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Using steric hindrance to manipulate and stabilize metal halide perovskites for optoelectronics

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The chemical instability of metal halide perovskite materials can be ascribed to their unique properties of softness, in which the chemical bonding between metal halide octahedral frameworks and cations is the weak ionic and hydrogen bonding as in most perovskite structures. Therefore, various strategies have been developed to stabilize the cations and metal halide frameworks, which include incorporating additives, developing two-dimensional perovskites and perovskite nanocrystals, etc. Recently, the important role of utilizing steric hindrance for stabilizing and passivating perovskites has been demonstrated. In this perspective, we summarize the applications of steric hindrance in manipulating and stabilizing perovskites. We will also discuss how steric hindrance influences the fundamental kinetics of perovskite crystallization and film formation processes. The similarities and differences of the steric hindrance between perovskite solar cells and perovskite light emission diodes are also discussed. In all, utilizing steric hindrance is a promising strategy to manipulate and stabilize metal halide perovskites for optoelectronics.

Introduction

Metal halide perovskites have become one of the most promising semiconductor materials during the past decade because of their excellent performances in charge carrier mobility, absorption coefficient, and photoluminescence quantum efficiency.^{1–4} The efficiencies of perovskite solar cells (PSCs) and

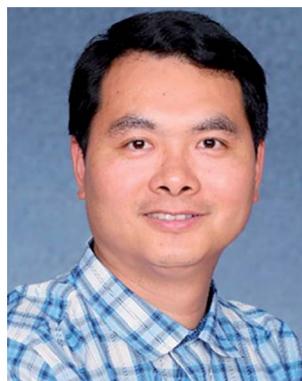
perovskite light-emitting diodes (PeLEDs) have reached impressive values of 25% (power conversion efficiency, PCE) and 23% (external quantum efficiency, EQE), respectively.^{5–10} However, the inferior operational stability caused by the weak ionic and hydrogen bonding in perovskite structures is one of the main obstacles hindering their further applications and potential commercialization.^{11,12} The most widely studied three-dimensional (3D) perovskites consist of frameworks organized with corner-sharing octahedra (*e.g.* [PbI₆]^{4–}) and small-sized cations localized in cages. Besides bonding effects, non-covalent steric interactions also influence the structure of perovskite crystals by repulsive forces between overlapping electron clouds. In chemical reactions, steric hindrance

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influences the active sites and reaction rate.^{13–17} Steric sizes of A, B and X in metal halide perovskites should also be suitable to meet an appropriate tolerance factor and octahedral factor to maintain the perovskite structure. Therefore, steric hindrance could help to tune and stabilize the crystal structure of perovskites.

Over the past few years, there have been some noticeable studies based on steric hindrance using anions and cations with different sizes, rigidity and functional groups to modify the crystal structure, film quality and device performance.^{18–22} Steric hindrance has been shown to have direct effects on the optical bandgap, tolerance factor, phase stability, trap passivation, hydrophobicity and conductivity of perovskite materials (Fig. 1).^{22–28} Several general strategies have been developed to obtain stable perovskite structures such as utilizing additives, and developing two-dimensional (2D) perovskites and perovskite nanocrystals (NCs). In these methods, some extra components such as pseudohalides, bulky organic cations and long chain ligands can determine the properties of perovskite devices as steric hindrance plays an important role in these processes.^{6,29–33} There is also some burgeoning research on rational design based on steric hindrance, which could improve both efficiency and stability.^{22,34–36} These features have made the perspective of steric hindrance very attractive to researchers to understand the fundamental kinetics of perovskite crystallization and film formation processes.

In this perspective, we mainly focus on the role of steric hindrance in manipulating and stabilizing perovskites. We will discuss steric hindrance related work, reveal the wide applicability of steric hindrance and promote the realization of commercialized stable perovskite devices. We also note that the properties of perovskite structures are not simply influenced by one factor but determined by the combination of several interactions between components such as ionic bonds, hydrogen bonds and steric hindrance. Therefore, this perspective will mainly discuss the role of steric hindrance in perovskites and illustrate how steric hindrance and other interactions co-determine the properties of perovskites.

Steric hindrance in 3D perovskites

3D perovskites have a general formula of ABX_3 where A is a small-sized cation such as $CH_3NH_3^+$ (MA^+), $CH(NH_2)_2^+$ (FA^+), and Cs^+ , B is a divalent metal cation such as Pb^{2+} or Sn^{2+} and X is a halide. The Goldschmidt tolerance factor (t) within the limit of 0.813–1.107 is usually used to determine whether a 3D perovskite structure can be formed by using the following equation:

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

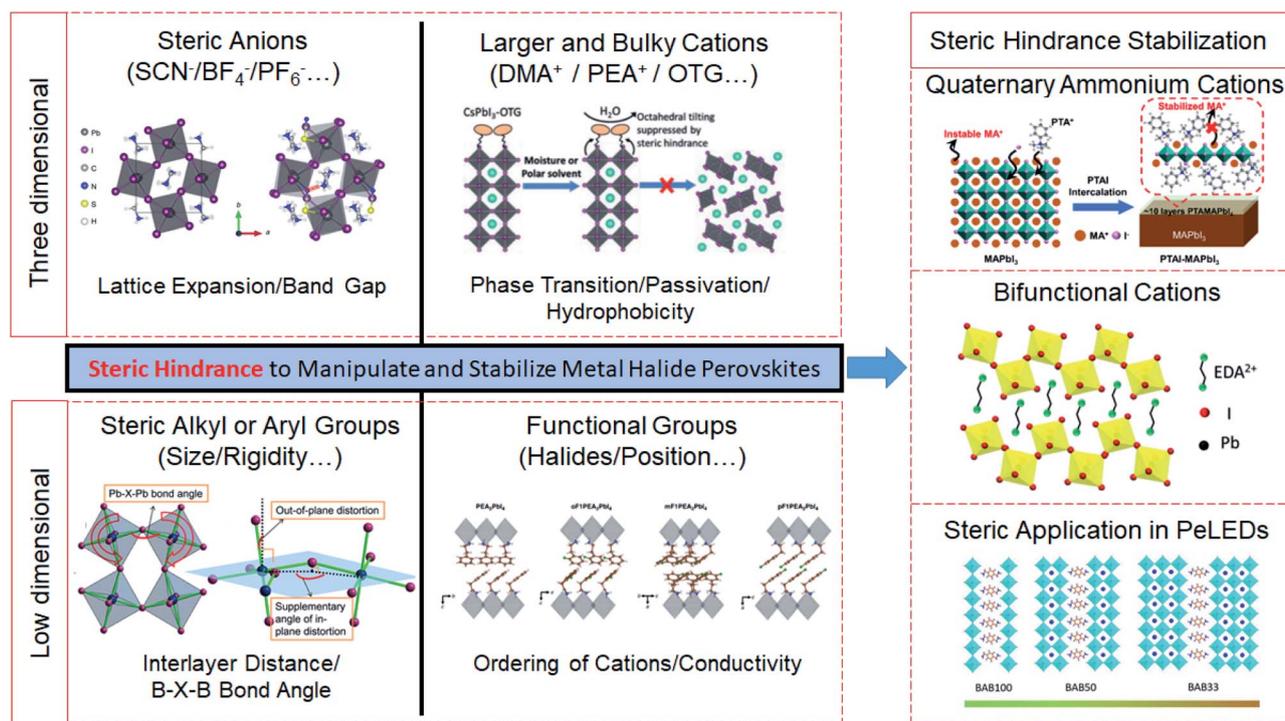


Fig. 1 Schematic illustration of the steric hindrance in three-dimensional perovskites and low-dimensional perovskites, and the steric hindrance induced stabilization of perovskites. Reproduced with permission from ref. 23. Copyright 2016 Springer Nature. Reproduced with permission from ref. 24. Copyright 2019 John Wiley and Sons. Reproduced with permission from ref. 25. Copyright 2017 American Chemical Society. Reproduced with permission from ref. 26. Copyright 2019 Springer Nature. Reproduced with permission from ref. 22. Copyright 2020 John Wiley and Sons. Reproduced with permission from ref. 27. Copyright 2018 American Association for the Advancement of Science. Reproduced with permission from ref. 28. Copyright 2019 American Association for the Advancement of Science.



r_A , r_B and r_X are the ionic radii of A, B and X, respectively.³⁷ Theoretical calculations and experiments have shown that the steric size of A, B and X will influence not only the tolerance factor, but also structural parameters such as metal–halide–metal (B–X–B) bond angles and B–X bond lengths which will determine the optical properties and stability of lattice structures. In this section, we will focus on the steric hindrance in 3D perovskites and review how the steric size of anions and cations influences the properties, especially the stability, of perovskite materials.

