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Green-Emitting Powders of Zero-Dimensional Cs_4PbBr_6 : Delineating the Intricacies of the Synthesis and the Origin of Photoluminescence

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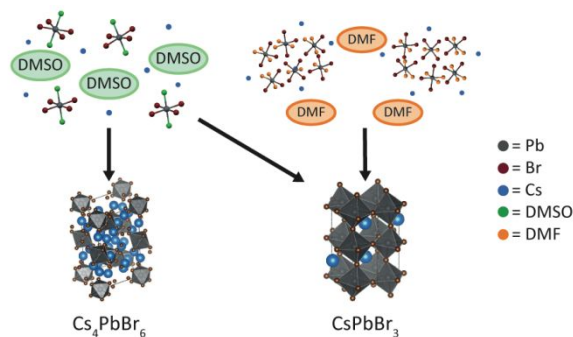
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ABSTRACT. A detailed investigation into the synthesis of green-emitting powders of Cs_4PbBr_6 and CsPbBr_3 materials by antisolvent precipitation from CsBr-PbBr_2 precursor solutions in dimethylformamide (DMF) and dimethyl sulfoxide (DMSO) is reported. Various solvated lead bromide and polybromide species (PbBr_2 , $[\text{PbBr}_3]^-$, $[\text{PbBr}_4]^{2-}$, and possibly $[\text{PbBr}_5]^{3-}$ or $[\text{PbBr}_6]^{4-}$) are detected in the precursor solutions by optical absorbance and emission spectroscopies. The solvodynamic size of the species in solution is strongly solvent-dependent: ~ 1 nm species were detected in DMSO, while significantly larger species were observed in DMF by dynamic light scattering. The solvodynamic size of the lead bromide species plays a critical role in determining the Cs-Pb-Br composition of the precipitated powders: smaller species favor the precipitation of Cs_4PbBr_6 , while larger species template the formation of CsPbBr_3 under identical experimental conditions. The powders have been characterized by ^{133}Cs and ^{207}Pb solid-state nuclear magnetic resonance and ^{133}Cs sensitivity toward the different Cs environments within Cs_4PbBr_6 is demonstrated. Finally, the possible origins of green emission in Cs_4PbBr_6 samples are discussed. It is proposed that a two-dimensional Cs_2PbBr_4 inclusion may be responsible for a green emission at ~ 520 nm in addition to the widely acknowledged CsPbBr_3 impurity, although we found no conclusive experimental evidence supporting such claim.

TOC



All-inorganic lead halide semiconductors have recently attracted renewed research interest due to the various stoichiometries that can be crystallized from the same precursors. Lead halide compounds demonstrate structural tunability based on the connectivity of the $[\text{PbX}_6]^{4-}$ ($\text{X} = \text{Cl}^-$, Br^- , or I^-) octahedra that can form three-dimensional (3D) corner-sharing structures (perovskites), layered 2D networks, chains of 1D, as well as 0D isolated octahedral clusters.¹⁻⁵ In case of the organic-inorganic lead halide compounds, the size of the organic cation has been found to be one of the main factors determining the dimensionality of the octahedral framework. However, in the all-inorganic Cs-Pb-Br compounds, all three dimensionalities can be formed with the same Cs cation. Mixing CsBr and PbBr_2 , in specific ratios and in appropriate solvents, leads to the formation of pure or mixed structures of CsPbBr_3 (3D), CsPb_2Br_5 (2D), and Cs_4PbBr_6 (0D).⁶ PbBr_2 -rich conditions lead to the 2D tetragonal CsPb_2Br_5 phase, characterized by a sandwich structure with Cs^+ ions in between layers of $[\text{Pb}_2\text{Br}_5]^-$. On the other hand, under CsBr-rich conditions, a trigonal 0D Cs_4PbBr_6 structure is formed, which consists of disconnected $[\text{PbBr}_6]^{4-}$ octahedra separated by Cs^+ ions. An equimolar amount or a slight excess of PbBr_2 with respect to CsBr results in formation of 3D CsPbBr_3 with an orthorhombic structure of corner-sharing $[\text{PbBr}_6]^{4-}$ octahedra, as in the organic-inorganic perovskites (MAPbX_3 and FAPbX_3 ; MA = methylammonium and FA = formamidinium) commonly used in optoelectronic applications.⁷

Cs_4PbBr_6 based materials have recently gained increased interest due to the reports of strong and stable green photoluminescence (PL) in their powder, thin-film, single crystal and nanocrystal forms.⁸⁻¹⁵ However, it was reported that Cs_4PbX_6 compounds are colorless¹⁶ with bandgaps in the UV region.¹² Specifically, Cs_4PbBr_6 has a matching experimental and theoretical bandgap around 3.95 eV.¹⁷⁻¹⁹ Hence, the origin of the green PL has been proposed to be either due to intrinsic defects within the wide bandgap Cs_4PbBr_6 ¹³⁻¹⁵ or due to the presence of 3D CsPbBr_3

nanocrystals.¹⁹⁻²⁴ Ling et al.²⁵ suggested that the 3D CsPbBr₃ phase is not the direct cause of the intense PL; instead, they proposed trap states at the interface between the two phases (3D and 0D) to be responsible for that intense green emission.

Herein, we explore reaction parameters governing the formation of Cs₄PbBr₆ and CsPbBr₃ phases from CsBr-PbBr₂ solutions and the origin of the green PL in the Cs₄PbBr₆ samples. The nature and solvodynamic size of the lead bromide species formed in solution varies with the coordination strength of the solvent and subsequently determines the composition of the formed Cs-Pb-Br powders: species with a ~1 nm solvodynamic radius (presumably isolated octahedra) were detected in dimethyl sulfoxide (DMSO) solutions of CsBr-PbBr₂ leading to the formation of Cs₄PbBr₆ phase upon addition of an antisolvent, while larger species were detected in dimethylformamide (DMF) and resulted in the formation of CsPbBr₃ phase. In addition to the solvent, the solvent-antisolvent pair and the absolute concentrations of the precursors are key parameters defining the structure of the precipitated material. For example, antisolvents such as tetrahydrofuran (THF) and dichloromethane (DCM) precipitate mainly Cs₄PbBr₆ and CsPbBr₃ phases, respectively, from the CsBr-PbBr₂ solutions in DMSO when used in 1:6 (solvent:antisolvent by volume) ratios. Increasing the concentration of CsBr-PbBr₂ while keeping their molar ratio fixed (1:1) from 0.05 M to 0.5 M in DMSO favors formation of CsPbBr₃ at higher concentrations. In addition to the powder X-ray diffraction (XRD) technique commonly used for a structural and compositional identification of the materials, we apply ¹³³Cs and ²⁰⁷Pb solid-state nuclear magnetic resonance (ssNMR) to the analysis of the various Cs-Pb-Br precipitates. We found that ¹³³Cs ssNMR is more sensitive toward detecting relatively low impurity content of the different phases compared to its ²⁰⁷Pb counterpart. The ¹³³Cs ssNMR results are in agreement with XRD data in terms of observed phases, but ¹³³Cs ssNMR could also distinguish between the two

different Cs crystallographic positions in the 0D crystal. Finally, based on the observed absorbance in the visible region, and the recurring green PL of Cs₄PbBr₆ samples, we hypothesize that a 2D Cs₂PbBr₄ inclusion (not observed experimentally) may be responsible for a green PL in addition to the widely acknowledged 3D impurity.

EXPERIMENTAL SECTION

Materials. CsBr (99.9%, Sigma-Aldrich), PbBr₂ (98%, Sigma-Aldrich), DMSO (99.9%, Sigma-Aldrich), DMF (99.8%, Sigma-Aldrich), THF (99.9%, Sigma-Aldrich) and DCM (99.8%, amylene stabilizer, Sigma-Aldrich).

Preparation of Cs-Pb-Br Powders. Generally, CsBr and PbBr₂ were dissolved in DMSO or DMF at the required ratio and at the required concentration by sonication for 1 hour. Then the solutions were filtered using PTFE filter with 0.2-mm pore size and the precipitation was carried out by quick injection of THF or DCM to the filtrate.

Characterization. The optical absorbance of the precursor solutions was recorded on a Varian Cary Eclipse UV-vis spectrophotometer. PLE and PL of precursor solutions were performed on a Varian Cary Eclipse fluorescence spectrophotometer. DLS was performed using a Zeta Sizer (Malvern Instruments) equipped with a 4.0 mW He-Ne laser operating at 633 nm and an avalanche photodiode detector, in order to determine the average size of the precursors in the mother solution.

