

Supporting Information

Thermally stable and hydrophilic CsPbBr₃/mPEG-NH₂ nanocrystals with enhanced aqueous fluorescence for cell imaging

Qi-Bao Yan^a, Ning Bao^b and Shou-Nian Ding^{a*}

^aJiangsu Province Hi-Tech Key Laboratory for Bio-medical Research, School of Chemistry and Chemical Engineering, Southeast University, Nanjing 211189, China

^bSchool of Public Health, Nantong University, 226019 Nantong, Jiangsu, China

*E-mail: snding@seu.edu.cn.

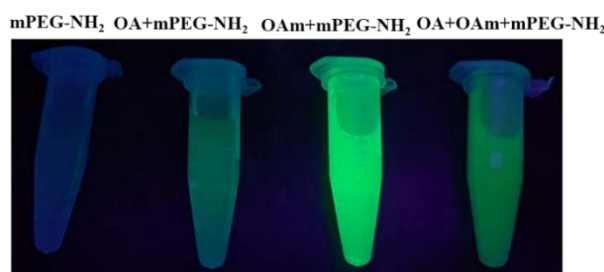


Figure S1. The PL image of CsPbBr₃ NCs aqueous solution (0.5 mg mL⁻¹) prepared via using different combinations of ligand.

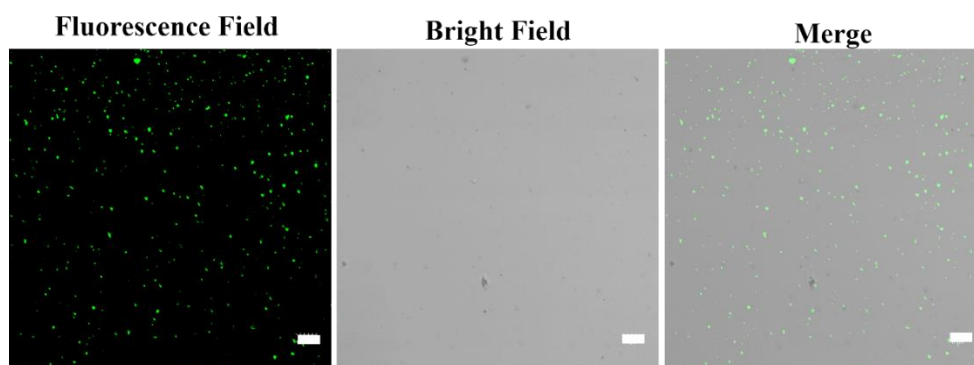


Figure S2. The confocal fluorescence image of CsPbBr₃/mPEG-NH₂ NCs. The scale bars are 5 μm.

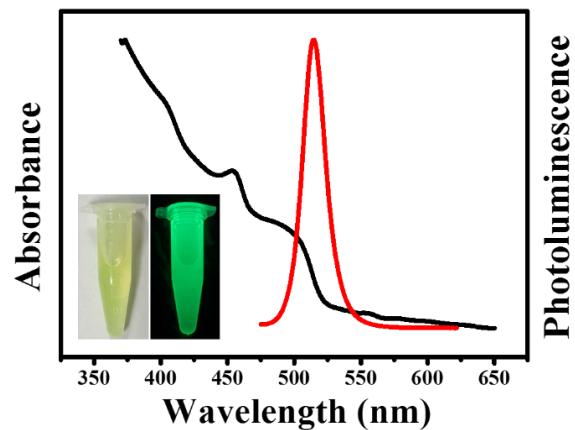


Figure S3. The UV/vis absorption and PL spectra of CsPbBr₃ NCs prepared by using oleic acid and oleylamine. The inset exhibited the optical image of corresponding sample in daylight (left) and under 365 nm UV light (right).

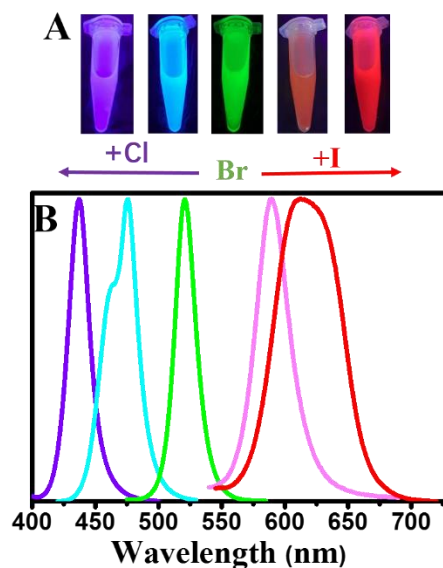


Figure S4. The (A) optical images of CsPbX₃/mPEG-NH₂ NCs with different halide compositions and the (B) PL spectra of the corresponding samples.