Anions

The 3D lead halide perovskite lattice structure is based on corner-sharing $[BX_6]^{4-}$ frames consisting of metal cations and halide anions. The bandgap of perovskite materials is mainly determined by the p and s orbitals of metal cations and p orbitals of halide anions.^{38–43} Halides such as I^- , Br^- and Cl^- are the most commonly used anions. However, the oxidation of halides is one of the main challenges for realizing stable metal halide perovskites. Halides in a perovskite lattice will form elementary X_2 and further accelerate the degradation of perovskites.^{44–47} Replacing some halide ions with other anions of appropriate sizes is an effective way to stabilize perovskites.⁴⁸ Pseudohalides such as SCN^- (217 pm), BF_4^- (218 pm) and PF_6^-

(255 pm) with a similar ionic radius to I^- (220 pm) are potential candidates for improving the structural stability of $APbI_3$ perovskites.^{49–51}

The lone pair of electrons in SCN^- ions not only has a stronger interaction with Pb^{2+} than with I^- , but also forms hydrogen bonds with MA.²³ The calculated formation constant for PbI_4^{2-} and $Pb(SCN)_4^{2-}$ are 3.5 and 7, respectively, which means SCN^- can improve the stability of perovskite frame structures. Jiang *et al.* replaced two thirds of I^- with SCN^- and the humidity stability was significantly improved.⁵² The PCE of $MAPbSCN_2I$ based PSCs only dropped from 8.3% to 7.4% after 14 days of storage. However, further studies reported that $MAPbSCN_2I$ is in fact a layered perovskite $MA_2PbSCN_2I_2$ with a bandgap of 1.53 eV (Fig. 2a).^{53,54} The stronger interaction between Pb^{2+} and SCN^- can also be applied to stabilize the $FAPbI_3$ perovskite by phase transition. Recent work by Lu *et al.* revealed that SCN^- from $MASCN$ or $FASCN$ vapor treatment can replace I^- on face-sharing δ - $FAPbI_3$ and induce the formation of corner-sharing α - $FAPbI_3$ with much reduced trap density and improved stability (Fig. 2b).⁵⁵ Our work reported that SCN^- incorporated into $CsPbBr_3$ nanocrystals will cause an abnormal blue PL shift (Fig. 2c). We revealed that the steric shape of rod-like SCN^- will induce the expansion of the $CsPbBr_3$ crystal lattice and increase the energy of the conduction band.⁵¹

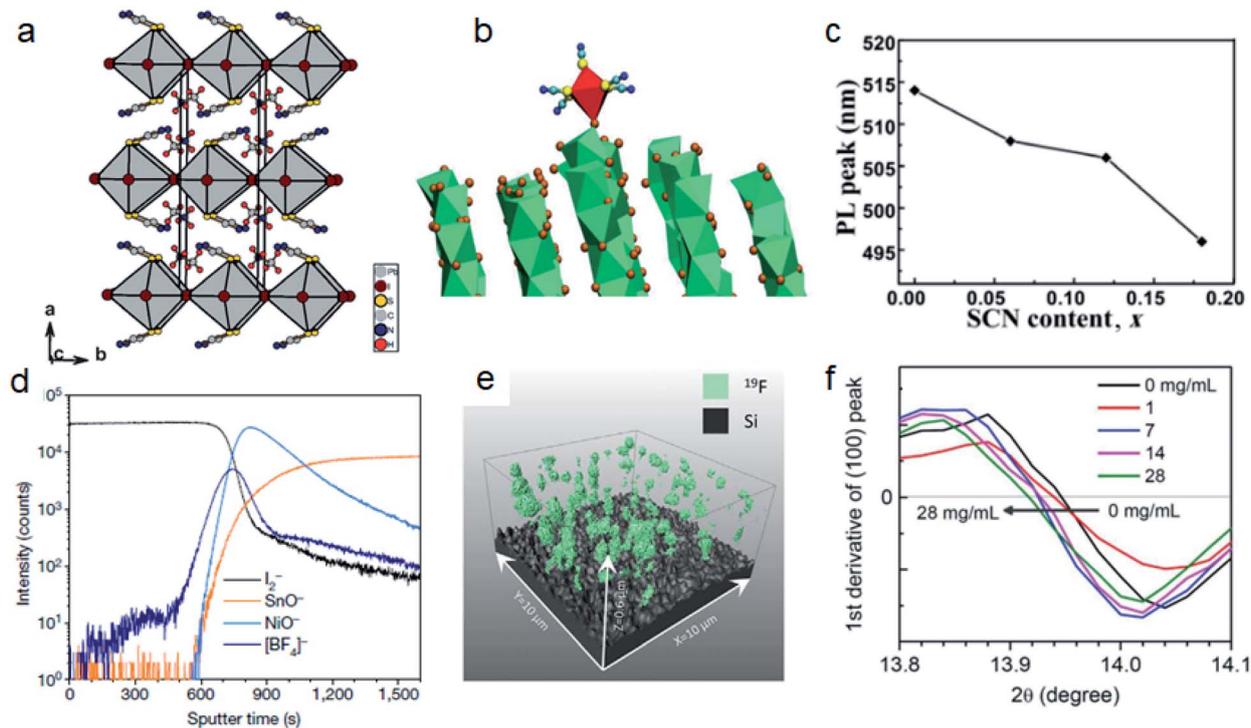


Fig. 2 (a) Crystal structure of $MA_2PbSCN_2I_2$. Reproduced with permission from ref. 53. Copyright 2015 John Wiley and Sons. (b) Molecular dynamics simulation of SCN^- induced face-sharing octahedra to the corner-sharing octahedra of the $FAPbI_3$ crystal structure. Reproduced with permission from ref. 55. Copyright 2020 American Association for the Advancement of Science. (c) PL peak positions of $CsPbBr_{3-x}(SCN)_x$ ($x = 0.06, 0.12, 0.18$) nanocrystals with different SCN contents. Reproduced with permission from ref. 51. Copyright 2018 Springer Nature. (d) ToF-SIMS depth profile of $[BF_4]^-$ in perovskite films. Reproduced with permission from ref. 56. Copyright 2019 Springer Nature. (e) 3D ToF-SIMS map of ^{19}F in perovskite films. Reproduced with permission from ref. 44. Copyright 2020 American Association for the Advancement of Science. (f) (100) peak positions in XRD patterns of $FA_{0.88}Cs_{0.12}PbI_3$ films post-treated with $FAPF_6$ with different concentrations. Reproduced with permission from ref. 58. Copyright 2018 John Wiley and Sons.



Different from SCN^- , BF_4^- has weak interaction with Pb^{2+} but can also improve the stability of perovskites by steric hindrance induced lattice strain relaxation. Zhang *et al.* added a small amount (0.5–2% molar ratio) of BF_4^- into $(\text{FAPbI}_3)_{0.83}(\text{MAPbBr}_3)_{0.17}$ precursor solutions.⁴⁹ They found that the XRD peak of (001) shifted from 14.20° to 14.14° , which means that BF_4^- had been incorporated into the perovskite structure and the elongated Pb– BF_4 bond can cause lattice relaxation. The PCE increased from 17.55% to 20.16%, and the device could retain 86% of the initial efficiency after 300 h under AM 1.5G illumination. Snaith and co-authors applied two kinds of BF_4^- based ionic liquids: 1-butyl-3-methylimidazolium tetrafluoroborate (BMIMBF_4) and 1-butyl-1-methylpiperidinium tetrafluoroborate (BMPBF_4) to modify $(\text{FA}_{0.83}\text{MA}_{0.17})_{0.95}\text{Cs}_{0.05}\text{Pb}(\text{I}_{0.9}\text{Br}_{0.1})_3$ and $\text{Cs}_{0.17}\text{FA}_{0.83}\text{Pb}(\text{I}_{0.77}\text{Br}_{0.23})_3$ perovskite films, respectively.^{44,56} In contrast, they found that BF_4^- in these two salts cannot be incorporated into the perovskite lattice and pairing with different cations will influence the distribution of BF_4^- . In the case of BMIMBF_4 , BF_4^- is located mainly at the buried interface (Fig. 2d), while BF_4^- of BMPBF_4 is distributed over the entire film (Fig. 2e). These two salts can improve both efficiency and stability of PSCs. BMIMBF_4 treated PSCs exhibited an increase of PCE from 18.5% to 19.8% and can maintain 95% efficiency under AM 1.5G at $\sim 75^\circ\text{C}$ for more than 1800 h. BMPBF_4 based PSCs reached a PCE of 20.1%, and the unencapsulated device can maintain 80% efficiency for 1010 hours at 60°C .

PF_6^- is another noncoordinating anion similar to BF_4^- .^{50,57} Chen *et al.* post treated a $\text{FA}_{0.88}\text{Cs}_{0.12}\text{PbI}_3$ film with FAPF_6 . The resulting XRD patterns show that the lattice constant increased, which means that some I^- ions were replaced by PF_6^- (Fig. 2f).⁵⁸ The larger steric size of PF_6^- leads to the expansion of the lattice and reduces the bandgap. The efficiency of these PSCs

increased from 17.79% to 19.25% and the PF_6^- treated device retained more than 80% PCE after 528 h under 50–70% humidity.

Cations

Larger cations for lattice doping. Small amount of larger cations such as $\text{CH}_3\text{CH}_2\text{NH}_3^+$ (EA^+), $(\text{CH}_3)_2\text{NH}_2^+$ (DMA^+) and $\text{C}(\text{NH}_2)_3^+$ (GA^+) can be doped into 3D perovskites to tune the crystal structure.⁵⁹ For example, incorporating larger EA^+ into MAPbI_3 can balance the lattice distortion strain and induce the tetragonal–cubic phase transition.⁶⁰ The secondary amine DMA can also reduce the lattice mismatch.^{61–64} When less than 20% DMA was used, the tetragonal–cubic phase transition was obvious without any sign of phase segregation.⁶⁴ The steric hindrance of larger DMA molecules in the framework stabilizes the I–Pb bond length and increases the rigidity of the perovskite lattice (Fig. 3a).⁶¹ The large size of DMA makes it incapable of forming 3D perovskites, but instead forms one-dimensional (1D) face-sharing perovskite DMAPbI_3 . Our group found that this 1D DMAPbI_3 is an appropriate intermediate phase for the realization of stable β -phase CsPbI_3 .⁶⁵

The incorporation of GA can stabilize the perovskite structure by reducing the degree of octahedral tilting. The lattice rigidity is enhanced and the defect formation energy is increased.^{62,66,67} The amount of larger cations incorporated into the lattice can be controlled by adjusting the sizes of the mixed A and X ions.⁶⁷ Zhou *et al.* found that the ratio of GA incorporation into a $\text{Cs}_{0.15}\text{FA}_{0.85}\text{Pb}(\text{I}_{0.85}\text{Br}_{0.15})_3$ based perovskite can be improved by increasing the amount of Cs and Br. This is because their smaller size can provide more space for the larger GA. GA introduced into the lattice can then interact with I and improve the stability of the perovskite and the PSC can maintain 80% efficiency under AM 1.5G for 8 h.

