



A facile, time-saving fabrication method of high purity CsPbBr₃ films for efficient solar cells

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ABSTRACT

All-brominated cesium lead bromide (CsPbBr₃) perovskite solar cells (PSCs) have drawn growing attention owing to unprecedented stability in the family of perovskites. It usually employs a two-step method to prepare CsPbBr₃ films owing to the low solubility of CsBr in the common solvents. The researchers have to provide enough Cs source through increasing the reaction time or repeat times of CsBr deposition so as to obtain the target CsPbBr₃ films. The long preparation period and tedious process increase the time and economic cost, which may limit the practical application of CsPbBr₃ solar cells. Herein, a facile method is proposed to prepare CsPbBr₃ films, in which appropriate CsBr is pre-introduced into PbBr₂ solutions in the first step, and CsBr/2-methoxyethanol solution (~35 mg mL⁻¹) is used as solution conversion medium in the second step. Benefitting from the pre-introduction of CsBr in the first step, a single phase of CsPbBr₃ films with full coverage can be converted for only once CsBr supplementation. After integrating the CsPbBr₃ films into a solar cell, it delivers a power conversion efficiency (PCE) of 7.48% basing on the configuration of FTO/TiO₂/CsPbBr₃/Carbon. This work largely shortens the preparation time of CsPbBr₃ films, which may provide a practical way for the large-scale production of CsPbBr₃ based devices.

1. Introduction

All-inorganic perovskite CsPbBr₃ attracts more attentions in optoelectronic devices such as solar cells, photodetectors and light-emitting devices, due to its superior optoelectronic properties and high stability in air conditions. The highly compatible with the carbon electrode in ambient condition makes CsPbBr₃ as a reasonable candidate to full solution-processed solar cell fabrication [1–4]. The big challenge to CsPbBr₃ films for solar cells is to fabricate pure and smooth thick perovskite layer due to the poor solubility of CsBr in traditional organic solvents. The common way is so-called two-step method, in which PbBr₂ films were firstly prepared, and then exposed to CsBr methanol solution for an optimized time, usually at least 10 min, to obtain desired CsPbBr₃ films [1–8]. However, the reaction time is so sensitive that either too short or too long reaction time could generate byproducts like CsPb₂Br₅

and Cs₄PbBr₆, which limits the photovoltaic performance and lead to other side effects [9–11].

To exclude the byproducts, Duan et al. [12] proposed a multi-steps solution processed technique to prepare high purity CsPbBr₃ films. In this method, a drop of CsBr/methanol was spin-coated onto the PbBr₂ films, followed by annealing at 250 °C for 5 min, then repeated this process for several cycles. According to their report, the components, crystal structure, and morphology of the as-prepared perovskite films are strongly dependent on the preparation cycles. Especially, it needs at least four cycles which cost no less than 20 min to obtain pure CsPbBr₃ films in their method. Obviously, it is a time-consuming method to prepare pure phase CsPbBr₃ films. Meanwhile, the methanol used to dissolve CsBr could decompose CsPbBr₃ during the treatment, which may resulting in poor morphology and more defects in the final products [13]. In order to eliminate methanol as the solvent, Feng et al. replaced

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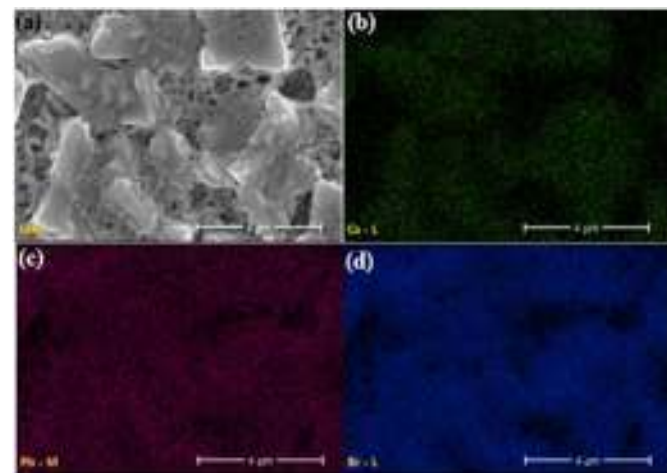


Fig. 3. (a) The SEM image of EDX result of the Cs-Pb-Br film prepared from the solution of 2:1; The mapping of the film elements, (b) Cs; (c) Pb; (d) Br.

a speed of 3000 rpm for 30 s, followed by an anneal treatment at 100 °C for 5 min in a N₂-filled glovebox. In the second step, 90 μ L CsBr/2-methoxyethanol solution (~ 35 mg mL⁻¹) (see Fig. S1) was dropped on the matrices and stay it for 60 s, then the substrates were spin-coated at 2000 rpm for 30 s in ambient air atmosphere. The matrices are further annealed at 250 °C for 5 min to form Cs-Pb-Br films. Finally, the carbon

electrode was deposited by using screen printing of carbon paste with drying at 100 °C for 10 min.

