



Cite this: *J. Mater. Chem. C*, 2018, **6**, 7149

## Enhancing the hydrophobicity of perovskite solar cells using C18 capped $\text{CH}_3\text{NH}_3\text{PbI}_3$ nanocrystals†

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An important limitation in the commercialisation of perovskite solar cells is lack of stability towards moisture due to fast degradation of the absorber perovskite layer. One approach to improve the stability is effective interface engineering by adding materials that can protect the underlying perovskite film. In this work, we look at the incorporation of C18 capped  $\text{CH}_3\text{NH}_3\text{PbI}_3$  nanocrystals (MAPI NCs) in perovskite solar cells with both standard and inverted architecture. Three different solution-processing techniques were investigated and compared. We show that solar cells with MAPI NCs integrated at the perovskite–Spiro interface can reach over 10% efficiency. The presence of long chain ligands bound to the MAPI NCs does not appear to damage hole extraction. Most importantly, the hydrophobicity of the surface is significantly enhanced, leading to a much higher device stability towards moisture.

Received 23rd April 2018,  
Accepted 14th June 2018

DOI: 10.1039/c8tc01939h

rsc.li/materials-c

## Introduction

Recent years have seen remarkable developments in perovskite solar cells (PSCs) due to their outstanding optoelectronic properties and the numerous cost-efficient processing techniques explored.<sup>1</sup> The great potential of perovskites has been reflected by a very rapid growth in power conversion efficiencies (PCEs), from 10.9% in 2012<sup>2</sup> to the current world record of 22.1% in small cells and 19.7% in 1 cm<sup>2</sup> cells.<sup>3</sup> The high power conversion efficiency, the cheap fabrication and the fast development achieved over very few years make this technology one of the most promising competitors for silicon based photovoltaics. Despite the high performance, there are currently several difficulties in implementing perovskite solar cells as a viable source of energy. The three most important factors preventing halide perovskite photovoltaics from being commercialised are their ecotoxicity, their lack of stability<sup>1</sup> and the current–voltage hysteresis effect.<sup>4</sup>

Standard perovskite solar cells use the highly toxic  $\text{Pb}^{2+}$  metal cation.<sup>5–7</sup> The use of Pb raises critical environmental concerns,<sup>8</sup> which are made worse by the device instability. The crystal structure and the chemistry of lead halide perovskite based materials are highly sensitive leading to fast degradation in the presence of moisture, heat, light and oxygen.<sup>9,10</sup> Long-term stability towards moisture is currently the biggest challenge.

When exposed to moisture, a hydration process occurs, which causes current and voltage losses.<sup>11</sup> This process is irreversible in high humidity environments, leading to dissolution of the methylammonium cation.<sup>11</sup> A lot of recent research has focused on improving the stability of perovskite devices towards high humidity. Alkyl ammonium salts have been used to improve the humidity tolerance of perovskite solar cells either by functionalising the perovskite surface<sup>12</sup> or by partly substituting the alkyl ammonium cation for methylammonium iodide in the precursor solution.<sup>13</sup> Li *et al.* used 2-aminoethanethiol (2-AET) as an additive to generate a barrier layer at the surface of perovskite films; this protected the perovskite crystal structure for over 10 minutes after immersion in water.<sup>14</sup> However, the use of 2-AET in a complete solar cell was not investigated. Major progress has been achieved by using a hydrophobic insulating fluoro-silane tunneling layer to make water-resistant devices.<sup>15</sup> Very recently perovskite solar cells encapsulated with an ultrathin plasma polymer showed unchanged photovoltaic performance after immersion in water for 60 s.<sup>16</sup> Reduced graphene oxide was used as hole transport material, replacing Spiro-OMeTAD and achieving longer stability under light illumination.<sup>17</sup> Another way of enhancing the moisture stability is through 2D/3D hybrid structures<sup>18–21</sup> and cells stable for over one year have already been reported.<sup>22</sup>

Another big challenge in the halide perovskite field is the hysteresis effect. Hysteresis in perovskite solar cells is believed to be caused by both ion migration and the nature of the contact layers. Careful selection of the hole and electron transport materials (HTM and ETM) can reduce the adverse phenomenon.<sup>23,24</sup> Optimization of interfacial charge transfer by interface engineering may also limit surface recombination and benefit the power conversion efficiency.

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8tc01939h



Small molecules are often deposited at the interface to passivate the surface, reducing defects that may act as trap states.<sup>25</sup> Several strategies have also been investigated to facilitate electron transfer between the perovskite and the ETM. Recently, CuInS<sub>2</sub> quantum dots (QDs) were applied on TiO<sub>2</sub> nanorod arrays, where they were believed to facilitate electron injection from the perovskite absorber layer to the TiO<sub>2</sub> scaffold.<sup>26</sup> However, less work has been done to optimise the interface between the perovskite and the HTM. In 2016, CH<sub>3</sub>NH<sub>3</sub>PbBr<sub>0.9</sub>I<sub>2.1</sub> QDs were incorporated at the CH<sub>3</sub>NH<sub>3</sub>PbI<sub>3</sub>-HTM interface, where it was suggested that they facilitated hole transfer.<sup>27</sup> Despite the power conversion efficiency being improved by introducing QDs at the interface, no higher stability towards moisture was observed in this study. Other interface engineering strategies used to improve hole transfer at the perovskite/HTM interface were through the use of thiophene derivatives to passivate surface defects and spray-coating of MoS<sub>2</sub> flakes to achieve higher stability and increased performance.<sup>28,29</sup>

