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Improvement in optical properties of Cs₄PbBr₆ nanocrystals using aprotic polar purification solvent

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We demonstrate the influence mechanism on the optical property of Cs₄PbBr₆ during purification of solution with different protonated levels and polarities. During the purification process, organic groups originating from oleic acid (OA) and PbBr impurity on the surface of Cs₄PbBr₆ nanocrystals can be removed using high polarity aprotic and protonic solvents, and the number of Br vacancies (V_{Br}) can be reduced. The protonic polar solvent can not only etch the organic groups on the surface of nanocrystals, causing surface reconstruction and particle growth of nanocrystals, but also enter into the lattice of Cs₄PbBr₆ and react with the embedded CsPbBr₃. However, aprotic polar solvent decreases the particle size of Cs₄PbBr₆ nanocrystals with the increase in the solvent polarity. The optical properties of Cs₄PbBr₆ can be effectively improved using aprotic polar solvents as a purification solvent, which is very significant to improve the luminescence efficiency of perovskites.

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Introduction

Inorganic perovskites have attracted extensive attention and have become one of the research hotspots in the photovoltaic field since their genesis, owing to excellent photoelectric characteristics, such as high extinction coefficient, long charge carrier diffusion lengths, precisely tunable bandgaps, high absorption coefficients, and high defect tolerance.^{1–7} It is well known, CsPbX₃ quantum dots (QDs) feature a fragile and unstable structure composed of [PbX₆]^{4–} octahedron with Cs⁺ cations filling. This structure has poor tolerance under moisture and heat because of the easy losing of halide anions. Diverse strategies have been used to overcome the above concerns, such as ligand passivation route⁸ and core-shelling method.^{9,10}

CsPbX₃ is very sensitive to polar solutions. For example, CsPbX₃ is easy to hydrolyze in water (high polarity) but can be preserved in *n*-hexane (low polarity). Anirban Dutta *et al.*¹¹ found that CsPbBr₃ tuned the phase upon changing the polarity of the reaction solvent from chloroform to hexane, while CsPbCl₃ retained the favorable cubic phase, but their shape was tuned from a cube to the platelet. In the process, solvent-polarity-induced precursor activation led to growth controls and different shapes and phase-controlled perovskite nanocrystals. Wu *et al.*¹² explored the effect of solvent polarity on the

morphology and crystallization of perovskite films during the process of synthesis and found that the root mean square (RMS) roughness decreases, while crystallization increases with the enhancement of alcohol solvent polarity. Li *et al.*¹³ found that the size of perovskite QDs reduced with the increase of solvent polarity. Recently, some studies reported that Cs₄PbX₆, a derived member of Cs–Pb–X compounds with large band gaps,¹⁴ can be synthesized by adding excessive cesium ions^{15–17} and has better stability than CsPbX₃. Zhang *et al.*¹⁸ found that the synthesized Cs₄PbX₆ phase 0-D nanocrystals could maintain good stability in some polar solvents (such as DMF) and air due to their high exciton binding energy (171 MeV). Chen *et al.*¹⁹ found that Cs₄PbBr₆ has excellent thermal stability, and the humidity resistance experiment shows that it still has good stability in the humidity environment. However, few scholars have studied the influence mechanism of different polar solutions on the optical performance of Cs₄PbBr₆ nanocrystals during the process of purification. Considering the polarity of the solution, we explore the influence on the optical properties of perovskite during the process of purification treatment from the perspective of polarity.

Here, we used common different polar reagents (*n*-hexane, ethyl acetate, acetone, and absolute ethanol) to purify Cs₄PbBr₆ nanocrystals. It was found that there were great changes in luminescent properties when the solvents with different protonated levels and polarities were used for purification. Then, we explored the influence mechanism of different protonated levels polar solutions on the optical properties of perovskites by analyzing the microstructure, morphology, defects, band gap, and so on. This work provides a reference for

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the purification of perovskites and lays a foundation for the application of this kind of material in the field of lighting.

Experimental section

Materials

Cesium bromide (CsBr, 99.5% metals basis), lead(II) bromide (PbBr₂, AR, 99.0%), oleylamine (OM, 80–90%) and dimethyl sulfoxide (DMSO) were purchased from Aladdin®. Ethyl acetate (EA), *n*-hexane (*n*-H), oleic acid (OA, AR), absolute ethanol and acetone were purchased from Sinopharm Chemical Reagent. All reagents were used as received without any further purification.

Synthesis of Cs₄PbBr₆ nanocrystals

Cs₄PbBr₆ nanocrystals were synthesized at room temperature without any protective gas. Briefly, 25 ml acetone was loaded into a 150 ml conical flask at 25 °C. Subsequently, 5 ml OA and 2.5 ml OM were added to the conical flask with stirring for 1 min to make it homogenized, and kept overnight at room temperature. After that, solid powders of 1 mmol PbBr₂ (0.367 g dissolved in 4 ml DMSO) and 1 mmol CsBr (0.213 g dissolved in deionized 1 ml water) were added to the flask with vigorous stirring (1200 rpm) for 10 min. Through ultraviolet irradiation, we could observe strong green fluorescence. As such, Cs₄PbBr₆ was synthesized successfully.

The purification process of Cs₄PbBr₆ nanocrystals

The crude solution was divided into four parts and centrifuged at 7000 rpm for 5 min. Nanocrystals were dispersed in 10 ml *n*-H, 10 ml EA, 10 ml acetone and 10 ml absolute ethanol individually, followed by ultrasonic cleaning for 5 min, and centrifuged at 7000 rpm for 5 min. Finally, they were dried in a vacuum oven for 12 h at room temperature, and the powder was prepared for characterization.