Characterization: The phase and morphology of PbBr₂ and Cs-Pb-Br films were measured by X-ray diffraction (PANalytical X'Pert³ Powder) and field-emission scanning electron microscopy (NovaTM NanoSEM 430), respectively. The UV-Vis spectrophotometer (Shimadzu UV-2550) was employed to get the absorption spectra of the PbBr₂ and Cs-Pb-Br films. A photoluminescence spectrometer (Edinburgh Instruments, LP 980) was utilized to obtain the photoluminescence (PL) spectra. The photovoltaic performance of solar cells was recorded using a Keithley 2400 under AM 1.5G simulated solar illumination. The incident photo-to-current collection efficiency spectrum (IPCE) of solar cells were performed an integrated system from Enlitech (QE-R3011).

3. Results and discussion

We first investigated the solubility of CsBr in PbBr₂/DMSO solution. As shown in Fig. 1, it is still a clear solution even the concentration of CsBr increases from 0 M to 1.0 M in the mixtures (Fig. 1a-c). However, the mixture becomes turbid when the concentration of CsBr increases to 1.5 M (Fig. 1d). In light of the dissolution state, we employ the solutions composed of different molar ratios between PbBr₂ and CsBr from 2:0, 2:0.5 and 2:1 to prepare the Cs-Pb-Br matrices in the first step for understanding the phase composition.

The as-prepared samples are examined by the X-ray diffraction (XRD). Fig. 2a shows the XRD patterns of the films prepared from



Fig. 4. (a) The normalized XRD curves of the Cs-Pb-Br films prepared from different solutions; Surface SEM images of Cs-Pb-Br films from different solutions; (b) 2:0; (c) 2:0.5; (d) 2:1. (e) $(Ah\nu)^2$ - $h\nu$ curve obtained from the UV absorption spectrum, and (f) the steady PL spectrum of the Cs-Pb-Br films film prepared from the solution of 2:1. The inset are the photograph of corresponding the Cs-Pb-Br films.



Fig. 5. (a) The device configuration of solar cells; (b) J - V curves of the best PSCs prepared from different solutions; (c) the steady output of best solar cells by applying a voltage at the maximum output point. (d) IPCE spectrum; (e) the J - V curves having highest V_{oc} ; (f) the distribution of PCE from a series of solar cells.

Table 1

The photovoltaic performance of best solar cells prepared from different solutions.

| Solution | V_{oc} (V) | J_{sc} (mA cm ⁻²) | FF(%) | PCE(%) |
|----------|--------------|---------------------------------|-------|--------|
| 2 : 0 | 1.13 | 1.70 | 51.84 | 0.99 |
| 2 : 0.5 | 1.27 | 4.02 | 73.05 | 3.73 |
| 2 : 1 | 1.41 | 7.15 | 74.07 | 7.48 |

different solutions. The film prepared from the solution of 2:0 are pure $PbBr_2$, which is confirmed by the characteristic peak at 18.5° corresponding to the (020) crystal plane [17]. In addition to having $PbBr_2$ in the film when 0.5 M CsBr is introduced into the solution, the obvious diffraction peaks can be indexed at 15.2° , 21.6° , 30.7° , 34.3° and 44.1° correspond to the (100), (110), (200), (210) and (220) planes of the $CsPbBr_3$ phase, respectively. Moreover, the film still shows the peaks at

11.7° , 23.4° , 29.4° and 47.9° correspond to the (002), (210), (213) and (420) planes of the $CsPb_2Br_5$ respectively [18], indicating the coexistence of $CsPb_2Br_5$, $CsPbBr_3$ and $PbBr_2$ in the Cs–Pb–Br films. When the concentration increased to 1.0 M, the $CsPbBr_3$ component increases in accompanied with the decreases of $PbBr_2$ and $CsPb_2Br_5$. We also study the surface morphology evolution by using scanning electron microscopy (SEM). Fig. 2b shows a SEM image of film prepared by pure $PbBr_2$ solution. The film exhibits an extremely smooth morphology with dense and compact structure, which is not beneficial for the permeation of CsBr solution in further step treatment. When including CsBr into the solution, the as-prepared films become much rougher with many pores and cracks on the surface (Fig. 2c and d). These rough morphologies have significant impacts on the permeation of CsBr/2-methoxyethanol solution in the second step treatment [19,20]. As shown in the insets of Fig. 2, the colours of the films evolves from transparent (Fig. 2b inset) to light yellow (Fig. 2d inset) when the films prepared from the solution

