

Stitching Triple Cation Perovskite by a Mixed Anti-Solvent Process for High Performance Perovskite Solar Cells

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Abstract

With the rapid development of organic-inorganic lead halide perovskite photovoltaics, increasingly more attentions are paid to explore the growth mechanism and precisely control the quality of perovskite films. In this study, we propose a “stitching effect” to fabricate high quality perovskite films by using chlorobenzene (CB) as an anti-solvent and isopropyl alcohol (IPA) as an additive into this anti-solvent. Because of the existence of IPA, CB can be efficiently released from the gaps of perovskite precursors and the perovskite film formation can be slightly modified in a controlled manner. More homogeneous surface morphology and larger grain size of perovskite films were achieved via this process. The reduced grain boundaries ensure low surface defect density and good carrier transport in the perovskite layer. Meanwhile, we also performed the Fourier transform infrared (FTIR) spectroscopy to investigate the film growth mechanism of unannealed and annealed perovskite films. Solar cells fabricated

by using the “stitching effect” exhibited a best efficiency of 19.2%. Our results show that solvent and solvent additives dramatically influenced the formation and crystallization processes for perovskite materials due to their different coordination and extraction capabilities. This method presents a new path towards controlling the growth and morphology of perovskite films.

Introduction

Organic-inorganic hybrid halide perovskite has recently emerged as one of the most promising materials for the low-cost third generation photovoltaic technology. Over the past few years, the power conversion efficiency (PCE) of perovskite solar cells (PSCs) has rapidly increased from 3.8% to more than 22.1%. [1-2] During this process, most works focus on deposition methods, interface engineering, hysteresis in the current-voltage curves of PSCs and the application of new functional materials. [3-12] In order to make it more conducive to practical application, the degradation mechanism of perovskite in air and the fabrication process of larger sized devices are investigated by a lot of research groups. [13-17] As a result, precisely controlling the growth of perovskite crystals has also become an interesting topic to explore the fundamental properties and growth dynamics of high quality perovskite films for efficient solar cells.

A lot of efforts has been devoted to improve the morphology, coverage, grain size of perovskite films, which play an important role in the performance of PSCs. [18-20] It is well known that grain size is a key factor related to the diffusion length of carriers, the rate of carrier recombination and charge generation. [19] To well control the fabrication of dense and uniform perovskite films with large grain size is challenging

and meaningful. Compared with other deposition methods (such as vapor deposition, vapor-assisted deposition, atomic layer deposition etc.), solution deposition is a simple and low-cost way to fabricate high efficiency PSCs. [5-6,21-31] Several researches have been reported in obtaining high quality perovskite films based on growth engineering, solvent engineering and so on. In 2016, M. Dar and his coworkers studied the impact of chloride on the conversion of lead halide into $\text{CH}_3\text{NH}_3\text{PbI}_3$. [21] The surface coverage of the perovskite film was effectively improved and material defects were reduced by adding 2% chloride in perovskite precursors. In order to optimize perovskite film quality, solvent engineering, especially mixed-solvent process, was widely investigated. [5-6] Various solvents were employed in the dissolved solution of perovskite precursors, solvent annealing, the solution of iodide methylammonium (MAI) and so on. Huang's group firstly proposed the solvent annealing on organic-inorganic perovskite materials in 2014. [22] DMF was used in the annealing of perovskite which successfully enhanced the grain size of $\text{CH}_3\text{NH}_3\text{PbI}_3$. Unlike DMF, DMSO molecules have strong coordination effect with Pb^{2+} so that DMSO was widely used to retard the crystallization of PbI_2 and improve the quality of perovskite films. [5] Using the DMSO solvent annealing process, a mesoporous structure of PbI_2 and large grain size of $\text{CH}_3\text{NH}_3\text{PbI}_3$ were achieved in our precious studies. In addition to strongly polar solvents, some weakly polar solvents (ethanol, acetonitrile, isopropanol, etc.) were also investigated. Li et al. introduced acetonitrile solvent to successfully mediate the nucleation and grain growth of perovskite films. [23] Wu et al. and Sun et al., respectively, employed DMF/CB and DMF/IPA solvent annealing to obtained smooth

and dense surface morphology of $\text{CH}_3\text{NH}_3\text{PbI}_3$. [24-25] Alcohol-vapor annealing treatment on $\text{CH}_3\text{NH}_3\text{PbI}_3$ was reported by Liu et al., aiming to improve the crystal growth and increase the grain size of perovskite crystals. [3] Actually, such low-polar solvents have the weak coordination ability with perovskite precursors which could play an important role in the growth of perovskite crystals. Note that, weak coordination ability was brought up here because of the poor solubility of lead ions and weak electron donating ability. Seok and his group reported a new one-step deposition method using a mixed solution of GBL and DMSO as the solvent of perovskite precursors and toluene was employed as the anti-solvent in this one-step procedure to achieve homogeneous interface morphology of perovskite. [26] After that, Park and his group used diethyl ether as the anti-solvent to remove DMF solvent molecules in their experiment. [27] They made highly reproducible $\text{CH}_3\text{NH}_3\text{Pb}_3$ solar cells with an average PCE of 18.3% and the best efficiency of 19.7%. Since then, anti-solvent deposition method has become a common way to fabricate high quality perovskite films and efficient PSCs. However, few works have investigated the effects of different anti-solvents in the nucleation kinetics of perovskite and the influence on triple cation perovskite crystal growth. It is thus very important and meaningful to reveal the influence of anti-solvent on perovskite crystal growth and film formation.

Precisely controlling growth of perovskite nano-films and understanding the mechanism of grain formation are always required for fabrication of high quality perovskite materials and their long-term application in optoelectronics. There are very few reports on the function and effect of weak coordination solvent in the anti-solvent

process.[23, 32] Especially for multi-ion perovskite materials, the growth control of perovskite films are important. Weak coordination solvents could effectively adjust the balance between the reduction of grain boundaries and the modification of films based on their low boiling point, good volatility and low solubility of perovskite relative to strong polar solvents. Here the weak coordination solvents are referred as solvent with poor solubility of lead ions, weak interaction between solvent molecular and Pb^{2+} , and weak electron donating ability.. Normally, chlorobenzene (CB) is often used as the anti-solvent for fabricating perovskite films. Chlorobenzene has a poor volatility and an insoluble property of perovskite and could remain in gaps of the perovskite film and cause cracks in the anti-solvent process. It is helpful and desired to introduce weak coordination solvent in the formation process of triple cations perovskite films. On the one hand, some weak coordination solvent could remove the residual CB quickly, which reduces the number of grain boundaries and enhances the grain size of perovskite films. However, weak coordination solvent could modify the surface via its low solubility of perovskite and perovskite precursors.

In this paper, we employed some weakly polar solvents (isopropanol, acetone) in the anti-solvent process designed to control the surface morphology and explore the growth mechanism of triple cation perovskite films. Here, we found a “stitching effect” of IPA/CB mixed anti-solvent in washing processes. This effect can be used to achieve large grain sizes of perovskite films by addition a proper amount of IPA in CB. Compared to CB, isopropyl alcohol offers a higher solubility for perovskite precursor. IPA molecules play an important role in stitching cesium-containing triple cation

perovskite grains so that more homogeneous morphology and lower defect density of perovskite films were achieved via this process. The best solar cell exhibited a stabilized PCE of 19.2%. We believed the stitching effect of mixed anti-solvent washing process can be very useful for understanding the formation mechanism of perovskite and fabricating high quality large-scale perovskite films.