Here, we have simultaneously addressed two important issues: the optimisation of the perovskite/HTM interface and the poor water/humidity resistance of halide perovskite films. C18 capped MAPI NCs were incorporated in both the standard and inverted perovskite architecture. To our surprise the C18 ligands did not appear to negatively affect charge transfer from the perovskite film to the HTM in the standard cells. In contrast, when incorporated in inverted structure devices, inverted hysteresis was observed, suggesting that the MAPI NCs might act as an extraction barrier at the CH<sub>3</sub>NH<sub>3</sub>PbI<sub>3</sub>/ETM interface. Standard mesoporous solar cells with MAPI NCs at the CH<sub>3</sub>NH<sub>3</sub>PbI<sub>3</sub>/HTM interface reached over 10% efficiency with almost no hysteresis. The perovskite film with integrated MAPI NCs was much more hydrophobic. The C18 long chain ligands bound to the MAPI NCs allowed the cells to have a higher water-resistivity and PSCs with capped C18 MAPI NCs showed much higher stability towards moisture with respect to the reference perovskite cells.

## Experimental

### Materials

CH<sub>3</sub>NH<sub>3</sub>I (MAI, Dyesol), PbI<sub>2</sub> (PbI<sub>2</sub>, 99%, Sigma-Aldrich), PbCl<sub>2</sub> (Sigma-Aldrich), *N,N*-dimethylformamide (DMF, anhydrous, Sigma-Aldrich). The electron transport material solution consisted of 200 mm titanium diisopropoxide bis(acetylacetonate) (Sigma-Aldrich) in ethanol. The hole transport material solution consisted of 85 mg ml<sup>-1</sup> Spiro OMeTAD (Sigma-Aldrich) in chlorobenzene (anhydrous, Sigma-Aldrich), with additives of 30 μm ml<sup>-1</sup> of *t*-butyl pyridine (Sigma Aldrich) and Li-TFSI.

Solvents used for the flow synthesis of the MAPI NCs: 1-octadecene (ODE, 90%, Acros Organics), 1-butanol (BuOH, 99%, Acros Organics), oleylamine (OLA, 80–90%, Acros Organics), oleic acid (OA, 90%, Alfa Aesar) and toluene (ACS reagent, VWR International).

### Preparation of MAPI NCs

PbI<sub>2</sub> was dissolved in ODE (sol. 1) and MAI was dissolved in a mixture of ODE and BuOH (sol. 2) with OA and OLA added

as ligands. The two solutions were injected through syringe pumps and mixed through a PFA T-junction. The mixed solution was then flowed through a 3 m long PTFE tube immersed in a water bath at 30 °C. The product was collected into an ice bath and centrifuged at 7000 rpm. The solid material was resuspended in toluene and centrifuged again for 10 min. The supernatant solution, the MAPI NCs suspended in toluene, was then further used to fabricate the devices.<sup>30</sup>

### Preparation of MAPI active layer

Perovskite deposition was performed in a nitrogen filled glove box.

**Method 1.** The standard perovskite precursor solutions were prepared by dissolving MAI, PbI<sub>2</sub> and PbCl<sub>2</sub> in DMF to get 40 wt% solution (MAI:PbI<sub>2</sub>:PbCl<sub>2</sub> = 4:1:1). 100 μl of precursor solution was spun onto the substrate at 4000 rpm for 30 s, then dried at 100 °C for 100 min. The MAPI NCs dispersed in toluene was spun coated onto the above perovskite layer at 4000 rpm for 30 s followed by annealing at 100 °C for 2 min, forming the perovskite/MAPI NCs film. In the reference perovskite film, no MAPI NCs dispersed in toluene were spin coated on top.

**Method 2.** The standard perovskite precursor solutions were prepared by dissolving MAI and PbI<sub>2</sub> in DMSO/GBL (3:7 vol ratio) to get 40 wt% solution (MAI:PbI<sub>2</sub> = 1:1). 100 μl of precursor solution was spun onto the substrate at 5000 rpm for 50 s. At the 25th second 1 ml of MAPI NCs dispersed in toluene was quickly dropped onto the center of the substrate and then dried at 100 °C for 10 min. The reference perovskite film was fabricated using pure toluene as anti-solvent.

**Method 3.** The PbI<sub>2</sub> precursor solutions were prepared by dissolving PbI<sub>2</sub> in DMF to get 40 w% solution. 100 μl of PbI<sub>2</sub> solution was spun onto the substrate at 3000 rpm for 30 s, then dried at 100 °C for 30 min. In the second step, substrates were dipped in a solution containing MAI for 30 min to allow the conversion of PbI<sub>2</sub> into CH<sub>3</sub>NH<sub>3</sub>PbI<sub>3</sub>. The dipping solution consisted of a mixture of MAPI NCs dispersed in toluene and MAI in anhydrous IPA (10 mg ml<sup>-1</sup>) (1:9 vol ratio). The reference perovskite film was fabricated using a dipping solution with MAI in IPA only.

### Preparation of devices

FTO glass TEC 15 (Sigma-Aldrich) was etched with zinc powder and HCl. It was then cleaned in 2 vol% Hellmanex solution in water, followed by deionised water, acetone, propan-2-ol and ethanol.

**Standard mesoporous cells.** A compact TiO<sub>2</sub> layer was deposited by spray pyrolysis, using a hand held atomiser to spray the solution onto the substrates held at 500 °C. Substrates were then sintered at this temperature for 10 minutes. A mesoporous TiO<sub>2</sub> layer (2:7 weight mixture of Dyesol's 30NRD TiO<sub>2</sub> paste in ethanol) was spun onto the TiO<sub>2</sub> compact layer and annealed for 15 min at 150 °C and then a further 30 min at 500 °C. After cooling, substrates were placed in a UV Ozone treatment for 30 min.

