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Magic guanidinium cations in perovskite solar cells: from bulk to interface

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Guanidinium (GA⁺) has been proven as the multifunctional cation in halide perovskite solar cells (PSCs), attracting considerable attention due to its improved power conversion efficiencies (PCE) and operational stability. This review introduces the elementary properties of the GA cation and GA-based perovskite. Then, we summarize the development history of utilizing quanidinium materials in PSCs, including composition engineering, additive engineering, and interface engineering, accompanied by their optimization mechanisms. Moreover, we propose a perspective on the potential opportunities in this popular field and challenges to the future design of guanidinium-based PSCs with enhanced performance.

1. Introduction

To deal with the issues of energy crisis and greenhouse effectinduced climate change, it is essential to exploit renewable energy technologies, in which photovoltaics (PV) has been one of the most promising candidates to generate inexhaustible and clean solar energy. Among the various PV technologies,

halide perovskite solar cells (PSCs) have attracted extensive interest due to their efficient power conversion efficiency (PCE), high voltage output, low fabrication cost, and simple processing techniques. 1-3 The optimization strategies towards perovskite solar cell devices such as composition engineering, additive engineering, and interface engineering help advance the performance improvements, which recently reached over 25% for typical FAPbI₃ PSCs.^{4,5} Nevertheless, the scalability and stability of such composition remain an issue because of the defects at the interfaces and grain boundaries and intrinsic phase segregation with ion migration.6-8

To overcome the abovementioned drawbacks, bulky or longchain organic cations such as butylammonium (C₄H₉NH₃ [BA⁺]), phenethylammonium (C₈H₉NH₃⁺ [PEA⁺]), and octylammonium (CH₃(CH₂)₇NH₃⁺ [OA⁺]) could be introduced to passivate the various trap states and enhance the stability of the



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Table 1 The ionic radius and t of common A-site cations ^{14,15,21–23}

A-site cations	Formula	Radius (pm)	t	μ _e (Debye)
Potassium	K ⁺	137	0.74	10.82
Ammonium	$[NH_4]^+$	146	0.76	0
Rubidium	Rb ⁺	152	0.78	11.48
Cesium	Cs^+	167	0.84	11.69
Methylammonium	$[(CH_3)NH_3]^+$	217	0.91	2.17
Azetidinium	$[(CH_2)_3NH_2]^+$	250	0.98	2.51
Formamidinium	$[NH_2(CH)NH_2]^+$	253	0.99	0.60
Dimethylammonium	$[(CH_3)_2NH_2]^+$	272	1.02	1.80
Ethylammonium	$[(C_2H_5)NH_3]^+$	274	1.03	4.49
Guanidinium	$\left[C(NH_2)_3\right]^+$	278	1.04	0
Tetramethylammonium	$[(CH_3)_4N]^+$	292	1.07	0
Ethylenediammonium	$[NH_3(CH_2)_2NH_3]_2^{2+}$	333	1.15	_
Butylammonium	$[(C_4H_9)NH_3]^+$	393	1.28	_
Phenylethylammonium	$[C_2H_4(C_6H_5)NH_3]^+$	431	1.35	1.5
Butane-1,4-diammonium	$[NH_3(CH_2)_4NH_3]_2^{2+}$	457	1.41	_
Hexane-1,6-diammonium	$[NH_3(CH_2)_6NH_3]_2^{-2+}$	582	1.67	_

Note: the t is calculated based on the chemical formula APbI₃; the ionic radius of Pb and I are 220 pm and 119 pm, respectively. "-" means the absence of relative information elsewhere.

perovskite. 9-11 These cations demonstrate significantly improved operational stability under multiple environments (e.g., moisture, thermal, and illumination). Unfortunately, their positive effects on the photovoltaic performance are not satisfying and are primarily limited by the hindered carrier transport because of the relatively low conductivity. 12,13 Additionally, these cations hardly form and diffuse into the 3D perovskite crystal structure due to the large ionic radius.

Among various kinds of cations, GA⁺ is substantially larger in ionic radius (278 pm) as compared to the commonly used MA⁺ (217 pm) and FA⁺ (253 pm) cations while smaller than BA⁺ (393 pm) and PEA+ (437 pm). Still, it does not form a 3D perovskite structure as the sole A-cation in terms of the t value (see detailed information in Table 1).14,15 Regardless of the

phase of the GA⁺-containing perovskites, positive interactions are usually found endowed by GA+, as long as it could be introduced in the system. GA has nearly zero dipole character, which has been proven to be effective in suppressing biasinduced ionic mobility and hysteresis characteristics. 16,17 In addition, GA⁺ can offer a lone pair of electrons to specifically passivate the imperfections in the surface and bulk perovskite films, i.e., ionic vacancies, uncoordinated Pb, and/or Pb clusters. 18 Most importantly, owing to the plentiful ammonium groups in the molecular structure, GA+ could provide more hydrogen bonds to coordinate with the surrounding halides, especially undercoordinated I⁻ ions firmly, thus increasing the formation energy of halide vacancies and improving the perovskite structural stability. 19,20 As a consequence, the charge carrier lifetime and photovoltaic performance, such as the cell's open circuit voltage (V_{OC}) and fill factor (FF) are expected to be ameliorated.

The GA cation-based material was first introduced in the dye/liquid electrolyte interface as an additive by Jeanbourquin et al.²⁴ to improve the performance of dye-sensitized solar cells (DSSCs) in 2014. The additive could eliminate the interfacial dye-I₂ complexes, increasing the collection efficiency and thus the photocurrent. Inspired by this, in 2016, Marco et al. 25 also added the GA-based additive firstly into the perovskite precursor to suppress nonradiative recombination defects of iodide vacancies in the bulk and iodine species at grain boundaries, resulting in superior performance with PCEs of more than 17% and a minor $V_{\rm OC}$ loss of 0.43 eV. Encouraged by the progress, GA-based materials seemed to be good candidates for PSCs, which have acquired rapid development in the past few years. In this passage, representative endeavors on the GA⁺ species for PSC developments have been summarized, which could be categorized into three strategies of compositional engineering, additive engineering, and interface engineering, among which



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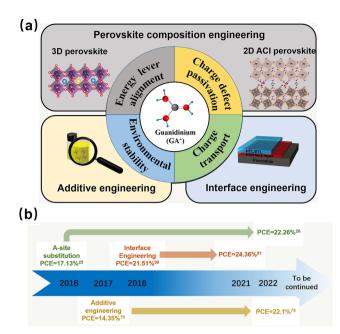


Fig. 1 (a) The schematic diagram of strategies for the GA+ cation in the PSCs. (b) The development of GA+-based PSCs.

the first one could be further classified into 3D compositional or 2D alternating-cation-interlayer (ACI) engineering types according to the perovskite matrix (Fig. 1). Although there have been reports of further PCE performances of over 24%, their simultaneous long-term stabilities against thermal, illumination or other harsh conditions have yet to be presented. 20,26 Therefore, how to simultaneously achieve both excellent efficiency and stability of the device would be the significant challenge in the future for GA-based PSCs. This systematic review could help us identify the fundamental mechanisms and the origins of the existing problems. Conversely, it further opens access to rational designs to overcome the challenges and fabricate more stable and efficient PSCs.

2. Structure and optical properties

Metal halide perovskites may crystallize into zero-dimensional (0D), 1D, 2D, or 3D perovskite structures, as shown in Fig. 2. The tolerance factor (t) defines the geometrical criteria for the formation of an ideal cubic perovskite structure:

$$t = \frac{(R_{\rm A} + R_{\rm B})}{\sqrt{2}(R_{\rm B} + R_{\rm X})}$$

where R_A , R_B , and R_X are the ionic radii for the corresponding ions of cations in A sites, metals in B sites, and halides in X sites, respectively.

When t is in the range of ~ 0.8 –1.0, a stable 3D perovskite structure can be formed. The 2D perovskite is preferred for inclusion in the case of t > 1, which generally adapts a formula of $(A')_m(A)_{n-1}B_nX_{3n+1}$. Here, A' is structurally featured with divalent (m = 1) or monovalent (m = 2) cations, and n indicates the number of layers consisting of $[BX_6]^{4-}$ octahedra. 1D perovskite micro/nanocrystals generally present a well-defined fiber or ribbon-like morphology. Distinct from the uniform 3D growth, the self-assembly of the precursors into a 1D nanostructure reacting in a particular solution or vapor environment, as well as the ordered texture aligned by external forces such as casting and evaporation techniques or the periodically confined nano-templates, consists of the two prerequisites for practical 1D perovskite applications.28 0D perovskites have the formula $A_4''BX_6$ consisting of A"—a monovalent organic or inorganic cation, and isolated BX₆ octahedral anions or separated clusters from each other by the surrounding inorganic or organic cations without certain composition and periodic bands' formation. Therefore, the t of 0D perovskite often deviates from 1. 29

Within the 3D framework, the inorganic layers of the $[BX_6]^{4-}$ octahedra are connected via sharing each of their six corners, while the A cations occupy voids between each layer.³⁰ In the case of B and X, respectively, representing pure Pb²⁺ and I⁻, the notorious B site oxidation and phase segregation problem can

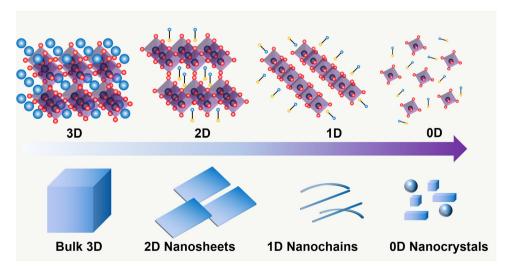


Fig. 2 Schematic representation of the halide perovskite with different dimensionalities from 3D to 0D at the structural level (top) and by morphology (bottom)

