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# A-site composition tuning in methylammonium-based metal halide perovskite colloidal nanocrystals†

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Utilizing the soft-lattice nature of metal halide perovskites, we employ post-synthetic cross-ion exchange to synthesize a series of narrow band-gap colloidal nanocrystals of methylammonium-based lead iodide solid solutions of composition  $\text{FA}_x\text{MA}_{1-x}\text{PbI}_3$ , as well as those of triple-cation composition  $\text{Cs}_x\text{FA}_y\text{MA}_{1-x-y}\text{PbI}_3$  (TCPbI<sub>3</sub>). The ability to finely tune the compositions not only helps in tailoring the optical properties in the near-infrared region, but also improves the stability of these colloidal nanocrystals towards moisture, which has been demonstrated as compared to their bulk counterparts. The thermal stability of these solid solutions is also comparable to that of the bulk, as evidenced by thermogravimetric studies. This study helps in expanding the composition space of stable 3D lead halide perovskites with band gaps suitable and relevant for photovoltaic applications.

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September 2019. His research interests are in the area of colloidal inorganic semiconducting nanomaterials, their synthesis, surface chemistry and photophysics. His current research focuses are on metal halide perovskite solar cells and colloidal quantum materials for optoelectronic applications.

## Introduction

No other semiconductor offers the ease and range of composition tuning found in metal halide perovskites (MHPs). Owing to the soft and ionic nature of the lattice, all the elemental sites in the unit cell of the three-dimensional structure  $\text{ABX}_3$  (X = halide) can be easily varied even under ambient conditions through simple chemistry, which in turn helps in tuning all the important optoelectronic properties such as optical band gap, photoluminescence, carrier life-time, *etc.*<sup>1–3</sup> However, the varying sizes of different elements puts a geometric constraint on  $\text{ABX}_3$  MHPs to retain their three-dimensional structure with a network of corner-shared  $\text{BX}_6$  octahedra. This geometric criterion, known as the Goldschmidt tolerance factor (GTF), provides a fair prediction of crystallographic arrangements in  $\text{ABX}_3$  compounds. Typically, a GTF value of 0.8–1.0 is considered favourable to form a 3D perovskite structure under ambient conditions with either ideal cubic geometry or with distorted corner-shared octahedra. Beyond this limit,  $\text{ABX}_3$  compounds crystallize in either the hexagonal phase or in the orthorhombic phase with face-shared or edge-shared octahedra, respectively. Although these MHP semiconductors came into the limelight after the realization of their potential in solar photovoltaics in 2009,<sup>4</sup> such materials were investigated as early as 1958.<sup>5</sup> This particular study introduced cesium plumbohalide ( $\text{CsPbX}_3$ ) semiconductor systems and reported their crystal structure and photoconductivity. Here, all the constituents in  $\text{ABX}_3$  structures are inorganic elements, and thus they can be termed all-inorganic lead halide perovskites. In 1978, it was first demonstrated that

organic compounds with suitable size can be incorporated into the A-site in  $ABX_3$ , forming a 3D perovskite structure.<sup>6</sup> This study reported a series of methylammonium lead halides,  $CH_3NH_3PbX_3$  or  $MAPbI_3$  (MAPI), where methylammonium ions occupied the cages in the 3D network of corner-shared  $PbX_6$  octahedra, showing the possibility of hybrid metal halide perovskites. The pioneering work by Miyasaka's group in 2009 demonstrated the feasibility of  $CH_3NH_3PbI_3$  as the effective photosensitizer in a solar cell achieving a photoconversion efficiency of  $\sim 3.8\%$ .<sup>4</sup> Since then, MHP material systems have been explored extensively, showing their potential not only in photovoltaics,<sup>7,8</sup> but also in almost all areas of optoelectronics such as LEDs,<sup>9</sup> lasing,<sup>10</sup> spintronics,<sup>11</sup> photodetectors,<sup>12</sup> etc. In photovoltaics, perovskite solar cells (PSCs) have already achieved a PCE of 26.1% for lab-scale devices within an unprecedented short period of time.<sup>13</sup>

In their colloidal nanocrystalline forms, MHPs have gained interest since their synthesis *via* hot injection was reported in 2015 by Kovalenko's group.<sup>14</sup> Since then, perovskite nanocrystals (PNCs) or perovskite quantum dots (PQDs) have taken the center stage of colloidal quantum dot (CQD) research, which was earlier solely dominated by metal chalcogenide and metal pnictide-based semiconductors. Like bulk MHPs, colloidal nanocrystalline MHPs have shown tremendous promise in various optoelectronic applications.<sup>15–19</sup> PNCs offer certain advantages as compared to the corresponding thin film/bulk MHPs in terms of the range of composition tunability and solvent compatibility. Composition tuning in MHPs, in general, has been one of the major focuses, particularly in solar-cell research involving lead halide perovskites, to shift the band gap towards lower energy than that of MAPI. The introduction of a larger organic cation, formamidinium (FA,  $CH(NH_2)_2^+$ ), at the A-site of  $APbX_3$  has resulted in lowering the band gap, but its larger ionic radius makes it difficult to stabilize the band gap in the required 3D perovskite structure under ambient conditions. On the other hand, all-inorganic perovskites, such as  $CsPbI_3$ , are ideally suited to achieve better thermal stability with a narrow band gap. However, the small ionic radius of  $Cs^+$  does not result in the favourable geometric factors required for crystallization in the 3D perovskite network. PNCs offer unique advantages here: surface-related strains in these small particles cause tilting of the  $PbX_6$  octahedra, thus compensating for the geometric constraints to remaining in 3D geometry.<sup>20,21</sup> Furthermore, many other MHP semiconductors with tunable A-, B- and X-site compositions can easily be synthesized by simple ion-exchange reactions in their colloidal nanocrystalline forms, which remain stable in the 3D crystal phase at room temperature and under ambient pressure.<sup>1–3</sup>

Considering the effective radii of methylammonium, lead and iodide ions, the composition of MAPI has the near-ideal GTF ( $\sim 0.9$ ) to be in the 3D cubic perovskite phase, and probably this is the reason why organic–inorganic hybrid perovskite research was started with this particular composition. Despite reports of numerous research works on bulk and thin films,<sup>4,7,20,22</sup> not much research has been carried out on

different aspects of MAPI in colloidal nanocrystalline form. One of the key challenges is the traditional hot-injection-based high-temperature synthesis of colloidally stable MAPI nanocrystals due to the lack of availability of suitably stable methylammonium salts. There have been reports demonstrating the synthesis of MAPI PNCs at room temperature using MA-THF in oleic acid (OA),<sup>23,24</sup> an MA-benzoyl iodide mixture,<sup>25</sup> and MAI.<sup>26</sup>

However, these syntheses resulted in the formation of MAPI PNCs with morphologies other than 3D cuboidal nanocrystals during the course of the reaction. At a slightly elevated temperature of 70 °C, MAPI PNCs have been synthesized using methylamine as the major precursor.<sup>23,25,27</sup> However, due to the high volatility of methylamine, it is challenging to control the reaction. Due to the lack of enough literature reports on MAPI, A-site composition tuning with the methylammonium ion as one of the components has also not gained much attention, thus limiting the compositional parameter space in lead halide perovskite nanocrystals.

Herein, we report an in-depth study on the optical and structural properties of colloidal MAPI nanocrystals and their A-site solid solutions. The MAPI nanocrystals have been synthesized *via* the hot injection method under inert conditions using methylamine acetate. A series of compounds of chemical formula  $FA_xMA_{1-x}PbI_3$  ( $0 < x < 1$ ) have been synthesized *via* post-synthetic A-site cross-ion exchange between the MAPI and  $FAPbI_3$  (FAPI) PNCs. In the case of thin-film lead halide perovskites, it has been shown that by incorporating three cations ( $Cs^+$ ,  $FA^+$  and  $MA^+$ ) at the A-site, it is possible to overcome the thermal and compositional stability issues. This so-called triple-cation perovskite, when used as an absorber, resulted in high-performance solar cells with enhanced stability.<sup>28,29</sup> We utilized the colloidal MAPI PNCs to demonstrate that in their colloidal nanocrystalline form, such triple-cation compositions  $Cs_xFA_yMA_{1-x-y}PbI_3$  (TCPbI<sub>3</sub>) can be achieved *via* ion exchange. All the nanocrystals are stable in their 3D perovskite phase at room temperature within an ambient environment. The nanocrystal solid solutions are found to be homogeneous without the formation of any secondary phase. The MAPI nanocrystal films demonstrate better moisture stability than those of the bulk MAPI films. We have also tried to elucidate the crystal structure of the colloidal MAPI nanocrystals from powder XRD patterns, which have been found to be closely matched with the orthorhombic (*Fmmm*) space group.