Results and Discussion

Cesium (Cs) has been studied in detail by Grätzel and his co-workers in the triple cation perovskite compositions.[33] $\text{Cs}_5\text{FA}_{79}\text{MA}_{16}\text{PbI}_{2.5}\text{Br}_{0.5}$ perovskite film exhibited improved stability and had the best performance. We employed this perovskite material in our study. Figure 1(a) exhibits the fabrication schematic of triple cation perovskite films prepared by an anti-solvent process. The $\text{Cs}_5\text{FA}_{79}\text{MA}_{16}\text{PbI}_{2.5}\text{Br}_{0.5}$ precursor was stirred more than 6 hours to ensure sufficient and uniform dissolution. Then this solution was spin-coated on a TiO_2 substrate, and different anti-solvents were poured on films with high speed rotation. In this process, we investigated eight different ratios between CB and IPA (1:0, 1:0.01, 1:0.02, 1:0.04, 1:0.06, 1:0.08, 1:0.1, 0:1) to control the interface morphology and grain size of perovskite films. The film was annealed at 110 °C for more than 1h. Such a long stirring time and high annealing temperature were employed to ensure the complete reaction between FAI, MABr, PbI_2 , PbBr_2 and CsI. We also used acetone as an additive solvent for CB. Although acetone is a strong solvent to halide perovskite, it has a much lower solubility of lead iodide and a much weaker coordination with lead ions than DMSO, DMF, GBL etc., and therefore acetone is also used here as one kind of weak coordination solvent for comparison. The J-V curve of

the PSC based on mixed solvent of acetone and CB washing process is showed in supplementary materials (FigureS1). The UV-visible (range from 300 nm to 1000 nm) absorption spectra of perovskite films prepared with different CB/IPA ratios were measured and shown in Figure 1(b). There are similar absorption spectra of perovskite films and based lines. However, it is also obvious that the absorption of perovskite films obtained via the mixed anti-solvent processes is slightly larger at ~760 nm than that of the perovskite films prepared using pure IPA or CB as the washing solvent. This illustrates the IPA/CB mixed anti-solvent improves the optical absorption of perovskite films. In order to study this optical absorption in more details, we intercept UV-Vis absorption spectrum from 700 nm to 800 nm in Figure1(c). Fig.1(c) shows three absorption steep slopes at 760 nm for perovskite films fabricated by the anti-solvent ratios of CB: IPA=1:0, 1:0.06, and 0:1 respectively. With 6% percent IPA addition, $\text{Cs}_5\text{FA}_{79}\text{MA}_{16}\text{PbI}_{2.5}\text{Br}_{0.5}$ film achieved the highest absorption at 760 nm. IPA (polarity= 4.3) is a polar solvent with a low boiling point and CB (polarity= 2.7) is a nonpolar solvent. It is well known that strong polar solvents such as DMF (polarity= 6.4), DMSO (polarity= 7.2) etc. are usually used to dissolve perovskites. Therefore perovskite precursors also have a very weak solubility in IPA. The addition of IPA actually resulted in a surface modification of perovskite films. Meanwhile, because of the lower boiling point, IPA also can take CB away from the surface efficiently, which enhances the perovskite grain size and crystallinity, and thereby improves the optical absorption of perovskite films. It's worth noting that the film with pure IPA washing process also exhibits an obviously absorption slope at 760 nm, which indicates the perovskite was

formed in this process. However, the lower absorption of perovskite should be owing to the different anti-solvent extraction properties between IPA and CB. Too much isopropanol could dissolve the perovskite and rearrange the films. All in all, with the ratio of CB: IPA=1:0.06, the highest optical absorption of the perovskite film was achieved. This ratio should be the best proportion for the mixed anti-solvent and “stitching effect”. The following J-V curves (Figure 1(e)) and parameter statistics of PSCs (Figure 2(a)-(d)) also confirm this conclusion.

Figure 1(d) presents the XRD spectra of perovskite films made by the anti-solvent ratios of CB: IPA=1:0, 1:0.06, 0:1. The XRD patterns with perovskite peaks at 14.16° , 19.8° , 24.4° , 28.1° , 31.55° , 34.67° , 40.17° , 42.7° , 49.68° and 51.91° , corresponding to the (111), (012), (003), (222), (123), (114), (024), (015), (006) and (135) planes, respectively, were observed in the figure. The perovskite peaks of the (111), (003), (123) and (015) are higher and sharper for the film prepared by the mixed anti-solvent process. Through calculating and fitting, the full width at half maximum (FWHM) of the (111) diffraction peak has been obtained. The FWHM of (111) peak for the sample prepared with mixed anti-solvent is 0.294 , which is lower than that (0.347) of the sample made with CB washing process. This suggests that suitable isopropyl alcohol anti-solvent additive could enhance the crystallinity of triple cation perovskite. Meanwhile, there is no XRD peak of δ -perovskite at 11.6° for the perovskite treated by IPA and mixed solvent washing, indicating that IPA is conducive to remove the δ phase of perovskite. A weak diffraction pattern of PbI_2 appeared when IPA was employed. This peak of PbI_2 should be ascribed to the different solubility of lead iodide and FA compounds in

isopropanol. The XRD measurement of perovskite prepared with pure IPA washing is consistent with this result. A much stronger diffraction peak of the PbI_2 (001) plane at 12.4° is shown in the XRD curve of perovskite based on pure IPA washing process. Although there are a lot of discussions about the function of residual PbI_2 , it is hard to tell the exact impact of such little residual PbI_2 on this mixed anti-solvent process for triple cation perovskite formation. It is very interesting to notice that the perovskite film fabricated with pure IPA washing also has much stronger diffraction peaks of the (003) and (006) planes at 24.42° and 49.68° , which shows another growth mode of perovskite films. The (003) and (006) planes are all parallel to the substrate. Isopropanol washing induced the parallel growth of perovskite and ensured a good crystallinity. Because of this, we can observe the perovskite (003) peak becomes slightly stronger and sharper when treated with the mixed anti-solvent than that prepared by the pure CB. Combined with the influence of chlorobenzene and isopropanol, the sample made with the mixed solvent washing has shown better UV-Vis absorption and film crystallinity.

The current density-voltage (J-V) curves of devices processed in different washing solvents with various mixing ratios between chlorobenzene and isopropanol are shown in Figure 1(e). The sample fabricated with the anti-solvent ratio of CB: IPA=1: 0.06 exhibits the best conversion efficiency which is 17.8%. This result is consistent with the previous UV-Vis absorption and XRD analyses. Actually, the sample via pure chlorobenzene washing process exhibits a PCE of 16.1%, not the lowest one among these devices. There is a tendency for the PCE to drop first and then rise again with increasing amount of IPA. Because of the difference (boiling point, density, solubility,

polarity etc.) between the CB and IPA, very little IPA couldn't evenly modify the films and affect CB. On the other hand, too much IPA would reduce the UV-Vis absorption of perovskite and change the growth mode of perovskite. **Therefore**, one needs to strike a delicate balance between the ratio of CB and IPA in order to form high quality perovskite films; in this study, the ratio CB: IPA=1: 0.06 was found to be the optimum proportion for preparation of high quality triple cation perovskite films. The PCE of the sample based on pure IPA washing process is exhibited in the supporting information.

Figure 2(a)-(d) show parameter (J_{SC} , V_{OC} , FF and PCE) statistics of devices fabricated using various anti-solvents with different CB/ IPA ratios. In addition to J_{SC} , it is clearly that V_{OC} , FF and PCE all exhibit the similar trend with the result illustrated in Figure 1(e). V_{OC} and FF reflect the junction properties of PSCs.[34] Higher V_{OC} and FF, and hence the enhancement of the power conversion efficiency (PCE), are due to the enhancement of the quality of perovskite films. However, as shown in Figure 2(a), the current density (J_{SC}) has been significantly improved along with adding IPA in anti-solvent. This result illustrates mixed washing solvent could reduce the recombination via “stitching” grains. Furthermore, the solar cells show good reproducibility of parameters, including J_{SC} , V_{OC} , FF and PCE.

To probe the “stitching effect” of the IPA and CB mixed washing solvent directly, we measured the surface morphology of the perovskite films by SEM and the SEM images are shown in Figure 3(a)-(c). Perovskite films fabricated by three CB: IPA ratios (1:0, 1:0.06, 0:1) were presented here. All three processes result in dense surface morphologies of perovskite films. It is particularly interesting to see that different grain

sizes were achieved through these three anti-solvents. The grain size statistics of perovskite films are also shown in Figure 3(d) (calculated by Nano Measurement software). Unlike the strong binding ability of DMSO with Pb^{2+} , IPA just has a weak coordination with triple perovskite precursors and chlorobenzene. Further analysis of washing process is as follows: isopropanol dissolved little perovskite and perovskite precursor due to their low solubility in IPA, then IPA also released more quickly than CB when films were washed by anti-solvent and annealed on the hot plate. Because of these two reasons, grain boundaries of perovskite were reduced by stitching small grains and the surface morphology of perovskite became smoother. Through this “stitching effect”, larger perovskite grains exhibit lower recombination and better junction properties as shown in Figure 3(e) and (f). From these two figures, the integrated steady photoluminescence (integrated PL) and plots of $-dV/dJ$ versus $(J_{\text{SC}} - J)^{-1}$ (including the linear fitting curves) are studied to investigate the recombination and heterojunction properties of solar cells. The samples contain compact TiO_2 , mesoporous TiO_2 and perovskite layers. From Figure 3(e), where the sample prepared with mixed solvent (CB: IPA= 1:0.06) achieved the highest integrated PL intensity. The intensity of steady PL are always associate with factors: nonradiative recombination and charge transfer at interface.[28,35-38] According to our device structure, the higher steady PL intensity means lower nonradiative recombination. Stitching effect of mixed anti-solvent results in high quality perovskite film with lower trap states. The PL spectrums are shown in Figure S8. We can see that intensities of integrated steady PL spectra also obey the trend of the V_{OC} , FF and PCE statistics. **Figure S11 shows the cross-sectional**

