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Local temperature reduction induced crystallization of MASnI₃ and achieving a direct wafer production[†]

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Despite the rapid advancement of perovskite solar cells, the fundamental drawback of toxicity in lead based materials has largely limited their large-scale development and application. Hence, we developed a local temperature reduction induced crystallization (LTRIC) method to directly obtain a lead-free perovskite wafer (CH₃NH₃Snl₃), which is 110 μ m-thick, in an oil bath. The XRD diffraction peaks of the MASnl₃ wafer are indexed to (001) and its parallel crystal planes, demonstrating its high orientation. The UV-vis-NIR discloses that the absorption onset of the MASnl₃ wafer is red-shifted to 1015 nm, corresponding to a narrow bandgap of 1.21 eV.

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Introduction

Lead halide based perovskite materials have attracted tremendous attention during the past few years because of their superior optoelectronic properties such as long carrier lifetime, high carrier mobility, long carrier diffusion length, high defect tolerance, *etc.*¹⁻¹² Even with a lot of grain boundaries existing in the microcrystalline perovskite films, there have been several kinds of high performance optoelectronic devices developed, such as solar cells,¹³⁻¹⁵ LEDs,¹⁶ lasers,¹⁷ and photodetectors.^{18,19} In particular, the efficiency of the perovskite based solar cells has been increased to as high as 22.1%, approaching that of the high efficiency of crystalline silicon based solar cells.²⁰⁻²²

To further improve performance for the optoelectronic devices, it is necessary to study defect-free single-crystalline perovskite materials.²³ Single crystalline perovskite with fewer defects is the best candidate to study its intrinsic properties. Huang developed a top seed solution growth method to produce single-crystalline CH₃NH₃PbI₃ with ultra-low trap state density ($3.6 \times 10 \text{ cm}^{-3}$) and ultra-long carrier life time (~90 µs) and diffusion length (>175 µm).⁴ Much more intriguingly, the absorption onset extended to 850 nm, around 50 nm red-shift compared to the microcrystalline films, which is better for sunlight utilization. Bakr developed an anti-solvent vapor-

assisted crystallization to prepare high quality singlecrystalline CH₃NH₃PbBr₃ and CH₃NH₃PbI₃, showing similar trap density as the Huang group reported.²⁴ However, the carrier mobility and carrier lifetime were much smaller. Furthermore, the Bakr and Liu independently developed a high temperature induced crystallization method to produce high quality singlecrystalline perovskites at high growth rate.25,26 Even though high performance photodetectors were designed and fabricated, it appears to be challenging to fabricate high efficient single-crystalline perovskite solar cells. The Bakr group used ultrasonic triggered crystallization to adjust the crystal growth process of CH₃NH₃PbBr₃ and obtained thickness controllable single-crystalline perovskite films.27 As reported in this intriguing work, the thickness plays an important role in determining the PCEs of single-crystalline perovskite solar cells. It is predicted that only <10 µm is needed to achieve a good photovoltaic performance. The PCE of single-crystalline perovskite solar cell will decrease dramatically as the film thickness exceed 10 µm. Wafer perovskite materials were needed while most of studies to date have been limited to bulk crystals which are difficult to be implemented for optoelectronic devices.

Herein, we report development of local temperature reduction induced crystallization (LTRIC) to prepare MASnI₃ wafer. The XRD results show the obtained wafer is well crystallized and shows strong orientation along [001]. MASnI₃ wafer shows an extended absorption spectrum with absorption onset reaching as long as 1015 nm, exceed 150 nm longer than 850 nm of the MAPbI₃ single crystal. Photoluminance (PL) peak is also redshifted by 160 nm to 930 nm compared to MAPbI₃. The great crystallinity and optical properties make the obtained wafer a good candidate to disclose its potential in optoelectronic devices.

