

## *Supporting Information*

### **A Universal Synthesis Strategy for Stable CsPbX<sub>3</sub>@Oxide Core-Shell Nanoparticles through Bridging Ligands**

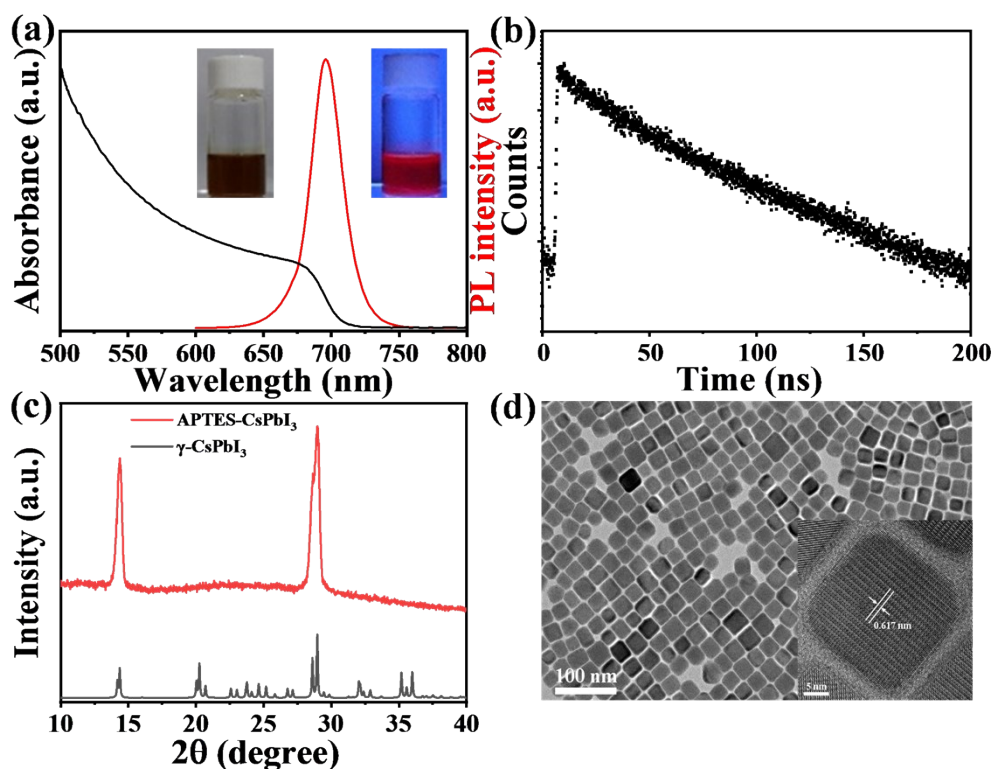
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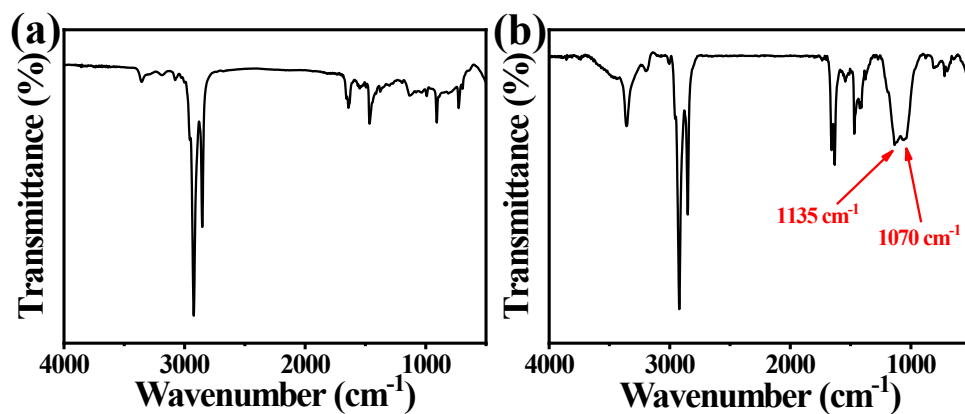
<sup>b</sup> Shenzhen Huazhong University of Science and Technology Research Institute, Shenzhen