SEM images of the perovskite films prepared by the CB, mixed anti-solvent (CB: IPA=1:0.06) and IPA washing processes. The achieved capping layer thicknesses show marginal difference and are 401 nm (CB), 392 nm (mixed solvent) and 433 (IPA) nm, respectively. To further investigate the photovoltaic characteristics, Figure 3(f) shows the plots of $-dV/dJ$ versus $(J_{sc}-J)^{-1}$ and the linear fitting curves to calculate the ideal factor of the heterojunction. The formula is expressed as:

$$-\frac{dV}{dJ} = \frac{NK_B T}{e} (J_{sc} - J)^{-1} + R_s \quad (1)$$

Where R_s is the series resistance, J_{sc} is the short circuit current density, J is the current density flowing through the external load, N is the ideality factor of a heterojunction, K_B is the Boltzmann constant, T is the absolute temperature, e is the elementary charge, and V is the DC bias voltage that applied to the cell. It is obviously that the device made through mixed solvent (CB: IPA=1:0.06) agrees best with this heterojunction solar cell analysis. In this case, the ideality factor is 1.64. Meanwhile, the calculation ranges from 0.03 to 0.06 (range=0.03) on the x-axis. Compared with the previous report, this range is similar with that reported by Meng's group (which ranges from 0.02 to 0.05). [34] For a well-behaved heterojunction solar cell, N is always between 1.3 and 2. For the samples based on other anti-solvent process of ratio CB: IPA= 1:0, 1:0.01, 1:0.02, 1:0.04, 1:0.08, 1:0.1, N values of 2.35, 2.95, 3.12, 2.54, 1.76 and 2.11 are achieved respectively. The improvement of the junction ability **should be owing** to the larger grain size and better quality of perovskite films. **This result is consistent with our SEM, XRD and PL measurements.** Again, the successful fabrication of high quality perovskite films by using the “stitching effect” of mixed solvent (CB:

IPA=1:0.06) process is confirmed. Figure 3(g) shows the cross-sectional SEM image of a complete sample fabricated by using the mixed anti-solvent (CB: IPA=1:0.06) process. The device was fabricated with the structure as follows: fluorine doped tin oxide (FTO)/compact TiO₂/mesoporous TiO₂/triple cation perovskite /spiro-OMeTAD (HTM)/Au contact. The thickness of the perovskite layer is about 417 nm. And we can clearly observe that the perovskite capping layer has high quality and good crystallization.

In order to further study the surface morphology and roughness of triple cation perovskite films based on CB, mixed solvent(CB:IPA=1:0.06) and IPA, respectively, AFM images are shown in Figure 4(a)-(c). Tapping mode was used in this measurement. It can be seen that there is a rough surface with large undulating grains of perovskite in Figure 4(a). Pure CB washing induced an uneven surface with the root mean square roughness (RMS) of 62 nm. Furthermore, there are some unclear image area in Figure 4(a). This is because the surface is very rough while AFM measurement is usually employed to study relatively smooth materials. [39-41] When the mixed solvent was used in the fabrication of perovskite, the grains became larger and connected. The RMS decreases to 53nm compared to that based on pure CB washing process. The surface morphology of perovskite prepared in pure IPA washing is smoother than that of the first two films. It shows a RMS of 43 nm, which is the lowest in these three samples. Height distribution of perovskite films are shown in Figure 4(d)-(f), which correspond to Figure 4(a)-(c), respectively. With the increase in the proportion of isopropanol in the mixed solvent, the height distribution is even more uniform and the intermediate

threshold is lower. The intermediate height threshold decreased from 179 nm to 112 nm when using IPA instead of CB as anti-solvent. In short, AFM measurements are consistent with the previous result that triple cation perovskite with less boundaries ascribed to the modified surface morphology via the “stitching effect”.

It is very necessary and interesting to reveal the mechanism of the “stitching effect” in the mixed solvent washing procedure. A schematic film growth process is shown in Figure 5(a). During normal CB washing process, CB quickly extracts DMF and DMSO from perovskite films as an anti-solvent. Chlorobenzene has a high boiling point of 132.2 °C and a bad volatility. At the same time, perovskite and perovskite precursors are insoluble in chlorobenzene. So, after CB extracting the solvents from the perovskite film in this washing process, some chlorobenzene remains on the moist perovskite film. These residual CB won't leave until the film begin to heat at 100 °C. Then an uneven film with many boundaries and a roughness surface is formed. When IPA is added into the anti-solvent, IPA removes the residual CB quickly through its much lower boiling point and better volatility. So some boundaries are disappeared because of this “stitching effect”. Meanwhile, IPA has a relatively stronger polarity than CB since IPA can dissolve a small amount of perovskite and perovskite precursors. However, this dissolution property is much lower than DMF, DMSO etc. Therefore, this influence of IPA is marginal.

To gain further insights into this “stitching effect” of mixed anti-solvent process, we perform Fourier transform infrared spectrometer (FTIR) measurements to analyze the interaction process in this film growth procedure. Here, we studied both annealed

perovskite films and unannealed perovskite films. The FTIR measurement used transmission mode in the range of 4000-400 cm^{-1} . Three cases include CB washing, IPA washing and mixed anti-solvent washing (CB: IPA= 1: 0.06) were considered in our experiment. Figure 5(b) shows the FTIR absorption spectra of annealed perovskite films. All the samples show a low transmission and the similar shape in the long wave number range from 3000 cm^{-1} to 4000 cm^{-1} . Such high baselines suggest strong light scattering and light absorption. However, the transmission between 3500 cm^{-1} and 3200 cm^{-1} represents the N-H stretch in FA cation.[42] Because of the high baseline, the N-H peaks seem very weak. The N-H stretch is more obvious in the FTIR absorption spectra of unannealed perovskite films, as shown in Figure 5(c). For these unannealed films, there is a much stronger transmission of the perovskite based on the mixed anti-solvent process. Some weak peaks of N-H stretch are shown in curves in the range from 3500 cm^{-1} to 3200 cm^{-1} . The peak of C-OH (belongs to IPA) shown between 3700 cm^{-1} and 3500 cm^{-1} is more obvious for the mixed anti-solvent case. Actually, we are more interested in the wavenumber between 1650 cm^{-1} to 1550 cm^{-1} which represents the C=C vibration only belongs to chlorobenzene. In Figure 5(b), C=C peaks only obviously exist in the sample fabricated by CB washing process. After annealing, samples based on IPA washing and mixed anti-solvent washing don't show any C=C peaks of chlorobenzene. This result clearly demonstrates that there is a small amount of residual CB in perovskite films. But if IPA is used in this process, this residual CB could be removed. While the sample prepared with CB shows strong C=C peaks in the wavenumber between 1650 cm^{-1} to 1550 cm^{-1} , such peaks becomes weaker when 6%

IPA was added into the anti-solvent. Because all the films are not annealed, we believe that the IPA is conducive to quickly remove residual CB in this washing process. Normally, there is no C=C peak in the curve of the sample with pure IPA washing. Confirmed by this result, IPA plays an important role in stitching perovskite films via the mixed anti-solvent process.

The unannealed perovskite films exhibited dramatically different textures as seen by metallographic microscope. (Figure 6 (a)-(d)) Without any anti-solvent treatment, the composite perovskite film shows a colorful surface covered with distributed needle-like crystals indicating a very inhomogeneous nucleation kinetic. After treatment by anti-solvent process (CB, mixed anti-solvent and IPA), radial needle shaped crystals grew in the relative uniform surface having different needle density. With the addition of isopropanol, the distribution of needle crystals is more homogeneous and these radiative crystals become much denser. The differences in crystal size and crystal density further confirm the “stitching effect” of IPA in the mixed anti-solvent. The nucleation kinetics is dominating in this process. [43] This conclusion is also in agreement with the absorption of unannealed perovskite films as shown in Figure S9. The UV-Vis absorption at ~760 nm of fresh unannealed perovskite films prepared in anti-solvent processes are very high. It reveals that the formation of perovskite had been completed before heating in agreement with the FTIR analysis of the unannealed perovskite films.