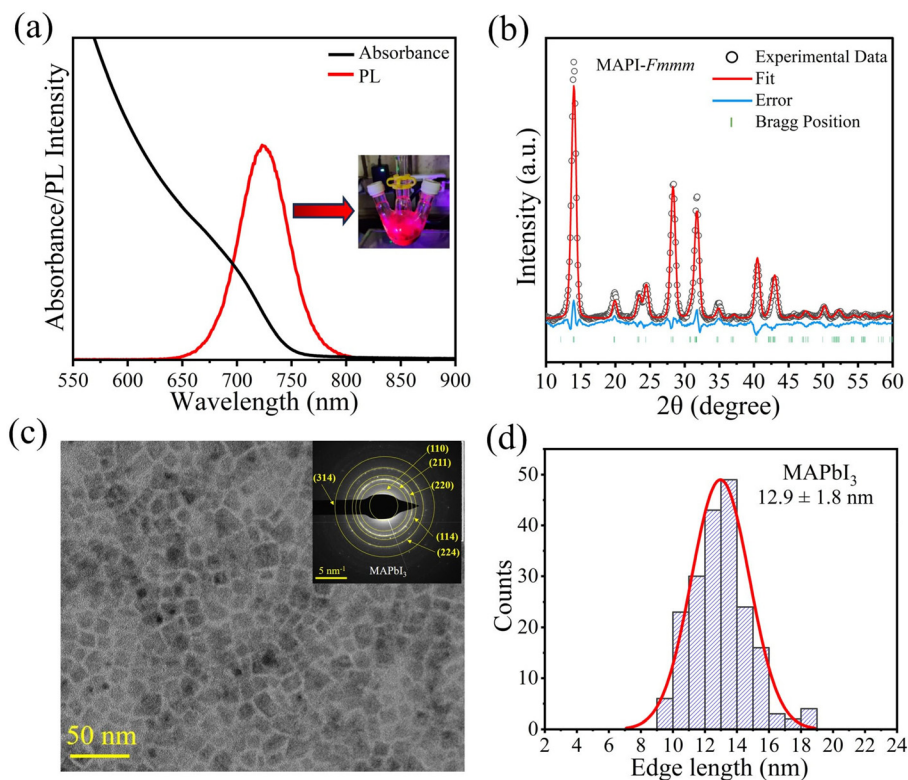
## Results and discussion

We employed methylamine acetate (MAOAc) salt as an MA precursor for the synthesis of colloidal MAPI PNCs. Unlike methylamine, which is a liquid with a boiling point of 40 °C, MAOAc is a solid at room temperature with a melting point of  $\sim 75$  °C. However, due to its hygroscopic nature, it needs to be handled under an inert atmosphere. MAOAc has been reacted with oleic acid (OA) to form MA-oleate, which has a relatively high boiling point suitable for the hot injection synthesis.

Using this MA-oleate, we followed two synthetic routes to the colloidal MAPI PNCs: in route I, MA-oleate was injected into a solution containing  $\text{PbI}_2$ , oleic acid (OA) and oleylamine (OLAm) in 1-octadecene (ODE) at a designated temperature, while route II involves three precursors where a solution of oleylammonium iodide (OLAmI) salt in toluene was injected into a mixture of Pb-oleate and MA-oleate precursors. In both these synthetic schemes, the reactions were quenched immediately after injection to control the growth of MAPI PNCs. The details of the synthesis procedure can be found in the Experimental section. The resulting PNCs exhibit good colloidal dispersion in nonpolar solvents as discussed in the following paragraphs.

The UV-Vis absorption and steady-state photoluminescence (PL) emission spectra of the MAPI PNCs, as shown in Fig. 1a, indicate a signature of quantum confinement with an absorption edge of  $\sim 750$  nm and a PL emission maximum of  $\sim 725$  nm, which are slightly blue-shifted ( $\sim 0.14$  eV) as compared to that of thin-film  $\text{MAPbI}_3$  (see Fig. S1 in the ESI† for comparison). Fig. S2† shows the powder XRD patterns of the MAPI PNCs, revealing that they crystallize in the 3D near-cubic perovskite structure where the MA cation occupies the body center of the cube with eight  $\text{PbI}_6$  octahedra occupying the eight corners of the cube. Previous studies have discussed and

debated the space group of bulk MAPI under ambient temperature and pressure conditions. While most of the literature confirms the tetragonal structure with  $I4cm$ ,<sup>30</sup>  $I4/mcm$ ,<sup>31</sup> and  $I4/m$ <sup>32</sup> space groups, a detailed crystallographic study by Karunadasa's group recently reported that MAPI at ambient pressure crystallizes in the orthorhombic  $Fm\bar{3}m$  space group.<sup>33</sup> In the colloidal nanocrystalline form, the MAPI PNCs have been indexed with a tetragonal geometry.<sup>23</sup> As shown in Fig. S2a in the ESI,† the XRD pattern matches closely with both orthorhombic ( $Fm\bar{3}m$ ) and tetragonal ( $I4cm$ ) symmetries. Moreover, closer observation at  $2\theta = 24.3^\circ$  (see Fig. S2b and S2c†) clearly reveals splitting of the cubic (1 1 1) reflection into (2 1 1) and (2 0 2) reflections despite small particle size-related peak broadening, indicating that the synthesized MAPI PNCs deviate from the highest symmetry cubic geometry. A tetragonal model ( $I4cm$ ) was tested but this resulted in a poor fit (see Fig. S3 in the ESI†). While it accounts for the splitting, it fails to reproduce the intensity accurately. In contrast, an orthorhombic model ( $Fm\bar{3}m$ ) provided a reasonable fit, as shown in Fig. 1b. Based on these observations, we propose that the structure of the MAPI PNCs is likely to adopt an orthorhombic symmetry. We note here that it is not straightforward to determine the crystallographic structure accurately for nanomaterials with small-range atomic ordering using traditional



**Fig. 1** (a) UV-Vis absorption and PL emission spectra of the MAPI PNCs; the inset shows the  $\text{MAPbI}_3$  reaction mixture under UV light illumination. (b) Rietveld refinement of the powder XRD pattern of the MAPI PNCs with reported crystallographic information for  $Fm\bar{3}m$  symmetry. (c) TEM image of the MAPI PNCs; the inset shows the SAED patterns of the MAPI PNCs; the planes were assigned according to the  $d$ -spacing value calculated from the powder XRD pattern. (d) Size distribution histogram calculated from the TEM image, indicating that the MAPI PNCs have an average edge length of  $12.9 \pm 1.8$  nm.

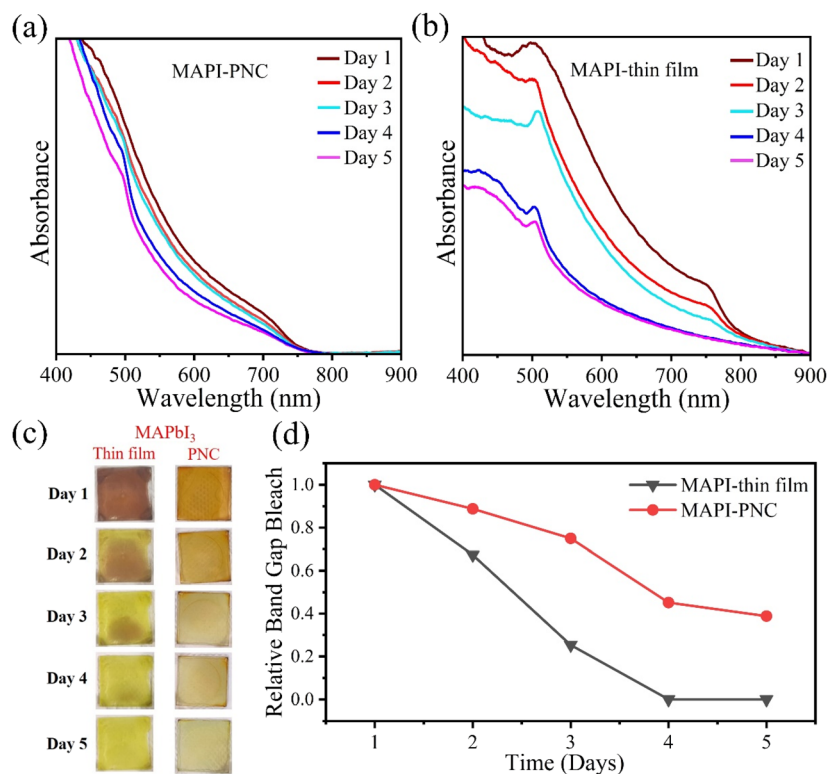
crystallographic methods due to inherent problems such as defects, size distribution and disorder.<sup>34–36</sup> Careful determination of the atomic structure for the PNCs requires total scattering approaches using synchrotron-based sources and neutron scattering, as well as local-to-average structure modeling based on a variety of other complementary techniques, such as atomically resolved high-resolution transmission emission microscopy (HRTEM) and extended X-ray absorption fine structure (EXAFS). Such approaches have been adopted for different nanomaterials for structure determination in different reports.<sup>37,38</sup>

The transmission electron microscopy (TEM) image shows the cuboidal shape of the MAPI PNCs (Fig. 1c) with an average edge length of  $12.9 \pm 1.8$  nm, while the SAED pattern reveals the crystalline nature of the MAPI NCs (Fig. 1d and the inset of Fig. 1c, respectively). The MAPI PNCs synthesized *via* route II exhibit similar optical and structural properties (see Fig. S4 in the ESI†).