**Corresponding Author**

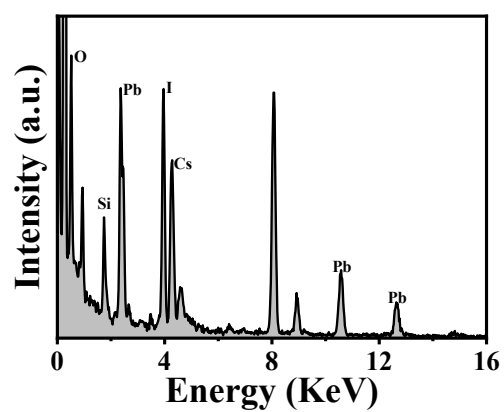
*\*E-mail: [yingma@hust.edu.cn](mailto:yingma@hust.edu.cn)*



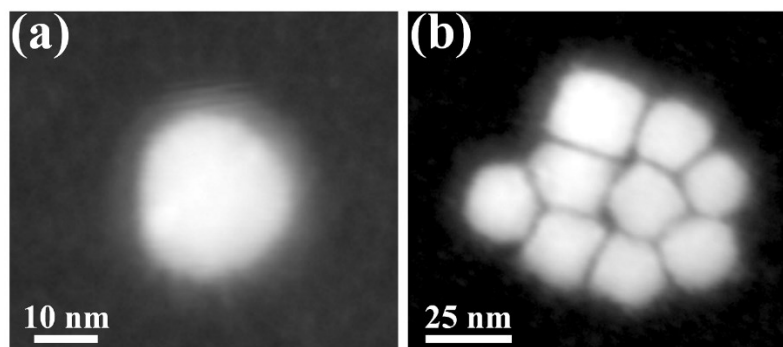
**Fig. S1** (a) PL spectrum and UV-Vis absorption spectrum of APTES-CsPbI<sub>3</sub> solution. The inset shows the APTES-CsPbI<sub>3</sub> solution under daylight and UV light irradiation ( $\lambda = 365$  nm). (b) Time-resolved PL decay, (c) XRD pattern and (d) TEM images of APTES-CsPbI<sub>3</sub> NCs.



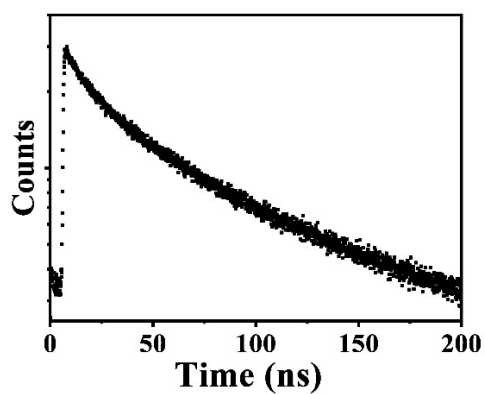
**Fig. S2** FT-IR spectra of (a) APTES-CsPbI<sub>3</sub> NCs and (b) CsPbI<sub>3</sub>@SiO<sub>2</sub> NCs.



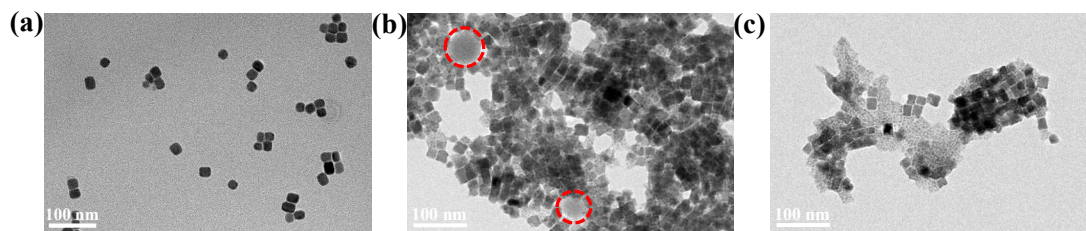
**Fig. S3** The energy-dispersive X-ray spectrum of the CsPbI<sub>3</sub>@SiO<sub>2</sub> NCs.



