

Cite this: *J. Mater. Chem. C*, 2025, 13, 4039

Room-temperature synthesis of highly luminescent methylammonium lead bromide nanocubes encapsulated in block copolymer micelles: impact of solvent choice on crystallization and stability†

Belda Amelia Junisu,  Ya-Sen Sun * and Bo-Cheng Zhao

Herein, we demonstrate a facile, room-temperature method for synthesizing highly luminescent methylammonium lead bromide (MAPbBr₃) nanocubes encapsulated within colloidal polystyrene-*block*-poly(2-vinylpyridine) (PS-*b*-P2VP) micelles in a non-polar 1,3,5-trimethylbenzene (TMB) solvent. The synthesis involves five distinct stages: micellization of the PS-*b*-P2VP block copolymer, dissolution and complexation of PbBr₂ precursors, coordination between bromoplumbate complexes and P2VP segments within the micelle core, multiple emulsion, and confined crystallization of perovskite nanocubes. Unlike conventional ligand-assisted methods, the colloidal micelles act as soft nanoreactors by controlling the nucleation and growth of the nanocubes through multiple emulsion. Moreover, the micelles serve as colloidal templates, preserving a significant concentration of [PbBr₃][−] ions formed during the complexation stage. We also demonstrate that the choice of solvent for transporting MA⁺ and Br[−] ions from the TMB phase to the micelle core significantly influences the dimensions of the resulting MAPbBr₃ perovskite. Utilizing methanol as the transport medium yields encapsulated cubic MAPbBr₃ nanocubes with a high photoluminescence quantum yield (PLQY) of ~77%, and these encapsulated nanocubes exhibit long-term stability with a superior PLQY of ~88%. In contrast, employing dimethylformamide (DMF) to solvate MA⁺ cations mainly yields MAPbBr₃ microcrystals with reduced PLQY, as the formation of encapsulated MAPbBr₃ nanocubes is hindered. This hindrance is attributed to the high entropic penalty associated with the diffusion and penetration of DMF-solvated MA⁺ cations through the PS shells. We believe that this versatile approach can be extended to synthesize a wide range of nanomaterials with well-defined morphologies, monodisperse sizes and long-term stability.

Received 10th September 2024,
Accepted 5th January 2025

DOI: 10.1039/d4tc03885a

rsc.li/materials-c

Department of Chemical Engineering, National Cheng Kung University, Tainan 701, Taiwan. E-mail: yssun@gs.ncku.edu.tw

† Electronic supplementary information (ESI) available: TEM image, EDS 2D maps, and EDS 1D profiles of the dried state of t-LHP₇ solution (Fig. S1); photos, UV-vis, and PL spectra of c-PER_{M20} solutions at various centrifugation speeds (Fig. S2); photos and PLQY of the c-PER_M solutions after being stored in an ambient environment for 3 months (Fig. S3); TEM image and EDS 1D profiles of the c-PER_{D20} solution in dried state (Fig. S4); UV-vis absorbance and PL spectra of fresh c-PER_{c7,M20} and c-PER_{c7,D20} solutions (Fig. S5); HR-TEM image and EDS 1D profiles of the fresh c-PER_{c7,M20} solution in dried state (Fig. S6); UV-vis absorbance and TEM image of c-PER_{c7,M20} solution with prolonged storage (Fig. S7); photos of solutions during the synthesis of perovskite in the absence of PS-*b*-P2VP (Fig. S8); characterization of solutions and its precipitates during the synthesis of perovskite without MABr dissolution (Fig. S9). See DOI: <https://doi.org/10.1039/d4tc03885a>

Introduction

Perovskites have received significant attention in recent years as promising materials across various fields. Their unique and flexible compositions enable precise tuning of optoelectronic properties, making them attractive for applications such as photovoltaics,^{1,2} light-emitting devices,^{3,4} and photodetectors.⁵ The desirable optical and electronic properties of hybrid perovskites can be attributed to their ABX₃ crystal structure, where “A” represents a monovalent cation, “B” is a bivalent cation, and “X” represents halide ions.^{6,7} While inorganic perovskites (*e.g.*, CsPbBr₃, CsPbI₃) offer superior thermal and environmental stability along with excellent optoelectronic properties,^{8–11} hybrid organic–inorganic perovskites (HOIPs) (*e.g.*, MAPbBr₃, MAPbI₃) provide a balance of good performance, easier processing, and greater flexibility at a lower capital cost.^{12,13}



Despite the promising properties of perovskite materials, challenges remain in their synthesis and processing. The most critical challenge is their stability under ambient conditions.^{14,15} In particular, the stability of perovskite nanocrystals (NCs) can be influenced by intrinsic factors such as morphology and size distribution. Numerous studies have shown the difficulty to obtain perovskite NCs with well-defined morphologies and monodisperse size distribution *via* solution processing.^{16,17} Perovskite NCs with different morphologies (such as nanocubes, nanoplatelets, and irregular shapes) and sizes, exhibit varying degrees of stability.^{18,19} Moreover, the size of the perovskite NCs is also crucial. Smaller nanocrystals (*e.g.* 4 nm) tend to be less stable compared to larger ones due to increased surface chemistry and reactivity.^{20,21}

Many bottom-up approaches for the synthesis of perovskite NCs employ ligands to control growth kinetics and prevent agglomeration.^{22–24} However, these ligands (*e.g.*, oleic acid, octylamine) mainly act as stabilizers without controlling morphology. Block copolymers (BCPs) represent a compelling alternative, capable of providing both stabilization and morphology control. In particular, BCP self-assembly can generate well-defined nanostructures that serve as templates or nano-reactors for the growth of perovskite NCs.²⁵ The thermodynamic incompatibility of the constituent polymer blocks drives microphase separation, leading to the formation of myriad ordered nanostructures.

By controlling the BCP composition, molecular weight, and solution processing conditions, a diverse array of morphologies can be obtained and offer soft colloidal templates to mediate the synthesis of inorganic nanomaterials.^{26–28} Polystyrene-*block*-poly(2-vinylpyridine) (PS-*b*-P2VP) BCPs have emerged as promising templates for enhancing the stability of perovskite NCs in both colloidal solutions^{29–32} and thin films.^{33–35} The efficient coordination between the P2VP block and metal precursor species promotes selective nucleation within the P2VP cores. For instance, Hou *et al.*³⁰ and Lee *et al.*³¹ successfully demonstrated the encapsulation of CsPbBr₃ perovskite NCs within PS-*b*-P2VP core-shell micelles, achieving remarkable stability against degradation in various polar solvents. Similarly, Hui *et al.*²⁹ utilized PS-*b*-P2VP micelles to synthesize MAPbX₃ nanoparticles exhibiting enhanced thermal stability, while Nah *et al.*³² further corroborated the efficacy of micelle encapsulation in producing stable MAPbX₃ perovskite NCs. However, a recurring limitation in these studies is the formation of micelles with less compact architectures, often accompanied by limited control over nanocrystal shape. Such non-uniformity increases the prevalence of defects and lattice distortions, ultimately leading to non-radiative recombination and reduced photoluminescence quantum yield (PLQY). Therefore, achieving monodisperse perovskite NCs with uniform shape is crucial for enhancing its PLQY.

Ligand-assisted re-precipitation (LARP) methods have been known as a more cost-effective alternative to produce perovskite NCs at room temperature.^{13,21} LARP leverages the principle of supersaturated recrystallization, whereby the solubility of a

highly saturated precursor solution is rapidly decreased—either by cooling, solvent evaporation, or the addition of a poor solvent (*i.e.*, an antisolvent, usually polar solvents).³⁶ This non-equilibrium condition induces rapid precipitation and crystallization until equilibrium is re-established. BCP colloidal templates function analogously to LARP by providing physical and chemical confinement for perovskite NCs. However, unlike LARP, BCP colloidal templates do not necessitate polar solvents for nanocrystal purification, thereby circumventing potential degradation and emission quenching associated with such polar solvents.²⁵

The growth of perovskite nanocrystals within BCPs is a multifaceted process involving several key steps: micellization of the BCP, solvation and salt complexation of precursors, coordination interactions between precursor species and BCP segments, and crystallization.^{29–31} Solvent quality significantly influences each of these steps. Notably, poor solvent quality can lead to collapsed or less organized BCP conformations. Furthermore, sufficient solvation is critical to ensure the formation of precursor salt complexes. For example, precursor complexes are typically produced by solvating PbX₂ precursors in an octadecene (ODE)^{37–39} or dimethylformamide (DMF) solvent.⁴⁰ In the initial stage of synthesis, precursor complexes should selectively interact with BCP active groups. For instance, coordination interactions between precursor species and pyridine groups in P2VP blocks are often driven by Lewis acid–base interactions. The lone pair of electrons on the nitrogen atoms of quarternized P2VP blocks can delocalize to the empty 6p-orbital of Pb-complex species.⁴¹ As the concentration of coordinated precursors increases, supersaturation conditions are eventually reached.⁴² Under such condition, rapid crystallization produces perovskite NCs encapsulated within micelle templates.^{29–31}

Despite the potential of BCP colloidal micelles for templated synthesis of hybrid organic–inorganic perovskite NCs, a comprehensive understanding of the underlying mechanisms remains elusive. This study aims to systematically understand how soft colloidal templating of using PS-*b*-P2VP micelles mediated the synthesis of encapsulated methylammonium lead bromide (MAPbBr₃) perovskite nanocrystals (NCs) in a non-polar solvent (1,3,5-trimethylbenzene). The synthesis of encapsulated perovskite NCs involved PS-*b*-P2VP micellization, formation of hierarchical emulsion, solvation and complexation of precursor salts, coordination interactions between PS-*b*-P2VP and salt ions, and nucleation and growth of perovskite crystals in a confined space. Furthermore, the influence of two different polar solvents (methanol and DMF) on MABr solvation and crystallization kinetics of the resulting MAPbBr₃ perovskite NCs were further examined.

Experimental section

Materials

Two polystyrene-*block*-poly(2-vinylpyridine) block copolymers, PS_{48.5}-*b*-P2VP₇₀ ($M_{n,PS} = 48.5 \text{ kg mol}^{-1}$, $M_{n,P2VP} = 70 \text{ kg mol}^{-1}$,



$M_w/M_n = 1.13$) and PS₄₉-*b*-P2VP₇₅ ($M_{n,PS} = 49 \text{ kg mol}^{-1}$, $M_{n,P2VP} = 75 \text{ kg mol}^{-1}$, $M_w/M_n = 1.03$), were purchased from Polymer Source Inc. PS₄₉-*b*-P2VP₇₅ was used as ligands for Fig. 1–7 and Fig. S2–S9 (ESI[†]). PS_{48.5}-*b*-P2VP₇₀ was used as ligands for Fig. S1 (ESI[†]). These two block copolymers have similar molecular weights and compositions. Therefore, they have the same micellization behavior in TMB. Lead bromide (PbBr₂) and methylammonium bromide (MABr) precursor were purchased from Sigma Aldrich and used as received. 1,3,5-Trimethylbenzene (TMB, C₆H₃(CH₃)₃) was purchased from Thermo Scientific. Methanol (CH₃OH) was purchased from Honeywell. Dimethyl formamide (DMF) was purchased from Sigma Aldrich.

Preparation of precursor solutions

The experimental procedures used in this study are depicted in Scheme 1. Perovskite NCs within BCP micelles were synthesized through three steps: micellization, precursor complexation, and confined crystallization. First, 0.5 wt% BCP colloidal solutions were prepared by dissolving 4.3 mg of PS-*b*-P2VP powder in 1 mL of 1,3,5-trimethylbenzene (TMB). The solutions were sonicated at 70 °C for 30 minutes and then cooled to room temperature. To ensure complete micellization, the BCP colloidal solutions were aged overnight. Next, precursor solutions were prepared by dissolving 10 mg of PbBr₂ in the block copolymer solutions. These precursor solutions were stirred at 800 rpm for 24 hours using a magnetic stirrer. The resulting turbid lead-halide precursor solutions were labeled as t-LHP solutions (Scheme 1B).

Preparation of MABr solutions

Due to MABr low solubility in TMB, MABr solutions were separately prepared in either methanol or DMF. These two solvents were selected for two reasons. First, MABr is highly soluble in methanol and DMF, ensuring complete ionization to produce MA⁺ and Br⁻ ions. Second, the high polarity index of both solvents (methanol = 5.1, DMF = 6.4)⁴³ facilitates the effective transport of ions from the polar solvent (methanol or DMF) to the P2VP core through the non-polar TMB phase. Specifically, 60 mg of MABr powder was dissolved in 200 μL of either methanol or DMF to produce 300 mg mL⁻¹ MABr solutions. The mixtures were then stirred for 30 minutes at 50 °C to ensure complete dissolution.