One of the major issues with lead halide perovskite semiconductors, in general, has been their poor stability towards air and moisture. Several mechanisms have been proposed to explain the degradation pathways of hybrid perovskites. Unlike in all-inorganic perovskites, such as CsPbI<sub>3</sub>, where degradation takes place *via* crystallographic phase change to the optically inactive  $\delta$ -phase,<sup>20,39</sup> hybrid perovskites undergo compo-

sitional change during interaction with air and moisture. For instance, methylammonium lead iodide, when it comes into contact with water, decomposes into lead iodide and methylamine/hydrogen iodide/ammonia.<sup>40–47</sup> We checked the stability of the MAPI PNC solid films under 50–70% relative humidity (RH), normal light and ambient conditions and compared their performance to that of MAPI polycrystalline (PC) thin films. As can be seen from Fig. 2c, the MAPI thin films start degrading from day 1 itself (appearance of yellow coloration). In contrast, the MAPI PNC films appear to become lighter with time without yellow coloration. This degradation can be better visualized by the extent of band gap bleaching in the corresponding absorption spectra, as shown in Fig. 2a and b; the relative absorbance at the exciton energy decreases faster in the MAPI thin film compared with that of the PNC film (Fig. 2d). The resistance of the MAPI PNC films towards moisture degradation can be attributed to the presence of hydrophobic surface ligands.

Post-synthetic ion exchange in colloidal PNCs has been shown to be an effective way to achieve compositional tunability that is beyond the reach of that in thin-film or bulk perovskites.<sup>1–3,48</sup> Such control, in turn, has helped in tuning optoelectronic and photovoltaic properties.<sup>48–50</sup> While X-site ion exchange is known to occur instantaneously at room temperature, for A-site ion-exchange reactions in PNCs, elevated



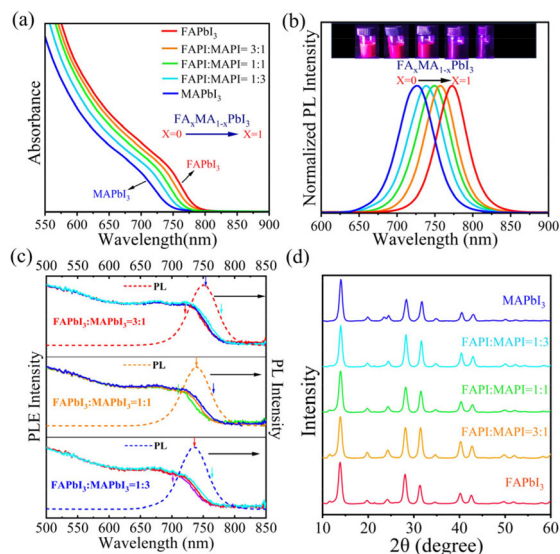
**Fig. 2** UV-vis absorption spectra of (a) the MAPI PNC film and (b) the MAPI thin film recorded from day 1 to day 5 after preparation in 50–70% relative humidity (RH) under normal light and ambient conditions. (c) Photographs of both the MAPI PNC and thin films showing how the colour of the film changes with time. (d) Comparison of the change of relative absorption of both films with time; the PNC film shows better phase stability compared with the thin film.

temperature/longer time/excess amounts of ligands are required.<sup>48,50</sup>

Following a similar A-site cross-exchange reaction between CsPbI<sub>3</sub> and FAPbI<sub>3</sub>,<sup>48,50,51</sup> we synthesized a series of PNCs with composition FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> by mixing the MAPI PNCs with the colloidal FAPI PNCs at different ratios. As shown in Fig. 3a and b, controlled variation of “x” resulted in tuning optical absorption and photoluminescence (PL) in the range of ~700 nm–800 nm, with larger values of “x” resulting in a red-shift in absorption and PL emission. We plotted the bandgap energy as well as the PL emission maximum energy of these solid solutions as a function of composition “x”, and both of these properties exhibited a linear relationship (Fig. S5†). Such a linear relationship of band gaps with composition “x” measured from diffuse reflectance has been reported in the case of bulk FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> solid solutions as well.<sup>52</sup> Apart from the steady-state PL emission, the time-resolved PL (TRPL) decay kinetics also exhibits monotonic variation with the introduction of more MA ions, resulting in faster decay, as can be seen from Fig. S6 in the ESI.† The absence of bimodal absorption in the UV-Vis absorption spectra and the narrow symmetric PL emission peaks indicate the formation of nearly homogeneous solid solutions all throughout the FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> composition range. Photoluminescence excitation (PLE) spectroscopy is a commonly used technique to study inhomogeneous broadening of PL emission in colloidal quantum dots, which primarily occurs due to size distribution in these quantum-confined particles, which means that this technique indirectly provides information on sample inhomogeneity.<sup>53–55</sup> In the weakly quantum-confined hybrid lead halide perovskite NC solid solutions, in the size range of 10 nm–15 nm, slight compositional variation is expected to result in inhomogeneous broadening of PL emission. We have collected PLE spectra of a few of the FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> compositions (Fig. 3c), and it has been observed that the PLE spectra at different emission energies overlap with each other in all the compositions, indicating negligible compositional inhomogeneity.

Powder XRD patterns (Fig. 3d) of the FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> PNCs indicate that all compositions retain the 3D perovskite phase even after the ‘A’-site cross-ion exchange reaction between MAPI and FAPI PNCs. Detailed crystallographic analyses of these solid solutions were carried out by performing Rietveld refinement. The results from the refinement are shown in Fig. S7 and Table S4.† While the FAPI PNCs crystallize in the cubic *Pm3m* space group, as also reported in the previous literature,<sup>56,57</sup> it has been found that introduction of even a small amount of FAPI results in transformation of the orthorhombic phase of pure MAPI into the cubic phase, and this cubic phase is retained in all the FA-containing compositions (*x* = 0.25, 0.5 and 0.75). Similar phase transformation has also been observed in the case of bulk FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub>, although from the tetragonal phase to the cubic phase.<sup>52,58,59</sup> However, in the case of the bulk, a higher FA content results in degradation of FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> from the cubic phase to the hexagonal phase with time.<sup>52</sup> In the case of our FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> PNCs, the stability of the cubic phase remains stable for months. From the calculated crystallographic information as shown in Table S1,† we observed that with an increase in FA content, the lattice parameters and unit cell volumes increase linearly (Fig. S8 in the ESI†). Such variations can also be found in the case of bulk FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> solid solutions.<sup>52,58</sup> Linear variation of the optical bandgap, PL emission energy maxima and lattice parameters as functions of “x”, all point towards the formation of homogeneous solids of FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> PNCs in the range of 0 ≤ *x* ≤ 1. The TEM images show the formation of cuboidal-shaped particles for all the FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> PNCs with an average edge length in the range of 13.2 ± 1.7–14.18 ± 1.9 nm, comparable to that of their parent MAPI (average edge length of 12.9 ± 1.8 nm) and FAPI (average edge length of 14.6 ± 2.3 nm, Fig. S9†) PNCs (see Fig. S10a–f† for complete TEM analysis of the FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> PNC alloys). Combining optical spectroscopic and structural data discussed above confirms the formation of homogeneous solid solutions of FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> PNCs in cuboidal 3D perovskite phases across all compositions.