Characterization

X-ray diffractometry (XRD) patterns were collected with an X-ray automatic diffractometer (Advance D8, BRUGG GROUP, Switzerland) using Cu K α radiation (wavelength 1.55406 Å). XRD patterns were recorded in the range of $10^\circ \leq 2\theta \leq 60^\circ$ with a scanning step of 0.02° and counting time 2 s per step. Transmission electron microscopy (TEM) was performed on a field emission scanning electron microscope (JEM-2100F, Japan). Photoluminescence (PL), photoluminescence excitation (PLE) spectra and PL decay spectra were acquired on a full-function fluorescence spectrometer (FLS-1000, Edinburgh, Britain) with a time range: 0–199.6 μ s; step: 0.40 μ s; frequency: 100.0 Hz. Ultraviolet and visible (UV-vis) absorption spectra were measured with a UV-3600 Ultraviolet-visible spectrophotometer (Shimadzu, Kyoto, Japan). X-ray photoelectron spectroscopy (XPS) was recorded with a K-ALPHA X-ray photoelectron spectrometer (Thermo Fisher Scientific, Britain) in the 1300–0 eV region. Fourier transform infrared reflection (FTIR) spectra were measured on a Nicolet instrument (iS5, Thermo, United States) in the region of 3800–400 cm^{−1}. Raman spectra were measured on a Lab RAM HR800 spectrometer with

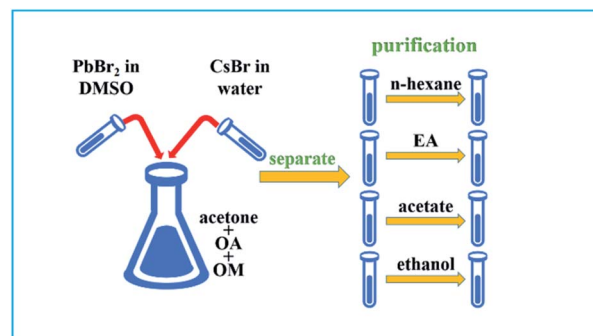
a 633 nm laser light at room temperature and recorded in the region of 50–500 cm^{−1}.

Results and discussion

After synthesis, the as-synthesized Cs₄PbBr₆ nanocrystal solution was divided into four equal parts and separated from the mother liquor by centrifugation. Then, the obtained Cs₄PbBr₆ nanocrystals were dispersed in different polar solutions (*n*-hexane, ethyl acetate, acetone and ethanol) for further purification treatment. Sample A was dispersed in *n*-H after centrifuging, while Sample B, C and D were dispersed in ethyl acetate, acetone and ethanol, respectively, and all samples were centrifuged again (solvent polarity: A < B < C < D) (see Scheme 1). The order of polarity is in accordance with previous report.²⁰

The precipitates of these samples were dried in a vacuum oven at room temperature. A significant color change of the nanocrystal powder occurred after purification, which indicated that the optical properties changed dramatically, as shown in Fig. 1. In our studies, we found that the optical photos of Cs₄PbBr₆ dispersed in toluene (aprotic inert solvent) were obviously different, as shown in Fig. 1a and b. Under daylight, the purified samples showed different degrees of yellow and the sample treated with acetone was even light green among these samples. PLE and PL (excitation: 360 nm) spectra obtained from powder are also shown in Fig. 1c and d. It can be seen that all PLE spectra consisted of an excitation peak at about 285 nm and a broad excitation band (320–500 nm), and the intensity first increased and then decreased with the increase of solvent polarity, as shown in Fig. 1c. We can also find, with the increase in the polarity, the PL peak position shifted from 512 nm (A) to 515 nm (B), 515 nm (C), 513 nm (D), and the PL intensity showed the same trend with PLE and PLQY as shown in Fig. 1e and f. These results suggested that the solvent polarity has a stronger influence on the PL peak position and PLQY optical properties of Cs₄PbBr₆ nanocrystals during the purification process. Considering the polarity of the solution, we explored the influence mechanism on the optical properties of perovskite Cs₄PbBr₆ nanocrystals during the process of solution treatment from the perspective of polarity.

To confirm the influence mechanism, we explored its principle through the characterization of the microstructure. The



Scheme 1 The synthesis and purification of Cs₄PbBr₆ nanocrystal.



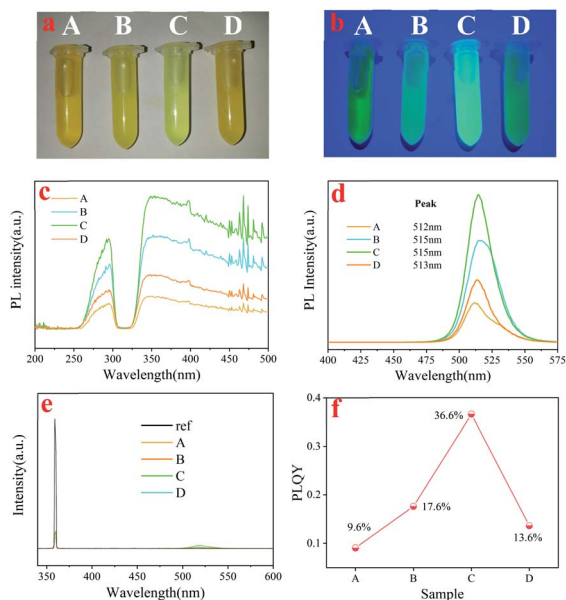


Fig. 1 Images of Cs_4PbBr_6 in toluene (10 mg mL^{-1}), washed by *n*-H (A), EA (B), acetone (C) and ethanol (D), respectively, under ambient light (a), and under UV lamp (b); the photoluminescence excitation (c) and the photoluminescence emission (d) spectra of sample A, B, C and D; (e and f) the PLQY of four samples.

TEM images shown in Fig. 2 demonstrate that Cs_4PbBr_6 nanocrystals show different sizes and morphologies after purifying with different polar solutions. It can be clearly seen that when the polarity is relatively low, the shape of nanocrystals is generally rectangular, and crystal dimension approximately equal to 10–20 nm, as shown in Fig. 2a. With the increase in polarity, the shape of nanocrystals changes accompanied by the appearance of a triangle, sphere, polygon and so on, and the size also decreased (5–10 nm), as shown in Fig. 2c. As the polarity continues to increase, nanocrystals turn into small spheres, and the size is also up to the quantum size (less than 5 nm), as shown in Fig. 2e. Then, as the polarity increases further, the morphology becomes is a regular rectangle, and the size increases sharply to more than 20 nm, as shown in Fig. 2g. However, the data of Hu *et al.* show that with the increase in the polarity of the solution, the particles gradually become smaller and uniform, and the phase of Cs_4PbBr_6 gradually transform into CsPbBr_3 . The size of nanocrystals in lower polarity solvents increases, compared to those with moderate polarity.¹¹ When sample D was magnified, we could clearly see that the square phase was formed by spherical particles agglomerating together with the organic matter, as shown in the inset in Fig. 2g, which is not found by the previous research institute.

HRTEM was further carried out to measure the lattice spacing of the products, as shown in Fig. 2b, d, f and h. The lattice fringes (sample A, B, and C) are clearly observed with an interplanar spacing of 0.195 nm, corresponding to the (600) plane of hexagonal Cs_4PbBr_6 nanocrystals ($d_{600} = 0.195 \text{ nm}$, JCPDS no.73-2478), while the interplanar spacing in sample D is 0.295 nm, corresponding the (223) plane of hexagonal Cs_4PbBr_6 nanocrystals ($d_{223} = 0.295 \text{ nm}$, JCPDS no.73-2478).