Synthesis of perovskite NCs

The final step involved the confined crystallization of MAPbBr₃ perovskite NCs within the BCP micelles. To initiate crystallization, aliquots (5–20 μL) of the MABr solutions were added dropwise to the t-LHP solutions. The mixtures were stirred continuously at 800 rpm for 30 minutes for the nucleation and growth of MAPbBr₃ nanocubes within the P2VP cores. The formation of MAPbBr₃ NCs was evidenced by a color change from white in the precursor solutions to yellow in the perovskite solutions (Scheme 1C). The resulting perovskite solutions were then centrifuged at 7000 rpm to separate the encapsulated MAPbBr₃ nanocubes from microstructures. The encapsulated MAPbBr₃ nanocubes were dispersed in supernatants (c-PER, Scheme 1D), while the microparticles existed as precipitates. The clear supernatants (briefly denoted as c-PER_{M5}, c-PER_{M10}

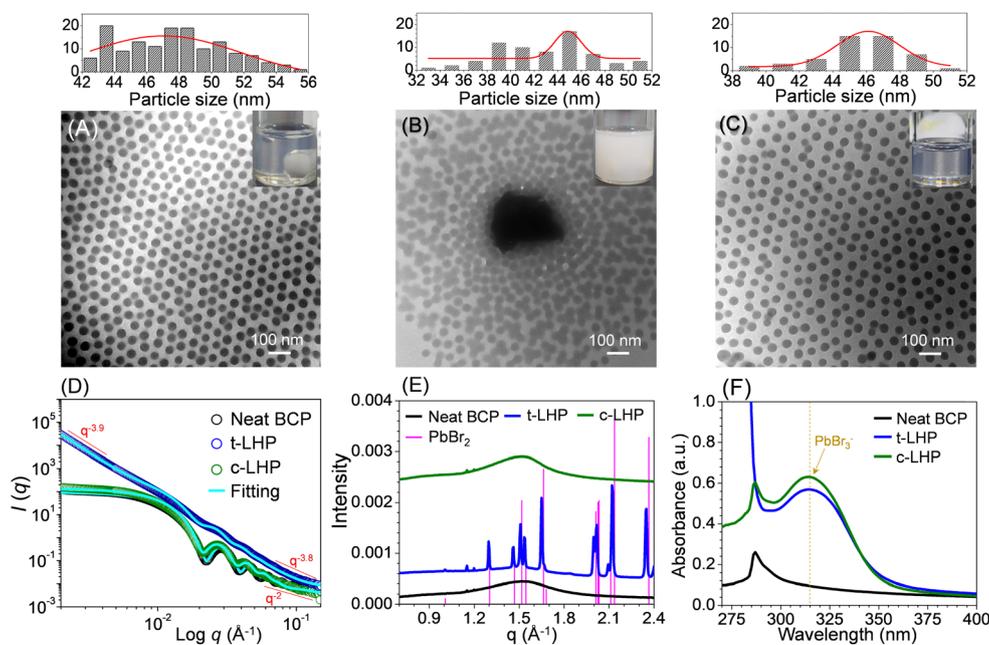


Fig. 1 (A)–(C) TEM images and corresponding (D) SAXS (symbols: experimental data and cyan lines: fitted data), (E) WAXD, and (F) absorbance profiles of BCP, t-LHP and c-LHP solutions. In (D), red lines at low- q and high- q regions that represent slope are plotted as a visual guide. Micelle size distributions from images (A)–(C) are shown in the upper panels. Inset in images (A)–(C) show photographs of the BCP, t-LHP and c-LHP solutions, respectively.



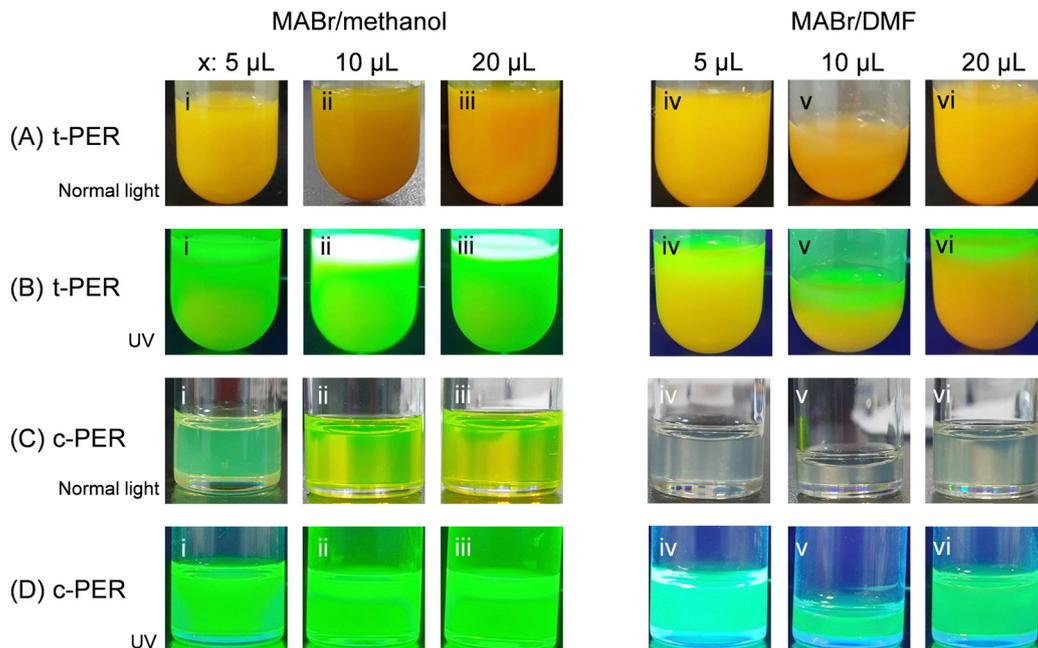


Fig. 2 Photos of ((A) and (B)) t-PER and ((C) and (D)) c-PER solutions. The solutions were prepared by adding different aliquots [(i, iv) 5 μL , (ii, v) 10 μL , and (iii, vi) 20 μL] of methanol-solvated (left panel) or DMF-solvated (right panel) MABr solutions into t-LHP solutions. Photos in (B) and (D) were taken under UV light exposure at $\lambda = 365$ nm.

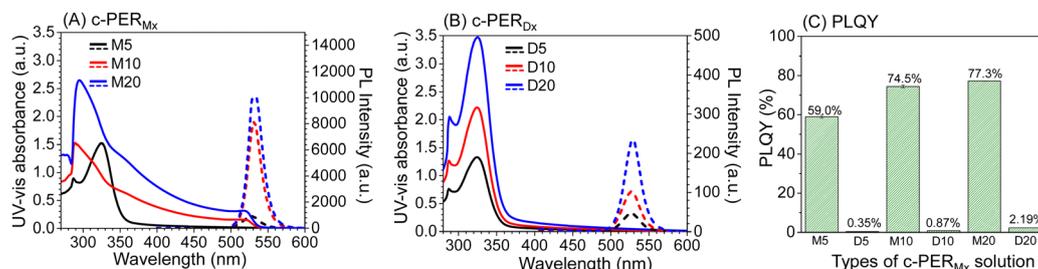


Fig. 3 UV-vis absorbance (solid lines) and PL (dashed lines) spectra of (A) c-PER_{Mx} and (B) c-PER_{Dx} solutions (x: 5, 10, and 20). (C) PLQY of the corresponding solutions described in (A) and (B). For UV-vis and PL measurements, the original solutions were diluted to a ratio of 1 : 10. For PLQY measurements, the dilution factor was 1 : 30.

and c-PER_{M20}) and precipitates were collected separately for further structural characterization. The same procedures were performed on c-PER_{D5}, c-PER_{D10} and c-PER_{D20} solutions that were prepared by adding 5, 10 or 20 μL of MABr aliquots in DMF to the t-LHP solutions, followed by centrifugation. The subscripts denote the used solvents (M: methanol; D: DMF) and the amount of added MABr aliquots. All solutions were prepared at room temperature.

We also prepared an additional set of BCP colloidal solutions to investigate the effect of prior centrifugation of the t-LHP solutions on the complexation and crystallization of perovskite NCs. The procedures for preparing the t-LHP solutions were identical to those described above, except that these solutions were stirred continuously for 7 days to produce t-LHP₇ solutions, where the subscript denotes the stirring time in days if specified (Scheme 1B'). After prolonged

stirring, the t-LHP₇ solutions were centrifuged at 7000 rpm for 10 minutes to remove undissolved PbBr₂ crystals, producing clear supernatant precursor solutions (denoted as c-LHP₇, Scheme 1C').

The centrifuged c-LHP₇ solutions were used for the synthesis of perovskite NCs. Specifically, 20 μL of either the MABr/methanol or MABr/DMF precursor solution was added dropwise to the stirred c-LHP₇ solutions. After stirring for 30 minutes at 800 rpm, the resulting solutions were labelled as t-PER_{c7} (Scheme 1D'). Here, the subscript "c" indicates prior centrifugation of the t-LHP₇ solution. These solutions were then centrifuged to separate the encapsulated perovskite NCs from unreacted precursors and by-products. The supernatants (c-PER_{c7}) and precipitates were collected separately for further characterization. A list of abbreviations in this study is shown in Table 1.



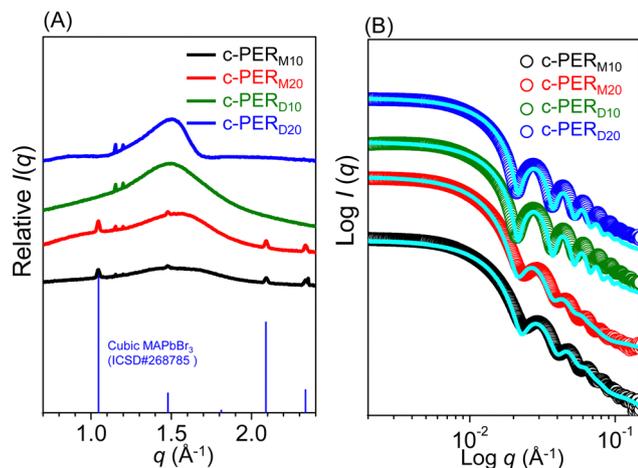


Fig. 4 (A) WAXD and (B) SAXS (symbols: experimental data; lines: fits) profiles of the c-PER_{M_x} and c-PER_{D_x} solutions (x: 10 and 20). Stick pattern in (A): cubic MAPbBr₃ crystals (ICSD#26785). The WAXD and SAXS curves were vertically shifted for better visualization.

Material characterization and instruments

UV-vis absorption spectra were measured with a JASCO V-770 UV-vis/near-IR spectrophotometer (JASCO Analytical Instruments), equipped with both a halogen and deuterium lamp. The fluorescence of the perovskite solutions was probed using a UV lamp (UVGL-25 4W compact UV lamp, $\lambda = 365$ nm). Photoluminescence (PL) spectra at room temperature were measured using an F-2500 FL Spectrophotometer (Hitachi), equipped with an Xe lamp ($\lambda_{\text{exc.}}$: 375 nm). Prior to UV-vis and PL measurements, all perovskite solutions were diluted to a ratio of 1 : 10 in TMB to accommodate detector resolution. PLQY were determined using an absolute PLQY measurement system (FluoroMax[®] spectrometer, HORIBA Scientific). Prior to the PLQY

measurements, all samples were diluted to a ratio of 1 : 30 in TMB.

Ten microliters of solutions were deposited on a carbon film-coated Cu grid. The grids were dried in an oven at 50 °C overnight before imaging. Morphologies of encapsulated perovskite nanocubes were characterized by low-resolution transmission electron microscopy (TEM) (H-7500) at National Cheng Kung University (NCKU), Taiwan. The low-resolution TEM was operated at 80 kV. High-resolution TEM (HR-TEM) was performed with a JEOL JEM-1400 (120 kV) at the NCKU Instrument Center, Taiwan. The analysis of elemental composition in perovskite nanocubes was performed using energy dispersive X-ray spectroscopy (EDS) attached to the HR-TEM. The elemental composition of N, Pb, and Br was quantified in each point scan. X-ray diffraction of precipitates was performed using Cu_{K α} radiation ($\lambda = 0.154$ nm) on a Rigaku Ultima IV-9407F701 X-ray diffractometer at 40 kV. Precipitates were carefully transferred from centrifuge tubes to glass substrates, which were then dried in an oven at 70 °C overnight. The 2θ scanning range was from 10° to 60° with a step size of 0.02° and a scan rate of 5° min⁻¹.