X-ray Diffraction analysis was performed using PANalytical Empyrean X-ray diffractometer equipped with a 1.8 kW Cu K α ceramic X-ray tube and a PIXcel3D 2 \times 2 area detector, operating at 45 kV and 40 mA. Cs-Pb-Br powders were studied under ambient conditions using a parallel beam Bragg Brentano geometry. XRD data was analysed using the HighScore 4.1 software from

PANalytical. Step sizes of 0.026° and 0.013° and scan speeds of 0.102 and $0.013^\circ/\text{s}$ were used for the standard and slow scans, respectively.

PL and PLQY measurements on dried powders were conducted on an Edinburgh Instruments FLS920 fluorescence spectrometer equipped with a calibrated integrating sphere exciting at 405 nm. Absorbance measurements for the powder samples were carried out in a Cary 5000 spectrometer using the diffuse reflectance accessory. Optical microscopy of the 4:1_DMSO powder was performed on a Nikon 80i microscope equipped with a grayscale digital camera and a high-pressure mercury lamp (Nikon) and optical filters. The sample was prepared by mixing a small amount of the finely ground powder with a drop ($\sim 5 \mu\text{l}$) of octadecene-1. The mixture was prepared on top of a rectangular microscope slide and covered with a thin glass coverslip. Confocal PL microscopy on the same sample was performed using a Nikon A1 confocal laser microscope with 488 nm excitation (Oxxius LBX-488 diode laser). The laser power setting was kept at 0.5%, and the resolution of a 32-channel spectral detector set at 2.5 nm during the experiments. The spectrally-resolved imaging data and the z-series used for a volume view reconstruction (Figure S8) were acquired (as *.nd2 files) using Nikon NIS-Elements High Content Software Analysis ver. 4.30.02 and viewed and processed in either Nikon NIS-Elements Viewer ver. 4.20.00 or ImageJ 1.51j8 as previously described.²⁶

XPS measurements were carried out on a Kratos Axis UltraDLD spectrometer, using a monochromatic Al K α source (15 kV, 20 mA). The spectra were taken on a $300 \times 700 \mu\text{m}^2$ area. Wide scans were collected with a constant pass energy of 160 eV and an energy step of 1 eV. High-resolution spectra were acquired at a constant pass energy of 10 eV and energy step of 0.1 eV. The binding energy scale was referenced to the C 1s peak at 284.8 eV. The spectra were

analyzed using the CasaXPS software (version 2.3.17). Samples were prepared by covering an indium substrate with the Cs-Pb-Br powder.

High-resolution transmission electron microscopy (HRTEM) analysis was carried out on a 200 kV TEM (JEOL JEM-2200FS) and a Bruker Quantax 400 system with an XFlash 5060 detector was used for EDS analysis. The sample was prepared by dispersing the powders in THF before depositing on an ultrathin carbon film supported on a holey carbon film coated 400 mesh TEM copper grids.

Solid-state NMR (ssNMR) measurements were collected on a 500 MHz *Bruker BioSpin* NMR Spectrometer AvanceTM 500, operating at a static field of 11.7 Tesla and equipped with a 4 mm CPMAS probe, by spinning the sample at the magic angle (MAS), with speed rates from 10 to 14 kHz. All samples were packed into Zirconia (ZrO₂) rotors and closed with Kel-F caps (80 μ L internal volume). The working frequency of the ¹³³Cs nucleus was 65.598 MHz and spectra were acquired by using a single pulse sequence (zg), with a spectral width of 104166 Hz, wide enough to ensure a homogeneous irradiation of the whole spectrum. The optimal spinning speed (MAS) was achieved at 10 KHz; however, comparative spectra were also acquired at different spinning rates (12 KHz and 14 KHz), to assign all sidebands in the spectrum and to exclude any possible signal impurity overlapping to the manifold SSB. The NMR parameters used for ¹³³Cs experiments were: 90° pulse length (p1) = 2.5 μ s, repetition time (d1) = 3-60 s, number of scans (ns) = 200-800. The working frequency of ²⁰⁷Pb nucleus was 104.63 MHz and spectra were acquired by using a single pulse sequence (*solidecho*, with an echo time of 7 μ s), with a spectral width of 1600 ppm (corresponding to 167.000 Hz), wide enough to allow an homogeneous irradiation of the whole spectrum, but not enough to cover the wide anisotropic chemical shift range, typical of such a nucleus. Different experiments were needed (with the same spectral width but changing the offset)

to cover the whole possible range of detectable signals. The spinning speed (MAS) was optimized at 10 kHz, a value that returned resolved spectra. The NMR parameters used for ^{207}Pb experiments were the following: 90° pulse length ($p1$) = 2 μs , repetition time ($d1$) = 0.5 s, ns = 28.000-105.000. ^{133}Cs and ^{207}Pb shifts were referenced to solid CsBr (δ_{iso} = 268.0 ppm vs. CsCl: δ_{iso} = 223.2 ppm) and $\text{Pb}(\text{NO}_3)_2$ (δ = -3490 ppm), respectively, which were used as external calibration standards.

RESULTS AND DISCUSSION

CsBr-PbBr₂ solutions in DMSO and DMF. PbBr₂-containing solutions in DMF and DMSO are precursors for the Cs₄PbBr₆ and CsPbBr₃ samples, thus it is anticipated that the nature of the solvated species has an effect on the resulting material obtained from them. Steady-state optical spectroscopy is a simple method for investigation of the lead bromide speciation. Absorbance, photoluminescence (PL), and photoluminescence excitation (PLE) spectra of 0.05 M solutions of CsBr:PbBr₂ mixtures (1:1 molar ratio) in DMSO and DMF were collected and are reported in Figure 1. The DMSO solution showed an absorbance peak at 285 nm that is attributed to PbBr₂, while the DMF solution had a clear absorbance peak at 310 nm corresponding to $[\text{PbBr}_3]^-$ (Figure 1a), consistent with previous studies.²⁷⁻²⁸ Both solutions showed an absorbance tail in the longer wavelength region (340 - 360 nm, inset in Figure 1a), suggesting the possible formation of $[\text{PbBr}_4]^{2-}$ species.²⁷ Both $[\text{PbBr}_3]^-$ and $[\text{PbBr}_4]^{2-}$ species are emissive (at ~600 nm and 560 nm, respectively)²⁷⁻²⁸ and their presence can be confirmed through selective wavelength excitation (Figure 1b). The DMSO solution emitted at 560 nm regardless of the excitation wavelength, indicating the presence of mainly $[\text{PbBr}_4]^{2-}$. On the other hand, the DMF solution exhibited emissions at 560 and 600 nm, confirming the presence of detectable amounts of both $[\text{PbBr}_3]^-$ and $[\text{PbBr}_4]^{2-}$ species. A relatively higher PL intensity of the DMF solution compared to DMSO under 360 nm excitation (Figure 1b) suggests relatively higher concentration of $[\text{PbBr}_4]^{2-}$ species in DMF

with respect to DMSO (consistent with a higher absorbance at around ~360 nm). The presence of an extra PLE peak at 374 nm in the DMF solution (Figure 1b) that is red shifted relative to that of $[\text{PbBr}_4]^{2-}$ in the DMSO solution suggests that higher-order lead polybromide complexes, such as $[\text{PbBr}_5]^{3-}$ or $[\text{PbBr}_6]^{4-}$ might be present in DMF (by analogy with their iodide counterparts).²⁹

We performed DLS measurements on both solutions to further investigate the discussed differences in lead bromide speciation between two solvents. It was found that the DMSO solution contained roughly 1 nm-sized lead bromide species while the DMF solution was composed of significantly larger clusters (Figure 1c). The large-sized clusters in the DMF solution could be due to an aggregation or polymerization of the lead bromide species, because of the different solvent-solute interaction between PbBr_2 and DMF compared to DMSO (similar to PbI_2 ²⁹⁻³¹). Our findings indicate that a change of solvent significantly alters the speciation and the solvodynamic size of the lead bromide species, as schematically summarized in Figure 1d. The results hold in the absence of CsBr, as summarized in the SI (see Figure S1 and an accompanying discussion). Previously, the changes in the solvodynamic size of the lead polyhalide species have been attributed to the relative ratio of salts (e.g. $\text{MAI}:\text{PbI}_2$; MA = methylammonium)³² or to the effect of additives such as hydrohalic acids.³³ This is of high significance to the perovskite photovoltaic community as the solvodynamic size of the species present in the precursor solution influences the composition of the resulting material (see below) and subsequent optoelectronic properties of the devices.³²⁻³³

