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## ZnX<sub>2</sub> mediated post-synthetic transformation of zero dimensional Cs<sub>4</sub>PbBr<sub>6</sub> nanocrystals for opto-electronic applications†

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Herein we demonstrate a facile approach for the synthesis of all inorganic cesium lead halide perovskite nanocrystal composites CsPbX<sub>3</sub> (X = Cl, Br, I) with high quantum yield by post-synthetic modulation of zero dimensional Cs<sub>4</sub>PbBr<sub>6</sub> nanocrystals with ZnX<sub>2</sub> salts. The transformation of Cs<sub>4</sub>PbBr<sub>6</sub> nanocrystals into CsPbBr<sub>3</sub> takes place in two steps, the first step being the surface modification of the Cs<sub>4</sub>PbBr<sub>6</sub> nanocrystals with Zn<sup>2+</sup> ions and the second step being extraction of CsBr by the Zn<sup>2+</sup> ions resulting in the formation of composite Cs<sub>4</sub>PbBr<sub>6</sub>/CsPbBr<sub>3</sub> nanocrystals. The transformed composite nanocrystals were found to have a PL QY exceeding 90% and the shape of the nanocrystals also changed from hexagonal to cubic shaped. Owing to the highly ionic nature of the nanocrystals, complete anion exchange could be also realized using ZnI<sub>2</sub> salt. In the case of the iodide post-treated samples, nanorods were obtained which exhibited bright red photoluminescence. Photodetectors based on the ZnI<sub>2</sub> treated Cs<sub>4</sub>PbBr<sub>6</sub> NCs were fabricated, and the photodetectors exhibited a high on/off ratio with a fast response time. The excellent optoelectronic properties make this treatment versatile for a wide range of functional optoelectronic devices like light emitting diodes and photovoltaic devices.

## Introduction

Lead halide perovskite NCs (especially CsPbX<sub>3</sub> where X = Cl, Br, I) have received massive attention in recent years as favorable optoelectronic materials due to their broad absorption,<sup>1</sup> intense photoluminescence (PL),<sup>1</sup> high photoluminescence (PL) quantum yield (QY),<sup>2</sup> very narrow emission line width,<sup>3</sup> high defect tolerance,<sup>4</sup> and wide-range band gaps which can be tuned

by controlling the composition<sup>5,6</sup> and morphology.<sup>7,8</sup> Due to these interesting properties exhibited by the lead halide perovskite NCs they have been examined in the context of their use in optoelectronic devices, such as photovoltaics,<sup>9,10</sup> light emitting diodes,<sup>11–15</sup> lasers<sup>16</sup> and photodetectors.<sup>17</sup> In the recent past, numerous other types of cesium lead halide NCs, with structural dimensionality different from the three dimensional orthorhombic/cubic CsPbX<sub>3</sub> have gained a lot of attention.<sup>18–25</sup> The structural dimensionality in these structures depends on the extent of coupling of PbX<sub>6</sub> octahedra in the crystal. If all the corners of the lead hexahalide octahedra are shared, then they are electronically coupled in all three dimensions and referred to as 3D perovskites. If all the corners of the lead hexahalide octahedra are not shared, they are referred to as 0D perovskites.<sup>25</sup>

It is extensively recognized that the excellent optical properties in lead halide based perovskite materials come from the high defect tolerance.<sup>4</sup> Even the highly luminescent perovskite NCs prepared by the hot injection method are really not defect-free, and these defects serve as trapping centers for exciton recombination and decrease the photoluminescence quantum yields (PLQYs) of the NCs.<sup>26–33</sup> Different groups have recently synthesized 0D colloidal Cs<sub>4</sub>PbBr<sub>6</sub> NCs and it has been found that these nanocrystals don't exhibit any photoluminescence.<sup>25</sup> However, the non-luminescent 0D Cs<sub>4</sub>PbBr<sub>6</sub> NCs can be converted to 3D CsPbBr<sub>3</sub> NCs using additives like water,<sup>34</sup> PbBr<sub>2</sub>,<sup>35</sup> ligands,<sup>36</sup> and Prussian blue.<sup>37</sup> The non-luminescent nature of Cs<sub>4</sub>PbBr<sub>6</sub> at room temperature is attributed to the origin of structural and point defects.<sup>38</sup> Nikl *et al.* observed the true emission of Cs<sub>4</sub>PbBr<sub>6</sub> at low temperatures (4 K) and the true emission is centered at 375 nm. In the case of thin films at room temperature, Cs<sub>4</sub>PbBr<sub>6</sub> is mixed with traces of the CsPbBr<sub>3</sub> phase, which in turn results in the emission at 545 nm. However in the case of nanocrystals, no emission has been observed so far for the pure Cs<sub>4</sub>PbBr<sub>6</sub> nanocrystals.

Recently it has been shown that addition of appropriate metal halide salts significantly enhances the photoluminescence intensity of CsPbBr<sub>3</sub> nanocrystals by improving

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the surface capping. Herein we demonstrate  $\text{ZnX}_2$  mediated post-synthetic transformation of  $\text{Cs}_4\text{PbBr}_6$  nanocrystals (NCs) into  $\text{CsPbX}_3$  nanostructures through an intermediate composite  $\text{Cs}_4\text{PbX}_6/\text{CsPbX}_3$ . We find that this procedure results in a highly reproducible increase of the PL QY of up to 90% for the  $\text{ZnX}_2$  treatment. The transformation takes place through the surface passivation of  $\text{Cs}_4\text{PbBr}_6$  nanocrystals with  $\text{ZnX}_2$  followed by anion exchange (in the case of  $\text{ZnI}_2$  and  $\text{ZnCl}_2$ ) resulting in the formation of an intermediate  $\text{Cs}_4\text{PbX}_6@/\text{CsPbX}_3$  passivated by  $\text{ZnX}_2$  followed by the extraction of  $\text{CsBr}$  from the composite nanostructures. Interestingly, as soon the  $\text{ZnX}_2$  is added to the nanocrystals, excitonic absorption resulting from the  $\text{Cs}_4\text{PbBr}_6$  particles passivated with  $\text{ZnX}_2$  is observed. This is the first time that the true emission of the  $\text{Cs}_4\text{PbX}_6$  crystal structure in the nanocrystalline phase was observed at room temperature.