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Results and discussion

Fig. 1 shows photograph of the experimental setup and a 1 cmsized MASnI₃ wafer prepared using LTRIC method. Thin MASnI₃ wafer could be found at the bottom of the precursor solution, as shown in Fig. 1a. The shape of the wafer looks like an imperfect hexagonal (Fig. 1b) that is common for general MAPbI₃/MASnI₃ perovskite crystals.^{26,28} More pictures of the MASnI₃ wafers are provided in Fig. S1.[†]

The crystallinity of the obtained MASnI₃ wafer is examined using XRD. Fig. 2a shows the XRD result taken on an as-grown wafer. It is surprising to see that essentially only four diffraction peaks indexed to (001), (002), (003) and (004) crystal planes, are observed. All of the four peaks are originated from (001) and its parallel planes, indicating that the wafer is highly orientated along [001] direction. To confirm that there is no impurity phases in the wafer, XRD measurement was taken after the wafer is grounded into fine powder. There are many more additional peaks shown in Fig. 2b, all of which can be assigned to MASnI₃, which agree well with the reported data (add a citation about the XRD results). Diffraction peaks from other impurity phases are not observed.

The thickness of wafer is $110 \ \mu m$ deduced from the crosssectional view of wafers (seen in Fig. 3a). There are some attachments on it as shown in Fig. 3b, which makes the surface rough. That should be the reason that the wafer looks dark rather than shiny as shown in Fig. 1b.

To find out the growth mechanism, a time dependent crystal growth process has been investigated and shown in Fig. 4a–d. As shown in Fig. 4b, small dark particle would firstly form on the surface of the precursor solution and would continue



Fig. 1 Photograph of MASnl₃ wafers in precursor solution (a) and the as-grown MASnl₃ wafer (b).



Fig. 2 X-ray diffraction of (a) the MASnl₃ wafer and (b) MASnl₃ powder grounded from a wafer.



Fig. 3 The cross-sectional view (a) and surface view (b) of a MASnl_3 wafer.

growing to be a large wafer (Fig. 4c). Finally, the wafer will sink into the bulk solution because of its larger gravity than the buoyancy supported by the solvent. Then, new seed crystal will form and the wafer growth process will repeat. After carefully observing the wafer growth process, we found that small water drops condensed on the cork, which should be cooler than bulk solution, would fall off the cork and dive into the solution. The local temperature of the spot into which the drop dived is reduced. Due to the normal temperature dependent solubility of MASnI₃, lowering the solution temperature will make MASnI₃ solution super saturation and induce the crystallization of MASnI₃. The cooled spot as we mentioned should induce the formation of MASnI₃ crystal seed as observed in Fig. 4b. Based on the observation and the solubility property of MASnI₃, a LTRIC process has been proposed to interpret the formation of MASnI₃ wafer, as shown in Fig. 4e-i. Due to the high solvent temperature, water would evaporate quickly to form hot vapor. As the hot vapor encountered with cold cork, it would condense to a small water drop. When the water drop became larger enough, it fell off the cold cork and dropped onto the hot solution surface (Fig. 4e). The local temperature of the spot where the drop fell onto would be reduced. And the reduction in temperature could generate a driving force for perovskite crystallization because of the normal temperature dependent solubility of MASnI₃. Then, small crystalline seed would be formed on the surface (Fig. 4g) as we observed during the wafer growth (Fig. 4b). Further increasing the reaction time, small seed will grow to a larger wafer, which would be sink into bulk solution as it is too heavy (Fig. 4h-j).

To further verify our hypothesis, anhydrous diethyl ether induced crystallization of $MASnI_3$ has been conducted as shown in Fig. S4.[†] First, two precursor solutions were prepared identically and immersed in 100 °C oil bath, as shown in Fig. S4a.[†] And the right one was used as a reference. An anhydrous diethyl ether drop was dripped into the left solution. As anhydrous diethyl ether cannot mix well with water, anhydrous diethyl ether will evaporate quickly as it contacted hot solvent surface due to its low evaporation temperature. During the evaporation, the local temperature should be reduced as evaporation process will carry energy away. Suddenly, a lot of smaller crystals were formed on the surface (Fig. S4b[†]). However, there is no any sign of crystal at the surface of reference solution (on the right). As the reaction process extended, a lot of wafers were formed on the surface of the solution, as shown in Fig. S4c.[†] At the same