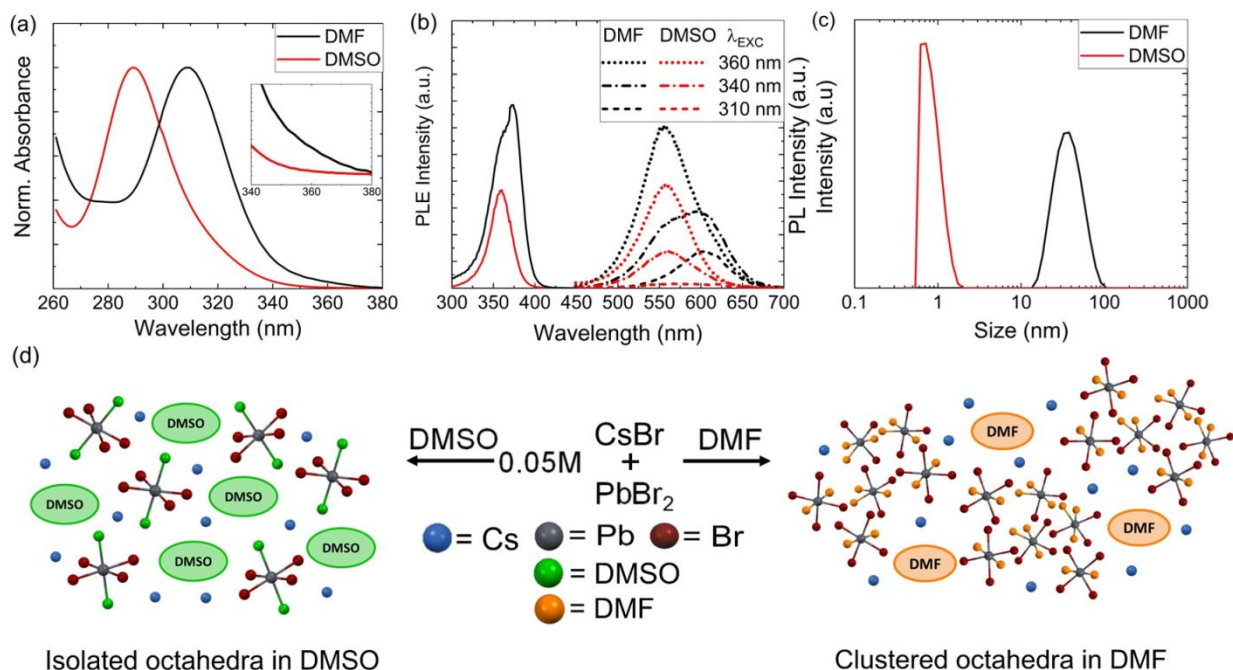


Figure 1. (a) Normalized Absorbance, (b) PLE (solid lines) and PL (dotted and dashed lines) spectra of 0.05 M 1:1 CsBr:PbBr₂ precursor solutions in DMF and DMSO. PLE is reported at the highest intensity emission (560 nm) while PL is recorded at different excitation wavelengths. (c) Corresponding DLS measurements. (d) Scheme illustrating the photoactive species formed upon dissolving 0.05 M 1:1 CsBr:PbBr₂ in DMSO and DMF.

Cs₄PbBr₆ and CsPbBr₃ formation. Next we investigated the differences in the composition of the powders precipitated from the abovementioned CsBr-PbBr₂ solutions in DMSO and DMF. Using the equimolar (0.05 M of each precursor) solutions in DMSO and DMF, the precipitation of the powders was done by a quick injection of 6 mL of tetrahydrofuran (THF) to 1 mL of the solution. Powders were then dried overnight in a vacuum oven at 40 °C. X-ray diffraction (XRD) patterns of the dried powders are plotted in Figure 2a along with the simulated patterns of Cs₄PbBr₆ and CsPbBr₃. It is observed that mainly the 0D trigonal phase was obtained from the DMSO solution, with only low intensity peaks corresponding to a 3D impurity, while 3D CsPbBr₃ was

the main precipitate from the DMF solution, with an extremely low contamination from the 0D phase (inset in Figure 2a). The 0D impurity was only detected upon performing a slow XRD scan (see experimental section). We could eliminate the 0D impurity by using a CsBr:PbBr₂ ratio of 1:1.2 in DMF. This is relevant for perovskite-based light-emitting devices, as we observed that even this extremely small amount of 0D impurity in the 3D sample could significantly enhance the PL peak (Figure S2). The differences in the compositions obtained from DMSO (mainly Cs₄PbBr₆) and DMF (mainly CsPbBr₃) under identical precipitation conditions can be rationalized on the basis of the DLS data of the precursor solutions (Figure 1c). The formation of clusters of the lead bromide species in the DMF solution possibly templates the growth of the 3D orthorhombic phase upon the addition of the antisolvent. On the other hand, the approx. one nm-sized particles in DMSO, which is in agreement with a previous report,³⁴ suggests that most of the complexes in solution are isolated octahedra, making the crystallization of the 3D phase less favorable. Coordination strength of the solvent is likely a main factor responsible for these differences. DMSO is known to have a stronger coordination with Pb²⁺ compared to DMF;³⁰ hence, the Cs-rich phase Cs₄PbBr₆ can be easily precipitated from DMSO solution while the composition of precipitates from DMF solution is closer to the stoichiometry of the feed solution (1:1 CsBr:PbBr₂). Nonetheless, 3D CsPbBr₃ could be precipitated out from a DMSO solution by changing the antisolvent or the absolute concentration of the precursors, as discussed next.

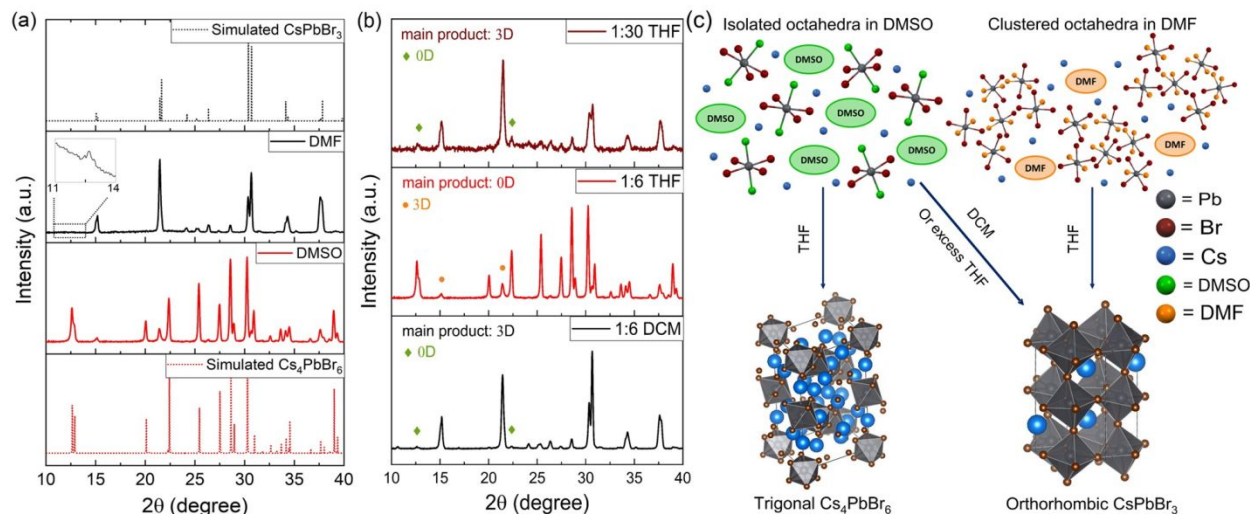


Figure 2. (a) XRD patterns of Cs-Pb-Br powders precipitated from 0.05 M 1:1 CsBr:PbBr₂ DMSO and DMF solutions along with simulated patterns for trigonal Cs₄PbBr₆ (ICSD-162158) and orthorhombic CsPbBr₃ (ICSD-97851) phases. (b) XRD of Cs-Pb-Br powders precipitated from different DMSO:antisolvent volume ratio. (c) Scheme illustrating the effect of the solvodynamic size and solvent-antisolvent pair on the formed phases.