The  $\text{Cs}_4\text{PbBr}_6$  nanocrystals were synthesized according to the hot injection strategy as described in the Experimental section. The  $\text{Cs}_4\text{PbBr}_6$  nanocrystal dispersion in hexane is completely transparent and as shown in the TEM images in Fig. 1b–d, the pristine  $\text{Cs}_4\text{PbBr}_6$  nanocrystals consist of hexagons of around 30 nm in size. The powder diffraction pattern indicated that the nanocrystals crystallized in the rhombohedral  $\text{Cs}_4\text{PbBr}_6$  phase in the space group  $R3c$ .

The pristine  $\text{Cs}_4\text{PbBr}_6$  nanocrystals (NCs) were post treated with a  $\text{ZnX}_2$ /hexane solution and used for further characterization. Upon addition of  $\text{ZnX}_2$  ( $\text{X} = \text{Cl}, \text{Br}, \text{I}$ ) solution, clear color change was observed (Fig. 2a). And the initially non-luminescent solution turned into a brightly luminescent solution under UV light after the post-treatment (Fig. 2b). To understand the reason for the observed color changes, powder diffraction analysis was carried out. Fig. 2c shows the X-ray diffraction patterns obtained for the pristine and  $\text{ZnX}_2$  treated

NCs. The XRD pattern of the parent NCs indicated that they are crystalline and matched well with the reported rhombohedral  $\text{Cs}_4\text{PbBr}_6$  phase. No characteristic reflections of  $\text{CsPbBr}_3$ ,  $\text{CsBr}$  or  $\text{PbBr}_2$  were observed. Upon treating with  $\text{ZnBr}_2$ , new reflections were observed along with the reflections from  $\text{Cs}_4\text{PbBr}_6$ . All the new reflections could be attributed to the  $\text{CsPbBr}_3$  phase and the post treatment resulted in the formation of the composite  $\text{Cs}_4\text{PbBr}_6/\text{CsPbBr}_3$ . Similar behavior was observed in the case of  $\text{ZnI}_2$  treated and  $\text{ZnCl}_2$  treated NCs wherein a composite of  $\text{Cs}_4\text{PbX}_6/\text{CsPbX}_3$  is formed. In the case of  $\text{ZnI}_2$  treated NCs, the pristine  $\text{Cs}_4\text{PbBr}_6$  reflections were shifted towards the lower  $2\theta$  indicating that anion exchange took place from the bromide to iodide resulting in the formation of  $\text{Cs}_4\text{PbI}_6$ . A slight shift towards higher  $2\theta$  was observed in the case of  $\text{ZnCl}_2$  treated NCs. Interestingly, when the  $\text{ZnX}_2$  post treated solutions were left without centrifugation for a few days, the luminescence of the solution was reduced in comparison to that of the samples which were centrifuged and washed after the  $\text{ZnX}_2$  post-treatment. We have carried out XRD analysis on those samples left without centrifugation and Fig. 2d shows the diffraction patterns obtained for the same samples. From the diffraction patterns it can be seen that the  $\text{ZnBr}_2$  left in the solution results in the complete transformation of 0D  $\text{Cs}_4\text{PbBr}_6$  NCs into 3D  $\text{CsPbBr}_3$  NCs. In the case of  $\text{ZnI}_2$  post-treatment, the reflections from 0D  $\text{Cs}_4\text{PbI}_6$  still exist, indicating that the transformation from 0D to 3D is not complete even after a few days.

The absorption spectra of the  $\text{ZnX}_2$  treated  $\text{Cs}_4\text{PbBr}_6$  NCs are shown in Fig. 3a. For the pristine nanocrystals, the absorption and the photoluminescence features are consistent with those in the previously reported studies. The absorption spectra of the as-prepared nanocrystals feature a strong and narrow absorption at 313 nm. The long tail in the absorption spectrum indicates that a certain degree of coupling exists between the  $\text{PbBr}_6$  octahedra. Fig. 2a and b show the photographs of the pristine  $\text{Cs}_4\text{PbBr}_6$  and  $\text{ZnX}_2$  treated NCs under room light and UV light. Upon treating the NCs with the  $\text{ZnBr}_2$ /hexane solution, the initially colorless solution turned green color. The same is observed in the UV-vis absorption spectrum with a red shift of the absorption band edge to 510 nm. The features are similar to the characteristic feature of the 3D  $\text{CsPbBr}_3$  NCs,<sup>1–10</sup> probably indicating the transformation of  $\text{Cs}_4\text{PbBr}_6$  into  $\text{CsPbBr}_3$  NCs. Along with this, another hump was observed at 420 nm in the  $\text{ZnBr}_2$  treated sample. This hump was not observed previously in the pure  $\text{CsPbBr}_3$  NCs,<sup>1–10</sup> the  $\text{ZnBr}_2$  stabilized  $\text{CsPbBr}_3$  NCs and the  $\text{Zn}^{2+}$  doped  $\text{CsPbBr}_3$  nanocrystals.<sup>39</sup> To see if similar behavior is observed with other zinc halides, the  $\text{Cs}_4\text{PbBr}_6$  NCs were also treated with  $\text{ZnI}_2$  and  $\text{ZnCl}_2$ .  $\text{ZnCl}_2$  treatment resulted in a slight blue shift in comparison to  $\text{ZnBr}_2$  treated NCs. However, in the case of  $\text{ZnCl}_2$  treated NCs, no hump in the absorption spectrum was evident. In the case of  $\text{ZnI}_2$ , an increased red shift (up to 670 nm) in comparison to that in the  $\text{ZnBr}_2$  treatment was observed. Along with the red shifted absorption, a sharp peak at 550 nm can be seen.