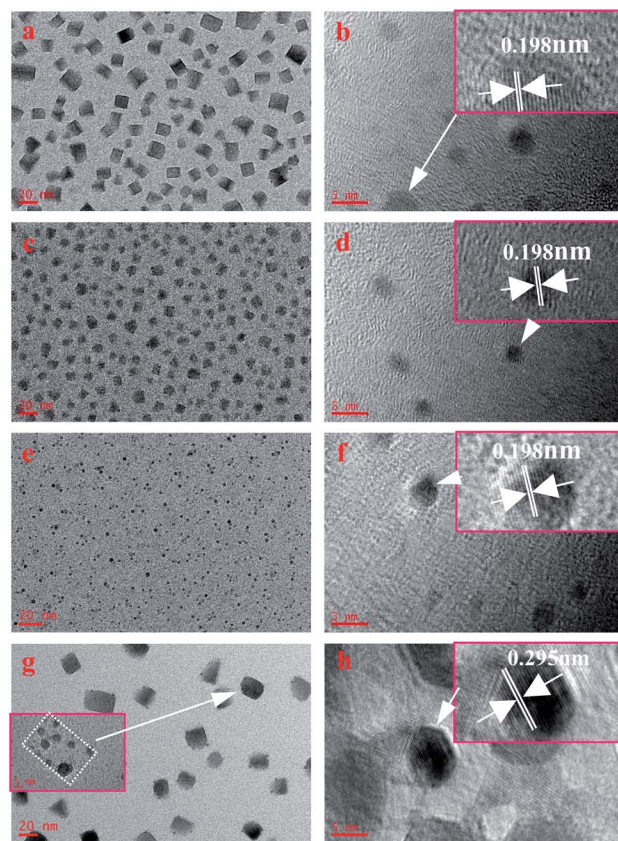


Fig. 2 TEM images of samples prepared with different polarity solvent: (a) A; (c) B; (e) C; (g) D. HRTEM images of samples prepared with different polarity solvent: (b) A; (d) B; (f) C; (h) D.

From the results of TEM and HRTEM, we found an interesting phenomenon, compared to the (600) dominant growth plane of the former three samples, the plane of sample D changes to the (223) plane. At the same time, with the increase in polarity, the particle size of the former three samples decreased, but in the last sample, it increased significantly. This phenomenon suggests that the interaction between ethanol and

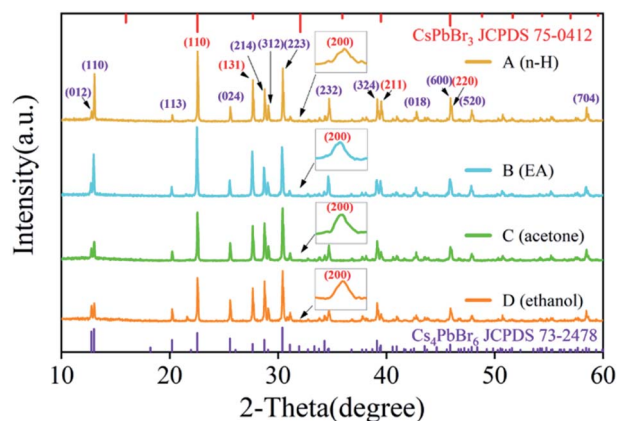


Fig. 3 XRD patterns of sample A, sample B, sample C and sample D. the inset is the peak at 31.9 degrees.

Table 1 The lattice parameter and lattice strain of Cs₄PbBr₆

| Sample | A | B | C | D |
|---|---------|---------|---------|---------|
| Lattice strain (10^{-4}) | 7.4153 | 9.4920 | 12.2000 | 7.9509 |
| Lattice parameter <i>a</i> and <i>b</i> (Å) | 13.7038 | 13.7286 | 13.7080 | 13.7050 |
| Lattice parameter <i>c</i> (Å) | 17.2870 | 17.3199 | 17.2851 | 17.2868 |

Cs₄PbBr₆ nanocrystals was different from that in *n*-hexane, ethyl acetate and acetone during the purification process. It is worth noting that the solvents *n*-hexane (inert solvent), acetone and ethyl (aprotic polar solvent) acetate, ethanol (proton polar solvent) are different. During the purification process, protonated ethanol can etch organic groups on the surface of nanocrystals, improve surface energy, and cause surface reconstruction and particle growth of nanocrystals, which is well agreeing with the result shown in the inset in Fig. 2g. This is why the dominant growth plane of perovskite Cs₄PbBr₆ nanocrystals changes and the particle size increases when purified by ethanol. As ethyl acetate and acetone are aprotic polar solvents, the etching effect on the nanocrystalline surface is very weak so that they cannot cause surface reconstruction. However, the solvent effect is improved due to polarity, and finally decreases the size of Cs₄PbBr₆ nanocrystals. Additionally, the size of sample B and sample C are smaller than those previously reported in the literature²¹ and can produce a quantum size effect, which can strongly influence the tuneable PL properties.²²

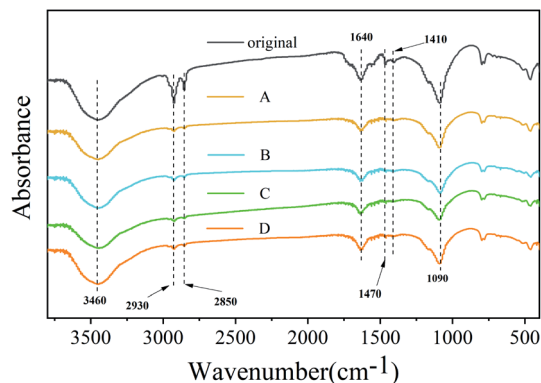
Fig. 3 shows X-ray diffraction (XRD) patterns of the above four samples. It can be seen that with the polarity increasing, the structure of Cs₄PbBr₆ nanocrystals does not change, which matched well with the standard diffraction peaks shown in JCPDS no. 73-2478. The illustration reports a detailed investigation of the peak at 31.9 degrees, which are matching well with the standard diffraction peaks shown in JCPDS no. 75-0412. These results confirmed the coexistence of Cs₄PbBr₆ (JCPDS no. 73-2478) and a tiny bit of CsPbBr₃ (JCPDS no. 75-0412), which is consistent with previous studies.^{23–25} These results also indicated that CsPbBr₃ are embedded in Cs₄PbBr₆ nanocrystals and are not easily removed by polar solvents during purification. We calculated the lattice parameter and lattice strain by Williamson–Hall and Scherrer's formulas, respectively, as shown in Table 1. It can be seen that the trends of the lattice parameter and the lattice strain are first increasing and then decreasing with the increase of the solution polarity. At the same time, the lattice strain can make PL red shift and the change of lattice strain is consistent with the change of PL peak shift, which indicates that the quantum size effect is not dominant in the process, and the change of PL peak shift is mainly controlled by the lattice strain.