We studied the effect of two different antisolvents on an equimolar (0.05 M) mixture of CsBr and PbBr₂ dissolved in DMSO. THF and dichloromethane (DCM) were chosen as antisolvents. XRD patterns of the powders are presented in Figure 2b. Simply by changing the antisolvent from THF (1:6 THF) to DCM (1:6 DCM), despite the absence of any large-sized clusters, 3D perovskite is the main precipitate (see sketches in Figure 2c). Hence, it is not only the solvodynamic size of the different lead polybromide in the precursor solution that controls the phase of the obtained precipitate but also the solvent-antisolvent pair. In a 1:1 CsBr:PbBr₂ DMSO solution, addition of THF favors the precipitation of the Cs-rich phase (Cs₄PbBr₆) possibly due to the higher solubility of PbBr₂, compared to CsBr, in this solvent-antisolvent pair. On the other hand, using DCM as an antisolvent resulted in the precipitation of the 3D phase as both precursors are relatively insoluble

in the DMSO-DCM pair. Nevertheless, when an excess amount of THF is added (1:30 THF), the 3D phase is the primary formed phase (see Figure 2b). It could be possible that excess THF forces full precipitation of the PbBr_2 present in the precursor solution, hence the stoichiometric 3D phase is formed. The effect of the solvodynamic size of the different lead polybromide species and solvent-antisolvent pair on the resulting Cs-Pb-Br phase is demonstrated in Figure 2c. We also explored a mixed DMSO/DMF solution and found that both THF and DCM leads to the precipitation of mainly the 3D phase with minor 0D impurity (Figure S3). Another parameter that we varied was the absolute concentration of the solution (0.05 M to 0.5 M) using DMSO-THF as the solvent-antisolvent pair, and interestingly discovered that the lower concentration favors the precipitation of majority 0D while the higher concentration favors a majority 3D perovskite (Figure S4). The 3D precipitated from the 0.5 M DMSO solution is likely due to the high concentration that results in significant interaction between adjacent octahedra after the addition of the antisolvent.³⁵ In fact, upon direct addition of the antisolvent, we observed a white precipitate, most probably 0D phase or ultra-small 3D clusters³⁶ that in few seconds turned into orange colored 3D perovskite.

XRD analysis of the precipitate from the lower absolute concentration (0.05 M) detected around 15% 3D CsPbBr_3 (see Note S1), hence, we used a 4:1 CsBr: PbBr_2 ratio in order to precipitate 3D-free 0D Cs_4PbBr_6 (Figure 3, blue pattern). We observed that the (121) peak at 21.5° (2theta) corresponding to the 3D orthorhombic phase completely disappeared. However, at this precursor ratio, peaks, especially the (110) peak at 29.5° (2theta), corresponding to cubic CsBr are detected. Importantly, powders precipitated from a 3.75:1 of CsBr: PbBr_2 contained detectable 3D impurity (Figure 3, green pattern). These results suggest that the formation of CsBr- and CsPbBr_3 -free

Cs_4PbBr_6 from DMSO and DMF solutions could be challenging. It is worth mentioning that the dried powders precipitated from all the different ratios were green emitting under the UV-lamp.

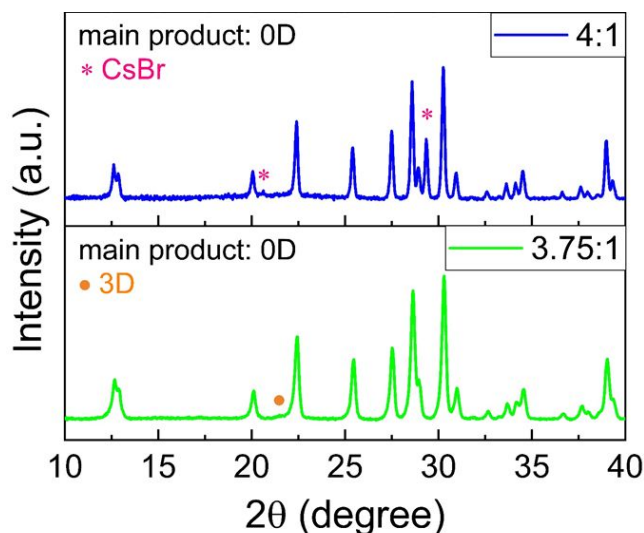


Figure 3. XRD patterns of powders precipitated from a 3.75:1 (green pattern) and a 4:1 (blue pattern) $\text{CsBr}:\text{PbBr}_2$ precursor solutions in DMSO.

ssNMR on Cs-Pb-Br powders. We carried out ^{207}Pb and ^{133}Cs solid-state nuclear magnetic resonance (ssNMR) analysis to explore the composition of our precipitates. ssNMR is a powerful characterization tool that can distinguish between different crystal structures of the same material,³⁷ and it can also identify the formation of amorphous and hydrated hybrid perovskite phases.³⁸⁻⁴⁰ Since it is well known that the main drawback for NMR is the intrinsic poor sensitivity, we first checked the sensitivity of ssNMR in detecting perovskite impurities in the Cs-based perovskites. For this purpose, two samples were used, specifically, a 3D CsPbBr_3 with very low 0D Cs_4PbBr_6 impurity (1:1_DMF) and 0D Cs_4PbBr_6 with very low 3D CsPbBr_3 impurity (3.75:1_DMSO) as suggested by their XRD patterns shown in Figure 2a (black pattern) and Figure 3 (green pattern), respectively. All ssNMR data are summarized in Table 1. The detection of the

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3 impurity phases was successful by ^{133}Cs ssNMR (Figure 4a) but not by ^{207}Pb ssNMR (Figure 4b),
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5 despite the first nuclide being quadrupolar ($I=7/2$, natural abundance = 100%) while ^{207}Pb is not
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7 ($I=1/2$, natural abundance = 22.6%), suggesting higher sensitivity of ^{133}Cs ssNMR compared to its
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9 ^{207}Pb counterpart in these high crystalline materials. It is worthwhile to notice that the Pb signals
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11 show a fine structure with well-defined components of the multiplet originating from the coupling
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13 between ^{207}Pb and the six $^{79/81}\text{Br}$ atoms ($I=3/2$ natural abundance ^{79}Br 50.54% and ^{81}Br 49.46%),
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15 surrounding the metal ions in an octahedral geometry, as already reported for similar Cl
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17 complexes.⁴¹ We found that ^{133}Cs ssNMR was also able to distinguish between two kinds of Cs
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19 atoms, lying in different environments in the 0D crystal structure⁴² with a ratio between 2:1 to 3:1,
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21 as expected from the 0D unit cell. Moving to the “3D-free” 0D sample (4:1_DMSO), the XRD
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23 pattern of which we report in Figure 3 (blue pattern), ^{133}Cs ssNMR identified the CsBr ($\delta_{\text{iso}} = 268$
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25 ppm), which was also detected by XRD. Importantly, ^{133}Cs ssNMR did not detect any 3D impurity
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27 in agreement with powder XRD.
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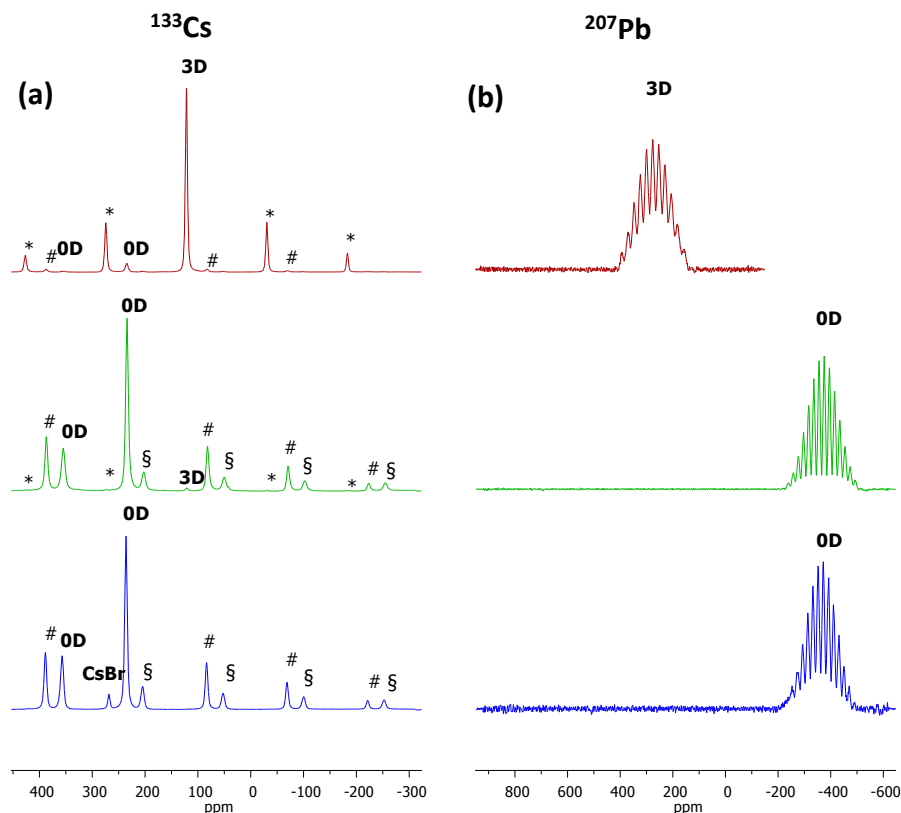


Figure 4. (a) ^{133}Cs and (b) ^{207}Pb ssNMR spectra (10 kHz MAS, 11.75 T, 312 K) of the samples 1:1_DMF (red trace), 3.75:1_DMSO (green trace), and 4:1_DMSO (blue trace). The asterisks (*) mark the spinning sidebands (SSB) of 3D phase signal, the section signs (§) mark the SSB of the less intense 0D peak while the hash signs (#) mark the SSB of the more intense 0D peak.