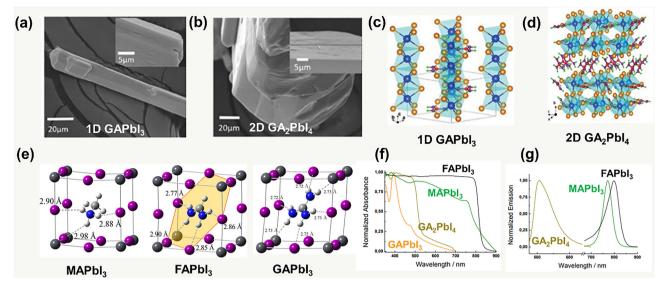


Fig. 3 SEM images of (a) GAPbI₃ and (b) GA₂PbI₄ (high-resolution SEM image as inset). Crystal structure of (c) 1D GAPbI₃ and (d) 2D GA₂PbI₄. Reproduced with permission. 32 Copyright 2021, ACS Publishing Group. (e) Pbl₆ inorganic framework and N-H···I hydrogen bonds in MAPbl₃, FAPbl₃, and GAPbl₃. Reproduced with permission.³³ Copyright 2018, ACS Publishing Group. (f) UV-vis absorption and (g) steady-state photoluminescence spectra of MAPbl₃, FAPbl₃, GAPbl₃, and GA₂Pbl₄. Reproduced with permission.³⁶ Copyright 2016, Wiley-VCH.

be avoided. When the A site is Cs^+ , MA^+ , FA^+ , and their alloys, t falls typically in the range of 0.8-1, which ensures a 3D cubic/ tetragonal structure. Compared to cations with moderate radii in the 3D perovskites, the A-site in a 2D perovskite occupied by large radii organic cations such as PEA⁺ and BA⁺ would result in t > 1. They cannot fit into the voids between the octahedral units. After insertion, the A-site cations expand the van der Waals gaps, making the inorganic layers discrete, along with the broadened band gaps.³¹

Different from MA⁺ and FA⁺, GA⁺ is highly symmetric and non-polarized, and the lattice can be stabilized by its abundant hydrogen bonds. However, because of the large cation radii leading to the high lattice stress, the 3D [PbI₆]⁴⁻ octahedra networks break into the 1D ribbon (Fig. 3a) of the GAPbI₃ form with edge-sharing octahedra or GA₂PbI₄, which is a 2D layer structure (Fig. 3b) with corner-sharing octahedra.³² The crystallographic data of GAPbI₃ and GA₂PbI₄ are summarized, as shown in Fig. 3c and d, and Table 2. In the GA-Pb-I system, GA in the unit cell is approximately orientated along the $(11\overline{1})$ plane and connects with six I ions to generate N-H···I hydrogen bonds within the PbI₆ octahedra, in the most stable structure. Furthermore, compared to MAPbI₃ and FAPbI₃ perovskites, the GA-based perovskite has more N-H···I hydrogen bonds in the unit cell (the number of N-H···I hydrogen bonds in MAPbI₃, FAPbI₃, and GAPbI₃ are 3, 4, and 6, respectively) (Fig. 3e).³³ Different conditions could regulate the formation of diverse dimensional GA-Pb-I perovskites. A solution process, such as spin-coating, is expected to deposit the 1D GAPbI₃ and 2D GA₂PbI₄ perovskite film with a stoichiometric ratio of PbI₂ and GAI.

The photophysical property of the GA-based perovskite, in contrast to MAPbI₃ and FAPbI₃, has an apparent blue shift of the absorption, which is also ascribed to the disruption of the

Table 2 Single-crystal data for the structures of GAPbl₃ and GA₂Pbl₄³⁷

Chemical formula	$GAPbI_3$	$\mathrm{GA}_{2}\mathrm{PbI}_{4}$
Formula weight	641.93	834.97
Crystal system	Monoclinic	Orthorhombic
Space group	Pnma	$P2_1/n$
a (Å)	11.9866(5)	9.2440(3)
b (Å)	4.4740(2)	26.9511(11)
c (Å)	20.8652(10)	12.7155(3)
α (deg)	90	90
β (deg)	90	91.482(2)
γ (deg)	90	90
Volume (ų)	1118.96(9)	3166.82(17)
Z	4	8
$ ho_{ m calc}$ (g cm $^{-3}$)	3.81	3.503

coherent 3D PbI₆ octahedra by the large size of the GA cation.³⁴ Thereby, GA-based perovskite, for example, GAPbI₃, provides a higher band gap (E_g) of 2.25 eV.³⁵ The photoluminescence spectra of both GAPbI₃ and GA₂PbI₄ also show the apparent shifts toward higher energy, which is in agreement with the results of the absorption spectra³⁶ (Fig. 3f and g). Due to the wide bandgap, the GA-based perovskite could not generate a sufficient visible light-induced carrier. Moreover, GAPbI3 and GA₂PbI₄ usually exhibit poor uniformity in film processing, which makes them ineffective for direct use as photoactive layers in PSCs. However, it would become effective in the case of GA as a supplementary composition, as introduced by the above-mentioned three strategies in Fig. 1.

Composition engineering

Perovskite compositional engineering could improve the PCE and stability of the PSCs. Until now, the commonly used

perovskite composition has been changed from the initial $MAPbI_3$ to the so-called triple cation $(MA_xFA_{1-x})_m$ $Cs_{1-m}Pb$ - $(I_{1-\nu}Br_{\nu})_3$ perovskite, which recently illustrated a comparable development in PCE to the record. Although it has a large radius, GA⁺ could also be partially introduced into the A-site, which brings more lattice stability, bulk defect passivation from lone-pair electrons and hydrogen bonds, band structure adjustment, among other properties. The alloying could drastically improve the charge extraction and transport, and resistance against harsh conditions, resulting in enhanced device performance and stabilities. 38,39 According to the dimension, the composition engineering will be divided into 3D perovskite composition engineering and 2D ACI composition engineering, whose PCEs have reached as high as 22.17% and 22.26%, respectively.

3.1 3D perovskite composition engineering

Although the MA-and FA-based PSCs commonly exhibit excellent PCEs, MA and FA cations suffer from the degradation of perovskite film under light and thermal exposure due to the low stability of the perovskite film and phase impurity. 40,41 Generally, the incorporation of GA, similar to Rb and Cs, could stabilize the perovskite phase, leading to enhanced performance and stability. 41,42

 $MA_{1-x}GA_xPbI_3$. When GA^+ partly replaces MA^+ in MAPbI₃, it can be inserted into the crystal lattice, enhancing device performance and stability. Marco et al.25 firstly mixed MAI and GAI as precursors in different ratios. They found that GA⁺ could effectively eliminate the under-coordinated iodine species at grain boundaries and within the bulk through their hydrogen bonding capability, and it could weaken orbital overlap between Pb s and I p orbitals. 46 Furthermore, Kubicki et al. 47 evidenced the rapid reorientation process for GA^+ ($t \le$ 18 \pm 8 ps) in the $GA_xMA_{1-x}PbI_3$ phase by solid-state magic angle spinning (MAS) NMR, much faster than that of MAPbI₃ (113 \pm 25 ps), which was dedicated to stabilization of the charge carriers for improving the PL lifetimes and device performance.

Besides the improved photovoltaic performance, the longtime thermal and photo instabilities are also of vital concerns. Jodlowski et al. 43 incorporated the GA⁺ into the MAPbI₃ perovskite film, forming the stable 3D perovskite structure. The SEM results showed that when x < 25%, the grain size was increased without unwanted phase segregation, whereas higher amounts of GA^+ (x > 25%) tended to form the 1D $GAPbI_3$ perovskite (Fig. 4a and b). Besides the improved film quality, the employed GA+ offered more H bonds within the inorganic framework to stabilize the 3D perovskite structure, resulting in improved efficiency and resistance to thermal and light corrosion.

Moreover, GA⁺ has a significant effect on the inhibition of ion migration, which is responsible for the hysteresis and poor stability of the device. 48,49 Ferdani et al. 50 reported that MAPbI₃ with 5% GA^+ substitutions had higher activation energy ($E_a =$ 0.78 eV) for iodide ion conduction than that of pristine MAPbI₃ $(E_a = 0.44 \text{ eV})$. Furthermore, Minussit et al.⁴⁹ found that GA^+ could slow down the relaxation process of space charge, which was two orders of magnitude longer, resulting in retarded ion migration. Meanwhile, incorporating GA+ has an impact on the low-temperature phase transitions of MAPbI3. When MA was substituted by 2 mol% GA^+ , the β (tetragonal) to α (cubic) phase transition happened under a reduced temperature of 278 K.

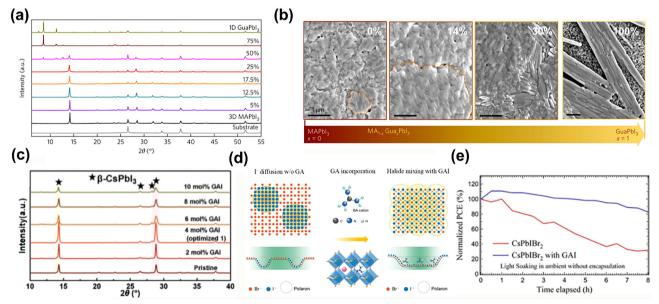


Fig. 4 (a) Normalized XRD patterns of perovskite films with different amounts of GA. (b) Top-view SEM images of $MA_{1-x}GA_xPbI_3$ with x = 0, 0.14, 0.30, and pure GAPbl₃. Reproduced with permission.⁴³ Copyright 2017, Nature Publishing Group. (c) XRD data for perovskite films with different GAI doping. Reproduced with permission. 44 Copyright 2022, Wiley-VCH. (d) Schematic illustration of the halide ion distribution in the $GA_xCs_{1-x}PbIBr_2$ lattice in $response \ to \ a \ low-power \ laser \ (1.27 \ mW). \ (e) \ Normalized \ PCEs \ of \ CsPblBr_2- \ and \ GA_{0.1}Cs_{0.9}PblBr_2- \ based \ PSCs \ under \ continuous \ illumination \ for \ 8 \ h \ with \ response \ various \ response \ various \ response \ response \ various \ response \ respon$ the AM 1.5G solar spectrum in the ambient environment (~55% relative humidity) without encapsulation. Reproduced with permission. 45 Copyright 2022, Wiley-VCH.