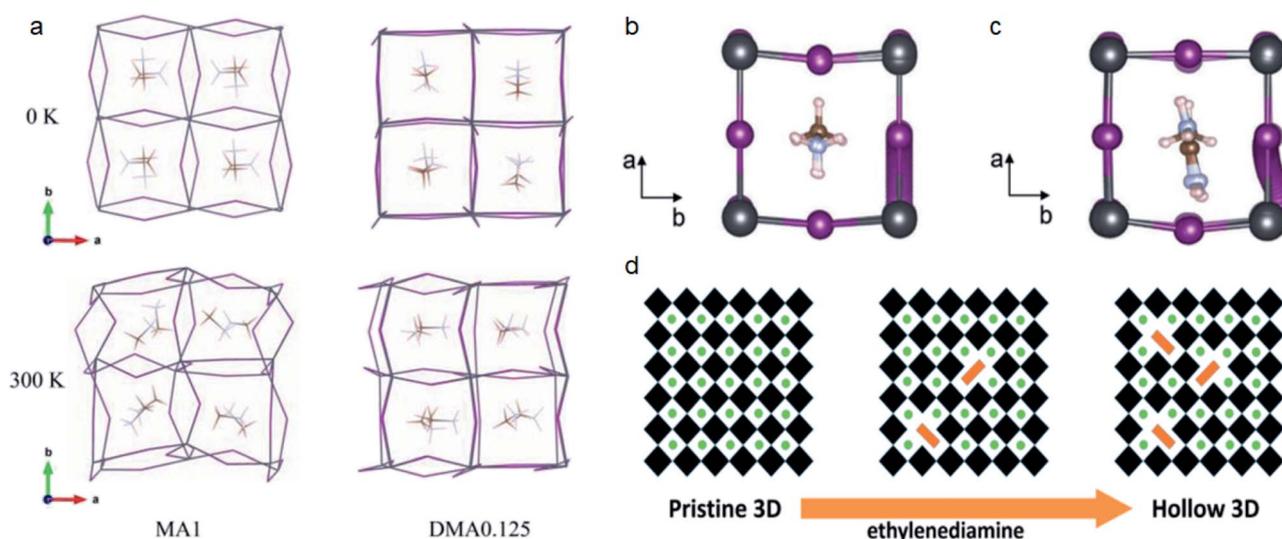


Fig. 3 (a) Rigidity changes for MAPbI_3 and $\text{DMA}_{0.125}\text{MA}_{0.875}\text{PbI}_3$ structures at different temperatures. Reproduced with permission from ref. 61. Copyright 2019 John Wiley and Sons. (b and c) Iodide migration pathway for MAPbI_3 and $\text{Ace}_{0.25}\text{MA}_{0.75}\text{PbI}_3$, respectively. Reproduced with permission from ref. 21. Copyright 2020 John Wiley and Sons. (d) Ethylenediamine induced formation of hollow 3D perovskites. Reproduced with permission from ref. 71. Copyright 2018 American Chemical Society.



Ion-migration in perovskites can be inhibited by the steric hindrance of larger cations.^{21,66} Tan *et al.* compared the activation energy for iodide migration in MAPbI₃ and CH₃C(NH₂)₂⁺ (Ace) incorporated perovskite Ace_{0.25}MA_{0.75}PbI₃.²¹ In the case of MA, I migrates directly from the lattice. When MA is partially replaced by Ace, the I migrated along an S-shaped curve because of the larger steric size of Ace (Fig. 3b and c). The inhibited I migration can help improve the stability of PSCs and the Ace based device can retain 55% of the initial efficiency after 2100 h under AM 1.5G.

Besides the all corner-sharing 3D structure, some larger cations can replace the position of Pb/I and form a deficient perovskite (d-perovskite).^{68–71} Leblanc *et al.* reported d-MAPbI₃ prepared by mixing MA⁺ and hydroxyethyl ammonium (HEA⁺) cations. These d-MAPbI₃ perovskites have similar absorption profiles to that of MAPbI₃ but with reduced Pb²⁺ concentration (Pb_{0.83}I_{2.83}), as well as improved flexibility and stability. Spanopoulos *et al.* also found that hollow 3D perovskite structures can be realized by introducing ethylenediammonium (EDA) cations (Fig. 3d).⁷¹ Addition of EDA can help improve the air stability of Sn based perovskites. The XRD pattern of MA_{0.6}-EDA_{0.4}Sn_{0.72}I_{2.84} remains unchanged for 9 days which indicates its greater stability than MASnI₃.

Bulky cations for surface passivation. Cations with bulky groups cannot partly replace native A-site cations of perovskites, but they show potential for surface and grain boundary defect passivation and can improve the hydrophobicity of the perovskite films.^{20,24,32,67,68,72–75} The transformation of CsPbI₃ from the cubic phase to orthorhombic phase can be suppressed by controlling the steric hindrance imposed by additives. Organic cations with a bulky alkyl chain as the organic terminal group

(OTG) can suppress octahedral tilting and increase the energy barrier of phase transition (Fig. 4a).²⁴ Bulky cation phenylethylammonium (PEA⁺) added into the precursor solution can also control nucleation for oriented crystallization and hinder phase transition *via* steric effects (Fig. 4b).²⁰ Besides addition to the precursor solution, bulky cations can also be used to post-treat perovskite films. Our group applied PEAI to post-treat CsPbI₃ films and found that PEAI would not interact with the CsPbI₃ film but form a defect-passivating organic termination layer (Fig. 4c).⁷⁶ This PEAI layer can inhibit the structural tilting through steric hindrance and improve the stability of CsPbI₃ films.

The alignment of bulky cations influences the water-resistivity of perovskite films. Wang *et al.* compared three phenylalkylamines: aniline, benzylamine, and phenethylamine. They found that benzylamine molecules are aligned perpendicularly on the film, therefore increasing the water penetration distance.⁷⁷ Post-treatment may also leave a crystalline layer of salt on the surface of the perovskite film, which could inhibit the charge transfer in the device. This unwanted aggregation can be prevented by applying cations with larger steric repulsion. For example, 4-*tert*-butyl-benzylammonium iodide (tBBAI) has a *tert*-butyl substitution, which could increase the steric hindrance and form an amorphous morphology on the perovskite film.⁷⁸

Some post-treatment cations, such as pyridine, not only passivate trap states but also may further react with perovskite films which could reduce the stability of PSCs. By rational molecular design, 2-Py with a long alkyl chain substituted at the *o*-position can effectively inhibit the further reaction between the perovskite and pyridine (Fig. 4d).⁷⁹ Since halogen migration

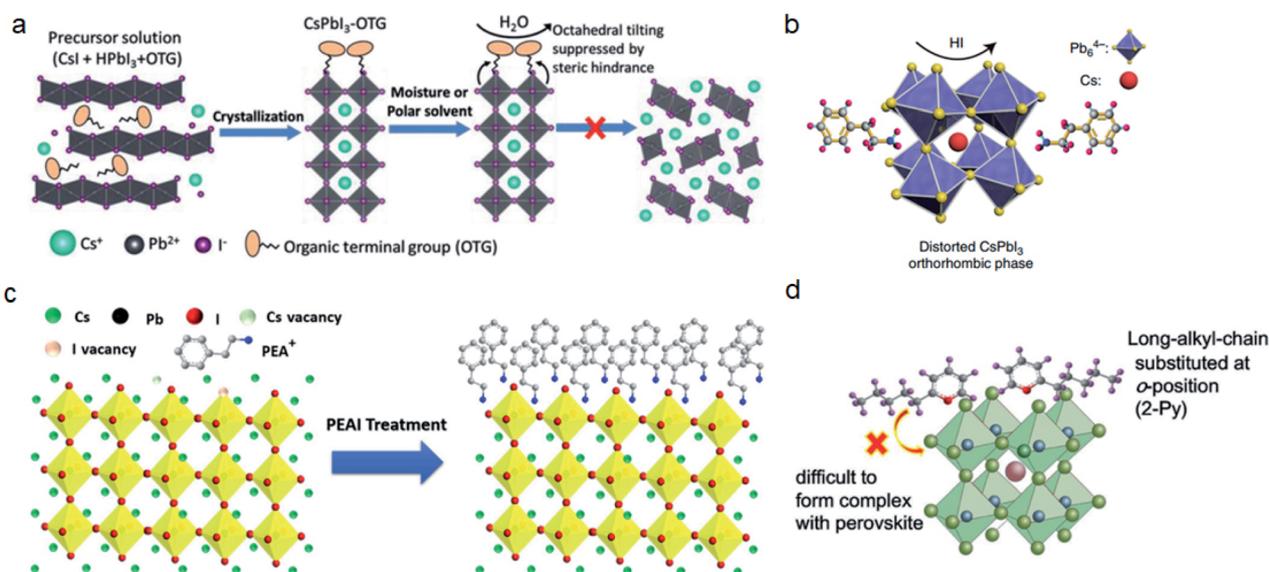


Fig. 4 (a) Octahedral tilting of CsPbI₃ is suppressed by the steric hindrance of OTG. Reproduced with permission from ref. 24. Copyright 2019 John Wiley and Sons. (b) PEAI additive induced formation of distorted CsPbI₃. Reproduced with permission from ref. 20. Copyright 2018 Springer Nature. (c) Surface termination of CsPbI₃ by PEAI post-treatment. Reproduced with permission from ref. 76. Copyright 2018 Elsevier. (d) The reaction of pyridine with perovskite is inhibited by using 2-Py with long alkyl chains. Reproduced with permission from ref. 79. Copyright 2016 John Wiley and Sons.



and its reaction with a metallic electrode is one of the main reasons for device degradation, using a hole-transporting layer (HTL) with high steric hindrance can restrain halogen migration and improve the stability of perovskite solar cells.⁸⁰ Li *et al.* compared two HTLs: HL-1 and HL-2, where HL-2 has a longer chain alkyl group compared with HL-1. They found that PSCs based on HL-2 show reduced iodide migration and improved stability.⁷³

Bulky cations are also commonly used in the synthesis of perovskite NCs. Perovskite NCs will undergo aggregation after long term storage; thus rational design of ligands with steric hindrance can help to maintain the colloidal and optical stability. Luo *et al.* reported a comparison between two kinds of branched ligands: (3-aminopropyl)triethoxysilane (APTES) and polyhedral oligomeric silsesquioxane PSS-[3-(2-aminoethyl)amino]propylheptaisobutyl substituted (NH_2 -POSS).⁸¹ The steric hindrance of APTES can inhibit the dissolution of NCs back into DMF, therefore preventing the penetration of protic molecules into the surface of perovskite NCs. As a result, NCs with uniform crystal size and improved long-term stability can be achieved and well-maintained.