To confirm if the peak observed at 420 nm arises from the sample, an aliquot of the  $\text{Cs}_4\text{PbBr}_6$  NCs was taken in a cuvette, different concentrations of  $\text{ZnBr}_2$  were added to the sample and

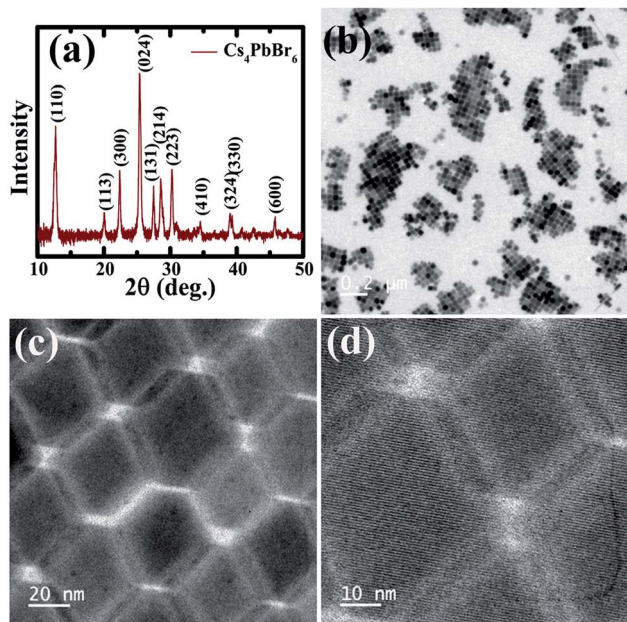


Fig. 1 (a) X-ray diffraction patterns obtained of the pristine  $\text{Cs}_4\text{PbBr}_6$  NCs. (b–d) TEM images of the pristine  $\text{Cs}_4\text{PbBr}_6$  NCs at different magnifications.



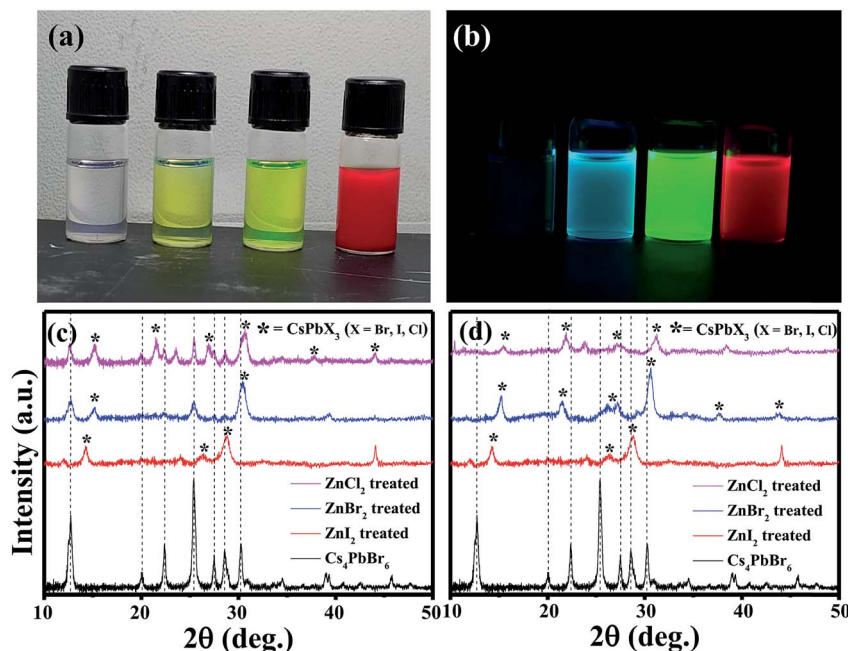


Fig. 2 Photographs of the solutions of Cs<sub>4</sub>PbBr<sub>6</sub>, ZnCl<sub>2</sub> treated, ZnBr<sub>2</sub> treated and ZnI<sub>2</sub> treated NCs, respectively under ambient light (a) and under UV light (b). (c) X-ray diffraction patterns obtained for the pristine and ZnX<sub>2</sub> treated NCs. (d) X-ray diffraction patterns obtained of the samples left without centrifugation after the addition of ZnX<sub>2</sub>.

the absorption spectrum of the ZnBr<sub>2</sub> treated NCs was measured. Fig. 3b shows the absorption spectra measured using different concentrations of the ZnBr<sub>2</sub>/hexane solution added to the cuvette containing NCs. A sharp absorption peak at 425 nm along with a long tail extending up to 510 nm was observed. The absorption intensity of the tail increased gradually and a sharp band edge at 510 nm, which is a characteristic peak of CsPbBr<sub>3</sub>, was observed. The peak at 425 nm is blue shifted to 420 nm and is masked (though visible) by the strong absorption of CsPbBr<sub>3</sub> in this region. Fig. S1† shows the comparison of the absorption spectra of the CsPbBr<sub>3</sub> nanocrystals synthesized and the ZnBr<sub>2</sub> treated Cs<sub>4</sub>PbBr<sub>6</sub> NCs. A similar concentration dependent study was carried out in the case of ZnI<sub>2</sub> treated Cs<sub>4</sub>PbBr<sub>6</sub> NCs. Fig. 3c shows the evolution of the absorption spectrum of ZnI<sub>2</sub> treated NCs with different amounts of ZnI<sub>2</sub> added. A sharp absorption at 540 nm is evident even with high amounts of added ZnI<sub>2</sub>. ZnBr<sub>2</sub> treated NC samples were found to be highly stable even after 2 months of maintaining under 70% RH conditions. Fig. S2† shows photographs under UV light for the ZnBr<sub>2</sub> treated NC sample and the as-prepared CsPbBr<sub>3</sub> NCs after storing under 70% RH conditions.

