



Cite this: *Mater. Adv.*, 2023,  
4, 4528

# First-principles study on the optoelectronic and mechanical properties of all-inorganic lead-free fluoride perovskites $ABF_3$ (A = Na, K and B = Si, Ge)<sup>†</sup>

Chol-Jin Pak, Un-Gi Jong, \* Chung-Jin Kang, Yun-Sim Kim, Yun-Hyok Kye and Chol-Jun Yu \*

In spite of extensive studies on halide perovskites for advanced photovoltaic applications, little attention has been paid to fluoride perovskites so far. Here, we present a systematic investigation of not only the electronic and optical properties but also the mechanical and chemical stabilities of all-inorganic lead-free fluoride perovskites  $ABF_3$  (A = Na, K and B = Si, Ge) in the cubic phase using first-principles calculations. We provide a reliable prediction of their lattice constants, band gaps, effective masses and exciton binding energies, revealing variation tendencies according to the selection of A- and B-site cations. Between the two different B-site fluoride perovskites, the silicon-based compounds  $ASiF_3$  (A = Na, K) are highlighted as being more suitable for a light absorber of perovskite solar cells, due to their appropriate band gaps, remarkably low exciton binding energies, and ultrahigh spectroscopic limited maximum efficiency reaching  $\sim 34\%$  with a layer thickness over  $1\ \mu\text{m}$ . Through the calculations of elastic constants, phase decomposition energies and phonon dispersion curves, we further find that the all-inorganic lead-free fluoride perovskites  $ABF_3$  are mechanically stable but only  $KGeF_3$  is chemically stable against phase decomposition into its chemical reactants and is also thermodynamically stable. This work highlights that all-inorganic lead-free fluoride perovskites have high potentiality to be used as a promising light absorber in less toxic and high-performance perovskite solar cells.

Received 24th July 2023,  
Accepted 25th August 2023

DOI: 10.1039/d3ma00457k

rsc.li/materials-advances

## 1 Introduction

Over the past decade, metal halide perovskites have triggered an enormous research interest in various fields such as solar cells, photodetectors, X-ray detectors, lasers and light emitting diodes.<sup>1</sup> In particular, perovskite solar cells (PSCs) have emerged as the most promising alternative to the conventional silicon-based solar cells, due to their high power conversion efficiency (PCE) exceeding  $25.5\%$ <sup>2</sup> and low cost of their scalable fabrication.<sup>3</sup> However, there still exist some challenges that need to be addressed for widespread commercialization of PSCs, such as poor stability<sup>4–6</sup> and toxicity of perovskite thin films.<sup>7,8</sup> Here, the poor stability is attributed to the intrinsic volatility of organic components in organic–inorganic hybrid

halide perovskites such as  $\text{MAPbI}_3$  (MA =  $\text{CH}_3\text{NH}_2$ ), while the toxicity originates from lead (Pb).<sup>9–11</sup> To address such problems and find alternatives to  $\text{MAPbI}_3$ , extensive studies have been performed by replacing the organic component with inorganic monovalent cations like alkali cations and toxic lead with nontoxic elements such as Sn and Ge in the same group.<sup>12–18</sup>

In particular, all-inorganic halide perovskites have been proved to exhibit long-term stability of PSCs against dampness, ultraviolet light and heat, concurrently providing an efficient pathway for removing toxicity.<sup>19,20</sup> As a typical example, all-inorganic cesium lead iodide  $\text{CsPbI}_3$  has been found to significantly improve the stability of PSCs compared with the organic–inorganic hybrid counterpart  $\text{MAPbI}_3$ , while maintaining high PCE comparable to that of hybrid PSCs.<sup>21–25</sup> In 2015, a  $\text{CsPbI}_3$ -based PSC was fabricated for the first time with an efficiency of  $2.9\%$  by Eperon and co-workers,<sup>25</sup> and soon after, a certified high efficiency of over  $18\%$  and excellent operational stability were achieved by using surface passivation and quantum dot engineering.<sup>24</sup> Moreover, Chang *et al.*<sup>21</sup> recently reported high-performance printable  $\text{CsPbI}_3$  solar cells with an efficiency of  $19\%$  *via* a scalable manufacturing technique under ambient conditions. They synthesized high quality  $\text{CsPbI}_3$  films by using