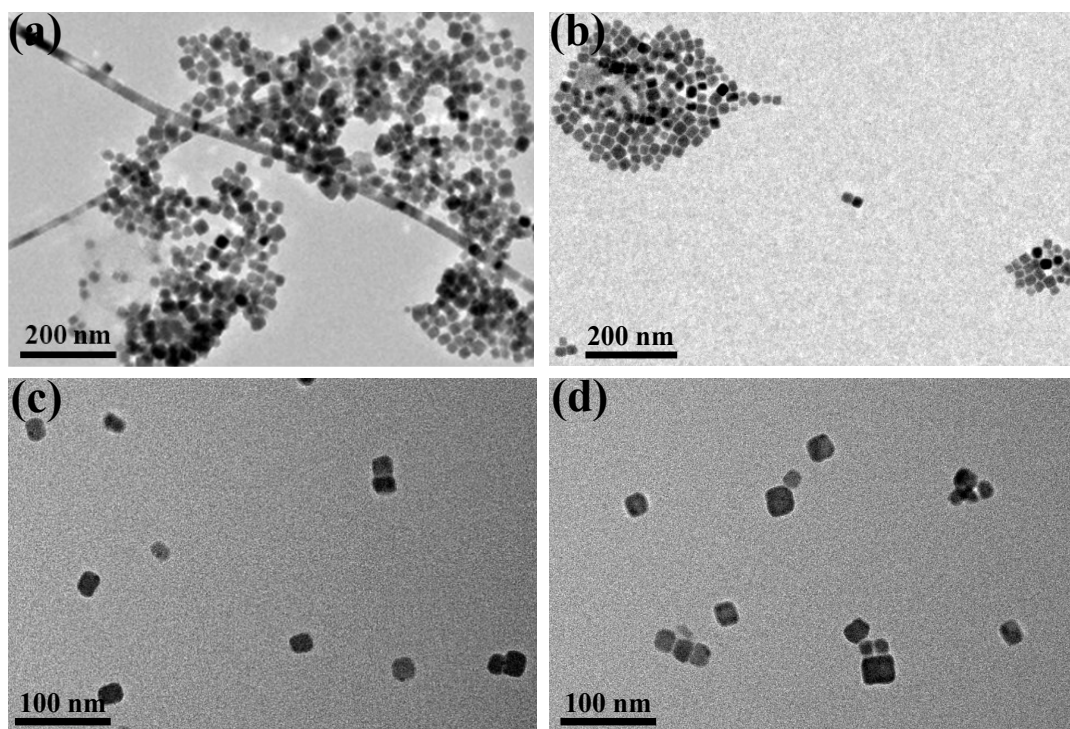
**Fig. S4** The HADF-STEM images of the CsPbI<sub>3</sub>@SiO<sub>2</sub> NCs.



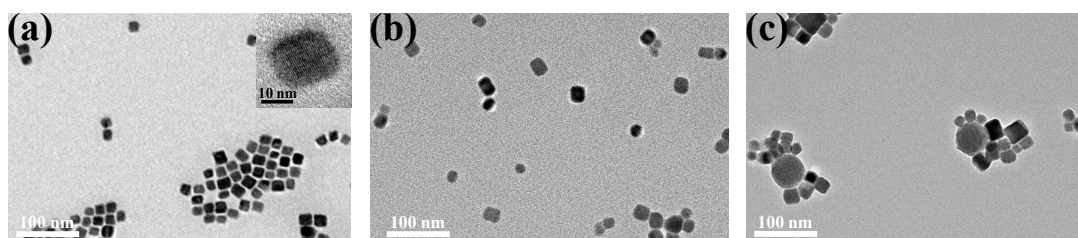
**Fig. S5** The time-resolved PL decay of the CsPbI<sub>3</sub>@SiO<sub>2</sub> NCs.