We have studied the ratio between DMF and DMSO which act as the solvent of perovskite precursors to improve the performance of our device. The result is shown in

supporting materials. After optimization, we achieved the best performance device based on mixed solvent (CB: IPA=1:0.06) process which exhibits a short-circuit current density (J_{SC}) of 24.47 mA cm^{-2} , an open-circuit voltage (V_{OC}) of 1130 mV, and a FF of 0.694, resulting in a PCE of 19.2%. The J-V curve is shown in Figure 7(a). The stability of our device was also considered. Figure 7(b) shows three PCEs were measured at 0 h, 12 h and 36 h after the fabrication, respectively. The perovskite solar cell shows a good stability. UV-Vis absorption of perovskite films using different anti-solvents after three days of preparation are shown in Figure S10. Each film has a similar high absorption from perovskite at 760 nm indicating good stability of perovskite. The best PCE was achieved at 12 h after fabrication and the performance didn't obviously decay for 36 h after fabrication. Note that, the device was stored in the atmospheric environment of a clean room (in dark, 23-26°C, 40%-60% relative humidity). The EQE and hysteresis analysis are shown in supporting materials. The perovskite with mixed anti-solvent process shows a wide photoresponse from 300 nm to 800 nm and the estimated J_{SC} value was in good agreement with the J_{SC} values from the J-V measurement. Meanwhile, the J-V curve of the sample based on mixed anti-solvent process has a slight hysteresis between forward scan mode and reverse scan mode.

Conclusion

By using solvent-solvent extraction process, different amounts of IPA were added to the anti-solvent. We found that IPA can take away residual CB rapidly in the film, which reduces the density of grain boundaries of perovskite films. Meanwhile, IPA also modifies the surface morphology via its weak solubility of perovskite and perovskite

precursor. In short, we propose a new “stitching effect” based on the mixed anti-solvent process, which results in a smooth surface morphology of perovskite with large grains. Solar cells fabricated using the anti-solvent process exhibit a best efficiency of 19.2%. This optimized process shows a new way to fabricate high quality perovskite films and high performance solar cells.

Experiment Section

Materials. PbI_2 , $\text{CH}_3\text{NH}_3\text{I}$, FAI, PbBr_2 , CsI, titanium diisopropoxide bis(acetylacetonate) and spiro-OMeTAD were purchased from Sigma-Aldrich. Titanium dioxide nanoparticles (20nm particle size) and solvents, 2-Propanol (IPA, 99.5%) and DMF, DMSO, and acetone, were provided by Dyesol and YouXuan respectively.

Device Fabrication. FTO glass substrates were cleaned sequentially in alkaline detergent, acetone, ethanol, and DI water for 15 minutes each, followed by drying with N_2 flow. A compact layer of TiO_2 was deposited via spray pyrolysis at 500°C from a precursor solution of titanium diisopropoxide bis (acetylacetonate) in ethanol (volume ratio 1:39). The films were annealed at 500°C for 20 minutes and then cooled down to room temperature. The mesoporous TiO_2 scaffold was spin-coated at 4000 rpm for 30s and annealed at 500°C for 20 minutes, in which the pristine paste was diluted in ethanol (weight ratio, 1:5). $\text{Cs}_5\text{FA}_{79}\text{MA}_{16}\text{PbI}_{2.5}\text{Br}_{0.5}$ was synthesized using the one-step spin coating method and the following experiments except for gold deposition were carried out in a glove box in nitrogen. The perovskite solution contains 517 mg PbI_2 , 171 mg FAI, 23.2 mg MABr, 73.4 mg PbBr_2 in 0.75 ml DMF and 0.25 ml DMSO. $39\mu\text{l}$ CsI

solution (390 mg/ ml in DMSO) was added into this perovskite solution and stirred more than 2 hours. 80 μ l perovskite solution was spin-coated at 1000 rpm for 10s and 5000 rpm for 45s. The anti-solvent was poured on the samples at the end of 10 seconds. The samples were annealed at 110 °C for 1 h. The substrates were subsequently covered with a 72.3 mg/ml solution of spiro-OMeTAD in 1 ml chlorobenzene with 28.8 μ l of added 4-tert-butylpyridine, 17.5 μ l of a stock solution of 520 mg ml⁻¹ lithium bis(trifluoromethylsulphonyl)imide in acetonitrile and 29 μ l of 520 mg ml⁻¹ tris (2-(1H-pyrazol-1-yl)-4-tert-butylpyridine) cobalt(III) bis (trifluoromethylsulphonyl) imide in acetonitrile. This solution was spin-coated at 4000 rpm for 30s. Finally, 80nm of a gold film was deposited using thermal evaporation.

Measurement and Characterization

X-ray diffraction (XRD) diagrams were carried out using a Bede D1 system with Cu K α radiation. Scanning electron microscope (SEM) images were obtained using field emission fitting SEM (FEI-Inspect F50, Holland). The UV-vis absorption spectrum were measured by an ultraviolet-visible (UV-vis) spectrophotometer (Schimadzu UV-3101 PC). Current-voltage measurements were carried out using Keithley 2400 under AM 1.5G illuminations (1000W/m²) from a solar simulator (Newport Oriel Solar 3A Class AAA, 64023A). The AM 1.5G sun light (100 mW/cm²) was calibrated using a standard Si-solar cell (Oriel, VLSI standards). Metallographic microscope images were carried out by MZ5000 (Jiangnan Yongxin). AFM figures were measured using 300HV scanning force microscope (SEIKO). Steady PL curves were measured by F-4600 Fluorescence

Spectrophotometer (HITACHI). FTIR iS50R was used to measure our Fourier transform infrared (FTIR) spectroscopies.

Acknowledgement

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Figure1

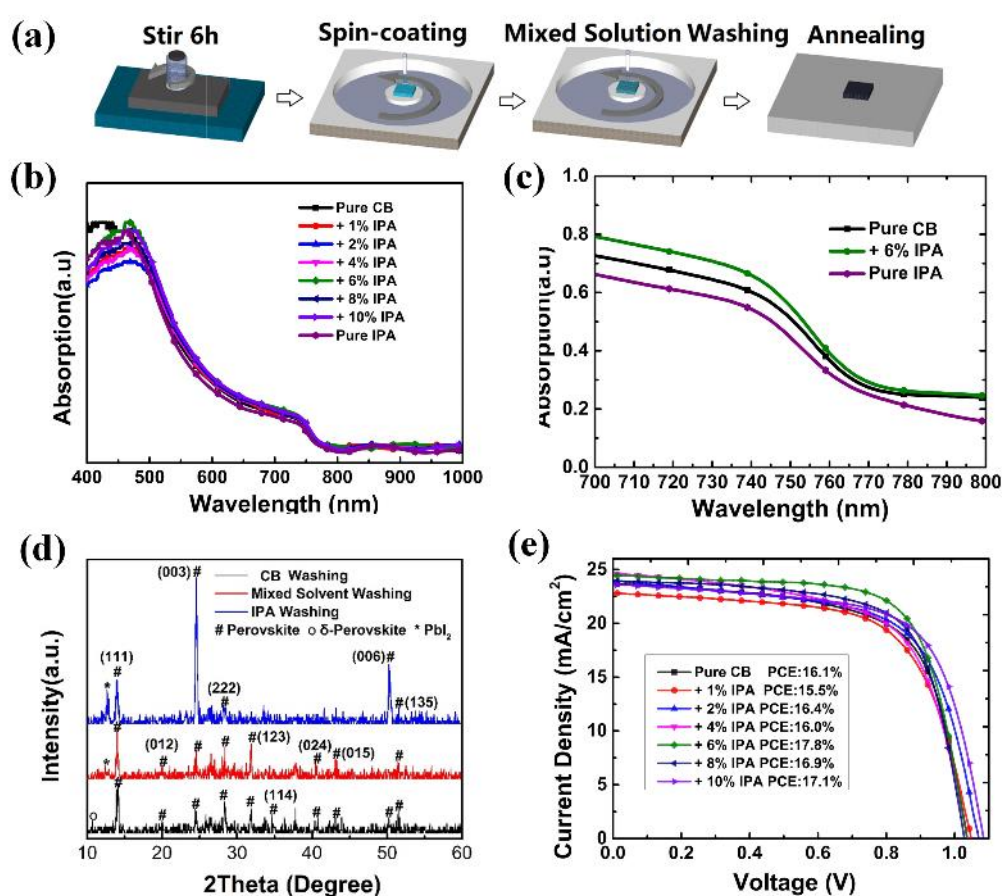


Figure1: (a) The schematic diagram of fabrication methods for PSCs; (b) UV-vis absorption spectra of perovskite using different anti-solvent (CB: IPA= 1:0, 1:0.01, 1:0.02, 1:0.04, 1:0.06, 1:0.08, 1:0.1, respectively) in fabrication (c) UV-vis absorption spectra (range from 700 nm to 800 nm) of perovskite based on pure CB, mixed solvent (CB:IPA=1:0.06) and pure IPA washing (d) XRD patterns of perovskite fabricated by

pure CB, mixed solvent (CB:IPA=1:0.06) and pure IPA washing method. (e) J-V curves of the solar cell samples with CB, mixed solvent and IPA washing processes.