Chair of Computational Materials Design (CMD), Faculty of Materials Science, Kim Il Sung University, PO Box 76, Pyongyang, Democratic People's Republic of Korea.  
E-mail: ug.jong@ryongnamsan.edu.kp, cj.yu@ryongnamsan.edu.kp

<sup>†</sup> Electronic supplementary information (ESI) available: Total energy convergence tests according to the kinetic energy cutoff and  $k$ -point mesh, comparison of the atomic forces, crystalline structures of cubic AF and orthorhombic  $\text{BF}_2$  and convergence of phonon dispersion curves according to the size of the supercell. See DOI: <https://doi.org/10.1039/d3ma00457k>



a combination of air-flow-assisted drying and the addition of organic Lewis acid  $\text{Zn}(\text{C}_6\text{F}_5)_2$ , demonstrating that the PSC degrades by only  $\sim 2\%$  after 700 h under air conditions without any encapsulation. However, perovskite  $\text{CsPbI}_3$  in the cubic phase has an intrinsic instability problem caused by phase transition, promoting the development of various methods such as nanostructuring,<sup>26–28</sup> ion substitution<sup>29</sup> and hybridization (e.g.,  $(\text{CsMA})\text{PbI}_3$ )<sup>12,30</sup>.

On the other hand, to mitigate the toxicity of lead halide perovskites, great research efforts have been devoted to replacing Pb with Sn or Ge, which has resulted in the development all-inorganic Sn- or Ge-based perovskites  $\text{CsSnI}_3$  or  $\text{CsGeI}_3$  and their solid solutions with tunable band gaps of around 1.3 eV, which are suitable for single-junction solar cells.<sup>31–34</sup> Promising progress has been reported with the solid-solution  $\text{CsSn}_{0.5}\text{Ge}_{0.5}\text{I}_3$  perovskite by Padture and co-workers.<sup>33</sup> They demonstrated that a native-oxide layer with full coverage can encapsulate and passivate the  $\text{CsSn}_{0.5}\text{Ge}_{0.5}\text{I}_3$  perovskite surface leading to a promising efficiency of up to 7.1% and high stability with less than 10% decay in efficiency after 500 h of continuous operation under ambient conditions. Recently, Lanzetta *et al.*<sup>4</sup> reported new mechanistic insights into the degradation of tin-based perovskites, proposing that dark hole withdrawal from the perovskite can delay the formation of  $\text{SnI}_4$  and its evolution to highly aggressive  $\text{I}_2$  and thus lead to improved stability of the tin perovskite film. In addition to the substitution of Pb with the same group elements of Sn and Ge, the replacement of Pb with monovalent and trivalent ions results in a double perovskite structure (e.g.,  $\text{Cs}_2\text{AgBiBr}_6$ )<sup>35,36</sup> which is a compositional variation within the perovskite lattice. So far, more than 350 all-inorganic lead-free double perovskites have been synthesized with good phase stabilities and a wide range of tunable optoelectronic properties, but they exhibited relatively lower PCEs below 5%.<sup>18</sup>