 $FA_{1-x}GA_xPbI_3$. GA^+ could also partly substitute FA to form $FA_{1-x}GA_xPbI_3$ to stabilize the α -FAPbI₃ phase. GA^+ matches better with FA⁺ than MA⁺ in ionic radius, which would help in not triggering the big change of lattice parameters and tfactors. Giorgi et al. 17 theoretically studied the thermodynamic stability and electronic properties of FA1-rGArPbI3 through density functional theory (DFT) calculations. The incorporation of GA⁺ could induce a zero dipole moment because of the D3h symmetry of the GA⁺ cation. The incorporation of GA⁺ into FAPbI₃ at a relatively low content would not interrupt the 3D perovskite structure, but the symmetry of the cuboctahedral volume (in which the FA+ cations reorients) had been lowered in the case of GA_{0.25}FA_{0.75}PbI₃.⁴⁷

 $Cs_{1-x}GA_xPbI_3$. Phase separation and transition into a photoinactive δ -phase is the dominant cause of relatively low PCE and instability of the Cs-based perovskite, which is mainly induced by ion migration in Cs-based systems. Incorporating GA⁺ could suppress this undesired tendency.

Wang et al.44 added GAI in the triple halide-mixed CsPbTh (Th = either I, Br, or Cl). The GA composition guaranteed the stabilization of the crystal lattice, achieving enhanced thermal stability (Fig. 4c). After that, the optimal device was obtained by PbAc2 introduction as a nucleating additive, accompanied with GCA spin-coated for interface passivation. Eventually, it delivered a PCE of 19.37% with a high FF of 82.7% and $V_{\rm OC}$ of 1.17 V, and the operational stability was also dramatically improved. A more profound mechanism of the I⁻ ion migration suppression

was proposed, suggesting that the strengthened hydrogen bonds between the GA⁺ and the [PbI₆]⁴⁻ octahedron were responsible for reducing the local lattice distortion, which was the origin of ion migration⁴⁵ (Fig. 4d). The retardation of light-induced ion migration via GA⁺ incorporation could retain over 80% under 8 h continuous illumination, showing superior light stability to the pristine CsPbIBr₂ (Fig. 4e).

Mutiple-APbX₃ (A= $MA^+/FA^+/Cs^+/GA^+$; X= $I^-/Br^-/CI^-$). It is significant to tune the tolerance factor by employing multications for improved performance and stability of the perovskite. 51,52 Similar to GA+ in MA_{1-x}GA_xPbI₃ and FA_{1-x}-GA_xPbI₃, GA⁺ in the multi-cation system could further mediate the tolerance factor and stabilize perovskite structure, along with the improvements in the charge carrier lifetime.

Hysteresis has raised the concern of accurate performance measurement in practice, while its origin remains an open question. GA+ participation could inhibit hysteresis, with diverse interpretations of the mechanisms. Wang et al. 53 added GAI in a FA/Cs-based perovskite to regulate the energy alignment and extend the PL lifetime. Consequently, the low content of GA⁺ in the FA/Cs-based perovskite induced a slight lattice expansion, but could reduce interfacial charge accumulation, thus removing the J-V hysteresis while improving PCE (Fig. 5a and b). On the other hand, Li et al. 54 explored the MA_{0.8}FA_{0.1-} GA_{0.1}PbI₃ system, and concluded that the GA⁺ incorporation into the perovskite structure could build the potential barrier of iodide ion migration and improve the perovskite crystalline,

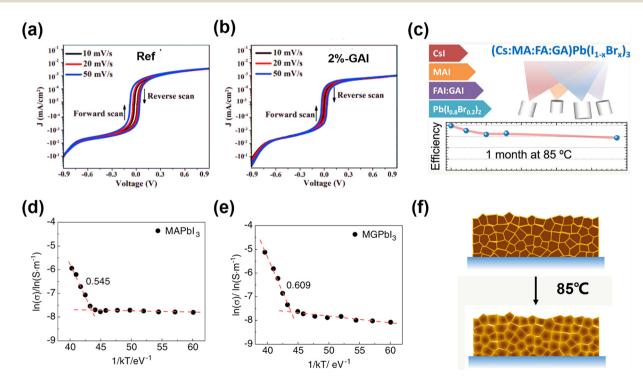


Fig. 5 Dark J-V curves in logarithm scaled current representation at different scan rates for (a) the control device and (b) the device with 2% GAI. Reproduced with permission. 53 Copyright 2019, Wiley-VCH. Temperature-dependent ion conductivity of the (d) MAPbI₃ (control) and (e) MA_{0.9}GA_{0.1}PbI₃ (MGPbl₃). Reproduced with permission.⁵⁴ Copyright 2022, Wiley-VCH. (c) J-V curves of quadruple-cation CsMAFAGA-based device at 85 °C. (f) Schematics of GA⁺ dynamic incorporation in vacuum-deposited, quadruple cation CsMAFAGA perovskite films. Reproduced with permission.⁵⁵ Copyright 2022, ACS Publishing Group

achieving the best PCE of over 22% and 19.18% for an active area of 14 cm² (Fig. 5d and e).

Due to the limited number of thermal sources and complexity of the Cs/MA/FA triple-cation-based perovskite, it is difficult to insert the GA⁺ into the lattice to form the CsMAFAGA quadruplecation perovskite via solution process and conventional vacuum process. However, Susic et al.55 realized the uniform alloying by sublimation of FA and GA in the same thermal source during vacuum deposition, given that the sublimation properties of FAI and GAI were similar. The GA+ initially accumulated at the grain boundaries. However, upon subsequent annealing, it progressively penetrated the crystal and induced a lower-symmetry phase perovskite structural change to relieve the strain stress. Ultimately, stabilization of the cubic phase with a larger unit cell volume was observed (Fig. 5c and f). The CsMAFAGA-based device could maintain the perovskite structure, and no parasitic phase (PbI₂, δ-CsPbI₃, and δ-FAPbI₃) formed at 85 °C after 37 days, demonstrating no degradation or segregation after stabilization by GA cations. Simultaneously, it kept over 75% original PCE at the 85 °C for about a month.

Although the GA⁺ in A-site compositional engineering plays an important role in the double-, triple- and even multiplecation-based perovskite system for the ion migration inhibition, perovskite structure stabilization, charge defects passivation, and band structure change, it inevitably increases the complexity of the perovskite system for processing. In addition, introducing a high concentration of GA+ (more than 20%) could facilitate low-dimensional perovskite impurities, such as 1D GAPbI₃ and 2D GA₂PbI₄ rather than uniform alloying in 3D structure, which would be imperfections in the interface and grain boundaries. The limited doping concentration becomes the bottleneck in composition engineering, so other guanidinium-involving approaches for perovskite modification would be the alternative for performance enhancement.

3.2 2D ACI composition engineering

Similar to the 2D Ruddlesden-Popper (RP) and Dion-Jacobson (DJ) perovskite, the GA-based 2D perovskites usually present excellent hydrophobicity and ambient stability. RP and DJ have the same chemical formula of $A_{m}'A_{n-1}B_{n}X_{3n+1}$. Here, A'

Table 3 The single-crystal data and structure refinement for various reported ACI perovskite^{34,56,57}

	$\mathrm{GAMAPbI}_4$	$\mathrm{GAMA}_2\mathrm{Pb}_2\mathrm{I}_7$	$\mathrm{GAMA_3Pb_3I_{10}}$	$GACsPbI_4$	$GACsPbBr_4$	$GACs_2Pb_2Br_7$	$\mathrm{GAFAPbI}_4$
Formula wt Temperature (K)	807.0 293	1426.9	2046.9	907.79 300	719.83	1299.66	819.95 298
Color Morphology	Yellow Plate Orthorhombic	Dark red	Black	Red None	Yellow None	Yellow None	Brown Platelet Monoclinic
Space group Volume (ų) Unit cell dimensions (Å)	Imma 1507.7(2) a = 6.4286(6)	Bmm2 2508.2(3) a = 6.3729(4) b = 12.5435(8) c = 31.376(2)	Imma 3473.8(5) a = 6.3520(7) b = 12.4378(9) c = 43.970(3)	()			C2/m $4678.7(7)$ $a = 26.948(2)$ $b = 12.8189(11)$ $c = 14.4080(12)$
Z $ ho_{ m calc}$ (g cm $^{-3}$) $ ho$ (mm $^{-1}$) $F(000)$ Crystal size (mm 3) $ ho$ range (deg) Index ranges	4 3.5551 19.349 1392 $0.1111 \times 0.0225 \times 0.0167$ $3.27-29.23$ $-8 \le h \le 8$ $-17 \le k \le 16$	3.7788 22.017 2424 $0.0511 \times 0.179 \times 0.0100$ $2.6-29.27$ $-8 \le h \le 8$ $-17 \le k \le 17$	3.9138 23.395 3464 $0.0733 \times 0.0372 \times 0.0193$ $2.78-29.29$ $-8 \le h \le 8$ $-17 \le k \le 16$		$-17 \le k \le 16$		$\begin{array}{c} 12\\ 3.492\\ 18.710\\ 4224\\ 0.32\times0.06\times\\ 0.04\\ 3.006-52.738\\ -33\leq h\leq 33\\ -16\leq k\leq 16\\ 18.16\leq k\leq 16$
Reflections collected	$-25 \le l \le 25$ 7172	$-42 \le l \le 42$ 11855	$-57 \le l \le 60$ 12119	$-1/ \le l \le 1/$ 33586	$-25 \le l \le 24$ 7142	$-16 \le l \le 16$ 11917	$-18 \le l \le 18$ 21674
Independent reflections	784 $[R_{\text{int}} = 0.1277]$	2595 $[R_{\text{int}} = 0.0563]$	1810 [$R_{\text{int}} = 0.1261$]	$4767 [R_{int} = 0.0637, R_{\sigma} = 0.0432]$	1094 [$R_{\text{int}} = 0.0406$, $R_{\sigma} = 0.0295$]	1814 [$R_{\text{int}} = 0.0473$, $R_{\sigma} = 0.0370$]	$5009 [R_{int} = 0.0386, R_{\sigma} = 0.0319]$
Completeness to θ	99%	99%	99%	None	None	None	None
Data/ restraints/ parameters	784/3/2	2595/9/82	1810/5/59	4767/18/109	1094/0/40	1814/0/58	5009/0/144
Goodness-of- fit on F^2	3.18	2.19	2.80	1.004	1.041	1.035	1.017
Final R indexes $[I \ge 2\sigma(I)]$ Final R indexes $[all\ data]$ Largest $[all\ data]$ Largest $[all\ data]$	$wR_2 = 0.1356$	$R_1 = 0.0555$ $wR_2 = 0.1099$ $R_1 = 0.0898$ $wR_2 = 0.1126$ 5.69/-1.65	$R_1 = 0.0597$ $wR_2 = 0.1254$ $R_1 = 0.0863$ $wR_2 = 0.1270$ $3.32/-3.20$	$R_1 = 0.0389$ $wR_2 = 0.0722$ $R_1 = 0.0835$, $wR_2 = 0.0888$ 1.36/-1.55	$\begin{array}{c} R_1 = 0.0223 \\ \text{w} R_2 = 0.0519 \\ R_1 = 0.0261 \\ \text{w} R_2 = 0.0535 \\ 0.84 / -0.88 \end{array}$	$R_1 = 0.0321$ $wR_2 = 0.0772$ $R_1 = 0.0407$ $wR_2 = 0.0823$ $2.21/-1.52$	$R_1 = 0.0409 \\ wR_2 = 0.0945 \\ R_1 = 0.0552 \\ wR_2 = 0.1016 \\ 1.87/-1.27$