Figure2

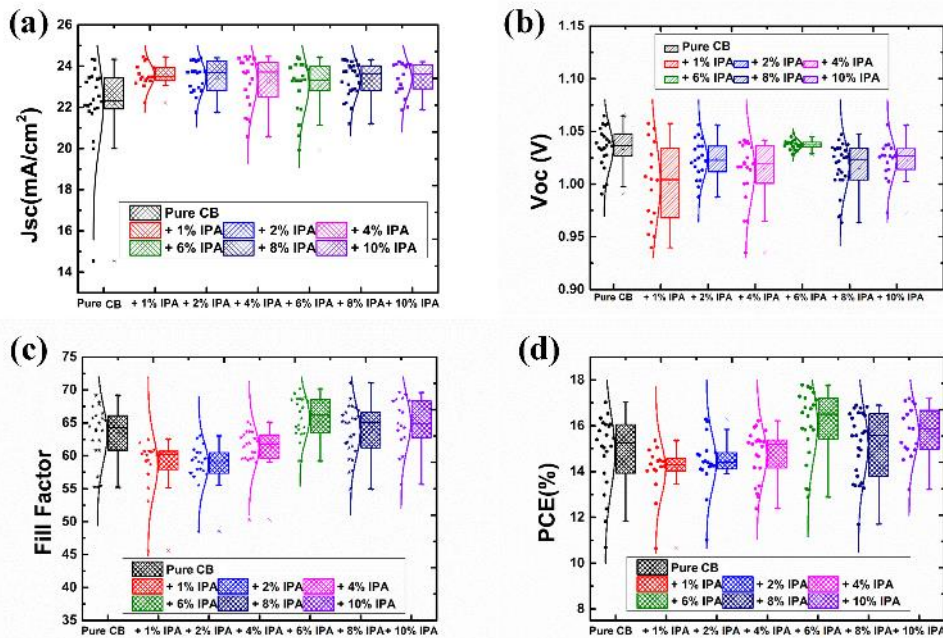


Figure.2 (a)-(d) Distribution of the J_{sc} , V_{oc} , FF and efficiency of the devices with different anti-solvent (CB: IPA= 1:0, 1:0.01, 1:0.02, 1:0.04, 1:0.06, 1:0.08, 1:0.1, respectively) process.

Figure3

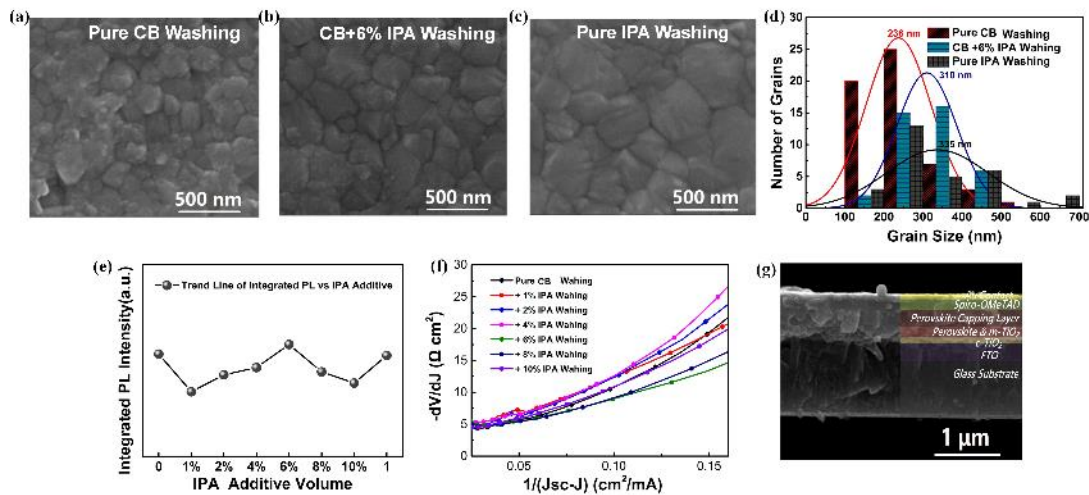


Figure3: (a)-(c) SEM top views of perovskite films treated by pure CB washing, mixed anti-solvent (CB: IPA=1:0.06) and pure IPA washing. (d) Grain size statistics of perovskite films based on CB washing, mixed anti-solvent (CB: IPA=1:0.06) and pure IPA washing. (e)-(f) Integrated Steady-State PL of perovskite and plots of $-dV/dJ$ versus $(J_{sc}-J)^{-1}$ and the linear fitting curves of PSCs based on different anti-solvent (CB: IPA= 1:0, 1:0.01, 1:0.02, 1:0.04, 1:0.06, 1:0.08 and 1:0.1) (g) SEM cross-section image of PSC based on mixed anti-solvent process.

Figure4

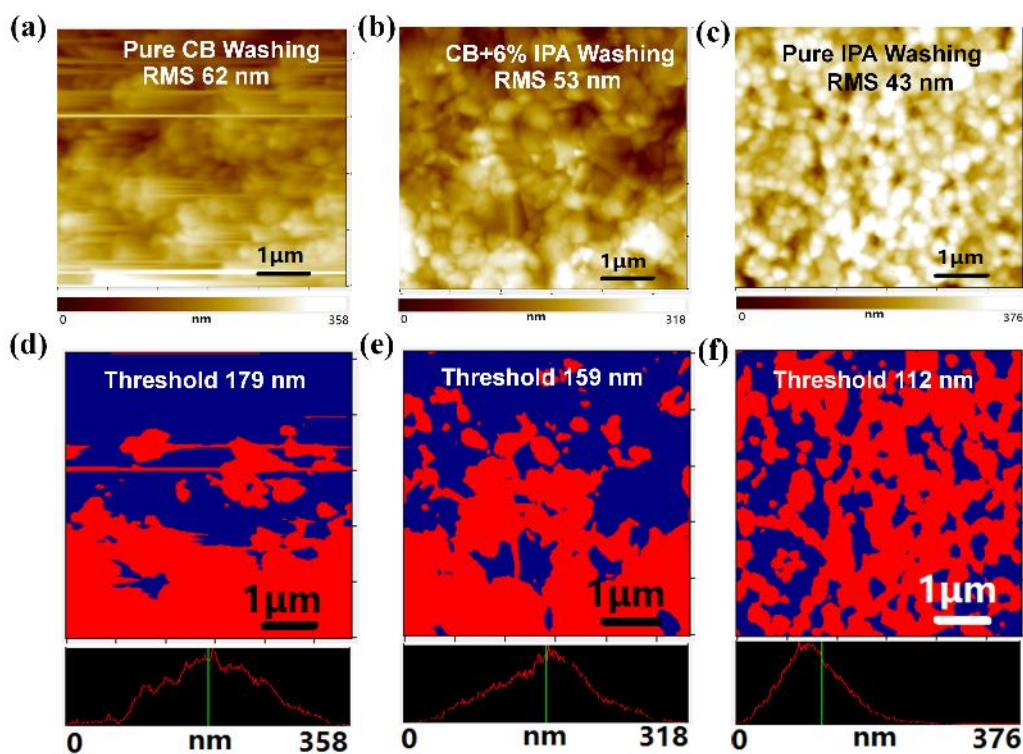


Figure4: (a)-(c) AFM top views of perovskite films and (d)-(f) the height distribution of perovskite films treated by pure CB washing, mixed anti-solvent (CB: IPA=1:0.06) and pure IPA washing.