Despite extensive research on all-inorganic lead-free halide perovskites, little attention has been paid to fluoride perovskites. Berastegui *et al.*<sup>37</sup> reported the relatively low temperature of cubic phase transition as 187 K for  $\text{CsPbF}_3$ . Li *et al.*<sup>38</sup> found that when X decreases from I to F in  $\text{ABX}_3$ , the temperature of cubic phase transition decreases from 634 K to 187 K, indicating that the perovskites with lower halogen element need less thermal energy to lift the symmetry to the cubic phase. Using the first-principles calculation, Yang *et al.*<sup>39</sup> confirmed the experimental finding that cubic  $\text{CSBX}_3$  perovskites (B = Sn, Pb) show improved stability with decreasing X. For all-inorganic fluoride perovskites, challenges may arise from the synthesis process and especially very large band gaps of over 4 eV,<sup>40</sup> hindering their use as solar absorbers. In this work, we propose another lead-free all-inorganic fluoride perovskite  $\text{ABF}_3$  (A = Na, K and B = Si, Ge) in order to address the band gap issue. Houari *et al.*<sup>41</sup> reported the theoretical work for the structural and optoelectronic properties of  $\text{KGeF}_3$ , but not for electronic and dynamical properties. Here, we report a systematic investigation of the structural, electronic, and optical properties and mechanical and thermodynamic stabilities of fluoride perovskites  $\text{ABF}_3$  using first-principles calculations.

## 2 Computational methods

All the DFT calculations in this work were carried out by applying the pseudopotential plane wave method as implemented in the Vienna *ab initio* simulation package (VASP).<sup>42,43</sup> For a description of the Coulomb interaction between the ionic cores and the valence electrons, we employed the projector augmented wave (PAW) potentials<sup>44,45</sup> provided in the package. The valence electronic configurations of atoms are  $\text{F-}2s^22p^5$ ,  $\text{K-}3p^64s^1$ ,  $\text{Na-}2p^63s^1$ ,  $\text{Ge-}4s^24p^2$  and  $\text{Si-}3s^23p^2$ . The Perdew–Burke–Ernzerhof (PBE) functional<sup>46</sup> within the generalized gradient approximation (GGA) was used to describe the exchange–correlation (XC) interaction among the valence electrons. For the cubic phase (see Fig. 1), structural optimizations and electronic structure calculations were performed with a plane-wave cutoff energy of 800 eV and a  $\Gamma$ -centered  $k$ -point mesh of  $8 \times 8 \times 8$ . These computational parameters provided an accuracy of total energy as 1 meV per formula unit (see Fig. S1, ESI<sup>†</sup>). We carried out variable cell structural optimizations until the atomic forces and pressures became less than  $10^{-4}$  eV  $\text{\AA}^{-1}$  and 0.003 GPa, respectively. The total energy convergence threshold was set to  $10^{-8}$  eV.

For the calculations of electronic density of states (DOS), a denser  $k$ -point mesh of  $12 \times 12 \times 12$  was adopted by imposing the partial occupancy on each orbital using the tetrahedra approach with Blöchl correction. To obtain more reliable band gaps, we also performed electronic structure calculations with the Heyd–Scuseria–Ernzerhof (HSE06) hybrid XC functional.<sup>47</sup> Here, 20% of the PBE functional was replaced with the exact Hartree–Fock exchange functional, which has been proven to yield band gaps in good agreement with the experimental values for inorganic halide perovskites.<sup>48,49</sup> The spin–orbit coupling (SOC) effect was taken into account only for electronic



Fig. 1 Polyhedral view for the crystalline structure of all-inorganic lead-free fluoride perovskite  $\text{ABF}_3$  (A = Na, K and B = Si, Ge) with a space group  $Pm\bar{3}m$ . The purple, blue and gray balls represent the A, B and F atoms, respectively.



structure calculations, resulting in negligible contribution to the band gap as the relativistic effect is very small for light atoms.

