

Supporting Information for

High-quality all-inorganic CsPbBr₃ single crystals prepared by a facile one-step solution growth method

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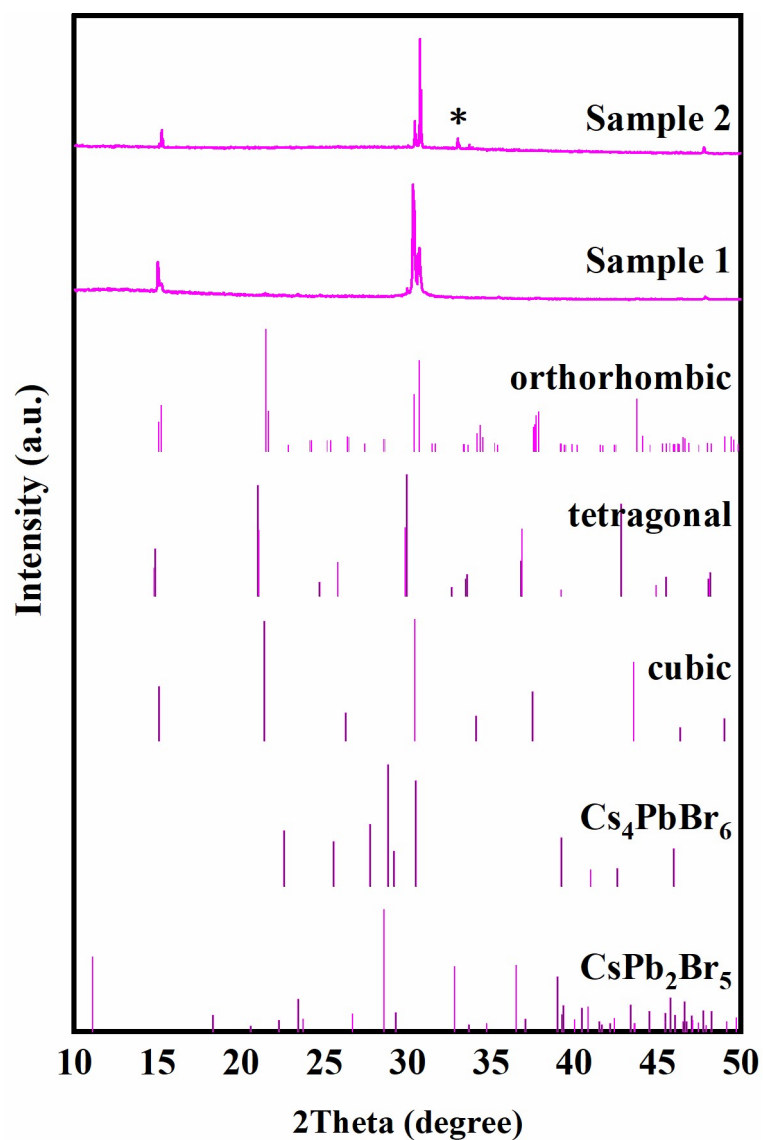


Figure S1. Powder XRD patterns of the as-grown CsPbBr₃ single crystal (**Sample 1**), *HPS* grown CsPbBr₃ single crystal (**Sample 2**), orthorhombic CsPbBr₃, tetragonal CsPbBr₃, cubic CsPbBr₃, Cs₄PbBr₆, and CsPb₂Br₅. The diffraction peak (labeled as *) in **Sample 2** is related to (310) of CsPb₂Br₅.

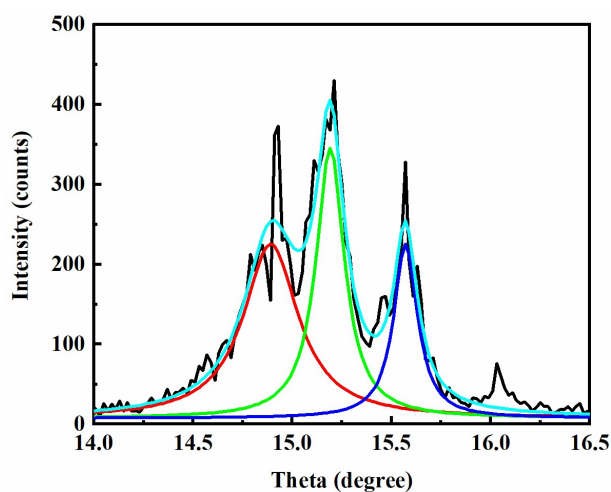


Figure S2. High-resolution XRD rocking curve spectrum (black) of CsPbBr₃ single crystals prepared using *HPS* method. The XRD rocking curve spectrum was fitted using the Lorentz function (shown in red, green, blue, and cyan lines). Here, the multi-diffraction peaks may be caused by complex multi-phase (CsPbBr₃/CsPb₂Br₅) of the sample, in which the peak at 15.09° is assigned to (002) of CsPbBr₃.