A perovskite active layer was deposited onto the mesoporous TiO<sub>2</sub> layer following the three different methods outlined in the Experimental section.



Spiro solution was spun onto the perovskite film at 3000 rpm for 30 s. To establish the contacts, a 2 mm wide channel of perovskite was removed from the centre of the substrate. Finally, a 100 nm thick layer of gold was thermally evaporated using evaporation masks to make 6 pixels of  $0.0625 \text{ cm}^2$ .

**Inverted cells.** 0.2 M solution of nickel acetate tetrahydrate (Sigma-Aldrich) in 2-methoxyethanol was prepared and stirred at  $60^\circ\text{C}$  until complete dissolution.  $12 \mu\text{l ml}^{-1}$  of ethanolamine was added to the solution, which was then filtered using a  $0.45 \mu\text{m}$  PTFE filter.  $\text{NiO}_x$  was spun coated onto the UV-ozone treated FTO substrates at 4000 rpm for 30 s and annealed at  $400^\circ\text{C}$  for 30 min. 1.25 : 1.25 molar solution of  $\text{PbI}_2$  and MAI in 4 : 1 vol of DMF : DMSO was prepared. 100  $\mu\text{l}$  of precursor solution was spun onto the substrate at 4000 rpm for 30 s. 10 s after the start of the spin, 200  $\mu\text{l}$  of ethyl acetate was deposited directly onto the centre of the spinning sample. Substrates were then dried at  $100^\circ\text{C}$  for 30 min. The MAPI NCs dispersed in toluene was spun coated onto the above perovskite layer at 4000 rpm for 30 s followed by annealing at  $100^\circ\text{C}$  for 2 min, forming the perovskite/MAPI NCs film. 100  $\mu\text{l}$  of PCBM solution (20  $\text{mg ml}^{-1}$  in chlorobenzene) was spun onto the perovskite layer at 3000 rpm for 30 s. Finally, a 100 nm thick layer of silver was thermally evaporated using evaporation masks to make 6 pixels of  $0.0625 \text{ cm}^2$ . In the control inverted devices, no MAPI NCs dispersed in toluene was spun coated on top.

### Characterisation

**PXRD.** Powder diffraction patterns were collected using a Bruker Advance D8 X-ray diffractometer with a Cu K $\alpha$  source. Measurements were taken from  $2\theta$  values of  $5^\circ$  to  $80^\circ$ .

**UV/vis spectroscopy.** Thin film optical transmission and reflectance measurements were performed on a Perkin-Elmer Lambda 750S UV/vis spectrometer, from 900 nm to 300 nm.

Absorption was calculated as  $\alpha = \log\left(\frac{1-R}{T}\right)$

**J-V curves.** J-V curves were measured using Keithley 2601A potentiostat, under 1 Sun intensity and AM 1.5. The voltage was swept at  $100 \text{ mV s}^{-1}$  from 1.2 V to 0 V and back to 1.2 V. The 6 pixels were measured independently using a mask to cover the non-active area.

**SEM, TEM and AFM images.** Top-view and cross section SEM images were taken on a JEOL SEM 6480LV, at an acceleration voltage of 5 kV. Samples for TEM imaging were prepared by dropping the toluene dispersions of the NCs onto carbon supported copper grids followed by drying overnight under vacuum. The TEM measurements were performed on a JEOL JEM-2100Plus transmission electron microscope with an acceleration voltage of 200 kV. AFM images were taken on a Nanosurf easyscan 2 FlexAFM system in Phase Contrast mode using a force of 2 nN. A ContAl-G Tip was used for measurements.

**PL fluorescence.** Photoluminescence spectra were obtained on a Gilden photonics fluoroSENS fluorimeter. The dispersions of MAPI NCs in toluene were placed in a 1 cm quartz cuvette and excited at 365 nm. The integrating time was 100 ms. All the data were obtained at room temperature and in the air atmosphere.

## Results and discussion

C18 capped MAPI NCs were synthesized in toluene with OLA and OA as coligands in a flow system reactor. A TEM image of the MAPI NCs is shown in Fig. 1(a). The nanoparticles produced were of an average size of  $10.5 \pm 1.5 \text{ nm}$  with a cubic morphology. Normal distribution of the particle size was observed, showing high homogeneity in the measured data.

The band gap of the MAPI NCs was estimated from the photoluminescence (PL) spectrum and the UV-vis absorption spectroscopy, which are both shown in Fig. 1(b). The emission peak and the onset absorption coincide at 745 nm, which corresponds to 1.66 eV.

Three different fabrication methods were investigated to incorporate MAPI NCs in the standard perovskite solar cell architecture, which are shown in Fig. 2. When using method 1 (M1), MAPI NCs were deposited on top of the pre-crystallised perovskite layer and, therefore, introduced just at the perovskite/HTM interface. In contrast, method 2 (M2) used MAPI NCs dispersed in toluene as anti-solvent. It is therefore likely that the NCs partially infiltrated the perovskite layer during crystallisation. Finally, when using method 3 (M3) MAPI NCs were mixed into the MAI solution during conversion of  $\text{PbI}_2$  into  $\text{CH}_3\text{NH}_3\text{PbI}_3$  and distributed throughout the material. MAPI NCs were also incorporated in inverted perovskite solar cells with  $\text{NiO}_x$  and PCBM used as HTM and ETM respectively. These devices were fabricated using method 1, where MAPI NCs were deposited on top of the crystallised perovskite, as shown in Fig. 2(d).