is structurally featured with bulky or long-chain monovalent organic cations for the 2D RP-perovskite, while divalent interlayer organic cations are suitable for the 2D DJ-perovskite. Unfortunately, the expanded interspaces filled with such bulky organic cations result in increased exciton binding energy and charge transfer barriers between the inorganic parts. The enlarged bandgap also brings the loss of optical absorption. Theoretically, it is hard to achieve a satisfying photovoltaic performance in the case of the normal 2D perovskite as a photoactive layer. Thus, partially replacing GA with a smaller molecule, forming the 2D ACI perovskite with a narrower band gap, has drawn considerable attention. 2D ACI follows the formula $(A')(A)_n B_n X_{3n+1}$, where the slightly larger A' cation (e.g., GA⁺) and smaller A cation (e.g., MA⁺) form an alternating arrangement in the interlayer space. So far, Cs, MA, FA, and otheres are typically regarded as the substitutable spacer to build 2D ACI perovskites together with GA cations. The crystallographic data of the ACI perovskites reported are presented in Table 3.

3.2.1 ACI crystal structure and physical property

 $(GA)(MA)_nPb_nI_{3n+1}$. In 2017, Soe et al. 56 firstly reported the ACI phase $(GA)(MA)_nPb_nI_{3n+1}$ series (n = 1, 2, and 3), in which GA and MA reside both inside the octahedral cage and between the layers, alternating with another cation (Fig. 6a). All 2D ACI perovskites demonstrated plate-like crystal morphology and crystallize in orthorhombic space groups, in which the ACI phase (n = 1 and 3) adopts the centrosymmetric *Imma* space group, while the ACI phase (n = 2) employs the polar noncentrosymmetric Bmm2 space group. The formation of the 2D ACI crystal structure is affected by the special molecular shapes of triangular GA and rod-like MA cations, leading to the "steric

hindrance". Thereby, the perovskite layers have to accommodate both cations between the two octahedral [PbI₆]⁴⁻ layers in a tight-packed arrangement. These 2D ACIs have a lower interlayer distance (minimum: ~ 3 Å), compared to that of 2D DJ $((3AMP)(MA)Pb_2I_7, 4 \text{ Å})$ and 2D RP $((BA)_2MAPb_2I_7, 7 \text{ Å})$ phase. The low strength of van der Waals forces could improve the stability of the 2D perovskite structures, while the intensive H bonds have a positive effect on ion fixation and thermal stability. The ACI exhibits a decreased bandgap as the n value increases; for example, $E_g = 2.27$ eV for n = 1, $E_g = 1.99$ eV for n = 12 and $E_g = 1.73$ eV for n = 3.

 $(GA)(Cs)_nPb_nX_{3n+1}$ (n = 1 and 2; X = I and Br). The GA/Cs-based ACI perovskites, including GACsPbI₄, GACsPbBr₄, and GACs₂ Pb₂Br₇, were reported by Nazarenko et al.⁵⁷ As for the GACsPbI₄ and GACsPbBr₄ crystal structure, a single PbX₆ octahedra layer and GA-contained interlayer were arranged alternatively in space, whereas GACs₂Pb₂Br₇ adopts a bilayer arrangement of the octahedra between each interlayer (Fig. 6b). Such GA/Csbased composition could stabilize the perovskite phase until 300 °C, delivering superior thermal stability. The GACs-based ACI perovskites have relatively wider bandgaps, for instance, 2.35 eV, 2.8 eV and 2.76 eV for GACsPbI4,GACsPbBr4, and GACs₂PbBr₇, respectively.

 $GAFAPbI_4$. So far, $GAFAPbI_4$ (n = 1) is the only reported GAFA-Based 2D ACI. Nazarenko et al. 34 found that it crystallized in the monoclinic space group and exhibited the stair-like morphology (Fig. 6c). The crystal structure of GAFAPbI₄ consists of corrugated Pb-I layers, and GA and FA cations are situated in the interlayer, where RP-like phases form. Similar to other ACI n = 1 phases,

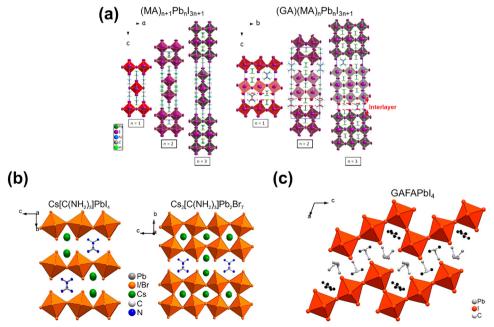


Fig. 6 Crystal structures of (a) $(MA)_{n+1}Pb_nI_{3n+1}$ and $(GA)(MA)_nPb_nI_{3n+1}$ (right), n=1, 2 and 3. Reproduced with permission. ⁵⁶ Copyright 2017, ACS Publishing Group. (b) $(GA)(Cs)_n Pb_n X_{3n+1}$ (n=1 and 2; X=I and Br) Reproduced with permission. ⁵⁷ Copyright 2017, ACS Publishing Group; (c) GAFAPbl₄. Reproduced with permission.³⁴ Copyright 2018, ACS Publishing Group.

GAFAPbI₄ has a wide bandgap of 2.44 eV. It is thermally stable up to 255 °C owing to the firm H bonds with the [PbI₆]⁴ framework.

In summary, the critical feather of 2D ACI is the alternatingcation interlayer configuration, which is distinguished from the DJ and RP phases, leading to chemical differences.

- (I) The octahedral arrangement in the structures. Due to the bulky or long-chain monovalent organic cation, the RP phase has half octahedral displacement between the adjacent 2D perovskite slabs. For DJ phase, divalent cations are vertically connected with 2D perovskite slabs. In contrast, an alternating cation arrangement is found in the ACI phase, which induces octahedral rotation in the crystal structure.
- (II) The interlayer distance (the distance between the terminal I ions and the connecting organic ammonium). The RP phase has the largest interlayer distance⁵⁸ because of the bilayer structure of the monovalent cation pairs. Compared with the RP phase, the DJ phase is shorter on account of the monolayer structure of its divalent organic cation.⁵⁹ Regarding the ACI phase, the small cations (e.g., MA and GA) form the tightly interacted interlayer, resulting in the lowest interlayer distance.⁵⁶
- (III) The interlayer interaction. Both the van der Waals gap and hydrogen bonding are present in the RP phase. Nevertheless, the strong interaction with adjacent 2D perovskite slabs weakens the van der Waals gap. As in the ACI phase, the van der Waals gap and hydrogen bonding are retained, but the hydrogen bonding is the main interaction. Therefore, the elimination or reduction of the van der Waals gaps may increase the structural stability of 2D.⁶⁰
- **3.2.2. 2D ACI-based PSCs.** Although the closer $[Pb_nI_{3n+1}]$ parts bring a narrower bandgap of GA-based ACI perovskites than that of pure GA-based 2D perovskites, it is still far from a satisfying extent for device application. To further boost the efficiency and stability of ACI-PSCs, most attempts have been focused on regulating the crystallization process and the

number of n. A finite but higher n represents more inorganic layers conjugated together between each interlayer, and their optical properties gradually become close to that of 3D ones. Up to now, only the cases of $n \ge 3$ combined with pristine I⁻ at the X-site have been reported. 26,61-63 Additive and solvent engineering are the key strategies to construct the desired 2D ACI perovskite film, whose device performances are shown in Table 4.

 $(GA)(MA)_3Pb_3I_{10}$ (n = 3). The $(GA)(MA)_3Pb_3I_{10}$ (n = 3) has a relatively narrow bandgap of 1.73 eV, which preserves both the excellent carrier transport properties of 3D perovskites and the environmental stability of 2D systems. GAMA₃Pb₃I₁₀ was first prepared by Soe et al. 56 and a PCE of only 7.26% was obtained. The unsatisfying efficiency is mainly ascribed to the phase distribution and poor crystallization of the 2D ACI perovskite film.