Note that, being similar to the phenomenon observed in previous studies,²⁶ Cs₄PbBr₆ is easily damaged under irradiation of focused electron beam, which may due to its low lattice energy. However, it can be seen that the phase transition in nanocrystals does not occur after purification with different polar solutions, which indicates that Cs₄PbBr₆ nanocrystals

prepared by us have better phase transition stability, suggesting that the obtained Cs₄PbBr₆ nanocrystals are a good candidate material for light-emitting applications.

Previous reports showed that purification could etch the surface of nanocrystals with the removal of the ligand.²⁷ Two organic surfactants were used during the synthesis; oleylamine enhanced the solubility of lead salts, whereas oleic acid acted as a hydrophobic ligand. Using FTIR, the presence of surface ligands was analyzed (Fig. 4). The peaks located at 1470 cm^{−1} (−COO− stretch) mainly originated from the groups in OA molecules. The peaks at 2930 cm^{−1} (−CH₂− stretch), 2850 cm^{−1} (−CH₂− stretch) and 1410 cm^{−1} (−CH₃ bending), which belong to OM and OA, are significantly weakened or even disappeared after purification using different polar solutions, suggesting that these groups can be eliminated easily by purification. The peaks appearing at 1090 cm^{−1} (−C−N stretch) and 1640 cm^{−1} (C=O or N−H stretch) originated from the groups in OM molecules, and their intensity change range is relatively small.²⁷ It is worth noting that the OH stretch caused by water is also apparent at about 3450 cm^{−1}, and its strength has hardly changed after purification with different polar solutions. The reason is that for the special structure of Cs₄PbBr₆, some small molecules can easily enter the lattice, which can influence the structure and bandgap, leading to the change of optical properties. Wang *et al.* found that hydroxyl has a deep influence on the structure and bandgap of Cs₄PbBr₆ to form a sub-bandgap of about 2.6 eV.²⁸ Pure Cs₄PbBr₆ nanocrystals have no photoluminescence efficiency in visible light emission. A very small amount of water, which is a proton polar solvent, can enter the lattice of Cs₄PbBr₆ nanocrystals and cause some Cs₄PbBr₆ nanocrystals to change into CsPbBr₃.

Fig. 5 shows the HRTEM of CsPbBr₃ embedded in Cs₄PbBr₆. The fast Fourier transformation (FFT) images inset at the bottom of the figure confirm that the cubic CsPbBr₃ and hexagonal Cs₄PbBr₆ exist together. The *d*-spacing for two red fringes was measured to be 2.9 Å, which belongs to (233) of hexagonal Cs₄PbBr₆ (JCPDS 74-2378). The *d*-spacing for brown fringes was measured to be 5.6 Å, belonging to (100) of cubic CsPbBr₃ (JCPDS 75-0412). It can be clearly seen that CsPbBr₃ is embedded in Cs₄PbBr₆, which indicates that the small molecules of water enter into the Cs₄PbBr₆ lattice and react to form

**Fig. 4** FTIR spectra of sample original, A, B, C and D.

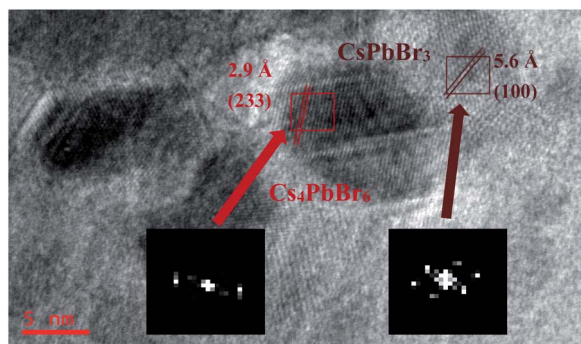


Fig. 5 The HRTEM of CsPbBr₃ embedded in Cs₄PbBr₆ in sample D.

CsPbBr₃. The existence of the -OH peak at 3460 cm^{-1} in FTIR spectra also proved the existence of small molecules of water, indirectly. The origin of Cs₄PbBr₆ green emission has always been controversial. As shown by Fig. 5, the mosaic structure between Cs₄PbBr₆ and CsPbBr₃ is more likely to be the cause of Cs₄PbBr₆ green emission, and lattice matching between the Cs₄PbBr₆ and CsPbBr₃ contributes to improved luminescent properties,²⁹ which show that the green emission of Cs₄PbBr₆ comes from the inside CsPbBr₃.

X-ray photoelectron spectroscopy (XPS) analysis was performed to investigate the surface chemical elements and valence states of Cs₄PbBr₆, as shown in Fig. 6. The full survey scan confirms the existence of Cs, Pb, O, and Br in Cs₄PbBr₆ (Fig. 6a). O may have contributed to -COO- , C=O or -OH groups in OA, OM, or water. High-resolution XPS spectra for the core level of Cs 3d, Pb 4f are shown in Fig. 6b and c. As shown in Fig. 6b, peaks at 723.9 and 737.9 eV can be attributed to Cs 3d_{5/2} and Cs 3d_{3/2}. There was no change in the position of the peak and the distance between the two peaks, indicating the chemical valence state of Cs did not change. The Pb 4f XPS spectrum also showed two peaks at 138.3 eV and 143.1 eV (Fig. 6c), which are assigned to Pb 4f_{7/2} and 4f_{5/2}. It can be seen that the full width at half maxima (FWHM) of the two peaks is first reduced and then increased with the increase of solvent polarity. The change of FWHM indicates that there are other lead-containing impurity phases on the surface of as-synthesized perovskite Cs₄PbBr₆ nanocrystals, which can be removed by high polar solvents. Pb 4f_{7/2} and Pb 4f_{5/2} peaks of Cs₄PbBr₆ in sample A can be fitted into two peaks with binding energies of 138.38 eV and 143.08 eV, 139.48 eV and 144.28 eV, respectively, suggesting that Pb in Cs₄PbBr₆ nanocrystals exists in two chemical environments. The spin-orbit splitting energy is 4.7 eV and 4.8 eV, referring to the Web database (<http://srdata.nist.gov/xps/>), which can be assigned to Pb²⁺ ion in PbBr₂ and Pb¹⁺ ion in PbBr, respectively, as shown in Fig. 6d, implying that there are many V_{Br} defects existing in sample A. From Fig. 6c, we found that the signal of Pb 4f in PbBr significantly reduced after purification using high polar solvents, suggesting that the V_{Br} defect reduced. The core-level Br 3d XPS scan was fitted into two peaks at the binding energies of 67.9 eV and 68.7 eV, as shown in Fig. 6e–h, which is very similar to findings from the previous studies.³⁰ The observed peak at about 67.9 eV can be attributed

