MAPbBr$_{3-x}$I$_x$ Crystals Improved by Accurate Solution-Grown Procedure for Alpha Particle Detection

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High-quality organic–inorganic halide perovskite brings an opportunity for ideal nuclear radiation detection due to its large carrier mobility, lifetime, and heavy atoms. However, the relatively low bulk resistivity restricts its development for alpha particles and X/γ-ray detectors. Here, we report on the MAPbBr$_{3-x}$I$_x$ (MPB:I) perovskite crystals grown using the modified inverse temperature crystallization method. Based on the measured supersolubility and solubility curves, the heating procedure was accurately designed to control the nucleation and growth rate. The resulting MPB:I single crystals exhibit higher bulk resistivity of $1.4 \times 10^9$ Ω cm, which is 10 times that obtained from the traditional inverse temperature crystallization method. Finally, an uncollimated 0.8 μCi $^{241}$Am @5.48 MeV alpha particles source was adopted to evaluate the MPB:I single crystals at room temperature condition. The electron mobility ($\mu_e$) and mobility-lifetime product ($\mu\tau_e$) of (264.6 ± 6.5) cm$^2$ V$^{-1}$ s$^{-1}$ and $(2.6 \pm 0.3) \times 10^{-3}$ cm$^2$/V·s, respectively, were obtained.

Keywords: perovskite, solution grown crystals, alpha particle detection, MAPbBr$_{3-x}$I$_x$, modified inverse temperature crystallization method

INTRODUCTION

Semiconducting nuclear detectors have attracted many interests to achieve the higher spatial and energy resolution due to the direct photoelectric conversion [1]. Hybrid metal halide perovskites, a new type of semiconductor material, have shown great competitive in photovoltaic and photoelectric application field because of the chemically robust, low trap density [2, 3], long carrier lifetime [2, 4, 5], and excellent charge transport performance [6]. Thus, both polycrystalline and single-crystalline perovskites have been used for photodetectors [7, 8], X-ray detectors [9–11], and solar cells [12–15].

Although organic–inorganic halide perovskites MAPbBr$_3$ (MPB) single crystals (SCs) has been proven to be a sensitive material for X-ray radiation detectors, the detection performance is still restricted by its large leakage current [16, 17]. Low bulk resistivity is significantly affected by the types and concentrations of defects in MPB SCs, which is mainly determined by the crystal growth. Some modified methods have been reported by different research groups, such as the modified antisolvent vapor-assisted crystallization methods [18], settled temperature and controlled antisolvent diffusion system [19], and seed-crystal inverse temperature crystallization (ITC) method [20]. The obtained resistivity of MPB SCs is mostly in the range of $1 \times 10^7$–5 $\times 10^8$ Ω cm, which is still limited for the development of nuclear detectors.
Therefore, currently, further increasing the resistivity is imperative. Huang's group [21] reported that the charge carrier concentration was reduced in the MPB bulk crystal by dopant compensation of chlorine so that the carrier mobility and lifetime have been improved. For I−-doped MPB:I, there are few literatures on the photoelectric performance, especially radiation detection [22, 23].

Here, we report the MPB:I perovskite crystals grown by a modified ITC (MITC) method. The heating rate is controlled accurately based on the obtained solubility and supersolubility curves. Then, the X-ray diffraction (XRD), scanning electron microscopy, and UV–visible–near infrared spectrum measurements are employed for the characterization of the crystal quality. Finally, the photoresponse and charge carrier transport behaviors are evaluated by light-emitting diode light and alpha particles source, respectively.

EXPERIMENTAL

Materials

The original materials, methylamine (CH₃NH₂) aqueous solution (30 wt-% in water), were purchased from Sinopharm Chemical Reagent Co., Ltd, China. Hydrobromide acid (40 wt-% in water) and N, N-dimethylformamide (DMF, 99.8%) were purchased from Kermel Reagent Ltd, China. Lead bromide (PbBr₂, 98%), and lead iodide (PbI₂, 98%) were purchased from Aladdin Reagent Ltd., China. All these original materials were employed directly without any further purification.