Table 1. Summary of powder XRD, and ^{133}Cs and ^{207}Pb ssNMR analysis on 3 different samples. In brackets the δ_{iso} (ppm) of the different phases is reported. For ^{207}Pb signal the scalar coupling constant $^1J_{\text{Pb-Br}}$ typical for these systems is also included.⁴³

Sample CsBr:PbBr ₂ _solvent	Powder XRD	^{133}Cs ssNMR	^{207}Pb ssNMR
1:1_DMF	3D + 0D impurity	3D (120 ppm) + 0D impurity	3D only (246 ppm, $^1J_{\text{Pb-Br}}=2425$ Hz)

3.75:1_DMSO	0D + 3D impurity	0D (354 ppm and 233.6 ppm) + 3D impurity	0D only (-373 ppm, $^1J_{\text{Pb-Br}}=2050 \text{ Hz}$)
4:1_DMSO	0D + CsBr	0D + CsBr (268 ppm)	0D only

Optical and compositional analysis of green-emitting Cs_4PbBr_6 powder. We focused on the “3D-free” 0D powder (4:1_DMSO). Although XRD (Figure 3 blue pattern) and ssNMR (Figure 4 blue trace) did not detect any 3D impurity in this sample, diffuse reflectance measurements, as plotted in Figure 5a, showed absorption throughout the green region with a matching PLE (Figure S5) and a single PL emission peak at 520 nm (Figure 5a). It is worth mentioning that in the case of our 0D powders with (3.75:1_DMSO) or without (4:1_DMSO) detectable 3D impurity, the as-precipitated wet powders are non-emissive under the UV-lamp. However, they turn green emitting after some time, whether the precipitate is left in solution, dried in a vacuum oven at 40 °C, or under N_2 at room temperature.

An inspection of the 4:1_DMSO sample with optical microscope under white light (Figure S6a) and UV light (Figure S6b) revealed that the green emission is localized in bright spots scattered across the sample. This observation was further confirmed by confocal PL microscope using 488 nm excitation (Figures 5b). The PL spectrum collected from a local area containing several emission centers peaked at ~520 nm (Figure S7), in agreement with the PL spectrum in Figure 5a. A spatial distribution of the emitting centers in three dimensions was obtained through the series of z slices of ~40×40×5 micron volume of the sample (Figure S8). It further confirms that the emission originates from the localized spots in the sample, which are unevenly distributed across the sample. The size of the emissive domains is sub-micron and resolving their shape is beyond the spatial resolution of the microscope used.

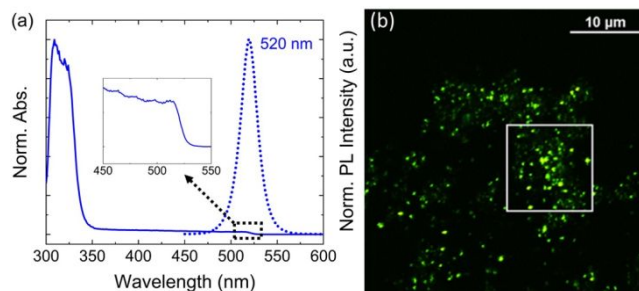


Figure 5. (a) Optical absorption (solid line) of 4:1_DMSO powder sample obtained by converting reflectance (R) to absorption using the Kubelka-Munk equation⁶: $F(R) = (1-R)^2/2R$ and PL emission (dotted line) at excitation wavelength of 405 nm. (b) Confocal PL microscopy image recorded with 488 nm excitation, the PL spectrum from an area highlighted by the white square is shown in Figure S7.

Elemental analysis of the powder was done using X-ray photoelectron spectroscopy (XPS) and Energy-dispersive X-ray spectroscopy (EDS) and the compositional data is listed in Table S1. Both XPS and EDS analyses reveal that the sample is Br-deficient. This Br-deficiency contradicts most reports on green-emitting 0D material where samples were reported to be Br-rich with Pb:Br ratios up to 1:7.4.⁴⁴ The bromine deficiency in our 0D sample could be due to incorporation of other phases with lower bromine content. Next, we measured the separation between the onset of valence band maxima (VBM) and Fermi level shown in Figure 6a. We observed only a small shift when compared to reported VBM onsets of 3D CsPbBr₃, suggesting a possible 3D or 3D-like impurity in our 0D sample.⁴⁵ Similarly, high-resolution transmission electron microscopy (HRTEM) analysis on the 0D sample detected a domain with lattice spacing of 5.8 Å that is indicative of 3D-like nature (Figure 6b,c). Based on elemental analysis of different particles in the 4:1_DMSO sample, Cs₄PbBr₆ and the impurity CsBr particles were observed as individual entities and not as a composite (Figure S9).

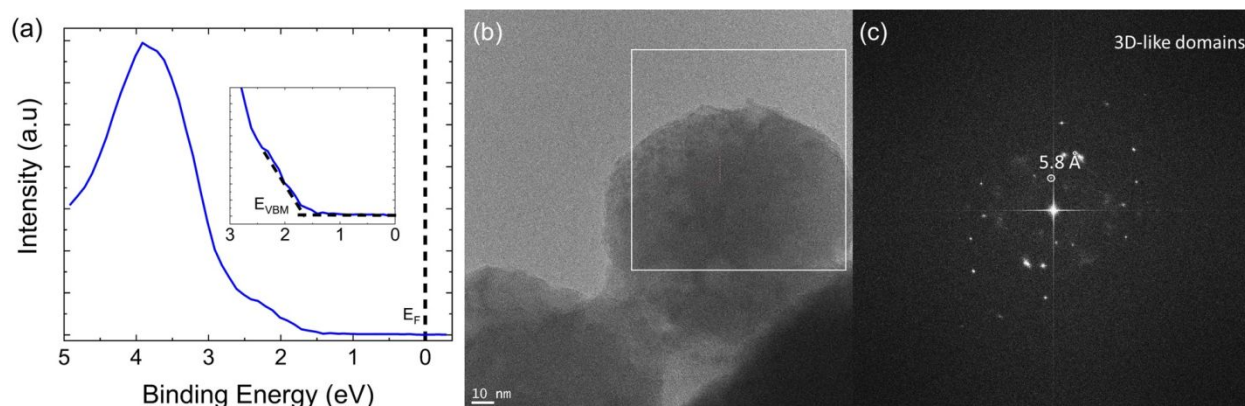


Figure 6. (a) XPS results on the 4:1_DMSO sample showing the separation between the onset of valence band maxima (VBM) and Fermi level. (b) HRTEM image and (c) corresponding Fast Fourier transform (FFT) of the white square area showing lattice spacing of 5.8 Å.

Origin of the green emission. While bromine vacancy (V_{Br}) transition level energy in the Cs_4PbBr_6 was calculated by Yin et al.¹³ to be 2.3 eV above the VBM, matching the green emission, recently, V_{Br} was instead suggested to form a shallow defect level.⁴⁶ In addition, as we have previously reported,¹² the intense and sharp emission and the absence of Stokes shift does not match with a deep-trap emission. Therefore, impurity emission would be more reasonable as the origin of the green emission. Impurity emission is in line with our confocal PL microscopy, VBM analysis, and HRTEM findings. 3D CsPbBr_3 could be the impurity source of the green PL emission, as non-quantum confined nanocrystals (> 8 nm) do indeed emit at around 2.39 – 2.41 eV.⁴⁷ However, in agreement with Wang et al.⁴⁸ it is expected that the 3D CsPbBr_3 impurity might not be the only origin of the green emission. For instance, a previous report detected quantum confined ~ 3 nm CsPbBr_3 nanocrystals embedded within larger-sized Cs_4PbBr_6 using HRTEM, however, the sample was green emitting at 520 nm,²² defying reported PL values for such small CsPbBr_3 nanocrystals.⁴⁷ In fact, PLQYs of our 0D samples with detectable 3D impurity (3.75:1_DMSO) and 3D-free (4:1_DMSO) were 25% and 50%, respectively.