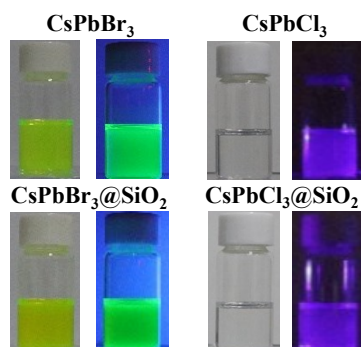
**Fig. S6** TEM images of the products prepared at different ammonia concentrations: (a) 3.4, (b) 6.7, and (c) 10 mmol L<sup>-1</sup>.



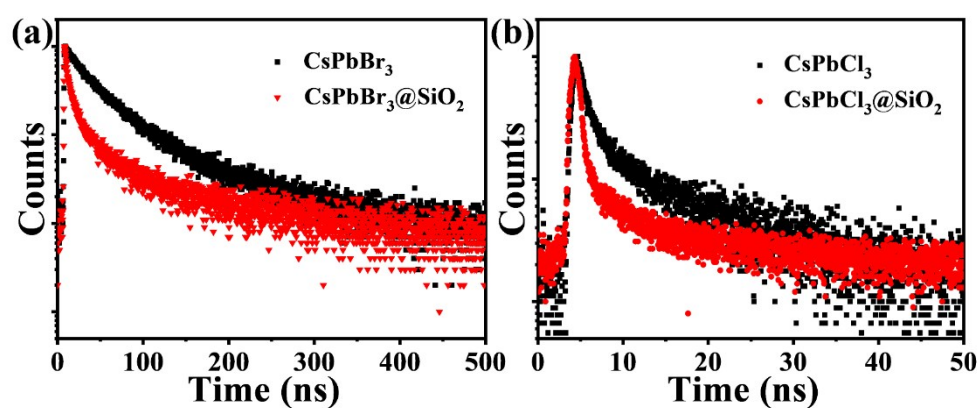
**Fig. S7** TEM images of the products prepared by varying ethanol quantities: (a) 0.5 mL, (b) 1 mL, (c) 2 mL, and (d) 3 mL.



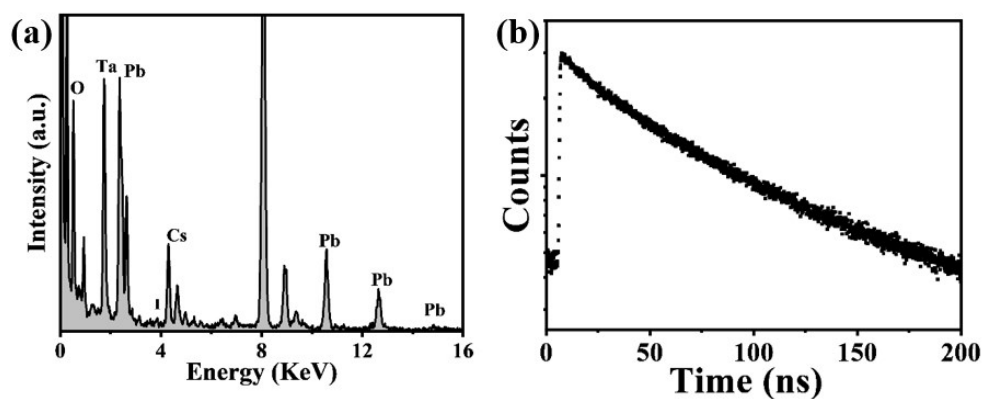
**Fig. S8** TEM images of the products prepared at different hydrolysis temperatures: (a) 20 °C, (b) 25 °C and (c) 30 °C.