Figure5

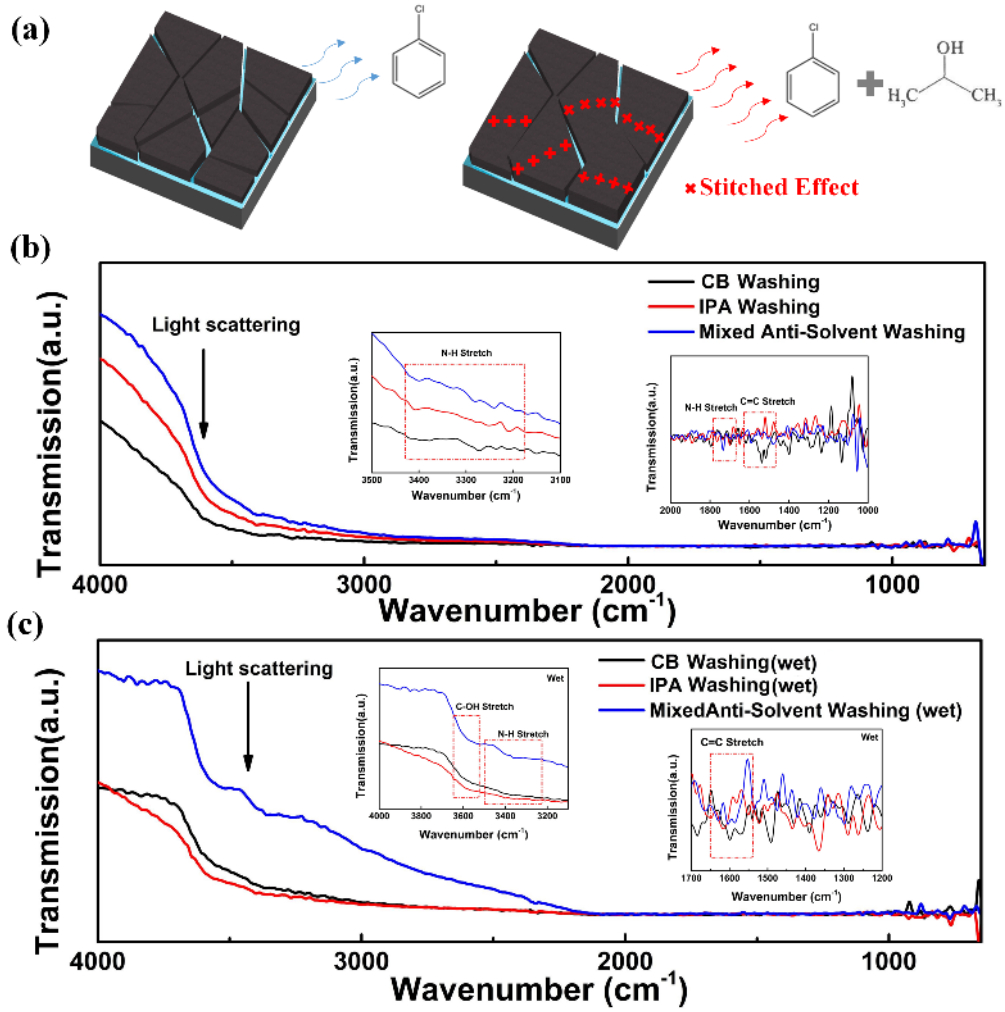


Figure5: (a) The schematic diagram of mixed-anti solvent mechanism. (b)-(c) The FTIR of annealed and unannealed perovskite films based on CB, IPA and mixed solvent (CB: IPA=1:0.06) washing processes.

Figure6

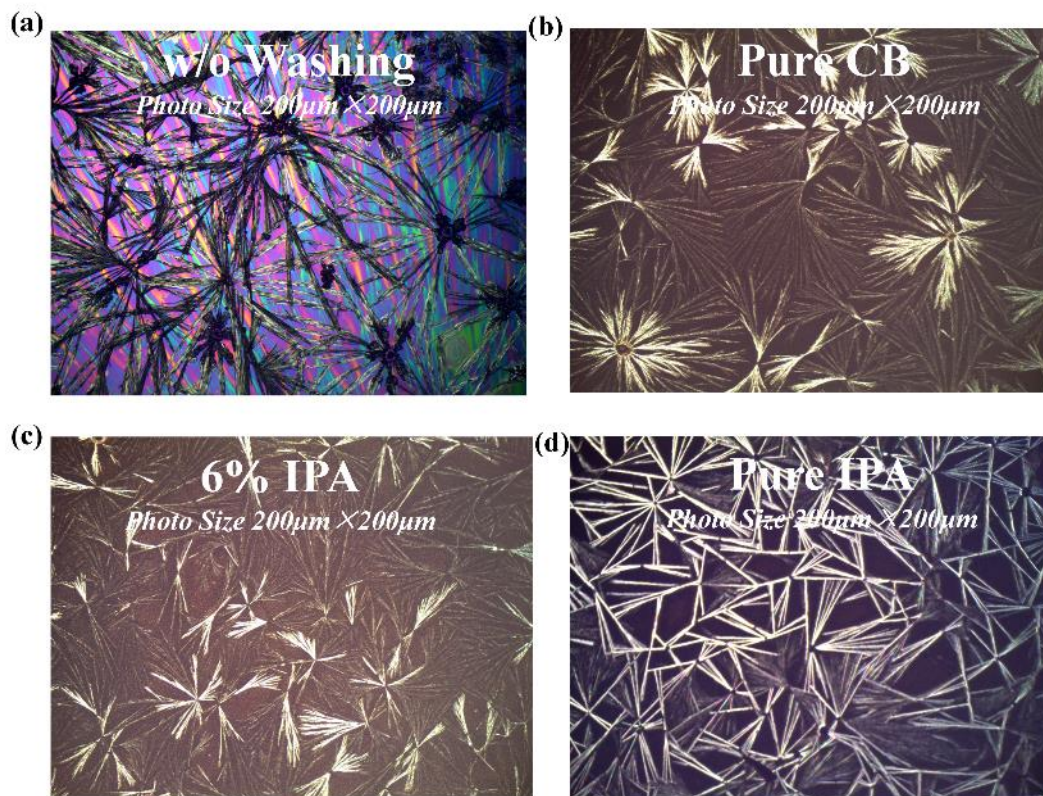


Figure6: (a)-(d) Metallographic microscope images of unannealed perovskite film that is not treated with an anti-solvent process and perovskite films using pure CB, mixed solvent (CB: IPA=1:0.06) and IPA washing processes, respectively.

Figure7

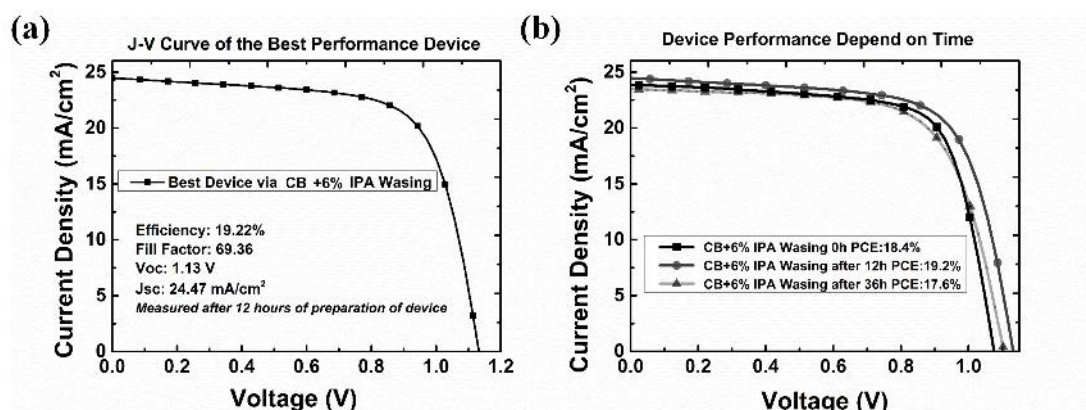


Figure7: (a) J-V curves of the best performance device obtained from mixed anti-solvent process; (b) J-V curves of the best performance device obtained from mixed