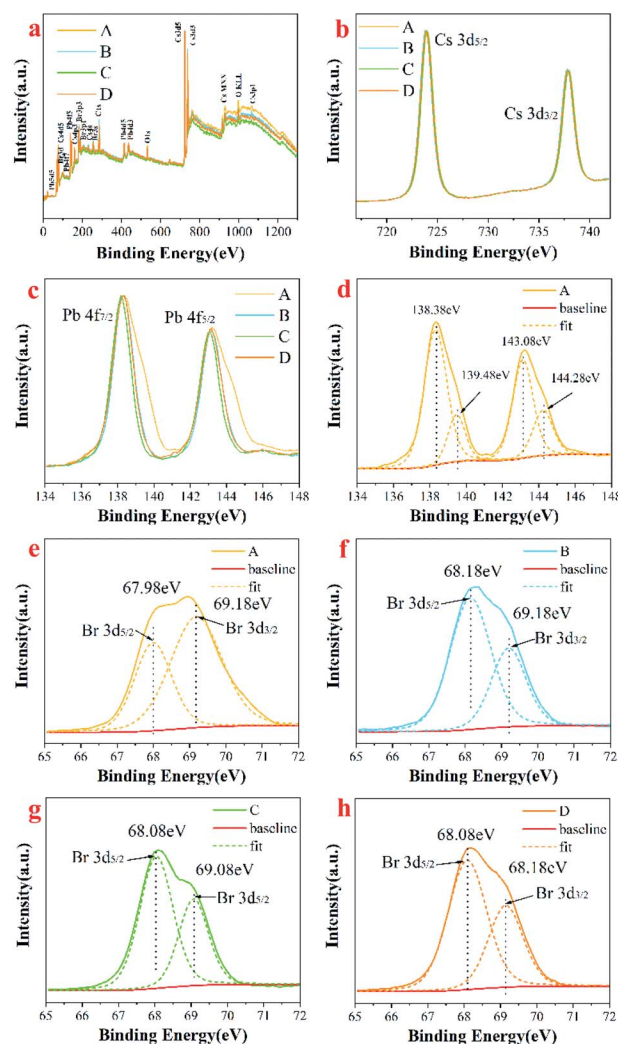


Fig. 6 Full survey scan of Cs₄PbBr₆ nanocrystals (a); high resolution XPS of (b) Cs 3d; (c) Pb 4f; the fitted high resolution XPS of (d) Pb 4f, (e–h) Br 3d.

to the Br atoms in isolated $[\text{PbBr}_6]^{4-}$ of Cs₄PbBr₆,²³ and redshifts first and then blueshifts with the increase in polarity. The peak at 68.7 eV correlates with the existence of Br atoms of CsBr impurity or corner-sharing $[\text{PbBr}_6]^{4-}$ of CsPbBr₃.²³ The results suggest that the CsPbBr₃ phase coexist in the synthesized Cs₄PbBr₆ phase. It can be seen that the intensity of peak at 68.7 eV increases first and is then reduced with the increase in the polarity of the solvent, indicating that the amount of CsBr impurity or corner-sharing $[\text{PbBr}_6]^{4-}$ of CsPbBr₃ reduces during purification with a high polarity solvent. In addition, we have noticed another strong peak at 75 eV, which can be attributed to Cs 4d₅ in CsBr.

Raman spectra of the Cs₄PbBr₆ nanocrystals provided the vibrational modes of the metal-halide sublattice, as shown in Fig. 7. Two Raman-active modes for Cs₄PbBr₆ nanocrystals were observed, with a peak at 86 cm^{-1} and a broad peak at 121 cm^{-1} . The peak at 86 cm^{-1} arises from the vibrational mode of the isolated $[\text{PbBr}_6]^{4-}$ octahedron in Cs₄PbBr₆ nanocrystals.³¹ The

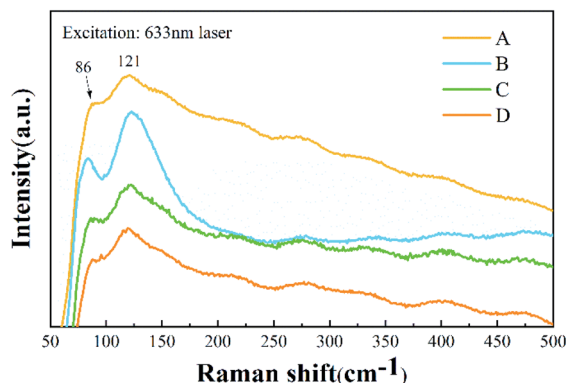


Fig. 7 Raman spectra of samples A, B, C and D.

broad peak comes from the Cs^+ cation surrounding the $[\text{PbBr}_6]^{4-}$ octahedron. Except for a little broadening, the Raman spectrum is similar to that previously reported for Cs_4PbBr_6 nanocrystals.³² According to a previous Raman study of CsPbBr_3 (ref. 16) and CsPbCl_3 crystals with a $Pnma$ phase,³³ the peaks assigned to the vibrational mode of $[\text{PbBr}_6]^{4-}$ octahedron and motion of Cs^+ cation are located at 72 cm^{-1} and 127 cm^{-1} , respectively. It can be seen that the two peaks that originated from CsPbBr_3 are submerged in the peaks of Cs_4PbBr_6 nanocrystals, which may be due to the small amount of CsPbBr_3 in Cs_4PbBr_6 nanocrystals, being consistent with the results of TEM and XRD.