Steric hindrances in low-dimensional perovskites

When larger or bulky cations are the main cations in the perovskite component, the perovskite structure cannot maintain a 3D framework and a low-dimensional perovskite may be formed.⁸² 2D perovskites with a corner-sharing inorganic layer structure, especially quasi-2D perovskites, have similar optical and electrical properties but much higher stability than 3D perovskites. This is because long chain cations can lock the unstable octahedral frame and inhibit the penetration of water molecules owing to the steric effect. As we mainly discuss the steric effect at the interface between organic layers and inorganic layers, we will combine the discussion on 2D perovskite, quasi-2D perovskite and 2D capping layers in this section.

Most commonly reported 2D perovskites are (100)-oriented, and they can be further classified into Ruddlesden–Popper (RP),

Dion–Jacobson (DJ) and alternating cations in the interlayer space (ACI) phases because of the different stacking modes and displacements of inorganic layers.^{83,84} RP phases have two spacer layers of large organic cations (Fig. 5a).⁸⁵ ACI phases have one layer of large size cations and one layer of small size cations. DJ phases have one spacer layer of organic cations between the inorganic layers.⁸⁶ Different structures are mainly affected by the chemical and spatial structure of cations. We will focus on the influence of steric hindrance on the structure and the optical and electrical-properties of 2D perovskites and perovskite based devices.

A large-sized spacer cation often consists of an ammonium and a terminal component such as an alkyl or aromatic group. Inorganic layers are bridged by cations by hydrogen bonding between the ammonium groups and octahedra, as well as van der Waals interactions between terminal groups. The steric effect caused by organic spacers will influence the configuration of inorganic layers and thereby the optical and electrical properties of 2D perovskite materials and optoelectronic devices. We will therefore discuss the steric effect on 2D perovskites with respect to the ammonium group, alkyl/aryl group and functional group.

Ammonium groups

Ammonium groups that can immobilize on the surface of inorganic layers by interacting with metal atoms have a direct influence on the properties of 2D perovskites. Stoumpos *et al.* studied different kinds of ammonium cations based on tin iodide.⁸⁷ They found that single ammonium $(\text{CH}_3)_m\text{NH}_{4-m}^+$ cations have very different structures. The number of methyl groups (m) on the N atom will influence the connectivity of the octahedra and face-sharing 1D perovskites will be formed when $m \geq 2$, such as in DMA^+ , $(\text{CH}_3)_3\text{NH}^+$ (TMA^+) and $(\text{CH}_3)_4\text{N}^+$ (NMe_4^+) (Fig. 5b). Very recently, our group reported that tetrabutyl ammonium (TBA^+) can intercalate into the CsPbI_3 film and form a 1D TBAPbI_3 layer which is quite hydrophobic.⁸⁸ TBAPbI_3 single crystals can remain unchanged even when immersed in water for 12 hours. In the case of $(\text{NH}_2)_n\text{CH}_{3-n}^+$,

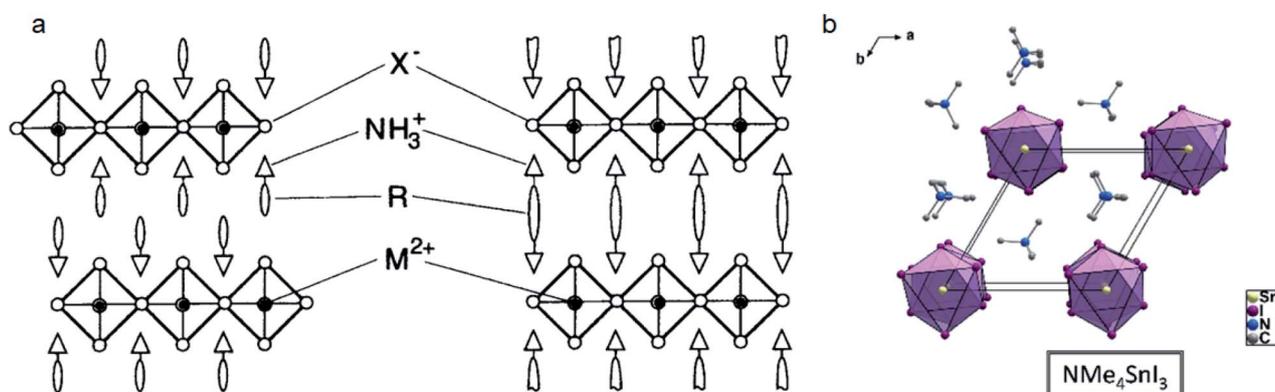


Fig. 5 (a) Crystal structure of the RP phase (left) and DJ phase (right) 2D perovskites. Reproduced with permission from ref. 85. Copyright 1999 John Wiley and Sons. (b) Crystal structures of NMe_4 based 1D perovskites. Reproduced with permission from ref. 87. Copyright 2017 American Chemical Society.



the number of ammonium groups will also influence the structure of 2D perovskites. These 2D perovskites will have a combination of corner-sharing and face-sharing structures which is defined as hexagonal perovskite polytypes such as GA^+ . When GA is mixed with MA , a $(GA)(MA)_nPb_{n-1}I_{3n+1}$ ($n = 1, 2, \text{ and } 3$) perovskite will form an ACI phase perovskite.^{89,90}

The position of the N atom will also influence the steric hindrance. Ruben and co-workers compared the substitution position of a N atom on pyridinium based cations.⁹¹ For example, 3BrPyH can form a 2D $(3BrPyH)_2PbI_4$ perovskite, while 2BrPyH and 4BrPyH will form 1D $(2BrPyH)PbI_3$ and $(4BrPyH)_4Pb_3I_{10}$ respectively. The steric size of the unconnected part will influence the face sharing or corner sharing mode and thereby define the structure of perovskites. Li *et al.* reported that 2-(aminomethyl)pyridine (2AMPY), 3-(aminomethyl)pyridine (3AMPY) and 4-(aminomethyl)pyridine (4AMPY) which have different N positions will form 2D, 0D and 1D perovskites, respectively.⁹² Mao *et al.* also studied DJ phase 2D perovskites based on 3-(aminomethyl)piperidinium (3AMP) and 4-(aminomethyl)piperidinium (4AMP) which have different N positions.⁸⁶ The position of N in the piperidinium organic backbone yields different hydrogen-bonding modes. The H-bonds in 3AMP and 4AMP are formed with terminal I^- and bridging I^- respectively. Therefore, the steric hindrance of the penetrated ring will induce different hydrogen-bonding modes and thereby influence Pb–I–Pb angles and optical band gaps.

Alkyl or aryl groups

Besides the ammonium groups that directly interact with inorganic layers, the steric hindrance of the alkyl or aryl groups connected with ammonium groups will also influence the perovskite properties.

Size of spacer cations. First of all, the size of cations can directly influence the interlayer distance and displacement between inorganic layers. Ishihara *et al.* investigated $(C_nH_{2n+1}NH_3)PbI_4$ with $n = 4, 6, 8, 9, 10, \text{ and } 12$, and found that the interlayer distance increases with longer carbon chains, from 15.17 Å ($n = 4$) to 24.51 Å ($n = 12$).⁹³ However, the optical properties remain almost unchanged, suggesting that there are weak interactions between inorganic layers with longer distances. Paritmongkol *et al.* found that the Pb–I tilting angle decreases as the number of CH_2 groups increases. They explained such an observation as the result of the stress between organic spacers being more absorbed by the longer chains and less transmitted to the inorganic slabs.⁹⁴

The interlayer distance can also directly influence the conductivity of 2D perovskites. Yu *et al.* compared the interlayer distances of cations with different sizes. The reduced distance can obviously improve the charge carrier mobility of perovskite films (Fig. 6a).⁹⁵ Polymer cations such as polyethylenimine (PEI) can provide a more compact organic layer between inorganic layers which can facilitate the charge transport of 2D films.⁹⁶

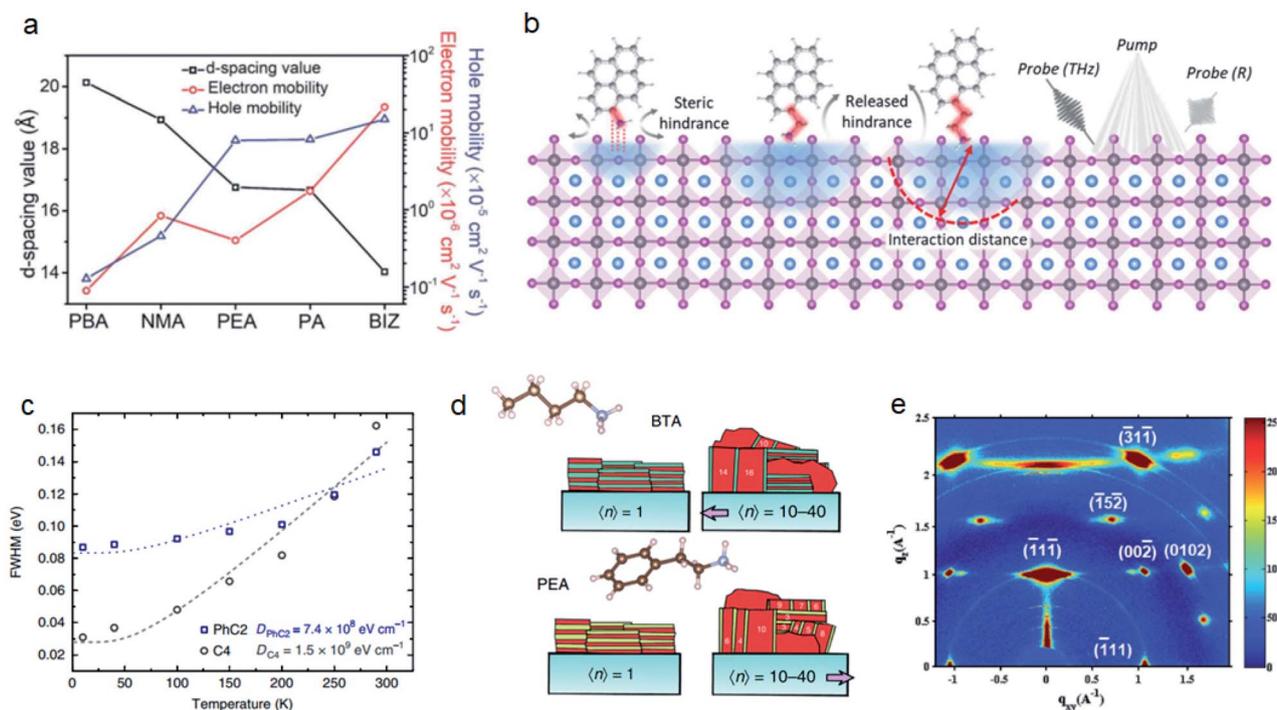


Fig. 6 (a) Mobilities of different 2D perovskites with different interlayer distances. Reproduced with permission from ref. 95. Copyright 2018 John Wiley and Sons. (b) Schematic illustration on steric hindrance of PRA, PRMA and PREA, their interaction distance and measurements by transient reflection spectroscopy. Reproduced with permission from ref. 36. Copyright 2021 American Association for the Advancement of Science. (c) The FWHM of PL spectra for PhC2 and C4 based 2D perovskites under different temperatures. Reproduced with permission from ref. 100. Copyright 2018 Springer Nature. (d) The orientations of BTA and PEA based 2D perovskites. Reproduced with permission from ref. 102. Copyright 2018 Springer Nature. (e) GIWAXS data of iso-C4 based 2D perovskites. Reproduced with permission from ref. 104. Copyright 2017 John Wiley and Sons.