Zhang et al. 65 optimized the crystallization process of the 2D ACI perovskite by solvent engineering-adopting a mixed solvent of dimethylformamide (DMF) and dimethyl sulfoxide (DMSO) at the volume ratio of 1:10. With the assistance of in situ grazingincidence X-ray scattering (GIWAXS) characterization, it was found that 2D GA₂PbI₄ was the crucial intermediate phase, which provided the scaffold for the ACI perovskite growth, and dominated the distribution of $GAMA_3Pb_3I_{10}$ (n = 3) with less of other n-value-phases and 1D GAPbI3 impurities. Additionally, the hot-casting method was used instead of traditional anti-solvent processing to effectively reduce the defect density. Consequently, the high-quality ACI perovskite film was deposited, yielding a greatly improved PCE of 14.7%.

Additive engineering also helps improve the crystallinity of the 2D GAMA₃PbI₁₀ (n = 3) perovskite films. Zhang et al.⁶⁶ introduced MASCN (0.3 molar ratio relative to PbI₂) in the perovskite precursor. Compared to the pristine 2D ACI perovskite film, the grain size in the MASCN-doped 2D ACI films was clearly enlarged,

Table 4 Summary of ACI perovskites and the corresponding device performance

Device structure	ACI formula	Strategy	PCE (%)	Stability	Ref.
ITO/PEDOT: PSS/PVSK/ PC ₆₁ BM/Al	(GA)MA ₃ Pb ₃ I ₁₀	None	7.26	None	56
ITO/PEDOT:PSS/PVSK/ PC ₆₁ BM/BCP/Ag	$(GA)MA_3Pb_3I_{10}$	Solvent engineering: DMSO	14.7	T_{88} = 240 d in the air with 30–40% RH at 25 $^{\circ}\mathrm{C}$	65
FTO/TiO ₂ /PVSK/spir- oOMeTAD/Au	$(GA)MA_3Pb_3I_{10}$	Additive engineering: MASCN	10.12	T_{80} = 30 d in air with 20% RH at 30 $^{\circ}\text{C};T_{50}$ = 3200 min in N_{2}	66
ITO/PEDOT:PSS/PVSK/PCBM/ LiF/Al	$(GA)MA_3Pb_3I_{10}$	None	16.65	$T_{80} = 2400 \text{ h in N}_2$	67
FTO/TiO ₂ /PVSK/spir- oOMeTAD/Au	(GA)MA ₃ Pb ₃ I ₁₀	Additive engineering: MACl	18.48	T_{95} = 131 d in the air with 30–40% RH at 25 °C; T_{79} = 65 h at 80 °C; T_{57} = 200 h at MPP in the air with 55% RH at 60 °C	64
ITO/PEDOT:PSS/PVSK/C ₆₀ / BCP/Ag	$(GA)MA_4Pb_4I_{13}$	Solvent engineering: DMSO	12.8	None	62
ITO/PEDOT:PSS/PVSK/ PC ₆₁ BM/BCP/Ag	$(GA)MA_4Pb_4I_{13}$	Post-treatment: GASCN and MACl	15.27	T_{75} = 480 h in air with 50% RH at 25 °C; T_{70} = 300 h in air with 50 \pm 5% RH at 25 °C	63
FTO/TiO ₂ /PVSK/spir- oOMeTAD/Au	$(GA)MA_5Pb_5I_{16}$	Solvent engineering: DMSO	19.18	T_{95} = \sim 123 d in the air with 30 \pm 10% RH at 25 \pm 5% $^{\circ}$ C; T_{80} = \sim 60 h in an inert atmosphere at 80 $^{\circ}$ C; T_{80} = \sim 240 h at 60%–70 RH	
FTO/TiO ₂ /PVSK/spir- oOMeTAD/Au	$(GA)MA_5Pb_5I_{16}$	Post-treatment: amphoteric imidazolium iodide	22.26	T_{93} = \sim 1200 h in air with 25% RH at 25 °C; T_{55} = \sim 60 h in an inert atmosphere at 85 °C	26

PVSK: perovskite. RH: relative humidity. RT: room temperature. MPP: maximum power point.

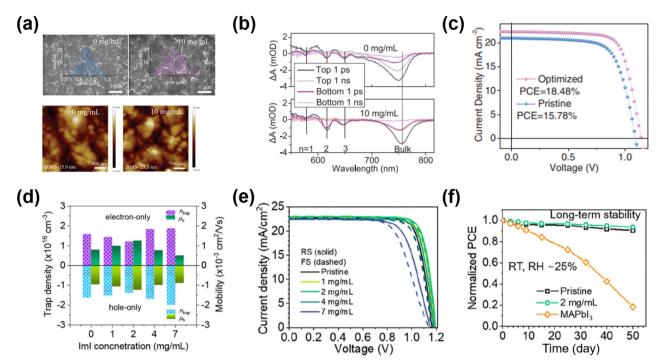


Fig. 7 The morphology of the 2D ACI film: (a) the top-view SEM images of the ACI films without/with MACI treatment (top); atomic force microscopy (AFM) images of the ACI films without/with MACI treatment. (b) Transient absorption (TA) spectra at different delay times for the ACI films without/with MACI treatment. (c) J-V curves under the reverse voltage scan of the best device ACI films without/with MACI treatment. Reproduced with permission.⁶⁴ Copyright 2019, Wiley-VCH. (d) Trap density (n_{trap}) and mobility (μ) of the GA(MA) $_n$ Pb $_n$ I $_{3n+1}$ (n=5) perovskite films with ImI treatment. (e) J-V curves under reverse- and forward-scan of PSCs with ImI treatment. (f) Long-term stability of PSCs without/with 2 mg mL⁻¹ ImI treatment at RT and 25% RH for 50 days. Reproduced with permission.²⁶ Copyright 2022, ACS Publishing Group.

which was accompanied by high crystallinity. Eventually, the PCE of the device with the MASCN additive was boosted from 9.05% (non-doped) to 10.12%. Suitable additives also controlled the distribution of the targeted ACI quantum wells (QWs, i.e., the n number).⁶⁷ Luo et al.⁶⁴ effectively promoted the charge transfer of 2D ACI (n = 3) by adding MACl in the precursor solution. The optimized thickness distribution of the QWs decreased the carrier-populating time to 0.8 ps, which enabled highly efficient transport/extraction in PSCs (Fig. 7a and b). Meanwhile, the morphology of the 2D ACI became smoother with the increased grain size. After optimization, the MACl-treated device delivered a record PCE of 18.48% (Fig. 7c).

 $(GA)(MA)_n Pb_n I_{3n+1}$ $(n \ge 4)$. As the *n* increases, the bandgap of 2D ACI approaches that of the 3D perovskites. Utilizing DMSO as a co-solvent to suppress lower n-phases could also be realized for n = 4 phase purification, according to the report by Ma et al.62 The DMSO in the precursor solution could obstruct the formation of the low-n (e.g., n = 1) phases in ACI perovskite films, resulting in reducing the hole-trap density and improving the charge transport. To give insight into the solvent influence on the crystallization of quasi-2D ACI, Yang et al.⁶¹ introduced a mixed solvent of DMSO/DMF (v:v = 1:10) to increase the $GAMA_5Pb_5I_{16}$ (n = 5) ACI formation. The stable intermediate phases can slow down the interposition of ions, which in turn retarded the crystallization of the ACI phases. Herein, the DMSO co-solvent led to a new intermediate solvate

phase of MA₂Pb₃I₈·2DMSO, whose solvation was immediately deprived along with the transition into ACI phases. The following thermal treatment completed the decomposition of intermediates and ACI crystallization with good film uniformity and surface coverage, leading to reduced charge recombination loss. Eventually, the unencapsulated PSCs realized a PCE of 19.8%.

Besides solvent engineering, post-treatment benefits the quasi-2D ACI phase arrangement. Zhang et al.63 proposed a sequential post-treatment method of modifying the GAMA4-Pb₄I₁₃ ACI film with GASCN and MACl. GASCN could improve the quality of the perovskite films through the coordination bonding between SCN⁻ and Pb²⁺, resulting in dense and well-aligned crystal grains. The sequential MACl treatment can suppress the charge defects on the perovskite surface and grain boundaries, which increased the electron/hole mobility. The PSCs based on sequential post-treatment demonstrated the best PCE of 15.27%, which was much higher than that of the pristine device (8.26%). Park et al.²⁶ employed a post-treatment reagent of amphoteric imidazolium iodide (ImI) on the top of the ACI n = 5 perovskite film. ImI posttreatment effectively reduced the vacancies/excess defects of the perovskite (Fig. 7d). Consequently, lower trap density and higher charge mobility were achieved, yielding a record-high PCE of 22.26% among the ACI-based 2D PSCs (Fig. 7e and f).

No research studies have been focused on GA-based ACI PSCs with further increased n, perhaps because of their negligible structural and optical differences from the GA-doped 3D species. Despite additive and solvent engineering optimization, the

parasitic 1D GAPbI₃ or 2D GA₂PbI₄ within the 3D perovskite matrix during ACI crystallization could not be eliminated. Their insufficiency in light capture, the role of the carrier trap center, as well as the resulting nonuniformity in the film morphologies are the critical barrier to achieving highly efficient and stable ACI-based devices. More technologies are needed to obtain the highly pure 2D ACI phases.

4. Additive engineering

Additive engineering is widely utilized to improve photovoltaic performance and long-term stability performances. The effects of prohibiting uncoordinated defects and ion migration by Ncontained lone-pair groups and the abundant hydrogen bonds from GA-based additives resemble that of GA-based dopants in Asite compositional engineering. Additionally, the ions could be inserted into the perovskite lattice and distort the perovskite structure in some cases, ameliorating structural and thermal stability. 68,69 Other distinctive functions are presented in the promotion of uniform and compact perovskite films by modulating the crystallization process such as nucleation and growth, and bandgap alignment through self-doping effect. 70,71 In most cases, the hydrophobic nature of the molecules also improves the ambient stability of the devices. GABr and GASCN are typically used as additives.⁷²⁻⁷⁴ The corresponding data are summarized in Table 5.