To corroborate the observed absorption behavior, photoluminescence (PL) studies were also carried out on the parent NCs and ZnX<sub>2</sub> treated NCs. Fig. 3d shows the PL spectra of the pristine NCs and ZnBr<sub>2</sub> treated NCs. No strong PL emission was observed, which is consistent with the previously observed decay of excitonic PL above temperatures of 100 K. Very weak PL emission centered at 515 nm was observed which can be attributed to the CsPbBr<sub>3</sub> phase embedded in the PL-inactive Cs<sub>4</sub>PbBr<sub>6</sub> crystalline matrix. With the ZnBr<sub>2</sub> post-treated NC solution, strong PL was observed at 518 nm. From the literature,

it is known that the excitonic absorption peak of pure CsPbBr<sub>3</sub> NCs is located at around 510 nm. So, it is reasonable to speculate that the strong excitonic absorption peak and strong PL peak at 518 nm in Fig. 3d originate from the CsPbBr<sub>3</sub> NCs, indicating that a transformation might have taken place from Cs<sub>4</sub>PbBr<sub>6</sub> to CsPbBr<sub>3</sub>.

We also carried out ZnBr<sub>2</sub> concentration dependent emission studies. With the addition of ZnBr<sub>2</sub>, two emission peaks were observed, one at 430 nm and the other at 510 nm, similar to the behavior observed in the absorption spectrum. The emission at 430 nm could be attributed to the 425 nm peak observed in the absorption spectrum and the peak at 510 nm could be attributed to the band edge emission corresponding to the absorption edge shown in Fig. 3e. Similar behavior was also observed in the case of the iodide counterpart, wherein the two emission peaks corresponds to the absorption peak at 550 nm and band edge absorption (Fig. 3f).

Together with the results obtained from the XRD studies and UV-vis and PL spectra, we speculate that CsPbX<sub>3</sub>/Cs<sub>4</sub>PbX<sub>6</sub> composites were formed after the post-treatment with ZnX<sub>2</sub>. In the case of ZnBr<sub>2</sub>, the excitonic CsPbBr<sub>3</sub> absorption was observed at 510 nm and the absorption peak observed at a lower wavelength could be attributed to the Cs<sub>4</sub>PbBr<sub>6</sub> phase. It has been previously reported that defect states created due to the surface halide vacancies could be responsible for quenching the true emission in Cs<sub>4</sub>PbBr<sub>6</sub> and were only observed at low temperatures (4 K). When the Zn<sup>2+</sup> salts were added to the Cs<sub>4</sub>PbBr<sub>6</sub> NCs, the surface halide defect states could be passivated and the true emission from the surface passivated/modified Cs<sub>4</sub>PbBr<sub>6</sub> nanocrystals could be observed at 430 nm. The true emission of Cs<sub>4</sub>PbBr<sub>6</sub> is observed for the first time at