The stronger steric hindrance of polymer cations can also make the 2D films more hydrophobic, thereby improving the moisture stability of perovskite solar cells. The length of the organic spacer will further influence the stability of perovskites.^{94,97} Koh *et al.* reported that $\text{CH}_3(\text{CH}_2)_3\text{NH}_3^+$ (C4) and $\text{CH}_3(\text{CH}_2)_7\text{NH}_3^+$ (C8) can both form 2D perovskites by post-treatment to the $\text{Cs}_{0.05}(\text{MA}_{0.17}\text{FA}_{0.83})_{0.95}\text{Pb}(\text{I}_{0.83}\text{Br}_{0.17})_3$ surface and C8 exhibits a better passivation effect and water repelling effect.⁹⁸ Spanopoulos *et al.* found that the higher tilting angles of $\text{CH}_3(\text{CH}_2)_4\text{NH}_3^+$ (C5) yield a more distorted structure. The resulting heat stability and light stability are both improved for C5 than for C4 and $\text{CH}_3(\text{CH}_2)_5\text{NH}_3^+$ (C6).⁹⁹ The length of spacer cations will influence cation penetration which influences the Pb–X–Pb bond angle. Du *et al.* synthesized several acene alkylamine cation based 2D perovskites.²⁵ They found that the number of methyl unit plays a more important role than the number of acene groups. 2-(2-Naphthyl)ethan ammonium (NEA) and PEA have larger NH_3 penetration and Pb–X–Pb bond angles than phenylmethyl ammonium (PMA) and 1-(2-naphthyl)methan ammonium (NMA). Very recently, Xue *et al.* compared three conjugated organic cations, pyrene-based ammonium (PRA), pyrene-based methyl ammonium (PRMA) and pyrene-based ethyl ammonium (PREA).³⁶ They found that the A-site cations can modify the band-edge state of perovskites. PRA without alkyl chains has shallow interaction depth because of the steric hindrance from the pyrene group. PREA with long alkyl chains can relieve the steric hindrance, thus leading to deeper vertical intercalation with inorganic layers (Fig. 6b).

Shape of spacer cations. The shape of spacer cations significantly changes the spatial distributions and thereby defines the properties of perovskite materials. Gong *et al.* explored several organic ammonium cations based on different aliphatic chain groups and phenyl groups: C4, $\text{CH}_3(\text{CH}_2)_{11}$ (C12), C_6H_5 (Ph), $\text{C}_6\text{H}_5\text{CH}_2$ (PhC), $\text{C}_6\text{H}_5(\text{CH}_2)_2$ (PhC2, also defined as PEA before) and $\text{C}_6\text{H}_5(\text{CH}_2)_3$ (PhC3).¹⁰⁰ They found that C4, C12, PhC and PhC2 can form 2D structures, while Ph and PhC3 result in non-2D structures. This phenomenon could be attributed to the steric effect caused by the bulky group which inhibits the corner connectivity of the octahedra as we discussed before.⁸⁷ A PhC2 based perovskite single crystal has much higher photoluminescence quantum efficiency (PLQE) than C4 because of the more rigid crystal structure and reduced electron–phonon interactions measured by the deformation potential (Fig. 6c). Recently, time-domain *ab initio* simulations also demonstrated that fluctuating $(\text{C}_4)_2\text{PbBr}_4$ has a much higher nonradiative recombination rate than rigid $(\text{PEA})_2\text{PbBr}_4$.¹⁰¹ During the formation of 2D perovskite films, the rigidity difference between C4 (defined as BTA in Fig. 6d) and PEA will also make C4 more easily redistributable which will increase the formation of in-plane distributed 2D perovskites.¹⁰² The structure of 3AMPY with aromatic rings is more rigid than 3AMP with the aliphatic ones and they have a reduced interlayer distance.¹⁰³ The exciton binding energy is also decreased because of the increased dielectric constant. The rigidity caused orientation difference is also studied by grazing incidence wide-angle X-ray scattering (GIWAXS) when replacing C4 with iso-C4, which has short branched chains (Fig. 6e).¹⁰⁴

Functional groups

Large size spacer cations can be further modified by other elements or groups such as halides (F/Cl/Br/I), –OH, MeO–, unsaturated bonds, and so on. These functional groups will also have steric interactions with the inorganic framework or between the organic layers.²⁵ Mitzi and co-workers compared the crystal structure of $(2\text{-XC}_6\text{H}_4\text{C}_2\text{H}_4\text{NH}_3)_2\text{SnI}_4$ (X: F, Cl, Br) with halides of different sizes.¹⁰⁵ They found that $(2\text{-ClC}_6\text{H}_4\text{C}_2\text{H}_4\text{NH}_3)_2\text{SnI}_4$ and $(2\text{-FC}_6\text{H}_4\text{C}_2\text{H}_4\text{NH}_3)_2\text{SnI}_4$ have a similar *gauche* conformation while $(2\text{-BrC}_6\text{H}_4\text{C}_2\text{H}_4\text{NH}_3)_2\text{SnI}_4$ exhibits *anti* conformation. The larger Br will cause structural distortion effects such as reduced Sn–I–Sn angles and blue shift on the absorption spectra (Fig. 7a). Their recent work revealed that functional group induced steric effects will influence the formation energies and thermal properties.¹⁰⁶ Knutson *et al.* also reported that the steric impact of organic cations can tune the bandgap of perovskites by more than 1 eV.¹⁰⁷ Steric bulk near the ammonium head such as 2-bromophenethyl ammonium (2-BrPEA), 2-trifluoromethylphenethylamine (2-CF₃PEA) and 1-pyreneethylamine (1-PYREA) will inhibit the penetration of ammonium into the perovskite sheet, thereby reducing in-plane distortion.

Functional groups with unsaturated bonds may form a cross-linked structure and improve the connectivity of organic layers, which can efficiently improve the conductivity and stability of 2D perovskites. For example, double bonds and triple bonds can lead to the formation of a cross-linked structure upon UV light or thermal annealing.^{108,109} The vinyl group at the *para* position of benzyl ammonium (BzA) can form new covalent bonds under UV light, which can improve the operational stability and reduce the hysteresis of PSCs (Fig. 7b).¹⁰⁸ Cross-linking combined with oxygen or iodine doping will also change the energy bandgap of organic layers from 3.0 to 1.4 eV and induce a thousand-fold conductivity increase.¹⁰⁹ In some conditions, in order to obtain tunable perovskite quantum wells, functionalized groups with methyl groups impart steric effects, which can inhibit intermolecular π – π interactions.¹¹⁰

Similar to the position of ammonium groups, the position of functional groups will also influence the steric effects. Recently 4-fluorophenethyl ammonium (4FPEA) was shown to have potential for high efficiency 2D perovskite solar cells because of its enhanced π -orbital overlap and decreased interlayer distance.^{26,111–114} Several groups have found that the position of monofluorination on PEA will influence the organic cation packing arrangements (Fig. 7c–f).^{26,115} The formation energy of 4FPEA is lower than that of other positions and will promote the formation of phase pure, flat and oriented 2D perovskites.

Steric hindrance stabilization

Quaternary cations

Compared with the commonly used ammonium cations, quaternary ammonium cations with larger steric hindrance were shown to have a significant effect on passivating the MA⁺ vacancy and improving the ambient stability of PSCs.^{35,116} Zheng *et al.*¹¹⁶ found that quaternary ammonium halides such as