Other than the known perovskite or perovskite-related phases of 3D (CsPbBr_3), 2D (CsPb_2Br_5) and 0D (Cs_4PbBr_6), Jizhong et al.⁴⁹ reported the synthesis of a Cs_2PbBr_4 material, which they described as a 2D sheet of corner sharing $[\text{PbBr}_6]^{4-}$ octahedra passivated by Cs^+ ions, exhibiting an absorbance and emission in the green region of the visible spectrum. This phase has not been previously reported in the phase diagram of this ternary Cs-Pb-Br system and the synthesis of a bulk sample of Cs_2PbBr_4 appears to be challenging, as the phase was suggested to be thermodynamically unstable.⁵⁰ However, this 2D sheet of corner-shared $[\text{PbBr}_6]^{4-}$ octahedra is in essence a monolayer of 3D CsPbBr_3 . Furthermore, the 2D Ruddlesden-Popper (RP) phase of $\text{Cs}_{n+1}\text{Pb}_n\text{Br}_{3n+1}$ with $n = 1$ or 2 has also been detected as an impurity within 3D CsPbBr_3 nanosheets.⁵¹ Theoretically, at $n = 1$, the Cs_2PbBr_4 layer is predicted to be a direct bandgap material, and upon introducing spin orbital coupling (SOC) combined with hybrid function (HSE), Yang et al.⁵² calculated its bandgap to be 2.29 eV.

In pure 2D perovskites such as 2D MA_2PbI_4 , the difference between the bandgaps of a monolayer and bulk material is theoretically calculated to be of only 0.01 eV.⁵³ Experimentally, unlike the quasi-2D e.g. $(\text{BA})_2(\text{MA})_{n-1}\text{Pb}_n\text{I}_{3n+1}$ (BA = butylammonium) that possess considerable blue shift in the PL peak position as number of layers (n) decreased,⁵⁴ pure 2D perovskites showed insignificant differences.⁵⁵⁻⁵⁶ Hence, a 2D perovskite structure with small Cs cation as an interlayer spacer could explain the narrow range of PL (515 nm to 524 nm)^{12, 57} reported for the emissive 0D single crystals, powders and nanocrystals. The proposed Cs_2PbBr_4 is different from 3-5 layered CsPbBr_3 nanosheets or nanoplatelets that usually emit between 440 nm to 460 nm.⁵⁸⁻⁵⁹ This discrepancy in the PL peak position could be attributed to the fact that, in case of nanosheets and nanoplatelets, their top and bottom surfaces are covered with large organic cations such as

oleylammonium or octylammonium, hence, they could be considered as quasi-2D perovskites, leading to the pronounced blue shift in their emissions.

This hypothesis of 2D Cs_2PbBr_4 inclusions being responsible for the absorbance and emission in the green region of the visible spectrum is in agreement with Liu et al.⁶⁰ where they suggested the formation of $[\text{PbBr}_4]^{2-}$ intermediates (in the form of sheet structures observed in TEM) during the transformation of CsPbBr_3 nanocrystals to Cs_4PbBr_6 nanocrystals. It is possible that the 0D matrix stabilizes this 2D $\text{Cs}_{n+1}\text{Pb}_n\text{Br}_{3n+1}$ with $n = 1$ (Cs_2PbBr_4) phase similar to what was detected in 3D nanosheets.⁵¹

CONCLUSION

To summarize, a mixture of CsBr - PbBr_2 salts dissolved in DMSO and DMF results in the formation of different lead bromide species with a solvent-dependent solvodynamic radius of the species. Solutions in DMSO feature smaller species (~ 1 nm solvodynamic size) compared to the solutions in DMF, likely due to the stronger coordination of DMSO to Pb^{2+} compared to DMF. The size of lead bromide species influences the composition of the Cs - Pb - Br precipitates upon addition of the antisolvent: smaller species favor formation of the Cs_4PbBr_6 phase, while larger species template the precipitation of CsPbBr_3 . Overall, 0D Cs_4PbBr_6 has a higher tendency to be precipitated out from solutions with stronger coordinating solvents to Pb^{2+} , lower absolute concentration of the precursors and higher $\text{CsBr}:\text{PbBr}_2$ ratios, compared to its 3D CsPbBr_3 counterpart. We have discussed a possible origin of the green emission in the wide bandgap Cs_4PbBr_6 and concluded that 3D impurities might not be the only source of the emission and high PLQY. Alternatively, an impurity of 2D Cs_2PbBr_4 has been proposed but no conclusive experimental evidence of its existence was obtained. Our work highlights the importance of the

less investigated small cation spacers (e.g. Cs or MA) in 2D perovskites. Future work would target understanding the formation mechanism of Cs_2PbBr_4 as an impurity within the Cs_4PbBr_6 .

ASSOCIATED CONTENT

Supporting Information. The Supporting Information is available free of charge on the ACS Publications website. PLE, PL, and DLS of PbBr_2 -only solutions. Optical images, confocal PL microscopy volume view, and high-angle annular dark-field (HAADF) and elemental analysis of the 4:1_DMSO sample.

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The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Notes

The authors declare no competing financial interest.

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REFERENCES

- (1) Saidaminov, M. I.; Mohammed, O. F.; Bakr, O. M. Low-Dimensional-Networked Metal Halide Perovskites: The Next Big Thing. *ACS Energy Lett.* **2017**, *2*, 889-896.
- (2) Lin, H.; Zhou, C.; Tian, Y.; Siegrist, T.; Ma, B. Low-Dimensional Organometal Halide Perovskites. *ACS Energy Lett.* **2017**, 54-62.
- (3) Shi, E.; Gao, Y.; Finkenauer, B. P.; Akriti; Coffey, A. H.; Dou, L. Two-Dimensional Halide Perovskite Nanomaterials and Heterostructures. *Chem. Soc. Rev.* **2018**, *47*, 6046-6072.
- (4) Gao, P.; Bin Mohd Yusoff, A. R.; Nazeeruddin, M. K. Dimensionality Engineering of Hybrid Halide Perovskite Light Absorbers. *Nat. Commun.* **2018**, *9*, 5028.
- (5) Yuan, Z.; Zhou, C.; Tian, Y.; Shu, Y.; Messier, J.; Wang, J. C.; van de Burgt, L. J.; Kountouriotis, K.; Xin, Y.; Holt, E.; Schanze, K.; Clark, R.; Siegrist, T.; Ma, B. One-Dimensional Organic Lead Halide Perovskites with Efficient Bluish White-Light Emission. *Nat. Commun.* **2017**, *8*, 14051.
- (6) Liu, M.; Zhao, J. T.; Luo, Z. L.; Sun, Z. H.; Pan, N.; Ding, H. Y.; Wang, X. P. Unveiling Solvent-Related Effect on Phase Transformations in CsBr-PbBr₂ System: Coordination and Ratio of Precursors. *Chem. Mater.* **2018**, *30*, 5846-5852.
- (7) Zhao, Y.; Zhu, K. Organic-Inorganic Hybrid Lead Halide Perovskites for Optoelectronic and Electronic Applications. *Chem. Soc. Rev.* **2016**, *45*, 655-89.
- (8) Adinolfi, V.; Yuan, M.; Comin, R.; Thibau, E. S.; Shi, D.; Saidaminov, M. I.; Kanjanaboos, P.; Kopilovic, D.; Hoogland, S.; Lu, Z. H.; Bakr, O. M.; Sargent, E. H. The in-Gap Electronic State Spectrum of Methylammonium Lead Iodide Single-Crystal Perovskites. *Adv. Mater.* **2016**, *28*, 3406-10.
- (9) Cha, J. H.; Han, J. H.; Yin, W.; Park, C.; Park, Y.; Ahn, T. K.; Cho, J. H.; Jung, D. Y. Photoresponse of CsPbBr₃ and Cs₄PbBr₆ Perovskite Single Crystals. *J. Phys. Chem. Lett.* **2017**, *8*, 565-570.
- (10) Yin, J.; Zhang, Y.; Bruno, A.; Soci, C.; Bakr, O. M.; Brédas, J.-L.; Mohammed, O. F. Intrinsic Lead Ion Emissions in Zero-Dimensional Cs₄PbBr₆ Nanocrystals. *ACS Energy Lett.* **2017**, *2*, 2805-2811.
- (11) Zhang, Y.; Saidaminov, M. I.; Dursun, I.; Yang, H.; Murali, B.; Alarousu, E.; Yengel, E.; Alshankiti, B. A.; Bakr, O. M.; Mohammed, O. F. Zero-Dimensional Cs₄PbBr₆ Perovskite Nanocrystals. *J. Phys. Chem. Lett.* **2017**, *8*, 961-965.
- (12) Akkerman, Q. A.; Abdelhady, A. L.; Manna, L. Zero-Dimensional Cesium Lead Halides: History, Properties, and Challenges. *J. Phys. Chem. Lett.* **2018**, *9*, 2326-2337.
- (13) Yin, J.; Yang, H.; Song, K.; El-Zohry, A. M.; Han, Y.; Bakr, O. M.; Bredas, J. L.; Mohammed, O. F. Point Defects and Green Emission in Zero-Dimensional Perovskites. *J. Phys. Chem. Lett.* **2018**, *9*, 5490-5495.