We applied the density functional perturbation theory (DFPT) method<sup>50</sup> and solved the Bethe-Salpeter equation within the GW method (BSE/GW)<sup>51–56</sup> to calculate the macroscopic frequency-dependent dielectric functions,  $\varepsilon(\omega) = \varepsilon_1(\omega) + i\varepsilon_2(\omega)$ , and estimated the light-absorption coefficients  $\alpha(\omega)$  as follows:<sup>13,57</sup>

$$\alpha(\omega) = \frac{2\omega}{c} \sqrt{\frac{\sqrt{\varepsilon_1^2(\omega) + \varepsilon_2^2(\omega)} - \varepsilon_1(\omega)}{2}} \quad (1)$$

where  $c$  is the light velocity in vacuum and  $\omega$  is the frequency of light waves. Within the single band approximation, the effective masses of electrons ( $m_e^*$ ) and holes ( $m_h^*$ ) were calculated by numerically processing the refined energy band structure around the  $R$  point of the Brillouin zone, at which both the valence band maximum (VBM) and the conduction band minimum (CBM) were found. Then, we estimated the exciton binding energy ( $E_b$ ) by using the following formula:<sup>13,57</sup>

$$E_b = 13.56 \frac{m_r^*}{m_e} \frac{1}{\varepsilon_s^2} \text{ (eV)} \quad (2)$$

where  $m_r^*$  is the reduced effective mass ( $1/m_r^* = 1/m_e^* + 1/m_h^*$ ) and  $\varepsilon_s$  is the static dielectric constant.

We calculated the spectroscopic limited maximum efficiency (SLME),<sup>58,59</sup> which is regarded as a good metric for determining the maximum efficiency that an absorber material can reach in a single-junction solar cell. Because it takes into account the nature of the band gap, the shape of absorption spectra, and the material-dependent non-radiative recombination losses,<sup>59</sup> SLME is an improved version of the Shockley and Queisser (SQ) efficiency.<sup>60</sup> Usually, the SLME is calculated as the ratio between the maximum power density and the incident power density, exceeding the Shockley–Queisser limit within the detailed balance framework. The standard solar spectrum, the energy band gap and the light-absorption coefficient were used as inputs for SLME calculation. In calculations, we employed the SL3ME package<sup>59</sup> for implementing SLME analysis of solar absorbers in Python3.

The mechanical stability of a polycrystalline solid can be estimated using elastic constants such as bulk ( $B$ ), shear ( $G$ ) and Young's ( $E$ ) moduli, which are determined from the elastic stiffness ( $C_{ij}$ ) and compliance ( $S_{ij}$ ) constants for a single crystal.<sup>61</sup> For a cubic structure, there exist three independent elastic constants ( $ij = 11, 12$  and  $44$ ), from which  $B$ ,  $G$  and  $E$  are calculated within the Voigt and Reuss approximation as follows:

$$B_V = \frac{C_{11} + 2C_{12}}{3}, \quad G_V = \frac{C_{11} - C_{12} + 3C_{44}}{5} \quad (3)$$

$$B_R = \frac{1}{3(S_{11} + 2S_{12})}, \quad G_R = \frac{5}{4S_{11} - 4S_{12} + 3S_{44}} \quad (4)$$

$$B = \frac{B_V + B_R}{2}, \quad G = \frac{G_V + G_R}{2}, \quad E = \frac{9BG}{3B + G} \quad (5)$$

The elastic moduli  $B$ ,  $G$  and  $E$  play an important role in describing the resistance of materials against mechanical deformation. Moreover, the Pugh criteria<sup>62</sup> indicate that crystals with Pugh's ratio  $B/G$  and Poisson's ratio  $\nu = (3B - 2G)/(6B + 2G)$  larger than 1.75 and 0.26, respectively, are considered as ductile materials, otherwise brittle materials.