Simultaneous small-angle X-ray scattering (SAXS) and wide-angle X-ray diffraction (WAXD) measurements of the BCP/perovskite solutions were conducted at the TPS 13A end station of the National Synchrotron Radiation Research Center (NSRRC, Hsinchu, Taiwan).⁴⁴ Data were collected at an energy of 15 keV ($\lambda = 0.827$ Å) using two detectors: Eiger X 9 M for SAXS and Eiger X 1 M for WAXD. Silver behenate and polyethylene standards were used to calibrate the sample-to-detector distance and absolute SAXS intensity, respectively. Solutions were sealed in capillary tubes ($\Phi = 2$ mm) and mounted on a modified Linkam force stage for measurements. Background subtraction was performed using scattering data from empty capillary tubes and tubes containing neat solvents or

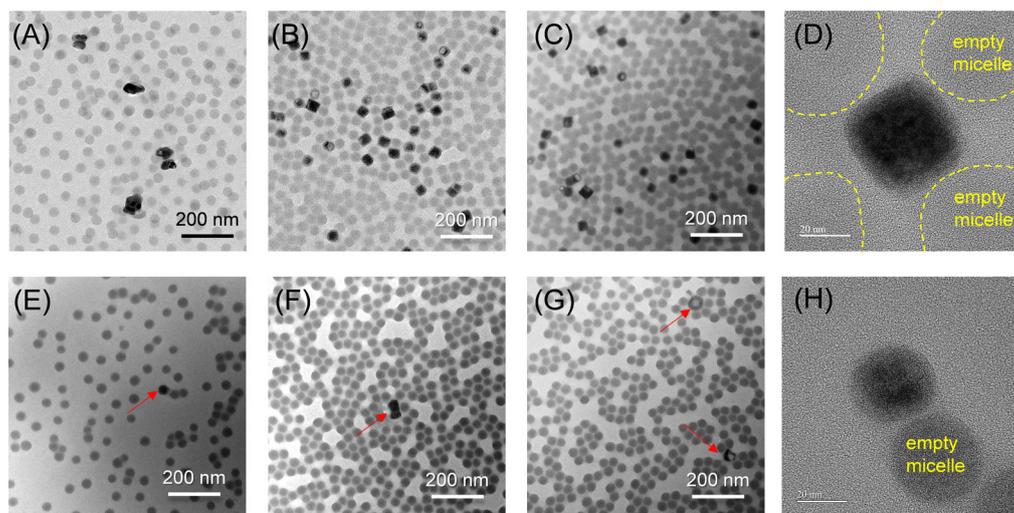


Fig. 5 TEM images collected in a dried state for the (A) c-PER_{M5}, (B) c-PER_{M10}, (C) c-PER_{M20}, (E) c-PER_{D5}, (F) c-PER_{D10} and (G) c-PER_{D20} solutions, respectively. Images (D) and (H) are the HR-TEM images of selected nanocube in c-PER_{M10} and nanocrystal in c-PER_{D20}, respectively. Yellow dashed lines in (D) indicate the periphery of micelles.



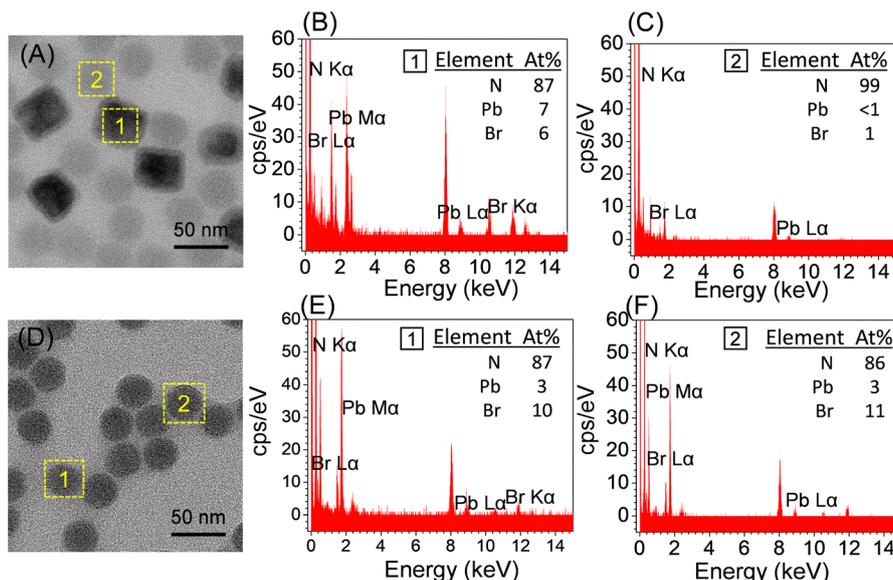


Fig. 6 ((A) and (D)) HR-TEM and ((B) and (C), (E) and (F)) EDS elemental analysis of dried c-PER_{M10} and c-PER_{D20} solutions. EDS spectra were selectively collected on two different positions of each specimen by scanning N, Pb, and Br elements. The positions are labelled with 1 and 2 in images (A) and (D), respectively.

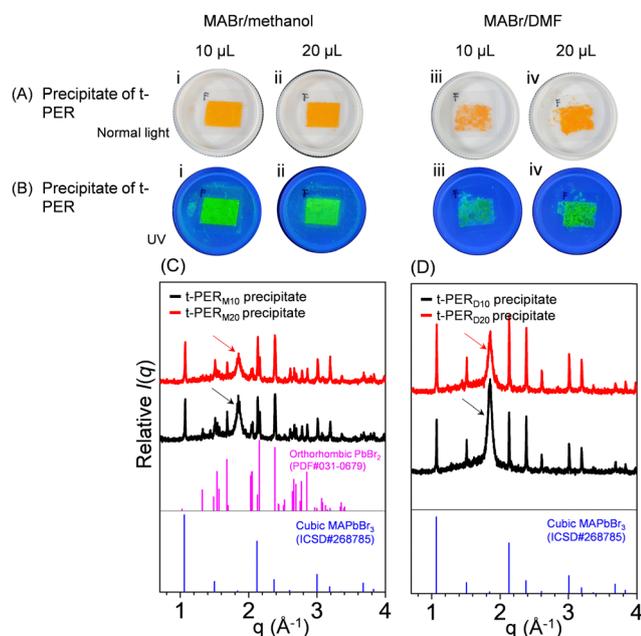


Fig. 7 ((A) and (B)) Photos and ((C) and (D)) WAXD profiles of dried precipitates separated from the (i) t-PER_{M10}, (ii) t-PER_{M20}, (iii) t-PER_{D10}, and (iv) t-PER_{D20} solutions after centrifugation at 7000 rpm for 10 min. Photos in (B) were taken under UV light exposure at $\lambda = 365$ nm. In (C) and (D), red and black arrows indicate the diffraction from the glass substrate. Stick patterns represent standard WAXD profiles of orthorhombic PbBr₂ crystals (PDF#031-0679) and cubic MAPbBr₃ crystals (ICSD#268785).

solvent/methanol mixtures. An exposure time of 10 s was used for all measurements. One-dimensional SAXS profiles were obtained from data reduction. Absolute intensity ($I(q)$), as a function of the scattering vector (q), was used to describe

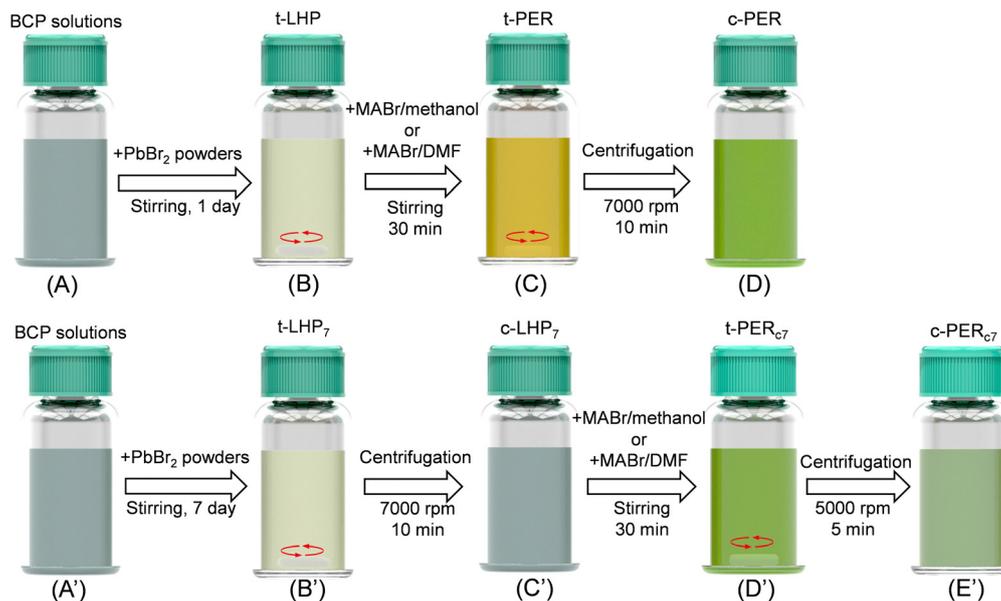
scattering features. q is defined as $q = \frac{4\pi}{\lambda} \sin \theta$, where λ is the X-ray wavelength and 2θ is the scattering angle. Structural parameters were determined by fitting the SAXS data using SASview software.⁴⁵ Two models were used: the two-level Beaucage model and the polydisperse core-shell sphere model.^{46–48} Based on the Beaucage model,⁴⁶ SAXS fitting quantifies hierarchical structures, with each level having structural parameters defined by the radius of gyration ($R_{g,i}$) and power-law exponent ($d_{i,i}$), where the subscript “ i ” represents the i -th level structure. $R_{g,i}$ indicates the size of the irregular structure, while $d_{i,i}$ reflects its dimensionality. For the polydisperse core-shell sphere model, SAXS fitting provides structural parameters, including core radius (r_c), shell thickness (r_s), size dispersity, and the scattering length density (ρ_c , ρ_s , and ρ_{sol}) of the core, shell, and solvent, respectively. Of the structural parameters, size dispersity is based on the Schultz size distribution function.⁴⁵

Results and discussion

Characterization of the BCP, t-LHP and c-LHP solutions

The synthesis of encapsulated perovskite nanocrystals (NCs) involves three stages in sequence: (1) PS-*b*-P2VP micellization, (2) precursor complexation in multiple emulsion, and (3) confined crystallization. The micellization of block copolymers (BCP) is governed by the interplay between the solubility parameters of the solvent and the constituent blocks. PS-*b*-P2VP exhibits strong micellization in TMB due to the selective solvation of the PS blocks. This selectivity arises because the solubility parameter of TMB ($\delta_{\text{TMB}} = 18 \text{ MPa}^{1/2}$)⁴⁹ is lower than





Scheme 1 Procedures of colloidal templated synthesis of perovskite nanoparticles in a solution added with PS-*b*-P2VP micelles. First set of samples were divided by four stages: (A) BCP solutions, (B) t-LHP solutions, (C) t-PER solutions and (D) c-PER solutions. Second set of samples were divided by five stages: (A') BCP solutions, (B') t-LHP₇ solutions, (C') c-LHP₇ solutions, (D') t-PER_{c7} solutions, and (E') c-PER_{c7} solutions. Note that white objects in (B), (C), (B'), and (D') represent the magnetic stirrer.