MAPI NC are capped by oleic acid (OA) and oleylamine (OLA) surface ligands. It has previously been demonstrated that the  $\text{COO}^-$  functional group in OA and the lone pair on the nitrogen in OLA can coordinate with  $\text{Pb}^{2+}$  present at the surface of the MAPI NCs.<sup>31–33</sup>

### Hydrophobic properties

In this work, pure MAPI perovskite material was investigated. Perovskite devices with compositions of mixed cations and mixed halides in the active absorber layer are well known to be much more efficient and stable than pure  $\text{CH}_3\text{NH}_3\text{PbI}_3$ .<sup>3,34</sup> The purpose of this work was not to make the highest efficiency cells, but to have a reproducible baseline where any changes due to the presence of MAPI NCs could clearly be seen. Therefore, the cells were not optimised for efficiency as it was more

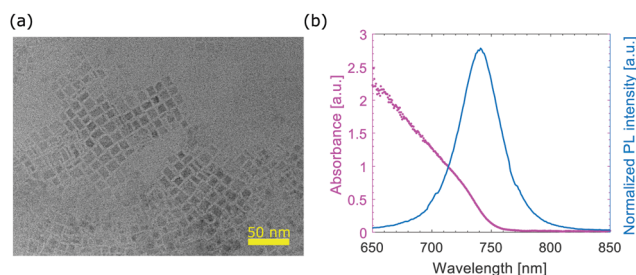


Fig. 1 (a) TEM images of C18 capped MAPI NCs; (b) absorbance and photoluminescence of MAPI NCs in toluene.





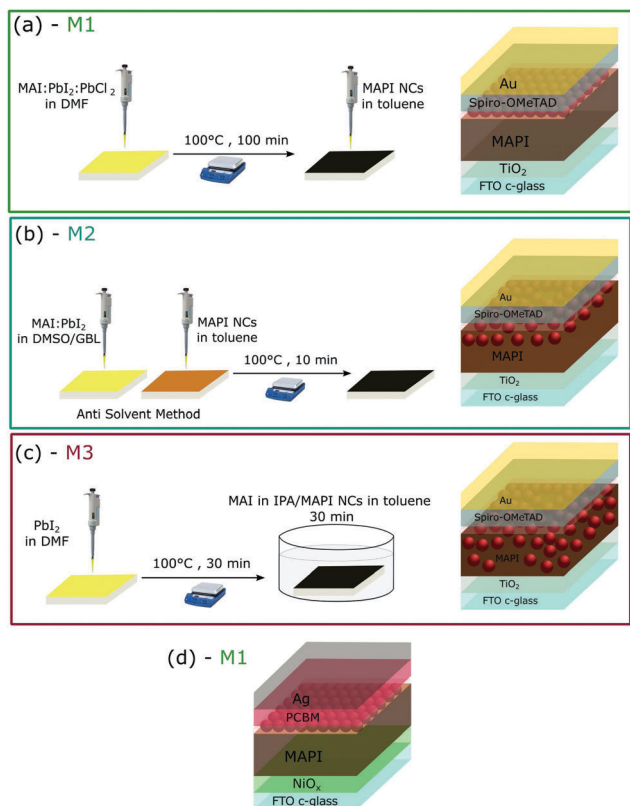


Fig. 2 Schematic illustration of the three fabrication procedures used for the processing of the perovskite solar cells with C18 capped MAPI NCs; (a) method 1 (M1) with MAPI NCs above the perovskite layer; (b) method 2 (M2) with MAPI NCs embedded in the perovskite layer using an anti-solvent deposition technique; (c) method 3 (M3) with MAPI NCs embedded in the perovskite layer using a 2-step deposition technique; (d) inverted architecture deposited using M1 with MAPI NCs above the perovskite layer at the MAPI/PCBM interface.

important to study the influence of MAPI NCs when incorporated in the standard and inverted architecture. Moreover, we were interested in seeing the effect of MAPI NCs on the stability towards moisture and on the water-resistivity of the surface. Therefore we decided to use the least stable perovskite material, which is pure CH<sub>3</sub>NH<sub>3</sub>PbI<sub>3</sub>.<sup>35</sup>

Fourier transform infrared (FTIR) spectroscopy was used to verify the presence of the C18 capped MAPI NCs in the perovskite films. All FTIR spectra, shown in Fig. S1 in the ESI,<sup>†</sup> showed N–H and C–H stretching and bending vibrations for MAPI at around 3130, 2800, 1580 and 1470 cm<sup>−1</sup>, consistent with previous reports in the literature.<sup>36</sup> When C18 capped MAPI NCs were incorporated in the perovskite film, a new peak at 2925 cm<sup>−1</sup> was observed, which was assigned to stretching of CH<sub>2</sub>,<sup>33</sup> typical of the C18 long chain ligands oleic acid and oleylamine used as surface ligands in the fabrication of MAPI NCs. The peak is particularly visible for films prepared using M1 and M2. The peak intensity is very small compared to the peaks for the bulk MAPI, but confirms the existence of C18 capped MAPI NCs in the materials. The hydrophobic properties of the perovskite films were investigated using contact angle measurements, which are shown in Fig. 3(a). We found that the