- (14) Seth, S.; Samanta, A. Fluorescent Phase-Pure Zero-Dimensional Perovskite-Related Cs_4PbBr_6 Microdisks: Synthesis and Single-Particle Imaging Study. *J. Phys. Chem. Lett.* **2017**, *8*, 4461-4467.
- (15) De Bastiani, M.; Dursun, I.; Zhang, Y.; Alshankiti, B. A.; Miao, X.-H.; Yin, J.; Yengel, E.; Alarousu, E.; Turedi, B.; Almutlaq, J. M.; Saidaminov, M. I.; Mitra, S.; Gereige, I.; AlSaggaf, A.; Zhu, Y.; Han, Y.; Roqan, I. S.; Bredas, J.-L.; Mohammed, O. F.; Bakr, O. M. Inside Perovskites: Quantum Luminescence from Bulk Cs_4PbBr_6 Single Crystals. *Chem. Mater.* **2017**, *29*, 7108-7113.
- (16) Wells, H. L. Über Die Cäsium- Und Kalium-Bleihalogenide. *Zeitschrift für anorganische Chemie* **1893**, *3*, 195-210.
- (17) Nikl, M.; Mihokova, E.; Nitsch, K.; Somma, F.; Giampaolo, C.; Pazzi, G. P.; Fabeni, P.; Zazubovich, S. Photoluminescence of Cs_4PbBr_6 Crystals and Thin Films. *Chem. Phys. Lett.* **1999**, *306*, 280-284.
- (18) Kondo, S.; Amaya, K.; Saito, T. Localized Optical Absorption in Cs_4PbBr_6 . *J. Phys.: Condens. Matter* **2002**, *14*, 2093-2099.
- (19) Akkerman, Q. A.; Park, S.; Radicchi, E.; Nunzi, F.; Mosconi, E.; De Angelis, F.; Brescia, R.; Rastogi, P.; Prato, M.; Manna, L. Nearly Monodisperse Insulator Cs_4PbX_6 (X = Cl, Br, I) Nanocrystals, Their Mixed Halide Compositions, and Their Transformation into CsPbX_3 Nanocrystals. *Nano Lett.* **2017**, *17*, 1924-1930.
- (20) Quan, L. N.; Quintero-Bermudez, R.; Voznyy, O.; Walters, G.; Jain, A.; Fan, J. Z.; Zheng, X.; Yang, Z.; Sargent, E. H. Highly Emissive Green Perovskite Nanocrystals in a Solid State Crystalline Matrix. *Adv. Mater.* **2017**, *29*, 1605945.
- (21) Xu, J.; Huang, W.; Li, P.; Onken, D. R.; Dun, C.; Guo, Y.; Ucer, K. B.; Lu, C.; Wang, H.; Geyer, S. M.; Williams, R. T.; Carroll, D. L. Imbedded Nanocrystals of CsPbBr_3 in Cs_4PbBr_6 : Kinetics, Enhanced Oscillator Strength, and Application in Light-Emitting Diodes. *Adv. Mater.* **2017**, *29*, 1703703.
- (22) Chen, X.; Zhang, F.; Ge, Y.; Shi, L.; Huang, S.; Tang, J.; Lv, Z.; Zhang, L.; Zou, B.; Zhong, H. Centimeter-Sized Cs_4PbBr_6 Crystals with Embedded CsPbBr_3 Nanocrystals Showing Superior Photoluminescence: Nonstoichiometry Induced Transformation and Light-Emitting Applications. *Adv. Funct. Mater.* **2018**, 1706567.
- (23) Xuan, T.; Lou, S.; Huang, J.; Cao, L.; Yang, X.; Li, H.; Wang, J. Monodisperse and Brightly Luminescent $\text{CsPbBr}_3/\text{Cs}_4\text{PbBr}_6$ Perovskite Composite Nanocrystals. *Nanoscale* **2018**, *10*, 9840-9844.
- (24) Riesen, N.; Lockrey, M.; Badek, K.; Riesen, H. On the Origins of the Green Luminescence in the "Zero-Dimensional Perovskite" Cs_4PbBr_6 : Conclusive Results from Cathodoluminescence Imaging. *Nanoscale* **2019**, *11*, 3925-3932.
- (25) Ling, Y.; Tan, L.; Wang, X.; Zhou, Y.; Xin, Y.; Ma, B.; Hanson, K.; Gao, H. Composite Perovskites of Cesium Lead Bromide for Optimized Photoluminescence. *J. Phys. Chem. Lett.* **2017**, *8*, 3266-3271.
- (26) Baranov, D.; Toso, S.; Imran, M.; Manna, L. Investigation into the Photoluminescence Red Shift in Cesium Lead Bromide Nanocrystal Superlattices. *J. Phys. Chem. Lett.* **2019**, *10*, 655-660.
- (27) Yoon, S. J.; Stamplecoskie, K. G.; Kamat, P. V. How Lead Halide Complex Chemistry Dictates the Composition of Mixed Halide Perovskites. *J. Phys. Chem. Lett.* **2016**, *7*, 1368-73.

- (28) Oldenburg, K.; Vogler, A., Electronic Spectra and Photochemistry of Tin(II), Lead(II), Antimony(III), and Bismuth(III) Bromide Complexes in Solution. In *Zeitschrift für Naturforschung B*, 1993; Vol. 48, p 1519.
- (29) Rahimnejad, S.; Kovalenko, A.; Forés, S. M.; Aranda, C.; Guerrero, A. Coordination Chemistry Dictates the Structural Defects in Lead Halide Perovskites. *ChemPhysChem* **2016**, *17*, 2795-2798.
- (30) Hamill, J. C.; Schwartz, J.; Loo, Y.-L. Influence of Solvent Coordination on Hybrid Organic-Inorganic Perovskite Formation. *ACS Energy Lett.* **2018**, *3*, 92-97.
- (31) Wu, Y.; Islam, A.; Yang, X.; Qin, C.; Liu, J.; Zhang, K.; Peng, W.; Han, L. Retarding the Crystallization of PbI₂ for Highly Reproducible Planar-Structured Perovskite Solar Cells Via Sequential Deposition. *Energy Environ. Sci.* **2014**, *7*, 2934-2938.
- (32) Yan, K.; Long, M.; Zhang, T.; Wei, Z.; Chen, H.; Yang, S.; Xu, J. Hybrid Halide Perovskite Solar Cell Precursors: Colloidal Chemistry and Coordination Engineering Behind Device Processing for High Efficiency. *J. Am. Chem. Soc.* **2015**, *137*, 4460-4468.
- (33) McMeekin, D. P.; Wang, Z.; Rehman, W.; Pulvirenti, F.; Patel, J. B.; Noel, N. K.; Johnston, M. B.; Marder, S. R.; Herz, L. M.; Snaith, H. J. Crystallization Kinetics and Morphology Control of Formamidinium-Cesium Mixed-Cation Lead Mixed-Halide Perovskite Via Tunability of the Colloidal Precursor Solution. *Adv. Mater.* **2017**, *29*, 1607039.
- (34) Wei, Z.; Perumal, A.; Su, R.; Sushant, S.; Xing, J.; Zhang, Q.; Tan, S. T.; Demir, H. V.; Xiong, Q. Solution-Processed Highly Bright and Durable Cesium Lead Halide Perovskite Light-Emitting Diodes. *Nanoscale* **2016**, *8*, 18021-18026.
- (35) Stevenson, J.; Sorenson, B.; Subramaniam, V. H.; Raiford, J.; Khlyabich, P. P.; Loo, Y.-L.; Clancy, P. Mayer Bond Order as a Metric of Complexation Effectiveness in Lead Halide Perovskite Solutions. *Chem. Mater.* **2017**, *29*, 2435-2444.
- (36) Xu, Y.; Zhang, Q.; Lv, L.; Han, W.; Wu, G.; Yang, D.; Dong, A. Synthesis of Ultrasmall CsPbBr₃ Nanoclusters and Their Transformation to Highly Deep-Blue-Emitting Nanoribbons at Room Temperature. *Nanoscale* **2017**.
- (37) Bureau, B.; Silly, G.; Buzaré, J. Y. CP-MAS ²⁰⁷Pb with ¹⁹F Decoupling NMR Spectroscopy: Medium Range Investigation in Fluoride Materials. *Solid State Nucl. Magn. Reson.* **1999**, *15*, 79-89.
- (38) Rosales, B. A.; Men, L.; Cady, S. D.; Hanrahan, M. P.; Rossini, A. J.; Vela, J. Persistent Dopants and Phase Segregation in Organolead Mixed-Halide Perovskites. *Chem. Mater.* **2016**, *28*, 6848-6859.
- (39) Askar, A. M.; Bernard, G. M.; Wiltshire, B. D.; Shankar, K.; Michaelis, V. K. Multinuclear Magnetic Resonance Tracking of Hydro, Thermal and Hydrothermal Decomposition of CH₃NH₃PbI₃. *J. Phys. Chem. C* **2017**, *121*, 1013-1024.
- (40) Rosales, B. A.; Hanrahan, M. P.; Boote, B. W.; Rossini, A. J.; Smith, E. A.; Vela, J. Lead Halide Perovskites: Challenges and Opportunities in Advanced Synthesis and Spectroscopy. *ACS Energy Lett.* **2017**, *2*, 906-914.
- (41) Wrackmeyer, B.; Horchler, K., ²⁰⁷Pb-NMR Parameters. In *Annual Reports on NMR Spectroscopy*, Webb, G. A., Ed. Academic Press: 1990; Vol. 22, pp 249-306.
- (42) Hu, M.; Ge, C.; Yu, J.; Feng, J. Mechanical and Optical Properties of Cs₄BX₆ (B = Pb, Sn; X = Cl, Br, I) Zero-Dimension Perovskites. *J. Phys. Chem. C* **2017**, *121*, 27053-27058.
- (43) Wrackmeyer, B.; Horchler, K., ²⁰⁷Pb-NMR Parameters. Webb, G. A., Ed. Annual Reports on NMR Spectroscopy: 1990; Vol. 22, pp 249-306.