Synthesis of MABr

The hydrobromide acid was added into the CH₃NH₂ with the molar ratio of 1:1.2. After chemical reaction and full cooling, the well-mixed solution was evaporated at ~70°C in the evaporator for ~6 h to obtain crude white MABr (MA = CH₃NH₂Br) powders. After washing by absolute ethanol (99.8%) and recrystallization in anhydrous diethyl ether (99.0%) for several times, the purified MABr crystalline powders were finally dried in a vacuum oven at 60°C for 24 h.

Crystallization of MPB:I

A MITC method was employed to grow MPB:I single crystals. The MABr, PbBr₂, and PbI₂ with the molar ratios of 7:6:1 were dissolved in single solvent DMF for ~60 min in a magnetic stirring apparatus. The solution concentration was controlled at 1.6, 1.7, 1.8, 1.9, 2.0, and 2.1 M in DMF. After filtering using a 0.22-μm pore size filter membrane, the solution was for standby. To reduce the crystal growth rate of MPB:I, the controlling temperature was increased very slowly at 1–2°C/day and the holding time (~48 h) for 42, 45, 50, 55, 60, and 65°C, respectively.

Characterization

XRD patterns of MPB:I single crystals were collected using D/Mx2500PC with Cu Kα1 from 10 to 60° (2θ) under 40 kV tube voltage and 40 mA tube current. Scanning electron microscopy and energy-dispersive spectral maps were measured using a JEOL (JSM-7500F) scanning electron microscope with an Oxford (X-Max50) energy-dispersive spectrometer. UV-VIS-NIR spectrum measurements were performed using a UV-visible spectrophotometry (UV-3150, Shimadzu, Japan). ZHD-300 high-vacuum resistance evaporation coating machine was used for Au electrodes preparation. Current–voltage (I–V) and current–time (I–t) measurements were evaluated using a Keithley 6517b electrometer/high resistance system. The charge transport properties of the MPB:I crystals were evaluated by an uncollimated 241Am @5.48 MeV particle source at room temperature in air. Moreover, an ORTEC 710 high voltage supply, a charge sensitive preamplifier (ORTEC 142), and a shaping amplifier (ORTEC 570) with an optimized shaping time of 2 μs were used during the measurement. Finally, a standard Imdetek AMCA-01 multichannel analyzer was employed for the pulse height spectra acquisition.

RESULTS AND DISCUSSION

Crystal Growth and Structure

Because the priorities of halogen solubility in precursor solution DMF are shown below: I > Br > Cl, the replacement of Br by I can maintain the driving force in nucleation and growth processes under higher supersaturation [22]. The supersolubility was measured by a series of solution under the initial concentrations from 1.6 to 2.1 M (2.1 M cannot be completely dissolved at room temperature). After being sealed completely, two batches of solution were placed in a temperature-controlled water tank, in which the temperature could be controlled with a maximum fluctuation of 0.1°C.

For supersolubility measurement, the solution was heated at a rather low speed (1–2°C/day). In the solution growth process, once we observed tiny crystals the first time, the concentration at this temperature was recognized as the supersolubility. Then, a series of the same concentration solution was chosen for solubility measurement. The solution was kept preserved at a certain constant temperature for a long holding time (~48 h) to ensure that the crystal growth process is completed. Based on the solute mass (the raw material weight), the initial solution volume, and the weight of crystals, the concentration of solubility at a temperature range from 40 to 65°C was obtained.

Three processing zones (nucleation zone, growth zone, and dissolution zone) were divided by the two curves (supersolubility and solubility curves), as shown in Figure 1A. First, in the dissolution zone (Part III), all the raw materials were dissolved in the solution. With the temperature increasing, the solubility of precursors decreases, and the solution enters the nucleation zone (Part I); the tiny crystals began to occur. Then, to avoid the new crystal nucleation, the solution is maintained in the growth zone (Part II).