We further extended the ‘A’-site composition tuning to synthesize phase-stable hybrid triple-cation perovskite NCs by introducing the ‘Cs’ cation. For hybrid lead halide perovskite thin films, it has been reported that incorporation of both Br<sup>-</sup> and I<sup>-</sup> at the X-site is essential to simultaneously accommodate Cs, MA, and FA at the A-site, resulting in the formation of

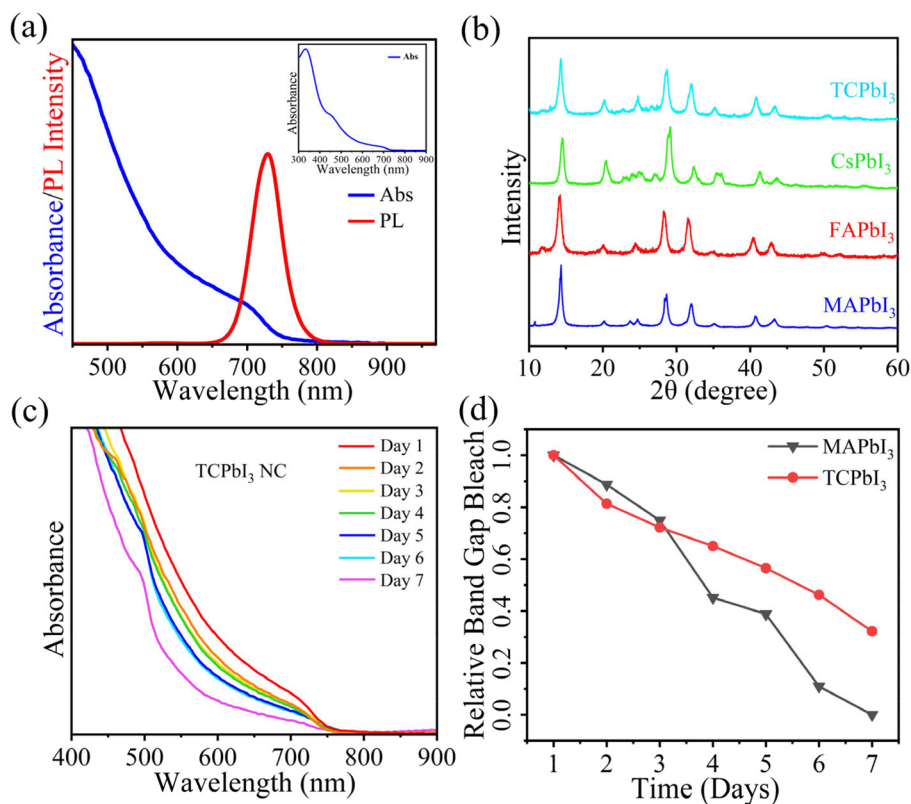


**Fig. 3** (a) UV-Vis absorption and (b) steady-state PL emission spectra of the FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> compositions showing tunability over the range of 720 nm–770 nm; the inset in figure b shows the digital photograph of the FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> alloys under UV illumination. (c) Photoluminescence excitation (PLE) spectra of the FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> NCs at different emission energies as indicated by the colour-coded arrow in the emission spectra, showing the absence of substantial inhomogeneous broadening of the PL emission due to alloying; the dashed line represents the PL spectra of the corresponding alloy. (d) Powder XRD patterns of the synthesized alloy compositions of formula FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> NCs along with their parent NCs; all of these diffraction patterns indicate the formation of the 3D perovskite phase.

3D triple-cation (TC) perovskites.<sup>29,60–62</sup> Although triple-cation perovskites provide structural stability and desirable band gap in a thin film, it is well known that the presence of both Br<sup>-</sup> and I<sup>-</sup> leads to halide phase segregation under constant light illumination.<sup>63–66</sup> In the colloidal nanocrystalline form, only a few reports are available for triple-cation PNCs. For instance, the ligand-assisted reprecipitation (LARP) approach to synthesize spherical-shaped CsFAMAPbBr<sub>3</sub> triple-cation PNCs has been reported.<sup>67</sup> Very recently, Manna's group demonstrated the formation of colloidal CsFAMAPbBr<sub>3</sub> triple-cation perovskite NCs *via* post-synthetic 'A'-site ion exchange reactions.<sup>51</sup> Following the synthesis of FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub>, as discussed above, we applied A-site cation cross-exchange to form I-only triple-cation PNCs of chemical composition Cs<sub>x</sub>FA<sub>y</sub>MA<sub>1-x-y</sub>PbI<sub>3</sub>, abbreviated as TCPbI<sub>3</sub>, which may act as a better alternative to Cs<sub>x</sub>FA<sub>y</sub>MA<sub>1-x-y</sub>Pb(Br<sub>z</sub>I<sub>1-z</sub>)<sub>3</sub>-based triple-cation perovskite thin films. 3D perovskite phase-stable triple-cation TCPbI<sub>3</sub> PNCs can be obtained in a number of different post-synthetic A-site ion-exchange reaction pathways: for instance, by mixing CsPbI<sub>3</sub>, FAPbI<sub>3</sub> and MAPbI<sub>3</sub> PNCs, or by mixing the CsPbI<sub>3</sub> PNCs with the FA<sub>x</sub>MA<sub>1-x</sub>PbI<sub>3</sub> PNCs described previously. To demonstrate the feasibility of this pathway, we mixed CsPbI<sub>3</sub> (see Fig. S11† for the details of CsPbI<sub>3</sub>), FAPbI<sub>3</sub> and MAPbI<sub>3</sub> PNCs in a 1:1:1 concentration ratio and allowed them to

undergo A-site ion exchange until a thermodynamic equilibrium was reached, resulting in a single symmetric PL peak centered at ~728 nm and a sharp absorption onset at ~750 nm, as can be seen in Fig. 4a. The powder X-ray diffraction pattern of TCPbI<sub>3</sub>, as can be seen from Fig. 4b, suggests the formation of a 3D perovskite phase with cubic/pseudo-cubic symmetry, which is also proved from the TEM images that show the resulting TCPbI<sub>3</sub> retains its cubic morphology after the extended 'A'-site ion-exchange reaction of its parent counterpart (see Fig. S12b†).

A closer look at the XRD data can qualitatively inform us about the effective A-site cation mixing in the TCPbI<sub>3</sub> PNCs. For instance, the diffraction peak of TCPbI<sub>3</sub>, at around  $2\theta = 14.29^\circ$ , shifts towards a lower angle as compared to that of pure CsPbI<sub>3</sub> (at  $2\theta = 14.53^\circ$ ), but when compared to that of pure FAPI ( $2\theta = 14.13^\circ$ ), this peak has slightly higher  $2\theta$  value (see Fig. S13 in the ESI†). This can be correlated with the effective A-site ionic radius in TCPbI<sub>3</sub>. Considering the Shannon ionic radii<sup>68,69</sup> of Cs, FA, and MA, the effective A-site ionic radius of TCPbI<sub>3</sub>, assuming a 1:1:1 ratio, can be calculated and is found to be 2.13 Å, which is larger than that of Cs but smaller than that of FA ions. This indicates lattice expansion in TCPbI<sub>3</sub> as compared to CsPbI<sub>3</sub> due to the incorporation of larger-sized FA and MA ions and lattice compression as



**Fig. 4** (a) UV-Vis absorption and PL emission spectra of the TCPbI<sub>3</sub> NCs; the inset shows the complete absorption spectrum over the 300 nm to 900 nm range. (b) Powder XRD pattern of the TCPbI<sub>3</sub> NCs along with their parent NCs, showing the diffraction pattern of the perovskite phase. (c) UV-Vis absorption spectra of the TCPbI<sub>3</sub> NC film recorded from day 1 to day 7 after preparation in 50–70% relative humidity (RH) under normal light and ambient conditions. (d) Comparison of the change of relative band gap bleaching of the TCPbI<sub>3</sub> NCs and the MAPI-PNC film with time; TCPbI<sub>3</sub> shows comparatively better stability than the MAPI PNC film.

compared to the unit cell of  $\text{FAPbI}_3$ . Since the estimated A-site ionic radius of  $\text{TCPbI}_3$  is almost similar to that of the  $\text{MA}^+$  ion, there is no significant shift in that particular diffraction peak ( $2\theta = 14.29^\circ$ ) of  $\text{TCPbI}_3$  as compared to that in  $\text{MAPI}$ . Moreover, the absence of the signature doublet peaks corresponding to the (211) and (022) diffraction planes of pure  $\text{MAPI}$  at around  $25^\circ$  in the case of the  $\text{TCPbI}_3$  PNCs further indicates that the crystallographic parameters between them are different, although the effective ionic radii are similar in both these compositions. We have further synthesized four different compositions of  $\text{TCPbI}_3$  NCs by mixing  $\text{CsPbI}_3$ ,  $\text{FAPbI}_3$  and  $\text{MAPbI}_3$  PNCs in different volume ratios. The optical and structural properties of these synthesized compositions are shown in Fig. S14.† All these compositions are formed in the black perovskite phase with linear variation of their optical properties.

The optical absorption and emission energy of the  $\text{TCPbI}_3$  PNCs is quite similar to that of  $\text{MAPI}$ , and the diffraction patterns are also very much similar, questioning the necessity of triple-cation perovskite compositions. However, if we compare the stability of these compositions under ambient conditions, the  $\text{TCPbI}_3$  PNCs are comparatively better than the  $\text{MAPI}$  PNCs. Fig. 4c shows the optical absorption spectra of solid films of  $\text{TCPbI}_3$  PNCs recorded for one week at 50–70% RH. If we compare their relative band gap bleaching with that of  $\text{MAPI}$  PNC solid films, we can clearly see that the  $\text{TCPbI}_3$  PNC composition still demonstrates good band-gap absorption performance at the same energy even after 7 days (Fig. 4d), which can also be visualized from the digital images of the films under normal light (see Fig. S15†).