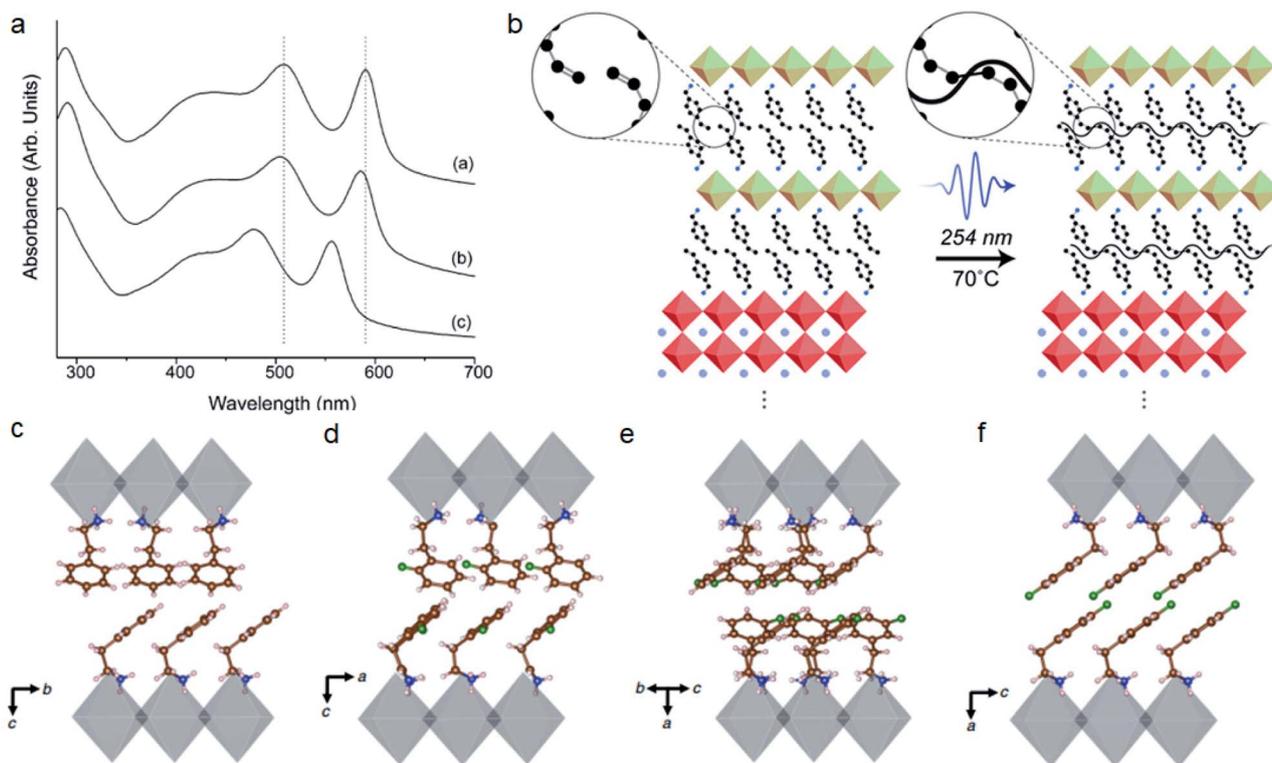


Fig. 7 (a) Absorption spectra of the $(2\text{-XC}_6\text{H}_4\text{C}_2\text{H}_4\text{NH}_3)_2\text{SnI}_4$ (X: F, Cl, Br) perovskite. Curve a, F; curve b, Cl and curve c, Br. Reproduced with permission from ref. 105. Copyright 2003 American Chemical Society. (b) Formation of new covalent bonds under UV light for BzA with a vinyl group at the *para* position. Reproduced with permission from ref. 108. Copyright 2019 American Chemical Society. (c–f) Organic cation packing arrangements of PEA₂PbI₄, 2FPEA₂PbI₄, 3FPEA₂PbI₄ and 4FPEA₂PbI₄, respectively. Reproduced with permission from ref. 26. Copyright 2019 Springer Nature.

choline iodide (CHI) and choline chloride (CHCl) can passivate both positive traps and negative traps on the surface of perovskites. This passivation can improve V_{oc} significantly for more than 100 mV. Our group also found that CHI can passivate the cracks and pinholes in $\beta\text{-CsPbI}_3$ films.⁶⁵ The PCE of all inorganic PSCs can be improved to 18.4%. The PSC can be retained at 92% of the initial PCE under 1 sun for 500 hours.

Several works have shown that light or temperature induced lattice expansion will improve the efficiency and stability of perovskite solar cells.^{117,118} As we have mentioned before, the steric size of ammonium will influence the tilting angles of Pb–I–Pb. This steric effect on the perovskite lattice can inhibit ion migration and improve the structural stabilities of perovskites. Recently, we reported that the passivation of phenyltrimethylammonium iodide (PTAI) to MAPbI₃ can form a mixed-2D PTAMAPbI₄ perovskite, which can construct a dynamic barrier for MA⁺ migration and thereby inhibit the degradation of perovskite films (Fig. 8a). The MAPbI₃ film exhibits exceptional stability with unchanged properties even when soaked in PTAI/IPA solution.²² The work by Yangi *et al.* also reported the formation of mixed-2D perovskites with cations of phenylformamidinium (PFA).¹¹⁹ We also discovered that bifunctional PTABr post-treatment to CsPbI₃ can further improve the phase stability by gradient Br doping and the PCE can reach 17.06%.¹²⁰ Furthermore, PTACL treatment to CsPbI₃

can passivate the surface traps effectively and boost the PCE to more than 19%.¹²¹

The steric hindrance provided by bulky quaternary cations can also prevent the interaction between the lone pair of electrons of water molecules and the perovskite surface. Yang *et al.* compared five bulky cations and classified them into three types: MA has one methyl group, tetra-methyl ammonium (TMA) and hexadecyl trimethyl ammonium (CTA) have three or four methyl groups, tetra-ethyl ammonium (TEA), TBA and tetra-hexyl ammonium (THA) have four longer alkyl groups. Among them, TEA with stronger steric hindrance can induce larger Pb–I tilt angles and reduce the tendency of water adsorption.⁷² The steric effect of quaternary cations is also certified by the fabricated PSCs based on the P3HT hole transport layer. Jung *et al.* prepared a wide-bandgap layer on a perovskite film by *n*-hexyl trimethyl ammonium bromide post-treatment (Fig. 8b).³⁵ The stability is significantly improved and the PSC can maintain 80% efficiency after 1008 h under a humidity of 85%.

Bifunctional cations

Diammonium cations. Diammonium cations with two ammonium groups have recently shown to have potential for efficient and stable PSCs as they can form DJ phase perovskites.^{27,86,122,123} As discussed before, 2D RP phase perovskites



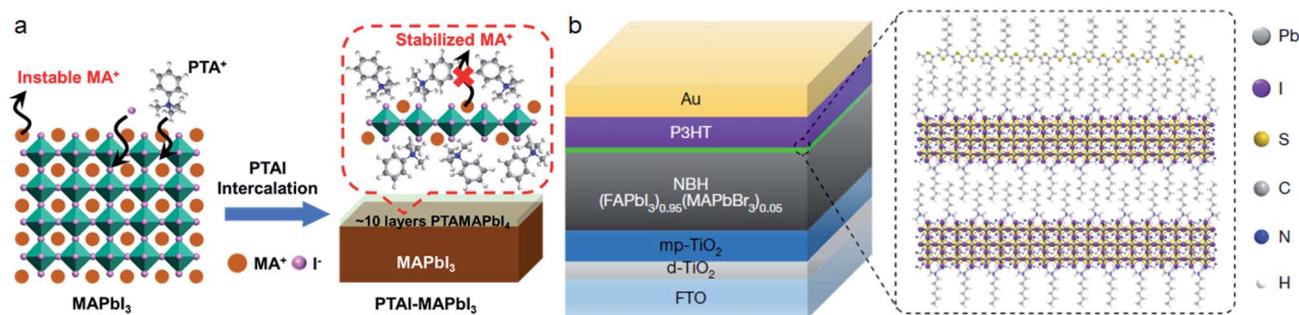


Fig. 8 (a) Schematic mechanism for the stabilization of MA⁺ by PTAI intercalation. Reproduced with permission from ref. 22. Copyright 2020 John Wiley and Sons. (b) PSC based on a P3HT hole transport layer with an *n*-hexyl trimethyl ammonium bromide (HTAB) based wide-bandgap perovskite as the interfacial layer. Reproduced with permission from ref. 35. Copyright 2019 Springer Nature.

based on single ammonium cations are over-layered by organic layers with van der Waals interactions which are weaker than ionic interactions. DJ phase perovskites based on diammonium cations can link two inorganic layers directly by ionic bonds, hence increasing the binding energy and dissociation energy (Fig. 9a).^{28,122,123} There is only one organic layer between inorganic layers in the DJ phase perovskite. Therefore, reducing the barrier length of quantum wells can improve the charge carrier mobility of the 2D perovskites.^{124,125} The conductivity comparison between RP phase BA₂PbI₄ and DJ phase BDAPbI₄ also shows that the DJ phase perovskite has better carrier transport and energy transfer properties (Fig. 9b).^{124,126,127} Ahmad *et al.* studied the moisture and light stability of a 1,3-propanediamine (PDA) based DJ phase perovskite. They found that eliminating the van der Waals energy gap in a propylamine (PA) based RP phase perovskite can significantly improve the stability of PSCs.¹²⁸

Black phase CsPbI₃ perovskites are believed to be unstable at room temperature as they tend to transform to the yellow phase perovskite that is unsuitable for photovoltaics.¹²⁹ In 2017, our group reported that EDA based 2D EDAPbI₄ can cross link CsPbI₃ crystals and the extrusion by the steric effect of EDA can inhibit the phase transition.²⁷ The film condition can keep unchanged upon annealing at 100 °C for a week (Fig. 9c). The DJ phase EDAPbI₄ can also maintain the conductivity of CsPbI₃ films and CsPbI₃·0.025EDAPbI₄ based PSC achieved a record PCE of 11.8% at that time.

Tin-based perovskites are attractive as nontoxic, efficient PSCs because of their lead-free composition and broader absorption properties. However, Sn(II) in tin based perovskite can be easily oxidized to Sn(IV) and induce the degradation of perovskites. The strong binding energy and rigid structure of DJ phase perovskites show potential for realizing stable tin based perovskites. Chen *et al.* reported that (4AMP)(FA)₃Sn₄I₁₃ based PSCs can obtain a PCE of 4.22% and retain 90% PCE for 100 h under AM 1.5G.¹³⁰ They suspected that the synergistic effect of reduced vacancy formation, improved binding energy and charge transport together improved the efficiency and stability of tin based PSCs. Li *et al.* reported that adding 1,4-butanediamine (BDA) ligands into a tin based perovskite can inhibit the oxidation of Sn²⁺.¹³¹ BDA has a large penetration depth between

NH₃⁺ and terminal iodides. This steric effect can reduce the electronic interaction and hydrogen bonding, thereby reducing the octahedral distortion, resulting in the DJ phase 2D perovskite having a similar band gap to 3D perovskites.^{124,131}

The energy quantum well (QW) distribution and orientation will influence the optical and electrical properties of DJ phase perovskites.^{132,133} Zheng *et al.* compared 1,3-propanediamine (PDA), BDA, 1,5-pentamethylenediamine (PeDA) and 1,6-hexamethylenediamine (HDA) with 3, 4, 5 and 6 carbon chains respectively.^{134,135} They found that PDA and BDA that have shorter chain lengths can yield oriented and shallow distributed QWs. The chain length of diammonium cations will also influence QW barrier thickness, charge carrier mobility and the 3D-to-2D transformation activation energy.^{122,136} Lu *et al.* compared the perovskite structures induced by PDA, HDA and EDA. They found that PDA and HDA will induce 3D to 2D transformation more easily than EDA because of their longer chain lengths (Fig. 9d).¹³⁷ Zhao *et al.* also compared a series of diammonium iodides, BDA, 1,8-octamethylenediamine (ODA) and 2,2'-(ethylenedioxy)bis(ethylamine) (EDBE).¹³⁶ They found that BDA and EDBE can induce the transformation of 3D MAPbI₃ to 2D perovskite by post treatment, while ODA only passivates the grain boundary without phase transformation. They suspected that the unique *anti-gauche* isomerization of ODA elevated the 3D to 2D activation energy.