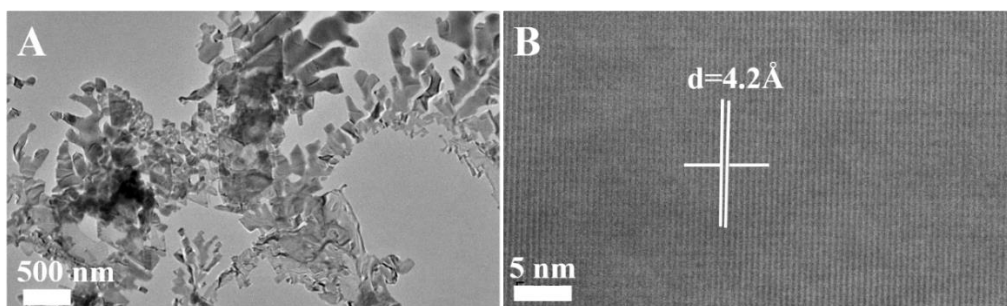


Figure S5. The (A) TEM and (B) HR-TEM images of CsPbBr₃/mPEG-NH₂ NCs aqueous solution after three days.

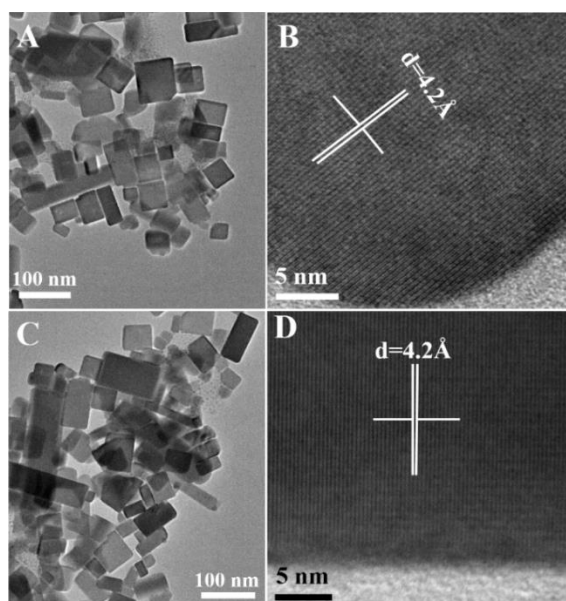


Figure S6. The TEM and HR-TEM images of CsPbBr₃/mPEG-NH₂ NCs toluene solution (A, B) at initial stage and (C, D) one week later.

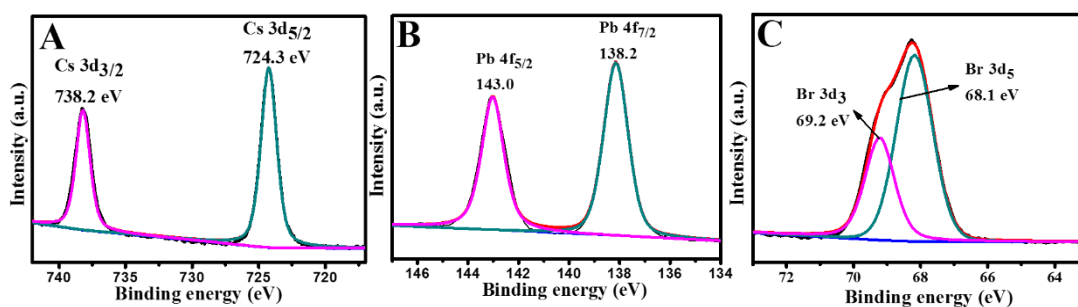


Figure S7. The high-resolution XPS analysis corresponding to (A) Cs 3d, (B) Br 3d and (C) Pb 4f, separately.

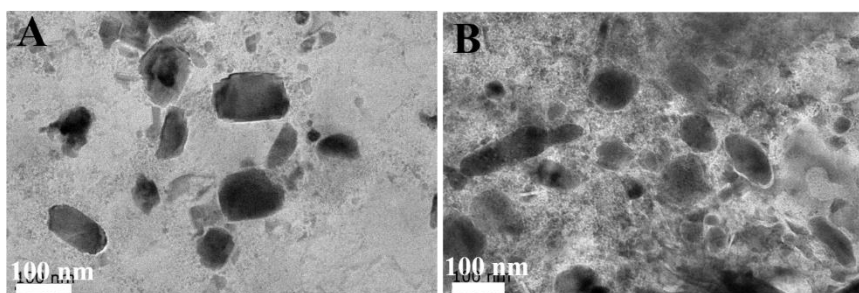


Figure S8. The negative stained TEM images of aqueous solution of CsPbBr₃/mPEG-NH₂ NCs synthesized with the optimum amount of mPEG-NH₂ at initial stage and after one week.