4.1 GABr additive

The multi-functional role of GABr was exemplified by Xu et al.⁷⁵ Due to the ionized ammonium group (-NH₃⁺) and unsaturated N atoms (-NH₂ or ==NH) in the GA⁺ cation, these two differently charged groups could be used to eliminate both positively charged (uncoordinated Pb2+) and negatively charged (I vacancy) defects to

reduce nonradiative recombination (Fig. 8a). More noticeably, the grain size in the perovskite film with the GABr additive was increased from 600 nm to about 1 µm (Fig. 8b). The high-quality crystallization contributed to the reduced surface charge defects, and the mechanisms of morphology modulation were attributed to the incorporation of bulky GA⁺ into the perovskite lattice. Last, the band alignment between the perovskite and HTL became better. In contrast to the pristine device, the 15 wt.% GABr-doped devices showed a higher built-in potential (V_{bi}) , which generated a strong driving force for intrinsic charge separation. These multiple factors together corresponded to the minimized $V_{\rm OC}$ loss. The FAPbBr₃based wide band-gap device ($E_{\rm g}\sim 2.3$ eV) with GABr reached the PCE of 8.92% with a high V_{OC} of 1.639 V.

Despite the great promise of the Sn-Pb binary perovskite for developing low band-gap PSCs ($E_g = 1.2-1.3$ eV), it suffers from the enormous challenge of Sn²⁺ oxidation, leaving Sn vacancies. The defect elimination caused by Sn2+ oxidation could also be achieved by GABr addition, as reported by Zhou et al. 76 In 2D GIXRD, the strongly dispersed secondary blots and bright rings in the perovskite film with 12% GABr additive demonstrated that the crystalline realms retained good orientation (Fig. 8d). Apart from the improved crystal quality and purity, GA+ was beneficial to the structural stability and restriction of the ion migration. Moreover, Br anions in the additive could act as a substituent on the X site to increase the tolerance factor and bandgap of the perovskites compared to those of the iodine-based ones. 77,85 As a result, the device with 12% GABr obtained a PCE of 20.63% with negligible J-V hysteresis and superior environment and thermal stability.

4.2 GASCN additive

GASCN has been proven as another effective additive in PSCs, in which GA⁺ reduces defect states at the surface and boundary

Table 5 Summary of PSCs with GA-based additives							
Additive	Perovskite component	Device configuration	PCE (%)	Stability	Ref.		
GABr	$FA_xMAyCs_zPbI_{3-y}Br_y$	FTO/SnO ₂ /PVSK/spiro-OMeTAD/Au	20.29	$T_{90} > 60$ d in the air at 25 °C with 25% RH; $T_{70} > 60$ d in the air at 80 °C with 40% RH; $T_{75} > 5$ h in the air under 1 sun illumination			
	FAPbBr ₃	ITO/SnO ₂ /PVSK/spiro-OMeTAD/Au	8.92	$T_{90} > 1024$ h in the air at RT with 50% RH	75		
	$FA_{0.7}MA_{0.3}Pb_{0.7}Sn_{0.3}I_3$	$\label{eq:total} ITO/EMICPEDOT:PSS/PVSK/S-acetylthiocholine chloride/C_{60}/BCP/Ag$	20.63	$T_{85}>1000~\rm h$ in the air at RT with 50–60% RH; $T_{90}>24~\rm h$ in the air at 80 $^{\circ}\rm C$ with 50–60% RH	76		
	$GA_{0.05}Cs_{0.15}FA_{0.8}$ Pb $(I_{0.85}Br_{0.15})_3$	FTO/SnO ₂ /PVSK/spiro-MeOTAD/Au	21.33	$T_{80} > 8$ h in N ₂ at MPP under 1 sun illumination	77		
GASCN	$(FASnI_3)_{0.6}(MAPbI_3)_{0.4}$	ITO/PEDOT:PSS/PVSK/C ₆₀ /BCP/Ag	20.5	$T_{80} = 220 \text{ h in N}_2 \text{ at MPP}$	78		
	(FASnI ₃) _{0.6} (MAPbI ₃) _{0.4}	ITO/PEDOT:PSS/PVSK/C ₆₀ /BCP/Ag	22.1 Tandem: 25.4	0.8 light soaking; $T_{90} = 500 \text{ h in air}$	79		
	$PEA_{0.1}FA_{0.9}SnI_3$	ITO/PEDOT:PSS/PVSK/PC ₆₁ BM/poly-ethylenimine/Ag	10.06	T_{91} = 720 h in N_2 at 25 °C; T_{45} = 15 h in the air at 25 °C with 25% RH; T_{55} = 2 h in air at 25 °C with 60% RH			
	FASnI ₃	ITO/NiOx/PVSK/ICBA/BCP/Ag	13.79	$T_{90} = 1200 \text{ h in N}_2$	81		
GACl	$CsMAFA/Pb/(I_{0.87}Br_{0.13})_3$	FTO/TiO ₂ /PVSK/spiro-OMeTAD/Au	17.9	None	82		
	MAPbI ₃	FTO/TiO ₂ /ZrO ₂ /PVSK/carbon layer	14.35	None	73		
GABF_4	$MAPbI_3$	ITO/NiOx/PVSK/PC ₆₁ BM/Ag	20.62	T_{81} = 45 d in the air at RT with 50–60% RH	83		
$GuaSO_3NH_2$	$Cs_{0.05}(MA_{0.12}FA_{0.88})_{0.95}$ Pb $(I_{0.88}Br_{0.12})_3$	FTO/TiO ₂ /PVSK/spiro-MeOTAD/Au	21.66	T_{90} = 750 h in the air at RT with 25% RH	84		
$GAPbI_2Br$	CsPbI ₂ Br	ITO/ZnO/PVSK/spiro-MeOTAD/Au	14.34	$T_{94} = 1000 \text{ h in N}_2$	85		

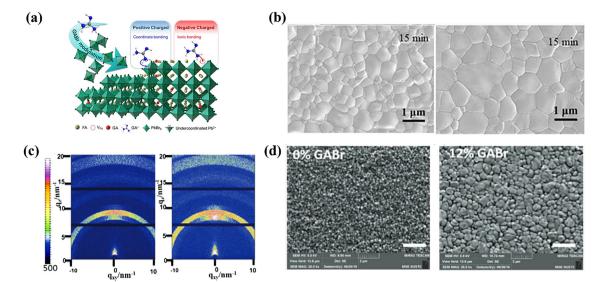


Fig. 8 (a) The schematic mechanism of GABr modulation on FAPbBr₃-based PSCs. (b) Top-view SEM images of the control (left) and 15% GABr film (right) annealed at 110 °C for 15 min. Reproduced with permission. 75 Copyright 2022, Elsevier. (c) 2D grazing-incidence X-ray diffraction (GIXRD) patterns of the perovskite films without (left) and with 12% GABr (right), respectively. (d) The SEM top images of the perovskite films without and with 12% GABr, respectively (the scale bar is 2 μm). Reproduced with permission.⁷⁶ Copyright 2020, Elsevier.

of the perovskite film due to the formation of hydrogen bonds. Meanwhile, SCN ions could regulate perovskite grain growth and again heal the defects in the grains of the perovskite film, owing to the Lewis-base nature of the S and N atoms in SCN. 86,87 Furthermore, SCN quickly is released from the perovskite films during the annealing process in the form of HSCN gas. Therefore, little residual GASCN remains.81,88 The

oxidation of Sn²⁺ could also be suppressed by GASCN.⁷⁸ In the charge-carrier dynamics, the (FASnI₃)_{0.6}(MAPbI₃)_{0.4} perovskite film modified with 7% GASCN exhibited drastically prolonged PL lifetimes ($\sim 1 \mu s$), which was comparable to that of pure Pbbased perovskite thin films due to the noticeable low defect density (Fig. 9a). Furthermore, the device with GASCN modification achieved a PCE of over 20% in single-junction cells and

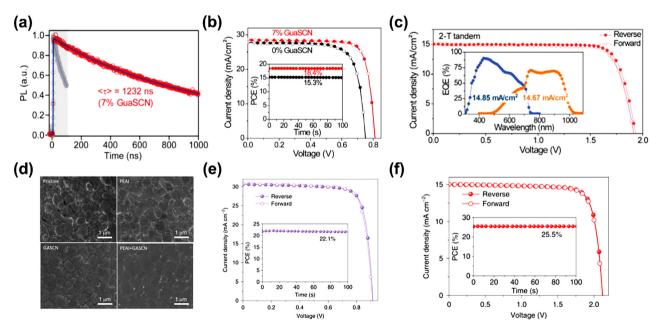


Fig. 9 (a) Time-resolved photoluminescence of the perovskite film with GASCN. J-V curves of (b) the champion single PSC; the inset shows the SPO efficiency near the maximum power point (MPP), and (c) the 2-T tandem cell. The inset shows the EQE spectra of the top and bottom subcells, respectively. Reproduced with permission.⁷⁸ Copyright 2019, AAAS. (d) Top-view SEM photograph of the perovskite film with PEAI and GASCN additives. J-V curves of (e) the single champion device with PEAI and GASCN additives along with the SPO efficiency, and (f) all-perovskite tandem PSCs along with the SPO. Reproduced with permission.⁷⁹ Copyright 2022, Nature Publishing Group.

25% in the all-perovskite tandem device, respectively, which benefited from the increase in carrier lifetime (Fig. 9b and c). Similar results were also reported by some other groups to stabilize pure Sn-based perovskites.80

Recently, PEASCN and GASCN additives have been employed in the Sn-Pb-based perovskite precursor.⁷⁹ The binary cation additives helped form the quasi-2D (PEA)2GAPb2I7 deposited at the surfaces and grain boundaries to suppress more charge traps than that caused by Sn2+ oxidation (Fig. 9d). The 2D additive engineering endowed the Sn-Pb-based single-junction cells and two-terminal tandems with efficiencies of 22.1% and 25.5%, respectively (Fig. 9e and f).