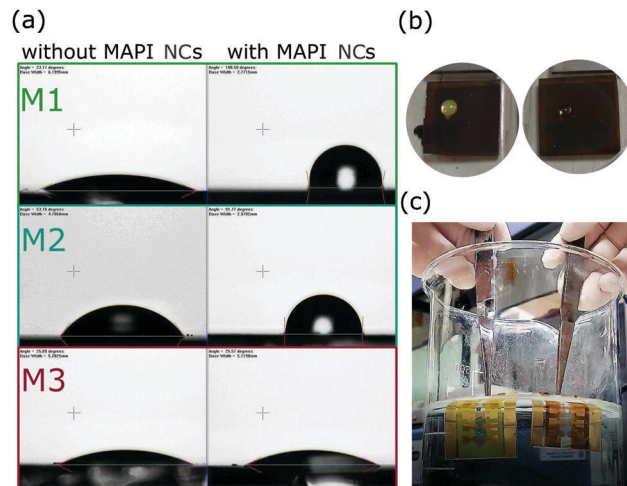


Fig. 3 (a) Contact angle measurements of perovskite films without and with MAPI NCs for perovskite films deposited with methods 1, 2 and 3; from the top to the bottom: 1 – 23° → 101°; 2 – 54° → 92°; 3 – 26° → 26°; (b) picture of perovskite films without and with MAPI NCs deposited on the surface (M1) when one droplet of water is put on the surface; (c) picture of solar cells without (left) and with (right) MAPI NCs after 30 seconds of immersion in water.

surface of perovskite films deposited using M1 and M2 were markedly more hydrophobic when the MAPI NCs were incorporated in the architecture. Contact angles higher than 100° were measured. In contrast, films deposited using M3 did not show improvements in the contact angle, indicating that MAPI NCs were not on its surface.

The presence of long chain C18 OA + OLA ligands stabilised NCs at the perovskite surface clearly enhanced hydrophobicity. In the fabrication of MAPI NCs, the initial concentration of C18 ligands is 15 vol%. The final product is centrifuged and the solid is re-dispersed in toluene in order to obtain equal volume. The exact amount of C18 ligands present in the final solution of MAPI NCs dispersed in toluene is unknown. However, it is expected to be less than 15 vol%. For comparison, the effect of depositing just the long chain ligands OA on the perovskite layer on the surface contact angle was also examined. This was done to investigate if the ligand alone would impart enhanced stability. A solution with different concentration of OA dispersed in toluene, ranging between 10 and 50 vol%, was deposited onto the perovskite film, using method 1. The contact angle, which is shown in Fig. S2 in the ESI,<sup>†</sup> increased when the long chain ligands were deposited onto the perovskite layer, reaching a maximum value of 60°. However, even when a solution of 50 vol% of OA in toluene was deposited onto the perovskite film, the surface was less hydrophobic than the one obtained by introducing MAPI NCs in the perovskite layer or on the surface (*via* M1 and M2).

Fig. 3(b) shows photographs of perovskite films deposited with method 1 without and with MAPI NCs on the surface when some water was dropped onto the surface. The perovskite films with MAPI NCs (on the right hand side) remained dark brown, while, the reference film became quickly yellow. Similar results were observed when the final solar cells were completely



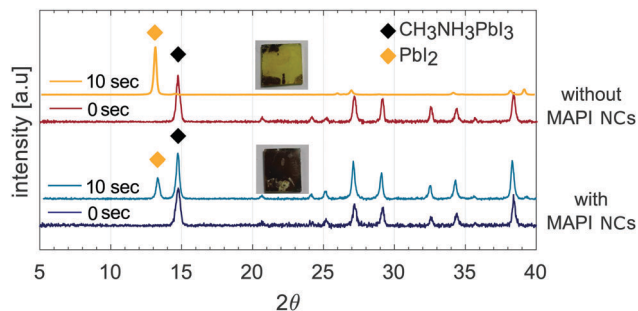


Fig. 4 XRD patterns of films with and without MAPI NCs as soon as deposited and after they have been immersed in water for 10 s.

immersed in water (Fig. 3(c)). The reference perovskite solar cell (left hand side) underwent a rapid colour change from dark brown to yellow, indicating a complete dissociation of the  $\text{CH}_3\text{NH}_3\text{PbI}_3$  phase into the  $\text{PbI}_2$  phase, while the cell with integrated MAPI NCs remained dark brown.

XRD patterns of perovskite films with and without MAPI NCs above the perovskite layer are shown in Fig. 4 (films fabricated using M1). After the reference perovskite film was immersed in water for 10 seconds, the  $\text{PbI}_2$  characteristic peak at  $13.2^\circ$  became the main diffraction peak and the  $\text{CH}_3\text{NH}_3\text{PbI}_3$  phase could no longer be detected. The amount of  $\text{PbI}_2$  also increased after the perovskite film with MAPI NCs was immersed in water for 10 seconds. However, the tetragonal  $\text{CH}_3\text{NH}_3\text{PbI}_3$  phase was still detected and its characteristic peak at  $14.7^\circ$  was identified as the main diffraction peak, indicating that the tetragonal  $\text{CH}_3\text{NH}_3\text{PbI}_3$  crystalline phase still dominated.