As is known, pure Cs_4PbBr_6 is an insulator with a large band gap of $\sim 3.9\text{ eV}$,^{14,34} and it itself does not show green

emission,^{19,35–38} whereas CsPbBr_3 is a semiconductor with a band gap of about 2.4 eV ,³⁹ showing a strong green emission. In order to study the influential mechanism of purification on luminescence properties of the as-synthesized Cs_4PbBr_6 nanoparticles, a diffuse reflectance absorption spectrum (DRS) was recorded and shown in Fig. 8a, and the corresponding Tauc plot was also measured as shown in Fig. 8b. It can be seen that the DRS was composed of a sharp absorption peak at about 315 nm and a broad absorption band ($320\text{--}550\text{ nm}$), while the Tauc plot showed two absorption band edges at about 2.3 eV and 3.7 eV , assigned as E_{g1} and E_{g2} , respectively. The strong absorption peak around 315 nm is related to Cs_4PbBr_6 , originating from the optical transitions between localized states within isolated PbBr_6^{4-} octahedra.^{14,34} It has been reported that the absorption peak of pure CsPbBr_3 is located at around 505 nm .⁴⁰ Notably, there is a sharp dip at $\sim 315\text{ nm}$ in the PLE spectrum (Fig. 1c) and a sharp peak in the absorption spectrum at the same wavelength. This is due to the strong absorption of Cs_4PbBr_6 at 315 nm ,¹⁴ and the absorption of CsPbBr_3 is comparably weak, which leads to a dip in the PL excitation spectrum. For comparison, band gaps (E_{g1}) of samples were determined and found to be approximately 2.31 eV (A), 2.33 eV (B), 2.32 eV (C) and 2.29 eV (D), as shown in Fig. 8c. Band gaps (E_{g2}) of samples were also determined and found to be about 3.69 eV (A), 3.7 eV (B), 3.7 eV (C) and 3.7 eV (D) (Fig. 8d). It can be seen that band gaps (E_{g1}) and (E_{g2}) are obviously smaller than those of Cs_4PbBr_6 nanoparticles. There are three factors that affect the change in the band gap. The first is that a small amount of CsPbBr_3 is embedded in Cs_4PbBr_6 nanocrystals, which leads to lattice distortion and eventual changes in the band gap of Cs_4PbBr_6 nanocrystals. The formation of CsPbBr_3 is triggered by water, which enters the lattice of Cs_4PbBr_6 nanocrystals according to the following transformation process during the synthesis: $\text{Cs}_4\text{PbBr}_6 \rightarrow \text{CsPbBr}_3 + \text{CsBr}$. Due to the high solubility of CsBr in water, water drives the reaction to the right. This phenomenon is consistent with the results of XRD and

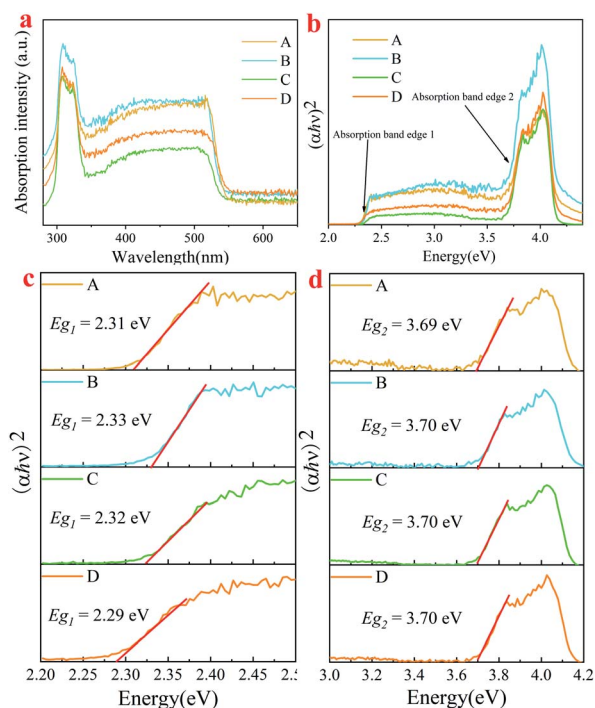


Fig. 8 (a) absorption spectra; (b) Tauc plot; local Tauc plot of absorption band edge 1 (c) and edge 2 (d) for absorption of samples A, B, C and D.

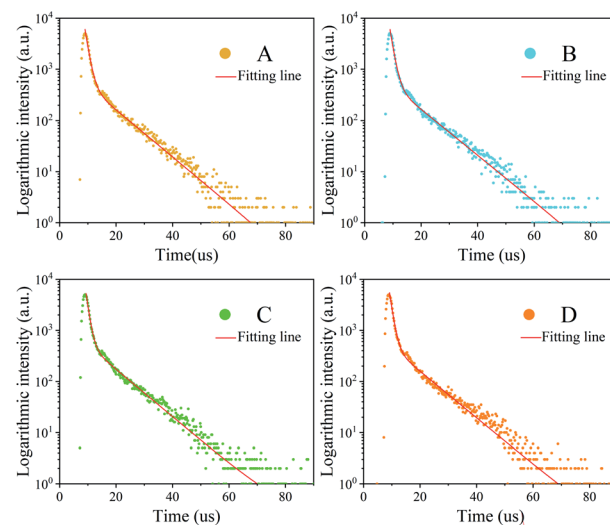


Fig. 9 The PL decays spectra of A, B, C and D.



Table 2 The fitted average lifetime of samples

| Sample | τ_1 | A_1 | τ_2 | A_2 | τ_{av} |
|--------|-----------------|-----------------|------------------|------------------|------------------|
| A | 1.18 ± 0.01 | 9.44 ± 0.10 | 6507.48 ± 12 | 529.59 ± 2.0 | 4.437 ± 0.06 |
| B | 1.18 ± 0.01 | 9.67 ± 0.10 | 6498.67 ± 12 | 527.68 ± 2.0 | 4.572 ± 0.06 |
| C | 1.02 ± 0.01 | 9.34 ± 0.10 | 5732.49 ± 13 | 549.59 ± 2.0 | 4.912 ± 0.05 |
| D | 1.09 ± 0.01 | 9.34 ± 0.10 | 5957.84 ± 12 | 501.55 ± 2.0 | 4.668 ± 0.05 |

HRTEM, as shown in Fig. 3 and 5, respectively. Secondly, previous researches have shown that green fluorescence emission is caused by defects in the Cs_4PbBr_6 crystal⁴¹ and surface ligands can act as passivation agents to reduce crystal defects.⁴² Considering that structural defects tend to be generated at the surfaces of particles, the emission behaviors of the products might be well expected to be influenced when using polar solutions for the purification, if the emission stems from the structural defects. The defects formed during the synthesis process or the purification mediated mid-band-gap states. This is in agreement with the long Urbach tail shown in Fig. 8a, which is common for disordered or poorly-crystalline semiconductors or defects such as V_{Br} . The third is that a small amount of water enters into Cs_4PbBr_6 nanocrystals, which leads to lattice distortion and reduces the band gap to about 3.7 eV.²⁸ At the same time, a small amount of water was ionized to form hydroxyl groups, which react with Cs_4PbBr_6 nanocrystals to form a sub band gap of the hydroxyl group. This speculation is confirmed by the band gap ($E_{\text{g}2}$), which also shows the change in band gap, as shown in Fig. 8d.