Fig. 4 Time dependent wafer process growth (a-d) and schematic illustration of the proposed LTRIC process (e-i).

time, just few small crystal seeds formed at the surface of the reference solution (Fig. S4c†). That means the quick evaporation of diethyl ether anhydrous drop accelerates the crystallization of MASnI₃, faster than the reference one. On the other hand, as the water on cork droplet does not easily fall off in the reference system, slower crystal formation and fewer seed crystal observed in Fig. S4c† are reasonable. Those results could be a direct prevents of our proposed LTRIC process. Cool water drop induced crystallization of MASnI₃ has also been done. Shiny MASnI₃ wafer can also be produced with good crystallinity, high orientation as well as much smoother surface as shown in Fig. S2 and S3.† Both of those model experiments demonstrate the local temperature reduction could be an effective way to trigger crystallization and induce direct wafer production.

The optical properties of the as grown crystals were investigated using UV-vis-NIR spectrum meter. Fig. 5a shows the UVvis-NIR absorption spectra of the MASnI₃ perovskite, whose absorption onset located at 1015 nm. UV-vis-NIR absorption



Fig. 5 UV-vis-NIR absorption spectrum, bandgap and PL spectra of MASnI₃ wafer.

spectrum of MASnI₃ wafer fabricated by cool water induced crystallization is showing in Fig. S5,† which present a similar light absorption curve. The MASnI₃ shows strong absorption across the entire visible spectrum and therefore it is seen as black in color. The band gap value was estimated on the basis of the Tauc plot as the intercept value of the plot of $(\alpha h\nu)^{1/m}$ or $(F(R \propto)h\nu)^{1/m}$ against photon energy $h\nu$. As MASnI₃ is a directband gap semiconductor material, the *m* value is 1/2.²⁹ By extrapolating the linear region of the $(F(R \propto)h\nu)^2$ ³⁰ to the energyaxis $h\nu$ intercept as shown in the inset, E_{g} of the obtained MASnI₃ can be established to be 1.21 eV, similar with the former reported value.28 The obtained MASnI3 exhibits PL peak at 930 nm as shown in Fig. 5b. It is interesting to see that the PL peak value is smaller than the absorption onsets, implying that the lights of PL due to the excitation can be extinguished by themselves. There is no doubt that it is advantageous for them to be applied in solar cells.²⁶ Considering no PL emission being observed from solution-grown MASnI3 wafer material determination at room temperature, the PL appears to be much more prominent in the present wafer samples.

To examine the thermal stability of the perovskite $MASnI_3$ wafer, thermogravimetric analysis (TGA) was carried out under nitrogen flow from room temperature to 570 °C with a rate of 5 °C min⁻¹. As shown in the TGA curve (Fig. 6), the decomposition of $MASnI_3$ could be classified to two steps, corresponding to 28% weight loss between 250 °C to 323 °C, corresponding to the CH_3NH_3I evaporation. 72% weight loss between 324 °C to 538 °C. The TGA result indicates the obtained $MASnI_3$ has higher thermal stability than $MAPbI_3$ crystal, which will be thermally decomposed at 240 °C.



Fig. 6 Thermal gravimetric (TGA) data for MASnl₃ wafer.



Fig. 7 The photo, X-ray diffraction, UV-vis-NIR absorption spectrum, bandgap and PL spectra of MAPbI₃ wafer.

Furthermore, MAPbI₃ wafer has also been produced by using LTRIC method as shown in Fig. 7a. A few of MAPbI₃ are floating at the surface of the solution. The XRD result of the obtained wafer is same to our previously reported results,²⁶ demonstrating the obtained wafers are MAPbI₃. Its optical absorption and emission properties were also investigated as shown in Fig. 7c and d. The wafers show strong light absorption across the whole UV-Vis-NIR region with a sharp cut-off around 830 nm and photoemission peak at 760 nm, which are similar with the reported data.