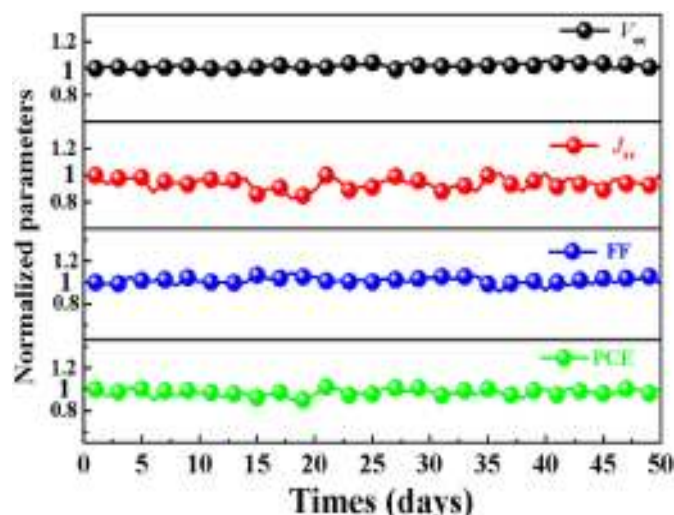


Fig. 6. The evolution of normalized parameters of solar cell at air atmosphere with temperature of $\sim 30^\circ\text{C}$ and relative humidity of $\sim 45\%$.

of 2:1, which is close to the intrinsic yellow of the CsPbBr_3 films. The UV/Vis absorption spectra of the films (see Fig. S2) also confirm the existence of CsPbBr_3 in the films, which is evidenced by an absorbance onset near 530 nm, corresponding to the bandgap of ~ 2.32 eV for CsPbBr_3 [21].

In order to learn the elements distribution in the films, we perform the energy-dispersive X-ray spectroscopy (EDX) experiments on the films prepared from the solution of 2:1 (Fig. S3). Besides the element from the substrates (Ti and O), Cs, Pb and Br element are also detected in the films, showing the existence of Cs in the films. These results strongly demonstrate that the element of Cs and Br are successfully introduced into the films. Fig. 3b–d are the elemental mapping of Cs, Pb and Br in the films, showing a uniform distribution of those elements in the films. Combined with the elemental ratio of EDX data (see Table S1), it is found that Cs:Pb:Br is close to 1:2:5 coincided with elemental ratio of the precursor solution of 2:1. Those results demonstrate that it is an effective way to supply the Cs source by introducing CsBr into the PbBr_2 solution in our first step treatment.

Fig. 4a is the normalized XRD curves of the Cs–Pb–Br films prepared from different solutions after reacting with CsBr/2-methoxyethanol solution, followed by an annealing treatment. Some PbBr_2 are still remained in the Cs–Pb–Br films prepared from the solution of 2:0. There are very weak diffraction peaks of CsPbBr_3 in the XRD pattern (blue line in Fig. 4a), indicating a tiny amount of CsPbBr_3 exists in the films due to Cs element deficiency. The phase of CsPbBr_3 begins to dominate in the Cs–Pb–Br films when 0.5 M CsBr is introduced into the PbBr_2 solution in the first step, but still exists PbBr_2 -riched CsPb_2Br_5 phase in the film (red line in Fig. 4a). Therefore, it is necessary to supply Cs source so as to obtain single phase CsPbBr_3 films. As expected, it produces pure CsPbBr_3 films when 1.0 M CsBr is added to the PbBr_2 solution (dark line in Fig. 4a). By checking with EDX analysis (see Fig. S4 and Table S2), it is found that Cs, Pb and Br are uniform distributed and the ratio is close to 1:1:3. Those results demonstrate that it is an effective way to get pure phase CsPbBr_3 . Fig. 4b to d show the morphology and related photos (inset figures) of the Cs–Pb–Br films after second step treatment. For the Cs–Pb–Br films prepared from the solution of 2:0, there are some small grains on the surface, which is different from the PbBr_2 films prepared in the first step (Fig. 2b). These small grains may be CsPbBr_3 grains, which is evidenced by the yellow dots on the films. There are apparent cracks in the Cs–Pb–Br films fabricated from the solution of 2:0.5 (Fig. 4c), which is consistent with the morphology of the mother matrix in the first step (Fig. 2c). The cracks in the Cs–Pb–Br films can act as the current-shunting pathways, leading to a serious current leakage and recombination loss [22–24]. Meanwhile, the film exhibits nonuniform yellow