During the traditional ITC growth, the rapid heating usually leads to higher nucleation rate; thus, the grains are in small sizes because of lots of nucleation. Besides, the larger supersaturation of MPB:I in the precursor solution also results in the fast nucleation. Therefore, the crystal size is limited, and the crystals are usually overlapped, as shown in Figure 1B.

To reduce the nucleation and growth rates, the heating process was taken into account. As a result, fewer numbers of MPB:I
tiny crystals were observed at the beginning of the crystal growth process by the MITC method. After \( \sim 10 \) days, less than four single crystals were observed in the precursor solution, with the average dimension over \( 5 \times 5 \times 2 \) mm\(^3\) (Figure 1C).

The color of MPB:I single crystals is dark red when iodide was introduced, from Figures 1B,C. Comparing Figures 1B,C, surface roughness and crystal morphology are improved by the MITC method, and the stacking-free MPB:I SCs are obtained.

Furthermore, the I content was characterized by energy dispersive spectroscopy, as shown in Figure 2A. The obtained ratio of Pb/Br/I is 13.45:36.96:0.57. The resulting value of \( x \) in MAPb\(_{3-x}\)I\(_x\) is 0.04. According to the molar ratio of the raw materials MABr, PbBr\(_2\), and PbI\(_2\) (7:6:1), the nominal doping ratio of \( x \) in MAPb\(_{3-x}\)I\(_x\) is 0.286. The actual I content is lower than the theory value, which may be attributed to the larger ionic radius of iodine \([ABX_3]; ionic radius r_X, 0.22 \text{ nm} (r_I) > 0.196 \text{ nm} (r_{Br})\), making it hard to enter into the crystal lattice in MAPbBr\(_3\) [24].

Figure 2B shows the XRD patterns of the MPB:I and MPB crystals. The sharp peaks in each curve suggest the pure phase of single crystals, without any impurity phase peaks. The (100) peaks of the MPB and MPB:I crystals appeared at 2\( \theta \) of 15.22 and 15.51\(^\circ\), respectively. Using the equation \( d = \lambda/(2\sin\theta) \), the corresponding lattice constants are obtained. Therefore, it clearly suggested that the lattice can be adjusted by the compensation of the doped iodine. Furthermore, the UV-VIS-NIR spectra of the MPB (thickness of 4.64 mm) and MPB:I (thickness of 3.62 mm) were measured from 200 to 800 nm, as shown in Figure 2C.

Based on the spectrum and Tauc law [25], the bandgaps of the direct-gap semiconductor were obtained, which decreased from 2.21 to \( \sim 2.19 \text{ eV} \), as seen in Figure 2D.

**Resistance and Photoresponse**

Here, two representative MPB:I crystals were adopted, named MPB:I-1 (traditional ITC method, 5.82 \( \times \) 5.11 \( \times \) 3.14 mm\(^3\)), MPB:I-2 (modified ITC method, 5.27 \( \times \) 4.73 \( \times \) 1.03 mm\(^3\)), respectively. Au/MPB:I/Au devices were fabricated by thermal evaporation with thickness of 80 nm Au electrode.

The \( I-V \) curves of the above MPB:I single crystals are shown in Figure 3A. The resistivity of MPB:I-1 is 1.4 \( \times \) 10\(^8\) \( \Omega \) cm. As comparison, the reported resistivity of solution-processed Pb-based perovskite (MPB and MPI) single crystals exhibits a large range from 10\(^7\) to 10\(^8\) \( \Omega \) cm [26–28]. The resistivity is enhanced by the MITC method, which reaches (1.4 \( \pm \) 0.02) \( \times \) 10\(^9\) \( \Omega \) cm in as-grown MPB:I-2 (as shown in Figure 3B).