Table 1 List of abbreviations

Abbreviation	Description
t-LHP	Turbid lead halide precursor solution (consists of BCP colloidal micelles and PbBr ₂) after 1 day of stirring
t-PER	Turbid perovskite solution after the addition of MABr aliquots
c-PER _{Mx}	Clear perovskite solution after subsequent centrifugation of t-PER solution synthesized by the addition of MABr/methanol aliquots to the t-LHP solution ($x = 5, 10, 20$)
c-PER _{Dx}	Clear perovskite solution after subsequent centrifugation of t-PER solution synthesized by the addition of MABr/DMF aliquots to the t-LHP solution ($x = 5, 10, 20$)
t-LHP ₇	Turbid lead halide precursor solution after 7 days of stirring
c-LHP ₇	Clear lead halide precursor solution after 7 days of stirring and subsequent centrifugation
t-PER _{c7}	Turbid perovskite solution after the addition of MABr aliquots to the c-LHP ₇ solution
c-PER _{c7,Mx}	Clear perovskite solution after subsequent centrifugation of t-PER _{c7} solution synthesized by the addition of MABr/methanol aliquot to the c-LHP ₇ solution ($x = 20$)
c-PER _{c7,Dx}	Clear perovskite solution after subsequent centrifugation of t-PER _{c7} solution synthesized by the addition of MABr/DMF aliquot to the c-LHP ₇ solution ($x = 20$)

those of both PS ($\delta_{PS} = 18.6 \text{ MPa}^{1/2}$)⁵⁰ and P2VP ($\delta_{P2VP} = 19.8 \text{ MPa}^{1/2}$).⁵¹

To quantitatively analyze the micellar structure, TEM and SAXS measurements were performed on dry films and original BCP solutions in TMB, respectively. The neat BCP solution appeared bluish due to the Rayleigh scattering of micellar nanostructures⁵² (inset of Fig. 1A). TEM image and size distribution analysis in Fig. 1A reveals well-defined dark spheres with an average diameter of 47 nm. The SAXS profile (the black curve in Fig. 1D) of the neat BCP solution exhibits a Guinier regime in the low- q region ($q < 0.008 \text{ \AA}^{-1}$), followed by form factor oscillations in the intermediate- q region ($0.008\text{--}0.08 \text{ \AA}^{-1}$). The oscillations locate at $0.0287, 0.0456, 0.0614$ and 0.0794 \AA^{-1} , respectively. In the high- q region ($q > 0.08 \text{ \AA}^{-1}$), a power-law intensity decay of $I \sim q^{-2}$ is observed. This intensity decay indicates polymer brushes forming within the micelle corona.⁴⁸

The black SAXS curve was fitted using the polydisperse core-shell spherical model to extract structural parameters for micelles in the BCP solution. The structural parameters are summarized in Table 2. A core radius of 19.59 nm and a shell thickness of $10.74 \pm 0.09 \text{ nm}$ were obtained from the fitting. The WAXD profile of the neat BCP solution shows a broad amorphous halo due to inter-chain or intra-chain correlations (the black curve in Fig. 1E).

Next, we characterized a t-LHP solution with visual examination and TEM. The t-LHP solution appeared turbid (the insert in Fig. 1B). The turbidity is attributed to Mie scattering from microstructures with dimensions comparable to the wavelength of visible light.⁵³ Fig. 1B shows a representative morphology of a microparticle surrounded by a collection of BCP micelles with an average diameter of $\sim 45 \text{ nm}$. This morphology is similar to that observed in Fig. S1A and E (ESI[†]),



Table 2 Structural parameters used to Fig. 1D

Model fitting	Solutions		
	Neat BCP	t-LHP	c-LHP
Beaucage model			
$R_{g,1}/\text{nm}$	—	152.07	—
$d_{f,1}$	—	3.89	—
$R_{g,2}/\text{nm}$	—	258.07	—
$d_{f,2}$	—	3.99	—
Core-shell sphere			
r_c/nm	19.59 ± 0.00	23.80 ± 0.05	20.59 ± 0.00
PDI_c	0.065	0.110	0.085
r_s/nm	10.74 ± 0.09	10.47 ± 0.01	11.40 ± 0.13
$\rho_{\text{core}} (\times 10^{-6} \text{ \AA}^{-2})$	8.720	8.705	8.720
$\rho_{\text{shell}} (\times 10^{-6} \text{ \AA}^{-2})$	8.111	8.072	8.090
$\rho_{\text{solvent}} (\times 10^{-6} \text{ \AA}^{-2})$	8.057	8.057	8.057

which was measured on a t-LHP₇ solution in its dried state. EDS 2D maps (Fig. S1B–D, ESI†) and 1D profile (Fig. S1G, ESI†) demonstrates that the microparticle is enriched with N, Pb and Br elements, suggesting that the PbBr₂ microparticle is adsorbed with PS-*b*-P2VP micelles. We believe that strong repulsions imparted by the PS shells effectively disperse the microparticles as a microemulsion within the TMB, causing the solution turbidity. Note that most micelles tend to cluster in the t-LHP solution (Fig. 1B). In comparison, the 1D profile (Fig. S1F, ESI†) of a selected PS-*b*-P2VP micelle reveals that each micelle contains Pb and Br elements with an atomic ratio of 1:3. This ratio suggests that PbBr₂ exists as complexes inside the PS-*b*-P2VP micelles.

The SAXS profile (the blue curve in Fig. 1D) of the t-LHP solution exhibits a low- q upturn with $I \sim q^{-3.9}$ in intensity at $q < 0.008 \text{ \AA}^{-1}$. This strong low- q upturn is likely due to PbBr₂ microparticles and micellar clustering in the t-LHP solution. Damped oscillations in the intermediate- q range ($0.008\text{--}0.08 \text{ \AA}^{-1}$) correspond to an increase in size dispersity of micelles. This fringe dampening is likely due to adsorption of P2VP cores onto the surface of microparticles for microscale emulsion, or micellar clustering. The surface adsorption or micellar clustering may cause micellar deformation or distortion, thus leading to the dampening of the fringes. The WAXD pattern of the t-LHP in Fig. 1E (blue curve) shows several diffractions, which can be assigned to orthorhombic PbBr₂ crystals (PDF #031-0679). This result indicates that microparticles are comprised of orthorhombic PbBr₂ crystals.

We further performed SAXS fitting on the blue curve shown in Fig. 1D for quantitative analysis. The fitting is based on a combination of the two-level Beaucage model and core-shell sphere model. The two-level Beaucage model fits the scattering of inorganic particles and micellar clustering, while core-shell sphere model fit the scattering of core-shell micelles. The fitted curve, based on structural parameters shown in Table 2, reveals that the inorganic particles are quantified with $R_{g,1} \sim 258.07 \text{ nm}$ and $d_{f,1} \sim 3.99$ and the micellar clustering is quantified with $R_{g,2} \sim 152.07 \text{ nm}$ and $d_{f,2} \sim 3.89$. The presence of these large particles indeed leads to an increase in the size distribution from 0.065 to 0.11 for the micelles in the t-LHP solution.

Yoon *et al.*⁵⁴ demonstrated the complexation of PbBr₂ when dissolved in DMF. In our study, PbBr₂ partially dissolves in TMB added with PS-*b*-P2VP, resulting in the formation of a microemulsion with BCP micelles. This microemulsion, however, is transient due to the large size of the PbBr₂ microcrystals. To further investigate how complexation occurred in the t-LHP solution, a c-LHP solution was prepared by centrifuging the t-LHP solution. The c-LHP solution (inset of Fig. 1C) exhibits a bluish color similar to the neat BCP solution, suggesting the removal of PbBr₂ microparticles. This is supported by the WAXD profile (the green curve in Fig. 1E) of the c-LHP solution, which closely resembles that of the neat BCP solution. Fig. 1C reveals a morphology of dark spheres with an average diameter of $\sim 46 \text{ nm}$. The dark spheres correspond to P2VP cores. Fitting the SAXS profile (the green curve in Fig. 2D) of the c-LHP solution shows that the radius ($\sim 20.6 \text{ nm}$) of the P2VP core and thickness ($\sim 11.4 \text{ nm}$) of the PS shell slightly increased for micelles in the c-LHP solution. Without PbBr₂ microparticles, the core-shell spherical micelles have a small size distribution at 0.085.

We further characterized the three solutions with UV-vis absorbance spectrometer. Interestingly, the UV-vis absorption spectra of the t-LHP and c-LHP solutions display an additional peak centered at $\sim 315 \text{ nm}$ (Fig. 1F). This broad band is assigned to the absorbance of $[\text{PbBr}_3]^-$ complexes bound with ligands,³⁸ suggesting coordination between PbBr₂ and 2-vinylpyridine segments within the micelle core. This coordination suggests nanoscale emulsions, by which $[\text{PbBr}_3]^-$ complexes are encapsulated within the PS-*b*-P2VP micelles.

Synthesis of perovskite nanocrystals

To investigate the influence of the precursor medium on the synthesis of encapsulated perovskite nanoparticles, varying volumes (5, 10, and 20 μL) of MABr aliquots (300 mg mL^{-1}) in either methanol or DMF were added to t-LHP solutions. The resulting solutions were labelled as t-PER_{Mx} and t-PER_{Dx} solutions. As shown in Fig. 2A_{i-iii} and A_{iv-vi}, the t-PER_{Mx} and t-PER_{Dx} solutions appeared milky yellow after 30 minutes of stirring, indicating perovskite formation. Under UV exposure (Fig. 2B_{i-vi}), a significant difference in luminescence was observed between the t-PER_{Mx} and t-PER_{Dx} solutions prepared with two types of MABr solutions. The t-PER_{Mx} solutions exhibited uniform green luminescence, while t-PER_{Dx} solutions displayed inhomogeneous luminescence, with only the top layer emitting green light (Fig. 2B_{iv-vi}). This inhomogeneity suggests a higher concentration of MAPbBr₃ microparticles with low fluorescence in the t-PER_{Dx} solutions.

Centrifugation at 7000 rpm for 10 minutes was performed to remove large particles. The resulting c-PER supernatants (c-PER_{Mx} and c-PER_{Dx}) exhibited uniform green luminescence (Fig. 2C and D). Notably, the c-PER_{Dx} solutions displayed a paler yellow coloration compared to c-PER_{Mx} solutions. A centrifugation speed of 7000 rpm was found to be optimal for producing uniform, high-quality MAPbBr₃ nanocrystals with superior optical properties. Deviations from this speed, whether too low or too high, failed to yield perovskite colloidal



solutions with comparable optical quality (Fig. S2, ESI[†]). Insufficient centrifugation left residual microcrystals in the solution, while excessive centrifugation likely disrupted the micelle encapsulation of the nanocrystals. The c-PER_{Mx} and c-PER_{Dx} solutions were further analyzed by UV-vis and PLQY.

Fig. 3A and B shows the optical properties of the c-PER_{Mx} and c-PER_{Dx} solutions, as measured by UV-vis absorbance and PL spectroscopy. The c-PER solutions are separated supernatants after centrifugation was performed on t-PER solutions to remove precipitates. By comparing the intensities of emitted and absorbed light during PL measurements, PLQY can be quantified (Fig. 3C). Several prominent features are evident from Fig. 3. With the exception of the c-PER_{M5} solution, the absorbance spectra of the c-PER_{M10} and c-PER_{M20} solutions show an Urbach tail extending from 350 to 500 nm followed by an absorbance edge at ~527 nm. These features indicate the formation of perovskite crystals in the c-PER_{M10} and c-PER_{M20} solutions (Fig. 3A). Due to the formation of perovskite crystals, the c-PER_{M10} and c-PER_{M20} solutions emit intense PL signals with superior PLQY of 74.5 and 77.3% (Fig. 3C). In comparison, the UV-vis absorbance of the c-PER_{M5} solution still shows an intense absorbance at 325 nm but a weak absorbance edge at ~527 nm. The absorbance at 325 nm can be assigned to the absorbance of [PbBr₃]⁻ complexes.³⁸ This result indicates that the c-PER_{M5} solution contained abundant [PbBr₃]⁻ complexes coexisting with a small amount of perovskite crystals. Because of the low content of perovskite crystals, the c-PER_{M5} solution emits a weak PL signal with a PLQY of approximately 59%. The weak PL and low PLQY are due to the insufficient MABr content. In contrast, the UV-vis absorbance spectra of the c-PER_{Dx} solutions show intense absorbance band at 325 nm and their PL spectra show extremely low signals with PLQY values of less than 3% (Fig. 3B and C). The result indicates that using DMF to solvate MA⁺ cations cannot effectively form perovskite crystals in the c-PER_{Dx} solutions. The comparison between the c-PER_{Mx} and c-PER_{Dx} solutions highlights the crucial role of solvent choice in the efficiency of perovskite crystallization and the resulting photoluminescence performance.

PLQY is a crucial parameter for quantitatively assessing the optical performance of perovskite nanocrystals. Table 3 summarizes the PLQY values of various perovskite materials encapsulated in BCP micelles reported in this work, alongside data from other studies.^{29–32} As shown in Table 3, our study

demonstrates a higher PLQY compared to other perovskite materials synthesized *via* BCP-templating approaches.