### Photovoltaic performance

Standard and inverted perovskite solar cells with and without MAPI NCs were fabricated using the methods described above. Fig. 5(a) shows the average performing standard mesoporous solar cells fabricated using the three different methods. To examine the reproducibility of photovoltaic performance, different samples were fabricated and measured. Table 1 lists the average values of the main photovoltaic parameters with 95% confidence intervals. The box plots showing the reproducibility of the performance are shown in Fig. S3 in the ESI.† Stabilised power output measurements were collected by monitoring the current when the cell was biased to the maximum power point voltage determined by the  $JV$  sweep and are shown in Fig. S5 (ESI†). All devices showed a stable output over 60 s. The PCEs of devices with MAPI NCs incorporated at the interface fabricated using M1 were similar to control cells. While, when methods M2 and M3 were used, the control cells showed a superior performance.  $JV$  curves showing both the forward and reverse scans are shown in Fig. S4 in the ESI.† Cells deposited with method M1 showed consistently reduced hysteresis with respect to cells deposited using M2 and M3. The solar cells deposited using the three different methods have similar  $V_{\text{OC}}$ , which is slightly larger than the  $V_{\text{OC}}$  measured in control cells. Films fabricated with the anti-solvent technique (M2) show considerably lower photocurrent density and FF, which are reflected in a decreased PCE.

The highest performance is obtained by fabricating cells using the first method M1. Cells with larger active area of  $0.25 \text{ cm}^2$  were also fabricated and shown in Fig. S8 in the ESI.† Devices deposited using method M1 showed PCE of 7.8%.

The most efficient cell with a mesoporous standard architecture and integrated MAPI NCs was fabricated using method M1 and gave  $V_{\text{OC}} = 0.74 \text{ V}$ ,  $J_{\text{SC}} = 20.89 \text{ mA cm}^{-2}$ ,  $\text{FF} = 68.52$  and  $\text{PCE} = 10.6\%$ . Cells fabricated using methods M2 and M3, where MAPI NCs are expected to be embedded in the perovskite active layer, showed consistently lower performance. Previous literature data from Cha *et al.* showed that the introduction of very thin layers of  $\text{MAPbBr}_{1-x}\text{I}_x$  QDs on top of  $\text{CH}_3\text{NH}_3\text{PbI}_3$  can change the band edge positions of the material, affecting charge extraction.<sup>27</sup> Similar to what we observe here, they found that QDs embedded in the  $\text{CH}_3\text{NH}_3\text{PbI}_3$  perovskite layer produced a PCE of only 6.93%, much lower than what measured when QDs were deposited just at the interface. They attributed the inferior performance to an increased number of recombination centers due to a misalignment of the band edge positions, the hydrophobicity of the films was not investigated.<sup>27</sup>

In the preparation of MAPI NCs, long chain ligands play a crucial role, preventing the aggregation of the perovskite nanocrystals. However, during the fabrication of quantum dot solar cells, organic ligands are usually removed because they can work as insulating barriers to charge transport between the NCs limiting the PCE of the device.<sup>37</sup> Surprisingly, the presence of C18 capped MAPI NCs did not appear to obstruct charge extraction in our standard architecture solar cells.

Photocurrent generation is directly related to the light-harvesting properties of the absorber layer. Gao *et al.* found that  $\text{CuInS}_2$  QDs integrated at the perovskite/ETM interface could assist light harvesting, increasing the absorbed light and the device performance.<sup>26</sup> To check whether the NCs had an effect on the absorption properties, UV-vis absorption of perovskite films with and without MAPI NCs was measured. The curves, which are shown in Fig. S9 of the ESI,† were found to be very similar, suggesting that the MAPI NCs did not affect light absorption in the active layer. This result is fully consistent with what was previously reported in the literature for perovskite films with  $\text{MAPbBr}_{1-x}\text{I}_x$  QDs on top.<sup>27</sup>

Recently, Yavari *et al.* observed that the incorporation of carbon nanoparticles into the perovskite active layer could boost the surface coverage of the film by enlarging grains compared to reference perovskite films.<sup>38</sup> We investigated possible changes in the morphology through top-view SEM and AFM and cross-sectional SEM images. Fig. 5(b) shows cross section SEM images of perovskite solar cells deposited using method 2 without and with C18 capped MAPI NCs. NCs can not be seen in the active layer and the film thickness does not appear to change. In Fig. 5(c) the surface of perovskite films with and without MAPI NCs deposited on top (M1) is compared through top-view SEM images. These films were deposited by using method 1 to see whether any changes due to the presence of NCs on the surface could clearly be seen. However, changes in the surface morphology are negligible. The influence of the concentration of MAPI NCs on the morphology was also