- (44) Yang, H.; Zhang, Y.; Pan, J.; Yin, J.; Bakr, O. M.; Mohammed, O. F. Room-Temperature Engineering of All-Inorganic Perovskite Nanocrystals with Different Dimensionalities. *Chem. Mater.* **2017**, *29*, 8978-8982.
- (45) Endres, J.; Egger, D. A.; Kulbak, M.; Kerner, R. A.; Zhao, L.; Silver, S. H.; Hodes, G.; Rand, B. P.; Cahen, D.; Kronik, L.; Kahn, A. Valence and Conduction Band Densities of States of Metal Halide Perovskites: A Combined Experimental–Theoretical Study. *J. Phys. Chem. Lett.* **2016**, *7*, 2722-2729.
- (46) Young-Kwang, J.; Joaquin, C.; Ji-Sang, P.; Sunghyun, K.; Lucy, W.; Aron, W., *Luminescence of Polybromide Defects in Cs₄PbBr₆*. ChemRxiv: 2019.
- (47) Almeida, G.; Goldoni, L.; Akkerman, Q.; Dang, Z.; Khan, A. H.; Marras, S.; Moreels, I.; Manna, L. Role of Acid-Base Equilibria in the Size, Shape, and Phase Control of Cesium Lead Bromide Nanocrystals. *ACS Nano* **2018**, *12*, 1704-1711.
- (48) Wang, K.-H.; Yang, J.-N.; Ni, Q.-K.; Yao, H.-B.; Yu, S.-H. Metal Halide Perovskite Supercrystals: Gold–Bromide Complex Triggered Assembly of CsPbBr₃ Nanocubes. *Langmuir* **2018**, *34*, 595-602.
- (49) Jizhong, S.; Leimeng, X.; Jianhai, L.; Jie, X.; Yuhui, D.; Xiaoming, L.; Haibo, Z. Monolayer and Few-Layer All-Inorganic Perovskites as a New Family of Two-Dimensional Semiconductors for Printable Optoelectronic Devices. *Adv. Mater.* **2016**, *28*, 4861-4869.
- (50) Li, J.; Yu, Q.; He, Y.; Stoumpos, C. C.; Niu, G.; Trimarchi, G. G.; Guo, H.; Dong, G.; Wang, D.; Wang, L.; Kanatzidis, M. G. Cs₂PbI₂Cl₂, All-Inorganic Two-Dimensional Ruddlesden-Popper Mixed Halide Perovskite with Optoelectronic Response. *J. Am. Chem. Soc.* **2018**, *140*, 11085-11090.
- (51) Yu, Y.; Zhang, D.; Yang, P. Ruddlesden–Popper Phase in Two-Dimensional Inorganic Halide Perovskites: A Plausible Model and the Supporting Observations. *Nano Lett.* **2017**, *17*, 5489-5494.
- (52) Yang, J. H.; Yuan, Q. H.; Yakobson, B. I. Chemical Trends of Electronic Properties of Two-Dimensional Halide Perovskites and Their Potential Applications for Electronics and Optoelectronics. *J. Phys. Chem. C* **2016**, *120*, 24682-24687.
- (53) Zhu, X.; Xu, Z.; Zuo, S.; Feng, J.; Wang, Z.; Zhang, X.; Zhao, K.; Zhang, J.; Liu, H.; Priya, S.; Liu, S. F.; Yang, D. Vapor-Fumigation for Record Efficiency Two-Dimensional Perovskite Solar Cells with Superior Stability. *Energy Environ. Sci.* **2018**, *11*, 3349-3357.
- (54) Stoumpos, C. C.; Cao, D. H.; Clark, D. J.; Young, J.; Rondinelli, J. M.; Jang, J. I.; Hupp, J. T.; Kanatzidis, M. G. Ruddlesden–Popper Hybrid Lead Iodide Perovskite 2D Homologous Semiconductors. *Chem. Mater.* **2016**, *28*, 2852-2867.
- (55) Dou, L.; Wong, A. B.; Yu, Y.; Lai, M.; Kornienko, N.; Eaton, S. W.; Fu, A.; Bischak, C. G.; Ma, J.; Ding, T.; Ginsberg, N. S.; Wang, L.-W.; Alivisatos, A. P.; Yang, P. Atomically Thin Two-Dimensional Organic-Inorganic Hybrid Perovskites. *Science* **2015**, *349*, 1518-1521.
- (56) Yin, H.; Jin, L.; Qian, Y.; Li, X.; Wu, Y.; Bowen, M. S.; Kaan, D.; He, C.; Wozniak, D. I.; Xu, B.; Lewis, A. J.; Shen, W.; Chen, K.; Dobereiner, G. E.; Zhao, Y.; Wayland, B. B.; Rao, Y. Excitonic and Confinement Effects of 2D Layered (C₁₀H₂₁NH₃)₂PbBr₄ Single Crystals. *ACS Appl. Energy Mater.* **2018**, *1*, 1476-1482.
- (57) Almutlaq, J. M.; Yin, J.; Mohammed, O. F.; Bakr, O. M. The Benefit and Challenges of Zero-Dimensional Perovskites. *J. Phys. Chem. Lett.* **2018**, *9*, 4131-4138.
- (58) Akkerman, Q. A.; Motti, S. G.; Srimath Kandada, A. R.; Mosconi, E.; D'Innocenzo, V.; Bertoni, G.; Marras, S.; Kamino, B. A.; Miranda, L.; De Angelis, F.; Petrozza, A.; Prato, M.;

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Manna, L. Solution Synthesis Approach to Colloidal Cesium Lead Halide Perovskite Nanoplatelets with Monolayer-Level Thickness Control. *J. Am. Chem. Soc.* **2016**, *138*, 1010-6.

(59) Shamsi, J.; Dang, Z.; Bianchini, P.; Canale, C.; Stasio, F. D.; Brescia, R.; Prato, M.; Manna, L. Colloidal Synthesis of Quantum Confined Single Crystal CsPbBr₃ Nanosheets with Lateral Size Control up to the Micrometer Range. *J. Am. Chem. Soc.* **2016**, *138*, 7240-3.

(60) Liu, Z.; Bekenstein, Y.; Ye, X.; Nguyen, S. C.; Swabeck, J.; Zhang, D.; Lee, S. T.; Yang, P.; Ma, W.; Alivisatos, A. P. Ligand Mediated Transformation of Cesium Lead Bromide Perovskite Nanocrystals to Lead Depleted Cs₄PbBr₆ Nanocrystals. *J. Am. Chem. Soc.* **2017**, *139*, 5309-5312.