color. In contrast, matrix prepared from the solution of 2:1 exhibits fully covered morphology, this feature is further confirmed in large scale SEM image of this film (Fig. S5). The photograph of the CsPbBr_3 film also exhibits uniform yellow color, demonstrating a uniform feature in a large scale. The UV–vis absorption spectrum of the Cs–Pb–Br film prepared from the solution of 2:1 shows an onset at ~ 530 nm (Fig. S6). Fig. 4e shows the Tauc plot of the film prepared from the solution of 2:1. The optical bandgap (E_g) is fitted to be ~ 2.3 eV, which is consistent with previous reports [10,25,26]. The characteristic bandgap of CsPbBr_3 can also be demonstrated by the steady PL spectrum (Fig. 4f), which shows an emission peak at ~ 536 nm, corresponding to an optical bandgap of ~ 2.3 eV. These results demonstrate that our method is an effective way to prepare high-purity CsPbBr_3 films through pre-introducing appropriate CsBr into PbBr_2 solution.

To examine the film performance in a full solution-processed device structure, we construct an HTL-free solar cell with a configuration of FTO/ TiO_2 / CsPbBr_3 /Carbon. Fig. 5a presents a cross-sectional SEM image of a solar cell sample, in which the CsPbBr_3 films are prepared from the solution of 2:1. It clearly indicates that the CsPbBr_3 grains are large enough to occupy the whole thickness of ~ 600 nm in active layer with dense arrangement. Fig. 5b shows the light J–V curves of the best PSCs prepared from different solutions. The PSC prepared from the solution of 2:1 has a power conversion efficiency (PCE) of 7.48%, with a short circuit current density (J_{sc}) of 7.16 mA cm^{-2} , a fill factor (FF) of 74.07%, and an open-circuit voltage V_{oc} of 1.41 V. Whereas the best PSCs fabricated from the solution of 2:0 and 2:0.5 exhibit very low PCE of 0.99% and 3.73%, respectively. Other photovoltaic parameters from different PSCs are summarized in Table 1. This performance comparison shows the importance of enhancing the purity and coverage of CsPbBr_3 phase in active layer of solar cell. Fig. 5c is the steady-state output near the maximum power point of the best PSC of the 2:1 device. A steady photocurrent density of 5.65 mA cm^{-2} and a steady PCE of 6.44% were obtained at a bias voltage 1.14 V. It is noted that the steady PCE is slightly lower than the value extracted from J–V curve under the reverse scan. The deviation might come from the hysteresis behavior which is usually observed in common PSC devices [11,27–30] (Fig. S7). The IPCE spectrum of the optimized solar cell is recorded and shown in Fig. 5d. The device shows strong light absorption ability in the range of 300– ~ 525 nm. The J_{sc} integrated from the IPCE spectra is 6.46 mA cm^{-2} , which is a little lower than that obtained from the J–V curve. This discrepancy might be related to the nonlinear properties of perovskite films [31]. It should be noted that a high V_{oc} of 1.5 V is achieved in a patch of PSCs prepared from the solution of 2:1. This value is one of the highest V_{oc} for the CsPbBr_3 solar cells based on unmodified configuration FTO/ TiO_2 / CsPbBr_3 /Carbon [32]. Fig. 5f summarizes the distribution of PCE collected from a series of PSCs. The average PCE is 5.86% of all samples demonstrating that our two-steps method is a reliable way to prepare efficient CsPbBr_3 solar cells.

The air stability of solar cells is also investigated. Fig. 6 shows the normalized photovoltaic parameters of an unencapsulated PSC under the open-air condition ($T \sim 30^\circ\text{C}$ RH $\sim 45\%$). The result shows that the photovoltaic parameters, including V_{oc} , J_{sc} , FF and PCE, are still stable over 50 days. The best-performance PSC still shows a PCE of 6.81%, V_{oc} of 1.41 V, J_{sc} of 7.21 mA cm^{-2} , and FF of 67% when it is stored in ambient condition for 90 days (see Fig. S8). The superior stability of CsPbBr_3 solar cells makes it become a competitive candidate to the commercialization of PSCs.

4. Conclusion

In summary, we fabricate high-quality CsPbBr_3 films by using a facile two-step spin-coating method, in which CsBr is introduced into PbBr_2 solution in the first step. The pre-incorporated CsBr in PbBr_2 solution can shorten the fabrication period and simplify the preparation process in comparison with the traditional dipping and multi-step solution processing methods. The CsPbBr_3 films are used to fabricate solar cell

devices with a configuration of FTO/TiO₂/CsPbBr₃/Carbon. The optimized samples can achieve a power conversion efficiency of 7.48%, with high stability in air conditions. These results demonstrate the efficient and time-saving method for preparation of CsPbBr₃ films in this work has a promising application in various optoelectronic devices.

Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.ceramint.2023.03.279>.

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