Experimental

Materials

Methylamine (CH_3NH_2) (40 wt% in water), hydroiodic acid (HI) (57 wt% in water, stable with 1.5 wt% hypophosphorous), and other basic materials were purchased from Sinopharm

Chemical Reagent Ltd., China. PbAc₂ was purchased from Xi'an Polymer Light Technology Corp. All these materials were used as received without further purification.

Wafer preparation

MASnI₃ wafer preparation. CH_3NH_3I (MAI) was synthesized by reaction of hydroiodic acid and methylamine with molar ratio of 1 : 1.2, as reported in literature.²⁶ Fresh synthesized MAI (2.3850 g, 0.015 mol) and SnO (2.02 g, 0.015 mol) were dissolved in a mixed solvent of HI (30 mL) and H_3PO_2 (15 mL) at 80 °C. To form a transparent yellow precursor solution, the mixture was actively stirred for 8 hours. It was then transferred into a round bottom flask and submerged into an oil bath set at 100 °C. To prevent solvent evaporation, the round-bottom flask was covered with a cork. The temperature of the oil bath was slowly reduced to 98 °C in 12 h. During this process, a few small crystalline seeds were formed at the surface of the solution. As the reaction time was extended, the small seed crystals grew into larger wafers. Finally, the wafer would sink into the solution as it became bigger.

MAPbI₃ wafer preparation. Fresh synthesized MAI (1.59 g, 0.01 mol) and PbAc₂ (2.61 g, 0.01 mol) were dissolved in a mixed solvent of HI (30 mL) and H_3PO_2 (15 mL) at 70 °C. To form a transparent yellow precursor solution, the mixture was actively stirred for 8 hours. It was then transferred into a round bottom flask and submerged into an oil bath set at 70 °C. To prevent solvent evaporation, the round-bottom flask was covered with a cork. The temperature of the oil bath was slowly reduced to 60 °C in 12 h. Then a drop of diethyl ether was added onto the precursor surface to induce the crystallization. As the reaction time was extended, the small seed crystals grew into larger wafers.

Characterization

Powder X-ray diffraction (XRD) was recorded using a DX-2700 Xray diffract meter operated at 40 kV and 30 mA. UV-Vis-NIR diffuse reflectance spectrum was measured at room temperature using an UV-Vis-NIR spectrometer (PerkinElmer Lambda 950) with an integrating sphere attachment. A highly refined barium sulfate (BaSO₄) powder plate was used as the reference (100% reflectance). The obtained reflectance data were used to estimate the band gap of the wafer by using the Kubelka–Munk equation: $\alpha/S = (1 - R)^2/(2R)$, where α is the absorption coefficient, *S* the scattering coefficient, and *R* the reflectance.³¹ PL measurement was performed on a Renishaw in *via* Raman microscope with a 532 nm laser as excitation source. A field emission scanning electron microscopy (FE-SEM, SU-8020, Hitachi) was used to investigate the surface morphology, cross-section of the obtained wafer.

Conclusions

In summary, we had successfully and directly obtained 110 μ mthick MASnI₃ wafer under an ambient atmosphere by LTRIC method. After primary characterizations, it has been found that the obtained MASnI₃ wafer shows good crystallinity and great orientation as well as gives a band gap of 1.21 eV, which is narrower than their corresponding thin-film material. After carefully observation and dedicated experiment design, local spot temperature reduction induced crystallization (LTRIC) has been proposed and further verified by the cold water and quick evaporation of diethyl ether anhydrous induced crystallization experiment. At the same time, MAPbI₃ wafer has been successfully prepared by using this method. Our technique may provide an effective strategy for growing wafer for demanding high quality, low cost device applications. Further controlling the number of crystal seed and getting rid of surface attachment should be effective ways control the crystallization and quality of MASnI₃ wafer.

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