In addition to PLQY, the long-term stability of perovskite solutions is crucial. After 90 days of ambient storage, no observable precipitate was found in the c-PER_{M5}, c-PER_{M10}, and c-PER_{M20} solutions (Fig. S3A, ESI[†]). Furthermore, the c-PER_{Mx} solutions continued to exhibit strong absorbance at 520 nm and photoluminescence (PL) at approximately 529.3 nm (Fig. S3B, ESI[†]), features attributed to the encapsulated MAPbBr₃ nanocubes. These findings demonstrate that the encapsulated MAPbBr₃ nanocubes in the c-PER_{Mx} solutions maintain good stability in air. Particularly, the c-PER_{Mx} solutions showed increased PLQY values of 61.2%, 87.9%, and 85.6%, respectively, after 90 days of ambient storage (Fig. S3C, ESI[†]).

Note that the freshly prepared c-PER_{M10} solution exhibited a PLQY of 77%, which is already the highest among the c-PER_{Mx} solutions. However, after three months of ambient storage, the PLQY increased to 88%, suggesting that the encapsulated MAPbBr₃ nanocubes underwent further passivation during storage. Prolonged storage likely allows residual ions or block copolymer chains in the solution to interact with the nanocube surfaces, passivating additional defects and further reducing non-radiative recombination pathways.

Structural characterization of perovskite nanocrystals

Next, we characterized the c-PER_{Mx} and c-PER_{Dx} supernatants using WAXD and SAXS. Fig. 4A shows the WAXD profiles of the c-PER_{Mx} and c-PER_{Dx} supernatants. All of the c-PER_{Mx} and c-PER_{Dx} supernatants reveal a broad halo in Fig. 4A. This broad halo corresponds to the inter-chain correlation of polymer chains within micelles.⁵⁵ Furthermore, the WAXD profiles of the c-PER_{Mx} supernatants display MAPbBr₃ diffractions, while those of the c-PER_{Dx} supernatants show no MAPbBr₃ diffractions. This discrepancy indicates that the c-PER_{Mx} supernatants contain a mixture of micelles and MAPbBr₃, whereas the c-PER_{Dx} supernatants contain only micelles with little to no MAPbBr₃.

Fig. 4B shows the SAXS profiles of the c-PER_{Mx} and c-PER_{Dx} supernatants. Several scattering features are prominent. First, they show a Guinier regime at $q < 0.008 \text{ \AA}^{-1}$ and a series of oscillations in the intermediate q regime ($0.02 < q < 0.08 \text{ \AA}^{-1}$), corresponding to form factor scattering of core-shell spheres.^{47,48,55} The series of oscillations indicates a monodisperse distribution of nanostructures. Second, the oscillations in the c-PER_{Mx} solutions damped more quickly than those in the c-PER_{Dx} solutions (Fig. 4B). This difference in oscillation prominence is attributed to the coexistence of PS-*b*-P2VP micelles and inorganic nanoparticles within the c-PER_{Mx} solutions. The dampened fringes observed in the c-PER_{Mx} solutions, compared to those in the c-PER_{Dx} solutions, reflect an increase in size dispersity.

We further fitted the SAXS data using a core-shell spherical model^{47,48} to extract the structural parameters of the micelles in both types of c-PER solutions, with the results summarized in Table 4. The fits based on the structural parameters in

Table 3 Comparison of PLQY performances of various perovskite materials synthesized using the wet chemical method with PS_m-*b*-P2VP_n block copolymer as a template, where the superscripts *m* and *n* represent molecular-weight quantities in kg mol⁻¹

BCP	Solvent	Perovskite materials	PLQY	Ref.
PS ₄₉ - <i>b</i> -P2VP ₇₅	TMB	MAPbBr ₃	77%	This work
PS ₇₅ - <i>b</i> -P2VP _{66.5}	Toluene, <i>o</i> -xylene	MAPbI ₃	—	29
PS ₅₇ - <i>b</i> -P2VP ₅₇	Toluene	CsPbBr ₃	51%	30
PS ₅₅ - <i>b</i> -P2VP ₅₀	Toluene	CsPbBr ₃	-	31
PS ₃₄ - <i>b</i> -P2VP ₁₈	Toluene	MAPbBr ₃	59%	32



Table 4 Structural parameters used to fit Fig. 4B

Model fitting	c-PER solutions			
	c-PER _{M10}	c-PER _{M20}	c-PER _{D10}	c-PER _{D20}
Core-shell sphere				
r_c /nm	20.55 ± 0.05	20.41 ± 0.05	20.61 ± 0.03	20.63 ± 0.03
PDI _s	0.090	0.100	0.045	0.052
r_s /nm	10.25 ± 0.11	10.19 ± 0.11	11.35 ± 0.06	11.44 ± 0.06
ρ_{core} ($\times 10^{-6} \text{ \AA}^{-2}$)	10.951	10.953	10.851	10.851
ρ_{shell} ($\times 10^{-6} \text{ \AA}^{-2}$)	8.208	8.208	8.112	8.114
ρ_{solvent} ($\times 10^{-6} \text{ \AA}^{-2}$)	8.057	8.058	8.057	8.057

Table 4 are consistent with the SAXS curves (Fig. 4B). Table 4 demonstrates that the c-PER_{Mx} solutions contain PS-*b*-P2VP micelles with a higher dispersity compared to the c-PER_{Dx} solutions. This comparison suggests that the coexistence of micelles and nanoparticles contributes to the dampening of the scattering fringes.

The c-PER solutions were characterized by TEM in a dried state. Fig. 5 shows TEM images of the dried c-PER solutions. While the addition of both types of MABr solutions to the t-LHP solutions produced MAPbBr₃ cubic crystals, the morphology and encapsulation behavior differed significantly. Using methanol as the solvent produced numerous monodispersed encapsulated MAPbBr₃ nanocubes in the c-PER_{M10} and c-PER_{M20} solutions (Fig. 5B and C). Adding a 5 μL MABr/methanol aliquot mainly produced distorted nanocubes (Fig. 5A). In contrast, using DMF as the solvent resulted in little to no formation of MAPbBr₃ nanocubes (Fig. 5E–G). This difference in crystal growth and encapsulation likely accounts for the observed variation in fluorescence intensity between the c-PER solutions (Fig. 3).

Note that not all PS-*b*-P2VP micelles encapsulated MAPbBr₃ nanocubes (Fig. 5A–D). A large portion of PS-*b*-P2VP micelles contained no nanocubes, indicating that the nanocubes grew inside the P2VP cores of only a portion of the micelles. Moreover, the growth of encapsulated MAPbBr₃ nanocubes was almost prohibited for the c-PER_{Dx} solutions. Little to no MAPbBr₃ nanocrystals can be found neighboring with empty micelles (Fig. 5E–H).

Note that there is a significant contrast in the appearance of empty micelles between c-PER_{Mx} and c-PER_{Dx} in Fig. 5. This discrepancy should be due to different contents of Pb and Br elements trapped within P2VP cores. To address this issue, we further performed HR-TEM and EDS characterization on the nanostructures formed in the c-PER_{M10} and c-PER_{D20} solutions. Fig. 6B, C and E, F show representative EDS elemental analysis imposed on the c-PER_{M10} and c-PER_{D20} solutions in a dried state. The atomic compositions of N, Pb, and Br were detected at two different positions, labeled 1 and 2 in Fig. 6A and D.

EDS elemental analysis reveals three key findings (Fig. 6B, C and E and F). First, the MAPbBr₃ nanocubes that formed in the c-PER_{M10} solution contained significant amounts of Pb and Br, although the Pb ratio did not precisely match the stoichiometric ratio of cubic MAPbBr₃ crystals. This stoichiometric imbalance suggests the presence of defects in the cubic

MAPbBr₃ crystals. Second, the spherical micelles in the c-PER_{M10} solution appeared to contain a low amount of Pb. One selected spherical micelle displayed a high Br ratio (Fig. 6C), indicating that each MAPbBr₃ nanocube in c-PER_{M10} likely grew at the expense of [PbBr₃][−] complexes released from multiple micelles. Third, the micelles in the dried c-PER_{D20} solution contained abundant Pb elements with an atomic ratio of approximately 1 : 3 (Fig. 6E and F). Analysis of the UV-vis absorbance spectrum (blue curve in Fig. 3B) and the WAXD profile (blue curve in Fig. 4A) indicates that the micelles in the c-PER_{D20} solution primarily retained [PbBr₃][−] complexes, with only a negligible amount of encapsulated MAPbBr₃ nanocubes. The Pb : Br ratio determined by EDS elemental analysis aligns closely with the stoichiometry of [PbBr₃][−] complexes. Since the formation of MAPbBr₃ nanocubes using DMF was significantly inhibited (Fig. 5G and 6D), it is likely that most of the [PbBr₃][−] complexes remained trapped within the spherical micelles (Fig. S4, ESI†).

Furthermore, a comparison of the EDS profiles indicates that the significant contrast in the appearance of empty micelles between c-PER_{Mx} and c-PER_{Dx} in Fig. 5 and 6A, D is attributed to the retention of [PbBr₃][−] complexes in the P2VP cores of the micelles in the c-PER_{Dx} solutions. In contrast, the perovskite-free micelles in the c-PER_{M10} solution contained negligible amounts of Pb and Br elements in their P2VP cores. As a result, the P2VP cores with negligible Pb and Br content appear as gray nanodomains (Fig. 5D and 6A), while those containing [PbBr₃][−] complexes appear as dark nanodomains (Fig. 5H and 6D) under high-magnification TEM imaging.

These findings raise two key questions: how do the different solvents influence the nucleation and growth of encapsulated MAPbBr₃ nanocubes? What is the underlying mechanism driving the selective encapsulation of MAPbBr₃ nanocubes within PS-*b*-P2VP micelles? Further investigation is required to elucidate the impact of solvent quality on MAPbBr₃ crystallization and its relationship to micelle encapsulation efficiency.

Fig. 4–6 demonstrate that MAPbBr₃ nucleation and growth differ in methanol and DMF. In methanol, MA⁺ cations effectively bind with P2VP-bound [PbBr₃][−] complexes, resulting in the formation of encapsulated MAPbBr₃ nanocubes. In contrast, in DMF, MA⁺ cations do not effectively bind with [PbBr₃][−] complexes, likely due to preferential interactions with more accessible PbBr₂ microparticles. To explore this hypothesis further, we characterized the precipitates separated from the



t-PER_{Mx} and t-PER_{Dx} solutions. Fig. 7A and B show that the dried precipitates appear orange and exhibit green fluorescence. However, their XRD patterns differ significantly. The XRD patterns of the precipitates separated from the t-PER_{Mx} solutions show diffractions corresponding to a mixture of cubic MAPbBr₃ perovskite (ICSD#268785) and orthorhombic PbBr₂ (PDF#031-0679) (Fig. 7C). In contrast, the precipitates from the t-PER_{Dx} solutions exhibit WAXD patterns with diffractions corresponding solely to cubic MAPbBr₃ crystals (Fig. 7D). Note that the additional peak at $q = 1.86 \text{ \AA}^{-1}$ is attributed to the diffraction of the glass substrate.⁵⁶ These results suggest that in methanol, MA⁺ cations easily migrate to the P2VP core to bind with encapsulated [PbBr₃][−] complexes through the PS shell, producing encapsulated MAPbBr₃ nanocubes *via* crystallization. Excess MA⁺ cations are further captured by PbBr₂ microcrystals, leading to the formation of MAPbBr₃ microparticles. However, the formation of MAPbBr₃ microparticles competes with the formation of MAPbBr₃ nanocubes encapsulated within P2VP cores. This competition explains why the precipitates from the t-PER_{Mx} solutions show diffractions corresponding to a mixture of MAPbBr₃ and PbBr₂. In contrast, in DMF, most MA⁺ cations preferentially bind with free PbBr₂ microparticles, leading to the formation of MAPbBr₃ microparticles rather than MAPbBr₃ nanocubes. The precipitates primarily consisted of these MAPbBr₃ microparticles. As a result, the precipitates obtained from the t-PER_{Dx} solutions exhibited visible green photoluminescence under UV light (Fig. 7B_{iii–iv}), while the supernatants separated from the t-PER_{Dx} solutions emitted PL with low quantum yields (Fig. 3B and C). The low quantum yields displayed by the c-PER_{Dx} solutions are attributed to the low density of encapsulated MAPbBr₃ nanocubes (Fig. 5E–H).