To investigate the photoresponse performance, the MPB:I-2 crystal was illuminated by a 375-nm light-emitting diode UV light with the power of 14.09 mW/cm\(^2\) at bias range from 1 to 50 V (as shown in Figure 3C). In Figure 3C, we can find that the values of dark current of 1 and 50 V are 2.4 and 315 nA,
respectively. There are more than 50 times difference between 1 and 50 V. The baseline shift phenomenon appeared as the bias increases. In our past research, the baseline shift phenomenon often happens on current–time curve measurements of 3D Pb-based perovskites samples. Maybe, Br ion migration leads to this problem.
According to the ion-beam-induced charge [29] and photoconductivity [10], the charge collection efficiency is not linear and reaches saturation gradually as the bias voltage increases. However, in this work, the \( \Delta I (I_{\text{on}} - I_{\text{off}}) \) is linear and not saturated even at bias of 50 V (as shown in Figure 3D).

### \( \alpha \) Particle Detection Characterization

Charge carrier mobility (\( \mu \)) and mobility-lifetime product (\( \mu \tau \)) of the MPB:1-2 sample are evaluated by the alpha particles. The measurements were carried out at room temperature and air condition. Generally, the incident depth of the \( \alpha \) particles is near the surface of the crystal detector, with the distance of dozens of micrometers (\(< 50 \mu \text{m}\)) [18, 30]. Therefore, we use the simplified single-carrier approximation Hecht equation (Equation 1) [29] to estimate the electron mobility-lifetime product (\( \mu \tau \)) of the MPB:1-2 sample.

\[
CCE = \frac{\mu \tau U}{d^2} \left( 1 - e^{-\frac{d^2}{2\mu \tau U}} \right)
\]

where \( U \) is the applied voltage, and \( d \) is the device thickness.

Simultaneously, we used time-of-flight technique to analyze pulse shape rise time and extract the carrier mobility (\( \mu \)) of MPB:1 detector. Figure 4A shows the transient pulse for the MPB:1-based detector under 50 V. Generally, a 10–90% amplitude rise time \( t_{10-90\%} \) was selected for further processing. Through analyzing the pulse height rise time distribution, the carrier drift time \( (t_{\text{dr}}) \) was acquired. Under the bias from 10 to 50 V, the \( t_{\text{dr}} \) were 5.01, 4.13, 3.16, 2.98, and 2.65 \( \mu \text{s} \), respectively. The mobility \( \mu \) is given by

\[
\mu = \frac{V_{\text{dr}}}{E} = \frac{d^2}{V \times t_{\text{dr}}}
\]

where \( V \) is the bias, \( t_{\text{dr}} \) is carrier drift time, and \( d \) is the thickness of the MPB:1 detectors.

The estimated electron mobility (\( \mu_e \)) for the MPB:1 detector is \((264.6 \pm 6.5) \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}\), seen in Figure 4B, which is higher than the reported \( \mu_e \) for MPB and MPI single crystals with the value in the range of \( 24–167 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}\) [2, 4, 31–35] using the transient photocurrent response measurements. The high mobility of MPB:1 may be attributed to the reduction in trap-state density and long carrier diffusion in organolead trihalide perovskite single crystals.

Figure 4C shows the pulse height spectra vs. applied bias for MPB:1-3 detector, which were illuminated by \( \text{^{241}Am} \) alpha particles source. With the applied bias increases, the cutoff edges of the curves reasonably shift to the high channel direction. The electron mobility-lifetime product (\( \mu \tau \)) is estimated to be \((2.6 \pm 0.3) \times 10^{-3} \text{ cm}^2/\text{V} \) (Figure 4D) by fitting the channel (cutoff edges of the curves) vs. the applied bias using the single-carrier Hecht equation.

### CONCLUSIONS

High-resistivity MPB:1 bulk crystals have been grown by the MITC method. Both supersolubility and solubility curves were acquired to tailor the growth rate by controlling the heating process. The Au/MPB:1/Au structure device exhibits a higher resistivity of \( 1.4 \times 10^9 \Omega \text{ cm} \). The electron mobility-lifetime (\( \mu \tau \)) products are \((2.6 \pm 0.3) \times 10^{-3} \text{ cm}^2/\text{V} \) by fitting the corresponding pulse height spectra as a function of the applied bias using the Hecht equation. The electron mobility is estimated to be \((264.6 \pm 6.5) \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}\), using a 0.8 \( \mu \text{Ci} \text{^{241}Am} \) alpha particle source at 5.48 MeV measured at room temperature under air environment.