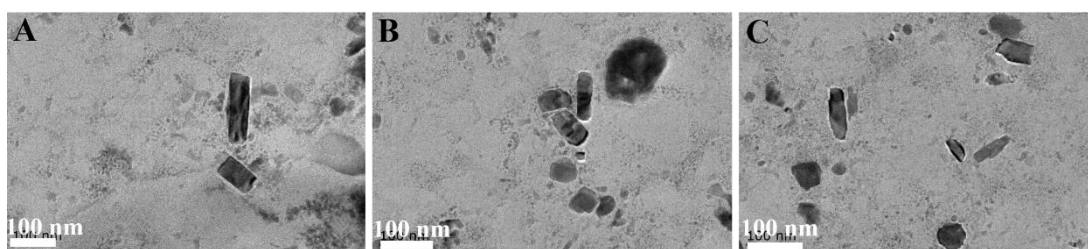


Figure S9. The negative stained TEM images of CsPbBr₃/mPEG-NH₂ NCs prepared by using different concentrations of mPEG-NH₂ (A: 6 mg mL⁻¹, B: 8 mg mL⁻¹, C: 10 mg mL⁻¹).

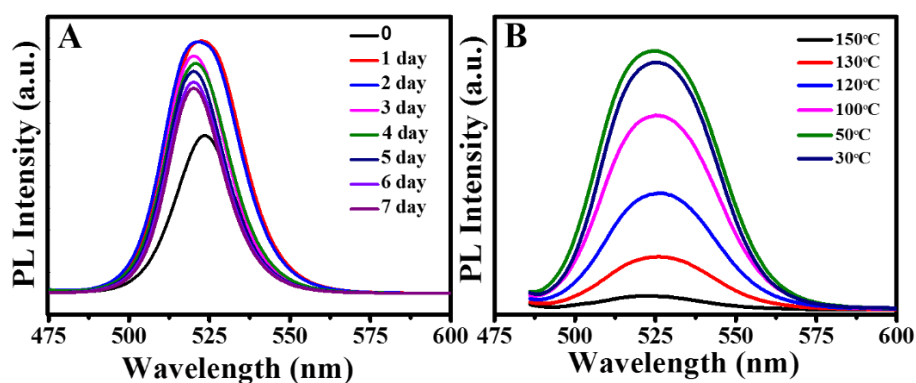


Figure S10. The (A) time-dependent PL spectra of 0.5 mg mL⁻¹ CsPbBr₃/mPEG-NH₂ NCs in water. The (B) temperature-dependent PL spectra of CsPbBr₃/mPEG-NH₂ NCs powders.

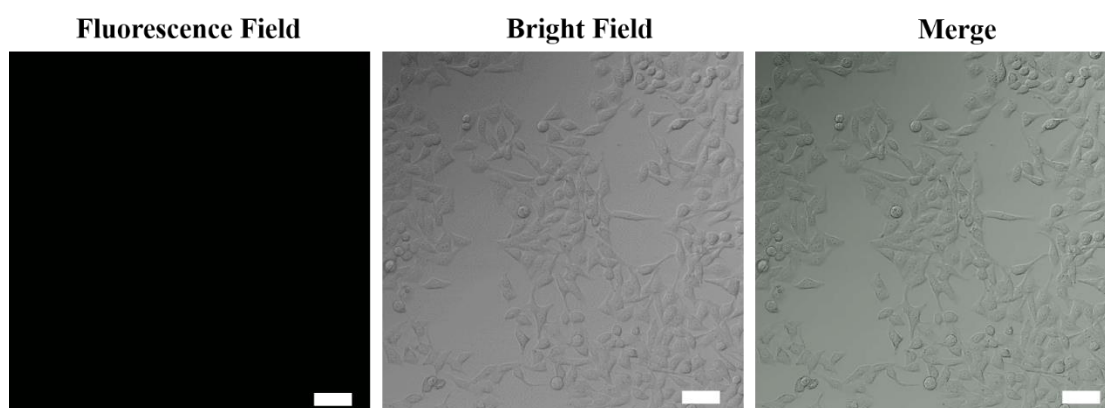


Figure S11. The confocal fluorescence image of HepG2 cells incubated without CsPbBr₃/mPEG-NH₂ NCs. The scale bars are 40 μm.