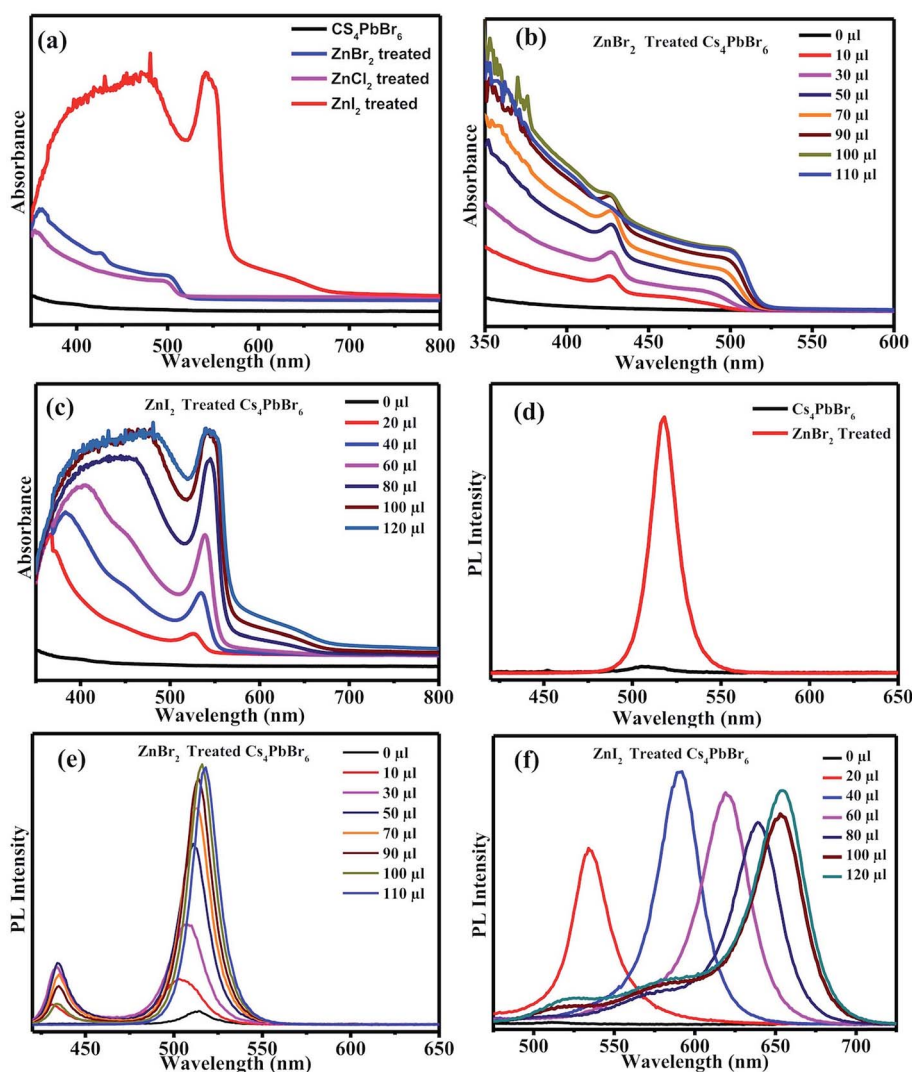


Fig. 3 (a) UV-vis absorption spectra of the  $\text{ZnX}_2$  treated  $\text{Cs}_4\text{PbBr}_6$  nanocrystals. (b) Absorption spectra measured using different concentrations of the  $\text{ZnBr}_2$ /hexane solution. (c) Absorption spectrum of  $\text{ZnI}_2$  treated NCs with the different concentrations of the  $\text{ZnI}_2$ /hexane solution. (d) PL spectra of the pristine NCs and  $\text{ZnBr}_2$  treated NCs. PL spectra of the  $\text{Cs}_4\text{PbBr}_6$  NCs with different concentrations of (e)  $\text{ZnBr}_2$  solution and (f)  $\text{ZnI}_2$  solution.

room temperature when  $\text{Zn}^{2+}$  salts are used to passivate the surface defects; however, the  $\text{Zn}^{2+}$  salts also triggered the transformation of 0D  $\text{Cs}_4\text{PbBr}_6$  NCs into 3D  $\text{CsPbBr}_3$  NCs. So, the intensity of the true emission of  $\text{Cs}_4\text{PbBr}_6$  NCs gradually decreased as the concentration of the zinc salt is increased. When the concentration of the zinc salt was kept constant at 50  $\mu\text{L}$  of  $\text{ZnBr}_2$ , the intensity gradually decreased indicating that the whole process takes place through a two-step mechanism. Similar studies carried out using other salts like  $\text{Cu}^{2+}$  and  $\text{Mn}^{2+}$  didn't result in the enhancement of the PL QY. In the case of  $\text{Mn}^{2+}$  a slight increase was observed but  $\text{Cu}^{2+}$  didn't show any improvement in the PL QY. Fig. S3† shows the PL of the  $\text{CuBr}_2$  and  $\text{MnBr}_2$  treated  $\text{Cs}_4\text{PbBr}_6$  NCs.

TEM images of the as-synthesized  $\text{Cs}_4\text{PbBr}_6$  NCs show hexagonal shaped particles of around 30–35 nm in size similar to the particles reported previously. Fig. 4 shows the TEM images of the as-synthesized  $\text{Cs}_4\text{PbBr}_6$  NCs and the  $\text{ZnBr}_2$  treated samples.

When the samples were treated with 50  $\mu\text{L}$  of  $\text{ZnBr}_2$ , initially hexagonal shaped particles were converted mostly to cubic shaped particles as shown in Fig. 4. More than 90% of the particles became cubic; however the cubic shaped particles did not have sharp edges as observed in the case of the as-prepared  $\text{CsPbBr}_3$  NCs (Fig. S4†). The HR-TEM image (Fig. 4d) shows a lattice spacing of 0.58 nm for the (100) plane of  $\text{CsPbBr}_3$  after  $\text{ZnBr}_2$  treatment. The edges of the particles, however, show that the lattice spacing is not the same as that in the center of the particles. The lattice spacing of 0.33 nm for the (411) plane corresponds to the presence of  $\text{Cs}_4\text{PbBr}_6$ , indicating the formation of a composite of  $\text{CsPbBr}_3/\text{Cs}_4\text{PbBr}_6$ . A plausible two step formation mechanism can be inferred from the observed TEM images. As the  $\text{Cs}_4\text{PbBr}_6$  nanocrystals were treated with the zinc halide salts, in the first step  $\text{Zn}^{2+}$  ions could passivate the surface of the nanocrystals and then result in the extraction of excess  $\text{Cs}^+$  ions, and an increase in the dimensionality from 0D to 3D is



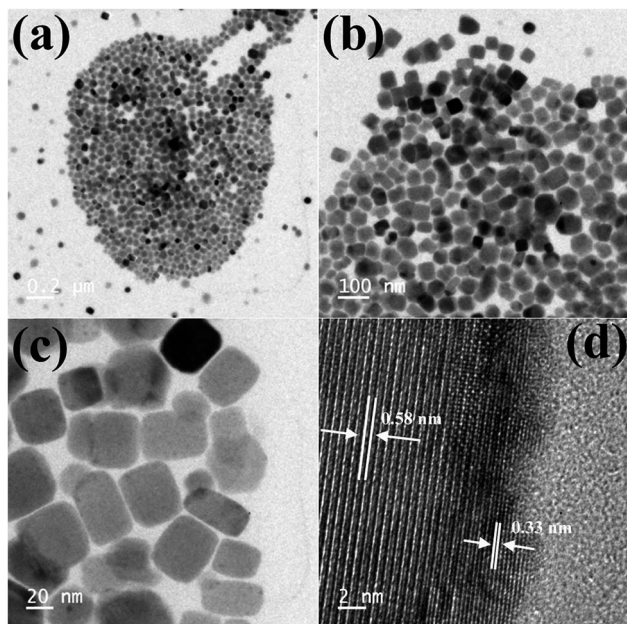


Fig. 4 TEM images of the  $\text{ZnBr}_2$  treated  $\text{Cs}_4\text{PbBr}_6$  NC samples (a–c) and HRTEM (d) image of the  $\text{ZnBr}_2$  treated  $\text{Cs}_4\text{PbBr}_6$  NC samples indicating the formation of a composite of  $\text{CsPbBr}_3/\text{Cs}_4\text{PbBr}_6$  NCs.