Other terminal groups. Besides the diammonium cations, some cations with multiple functional groups, such as phosphonic acid ammonium and amino acid, can also form a crosslinked structure.^{138–140} Cross-linked structures can improve the binding energy and charge carrier transfer between perovskite crystals. Li *et al.* applied bifunctional butylphosphonic acid 4-ammonium chloride (4-ABPACl) to modify the surface of perovskite crystals.¹³⁸ Both the ammonium group and phosphonic acid group will interact with the perovskite surface and crosslink perovskite crystals (Fig. 10a). This kind of cross-linked perovskite can help the formation of smooth perovskite films and facilitate the contact between the perovskite and TiO₂ substrate. 4-ABPACl based devices can maintain 80% of the initial PCE at 85 °C for 350 h.

Han and co-workers applied 5-ammoniumvaleric acid iodide (5-AvaI) in MAPbI₃ based mesoscopic perovskite solar cells and



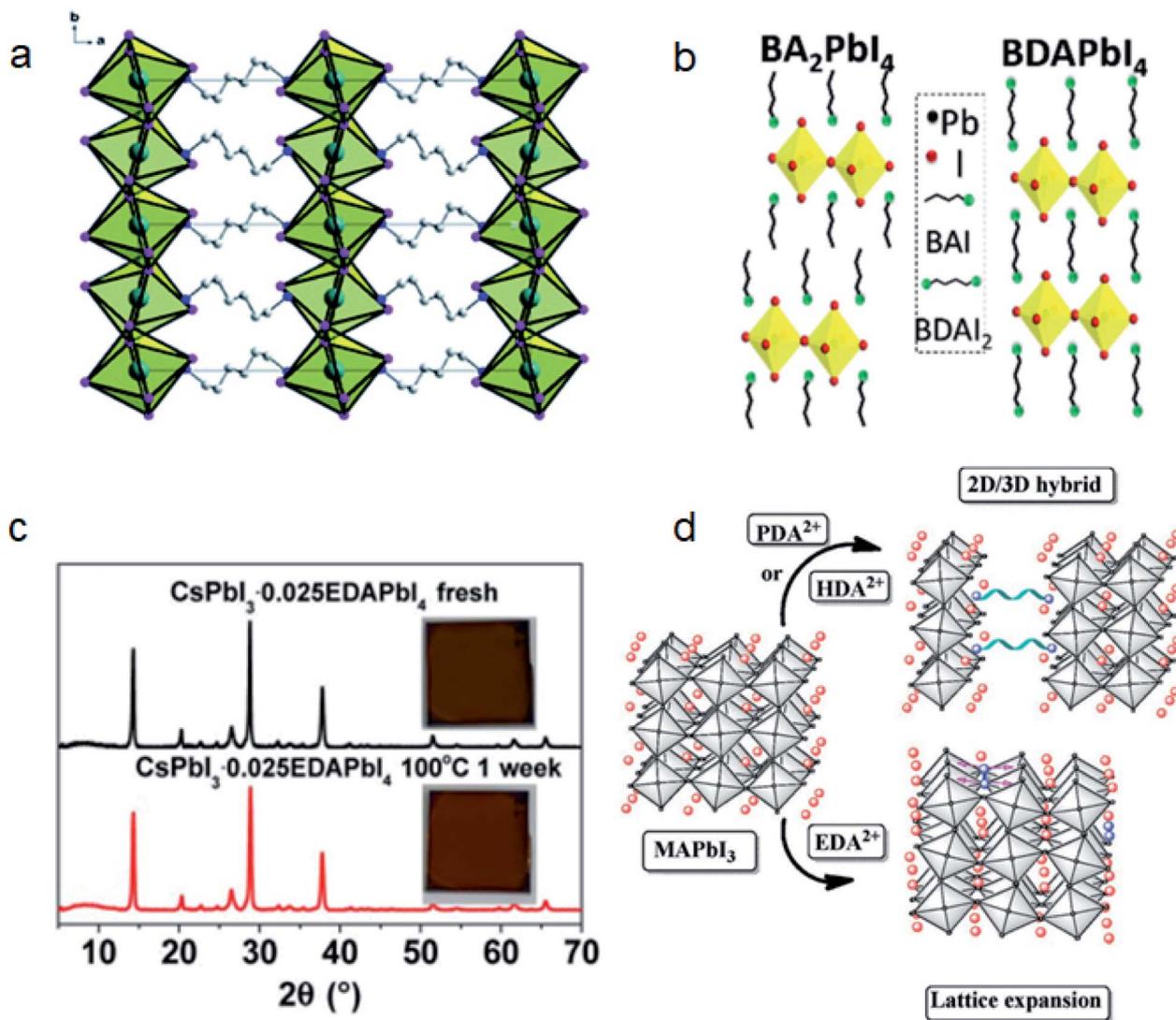


Fig. 9 (a) Schematic structure of HDAPbI₄. Reproduced with permission from ref. 122. Copyright 2016 Royal Society of Chemistry. (b) Schematic structures of BA₂PbI₄ and BDAPbI₄. Reproduced with permission from ref. 126. Copyright 2018 IOP Publishing Ltd. (c) XRD patterns of fresh and aged CsPbI₃-0.025EDAPbI₄ perovskite films at 100 °C for 1 week. Reproduced with permission from ref. 27. Copyright 2017 American Association for the Advancement of Science. (d) Effect of diammonium cations with different lengths on the formation of 2D perovskites or lattice expansion. Reproduced with permission from ref. 137. Copyright 2017 John Wiley and Sons.

achieved a PCE of 12.8%.¹⁴⁰ The effect of 5-AvaI was considered to be the result of passivating traps and improving the contact with mesoporous TiO₂. Our group applied 5-Ava in the MAPbBr₃ system and further revealed that the COO⁻ and NH₃⁺ groups on 5-Ava can cross-link perovskites *via* Pb-COO bonds and the interaction between NH₃⁺ and [PbBr₆]⁴⁻ units. We found that 5-AvaBr will be incorporated into the perovskite lattice and forms the RP phase Ava_xMA_{1-x}PbBr₃ structure as reported in previous work.¹⁴⁰⁻¹⁴² In contrast, 5-Ava will form the crosslinked DJ phase Ava(MAPbBr₃)_n structure because the COOH on 5-AvaBr will not form such Pb-COO bonds (Fig. 10b and c). This cross-linked structure will sterically reduce the interlayer distance and improve the PLQE of the *in situ* formed MAPbBr₃ QD films up to 80% (Fig. 10d).¹³⁹ The 5-Ava also shows potential for cross-linking and stabilizing the FASnI₃ based perovskite by reducing the Sn⁴⁺ content.¹⁴³

Steric hindrance design for PeLEDs

Perovskite materials are also attractive for light-emitting diodes benefiting from the high PLQE and high emission purity. PSCs and PeLEDs have a similar layered device structure, *i.e.*, anode/HTL/perovskite layer/ETL/cathode. The generation of the excited state is the key step for them. They all require low trap density and reduced nonradiative recombination. That is to say, efficient PSCs are usually efficient PeLEDs. There are also some different requirements between PSCs and PeLEDs because the excited states are separated by an electrical field in PSCs, but are recombined to emit photons in PeLEDs. Therefore, tuning of steric hindrance in perovskite materials can enable us to design specific films for different optoelectronic devices.

In order to separate charge carriers effectively, PSCs usually have a type-II band alignment which can drive electrons and