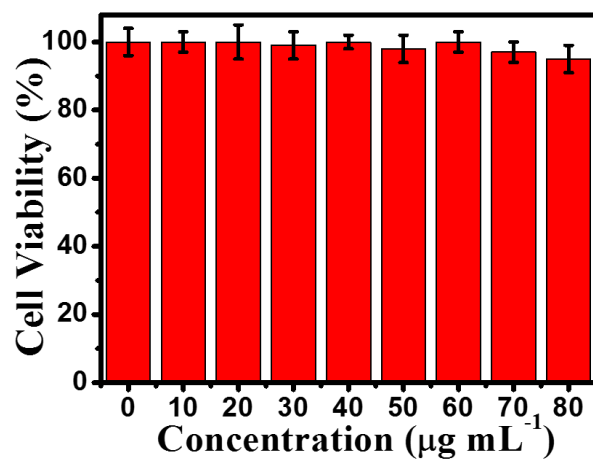


Figure S12. Viability of HepG2 cells at various CsPbBr₃/mPEG-NH₂ NCs concentrations.

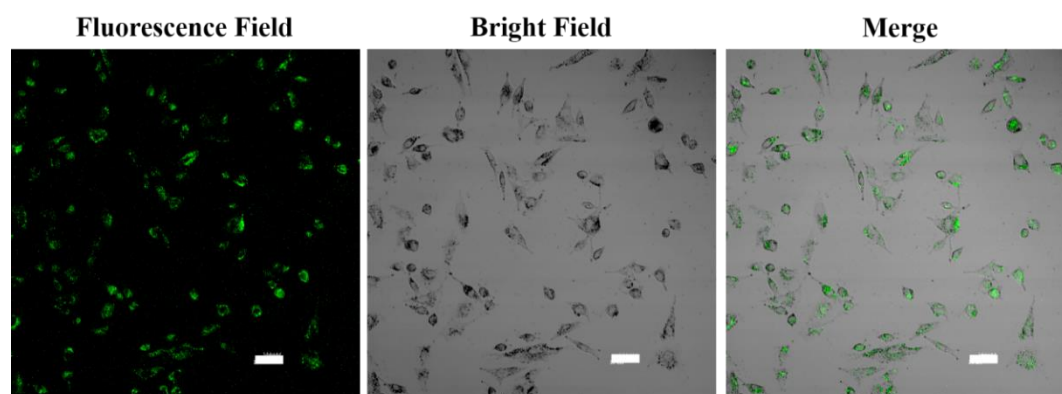


Figure S13. The confocal fluorescence image of A549 cells incubated with CsPbBr₃/mPEG-NH₂ NCs for 24 h. The scale bars are 50 μm .

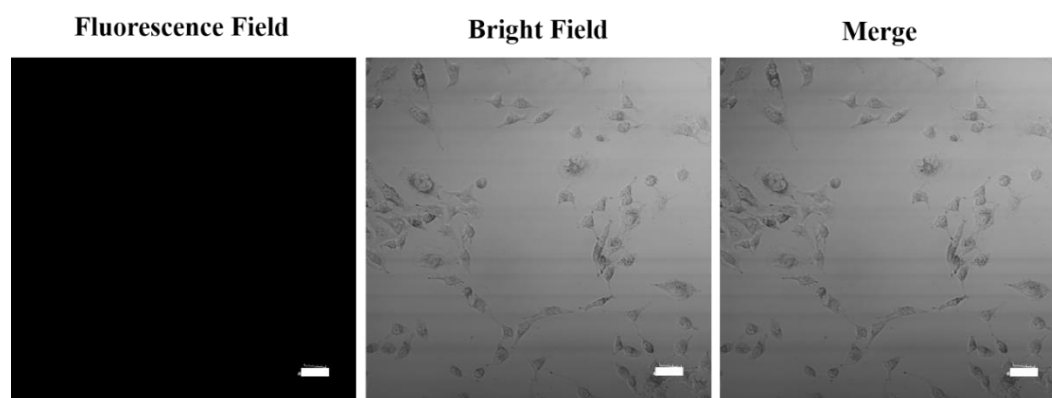


Figure S14. The confocal fluorescence image of A549 cells incubated without CsPbBr₃/mPEG-NH₂ NCs for 24 h. The scale bars are 50 μm .