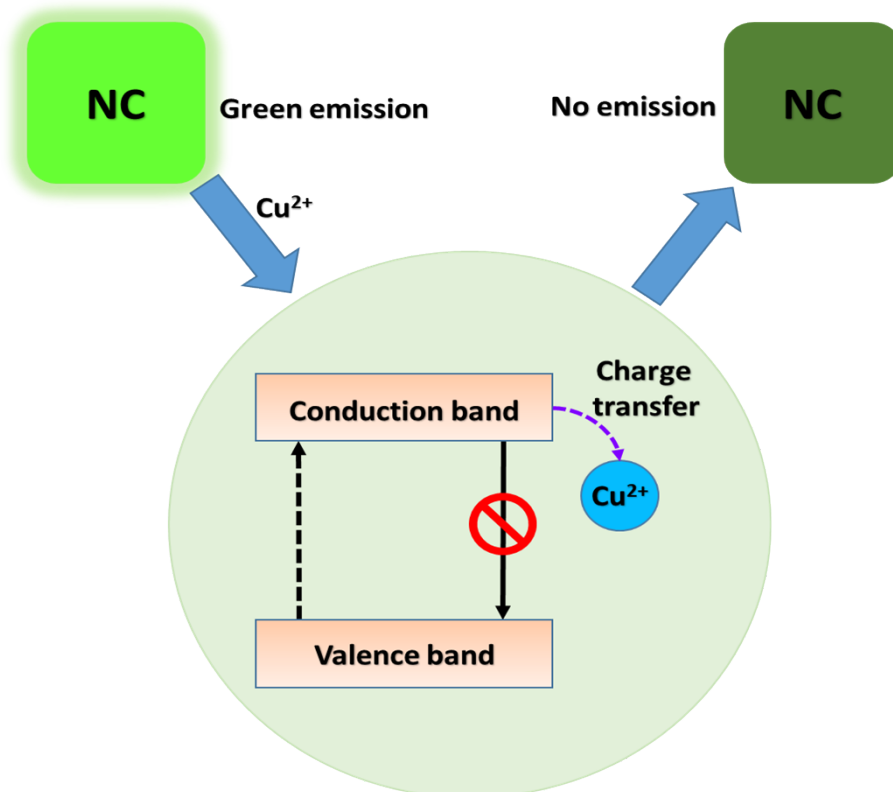


Figure S12. Schematic representation of charge transfer mechanism in NCs in presence of Cu^{2+} -ions.