For a direct estimation of material stability of fluoride perovskites  $ABF_3$ , we calculated the cohesive energy  $\Delta E_c$  and formation energy  $\Delta E_f$  of  $ABF_3$  with respect to the chemical reaction  $ABF_3 \rightarrow AF + BF_2$  (see Fig. S2, ESI<sup>†</sup>) as follows:<sup>57,63</sup>

$$\Delta E_c = E(ABF_3) - \sum_i E_i^{\text{atom}} \quad (6)$$

$$\Delta E_f = E(ABF_3) - [E(AF) + E(BF_2)] \quad (7)$$

where  $E(ABF_3)$ ,  $E(AF)$  and  $E(BF_2)$  are the DFT total energies for cubic  $ABF_3$  and  $AF$  with the  $Pm\bar{3}m$  space group and orthorhombic  $BF_2$  with the  $P212121$  space group,<sup>64</sup> respectively, while  $E_i^{\text{atom}}$  is for the isolated atom of  $i$ -th species. The positive or negative formation energy  $\Delta E_f$  directly indicates whether the phase decomposition of  $ABF_3$  into  $AB$  and  $BF_2$  is thermodynamically favorable or unfavorable.

To proceed with calculations of phonon dispersion curves and the corresponding phonon DOS, we utilized the supercell method, as implemented in the ALAMODE code.<sup>65</sup> We applied the finite displacement approach by the use of  $3 \times 3 \times 3$  supercells including 135 atoms (see Fig. S4, ESI<sup>†</sup>). For these supercell calculations, a reduced  $k$ -point mesh of  $3 \times 3 \times 3$  and a cutoff energy of 400 eV were used in accordance with the larger size of the supercell. A  $q$ -point mesh of  $50 \times 50 \times 50$  was used for the calculations of phonon DOS. For the harmonic and quartic interatomic force constants (IFCs), we prepared 5 and 60 different configurations where all atoms were randomly displaced by 0.01 and over 0.06 Å from their equilibrium positions, and then computed the atomic forces for each displaced configuration by performing precise DFT calculations. In these calculations, the VASP code was used as a force calculator and IFCs were extracted by performing the compressive sensing lattice dynamics (CSLD),<sup>66</sup> as implemented in the ALAMODE code.<sup>65</sup> Using these IFCs, the atomic forces were reproduced within the relative errors less than 3% compared to the DFT-calculated ones (see Fig. S3, ESI<sup>†</sup>). With the harmonic and quartic IFCs, we finally calculated the phonon dispersion curves, and total and atom-projected phonon DOSs at 0 K and finite temperatures ranging from 100 to 900 K with a step of 100 K.

### 3 Results and discussion

First we assessed the possibility of fluoride perovskite  $ABF_3$  formation in the cubic phase by evaluating the geometric factors such as the Goldschmidt tolerance factor ( $m_c^*$ ) and the octahedral factor  $t_o = r_B/r_X$ , where  $r_i$  is the ionic radius of the  $i$ -th species.<sup>71,72</sup> In fact, the Goldschmidt tolerance factor  $t_G$



**Table 1** Geometric factors including the Goldschmidt tolerance factor ( $t_G$ ) and octahedral factor ( $t_o$ ), lattice constant of the cubic phase, band gap, effective masses of electrons ( $m_e^*$ ) and holes ( $m_h^*$ ), reduced mass  $m_c^*$ , static dielectric constant ( $\epsilon_s$ ), and exciton binding energy ( $E_b$ ) in fluoride perovskites ABF<sub>3</sub> (A = Na, K and B = Si, Ge), calculated with PBE and HSE06 XC functionals. For comparison, those for typical all-inorganic halide perovskites such as CsPbI<sub>3</sub>, CsGeI<sub>3</sub>, RbGeI<sub>3</sub> and KGeI<sub>3</sub> are provided