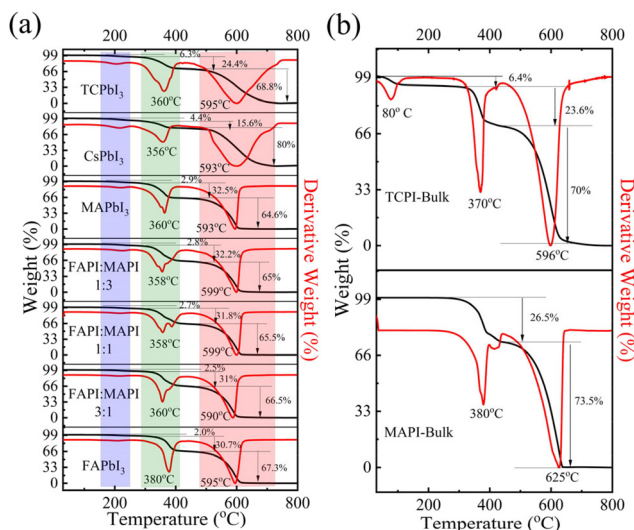
Furthermore, such triple-cation PNC compositions can also be synthesized by cation-exchange reactions between  $\text{CsPbI}_3$  and  $\text{FA}_x\text{MA}_{1-x}\text{PbI}_3$  PNCs under ambient conditions. For example, by reacting the three  $\text{FA}_x\text{MA}_{1-x}\text{PbI}_3$  compositions discussed in the previous section with  $\text{CsPbI}_3$  at similar concentrations, stable homogeneous solid solutions of chemical formula  $\text{Cs}_x\text{FA}_y\text{MA}_{1-x-y}\text{PbI}_3$  can be obtained with tunable optical properties as shown in Fig. S16a–f.† All of these  $\text{TCPbI}_3$  PNCs retain their colloidal stability and structural integrity for over a month.

Apart from moisture stability, thermal stability is another important aspect of perovskite nanocrystals in their practical applications to achieve high-performing, repeatable, and stable optoelectronic devices. Several studies in the past highlighted the thermal decomposition pathways of lead halide perovskite thin films and single crystals using thermogravimetric analysis (TGA).<sup>70–73</sup> For example, in the case of  $\text{MAPI}$  thin films, two weight-loss events have been described: the first one, between  $250^\circ\text{C}$  and  $380^\circ\text{C}$ , corresponds to the decomposition of the organic component ( $\text{MAI}$ ) while the second one, until  $450^\circ\text{C}$ , corresponds to the decomposition of  $\text{PbI}_2$ . In the case of  $\text{FAPI}$  thin films, on the other hand, the first weight loss, corresponding to the decomposition of  $\text{FAI}$ , occurs at the much higher temperature of  $420^\circ\text{C}$ , which has been ascribed to the stronger interaction of  $\text{FA}^+$  with the inorganic  $[\text{PbI}_6]^{4-}$  octahedral matrices as compared to that of

$\text{MA}^+$ . As discussed earlier, the decomposition of  $\text{FA/MA}$ -based perovskites occurs *via* compositional change.

Ma *et al.* have shown, using the TGA-FTIR technique, that at lower temperatures,  $\text{MAPI}$  decomposed *via* the release of  $\text{CH}_3\text{I}$  and  $\text{NH}_3$  gases, and at high temperatures, the decomposition of  $\text{MAPI}$  leads to the formation of  $\text{CH}_4$ . However, the decomposition of  $\text{FAPI}$  leads to the formation of s-triazine ( $\text{HCN}$ )<sub>3</sub> and  $\text{NH}_3$  at low temperatures, and  $\text{HCN}$  and  $\text{NH}_3$  at high temperatures.<sup>73</sup> A few groups have reported the decomposition of  $\text{MAPI}$  *via* the formation of  $\text{HI}$  and  $\text{CH}_3\text{NH}_2$ .<sup>70,71</sup>

To check the thermal stability and decomposition temperatures of the PNCs, we carried out the TGA study by heating samples up to  $800^\circ\text{C}$  under nitrogen at a ramp rate of  $10^\circ\text{C min}^{-1}$ . Unlike the thin films, all the PNC compositions discussed above display three-step weight-loss events, as can be seen from the TGA and their derivative plots shown in Fig. 5a: (a) initial loss ( $\sim 2$ – $6\%$ ) below  $200^\circ\text{C}$ , which may be attributed to the decomposition of organic surface ligands such as  $\text{OA}$  and  $\text{OIAM}$ ; (b) the second step of mass loss, occurring above  $300^\circ\text{C}$ , may be attributed to the decomposition of organic A-site cations and the remaining organic surface ligands; and (c) the final step, above  $500^\circ\text{C}$ , where the major mass loss occurs due to decomposition of  $\text{PbI}_2$ . A similar three-step degradation process has also recently been reported by Li's group for  $\text{Cs}_x\text{FA}_{1-x}\text{PbI}_3$  PNCs.<sup>74</sup> This noted that, unlike  $\text{MAI}$  and  $\text{FAI}$ ,  $\text{CsI}$  decomposes at very high temperatures, above  $600^\circ\text{C}$ , hence in the second step, only  $\sim 15\%$  of the total weight-loss event occurs in the case of pure  $\text{CsPbI}_3$  PNCs as opposed to  $\sim 30\%$  in the PNCs with organic-only A-site cations. As also observed in hybrid lead halide perovskite thin films, in the case of pure  $\text{FAPI}$  PNCs, the second weight-loss event occurs at a slightly higher temperature ( $378^\circ\text{C}$ ) than that in



**Fig. 5** TG curves and their corresponding DTG plots of (a)  $\text{TCPbI}_3$ ,  $\text{CsPbI}_3$ ,  $\text{FAPbI}_3$ ,  $\text{MAPbI}_3$  and  $\text{FA}_x\text{MA}_{1-x}\text{PbI}_3$  PNCs, and (b) the  $\text{MAPbI}_3$  and  $\text{TCPbI}_3$  bulk thin-film perovskites. The percentage mass loss in the three different steps is shown by the arrow along with the corresponding value.

pure MAPI PNCs (360 °C) due to the stronger interaction of FA cations with the  $[\text{PbI}_6]^{4-}$  octahedral units compared to the MA cation.<sup>73</sup> All the solid solutions of  $\text{FA}_x\text{MA}_{1-x}\text{PbI}_3$  and  $\text{TCPbI}_3$  PNCs followed similar three-step weight-loss events. However, when we compare the thermal degradation of the  $\text{TCPbI}_3$  PNC composition with the MA- and FA-based PNC compositions, we can see that the percentage mass loss below 400 °C is ~28% for  $\text{TCPbI}_3$ , which is lower when compared to that of the FA- and MA-based PNCs where we observed 32–35% mass loss, indicating better thermal stability of  $\text{TCPbI}_3$  due to the incorporation of Cs into the A-site. When compared to the degradation of the  $\text{TCPbI}_3$  PNCs with the corresponding single crystals of similar compositions, we observed that both exhibit comparable thermal stability, although the second mass-loss event occurs at a slightly lower temperature in the case of the PNCs. However, in the case of bulk  $\text{TCPbI}_3$ , there is substantial mass loss below 100 °C, which is not observed in the case of the nanocrystals. Such low-temperature decomposition may occur *via* the release of  $\text{CH}_3\text{NH}_2$  and HI, which is reported for MAPI thin films.<sup>75</sup> It is interesting to note here that in their thin-film form, both MAPI and FAPI undergo temperature-dependent crystallographic phase transformations, as reported in multiple literature reports.<sup>52,58,76</sup> For instance, Ambra Pisanu *et al.* reported that the MAPI thin film undergoes a tetragonal-to-cubic phase transition at ~57 °C, while the as-prepared yellow FAPI undergoes a hexagonal-to-cubic phase transition at ~165 °C; moreover, the  $\text{FA}_x\text{MA}_{1-x}\text{PbI}_3$  compositions do not undergo such crystallographic phase transitions. Unlike in thin films, FAPI PNCs crystallize as the black cubic perovskite phase at room temperature, and temperature-dependent *in situ* XRD and thermogravimetric studies in the recent literature<sup>74</sup> prove that FAPI PNCs directly decompose into  $\text{PbI}_2$  without undergoing any intermediate crystal phase transformation. However, similar studies on methylammonium-based PNCs have not been reported so far. Our TGA results do not reveal any signature of temperature-dependent crystallographic phase transformations in any of the PNC compositions including FAPI and MAPI.