The PL decay curve can be fitted well with a biexponential function (S1), where A_1 and A_2 are constants, t is time. τ_1 , τ_2 in eqn (S1) represent the short-lived and long-lived lifetimes relating to the trap-assisted recombination and carrier radiative recombination, separately.⁴³ Besides, the average lifetime (τ_{av}) can be obtained according to eqn (S2) for different nanocrystals.⁴⁴

The fluorescence intensity $I(t)$ of the sample is fitted with a two exponential decay model described as:

$$I(t) = A_1 \exp(-t/\tau_1) + A_2 \exp(-t/\tau_2) \quad (\text{S1})$$

Equation used for the average lifetime calculation:

$$\tau_{\text{av}} = \frac{A_1 \times \tau_1^2 + A_2 \times \tau_2^2}{A_1 \times \tau_1 + A_2 \times \tau_2} \quad (\text{S2})$$

It can be easily found that the trend of a quantum lifetime is first increasing and then decreasing, and there is a peak for sample C. The reagents used to remove excessive ligands (oleic acid and oleic amine in this synthesis), which may also remove some ligands on the nanocrystal surface. Once the surface ligands are removed, the defect-trapping state will increase and the trapping time will be shortened.²⁷ Then, the electron-hole recombination time has also changed as shown in Fig. 9. To a certain extent, the longer lifetime of photoluminescence transition indicates the decrease of the defect concentration and the increase of crystallinity.^{45,46}

The average PL decay lifetime of the synthesized Cs_4PbBr_6 nanocrystals from 4.437 μs to 4.912 μs was derived (Table 2). The long decay lifetime reveals high ratios of radiative recombination to nonradiative recombination and less transition to defect states. The results (long lifetime) can be attributed to their less nonradiative energy transfer to the trap-defect and surface state of the Cs_4PbBr_6 nanocrystals. The small value of τ_1 indicates good intrinsic excitonic feature with fewer defect trappers. From Table 2, it can be seen that the average PL decay lifetime is improved by the aprotic polar solvent.

Conclusions

In summary, we demonstrate the influence mechanism on the optical properties of Cs_4PbBr_6 during purification with solutions of different protonation capabilities and polarity. During the purification process, the organic groups originating from oleic acid (OA) and the PbBr impurity on the surface of Cs_4PbBr_6 nanocrystals can be removed by high polarity aprotic and proton solvents, and the concentration of Br vacancy (V_{Br}) can be reduced. The protonic polar solvent can not only etch the organic groups from the surface of nanocrystals, causing the surface reconstruction and particle growth of nanocrystals, but also enter into the lattice of Cs_4PbBr_6 , and react with the embedded CsPbBr_3 . However, aprotic polar solvent decreases the particle size of Cs_4PbBr_6 nanocrystals with increasing solvent polarity. The optical properties of Cs_4PbBr_6 can be effectively improved by using aprotic polar solvents as purification solvents, which is very significant to improve the luminescence efficiency of perovskites.

Conflicts of interest

The authors declare no conflict of interest.

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Notes and references

- 1 T. C. Sum and N. Mathews, *Energy Environ. Sci.*, 2014, 7, 2518–2534.
- 2 T. M. Brenner, D. A. Egger, A. M. Rappe, L. Kronik, G. Hodes and D. Cahen, *J. Phys. Chem. Lett.*, 2015, 6, 4754–4757.