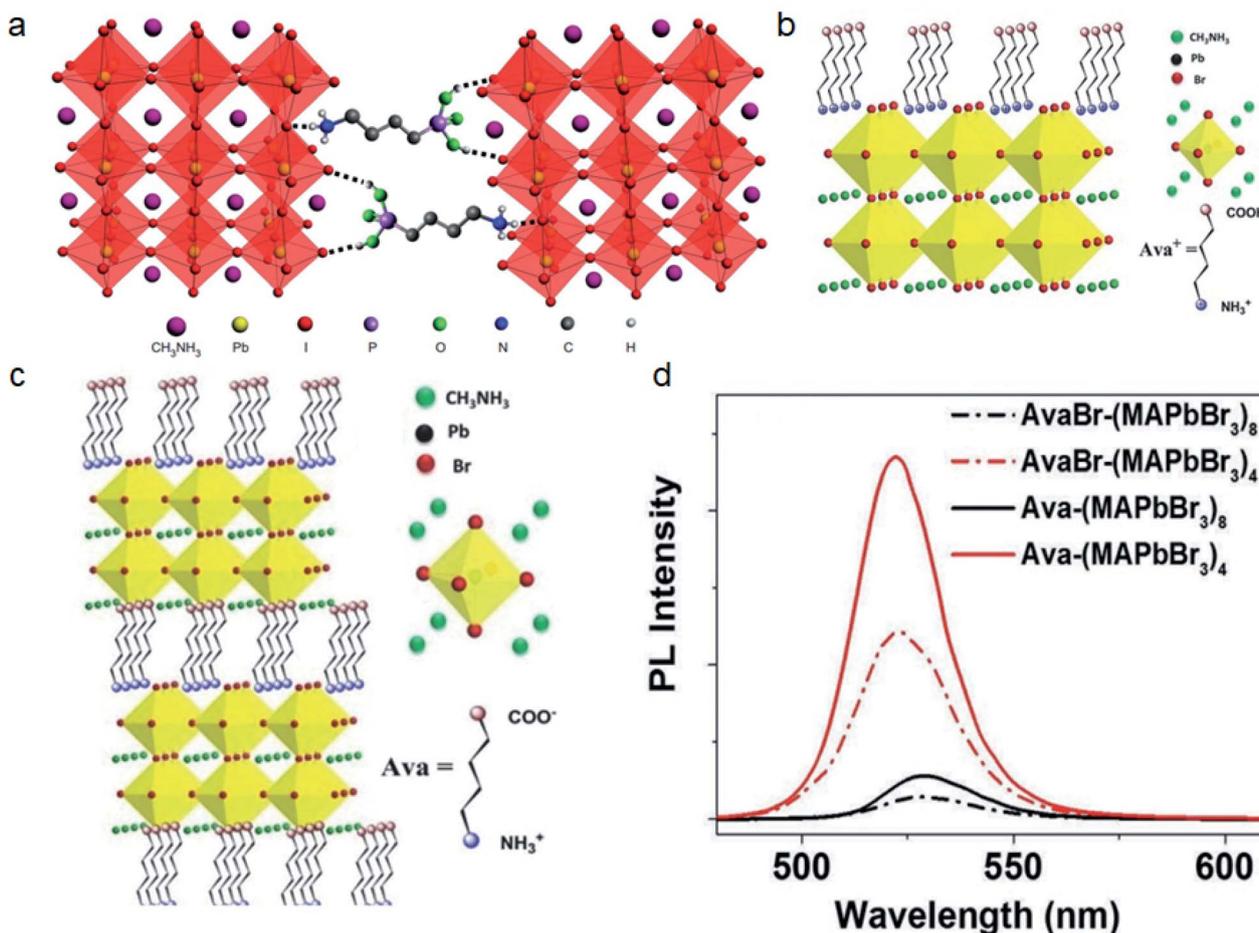


Fig. 10 (a) 3D perovskites cross-linked by 4-ABPAI. Reproduced with permission from ref. 138. Copyright 2015 Springer Nature. (b) 5-Ava perovskites cross-linked by Pb–COO bonds and interactions between NH₃⁺ and [PbBr₆]⁴⁻ units. (c) Single linked 5-AvaBr on 3D perovskites. (d) PL spectra of 5-Ava and 5-AvaBr treated perovskite films. Reproduced with permission from ref. 139. Copyright 2017 John Wiley and Sons.

holes to different electrodes.¹⁴⁴ In the case of LEDs, a type-I band alignment can accumulate electrons and holes to increase the charge carrier density and to improve the radiative recombination rate and PLQE.^{145–147} Yuan *et al.* changed the n value in $\text{PEA}_2(\text{CH}_3\text{NH}_3)_{n-1}\text{Pb}_n\text{I}_{3n+1}$ and found that a film prepared with a certain n value will consist of a variety of quasi-2D perovskites with different n values.¹⁴⁸ This type-I structure can act as an energy funnel and significantly improve PLQE (Fig. 11a). Besides the addition of different kinds of cations, the steric hindrance induced band alignment can also be modified by changing the concentration of cations. Quintero-Bermudez *et al.* found that the ligand density at the surface influences the work function of the material and shows tunable type-I or type-II properties.¹⁴⁶

The bifunctional effect of 5-Ava is also applied in the fabrication of high efficiency PeLEDs. Cao *et al.* found that 5-Ava not only reduces the trap density of perovskite films but also undergoes a dehydration reaction with ZnO/polyethylenimine ethoxylated (PEIE) to form an organic layer which reduces the leakage current, thus boosting the EQE up to 20.7%.⁸ They reported that 6-aminocaproic acid (6-ACA) and 7-aminoheptanoic acid (7-AHA) with longer chain lengths can also improve the

efficiency of PeLEDs. Wang *et al.* found that γ -aminobutyric acid (3-ABA) which has a similar structure to 5-Ava can control the distribution of n values in the $\text{PEA}_2\text{Cs}_{n-1}\text{Pb}_n\text{Br}_{3n+1}$ perovskite and suppress the formation of thicker quantum wells.¹⁴⁹ Bifunctional 3-ABA can also passivate trap states of perovskite films and achieve a high EQE of 6.3% for blue PeLEDs.

It's more difficult to realize stable PeLEDs because of the higher current density and internal electrical field under working conditions. 2D perovskite materials with bulky cations can significantly improve the stability of PeLEDs. Wang *et al.* reported that adding 1-naphthylmethylamine iodide (NMAI) into FAPbI_3 precursor solution can also realize quasi-2D perovskite films with multiple quantum wells.¹⁵⁰ The PeLED reaches a T_{50} lifetime (time to 50% of the initial brightness) of 2 h under a current density of 10 mA cm^{-2} , while the control device without NMAI can only work for 1 min. The steric hindrance from large cations contribute to both type-I energy transport and improved stability. DJ-phase 2D perovskites with cross-linked inorganic layers also show potential for stable PeLEDs. Shang *et al.* fabricated 2D PeLEDs with 1,4-bis(aminomethyl)benzene (BAB) and PEA.²⁸ BAB based DJ phase perovskites exhibit higher molecule dissociation energy than



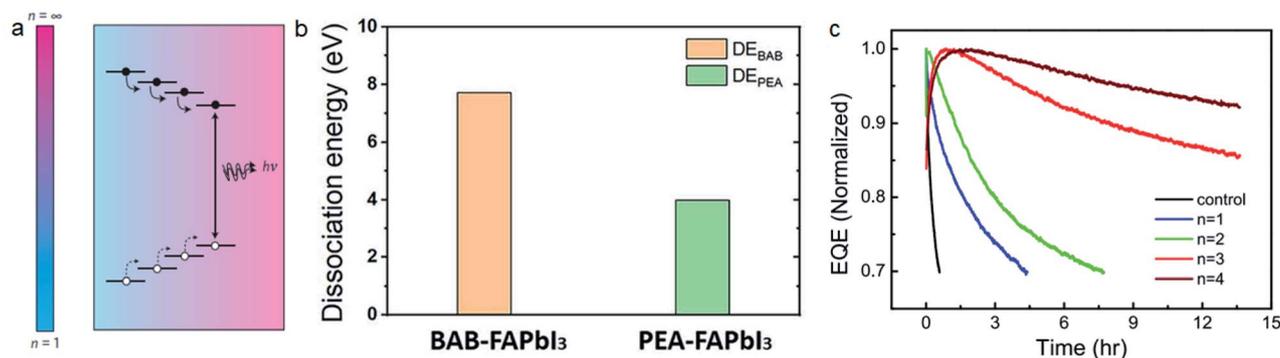


Fig. 11 (a) Charge transport in MQW based films. Reproduced with permission from ref. 148. Copyright 2016 Springer Nature. (b) Molecule dissociation energy of BAB and PEA based perovskites. Reproduced with permission from ref. 28. Copyright 2019 American Association for the Advancement of Science. (c) Stability of PeLEDs with different phenylalkylammonium iodide molecules. Reproduced with permission from ref. 34. Copyright 2021 Springer Nature.

PEA based RP phase perovskites (Fig. 11b). BAB based PeLEDs have a T_{50} lifetime of 100 h under a current density of 25 mA cm^{-2} . Surface passivation is another way for improving the stability of PeLEDs while keeping the high conductivity of 3D perovskite materials.¹⁵¹ Guo *et al.* passivated FAPbI₃ films with four kinds of phenylalkylammonium iodide molecules of different alkyl chain lengths (phenylmethylammonium iodide (PMAI, $n = 1$), phenethylammonium iodide (PEAI, $n = 2$), phenpropylammonium iodide (PPAI, $n = 3$), and phenylbutanammonium iodide (PBAI, $n = 4$)).³⁴ They found that the lifetime of PeLEDs is significantly improved with longer alkyl chain lengths (Fig. 11c). Density functional theory (DFT) calculations show that phenylalkylammonium with longer chain lengths exhibits larger steric hindrance and inhibits the iodide migration. The PPAI treated PeLED reached a peak EQE of 17.5% and a record T_{50} lifetime of 130 h under a current density of 100 mA cm^{-2} .

Conclusion and outlook

In conclusion, the past decade has witnessed the rapid development of metal halide perovskite materials and perovskite-based optoelectronic devices. The key structure for metal halide perovskites is the [BX]₆⁴⁻ octahedron. This structure endows perovskites with “flexible” properties and can be easily formed. However, this “flexibility” also makes the perovskite structure more prone to degradation. Among various strategies to stabilize the perovskite structure, steric hindrance mainly stabilizes the cations and halide anions by inhibiting their migration. Other than the well-known tolerance factor tuning, steric hindrance can relieve the lattice strain of the crystal structures of 3D perovskites, which effectively improves the intrinsic phase stability of perovskites. Surface termination and incorporation of additives in 3D perovskites by large-sized cations can also passivate surface defects and suppress the degradation of perovskites caused by humidity. In low-dimensional perovskites, the steric hindrance of spacer cations can significantly change the spatial distance, orientation, conductivity and optoelectronic properties of layered perovskites. There are several specific cations that can

significantly stabilize perovskites such as quaternary cations, diammonium cations and bifunctional cations. Steric hindrance has proved to be a promising strategy for both efficient and stable perovskite optoelectronic devices. However, the current development of steric hindrance in perovskites is still in the early stages without enough rational molecular design and detailed fundamental mechanism understanding.

The concept of steric hindrance has been widely used in selective chemical reactions, especially asymmetric synthesis, and tremendous molecular design samples and theoretical rules have been developed in this field.^{152–154} Therefore, we believe that there is still enough potential for making good use of steric hindrance with a rational molecular design strategy to further improve the efficiency and stability of perovskite devices. Especially, the combination of previous knowledge on asymmetric syntheses based on steric hindrance and the latest advances in *in situ* material characterization and theoretical investigations would help us to further explore the mechanism of steric hindrance. With ongoing fundamental investigations and interdisciplinary studies, we believe that more and more molecular and crystal structure design strategies based on steric hindrance will be realized to help achieve the goal of high efficiency and high stability for perovskite materials and their optoelectronic applications.

Author contributions

Y. Z. conceived the idea of the perspective. Y. M., Y. C., H. C. and X. W. wrote and edited the initial manuscript. Y. M. and Y. Z. revised the manuscript.

Conflicts of interest

There are no conflicts to declare.

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