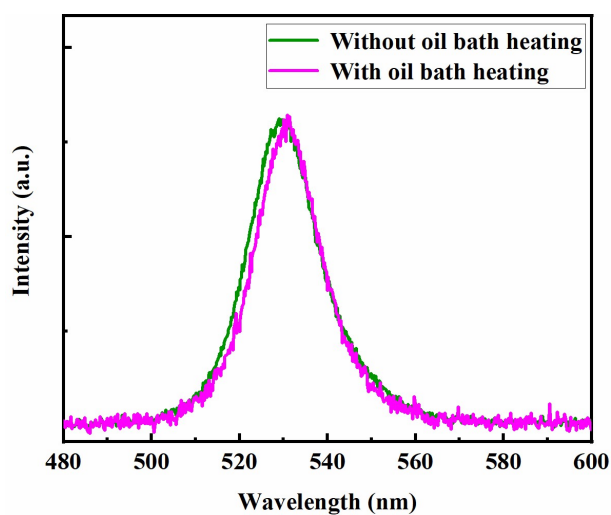


Figure S3. PL spectra of CsPbBr₃ single crystals prepared with and without oil bath heating.

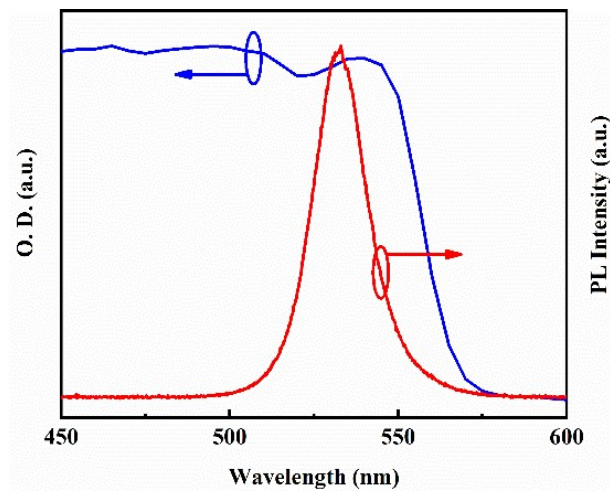


Figure S4. PL and absorption spectra of CsPbBr₃ single crystal.

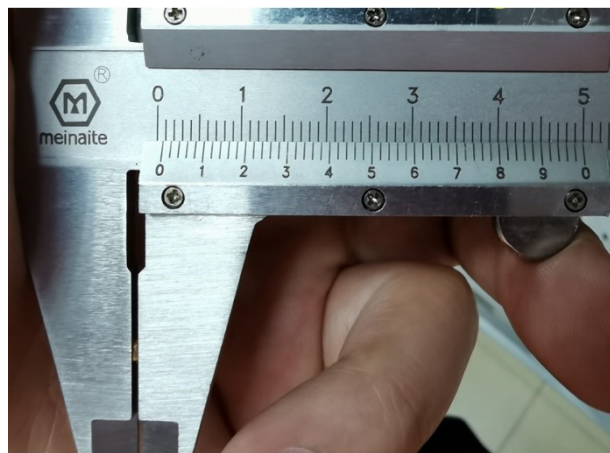


Figure S5. The thickness of the selected CsPbBr₃ single crystals for I - V measurements.

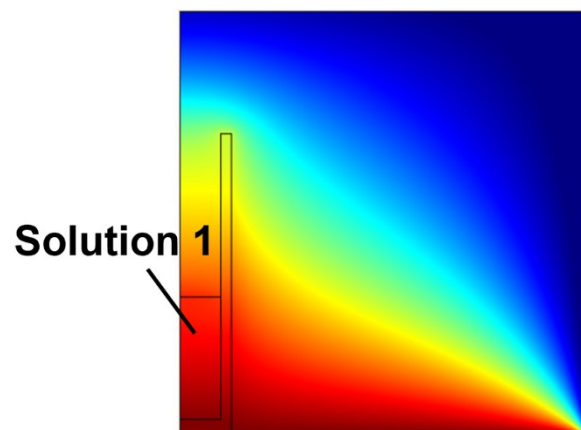


Figure S6. Calculated steady-state temperature field distribution of the precursor solution (labeled as solution 1) heated on the hot table.

Table S1. Comparison of trap density of CsPbBr₃ single crystal reported in this work and related results reported previously.

Material	Method	Trap density (cm ⁻³)	Reference
CsPbBr ₃	Inverse temperature crystallization	1.1×10 ¹⁰	1
CsPbBr ₃	Bridgman	1.9×10 ⁹	2
CsPbBr ₃	Bridgman	6.3×10 ¹⁰	3
CsPbBr ₃	Antisolvent vapor	2.8×10 ¹⁰	4
CsPbBr ₃	Low-temperature crystallization in water	1.7×10 ¹⁰	5
CsPbBr ₃	Inverse temperature crystallization	7.6×10 ¹⁰	5
CH ₃ NH ₃ PbBr ₃	seed-induced heterogeneous nucleation	2.6×10 ¹⁰	6
CH ₃ NH ₃ PbBr ₃	cavitation-triggered asymmetric crystallization	~ 10 ¹¹	7
CH ₃ NH ₃ PbI ₃	Inverse temperature crystallization with ligand regulation	7×10 ¹⁰	8
CH ₃ NH ₃ PbBr ₃	liquid-diffused separation induced crystallization	4.4×10 ⁹	9
CH ₃ NH ₃ PbI ₃	Inverse temperature crystallization	1.4×10 ¹⁰	10
CH ₃ NH ₃ PbBr ₃	Inverse temperature crystallization	3×10 ¹⁰	10
CsPbBr ₃	HPS with an oil bath heating	5.1×10 ⁹	This work

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