- 3 P. C. Harikesh, B. Wu, B. Ghosh, R. A. John, S. Lie, K. Thirumal, L. H. Wong, T. C. Sum, S. Mhaisalkar and N. Mathews, *Adv. Mater.*, 2018, **30**, 1802080.
- 4 R. Mayengbam, S. K. Tripathy and G. Palai, *J. Phys. Chem. C*, 2018, **122**, 28245–28255.
- 5 N. M. Rahman, M. Adnaan, D. Adhikary, M. Islam and M. K. Alam, *Comput. Mater. Sci.*, 2018, **150**, 439–447.
- 6 N. Li, Z. Zhu, J. Li, A. K. Y. Jen and L. Wang, *Adv. Energy Mater.*, 2018, **8**, 1800525.
- 7 F. Zhang, B. Yang, X. Mao, R. Yang, L. Jiang, Y. Li, J. Xiong, Y. Yang, R. He, W. Deng and K. Han, *ACS Appl. Mater. Interfaces*, 2017, **9**, 14827–14832.
- 8 J. Li, L. Xu, T. Wang, J. Song, J. Chen, J. Xue, Y. Dong, B. Cai, Q. Shan, B. Han and H. Zeng, *Adv. Mater.*, 2017, **29**, 1603885.
- 9 Q. Zhong, M. Cao, H. Hu, D. Yang, M. Chen, P. Li, L. Wu and Q. Zhang, *ACS Nano*, 2018, **12**, 8579–8587.
- 10 B. Qiao, P. Song, J. Cao, S. Zhao, Z. Shen, D. Gao, Z. Liang, Z. Xu, D. Song and X. Xu, *Nanotechnology*, 2017, **28**, 445602.
- 11 A. Dutta, R. K. Behera and N. Pradhan, *ACS Energy Lett.*, 2019, **4**, 926–932.
- 12 M. Wu, D. Zhao, Z. Wang and J. Yu, *Nanoscale Res. Lett.*, 2018, **13**, 128.
- 13 G. Li, H. Wang, T. Zhang, L. Mi, Y. Zhang, Z. Zhang, W. Zhang and Y. Jiang, *Adv. Funct. Mater.*, 2016, **26**, 8478–8486.
- 14 Q. A. Akkerman, S. Park, E. Radicchi, F. Nunzi, E. Mosconi, F. De Angelis, R. Brescia, P. Rastogi, M. Prato and L. Manna, *Nano Lett.*, 2017, **17**, 1924–1930.
- 15 J. Yin, Y. Zhang, A. Bruno, C. Soci, O. M. Bakr, J.-L. Bredas and O. F. Mohammed, *ACS Energy Lett.*, 2017, **2**, 2805–2811.
- 16 J.-H. Cha, J. H. Han, W. Yin, C. Park, Y. Park, T. K. Ahn, J. H. Cho and D.-Y. Jung, *J. Phys. Chem. Lett.*, 2017, **8**, 565–570.
- 17 M. I. Saidaminov, J. Almutlaq, S. Sarmah, I. Dursun, A. A. Zhumekenov, R. Begum, J. Pan, N. Cho, O. F. Mohammed and O. M. Bakr, *ACS Energy Lett.*, 2016, **1**, 840–845.
- 18 Y. Zhang, L. Sinatra, E. Alarousu, J. Yin, A. M. El-Zohry, O. M. Bakr and O. F. Mohammed, *J. Phys. Chem. C*, 2018, **122**, 6493–6498.
- 19 D. Chen, Z. Wan, X. Chen, Y. Yuan and J. Zhong, *J. Mater. Chem. C*, 2016, **4**, 10646–10653.
- 20 C. Reichardt, *Chem. Rev.*, 1994, **94**, 2319.
- 21 S. Li, G. Liu, Q. Liu, L. Nie, G. Yao, F. Zeng, Y. He and W. Xiang, *ACS Sustainable Chem. Eng.*, 2020, **8**, 10646–10652.
- 22 S. Yan, S. Yang, L. He, C. Ye, X. Song and F. Liao, *Synth. Met.*, 2014, **198**, 142–149.
- 23 X. Chen, F. Zhang, Y. Ge, L. Shi, S. Huang, J. Tang, Z. Lv, L. Zhang, B. Zou and H. Zhong, *Adv. Funct. Mater.*, 2018, **28**, 1706567.
- 24 M. He, W. Chunyun, J. Li, J. Wu, S. Zhang, H.-C. Kuo, L.-Y. Shao, S. Zhao, J. Zhang, F. Kang and G. Wei, *Nanoscale*, 2019, **11**, 22899–22906.
- 25 M. De Bastiani, I. Dursun, Y. Zhang, B. A. Alshankiti, X.-H. Miao, J. Yin, E. Yengel, E. Alarousu, B. Turedi, J. M. Almutlaq, M. I. Saidaminov, S. Mitra, I. Gereige, A. AlSaggaf, Y. Zhu, Y. Han, I. S. Rogan, J.-L. Bredas, O. F. Mohammed and O. M. Bakr, *Chem. Mater.*, 2017, **29**, 7108–7113.
- 26 Y. Zhang, M. I. Saidaminov, I. Dursun, H. Yang, B. Murali, E. Alarousu, E. Yengel, B. A. Alshankiti, O. M. Bakr and O. F. Mohammed, *J. Phys. Chem. Lett.*, 2017, **8**, 961–965.
- 27 L. Yuan, R. Patterson, X. Wen, Z. Zhang, G. Conibeer and S. Huang, *J. Colloid Interface Sci.*, 2017, **504**, 586–592.
- 28 X. Wang, J. Yu, M. Hu, Y. Wu, L. Yang, W. Ye and X. Yu, *J. Lumin.*, 2020, **221**, 116986.
- 29 L. N. Quan, R. Quintero-Bermudez, O. Voznyy, G. Walters, A. Jain, J. Z. Fan, X. Zheng, Z. Yang and E. H. Sargent, *Adv. Mater.*, 2017, **29**, 1605945.
- 30 P. Uthirakumar, H. Yun, M. Devendiran, W. W. Lee and I.-H. Lee, *J. Lumin.*, 2019, **209**, 163–169.
- 31 V. Dracopoulos, D. T. Kastrissios and G. N. Papatheodorou, *Polyhedron*, 2005, **24**, 619–625.
- 32 M. Velazquez, A. Ferrier, S. Pechev, P. Gravereau, J.-P. Chaminade, X. Portier and R. Moncorge, *J. Cryst. Growth*, 2008, **310**, 5458–5463.
- 33 D. M. Calistru, L. Mihut, S. Lefrant and I. Baltog, *J. Appl. Phys.*, 1997, **82**, 5391–5395.
- 34 M. Nikl, E. Mihokova, K. Nitsch, F. Somma, C. Giampaolo, G. P. Pazzi, P. Fabeni and S. Zazubovich, *Chem. Phys. Lett.*, 1999, **306**, 280–284.
- 35 J. Almutlaq, J. Yin, O. F. Mohammed and O. M. Bakr, *J. Phys. Chem. Lett.*, 2018, **9**, 4131–4138.
- 36 J. Xu, W. Huang, P. Li, D. R. Onken, C. Dun, Y. Guo, K. B. Ucer, C. Lu, H. Wang, S. M. Geyer, R. T. Williams and D. L. Carroll, *Adv. Mater.*, 2017, **29**, 1703703.
- 37 F. Palazon, C. Urso, L. De Trizio, Q. Akkerman, S. Marras, F. Locardi, I. Nelli, M. Ferretti, M. Prato and L. Manna, *ACS Energy Lett.*, 2017, **2**, 2445–2448.
- 38 M. Shin, S.-W. Nam, A. Sadhanala, R. Shivanna, M. Anaya, A. Jimenez-Solano, H. Yoon, S. Jeon, S. D. Stranks, R. L. Z. Hoyer and B. Shin, *ACS Appl. Energy Mater.*, 2020, **3**, 192–199.
- 39 S. Ananthakumar, J. R. Kumar and S. M. Babu, *J. Photon. Energy*, 2016, **6**, 042001.
- 40 X. Li, Y. Wu, S. Zhang, B. Cai, Y. Gu, J. Song and H. Zeng, *Adv. Funct. Mater.*, 2016, **26**, 2435–2445.
- 41 R.-T. Liu, X.-P. Zhai, Z.-Y. Zhu, B. Sun, D.-W. Liu, B. Ma, Z.-Q. Zhang, C.-L. Sun, B.-L. Zhu, X.-D. Zhang, Q. Wang and H.-L. Zhang, *J. Phys. Chem. Lett.*, 2019, **10**, 6572–6577.
- 42 F. Gao, Y. Zhao, X. Zhang and J. You, *Adv. Energy Mater.*, 2020, **10**, 1902650.
- 43 S. Sun, D. Yuan, Y. Xu, A. Wang and Z. Deng, *ACS Nano*, 2016, **10**, 3648–3657.
- 44 Z. Liang, S. Zhao, Z. Xu, B. Qiao, P. Song, D. Gao and X. Xu, *ACS Appl. Mater. Interfaces*, 2016, **8**, 28824–28830.
- 45 X. Yang, X. Zhang, J. Deng, Z. Chu, Q. Jiang, J. Meng, P. Wang, L. Zhang, Z. Yin and J. You, *Nat. Commun.*, 2018, **9**, 570.
- 46 L. Zhang, X. Yang, Q. Jiang, P. Wang, Z. Yin, X. Zhang, H. Tan, Y. M. Yang, M. Wei, B. R. Sutherland, E. H. Sargent and J. You, *Nat. Commun.*, 2017, **8**, 15640.