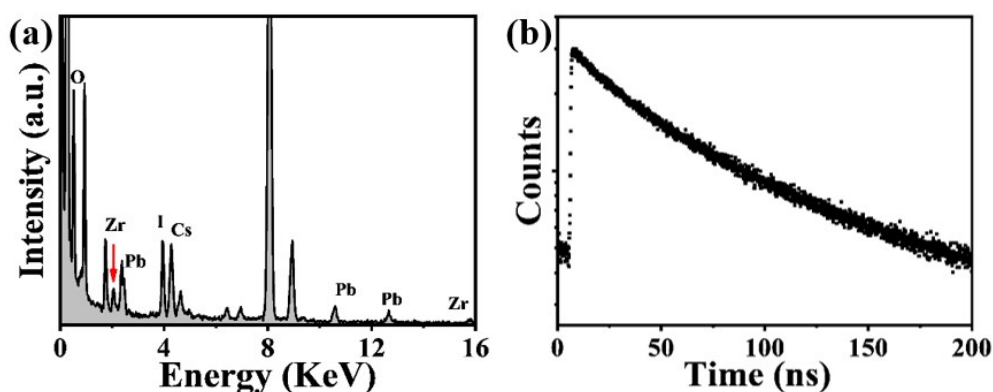
**Fig. S9** Photographs of the CsPbBr<sub>3</sub> and CsPbCl<sub>3</sub> NCs before and after SiO<sub>2</sub> shell growth under daylight and UV light irradiation ( $\lambda = 365$  nm).



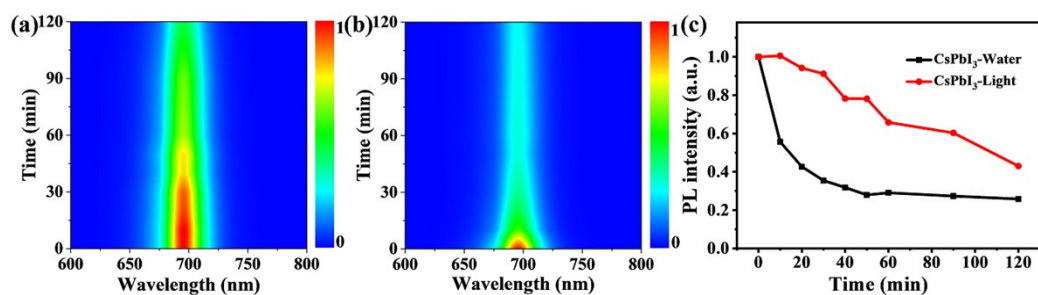
**Fig. S10** Time-resolved PL decay curves of (a) CsPbBr<sub>3</sub> and (b) CsPbCl<sub>3</sub> NCs before and after SiO<sub>2</sub> shell growth.



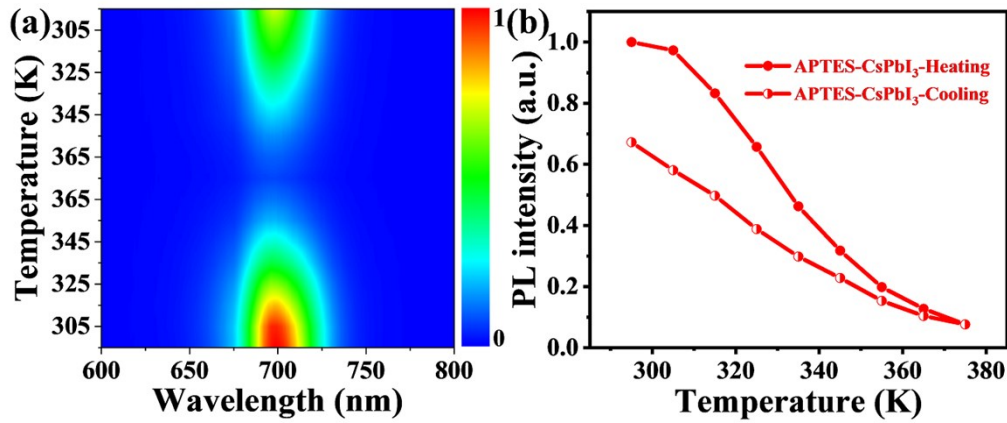
**Fig. S11** The energy-dispersive X-ray spectrum (a) and the time-resolved PL decay (b) of the CsPbI<sub>3</sub>@Ta<sub>2</sub>O<sub>5</sub> NCs.