observed. In the second step we believe that an inside out diffusion of  $\text{Cs}^+$  ions takes place resulting in the formation of  $\text{CsPbBr}_3/\text{Cs}_4\text{PbBr}_6$  composites.

When the  $\text{Cs}_4\text{PbBr}_6$  NCs were treated with 40  $\mu\text{L}$  of  $\text{ZnI}_2$  solution, the shape of the particles changed from hexagonal to rod shaped. As seen in the TEM images shown in Fig. 5b the corners of the rods still have hexagonal facets. With the increase in the concentration of the  $\text{ZnI}_2$  solution to 100  $\mu\text{L}$  the length of the rods increased to nearly 250 nm and the width increased to 20–30 nm.

To see if any zinc is present on the surface of the nanocrystals, resulting in the enhanced PL QY, X-ray photoelectron spectroscopy (XPS) measurements have been used to assess the surface modification of the NCs and to understand the mechanism of PL enhancement. After measurements, all core level spectra were calibrated using the C 1s peak at 285.0 eV. The high-resolution XPS spectrum of untreated NCs for Pb 4f yields 2 peaks as shown in Fig. S5†. Two intense peaks are observed at 138 eV ( $4f_{7/2}$ ) and 143 eV ( $4f_{5/2}$ ) with a spin-orbit splitting of nearly 5 eV, which corresponds to Pb in the 2+ oxidation state. Quantitative XPS analysis indicates that  $\text{Cs} : \text{Pb} : \text{Br} = 1.5 : 1 : 4.2$  for untreated  $\text{Cs}_4\text{PbBr}_6$  NCs, which is Pb rich in composition. However, for  $\text{ZnBr}_2$ - $\text{Cs}_4\text{PbBr}_6$  NCs, the composition becomes anion-rich ( $\text{Cs} : \text{Pb} : \text{Br} = 0.6 : 1 : 5.5$ ), which was also repeatedly observed (Fig. S6†). A small fraction of the  $\text{Zn}^{2+}$  was also observed in the  $\text{ZnBr}_2$  treated  $\text{Cs}_4\text{PbBr}_6$  NCs, with the ratio of  $\text{Pb} : \text{Zn} = 1 : 0.08$ . This also conforms with the previous observations that the  $\text{Zn}^{2+}$  ions modify the surface of the nanocrystals, which then results in the extraction of excess  $\text{Cs}^+$  ions from the  $\text{Cs}_4\text{PbBr}_6$  NCs. Furthermore, the Pb/Cs ratio increased from 0.66 to 1.66 in  $\text{ZnBr}_2$  treated  $\text{Cs}_4\text{PbBr}_6$  NCs compared with pristine- $\text{Cs}_4\text{PbBr}_6$  NCs indicating that  $\text{ZnBr}_2$  extracted excess  $\text{Cs}^+$  ions present. This is also consistent with the XRD analysis wherein the  $\text{Cs}_4\text{PbBr}_6$  is converted to  $\text{CsPbBr}_3$  after the extraction of cesium ions. We have also carried out XPS analysis of the  $\text{ZnI}_2$  treated  $\text{Cs}_4\text{PbBr}_6$  NCs and found that the ratio of  $\text{Pb} : \text{Zn}$  (1 : 0.15) is much higher compared to that in the bromine case (Fig. S7†).

Owing to the one-dimensional nature of the nanorods in the case of  $\text{ZnI}_2$  treated  $\text{Cs}_4\text{PbBr}_6$  NCs, we have explored their potential application in optoelectronic devices. To this end, we have fabricated photodetectors by dropcasting the  $\text{ZnI}_2$  treated NCs in hexane between two fluorine doped tin oxide (FTO)