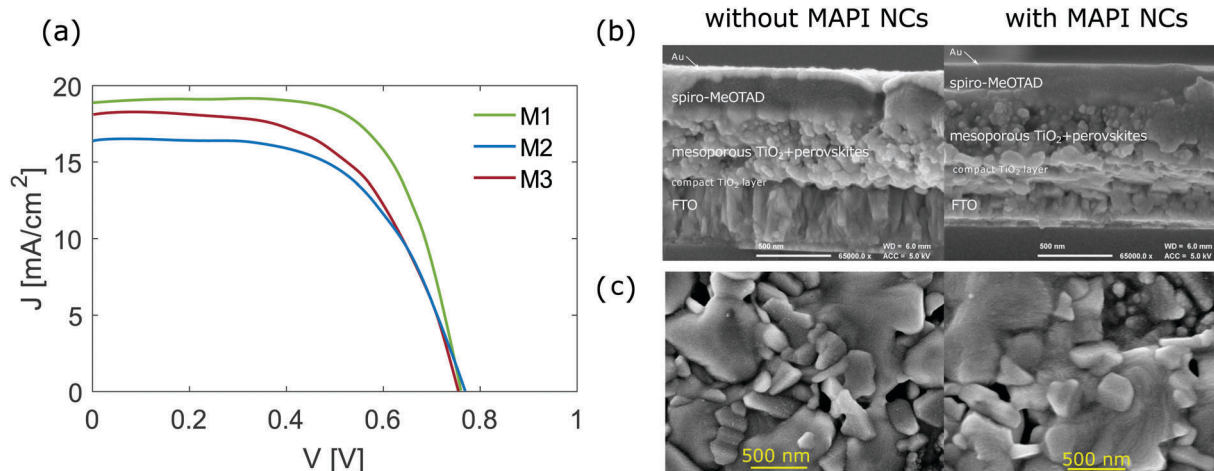


Fig. 5 (a) *JV* curves of average performing solar cells fabricated using the three different deposition method (only the reverse scan is showed, the forward scan is in Fig. S4 in the ESI†); (b) cross-section SEM images of perovskite films without and with C18 capped MAPI NCs. Perovskite solar cells fabricated using the second method M2 via the anti-solvent technique, where the NCs are embedded in the perovskite film; (c) top-view SEM images of perovskite films without and with C18 capped MAPI NCs. Perovskite films deposited using the first method M1, where the NCs are above the perovskite film.

Table 1 Average PV parameters for mesoporous cells deposited using the three different methods with 95% confidence intervals

Param.	Control	M1	M2	M3
$J_{SC}$ [ $\text{mA cm}^{-2}$ ]	$19.98 \pm 0.26$	$18.48 \pm 0.76$	$15.40 \pm 1.01$	$18.21 \pm 1.08$
$V_{OC}$ [V]	$0.73 \pm 0.01$	$0.76 \pm 0.02$	$0.76 \pm 0.01$	$0.76 \pm 0.02$
FF [%]	$68.57 \pm 4.24$	$68.22 \pm 1.37$	$61.88 \pm 2.99$	$63.54 \pm 2.70$
PCE [%]	$10.05 \pm 0.55$	$9.61 \pm 0.32$	$7.19 \pm 0.21$	$8.80 \pm 0.74$

investigated and showed in Fig. S11 in the ESI† through AFM images (films deposited using method 1).

To further investigate the effect that MAPI NCs might have at the interface, different contact layers were used. MAPI NCs were integrated in an inverted structure with  $\text{NiO}_x$  and PCBM used as HTM and ETM respectively by depositing them above the perovskite layer as described in the Experimental section. Fig. 2(d) presents a schematic of the inverted device. Fig. 6(a) shows the *JV* curves of an inverted solar cell with integrated MAPI NCs measured under forward and reverse scan. Interestingly, inverted hysteresis was observed, where the  $V_{OC}$  measured during forward scan was much larger than the one measured during the reverse scan. A similar phenomenon was observed for all the inverted cells with MAPI NCs fabricated for this study. Fig. 6(b) shows the  $V_{OC}$

measured by the reverse and forward scan for over 30 different devices fabricated in different days. In contrast, almost no hysteresis was observed in the inverted control devices without MAPI NCs at the  $\text{CH}_3\text{NH}_3\text{PbI}_3/\text{PCBM}$  interface. A typical *JV* curve and box plots of the  $V_{OC}$  under forward and reverse scan are shown in the ESI† in Fig. S13. The champion control inverted cell without MAPI NCs exhibited  $V_{OC} = 0.93$  V,  $J_{SC} = 16.74$   $\text{mA cm}^{-2}$ , FF = 69.71 and PCE = 10.89%, with very low hysteresis. While, very high inverted hysteresis was measured for the solar cells with MAPI NCs. The most efficient cell with an inverted architecture and integrated MAPI NCs exhibited  $V_{OC} = 0.85$  V,  $J_{SC} = 18.61$   $\text{mA cm}^{-2}$ , FF = 68.03 and PCE = 10.75% during reverse scan and  $V_{OC} = 0.95$  V,  $J_{SC} = 17.67$   $\text{mA cm}^{-2}$ , FF = 43.01 and PCE = 7.18% during forward scan.

Recent studies have associated the inverted hysteresis phenomenon with the presence of an extraction barrier at the perovskite/ETM interface.<sup>39,40</sup> As suggested by Cha *et al.*, very thin layers of MAPI NCs might change the surface properties of  $\text{CH}_3\text{NH}_3\text{PbI}_3$ , affecting the charge extraction.<sup>27</sup> Electrochemical experiments can be very useful in understanding the chemical changes occurring at the perovskite/HTM interface.<sup>41</sup> The anodic part of CV scans of a perovskite film and a perovskite/MAPI NCs film are shown in Fig. S14(a) in the ESI†. When MAPI NCs were deposited on the surface, the peak slightly shifted to lower voltages. The position of the first oxidation peak of the  $\text{CH}_3\text{NH}_3\text{PbI}_3$  and  $\text{CH}_3\text{NH}_3\text{PbI}_3/\text{MAPI NCs}$  films was calculated from the onset potential of the redox waves and it was found to be  $-5.32$  eV and  $-5.27$  eV respectively. Very recently, Samu *et al.* looked at the reduction and oxidation events that occur in halide perovskites and concluded that CV scans alone can not be used to determine the band edge positions of  $\text{CH}_3\text{NH}_3\text{PbI}_3$ . Other chemical processes occur in the material within the electrochemical window, therefore the measured currents can not be related to actual carrier injection.<sup>42</sup> Even though the exact band edge positions of  $\text{CH}_3\text{NH}_3\text{PbI}_3$  and MAPI NCs can not be determined from CV scans, a consistent difference between the

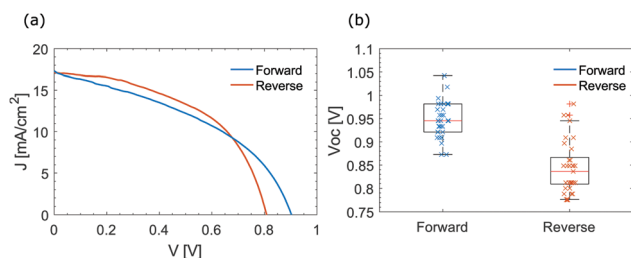


Fig. 6 (a) *JV* curves of an inverted structure perovskite solar cell with MAPI NCs incorporated at the  $\text{CH}_3\text{NH}_3\text{PbI}_3/\text{ETM}$  interface; (b) comparison of the open circuit voltages measured during forward and reverse scan for over 30 devices.