4.3 Other GA-based additives

Anions such as Cl⁻, tetrafluoroborate (BF₄⁻), and sulfamate (H₂N-SO₃⁻) in GA-based additives play a crucial and synergistic role, similar to SCN-. These anions could regulate the grainsize growth of perovskite films and passivate defects such as undercoordinated Pb2+ and Pb0, thus affecting the charge transfer behaviors and photovoltaic properties.

Given that the O and N atoms of H₂N-SO₃(SM) could bond with uncoordinated Pb2+ because of the lone electron pairs in O and N, Liu et al.84 introduced GASM into the [Cs_{0.05}(MA_{0.12}- $FA_{0.88})_{0.95}Pb(I_{0.88}Br_{0.12})_3$ perovskite precursor to passivate charge traps at the surface and grain boundary effectively. With the dual-function synergy effect of GA⁺ and SM, the grain size of the perovskite film was enhanced, and the PL intensity of perovskite film became stronger. The optimal device yielded the best PCE of 21.66%, and kept 90% of its original efficiency after 800 h storage in ambient conditions.

Similar to GA-based dopants in A-site composition engineering, the content of GA-based additives is usually less than 20% (molar ratio vs. Pb species). A high concentration of GA-based

additives would cause the formation of the impure phases, and the disruption of high-quality crystallization. Moreover, the anions could influence the GA incorporation in the perovskite lattice, which expands the lattice constant. A detailed mechanism of the synergistic effect of the GA⁺ cation and anion on the perovskite is still unclear, and requires further research.

Interface engineering

The interfaces of the ETL/perovskite and perovskite/HTL determine the efficiency of the charge transport, perovskite crystallization, and energy level adjustment. 89,90 Such interface properties could be engineered by the GA-contained interface introduction, which functions as a conventional passivator or 2D perovskite passivation layer. 91-93 Performances of PSCs with GA-based interface engineering are presented in Table 6.

5.1 GA-cation passivation layer

GA-based salts are considered suitable alternatives for interface engineering because the GA cation could bond with the uncoordinated halogen by hydrogen bonds and uncoordinated Pb²⁺ or Pb clusters by lone pair electrons provided by N atoms, resulting in reduced trap sites. 94,96 The added ion doping into the perovskite lattice was also observed in some cases, whose effects have attracted more comprehensive research. GA-based salts that have been reported as passivation agents so far are GA-halide salts (e.g., GAI, GABr, and GACl) and GASCN.

Luo et al.94 spin-coated the multi-functional GABr on the surface of (FA_{0.95}PbI_{2.95})_{0.85}(MAPbBr₃)_{0.15} perovskite film in the inverted PSCs. With the GABr passivation, the bright spots on the surface and boundaries disappeared, indicating that the PbX₂ impurity was removed (Fig. 10a and b). Moreover, the GABr additive made the Fermi levels of the perovskite films

Table 6 Representative PSCs with GA-based interface engineering

Interface engineering layer	3D perovskite layer	Device structure	PCE (%)	Stability	Ref.
- Idyer	3D perovskite layer	Bevice structure	T CE (70)	Beability	101.
GACl	$MAPbI_3$	ITO/PTAA/PVSK/PC61BM/LiF/Al	18.65	None	96
GACl	$MAPbI_{2.7}Cl_{0.3}$	FTO/TiO ₂ /PVSK/spiroOMeTAD/Ag	18.46	$T_{95} > 22 \text{ d in N}_2$	91
GABr	CsPbI ₃	FTO/TiO ₂ /PVSK/PTAA/Au	18.02	$T_{92} > 30 \text{ d}$ at 20–30% RH	107
GABr	$FA_{0.8}Cs_{0.2}Pb(I_{0.7}Br_{0.3})_3$	ITO/PTAA/PVSK/C ₆₀ /BCP/Ag	18.19	$T_{72} > 70$ h at RT and $> 50\%$ RH in air	74
GABr	$(FA_{0.95}PbI_{2.95})_{0.85}(MAPbBr_3)_{0.15}$	ITO/PTAA/PVSK/PC ₆₁ BM/C ₆₀ /BCP/Cu	21.51	$T_{95} = 500 \text{ h at } 85 ^{\circ}\text{C in N}_2$	94
GAI	MAPbI ₃	ITO/SnO ₂ /PVSK/spiroOMeTAD/Au	18.54	T_{85} = 30 d in dry condition; T_{50} = 15 d in 60 ± 10% RH.	52
GAI	FAPbI ₃	FTO/TiO ₂ /PVSK/spiroOMeTAD/MgF ₂ /Au	24.36	None	95
GASCN	$MAPbI_3$	FTO/TiO ₂ /PVSK/spiroOMeTAD/Au	21.26	T_{80} = 1000 h in the air at 20% RH	97
1D GAPbI ₃	$(FA_{0.9}Cs_{0.07}MA_{0.03}Pb(I_{0.92}Br_{0.08})_3$	FTO/TiO ₂ /PVSK/spiroOMeTAD/Au	21	T_{80} = 550 h in N ₂ at MPP under RT	
2D GA ₂ PbI ₄	CsPbI ₂ Br	FTO/SnO ₂ /PVSK/spiroOMeTAD/Au	15.6	T_{88} = 60 d in the air at 25 °C with 25% RH; T_{76} = 30 d in the air at 80 °C with 25% RH; T_{90} = 5 h in the air at MPP	103
2D GA ₂ PbI ₄	n-i-p: Cs _{0.05} FA _{0.85} MA _{0.1} PbI ₃ p-i-n: Cs _{0.3} FA _{0.6} DMA _{0.1} PbI _{2.4} Br _{0.6}	n-i-p: ITO/SnO ₂ /PVKS/spiroOMeTAD/Au p-i-n: ITO/NiOx/PVKS/PC ₆₁ BM/BCP/Ag	n-i-p: 20.1 p-i-n: 22.9	$T_{90} > 30 \text{ d in N}_2$	99
2D GA ₂ PbI ₄		FTO/SnO ₂ /PVKS/spiroOMeTAD/Au	22.53	$T_{90}>30$ d in the air with 25% RH; $T_{80}>30$ d in the air with 75 \pm 5% RH; $T_{80}>30$ d in N ₂ under light illumination	98

PVSK: perovskite; RH: relative humidity; T_x is when PCE degrades to x% of the initial value.

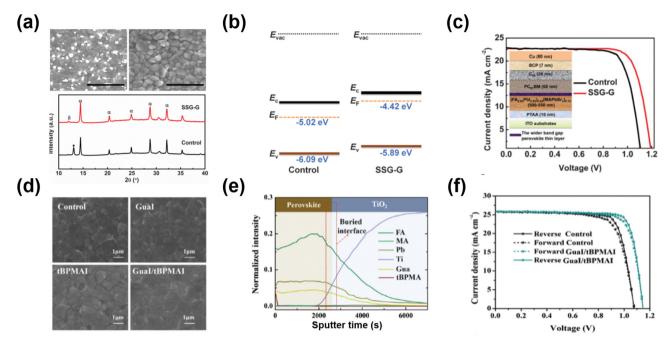


Fig. 10 (a) Top view SEM images of the perovskite film with/without GABr passivation (upper panel). XRD patterns (lower panel) of the perovskite film with/without GABr treatment (α: black perovskite phase; β: non-perovskite phase). (b) Schematic energy-level diagrams of the control and GABr-treated films. (c) J-V curves under the reverse scan of devices with/without GABr passivation. Reproduced with permission. 94 Copyright 2018, AAAS. (d) Top view SEM images of the perovskite film with/without binary GAI/tBPMAI treatment. (e) The time-of-flight secondary-ion mass spectrometry (ToF-SIMS) of the binary GAI/tBPMAI-treated perovskite film. (f) The forward/reverse scan J-V curves of the best devices with/without the binary GAI/tBPMAI treatment. Reproduced with permission.⁹⁵ Copyright 2021, Royal Society of Chemistry.

shift toward the vacuum (400 mV), which was higher than the valence band upshift (200 mV), indicating more n-type nature after GABr treatment (Fig. 10c). The more n-type perovskite film could result in a larger fraction of occupied versus vacant traps, and a reduction in the rate of trap-assisted recombination. Additionally, Br enrichment was observed near the interface after the addition, leading to a wider bandgap (80 meV) in this region. It is energetically favorable for one or both charge carriers to reside predominantly within the bulk of the film, away from the surface defects. The device with GABr passivation delivered a PCE of 21.51% with a record $V_{\rm OC}$ of 1.21 V, resulting from the suppression of trap-induced recombination

Sometimes, GA-based passivation agents were simultaneously post-treated along with other salts for more progress. Zhang et al. 95 addressed the complete passivation from the interfaces to bulk in the perovskite film by combining GAI with 4-tert-butylphenylmethylammonium iodide (tBPMAI) as the passivation agent. Given the difference between GAI and tBPMAI in ionic radius, GA could diffuse into the perovskite lattice. At the same time, the tBPMAI gathered on the surface of the perovskite, simultaneously modulating defects in perovskites from bulk to interfaces, which were confirmed by ToF-SIMS (Fig. 10f). More interestingly, it showed a positive effect on crystallization modulation after the post-treatment of this composite layer. The perovskite grains were regularly arranged, leading to the compact and smooth morphology of the perovskite (Fig. 10e). Finally, the device with individual GAI or tBPMAI treatment showed a low PCE of 21.79%. In contrast, the device with binary passivation obtained the champion PCE of 24.36% with a lower voltage loss of 0.388 V (Fig. 10g).

Better crystal growth with fewer trap states also can be regulated by the passivator itself and its solvent. Yang et al. 97 selected GASCN dissolved in chlorobenzene (CB) anti-solvent to passivate the MAPbI₃ surface effectively. The GA cation could stabilize the black phase, anchor the lattice ion, and inhibit their migration due to the hydrogen bond formation. Results from different anti-solvents such as ethyl acetate (EA), i-propanol (IPA), and cyclohexane (CH) were also discussed here. Compared with other anti-solvents, CB with low polarity could cause minor damage to the perovskite grain and slow down the crystallization process. As a synergistic result of GASCN and CB, the grain size was dramatically increased. This binary strategy increased the PCE from 20.14% to 21.26%, and the thermal and moisture stability were enhanced.