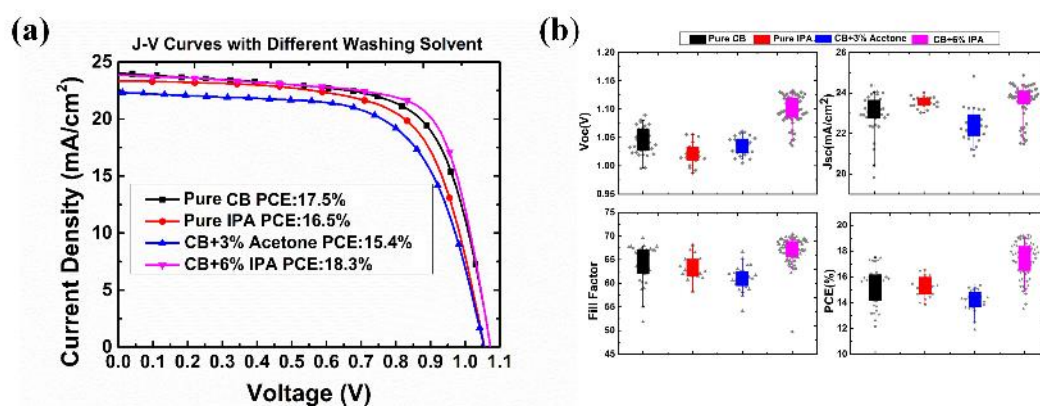
anti-solvent process measured at 0 h, 12 h and 36 h after fabrication.

Supporting Information

In our study, acetone was also used for “stitching effect”. As shown in Figure S1 (a)-(b), J-V curves and statistics of parameters of solar cells based on CB, IPA, CB+6% IPA and CB+3% acetone anti-solvent processes are exhibited. Compared with pure CB, adding 3% volume acetone couldn't improve the performance of the solar cell. Acetone could dissolve more perovskite and perovskite precursor that it may seriously rearrange the perovskite film. We also investigate the ratio between DMF and DMSO as the solvent of perovskite to improve the performance of the perovskite solar cell. DMF: DMSO=75: 25 is the best ratio for the deposition of perovskite films in our study. The EQE, integrated current density and its corresponding J-V curve are shown in Figure S3 (a)-(b). It is obvious that results of EQE and integrated current density are in good agreement with J_{sc} from J-V measurement. Figure S4 shows the AFM phase diagrams of perovskite using pure CB, mixed anti-solvent (CB+ 6%IPA) and pure IPA washing, which are consistent with the previous AFM images. Figure S5 shows photographs of perovskite films based on pure CB, mixed anti-solvent (CB+ 6%IPA) and pure IPA washing. Metallographic microscope graphs of perovskite films using different ratios between CB and IPA (CB: IPA= 1:0, 1:0.01, 1:0.02, 1:0.04, 1:0.06, 1:0.08, 1:0.1 and 0:1) in anti-solvent washing process are exhibited in Figure S7. These measurements of surface morphology demonstrate the different effects on perovskite films through different anti-solvent processes. We also study the hysteresis and find that the J-V curve

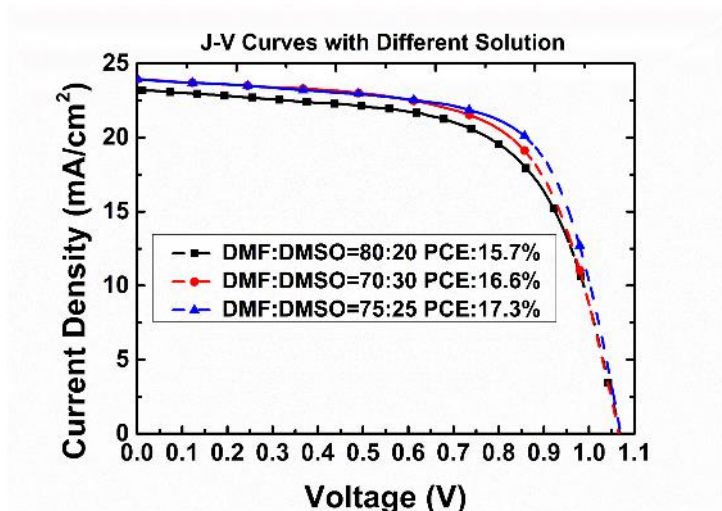
of the sample based on mixed anti-solvent process has a very slight hysteresis contrast to forward scan mode and reverse scan mode as shown in Figure S6. The steady PL intensity spectroscopy shown in Figure S8 display the trend and difference in luminescent intensities of perovskite films fabricated from different mixed anti-solvent (CB: IPA= 1:0, 1:0.01, 1:0.02, 1:0.04, 1:0.06, 1:0.08, 1:0.0 and 0:1). Figure S9 and S10 shows the UV-Vis absorption of fresh unannealed perovskite films and annealed perovskite films after loading in glove box for about three days.

FigureS1



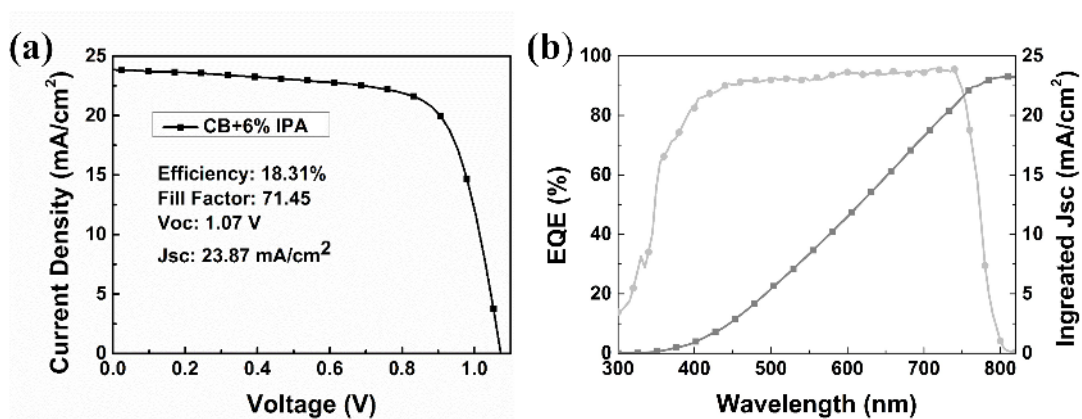
FigureS1: (a)-(b) J-V curves and statistics of J_{sc}, V_{oc}, FF and efficiency of the solar cells obtained from pure CB, pure IPA, mixed solvent of CB and 3% acetone, and mixed solvent of CB and 6% IPA anti-solvent process.

FigureS2



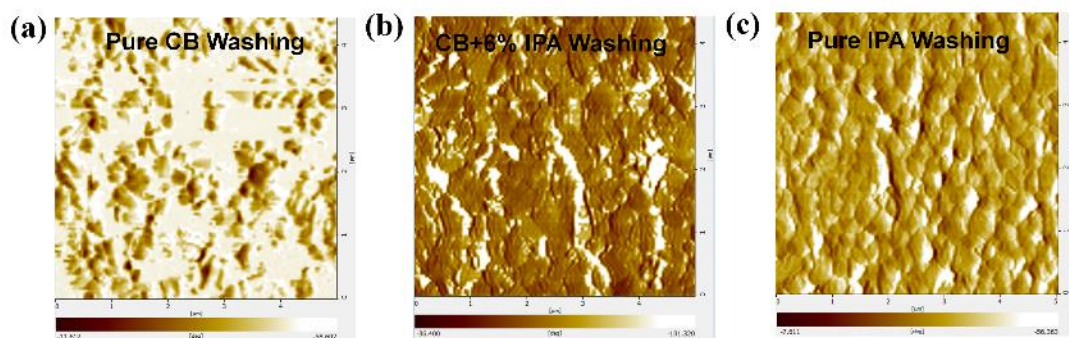
FigureS2: J-V curves of the device using different ratio between DMF and DMSO (80:20, 70:30 and 75:25, respectively) which act as the solvent of perovskite precursor.