Compound	Geometric factor		Lattice constant (Å)		Band gap (eV)			Effective mass ( $m_e$ )			$\epsilon_s$	$E_b$ (meV)
	$t_G$	$t_o$	This	Prev.	PBE	HSE06	Prev.	$m_h^*$	$m_e^*$	$t_G = (r_A + r_X) / [\sqrt{2}(r_B + r_X)]$		
NaSiF <sub>3</sub>	1.03	0.43	4.27	—	0.81	1.24	1.15 <sup>b</sup>	0.26	0.41	0.16	16.23	8.50
NaGeF <sub>3</sub>	0.94	0.57	4.39	—	1.93	2.57	—	0.54	0.65	0.29	21.10	9.02
KSiF <sub>3</sub>	1.13	0.43	4.34	—	0.85	1.37	—	0.35	0.44	0.19	15.28	11.19
KGeF <sub>3</sub>	1.02	0.57	4.45	4.46 <sup>a</sup>	1.97	2.61	2.00 <sup>a</sup>	0.58	0.89	0.35	20.94	10.98
CsPbI <sub>3</sub> <sup>c</sup>	0.85	0.54	6.30	—	1.76	2.24	1.73	0.15	0.20	—	4.7	54
CsGeI <sub>3</sub> <sup>d</sup>	0.93	0.40	5.94	—	1.19	1.64	1.63	0.15	0.16	0.08	19.82	19.24
RbGeI <sub>3</sub> <sup>d</sup>	0.90	0.40	5.91	—	1.31	1.78	—	0.16	0.17	0.08	23.01	13.49
KGeI <sub>3</sub> <sup>e</sup>	0.87	0.34	—	—	0.85	1.24	—	0.60	0.13	0.11	—	—

<sup>a</sup> The lattice constant and band gap were calculated with PBE functional.<sup>41</sup> <sup>b</sup> The band gap were calculated with HSE06 functional.<sup>67</sup> <sup>c</sup> Our previous first-principles work using Quantum ESPRESSO code.<sup>68</sup> <sup>d</sup> Our previous first-principles work using VASP code.<sup>69</sup> <sup>e</sup> Our previous first-principles work for tetragonal phase using VASP code.<sup>70</sup>

is used to check whether the A-site cation can fit between the BX<sub>6</sub> octahedra, while the octahedral factor  $t_o$  is used to assess whether the BX<sub>6</sub> octahedron is stable. According to the empirical criteria, a cubic perovskite structure is formed when  $1.0 \geq t_G \geq 0.8$  and  $t_o \geq 0.41$ .<sup>71,72</sup> These factors have been widely applied to check the formability of oxide and halide perovskites.<sup>73–78</sup> Table 1 lists the evaluated geometric factors of fluoride perovskites ABF<sub>3</sub> (A = Na, K and B = Si, Ge) using the ionic radii of  $r_{Na} = 1.39$  Å,  $r_K = 1.64$  Å,  $r_{Si} = 0.55$  Å,  $r_{Ge} = 0.73$  Å, and  $r_F = 1.285$  Å. Except KSiF<sub>3</sub>, all the compounds under study were found to have  $t_G$  values in the range of  $1.0 \geq t_G \geq 0.8$  for a stable cubic phase, while the  $t_o$  values ranged from 0.43 to 0.57, indicating that BF<sub>6</sub> octahedra are stable in these compounds. Therefore, the fluoride compounds ABF<sub>3</sub> considered in this work are expected to crystallize in a stable perovskite structure. Based on such analysis of geometrical factors, we supposed that the fluoride perovskites ABF<sub>3</sub> stabilize in the cubic phase with the  $Pm\bar{3}m$  space group like the inorganic iodide perovskites.

Through variable-cell structural optimization by using the PBE functional, we determined the equilibrium lattice constants of these lead-free inorganic fluoride perovskites ABF<sub>3</sub> (A = Na, K and B = Si, Ge). The calculated lattice constants are given in Table 1. To the best of our knowledge, there are no experimental data for lattice constants of these fluoride perovskites. Only for the case of KGeF<sub>3</sub>, Houari *et al.*<sup>41</sup> reported a lattice constant of 4.46 Å calculated with the same PBE functional, which coincides with our lattice constant of 4.45 Å. As the ionic radii of A- and B-site cations increase, we found a slight increase in the lattice constant in accordance with common knowledge.