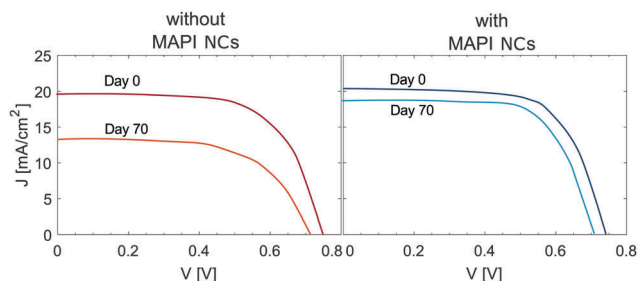


Fig. 7 Shelf-stability of the perovskite devices fabricated using method 1. *JV* curves of solar cells without and with MAPI NCs as soon as fabricated and after 70 days of ageing in ambient atmosphere (method 1 M1 was used to fabricate these cells). Without MAPI NCs – day 0: PCE = 9.57%, day 70: PCE = 5.81%; with MAPI NCs – day 0: PCE = 10.47%, day 70: PCE = 9.27%.

onset potential of the redox events was detected for the studied films, highlighting different material properties. The inverted hysteresis observed in the inverted architecture with MAPI NCs suggested that such interlayer may act as an extraction barrier at the  $\text{CH}_3\text{NH}_3\text{PbI}_3/\text{ETM}$  interface. When the same NC layer was used at the  $\text{CH}_3\text{NH}_3\text{PbI}_3/\text{HTM}$  interface no change in the shape of the *JV* curve was observed suggesting that the NCs are not detrimental at this interface.

### Moisture stability

To investigate the stability of the perovskite solar cells, the PCE of two devices with and without C18 capped MAPI NCs was measured 70 days after fabrication. During this time period, the cells were left in an ambient atmosphere without encapsulation, in the dark, at open circuit with relative humidity (RH) measured to be  $60 \pm 10\%$  (also known as shelf-stability). Two sets of cells were fabricated on the same day and exposed to the same ageing process. Two solar cells with similar initial PCEs were evaluated, the *JV* curves measured on the day of fabrication and after 70 days are shown in Fig. 7. The PCE of the reference cell decreased by over 40%. The main parameter affecting the reduced PCE was the current density. This might be due to a degradation of the perovskite  $\text{CH}_3\text{NH}_3\text{PbI}_3$  to  $\text{PbI}_2$ , due to the presence of moisture which caused a decreased light harvesting. In contrast, the cell with integrated MAPI NCs decreased by only 10%, showing what we believe to be enhanced stability towards moisture due to the hydrophobic nature of  $\text{CH}_3\text{NH}_3\text{PbI}_3/\text{MAPI NCs}$  layers.

## Conclusions

We developed standard mesoporous and inverted perovskite solar cells with integrated C18 capped MAPI NCs. Inverted hysteresis was observed in the inverted architecture, suggesting that the MAPI NCs introduced an extraction barrier at the  $\text{CH}_3\text{NH}_3\text{PbI}_3/\text{ETM}$  interface. In contrast, the hole extraction was not affected when the MAPI NCs were deposited at the  $\text{CH}_3\text{NH}_3\text{PbI}_3/\text{HTM}$  interface in standard mesoporous cells.

Three different deposition methods were investigated. MAPI NCs were deposited just at the  $\text{CH}_3\text{NH}_3\text{PbI}_3/\text{HTM}$  interface and

embedded in the perovskite active layer. When using method 1, in which MAPI NCs were above the perovskite layer, solar cells with PCEs over 10% were measured. While, when using method 2 with MAPI NCs embedded in the perovskite layer, devices with only 7% conversion efficiency were obtained.

MAPI NCs did not contribute to the light-harvesting, however they gave the perovskite films higher hydrophobicity with surface water contact angles that exceeded  $100^\circ$ . XRD measurements showed that the tetragonal  $\text{CH}_3\text{NH}_3\text{PbI}_3$  phase was retained after the film was immersed in water for 10 s. Such hydrophobic properties allowed solar cells with MAPI NCs to be much more stable towards moisture with respect to the reference perovskite solar cells.

MAPI NCs were fabricated in a flow reactor, in ambient conditions and at room temperature, therefore, no complex production steps were added to the PSC fabrication. Flow chemistry allows fast reactions, clean products and it is a technique that is very easy to scale-up, which is crucial for viable and commercial photovoltaic technologies.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

We acknowledge the MAS staff at University of Bath for their assistance in SEM acquisition. We would like to thank the EPSRC Centre for Doctoral Training in Sustainable Chemical Technologies (EP/L016354/1). This project has received funding from the European Union's Horizon 2020 research and innovation programme H2020-MSCA-CO-FUND-2014 (# 665992, MSCA FIRE: Fellows with Industrial Research Enhancement).

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