The functions of constructing a GA-based interface (defect passivation, energy level alignment, and crystallization regulation) are similar to that of GA-based additives. However, this approach induces two individual layers, different from the doping behavior of additives in the perovskite layer during additive engineering. So, the limits of the GA molar ratio (vs. overall A-site cation content) considering the possible interference on perovskite crystallization are not involved here. However, the limits lie in its low solubility in CB. Generally, IPA is chosen to dissolve the GA-based agents, but as mentioned before, IPA could erode the surface of the perovskite film owing to the relatively high polarity.

Searching for a low-polarized anti-solvent that is compatible with GA salts or exploring soluble GA salts in CB should be the future task for managing GA-contained interfaces.

5.2 2D perovskite passivation layer

Recently, the PCE of PSCs with 2D/3D structure has surprisingly achieved 25%, comparable to that of crystalline silicon-based commercial PV products. The 2D perovskite passivation layer has been produced for various functions, from passivating surface defects of 3D perovskite and energy level alignment to improving resistance to environmental stress. 100-102 Given the high hydrophobicity of 2D $GA_{n+1}Pb_nI_{3n+1}$, devices comprising the 2D/3D form exhibit significantly improved environmental stability compared to devices containing only the 3D perovskite. 103,104 2D passivation layers could be prepared by solution processing, immersion processing, vacuum deposition, and mechanical press as standard, 105 but unconventionally, it was a new phase induced by the interaction between dopants and the surface perovskite layers.

Similar to the dual optimization scheme introduced previously,95 Zhang et al.98 treated the perovskite with GAI and GACl to realize surface and bulk modification. Theoretical calculations proposed that among GAX salts, the Cl doping system exhibited the highest formation energy of iodine vacancy and the longest excited state lifetime. Therefore, GACl was expected to be an appropriate passivator for bulk doping, and the outperformed PCE of GACl (20.07%) was higher than others

(17.71% for GABr and 18.01% for GAI). However, the smallest ionic radius of Cl⁻, showing the lowest activation energy barrier for X ion migration, infers the need to suppress its bulk diffusion to the surface. Co-doping with GAI was suggested to form a stable capping layer owing to its larger activation energy barrier (Fig. 11a and b). After co-doping, GAI was inserted into the 3D perovskite lattice forming the 2D GA₂PbI₄ layer for surface passivation (Fig. 11c). The incorporation of large GA cations also modulated the lattice tensile strain towards minimizing defect centers. Benefiting from the strategy for reducing nonradiative recombination centers from both bulk and interfacial regions and the energy level alignment after 2D layer insertion, the PCE of the device with a 2D/3D perovskite structure exceeded 22.53%. Moreover, the PCE maintained more than 90% after 30 days at 40% R.H. due to the enhanced hydrophobicity of the 2D layers.

The 2D perovskite layer not only passivates the 3D perovskite layer, but also shows the great potential of passivating buried side. Chen et al. 99 introduced the 2D GA₂PbI₄ passivation layer in the inverted PSCs between NiO_r HTL and the perovskite layer. The GA in 2D GA₂PbI₄ could connect with NiO_x by forming hydrogen bonds, suppressing the migration of oxygen vacancies on the NiOx surface. In addition, the symmetric structure of GA promoted its accumulation onto the NiOx surface to suppress non-radiative recombination (Fig. 11e). It was special when other cations such as n-butylammonium (BA), ethylammonium (EA), and dimethylammonium (DMA) were

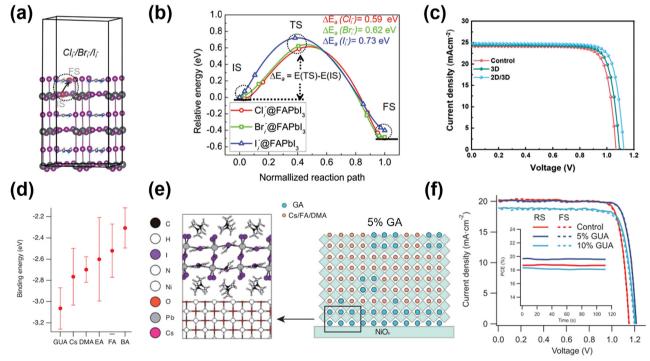


Fig. 11 (a) Illumination of the halogen ion interstitial ($Cl_i^-/Br_i^-/l_i^-$) migration from the perovskite bulk (defined as initial state, IS) to the surface (defined as the final state, FS), along with (d) the energy profile. (b) The activation energy barrier of the halogen ion. (c) J-V curves under the reverse scan of devices with/without 2D perovskite passivation. Reproduced with permission. 98 Copyright 2022, Wiley-VCH. (d) The average binding energy of different cations with NiO_x surface by DFT. (e) Structural model and corresponding schematics of the crystallization process for films with varying concentrations of GA on NiO_x . (f) The forward/reverse scan J-V curves of the champion devices with/without 2D perovskite passivation. Reproduced with permission. Copyright 2021, Wiley-VCH.

also added to the precursor. It was found that DMA and EA could not form the 2D perovskite. Furthermore, the 2D BAbased perovskite phase was not present in the NiOx/perovskite interface, but was located in bulk. The authors ascribed the unique phenomenon to the GA cation's higher binding energy and polarity action (Fig. 11d). Finally, the inverted device boosted the PCE up to 22.9% with an increase of 65 mV in $V_{\rm OC}$ (Fig. 11f).

It should not be ignored that in some cases, other lowdimensional phases would form when a GA-contained solution is spin-coated on top of a pre-formed 3D perovskite layer, which also shows a good passivation effect and interfacial band alignment. For instance, Alharbi et al. 106 utilized solid-state magic angle spinning (MAS) NMR to verify the coexistence of 1D GAPbI₃ and 2D GA₂PbI₄ on the surface of FA_{0.93}Cs_{0.07}PbI₃. Similarly, Chavan et al. 104 spin-coated GAI/IPA on the surface of a 3D perovskite film and detected the 1D δ-FAPbI₃-like phase. Empirically, the formation of the GA-based 2D perovskite passivation interface is probably accompanied by the complicated mixture of the original GA-based precursors, 3D perovskite constitutes, or other 2D species with different QWs. However, it does not mean that there are positive contributions from every impurity, or in many cases. Moreover, the inefficiency in charge transport across the organic interlayer and the light conversion due to the wider bandgap becomes intrinsic drawbacks of 2D materials. Delicate control for pristine or specific composites in 2D interlayers, as well as the design of innovative 2D materials with better intrinsic properties, should be valued in the future.

6. Summary and perspective

In summary, in light of the processing approaches for introducing the guanidinium cation, we systematically reviewed recent research and development on the application of GA in PSCs, including A-site substitution in either 3D perovskite or 2D ACI perovskite, additive, and interface engineering. The series of breakthroughs showed that GA cations exhibited unique potential in addressing the coexistence of long-term operational stability issues and effective photovoltaic performance in PSCs. Despite these merits, efforts toward some intractable challenges remain to be emphasized as follows:

- (1) Choosing the proper anion in GA salts for improving the performance of PSCs. Nowadays, GA-based salts have been widely used in the field of PSCs, and the anion is another factor for defect passivation, energy alignment, and structure stabilization. For example, Cl⁻ could tune the crystallization process, and Br could change the charge distribution and bandgap of the perovskite surface. SCN and BF₄ could effectively passivate defects such as undercoordinated Pb ions in the bulk and interface due to lone pair electrons. As a result, a suitable anion is a key aspect to eliminate charge defects.
- (2) Developing a more moisture-resistant GA-based perovskite. Due to the lack of long hydrophobic chains or ring structures, the GA-based perovskite film degrades faster than DJ or RP perovskites under moisture and thermal stresses. Therefore, it should be

taken into consideration to introduce the new GA-based derivatives for long-term operation stability, but the relevant papers are limited. 108,109 In addition, it is an alternative way to take the place of GA by long-chain cations as the interlayer cations phase to endow more hydrophobic properties, thereby increasing moisture repellence and stability for ACI perovskites. 110 Few reports are applying monovalent cations (e.g., Cs and FA) to replace MA. Regarding the effect of steric hindrance on the perovskite structure, the cations with appropriate ionic radius are expected to be better candidates to alternate with GA than MA.

- (3) Constructing the new 2D GA-based ACI capping layer on top of the 3D perovskite. Although GA2PbI4 was shown to passivate the surface defects of the 3D perovskite and prevent moisture from damaging the perovskite structure, its poor charge mobility inhibits efficient device operation. The 2D ACI layer with large n is expected to be a competitive alternative as the capping layer due to the high carrier mobility. It is noted that when the 2D ACI perovskite is deposited on the surface of 3D perovskite, the formation of multiple-perovskite phases would be induced, including $(GA)_x(MA)_{1-x}PbI_3$, GA_2PbI_4 , or other GA-based ACI with different quantum wells (n), which are possible trap sites on the surface. More precise control towards n should be valued in the future.
- (4) Further investigating the photophysical behavior in GA-based perovskites, especially ACI systems. The mechanisms of charge carrier dynamics, phase degradation, and ion migration in 2D ACI have not been systematically studied. A comprehensive understanding of these evolutions would provide critical guides for improving photovoltaic performances. Moreover, it would help ACI materials to become competitive candidates for application in other semiconducting fields, including photodetectors, light-emitting diodes, or field-effect transistors.
- (5) Application in other photoelectric devices. In addition to PSCs, the GA-based perovskite materials demonstrated excellent performance in the area of the optoelectrical device including light-emitting diodes (LEDs) and photodetectors. 111,112 Similar to solar cells, the GA could treat the surface of nanocrystals or colloidal quantum dots^{113,114} to passivate the defects and provides lattice-stabilizing effect via hydrogen bonds, reducing nonradiative recombination. Further research studies are still needed to define the "insight" mechanism.

Conflicts of interest

The authors declare no conflict of interest.

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