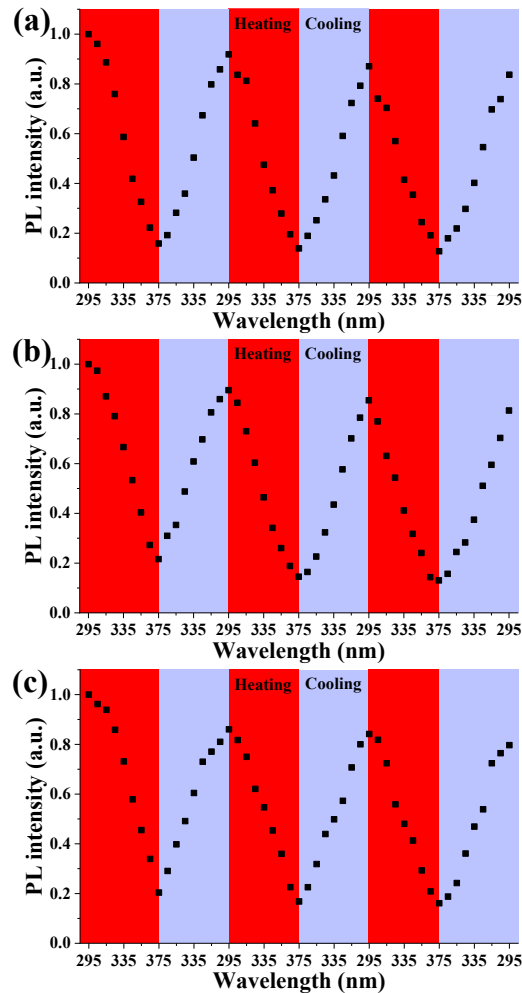
**Fig. S12** The energy-dispersive X-ray spectrum (a) and the time-resolved PL decay (b) of the CsPbI<sub>3</sub>@ZrO<sub>2</sub> NCs.



**Fig. S13** Pseudocolor map of PL spectra for APTES-CsPbI<sub>3</sub> NCs recorded at different time intervals after toluene dispersion is floated onto water forming a toluene/water interface. Pseudocolor map of PL spectra for APTES-CsPbI<sub>3</sub> NCs recorded at different time intervals after UV-light irradiation. (c) The corresponding integrated PL intensity as a function of time. The integrated PL intensity is normalized to the initial value.



**Fig. S14** Pseudocolor map of temperature-dependent PL spectra for APTES-CsPbI<sub>3</sub> NCs via heating/cooling cycles from room temperature to 375 K. (b) The corresponding integrated PL intensity as a function of temperature. The integrated PL intensity is normalized to the initial value.



**Fig. S15** Thermal cycling measurements for (a) CsPbI<sub>3</sub>@SiO<sub>2</sub> NCs, (b) CsPbI<sub>3</sub>@Ta<sub>2</sub>O<sub>5</sub> NCs and (c) CsPbI<sub>3</sub>@ZrO<sub>2</sub> NCs. The integrated PL intensity is normalized to the initial value.

**Table S1** PLQY for CsPbX<sub>3</sub> NCs and CsPbX<sub>3</sub>@Oxide core-shell NCs.

<b>Sample</b>	<b>PLQY</b>	<b>Sample</b>	<b>PLQY</b>
CsPbI <sub>3</sub>	66.9%	CsPbBr <sub>3</sub>	85.3%
CsPbI <sub>3</sub> @SiO <sub>2</sub>	51.1%	CsPbBr <sub>3</sub> @SiO <sub>2</sub>	66.9%
CsPbI <sub>3</sub> @Ta <sub>2</sub> O <sub>5</sub>	57.7%	CsPbCl <sub>3</sub>	1.22%
CsPbI <sub>3</sub> @ZrO <sub>2</sub>	59.7%	CsPbCl <sub>3</sub> @SiO <sub>2</sub>	1.03%