Effect of centrifugation to remove PbBr₂ microparticles

The first section demonstrates that the hierarchical emulsion of PbBr₂ in TMB with PS-*b*-P2VP forms dispersed PbBr₂ microparticles and [PbBr₃][−] complexes. During the micro-scale emulsion process, the PbBr₂ microparticles are coated with PS-*b*-P2VP micelles, where the P2VP cores adhere to the PbBr₂ microparticles, and the PS shells swell in TMB. The [PbBr₃][−] complexes are encapsulated within the P2VP cores through nanoscale emulsion. These encapsulated [PbBr₃][−] complexes are well-dispersed in TMB due to the strong repulsive forces provided by the swollen PS shells. As a result, the hierarchical emulsion of PbBr₂ in TMB with PS-*b*-P2VP offers two key resources—dispersed PbBr₂ microparticles and [PbBr₃][−] complexes—for the formation of MAPbBr₃. To eliminate the influence of dispersed PbBr₂ microparticles on the formation of encapsulated MAPbBr₃ nanocubes, centrifugation was performed on the t-LHP solutions before adding MABr aliquots.

To saturate the loading of [PbBr₃][−] complexes within the P2VP cores, lead halide precursor solutions (denoted as t-LHP₇ solutions) were prepared by mixing PbBr₂ with PS-*b*-P2VP in TMB, followed by prolonged stirring for 7 days. After 7 days, excess PbBr₂ microparticles were removed by centrifugation at 7000 rpm for 10 minutes, resulting in clear supernatants

(denoted as c-LHP₇ solutions). This centrifugation step ensured that MA⁺ and Br[−] ions interacted exclusively with [PbBr₃][−] complexes, eliminating the influence of excess PbBr₂ microparticles while preventing the disruption of micellar encapsulation efficiency by harsh centrifugation.

The c-LHP₇ solutions were then added with 20 μL of MABr aliquots to produce clear perovskite solutions (denoted as c-PER_{c7,M20} or c-PER_{c7,D20}). The c-PER_{c7,M20} solution appeared fluorescent green and emitted strong PL (Fig. S5A and B, ESI[†]), whereas the c-PER_{c7,D20} solution appeared light yellow and emitted weak PL (Fig. S5C and D, ESI[†]). The absorbance spectrum of the c-PER_{c7,D20} solution exhibited a strong absorbance band at ~320 nm, indicating the presence of abundant [PbBr₃][−] complexes (red curve in Fig. S5E, ESI[†]). The c-PER_{c7,M20} solution showed a prominent band at ~310 nm, followed by a Urbach tail in the range of 340–500 nm and an absorbance edge at 528 nm. A review article has demonstrated that the Urbach tail is associated with defects in semiconductors.⁵⁷ Therefore, the Urbach tail and the band at 528 nm indicate the formation of perovskite crystals with defects in the c-PER_{c7,M20} solution. The 310-nm band is more pronounced in the c-PER_{c7,M20} solution (black line in Fig. S5E, ESI[†]) compared to the c-PER_{M20} solution (blue line in Fig. 3A). Previous studies have shown that the 310-nm band arises from [PbBr₆]^{4−} octahedral clusters.^{58–64} We believe that adding MABr to the c-LHP₇ solution after the removal of PbBr₂ microparticles may also lead to the formation of excess [PbBr₆]^{4−} octahedral clusters coexisting with MAPbBr₃ nanocubes.

Fig. S6A and B (ESI[†]) shows that the c-PER_{c7,M20} solution forms encapsulated MAPbBr₃ nanocubes with a slightly increased content. However, most of the PS-*b*-P2VP micelles remain a spherical shape without MAPbBr₃ nanocubes. EDS analysis of three different regions enriched with spherical micelles demonstrates that the spherical micelles contain high Br residues but low Pb residues (Fig. S6C–E, ESI[†]), indicative of [PbBr₆]^{4−} complexes. While the optical properties of c-PER_{c7,M20} and c-PER_{M20} are comparable, the former exhibits reduced long-term stability. After three months of storage, the c-PER_{c7,M20} solution became transparent due to perovskite degradation (Fig. S7, ESI[†]).

Mechanistic insights on micellization, hierarchical emulsion, PbBr₂ complexation, and formation of encapsulated MAPbBr₃ nanocubes

It is generally understood that the formation of perovskite colloidal NCs occurs *via* ion exchange through a crystalline state.⁶⁵ The mechanism of ion exchange allows us to explain how dispersed PbBr₂ microparticles bind with MA⁺ cations to form MAPbBr₃ microparticles. However, our findings indicate that amorphous [PbBr₃][−] complexes within the P2VP cores are the primary species to grow encapsulated perovskite nanocubes. This mechanism of the nucleation and growth of perovskite nanocubes in our system differs from the classical model of ion exchange.

Based on the results in Fig. 1–7, we propose an alternative mechanism to explain how PS-*b*-P2VP colloidal micelles



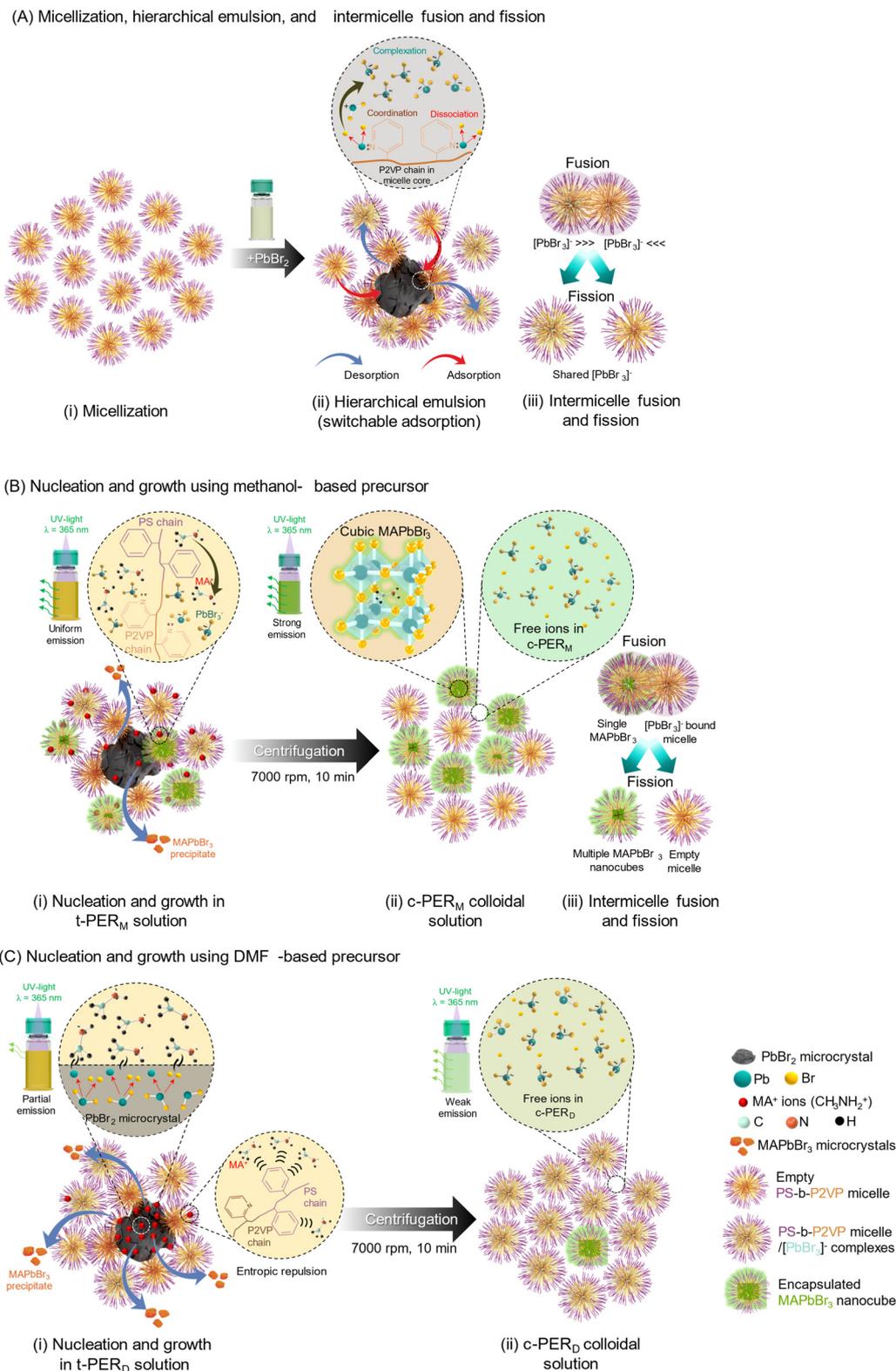


Fig. 8 Schematic illustration of structural evolutions: (A) micellization, hierarchical emulsion, and inter-micelle fusion and fission in a t-LHP solution, and the nucleation and growth of MAPbBr₃ in a t-PER solution added with (B) methanol-based or (C) DMF-based MABr aliquots.

mediate PbBr₂ complexation and MAPbBr₃ crystallization in TMB when MABr aliquots, either in methanol or DMF, are added. In TMB, a good solvent for PS, the micellization of PS-*b*-P2VP produces core-shell micelles (Fig. 8A_i). Two types of



emulsions occur at different scales: microscale and mesoscale (Fig. 8A_{ii}). At the microscale, P2VP chains preferentially adsorb onto PbBr₂ microparticles while PS chains extend into TMB. At the mesoscale, PbBr₂ undergoes dissociation and complexation, followed by coordination within the P2VP cores. This process results in a hierarchical emulsion comprising a microemulsion of dispersed PbBr₂ microparticles and a nanoemulsion of [PbBr₃]⁻ complexes within P2VP cores. It is important to note that the surface adsorption of PS-*b*-P2VP micelles is a dynamic and transient process. Frequent exchanges between complex-captured micelles and complex-free micelles occur at the interface of PbBr₂ microcrystals and PS-*b*-P2VP micelles due to Brownian motion in the solution (see Fig. 8A_{ii} where desorption and adsorption are depicted by red and blue arrows, respectively). This dynamic adsorption behavior contributes to a more uniform distribution of [PbBr₃]⁻ complexes throughout all P2VP cores (Fig. 8A_{iii}). Without BCP micelles, hierarchical emulsion could not form. Instead, adding a MABr/methanol aliquot led to the complete conversion of PbBr₂ into MAPbBr₃ microcrystals with poor dispersion in TMB (Fig. S8, ESI[†]).

Next, we explain how encapsulated MAPbBr₃ nanocubes form at the expense of [PbBr₃]⁻ complexes inside P2VP cores. Separately dissolving MABr in methanol is crucial for the formation of encapsulated MAPbBr₃ nanocubes in non-polar solvents. This dissolution step generates solvated MA⁺ and Br⁻ ions, which are necessary for the subsequent nucleation and growth of perovskite nanocubes confined within P2VP cores. Without this solvent-mediated ion formation, MABr would predominately bind with dispersed PbBr₂ microparticles in TMB to form MAPbBr₃ microparticles, thus preventing the formation of encapsulated MAPbBr₃ nanocubes inside P2VP cores (Fig. S9, ESI[†]). As a result, the P2VP cores still retain a high content of [PbBr₃]⁻ complexes. When using methanol to dissolve MABr, solvated MA⁺ cations can diffuse from the non-polar phase into the P2VP cores filled with [PbBr₃]⁻ complexes. The coordination between solvated MA⁺ cations and [PbBr₃]⁻ complexes initiates the crystallization of MAPbBr₃ (Fig. 8B_i). This mechanism facilitates the formation of encapsulated MAPbBr₃ nanocubes.

The confinement of MAPbBr₃ crystallization within P2VP cores imparts a space for homogeneous nucleation. This nucleation and growth promote the formation of stable cubic crystals (Fig. 8B_{ii}). The stability of perovskite structures can be predicted using the Goldschmidt tolerance factor (t) and the octahedral factor (μ), which are determined by the ionic radii of the constituent atoms.⁶⁶ The values of t and μ are calculated using eqn (1) and (2), respectively:

$$t = \frac{r_A + r_X}{\sqrt{2}(r_B + r_X)} \quad (1)$$

$$\mu = \frac{r_B}{r_X} \quad (2)$$

where r_A , r_B , and r_X represent the ionic radii of the A cation

(MA⁺), B cation (Pb²⁺), and X anion (Br⁻), respectively. A stable perovskite structure typically forms when $0.8 < t < 1$ and $0.44 < \mu < 0.9$. A cubic perovskite structure is expected when $0.9 < t < 1$. Using the effective ionic radii of MA⁺ (0.22 nm),⁶⁷ Pb²⁺ (0.13 nm),⁶⁸ and Br⁻ (0.18 nm),⁶⁸ we estimated $t \approx 0.91$ and $\mu \approx 0.72$ for MAPbBr₃. These values fall within the ranges for a stable cubic perovskite structure, consistent with our results.