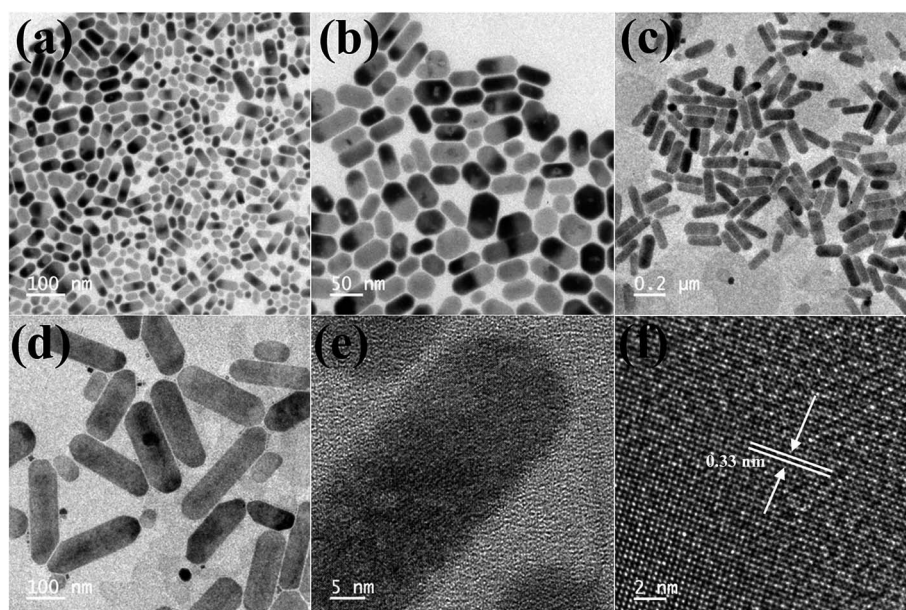


Fig. 5 TEM images obtained after the 40  $\mu\text{L}$   $\text{ZnI}_2$  post treatment (a and b) and 100  $\mu\text{L}$   $\text{ZnI}_2$  post treatment (c–f). HRTEM images showing the lattice spacing after the post treatment with 100  $\mu\text{L}$   $\text{ZnI}_2$  (e and f).



electrodes. Fig. 6a shows the schematic of the device architecture that is employed for the fabrication of the devices. The dark current observed was in the order of a few picoamperes, within the detection limit of our instrument. The current of the same device increased under illumination using a diode source of 617 nm, demonstrating their potential as an efficient photo-detector. Fig. 6b shows the IV curves obtained in the dark and in the presence of the light diodes. The ratio of the photocurrent to dark current could reach over two orders of magnitude. Time-dependent photocurrent measurements were carried out by pulsing the diode at a fixed light intensity with an applied bias voltage of 2 V. Prompt and reproducible photocurrent response to the pulsing was observed indicating the excellent optical switching and stability of the photodetectors. The rise time and decay time were also determined from the pulsing experiments and they are in the order of a few milliseconds as shown in Fig. 6d. When the photodetectors were fabricated using  $\text{ZnBr}_2$  treated nanocrystals, the currents obtained were lower compared to those using the  $\text{ZnI}_2$  treated samples. Fig. S8† shows the performance of the photodetectors using  $\text{ZnBr}_2$  treated NCs in the same configuration. This indicates that the performance of the photodetectors could be improved if the length of the nanorods could be made longer than the width of the channel used and also by removing the surface capping ligands by appropriate treatments.

In conclusion, we have developed a simple and convenient post synthetic strategy for the transformation of 0D  $\text{Cs}_4\text{PbBr}_6$  into highly luminescent  $\text{CsPbBr}_3/\text{Cs}_4\text{PbBr}_6$  composites using  $\text{ZnBr}_2$  treatment. The transformation with  $\text{ZnBr}_2$  treatment proceeds through a two-step process, where in the first step, zinc doped/modified  $\text{Cs}_4\text{PbBr}_6$  NCs are formed which then results in the conversion to  $\text{CsPbBr}_3/\text{Cs}_4\text{PbBr}_6$  nanocomposites. The composite of  $\text{CsPbBr}_3/\text{Cs}_4\text{PbBr}_6$  NCs is found to be very

highly luminescent with a PL QY of 90%. In the case of  $\text{ZnI}_2$  treatment, the post synthetic treatment also resulted in morphological transformation from the hexagonal shaped nanocrystals to nanorods. The length of the rods could be varied depending on the concentration of the nanocrystals and the zinc iodide salt added. In both the iodide and bromide case an intermediate is formed upon treatment with the zinc salt. The intermediate was observed in the time/concentration dependent UV-vis and PL studies, and from the X-ray diffraction analysis, we presume that it is the 0D structure modified/doped by the zinc cations. Photodetectors based on the  $\text{ZnI}_2$  treated  $\text{Cs}_4\text{PbBr}_6$  NCs were fabricated, which exhibit a high on/off ratio with a fast response time. The excellent optoelectronic properties make this treatment versatile for a wide range of functional optoelectronic devices like light emitting diodes and photovoltaic devices.

## Experimental section

### Chemicals

Cesium carbonate ( $\text{Cs}_2\text{CO}_3$ , 99.9%), lead(II) bromide (98%), zinc(II) iodide (98%), zinc(II) chloride (99%), oleylamine (OLA, 70%), oleic acid (90%), and 1-octadecene (90%) were purchased from Sigma Aldrich. Zinc(II) bromide (98%) was purchased from Alfa Aesar.

### Synthesis of cesium oleate

Cesium carbonate ( $\text{Cs}_2\text{CO}_3$ , 160 mg), oleic acid (1 mL), and octadecene (16 mL) were added to a three-neck flat bottom flask and kept under vacuum for 30 min at 120 °C. After 30 minutes the temperature was increased to 150 °C and the system was