The observed enhanced stability of the colloidal PNCs can be attributed to the hydrophobic surface ligands that resist the penetration of water molecules, which in thin-film lead halide perovskites initiates chemical reactions to disintegrate the chemical compositions forming individual components such as lead halides, methylamine, and/or ammonia. By incorporating multiple cations at the A-site, particularly the inorganic cation Cs, such moisture stability is further improved most likely due to reduced reactivity of the inorganic component towards moisture. Inclusion of these inorganic cations also helps in improving the thermal stability as compared to the simple MAPI composition where the organic A-site cation is known to be very volatile. Surface ligand binding strength with different A-site cations also plays a crucial role in the enhancement of moisture and thermal stability. However, further theoretical studies are required for a better understanding.

## Conclusions

In conclusion, we have demonstrated the versatility of A-site composition tuning in colloidal lead halide perovskite nanocrystals by post-synthetic cross-ion exchange that involves methylammonium ions. Such ability has not only provided freedom to tune the optoelectronic properties, but also helped to achieve better moisture stability and thermal stability comparable to that of the bulk/thin-film lead halide perovskites. We have also demonstrated the synthesis of triple-cation iodide-only perovskite PNCs, expanding the compositional space of lead halide perovskite NCs. Such fine control over composition tuning may help to further strengthen perovskite research in general by developing efficient and stable perovskite optoelectronic devices. While in colloidal form, PNC inks are stable for months without losing phase stability, the long-term stability of their bare films under ambient conditions is still not good enough for practical purposes. However, it is to be noted that in real optoelectronic devices, the active layer deposited from the PNC inks is sandwiched between other charge transport layers and electrodes. There have been numerous reports on PNC-based encapsulated optoelectronic devices with operational stability up to hundreds of hours. So, the stability of the PNC films demonstrated in this study does not necessarily reflect their utilization in stable optoelectronic devices. We envisage that decorating the surface of the methylammonium-based PNCs with suitable ligands, such as zwitterionic ligands and amino acid ligands, as well as encapsulating these PNCs with suitable matrices, such as polymers and silica, can further enhance the stability of PNC inks leading to stable optoelectronic devices.

## Experimental section

### Chemicals

Cesium carbonate ( $\text{Cs}_2\text{CO}_3$ ; 99.9%), oleic acid (OA; technical grade, 90%), oleylamine (OAm; technical grade, 70%), 1-octadecene (1-ODE; technical grade, 90%), formamidinium acetate (FA-acetate, 99%), methylamine acetate (MA-acetate; >97.0%), methyl acetate (MeOAc, anhydrous, 99.5%),  $\text{PbI}_2$  (99.99%, trace metal basis),  $\text{Pb}(\text{acetate})_2 \cdot 3\text{H}_2\text{O}$ , hexane (reagent grade,  $\geq 95\%$ ), toluene (AR/ACS, 99.5%), ethanol (Sigma-Aldrich, used without further purification), hydroiodic acid (HI) (57% aq. solution), and diethyl ether (extrapure, 99%) were used in this study.

### Synthesis of $\text{CsPbI}_3$ PNCs

$\text{CsPbI}_3$  was synthesized by following a reported method where a mixture of 1.08 mmol (0.5 g)  $\text{PbI}_2$  and 25 ml of 1-ODE was degassed at 120 °C for 30 minutes. Then a mixture of 2.5 ml of OA and 2.5 ml of OAm, which was preheated at 120 °C, was injected into the  $\text{PbI}_2$  precursor solution. The reaction vessel was further degassed to produce a clear solution of dissolved  $\text{PbI}_2$ . Then it was heated to 180 °C under a continuous  $\text{N}_2$  flow. At a temperature of 175 °C, 2 ml of Cs-oleate stock solution

(the synthesis procedure is mentioned in the ESI†) was swiftly injected into the  $\text{PbI}_2$  precursor solution and the reaction mixture was quenched in an ice bath within 5 seconds to slow down the growth steps. The crude reaction mixture was transferred to a centrifuge tube and washed with 70 ml of MeOAc, followed by centrifugation at 8000 rpm for 10 minutes. The resulting precipitate was dispersed in 3 ml of hexane and stored for further reaction.

### Synthesis of FAPbI<sub>3</sub> PNCs

For FAPbI<sub>3</sub> synthesis, 0.74 mmol (0.344 g)  $\text{PbI}_2$  and 20 ml of 1-ODE were loaded into a 3-necked round bottom flask and degassed at 120 °C for 30 minutes. Then, a mixture of 4 ml of OA and 2 ml of OlAm, preheated at 120 °C, was injected to dissolve the  $\text{PbI}_2$ . The reaction mixture was further degassed to completely dissolve  $\text{PbI}_2$  and form a clear solution. Then the mixture was transferred into an  $\text{N}_2$  atmosphere and the temperature was reduced to 80 °C. At 80 °C, 5 ml of FA-oleate stock solution (see the ESI† for FA-oleate preparation) was swiftly injected into the  $\text{PbI}_2$  precursor and the reaction mixture was quenched in an ice water bath after 20 seconds. Then, 5 ml of MeOAc was added to the reaction mixture. Then it was centrifuged at 8000 rpm for 30 minutes. After that, the precipitate was dispersed in 3 ml of hexane and stored for the ion-exchange reaction.

### Synthesis of MAPbI<sub>3</sub> PNCs via the two-precursor method using PbI<sub>2</sub> (route I)

MAPbI<sub>3</sub> NC was synthesized by transferring 0.37 mmol (0.172 g)  $\text{PbI}_2$  and 10 ml of 1-ODE into a 3-necked round bottom flask, which was then degassed at 120 °C for 30 minutes. Then a mixture of 2 ml of OA and 1 ml of OlAm, preheated to 120 °C, was injected into the  $\text{PbI}_2$  precursor solution. Then it was further degassed for complete dissolution of  $\text{PbI}_2$ . The reaction mixture was transferred into an  $\text{N}_2$  atmosphere and the temperature was decreased to 80 °C. At 80 °C, 5 ml of MA-oleate stock solution (synthesis of MA-oleate is provided in the ESI†) was swiftly injected and the reaction mixture was quenched after 15 seconds in an ice water bath. The crude product was transferred to a centrifuge tube and 1 ml of hexane and 3 ml of MeOAc were added. Then it was centrifuged at 8000 rpm for 15 minutes and the resulting precipitate was dispersed in 2 ml of hexane and stored for further use.

### Synthesis of MAPbI<sub>3</sub> PNCs via the three-precursor method using OAmI (route II)

$\text{Pb}(\text{acetate})_2 \cdot 3\text{H}_2\text{O}$  (0.152 g, 0.4 mmol, Aldrich, 99.99%), MA-acetate (0.136 g, 1.5 mmol), ODE (16 mL, dried), and OA (4 mL, dried) were transferred into a 100 ml 3-necked round bottom flask and dried under vacuum for 30 min at 60 °C. Under these conditions, all the salts were dissolved in ODE. Under  $\text{N}_2$ , the resulting mixture was heated to 80 °C, and subsequently into it 1.2 mmol OAmI in 4 ml of toluene was swiftly injected (synthesis of OAmI is described in the ESI†). The mixture was immediately placed in an ice water bath and the reaction was quenched within 20 s. The crude solution was

divided into two equal parts in two centrifuge tubes and, to each part, 10 ml of MeOAc was added. Then each tube was centrifuged at 8000 rpm for 10 min, the resulting precipitate was dispersed in 3 ml of *n*-hexane and stored in a glass vial. After 1 hour, a small amount of precipitate was observed at the bottom of the glass vial. The upper part of the solution was isolated and centrifuged at 3000 rpm for 5 minutes. The supernatant was then stored for further use.

### Synthesis of PNC solid solution via A-site cation exchange

To synthesize the solid solution of NCs of FAPbI<sub>3</sub> and MAPbI<sub>3</sub>, their concentration was first matched by adjusting their absorption value in the UV-Vis absorption spectra. Then the colloidal solution of FAPbI<sub>3</sub> and MAPbI<sub>3</sub> (synthesized *via* route I) was mixed at different volume ratios at room temperature to make an alloy of  $\text{FA}_x\text{MA}_{1-x}\text{PbI}_3$ . The triple-cation PNCs were synthesized by reacting CsPbI<sub>3</sub> with different compositions of  $\text{FA}_x\text{MA}_{1-x}\text{PbI}_3$ . In this case also the absorption spectra of individual samples were recorded and their concentration was matched in a similar way to that mentioned before. Then the colloidal solution of CsPbI<sub>3</sub> and  $\text{FA}_x\text{MA}_{1-x}\text{PbI}_3$  was mixed in a 1 : 1 ratio at room temperature. In an alternative method, all the three parent NCs, *i.e.*, CsPbI<sub>3</sub>, FAPbI<sub>3</sub> and MAPbI<sub>3</sub>, were mixed in different volume ratios at room temperature to form TCPbI<sub>3</sub> NCs. In all of these ion exchange reactions, the parent NCs were cleaned only once with MeOAc after synthesis, and used for 'A' site composition tuning at room temperature under ambient conditions. The equilibrium of these reactions is reached within ~2 minutes, as evidenced by the homogeneous PL emission spectra that do not change over time in peak energy and width.