FigureS3



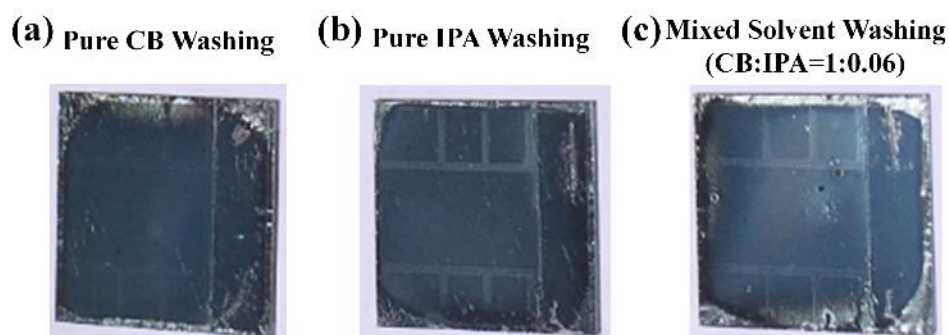
FigureS3: (a)-(b) J-V curve, EQE and integrated current density of the device using mixed anti-solvent (CB+ 6%IPA).

FigureS4



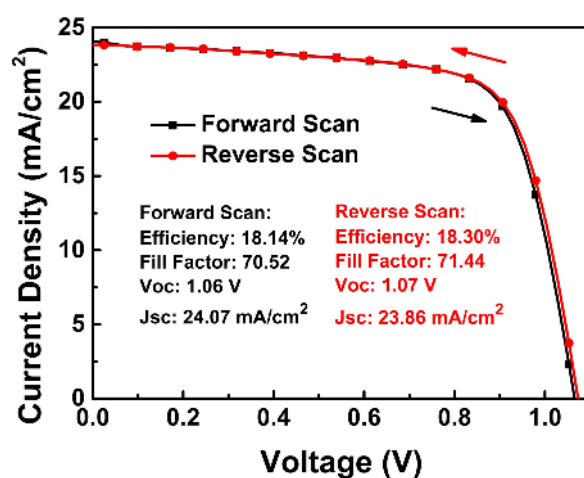
FigureS4: (a)-(c) AFM phase diagrams of perovskite using pure CB, mixed anti-solvent (CB+ 6% IPA) and pure IPA washing.

FigureS5



FigureS5: (a)-(c) Photos of perovskite films using pure CB, pure IPA and mixed anti-solvent (CB+ 6% IPA) washing.

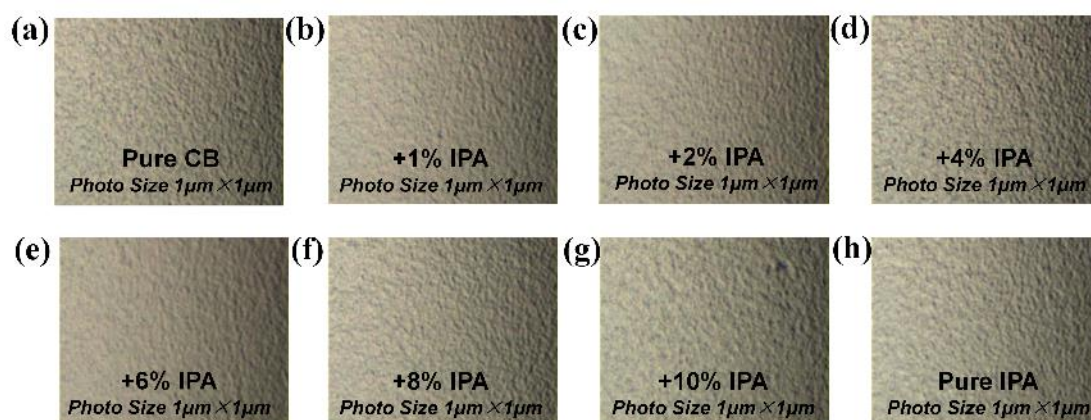
FigureS6



FigureS6: J-V curves (forward and reverse scan) of triple cations perovskite sola cell

using mixed anti-solvent (CB+ 6% IPA) washing process.

FigureS7



FigureS7: Metallographic microscope graphs of perovskite films using different ratios between CB and IPA (CB: IPA= 1:0, 1:0.01, 1:0.02, 1:0.04, 1:0.06, 1:0.08, 1:0.1 and 0:1) in anti-solvent washing process.

Figure S8

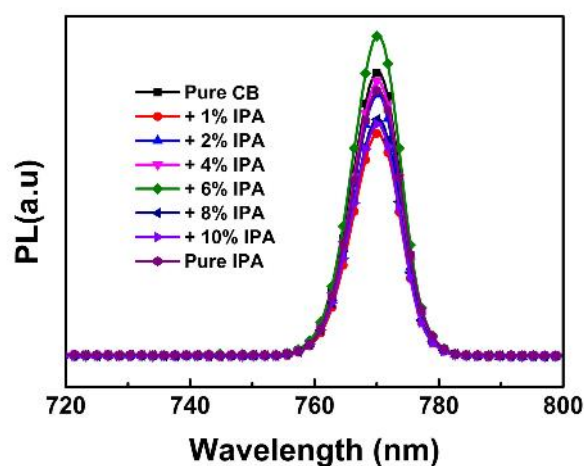


Figure S8: Steady-State PL of perovskite and plots of $-dV/dJ$ versus $(J_{sc}-J)^{-1}$ and the linear fitting curves of PSCs based on different anti-solvent (CB: IPA= 1:0, 1:0.01, 1:0.02, 1:0.04, 1:0.06, 1:0.08 and 1:0.1)

Figure S9

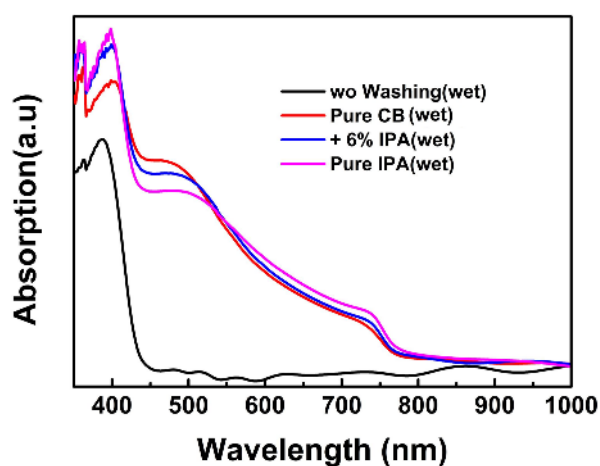


Figure S9: UV-Vis absorption of unannealed perovskite films without anti-solvent process, with CB washing, mixed solvent washing and IPA washing process, respectively.

Figure S10

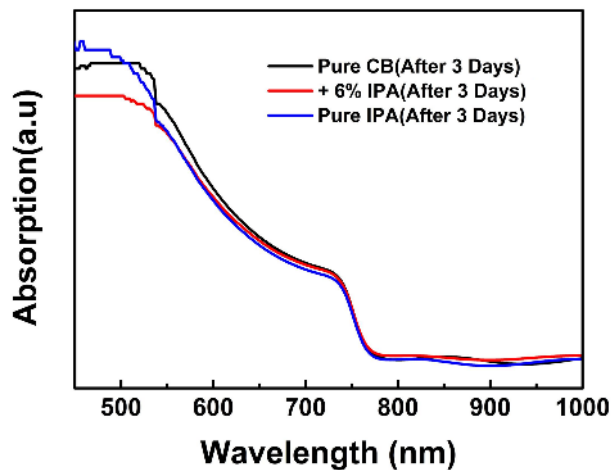


Figure S10: UV-Vis absorption of perovskite films based on CB, mixed anti-solvent and IPA washing on the third days after the preparation.

Figure S11

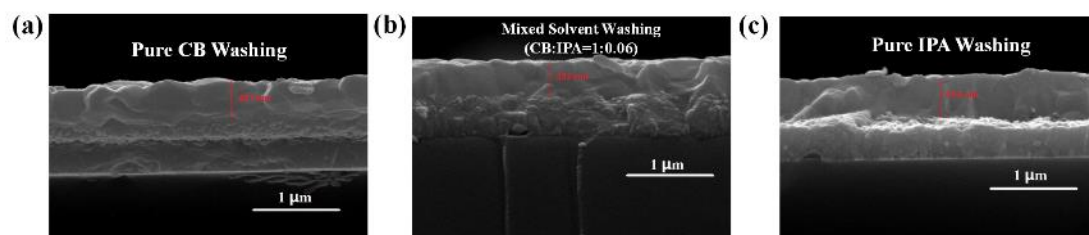


Figure S11: Cross section SEM images of perovskite films based on CB, mixed anti-solvent and IPA washing process.