### DATA AVAILABILITY STATEMENT

All datasets generated for this study are included in the article/supplementary material.

### AUTHOR CONTRIBUTIONS

XL and YX conceived the idea. XL, BZ, and YX designed the experiments. JF and YH synthesized MABr, the perovskite precursors, and crystals growth. MX carried out scanning electron microscopy and EDS measurements. XL and DZ performed the UV-VIS-NIR spectrum measurements, I-V measurements, and I-t measurements. XL and HZ performed alpha particle detection measurements. XL and YX analyzed the data. XL, DZ, and YX drafted the manuscript. BZ, WJ, and YX advised on the overall experiments with critical comments, and finalized the manuscript. All the authors contributed to interpretation of the results and revision of the manuscript.

### FUNDING

This work was supported by the National Natural Science Foundations of China (Nos. U1631116, 51872228, and 51802262). This project was also supported by the National Key Research and Development Program of China (2016YFE0115200) and the Natural Science Basic Research Plan in Shaanxi Province of China (2019ZDLGY04-07), the Natural Science Foundation of Shaanxi Province (2019JQ-459), and the Fundamental Research Funds for the Central Universities (3102018jcc036, 3102019TS0408).

### REFERENCES

1. Szeles C. CdZnTe and CdTe materials for X-ray and gamma ray radiation detector applications. Phys Status Solidi B. (2004) 241:783–90. doi: 10.1002/pssb.200304296

2. Shi D, Adinolfi V, Comin R, Yuan M, Alarousu E, Buin A, et al. Low trap-state density and long carrier diffusion in organolead trihalide perovskite single crystals. Science. (2015) 347:519–22. doi: 10.1126/science.aai2725

3. Lian Z, Yan Q, Gao T, Ding J, Lv Q, Ning C, et al. Perovskite CH\textsubscript{3}NH\textsubscript{3}Pb\textsubscript{3}I\textsubscript{6} (CJ) single crystals: rapid solution growth, unparalleled crystalline quality,
and low trap density toward 108 cm$^{-3}$. J Am Chem Soc. (2016) 138:9409–12. doi: 10.1021/jacs.6b05683

4. Dong Q, Fang Y, Shao Y, Mulligan P, Qiu J, Cao L, et al. Electron-hole diffusion lengths$\sim$175 nm in solution-grown CH$_3$NH$_3$PbI$_3$ single crystals. Science. (2015) 347:967–70. doi: 10.1126/science.aaa5760

5. Bi Y, Hutter EM, Fang Y, Dong Q, Huang J, Savenije TJ. Charge carrier lifetimes exceeding 15 ms in methylammonium lead iodide single crystals. J. Phys Chem Lett. (2016) 7:923–8. doi: 10.1021/acs.jpclett.6b00269

6. Heo JH, Im SH, Noh JH, Mandal TN, Lim C-S, Chang JA, et al. Efficient organic–inorganic hybrid heterojunction solar cells containing perovskite compound and polymeric hole conductors. Nat Photonics. (2013) 7:486–91. doi: 10.1038/nphoton.2013.80

7. Liu Y, Zhang Y, Zhao K, Yang Z, Feng J, Zhang X, et al. A 1300 mm$^{-2}$ ultra-high-performance digital imaging assembly using high-quality perovskite single crystals. Adv Mater. (2018) 30:1707314. doi: 10.1002/adma.201707314

8. Ahmadi M, Wu T, Hu B. A review on organic–inorganic halide perovskite photodetectors: device engineering and fundamental physics. Adv Mater. (2017) 29:1605242. doi: 10.1002/adma.201605242

9. Yakunin S, Sytyyk M, Krieger D, Shrestha S, Richter MJ, et al. Detection of X-ray photons by solution-processed organic-inorganic perovskites. Nat Photonics. (2015) 9:444–9. doi: 10.1038/nphoton.2015.82