### Characterization

UV-Vis absorption spectra were recorded using a Cary 5000 UV-Vis-NIR spectrophotometer while steady-state PL emission and PLE spectra were recorded using a Cary Eclipse fluorescence spectrometer. Photoluminescence lifetime measurements were carried out using a Fluoro Log3-Triple Illuminator and an IBH Horiba Jobin Yvon instrument and the samples were excited with a 484 nm picosecond-light-emitting diode laser (NanoLED). Powder XRD measurements were performed on drop-cast QD films using a Bruker D-8 Advance powder X-ray diffractometer with Cu K $\alpha$  radiation ( $\lambda = 1.5406 \text{ \AA}$ ). TEM images were captured by using an FEI-Talos 200 $\times$  electron microscope operated at 200 kV. All TGA measurements were carried out using a TGA Q500 instrument. The samples were heated in a temperature range of 25–800 °C at a heating rate of 10 °C min<sup>-1</sup> under an inert atmosphere. For the TGA measurements, the samples were washed two times with MeOAc and 20–30 mg of each solid NC sample was used.

### Data availability

The data supporting this article have been included as part of the ESI.†

## Conflicts of interest

There are no conflicts of interest to be declared.

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## References

- G. Nedelcu, L. Protesescu, S. Yakunin, M. I. Bodnarchuk, M. J. Grotevent and M. V. Kovalenko, *Nano Lett.*, 2015, **15**(8), 5635–5640.
- Q. A. Akkerman, V. D'Innocenzo, S. Accornero, A. Scarpellini, A. Petrozza, M. Prato and L. Manna, *J. Am. Chem. Soc.*, 2015, **137**(32), 10276–10281.
- M. Suri, A. Hazarika, B. W. Larson, Q. Zhao, M. Vallés-Pelarda, T. D. Siegler, M. K. Abney, A. J. Ferguson, B. A. Korgel and J. M. Luther, *ACS Energy Lett.*, 2019, **4**, 1954–1960.
- A. Kojima, K. Teshima, Y. Shirai and T. Miyasaka, *J. Am. Chem. Soc.*, 2009, **131**(17), 6050–6051.
- C. K. Møller, *Nature*, 1957, **180**, 981–982.
- D. Weber, *Z. Naturforsch., B*, 1978, **33**(12), 1443–1445.
- M. M. Lee, J. Teuscher, T. Miyasaka, T. N. Murakami and H. J. Snaith, *Science*, 2012, **338**(6107), 643–647.
- K. Wang, Z. Jin, L. Liang, H. Bian, D. Bai, H. Wang, J. Zhang, Q. Wang and S. Liu, *Nat. Commun.*, 2018, **9**(1), 4544.
- W. Xu, Q. Hu, S. Bai, C. Bao, Y. Miao, Z. Yuan, T. Borzda, A. J. Barker, E. Tyukalova, Z. Hu, *et al.*, *Nat. Photonics*, 2019, **13**(6), 418–424.
- M. Saliba, S. M. Wood, J. B. Patel, P. K. Nayak, J. Huang, J. A. Alexander-Webber, B. Wenger, S. D. Stranks, M. T. Hörantner, J. T.-W. Wang, *et al.*, *Adv. Mater.*, 2016, **28**(5), 923–929.
- V. V. Belykh, D. R. Yakovlev, M. M. Glazov, P. S. Grigoryev, M. Hussain, J. Rautert, D. N. Dirin, M. V. Kovalenko and M. Bayer, *Nat. Commun.*, 2019, **10**(1), 673.
- X. Geng, F. Wang, H. Tian, Q. Feng, H. Zhang, R. Liang, Y. Shen, Z. Ju, G.-Y. Gou, N. Deng, *et al.*, *ACS Nano*, 2020, **14**(3), 2860–2868.
- <https://www.nrel.gov/pv/cell-efficiency.html> (accessed 2024 May 25).
- L. Protesescu, S. Yakunin, M. I. Bodnarchuk, F. Krieg, R. Caputo, C. H. Hendon, R. X. Yang, A. Walsh and M. V. Kovalenko, *Nano Lett.*, 2015, **15**(6), 3692–3696.
- J. Song, J. Li, X. Li, L. Xu, Y. Dong and H. Zeng, *Adv. Mater.*, 2015, **27**(44), 7162–7167.
- Y. Wang, X. Li, J. Song, L. Xiao, H. Zeng and H. Sun, *Adv. Mater.*, 2015, **27**(44), 7101–7108.
- J. Pan, S. P. Sarmah, B. Murali, I. Dursun, W. Peng, M. R. Parida, J. Liu, L. Sinatra, N. Alyami, C. Zhao, *et al.*, *J. Phys. Chem. Lett.*, 2015, **6**(24), 5027–5033.
- J. Pan, L. N. Quan, Y. Zhao, W. Peng, B. Murali, S. P. Sarmah, M. Yuan, L. Sinatra, N. M. Alyami, J. Liu, *et al.*, *Adv. Mater.*, 2016, **28**(39), 8718–8725.
- Q. Zhao, A. Hazarika, X. Chen, S. P. Harvey, B. W. Larson, G. R. Teeter, J. Liu, T. Song, C. Xiao, L. Shaw, *et al.*, *Nat. Commun.*, 2019, **10**(1), 2842.
- A. Swarnkar, A. R. Marshall, E. M. Sanehira, B. D. Chernomordik, D. T. Moore, J. A. Christians, T. Chakrabarti and J. M. Luther, *Science*, 2016, **354**(6308), 92–95.
- Q. Zeng, X. Zhang, Q. Bing, Y. Xiong, F. Yang, H. Liu, J.-Y. Liu, H. Zhang, W. Zheng, A. L. Rogach, *et al.*, *ACS Energy Lett.*, 2022, **7**(6), 1963–1970.
- G. E. Eperon, S. D. Stranks, C. Menelaou, M. B. Johnston, L. M. Herz and H. J. Snaith, *Energy Environ. Sci.*, 2014, **7**, 982–988.
- O. Vybornyi, S. Yakunin and M. V. Kovalenko, *Nanoscale*, 2016, **8**(12), 6278–6283.
- S. Das and A. Samanta, *ACS Energy Lett.*, 2021, **6**(11), 3780–3787.
- M. Imran, V. Caligiuri, M. Wang, L. Goldoni, M. Prato, R. Krahné, L. De Trizio and L. Manna, *J. Am. Chem. Soc.*, 2018, **140**(7), 2656–2664.
- F. Zhang, H. Zhong, C. Chen, X. G. Wu, X. Hu, H. Huang, J. Han, B. Zou and Y. Dong, *ACS Nano*, 2015, **9**(4), 4533–4542.
- Y. Zhang, T. D. Siegler, C. J. Thomas, M. K. Abney, T. Shah, A. De Gorostiza, R. M. Greene and B. A. Korgel, *Chem. Mater.*, 2020, **32**(13), 5410–5423.
- M. Saliba, T. Matsui, J.-Y. Seo, K. Domanski, J.-P. Correa-Baena, M. K. Nazeeruddin, S. M. Zakeeruddin, W. Tress, A. Abate, A. Hagfeldt, *et al.*, *Energy Environ. Sci.*, 2016, **9**(6), 1989–1997.
- Y. Sun, J. Peng, Y. Chen, Y. Yao and Z. Liang, *Sci. Rep.*, 2017, **7**(1), 46193.
- C. C. Stoumpos, C. D. Malliakas and M. G. Kanatzidis, *Inorg. Chem.*, 2013, **52**(15), 9019–9038.
- J. H. Noh, S. H. Im, J. H. Heo, T. N. Mandal and S. I. Seok, *Nano Lett.*, 2013, **13**(4), 1764–1769.
- M. I. Saidaminov, A. L. Abdelhady, B. Murali, E. Alarousu, V. M. Burlakov, W. Peng, I. Dursun, L. Wang, Y. He, G. Maculan, *et al.*, *Nat. Commun.*, 2015, **6**(1), 7586.
- A. Jaffe, Y. Lin, C. M. Beavers, J. Voss, W. L. Mao and H. I. Karunadasa, *ACS Cent. Sci.*, 2016, **2**(4), 201–209.
- A. S. Masadeh, E. S. Božin, C. L. Farrow, G. Paglia, P. Juhas, S. J. L. Billinge, A. Karkamkar and M. G. Kanatzidis, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2007, **76**(11), 115413.
- F. Bertolotti, D. Moscheni, A. Guagliardi and N. Masciocchi, *Eur. J. Inorg. Chem.*, 2018, **34**, 3789–3803.

