Supporting Information

Improved Stability and Photoluminescence Yield of Mn²⁺-Doped CH₃NH₃PbCl₃ Perovskite Nanocrystals

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Table S1. Elemental analysis of CH₃NH₃Pb_xMn_{1-x}Cl₃ PNCs passivated with APTES under different doping concentration using ICP-AES.

Nominal concentration (C _{Mn} /C _{Mn} +C _{Pb})	25 at. %	50 at. %	75 at. %	90 at. %
Detected concentration $(C_{Mn}/C_{Mn}+C_{Pb})$	10.1 at. %	14.8 at. %	30.0 at. %	59.1 at. %



Figure S1. Photographs of 50 at. % Mn-doped CH₃NH₃PbCl₃ PNCs passivated with octylamine (left) and APTES (right) under room light (A) and 365 nm UV light (B).



Figure S2. TEM image of CH₃NH₃PbCl₃ PNCs passivated with APTES.



Figure S3. (a) Absorption and (b) PL spectra of solutions prepared using MnCl₂ and APTES precursors (black line), and using MnCl₂, APTES, PbCl₂ and CH₃NH₃Cl precursors (red line). (c) PL spectra of MnCl₂ (λ_{ex} = 360 nm).



Figure S4. PLE spectra of Mn doped CH₃NH₃PbCl₃ PNCs with different doping concentration $(\lambda_{em} = 610 \text{ nm}).$



Figure S5. PL spectra of 50 at. % Mn-doped CH₃NH₃PbCl₃ PNCs solutions with different toluene/precursor volume ratios.



Figure S6. Air stability test of Mn-doped CH₃NH₃PbCl₃ PNCs passivated with octylamine and APTES.