10. Pan W, Wu H, Luo J, Deng Z, Ge C, Chen C, et al. Cs$_2$Ag$_x$Br$_{3-y}$ single-crystal X-ray detectors with a low detection limit. Nat Photonics. (2017) 11:726–32. doi: 10.1038/s41566-017-0012-4

11. Panneerselvam DM, Kabir MZ. Evaluation of organic–inorganic photodetectors for direct conversion X-ray imaging detectors. J Mater Sci Mater El. (2017) 28:1–8. doi: 10.1007/s10854-016-6409-5

12. Yin W-J, Yang J-H, Kang J, Yan Y, Wei S-H. Halide perovskite materials for solar cells: a theoretical review. J Mater Chem A. (2015) 3:8926–42. doi: 10.1039/C5TA05333A

13. Kim HS, Seo JY, Park NG. Material and device stability in perovskite solar cells. ChemSusChem. (2016) 9:2528–40. doi: 10.1002/cssc.201600915

14. Grancini G, Roldáncarmona C, Zimmermann I, Mosconi E, Lee X, Martineau D, et al. One-year stable perovskite solar cells by 2D/3D interface engineering. Nat Commun. (2017) 8:15684. doi: 10.1038/ncomms15684

15. Yang D, Sano T, Yaguchi Y, Sun H, Sasabe H, Kido J. Achieving 20% efficiency for low-temperature-process inverted perovskite solar cells. Adv Funct Mater. (2018) 28:1807556. doi: 10.1002/adfm.201807556

16. Onoda-Yamamuro N, Matsuo T, Suga, H. Dielectric study of CH$_3$NH$_3$PbI$_3$ ($X = $ Cl, Br, I) J Phys Chem Solids. (1992) 53:935–9. doi: 10.1016/0022-3697(92)90121-S

17. Meloni S, Moehl T, Tress W, Frankevicius M, Saliba M, Lee YH, et al. Ionic polarization-induced current–voltage hysteresis in CH$_3$NH$_3$PbI$_3$ perovskite solar cells. Nat Commun. (2017) 8:15684. doi: 10.1038/ncomms15684

18. Wang L, Yuan G, Duan R, Huang F, Wei T, Liu Z, et al. Tunable bandgap in hybrid perovskite CH$_3$NH$_3$PbI$_3$Cl$_x$ single crystals and photodetector applications. AIP Adv. (2016) 6:045115. doi: 10.1063/1.4948312

19. Zhang Y, Liu Y, Li Y, Yang Z, Liu SF. Perovskite CH$_3$NH$_3$Pb(Br$_x$I$_{1−x}$)$_3$ single crystals with controlled composition for fine-tuned bandgap towards optimized optoelectronic applications. J Mater Chem C. (2016) 4:9172–8. doi: 10.1039/C6TC03592B

20. Chonghe Lee, Xionggang L, Weizhong D, Liming F, Yonghui G, Ming Z. Formability of ABX$_3$ ($X = $ F, Cl, Br) halide perovskites. Acta Crystallogr. (2008) 64:702–7. doi: 10.1107/S0021889808032734

21. Tauc J, Menth A. States in the gap. J Non-Cryst Solids. (1972) 8:569–85. doi: 10.1016/0022-3093(72)90194-9

22. Saidaminov MI, Haque MA, Almutlaq I, Sarmah S, Miao XH, Begum R, et al. Inorganic lead halide perovskite single crystals: phase-selective low-temperature growth, carrier transport properties, and self-powered photodetection. Adv Opt Mater. (2017) 5:1600704. doi: 10.1002/adom.201600704

23. Rakita Y, Kedem N, Gupta S, Sadhanala A, Kalchenko V, Böhm ML, et al. Low-temperature solution-grown CsPbBr$_3$ single crystals and their characteristics. Cryst Growth Des. (2016) 16:5717–25. doi: 10.1021/acs.cgd.6b01119

Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest. The handling editor declared a past co-authorship with the authors BZ and YX.