- 36 S. J. L. Billinge and I. Levin, *Science*, 2007, **316**(5824), 561–565.
- 37 S. Tao, J. Billet, J. De Roo and S. J. L. Billinge, *Chem. Mater.*, 2024, **36**(21), 10912–10921.
- 38 X. Yang, A. S. Masadeh, J. R. McBride, E. S. Božin, S. J. Rosenthal and S. J. L. Billinge, *Phys. Chem. Chem. Phys.*, 2013, **15**(22), 8480–8486.
- 39 T. Moot, D. R. Dikova, A. Hazarika, T. H. Schloemer, S. N. Habisreutinger, N. Leick, S. P. Dunfield, B. A. Rosales, S. P. Harvey, J. R. Pfeilsticker, *et al.*, *Chem. Mater.*, 2020, **32**(18), 7850–7860.
- 40 J. M. Frost, K. T. Butler, F. Brivio, C. H. Hendon, M. van Schilfgaarde and A. Walsh, *Nano Lett.*, 2014, **14**(5), 2584–2590.
- 41 B. Hailegnaw, S. Kirmayer, E. Edri, G. Hodes and D. Cahen, *J. Phys. Chem. Lett.*, 2015, **6**(9), 1543–1547.
- 42 W. Huang, J. S. Manser, P. V. Kamat and S. Ptasincka, *Chem. Mater.*, 2016, **28**(1), 303–311.
- 43 E. Smecca, Y. Numata, I. Deretzis, G. Pellegrino, S. Boninelli, T. Miyasaka, A. La Magna and A. Alberti, *Phys. Chem. Chem. Phys.*, 2016, **18**, 13413–13422.
- 44 N. Aristidou, C. Eames, M. S. Islam and S. A. Haque, *J. Mater. Chem. A*, 2017, **5**, 25469–25475.
- 45 W. Wei and Y. H. Hu, *Int. J. Energy Res.*, 2017, **41**(7), 1063–1069.
- 46 J. A. McLeod and L. Liu, *J. Phys. Chem. Lett.*, 2018, **9**(9), 2411–2417.
- 47 C. Zheng and O. Rubel, *J. Phys. Chem. C*, 2019, **123**(32), 19385–19394.
- 48 A. Hazarika, Q. Zhao, E. A. Gaulding, J. A. Christians, B. Dou, A. R. Marshall, T. Moot, J. J. Berry, J. C. Johnson and J. M. Luther, *ACS Nano*, 2018, **12**(10), 10327–10337.
- 49 S. Yakunin, L. Protesescu, F. Krieg, M. I. Bodnarchuk, G. Nedelcu, M. Humer, G. De Luca, M. Fiebig, W. Heiss and M. V. Kovalenko, *Nat. Commun.*, 2015, **6**(1), 8056.
- 50 M. Hao, Y. Bai, S. Zeiske, L. Ren, J. Liu, Y. Yuan, N. Zarrabi, N. Cheng, M. Ghasemi, P. Chen, *et al.*, *Nat. Energy*, 2020, **5**(1), 79–88.
- 51 C. Otero-Martínez, M. Imran, N. J. Schrenker, J. Ye, K. Ji, A. Rao, S. D. Stranks, R. L. Z. Hoyer, S. Bals, L. Manna, *et al.*, *Angew. Chem., Int. Ed.*, 2022, **61**, e202205617.
- 52 A. Pisanu, C. Ferrara, P. Quadrelli, G. Guizzetti, M. Patrini, C. Milanese, C. Tealdi and L. Malavasi, *J. Phys. Chem. C*, 2017, **121**(16), 8746–8751.
- 53 D. J. Norris and M. G. Bawendi, *Phys. Rev. B:Condens. Matter Mater. Phys.*, 1996, **53**(24), 16338–16346.
- 54 O. I. Micic, C. J. Curtis, K. M. Jones, J. R. Sprague and A. J. Nozik, *J. Phys. Chem.*, 1994, **98**(19), 4966–4969.
- 55 S. Ithurria and D. V. Talapin, *J. Am. Chem. Soc.*, 2012, **134**(45), 18585–18590.
- 56 J. A. Vigil, A. Hazarika, J. M. Luther and M. F. Toney, *ACS Energy Lett.*, 2020, **5**(8), 2475–2482.
- 57 J. A. Vigil, B. M. Wieliczka, B. W. Larson, M. Abdelsamie, N. S. Dutta, A. M. Haque, A. Hazarika, J. M. Luther and M. F. Toney, *Chem. Mater.*, 2023, **35**(23), 9924–9934.
- 58 O. J. Weber, B. Charles and M. T. Weller, *J. Mater. Chem. A*, 2016, **4**, 15375–15382.
- 59 A. Francisco-López, B. Charles, M. I. Alonso, M. Garriga, M. Campoy-Quiles, M. T. Weller and A. R. Goñi, *J. Phys. Chem. C*, 2020, **124**(6), 3448–3458.
- 60 M. Thambidurai, S. Foo, K. M. Muhammed Salim, P. C. Harikesh, A. Bruno, N. F. Jamaludin, S. Lie, N. Mathews and C. Dang, *J. Mater. Chem. C*, 2019, **7**(17), 5028–5036.
- 61 I. M. Pavlovets, M. C. Brennan, S. Draguta, A. Ruth, T. Moot, J. A. Christians, K. Aleshire, S. P. Harvey, S. Toso, S. U. Nanayakkara, *et al.*, *ACS Energy Lett.*, 2020, **5**(9), 2802–2810.
- 62 N. Landi, E. Maurina, D. Marongiu, A. Simbula, S. Borsacchi, L. Calucci, M. Saba, E. Carignani and M. Geppi, *J. Phys. Chem. Lett.*, 2022, **13**(40), 9517–9525.
- 63 E. T. Hoke, D. J. Slotcavage, E. R. Dohner, A. R. Bowring, H. I. Karunadasa and M. D. McGehee, *Chem. Sci.*, 2015, **6**(1), 613–617.
- 64 S. Draguta, O. Sharia, S. J. Yoon, M. C. Brennan, Y. V. Morozov, J. S. Manser, P. V. Kamat, W. F. Schneider and M. Kuno, *Nat. Commun.*, 2017, **8**(1), 200.
- 65 A. J. Barker, A. Sadhanala, F. Deschler, M. Gandini, S. P. Senanayak, P. M. Pearce, E. Mosconi, A. J. Pearson, Y. Wu, A. R. Srimath Kandada, *et al.*, *ACS Energy Lett.*, 2017, **2**(6), 1416–1424.
- 66 A. J. Knight, A. D. Wright, J. B. Patel, D. P. McMeekin, H. J. Snaith, M. B. Johnston and L. M. Herz, *ACS Energy Lett.*, 2019, **4**(1), 75–84.
- 67 P. Vashishtha, S. A. Veldhuis, S. S. H. Dintakurti, N. L. Kelly, B. E. Griffith, A. A. M. Brown, M. S. Ansari, A. Bruno, N. Mathews, Y. Fang, *et al.*, *J. Mater. Chem. C*, 2020, **8**, 11805–11821.
- 68 R. Shannon, *Acta Crystallogr., Sect. A*, 1976, **32**(5), 751–767.
- 69 G. Kieslich, S. Sun and A. K. Cheetham, *Chem. Sci.*, 2015, **6**(6), 3430–3433.
- 70 A. Dualeh, P. Gao, S. I. Seok, M. K. Nazeeruddin and M. Grätzel, *Chem. Mater.*, 2014, **26**(21), 6160–6164.
- 71 A. E. Williams, P. J. Holliman, M. J. Carnie, M. L. Davies, D. A. Worsley and T. M. Watson, *J. Mater. Chem. A*, 2014, **2**(45), 19338–19346.
- 72 E. J. Juarez-Perez, Z. Hawash, S. R. Raga, L. K. Ono and Y. Qi, *Energy Environ. Sci.*, 2016, **9**(11), 3406–3410.
- 73 L. Ma, D. Guo, M. Li, C. Wang, Z. Zhou, X. Zhao, F. Zhang, Z. Ao and Z. Nie, *Chem. Mater.*, 2019, **31**(20), 8515–8522.
- 74 S. Wang, Q. Zhao, A. Hazarika, S. Li, Y. Wu, Y. Zhai, X. Chen, J. M. Luther and G. Li, *Nat. Commun.*, 2023, **14**(1), 2216.
- 75 A. Cicciooli and A. Latini, *J. Phys. Chem. Lett.*, 2018, **9**(13), 3756–3765.
- 76 W. T. M. Van Gompel, R. Herckens, G. Reekmans, B. Ruttens, J. D’Haen, P. Adriaensens, L. Lutsen and D. Vanderzande, *J. Phys. Chem. C*, 2018, **122**(8), 4117–4124.