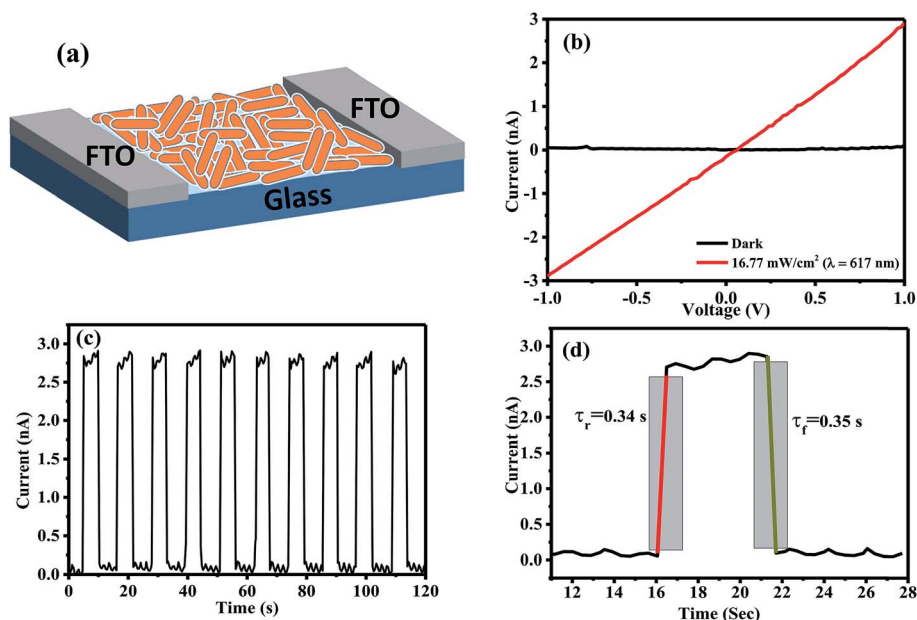


Fig. 6 (a) Schematic of the photo-detector architecture. (b)  $I$ - $V$  curves in the dark and in the presence of 617 nm light. (c) Photocurrent time characteristics of the photo detector in the dark and under light (617 nm, 1 volt). (d) Rise and decay time.





kept under a N<sub>2</sub> atmosphere until a clear solution of cesium oleate was obtained.

### Synthesis of Cs<sub>4</sub>PbBr<sub>6</sub> nanoparticles

Octadecene (10 mL), oleylamine (1 mL), oleic acid (1 mL) and lead bromide (0.36 mmol, 73.4 mg) were loaded into a three-neck round bottom flask and degassed for 30 min at 120 °C. The flask was then filled with N<sub>2</sub> and the temperature was increased to 140 °C. After that, 4.4 mL of hot cesium oleate solution was injected quickly. After 5 s, the reaction mixture was quenched in an ice-water bath.

### Synthesis of CsPbBr<sub>3</sub> nanoparticles

Lead bromide (207 mg) and octadecene (15 mL) were loaded into a three-neck round bottom flask and degassed for 1 hour at 120 °C. The flask was then filled with N<sub>2</sub> and oleylamine (1.5 mL) and oleic acid (1.5 mL) were added. After complete solubilisation of PbBr<sub>2</sub>, the temperature was increased to 165 °C and finally the as-prepared Cs-oleate (1.2 mL) was injected. The reaction was immediately quenched in an ice-water bath.

### Purification of nano particles

The crude solution of nanoparticles was first centrifuged at 6000 rpm for 10 min. After that, the supernatant was discarded and the precipitate was redispersed in hexane solution. Then again, the large particles present in the solution were removed by centrifugation at 6000 rpm for 10 min. The supernatant was discarded and a colloidal solution of nanoparticles was obtained in hexane.

### ZnX<sub>2</sub> (X = Br, I and Cl) treatment

The NCs dispersed in hexane (~8 mg mL<sup>-1</sup>) were treated with the pre-made zinc halide solution. Zinc halide solutions were prepared by dissolving zinc halides (ZnX<sub>2</sub>, X = I, Br, Cl) (0.05 M) in the mixture of hexane (2 mL) and oleylamine (40 μL) at room temperature. Then the required zinc halide solution was added to 1 mL of the Cs<sub>4</sub>PbBr<sub>6</sub> NCs in hexane solution (8 mg mL<sup>-1</sup>) under constant stirring. Excess zinc halide salts are removed by centrifugation and the NCs can be stored in organic solvents like hexane, toluene, *etc.* for a few months.

### Characterization

High resolution transmission electron microscope images were obtained using a Tecnai G2 F30 300 kV transmission electron microscope. X-ray diffraction spectra were obtained by using a Rigaku Japan SmartLab X-ray diffractometer (Cu Kα, λ = 1.541). PL spectra were collected by using a Horiba FluoroMax spectrometer with 400 nm excitation wavelength, while UV-vis absorption measurement was performed using a UV-vis-NIR spectrometer (PerkinElmer LAMBDA 950). The room-temperature PLQY was calculated according to the following equation<sup>38</sup>

$$QY_{\text{sample}} = QY_{\text{ref}} \left( \frac{\text{PLArea}_{\text{sample}}}{\text{PLArea}_{\text{ref}}} \right) \left( \frac{I_{\text{ref}}}{I_{\text{sample}}} \right) \left( \frac{n_{\text{sample}}^2}{n_{\text{ref}}^2} \right)$$

In this equation, QY<sub>sample</sub> is the PLQY of the sample. PLArea<sub>sample</sub> is the integrated emission spectrum curve area for the sample. I<sub>sample</sub> is the absorption value at the excitation wavelength for the sample. n<sub>sample</sub> is the refractive index of the employed solvent. QY<sub>ref</sub>, PLArea<sub>ref</sub>, I<sub>ref</sub> and n<sub>ref</sub> are those for the standard reference respectively. In our experiment, fluorescein sodium salt in water and Rhodamine 6G in water (PLQY = 0.95) were chosen to be the standards for green and red spectral regions and the standard solutions should be freshly prepared. Attention was paid to keeping the optical densities of the sample and standard below 0.1 at the excitation wavelength.

### Photodetector fabrication

Laser patterned FTO coated glass with an etched area was used as the photodetector. Before deposition, the laser patterned FTO was cleaned thoroughly by using a soap solution, DI water, acetone, and ethanol for 10 minutes, respectively, by ultrasonication. Perovskite nanocrystals in hexane were drop cast on the etched area of the FTO coated glass by masking the unetched area with scotch tape. The drop cast substrate was then subjected to vacuum drying. Then the photoresponse was measured under a 617 nm LED light.

## Conflicts of interest

There are no conflicts to declare.

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