Interestingly, the observation that nanocubes are encapsulated within some individual micelles, while others remain empty, suggests that inter-micelle fusion and fission events occur during MAPbBr₃ crystallization (Fig. 8B_{iii}). This dynamic process could explain the observed distribution of nanocubes, where some micelles encapsulate multiple nanocubes, while others do not contain any [PbBr₃]⁻ complexes. The uniform capture of [PbBr₃]⁻ complexes by micelles in the t-LHP solution further supports this mechanism.

The choice of solvent significantly influences the crystallization kinetics of MAPbBr₃. Using methanol yields MAPbBr₃ nanocrystals with narrow size dispersity, well dispersed in the supernatants. In contrast, using DMF quickly results in MAPbBr₃ microcrystals (Fig. 8C_i), which are found only in the precipitates. Only a few MAPbBr₃ nanocubes can form within P2VP cores when DMF is used to solvate MA⁺ and Br⁻ ions (Fig. 8C_{ii}). This difference arises from the solvents' impact on MA⁺ ion diffusion through the PS shells. Methanol facilitates MA⁺ diffusion through the PS shells, promoting coordination with [PbBr₃]⁻ complexes within the P2VP cores, allowing the encapsulated nanocubes to be well dispersed in the supernatants. Conversely, DMF-solvated MA⁺ cations have difficulty penetrating and diffusing through the PS shells, thus limiting their access to the [PbBr₃]⁻ complexes. Instead, DMF-solvated MA⁺ cations preferentially bind to PbBr₂ microcrystals, leading to the rapid growth of MAPbBr₃ microcrystals, which then exist as precipitates after centrifugation (Fig. 7).

Several factors can explain the observed differences in crystallization kinetics. The first factor is the distinct precursor-solvent interactions, quantified by Guttmann's donor number (DN).⁶⁹ A significant difference in DN values of methanol (DN = 19)⁷⁰ and DMF (DN = 27)⁷⁰ reflects their electron-donating abilities. Previous work by Hamill *et al.*⁷¹ demonstrated that DN serves as a strong predictor of a solvent's ability to solvate perovskite precursors. Solvents with higher DN values can more effectively compete with halide ions for coordination sites around the Pb²⁺ center. This competition might suppress the formation of ion complexes, potentially retarding perovskite nucleation and growth due to a reduced concentration of precursor ion complexes. In our case, MA⁺ cations strongly coordinate with DMF to form solvated species. While DMF's higher DN could explain its stronger coordination with MA⁺ ions compared to methanol, the high DN value does not fully account for the preferential formation of MAPbBr₃ microcrystals in t-PER_{DX} solutions. The interaction between MA⁺ cations and PbBr₂ microcrystals, which governs microcrystal formation, should not be directly affected by DN. Therefore, we can exclude the difference in Guttmann's donor number as the



principal cause for the distinctive crystallization pathways observed in the two solvent systems.

Here, it is important to consider that the swollen PS shells act as barriers to the diffusion of solvated MA⁺ cations from the liquid matrix into the P2VP cores. As a result, the diffusion rate of solvated MA⁺ cations is significantly faster in the liquid matrix than within the PS shells. The diffusion rate of solvated MA⁺ cations dispersed in a solvent can be quantified by:

$$D_o = \frac{\kappa_B T}{6\pi\mu R_{sc}} \quad (3)$$

where κ_B is the Boltzmann constant, T is the absolute temperature (in K), μ is the viscosity of the solvent, and R_{sc} denotes the effective radius of the solvated MA⁺ cations. Given that only small aliquots of MABr/DMF and MABr/methanol solutions were added to the precursor solutions containing a mixture of PbBr₂ and PS-*b*-P2VP, the viscosity of the solvent is primarily determined by the viscosity of TMB. Consequently, the diffusion of solvated MA⁺ cations is mainly influenced by their effective radius. A neat MA⁺ cation has an approximate radius of 0.22 nm.⁶⁷ Furthermore, methanol exhibits a smaller van der Waals volume compared to DMF (~34.6 Å³ for methanol and ~77.6 Å³ for DMF).⁷² It is believed that a DMF-solvated MA⁺ cation has a larger effective radius compared to a methanol-solvated MA⁺ cation.

In solution, methanol-solvated MA⁺ cations can more quickly promote the formation of MAPbBr₃ microcrystals at the expense of PbBr₂ microcrystals adsorbed by PS-*b*-P2VP micelles, compared to DMF-solvated MA⁺ cations. Microscale emulsion disperses the PbBr₂ microcrystals, which are partially adsorbed by PS-*b*-P2VP micelles. However, there should be sufficient voids that are free from PS-*b*-P2VP micelle adsorption. These BCP-free voids allow the solvated MA⁺ cations to bind with PbBr₂ microcrystals, facilitating the rapid formation of MAPbBr₃ microcrystals. This observation explains why the precipitates are predominantly composed of cubic MAPbBr₃ microcrystals.

For the formation of encapsulated MAPbBr₃ nanocubes within P2VP cores through nanoscale emulsion, the additional barriers imparted by PS shells must be carefully considered. The penetration and diffusion of solvated MA⁺ cations are analogous to the permeation of gas through a polymer.⁷³ Two factors affect permeability: the solubility and diffusivity of the permeating species through the polymer along the concentration gradient. In our system, solubility should not be a key factor as the contrast in solubility parameters between methanol (29.6 MPa^{1/2})⁴⁹ and TMB (18 MPa^{1/2})⁴⁹ is greater than the contrast in solubility parameters between DMF (24.8 MPa^{1/2})⁴⁹ and TMB. Therefore, we believe that diffusivity plays a more critical role in the permeation of solvated MA⁺ cations through PS shells (18.6 MPa^{1/2}).⁵⁰ Diffusivity is influenced by polymer packing and the kinetic diameter of the diffusing molecule itself.

Quantitative analysis of the SAXS data demonstrates that the addition of DMF or methanol has no significant effect on the radii of the P2VP core and PS shell, indicating that the spatial

packing of the shell-forming corona remains unchanged. In other words, the polymer packing of PS shells swollen in TMB-rich solutions is consistent, regardless of whether DMF or methanol is added.

In this study, we hypothesize that the inhibition of encapsulated MAPbBr₃ nanocube formation is linked to the possibility that DMF-solvated MA⁺ cations have a larger kinetic diameter. As a result, the penetration of PS shells with a brush conformation by DMF-solvated MA⁺ cations may cause a significant entropic penalty (Fig. 8C_i), which is unfavorable for the diffusion of DMF-solvated MA⁺ cations through the PS shell. This could explain why [PbBr₃]⁻ complexes remained within the P2VP cores without forming MAPbBr₃ nanocubes when DMF was used to produce solvated MA⁺ and Br⁻ ions.

Next, we explain why the c-PER_{c7,M20} solution exhibits lower PL stability compared to the c-PER_{M20} solution. PbBr₂ microcrystals likely serve as a reservoir, continuously replenishing [PbBr₃]⁻ complexes. Without PbBr₂ microcrystals, Br⁻ ions are left unbound, leading to a significant reduction in [PbBr₃]⁻ complexes. This is supported by the distinct optical properties in freshly prepared c-PER_{c7,M20} solutions compared to c-PER_{M20} (see Fig. S5 (ESI[†]) and Fig. 3). Unlike the broad shoulder displayed by the c-PER_{M20} solution, a sharp peak at ~308 nm suggests the presence of abundant [PbBr₆]⁴⁻ octahedral clusters, likely due to excess Br⁻ ions in the c-PER_{c7,M20} solution. A high residue of [PbBr₆]⁴⁻ octahedral clusters may lead to the degradation of MAPbBr₃ nanocubes (see Fig. S7, ESI[†]). This explains why the PL performance of the c-PER_{c7,M20} solution degraded after three months of storage at room temperature.

Conclusions

We have demonstrated a straightforward method for synthesizing MAPbBr₃ nanocubes encapsulated within polymeric micelles. This approach capitalizes on the interaction between [PbBr₃]⁻ complexes and solvated MA⁺ cations within a confined micellar environment. The choice of solvent to solvate MA⁺ cations significantly influences the crystallization kinetics and the quality of the nanocubes. Methanol, with its smaller molecular size, facilitates faster diffusion of solvated MA⁺ ions, promoting the formation of well-defined nanocubes. In contrast, DMF leads to slower diffusion of solvated MA⁺ ions and the formation of less desirable microcrystals. The MAPbBr₃ colloidal solutions synthesized using methanol-based aliquots exhibit remarkable stability. After 90 days of storage under ambient conditions, no decrease in PLQY was observed. In fact, the PLQY measurements indicated a 14% increase compared to the freshly prepared solutions.

Furthermore, our findings highlight the crucial role of excess PbBr₂ microcrystals in synthesizing encapsulated nanocubes and maintaining their long-term stability. Although removing these microcrystals through centrifugation initially produces solutions with comparable optical properties, it disrupts the stoichiometric balance of MA, Pb, and Br elements necessary for forming MAPbBr₃ crystals. This stoichiometric



imbalance ultimately undermines the stability of the nanocubes, leading to degradation over time. These insights underscore the importance of carefully controlling crystallization conditions and understanding the underlying stoichiometric equilibria to achieve stable, high-quality perovskite nanomaterials for optoelectronic applications.

Author contributions

Belda Amelia Junisu: conceptualization, writing – original draft, data curation, methodology, investigation, visualization; Ya-Sen, Sun: conceptualization, writing – review and editing, funding acquisition, resources, supervision; Bo Cheng Zhao: investigation

Data availability

The data that support the findings of this study are available from the corresponding author upon reasonable request.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Financial support from the National Science and Technology Council (grant numbers: NSTC 111-2221-E-006-232-MY3 and 112-2811-E-006-042-MY2) is acknowledged. We further thank Dr Orion Shih, Dr Yi-Qi Yeh, Mr Je-Wei Chang, and Ms Kuei-Fen Liao for SAXS and WAXD experiments at TPS 13A. We also express our gratitude for the use of EM025800 (NSTC 113-2740-M-006-002) at the Core Facility Center of National Cheng Kung University (NCKU). Additionally, we thank Mr Yu-Hsuan Cheng for performing the TEM and EDS measurements at NCKU.

References

- J. Y. Kim, J.-W. Lee, H. S. Jung, H. Shin and N.-G. Park, *Chem. Rev.*, 2020, **120**, 7867–7918.
- J. J. Yoo, G. Seo, M. R. Chua, T. G. Park, Y. Lu, F. Rotermond, Y.-K. Kim, C. S. Moon, N. J. Jeon, J.-P. Correa-Baena, V. Bulović, S. S. Shin, M. G. Bawendi and J. Seo, *Nature*, 2021, **590**, 587–593.
- S. D. Adhikari, A. F. G. Reyes, S. Paul, J. Torres, B. Escuder, I. Mora-Seró and S. Masi, *Chem. Sci.*, 2023, **14**, 8984–8999.
- D. Park, W. Kim, C. Park, J. Choi, A. Ghorai, G. Lee, S. Choi, W. Moon and U. Jeong, *Small*, 2024, **20**, 2402281.
- F. Wang, X. Zou, M. Xu, H. Wang, H. Wang, H. Guo, J. Guo, P. Wang, M. Peng, Z. Wang, Y. Wang, J. Miao, F. Chen, J. Wang, X. Chen, A. Pan, C. Shan, L. Liao and Q. Hu, *Adv. Sci.*, 2021, **8**, 2100569.
- Y. Liu, Y. Liu and Y. Guo, *Mater. Chem. Front.*, 2023, **7**, 5215–5246.
- S. Luo and W. A. Daoud, *J. Mater. Chem. A*, 2015, **3**, 8992–9010.
- B. W. Boote, H. P. Andaraarachchi, B. A. Rosales, R. Blome-Fernández, F. Zhu, M. D. Reichert, K. Santra, J. Li, J. W. Petrich, J. Vela and E. A. Smith, *Chem. Phys. Chem.*, 2019, **20**, 2647–2656.
- D. Han, K. Yang, C. Bai, F. Chen, Z. Sun, Y. Wang, H. Ji, Z. Yang and X. Tang, *J. Chem. Eng.*, 2023, **475**, 146209.
- D. Liu, Z. Shao, C. Li, S. Pang, Y. Yan and G. Cui, *Small Struct.*, 2021, **2**, 2000089.
- A. Ghorai, A. Das, Y. Choi, B. Roy and U. Jeong, *Small*, 2024, **20**, 2404384.
- Y. Zhao and K. Zhu, *Chem. Soc. Rev.*, 2016, **45**, 655–689.
- W. K. Chan, J. Chen, D. Zhou, J. Ye, R. J. Vázquez, C. Zhou, G. C. Bazan, A. Rao, Z. Yu and T. T. Y. Tan, *Nanomaterials*, 2023, **13**, 815.
- J. You, Y. M. Yang, Z. Hong, T.-B. Song, L. Meng, Y. Liu, C. Jiang, H. Zhou, W.-H. Chang, G. Li and Y. Yang, *Appl. Phys. Lett.*, 2014, **105**, 183902.
- T. A. Berhe, W.-N. Su, C.-H. Chen, C.-J. Pan, J.-H. Cheng, H.-M. Chen, M.-C. Tsai, L.-Y. Chen, A. A. Dubale and B.-J. Hwang, *Energy Environ. Sci.*, 2016, **9**, 323–356.
- M. L. Petrus, J. Schlipf, C. Li, T. P. Gujar, N. Giesbrecht, P. Müller-Buschbaum, M. Thelakkat, T. Bein, S. Hüttner and P. Docampo, *Adv. Energy Mater.*, 2017, **7**, 1700264.
- W. Zhao, D. Yang and S. F. Liu, *Small*, 2017, **13**, 1604153.
- X. Sheng, G. Chen, C. Wang, W. Wang, J. Hui, Q. Zhang, K. Yu, W. Wei, M. Yi, M. Zhang, Y. Deng, P. Wang, X. Xu, Z. Dai, J. Bao and X. Wang, *Adv. Funct. Mater.*, 2018, **28**, 1800283.
- J. A. Laranjeira, S. A. Azevedo, G. S. Fabris, J. R. Sambrano and M. M. Ferrer, *Comput. Mater. Sci.*, 2024, **239**, 112977.
- H. Li, L. Xiong, J. Li, Y. Lu, Z. Shen, D. Song, S. Zhao, Z. Xu, Z. Liang and B. Qiao, *J. Phys. Chem. Lett.*, 2023, **14**, 5481–5488.
- S. Wang, A. A. Yousefi Amin, L. Wu, M. Cao, Q. Zhang and T. Ameri, *Small Struct.*, 2021, **2**, 2000124.
- M. Meyns, M. Perálvarez, A. Heuer-Jungemann, W. Hertog, M. Ibáñez, R. Nafria, A. Genç, J. Arbiol, M. V. Kovalenko, J. Carreras, A. Cabot and A. G. Kanaras, *ACS Appl. Mater. Interfaces*, 2016, **8**, 19579–19586.
- H. Sun, Z. Yang, M. Wei, W. Sun, X. Li, S. Ye, Y. Zhao, H. Tan, E. L. Kynaston, T. B. Schon, H. Yan, Z. H. Lu, G. A. Ozin, E. H. Sargent and D. S. Seferos, *Adv. Mater.*, 2017, **29**, 1701153.
- H. Kim, N. Hight-Huf, J. H. Kang, P. Bisnoff, S. Sundararajan, T. Thompson, M. Barnes, R. C. Hayward and T. Emrick, *Angew. Chem., Int. Ed.*, 2020, **59**, 10802–10806 (*Angew. Chem.*, 2020, **132**, 10894–10898).
- Y. He, Y. J. Yoon, Y. W. Harn, G. V. Biesold-McGee, S. Liang, C. H. Lin, V. V. Tsukruk, N. Thadhani, Z. Kang and Z. Lin, *Sci. Adv.*, 2019, **5**, eaax4424.
- Y. Mai and A. Eisenberg, *Chem. Soc. Rev.*, 2012, **41**, 5969–5985.
- Y. Zou, X. Zhou, J. Ma, X. Yang and Y. Deng, *Chem. Soc. Rev.*, 2020, **49**, 1173–1208.



- 28 C. Li, Q. Li, Y. V. Kaneti, D. Hou, Y. Yamauchi and Y. Mai, *Chem. Soc. Rev.*, 2020, **49**, 4681–4736.
- 29 L. Hui, C. Beswick, A. Getachew, H. Heilbrunner, K. Liang, G. Hanta, R. Arbi, M. Munir, H. Dawood, N. Isik Goktas, R. LaPierre, M. C. Scharber, N. S. Sariciftci and A. Turak, *ACS Appl. Nano Mater.*, 2019, **2**, 4121–4132.
- 30 S. Hou, Y. Guo, Y. Tang and Q. Quan, *ACS Appl. Mater. Interface*, 2017, **9**, 18417–18422.
- 31 K. Lee, J. Y. Kim and B.-H. Sohn, *RSC Adv.*, 2022, **12**, 6389–6395.
- 32 Y. Nah, D. Jang and D. H. Kim, *Chem. Commun.*, 2021, **57**, 1879–1882.
- 33 M. G. Greiner, A. Singldinger, N. A. Henke, C. Lampe, U. Leo, M. Gramlich and A. S. Urban, *Nano Lett.*, 2022, **22**, 6709–6715.
- 34 V. A. Hintermayr, C. Lampe, M. Löw, J. Roemer, W. Vanderlinden, M. Gramlich, A. X. Böhm, C. Sattler, B. Nickel, T. Lohmüller and A. S. Urban, *Nano Lett.*, 2019, **19**, 4928–4933.
- 35 Q. Jiang, Y. Ren, Z. Cui, Z. Li, L. Hu, R. Guo, S. Duan, F. Xie, G. Zhou and S. Xiong, *ACS Appl. Nano Mater.*, 2023, **6**, 8655–8667.
- 36 C. Collantes, W. Teixeira, V. G. Pedro, M.-J. Bañuls and Á. Maquieira, *Appl. Mater. Today*, 2023, **31**, 101775.
- 37 H. Zamani, T.-H. Chiang, K. R. Klotz, A. J. Hsu and M. M. Maye, *Langmuir*, 2022, **38**, 9363–9371.
- 38 S. K. Balakrishnan and P. V. Kamat, *Chem. Mater.*, 2018, **30**, 74–78.
- 39 M. Wang, Q. Yu, T. Yu, S. Zhang, M. Gong and Y. Liu, *RSC Adv.*, 2023, **13**, 5158–5167.
- 40 K. G. Stamplecoskie, J. S. Manser and P. V. Kamat, *Energy Environ. Sci.*, 2015, **8**, 208–215.
- 41 H. Han, J. W. Oh, H. Lee, S. Lee, S. Mun, S. Jeon, D. Kim, J. Jang, W. Jiang, T. Kim, B. Jeong, J. Kim, D. Y. Ryu and C. Park, *Adv. Mater.*, 2024, **36**, 2310130.
- 42 N. Pradhan, *ACS Phys. Chem. Au.*, 2022, **2**, 268–276.
- 43 P. C. Sadek, *The HPLC solvent guide*, Wiley-Interscience, 2002, ISBN: 978-0.
- 44 O. Shih, K.-F. Liao, Y.-Q. Yeh, C.-J. Su, C.-A. Wang, J.-W. Chang, W.-R. Wu, C.-C. Liang, C.-Y. Lin, T.-H. Lee, C.-H. Chang, L.-C. Chiang, C.-F. Chang, D.-G. Liu, M.-H. Lee, C.-Y. Liu, T.-W. Hsu, B. Mansel, M.-C. Ho, C.-Y. Shu, F. Lee, E. Yen, T.-C. Lind and U. Jenga, *J. Appl. Cryst.*, 2022, **55**, 340–352.
- 45 M. Doucet, J. H. Cho, G. Alina, Z. Attala, J. Bakker, P. Beaucage, W. Bouwman, R. Bourne, P. Butler, K. Campbell, T. Cooper-Benun, C. Durniak, L. Förster, P. Gilbert, M. Gonzalez, R. Heenan, A. Jackson, S. King, P. Kienzle, J. Krzywon, T. Nielsen, L. O'Driscoll, W. Potrzebowski, S. Prescott, R. Ferraz Leal, P. Rozyczko, T. Snow, A. Washington and C. Wolf, *SasView (version 5.0.3)*, 2020.
- 46 G. Beaucage, *J. Appl. Crystall.*, 1995, **28**, 717–728.
- 47 S. Förster, A. Timmann, M. Konrad, C. Schellbach, A. Meyer, S. Funari, P. Mulvaney and R. Knott, *J. Phys. Chem. B*, 2005, **109**, 1347–1360.
- 48 I. Akiba and K. Sakurai, *J. Polym.*, 2021, **53**, 951–973.
- 49 C. M. Hansen, *Hansen solubility parameters: a user's handbook*, CRC press, 2007.
- 50 A. F. M. Barton, *Chem. Rev.*, 1975, **75**, 731–753.
- 51 J. G. Kennemur, *Macromolecules*, 2019, **52**, 1354–1370.
- 52 J.-Y. Liou and Y.-S. Sun, *Macromolecules*, 2012, **45**, 1963–1971.
- 53 C. M. Septani, M.-F. Ku, C.-Y. Chen, J.-M. Lin and Y.-S. Sun, *Colloids Surf., A*, 2022, **652**, 129857.
- 54 S. J. Yoon, K. G. Stamplecoskie and P. V. Kamat, *J. Phys. Chem. Lett.*, 2016, **7**, 1368–1373.
- 55 R.-J. Roe, *Methods of X-ray and neutron scattering in polymer science*, Oxford University Press, 2000, pp. 140–145, pp 160–162.
- 56 P. Ponce-Peña, M. A. González-Lozano, M. Á. Escobedo-Bretado, D. M. Núñez-Ramírez, A. Rodríguez-Pulido, Z. V. Quiñones Jurado, M. Poisot and B. Sulbarán-Rangel, *Crystals*, 2022, **12**, 574.
- 57 L. Wang, H. Liu, Y. Zhang and O. F. Mohammed, *ACS Energy Lett.*, 2020, **5**, 87–99.
- 58 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.
- 59 Z. Liu, Y. Bekenstein, X. Ye, S. C. Nguyen, J. Swabeck, D. Zhang, S.-T. Lee, P. Yang, W. Ma and A. P. Alivisatos, *J. Am. Chem. Soc.*, 2017, **139**, 5309–5312.
- 60 M. Nikl, E. Mihokova, K. Nitsch, F. Somma, C. Giampaolo, G. Pazzi, P. Fabeni and S. Zazubovich, *Chem. Phys. Lett.*, 1999, **306**, 280–284.
- 61 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.
- 62 S. K. Sharma, C. Phadnis, T. K. Das, A. Kumar, B. Kavaipatti, A. Chowdhury and A. Yella, *Chem. Mater.*, 2019, **31**, 3111–3117.
- 63 S. Seth and A. Samanta, *J. Phys. Chem. Lett.*, 2017, **8**, 4461–4467.
- 64 D. Chen, Z. Wan, X. Chen, Y. Yuan and J. Zhong, *J. Mater. Chem. C*, 2016, **4**, 10646–10653.
- 65 H. Jiang, S. Cui, Y. Chen and H. Zhong, *Nano Select*, 2021, **2**, 2040–2060.
- 66 E. Akman, T. Ozturk, W. Xiang, F. Sadegh, D. Prochowicz, M. M. Tavakoli, P. Yadav, M. Yilmaz and S. Akin, *Energy Environ. Sci.*, 2023, **16**, 372–403.
- 67 G. Kieslich, S. Sun and A. K. Cheetham, *Chem. Sci.*, 2015, **6**, 3430–3433.
- 68 T. J. Jacobsson, J.-P. Correa-Baena, M. Pazoki, M. Saliba, K. Schenk, M. Grätzel and A. Hagfeldt, *Energy Environ. Sci.*, 2016, **9**, 1706–1724.
- 69 V. Gutmann and V. Gutmann, *The donor-acceptor approach to molecular interactions*, Springer, 1978.
- 70 F. Cataldo, *Eur. Chem. Bull.*, 2015, **4**, 92–97.
- 71 J. C. Hamill Jr, J. Schwartz and Y.-L. Loo, *ACS Energy Lett.*, 2018, **3**, 92–97.
- 72 Y. H. Zhao, M. H. Abraham and A. M. Zissimos, *J. Org. Chem.*, 2003, **68**, 7368–7373.
- 73 L. H. Sperling, *Introduction to Physical Polymer Science*, John Wiley & Sons, 2015.