Using the optimized structures, we then calculated their electronic band structures and atomic-resolved DOS by the use of both PBE and HSE06 XC functionals. It was established that the GGA-PBE functional could yield band gaps in good agreement with the experimental values for organic–inorganic hybrid iodide perovskite MAPbI<sub>3</sub> due to a fortuitous error cancellation between the GGA underestimation and overestimation by ignoring the spin–orbit coupling (SOC) effect.<sup>57,79</sup>

For all-inorganic halide perovskites, however, the PBE functional underestimated the band gaps compared to the experimental data, whereas the HSE06 hybrid functional without SOC provided precise band gaps in line with experiments.<sup>68,69</sup> Based on the previously established facts, we adopted the HSE06 hybrid functional without the SOC effect to obtain sufficiently reliable band gaps, being the most important material property of light absorbers to assess the light-to-electricity conversion efficiency of solar cells, and also the PBE functional only for comparison. Fig. 2 shows the electronic band structures and atom-projected DOSs calculated with HSE06 (solid lines) and PBE (dashed lines) functionals for the fluoride perovskites ABF<sub>3</sub> under study.

Our calculations indicate that the electron excitation from the valence band maximum (VBM) to the conduction band minimum (CBM) occurs in a direct way at the edge point *R* (0.5, 0.5, 0.5) of the Brillouin zone (BZ), similar to other halide perovskites in the cubic phase. Obviously, such direct transition is useful for the generation of charge carriers including electrons in the conduction band and holes in the valence band by absorbing light compared with indirect way. We also confirmed that the HSE06 hybrid functional yielded remarkably larger band gaps compared to the PBE functional for the fluoride perovskites due to an upward shift of conduction bands and a downward shift of valence bands. Experimental and theoretical data are not yet available for these fluoride perovskites, except KGeF<sub>3</sub> (NaSiF<sub>3</sub>), for which the band gap was calculated to be 2.00 (1.15) eV with the PBE (HSE06) functional<sup>41,67</sup> in accordance with our calculation of 1.97 (1.24) eV with the same XC functional (see Table 1). As far as believing the HSE06 calculations, the Si-based fluoride perovskites ASiF<sub>3</sub> (A = Na, K) were found to be suitable for visible light absorber applications because of their appropriate band gaps of 1.24 eV (A = Na) and 1.37 eV (A = K). These values are comparable to those of CsPbI<sub>3</sub> (1.76 eV),<sup>68</sup> CsGeI<sub>3</sub> (1.19 eV) and RbGeI<sub>3</sub> (1.31 eV),<sup>69</sup> which have been widely used as light absorbers for PSCs. Meanwhile, Ge-based fluoride perovskites AGeF<sub>3</sub> have a significantly larger band gap over 2.5 eV, implying





Fig. 2 (a)–(d) Electronic band structure and atom-projected density of states (DOS) calculated with PBE (dashed lines) and HSE06 (solid lines) functionals, and (e) and (f) isosurface plot of electronic charge density corresponding to the valence band maximum (VBM) and conduction band minimum (CBM) at a value of  $0.02 |e|/\text{\AA}^3$  in fluoride perovskites  $\text{ABF}_3$  ( $A = \text{Na}, \text{K}$  and  $B = \text{Si}, \text{Ge}$ ). The blue and red lines in the band structures indicate valence and conduction bands, respectively.

their other possible applications such as charge carrier conductors and light emitting diodes. It is worth noting that in accordance with the lattice constant analysis, the larger the ionic radii of A- and B-site cations are, the larger band gaps the fluoride perovskites  $\text{ABF}_3$  provide, which is similar to the halide perovskites.<sup>68,69</sup> Through the analysis of atom-projected DOS, we found that the valence bands are dominated by F-p states coupling with the B-s state while the conduction bands are dominated by F-p and A-p states with negligible contribution from the A-site cations. Moreover, we observed that the VBM is characterized by B-s and F-p states and the CBM has characteristics of B-s and F-s states, as shown in Fig. 2(e) and (f). Similar findings were observed for the all-inorganic and organic-inorganic hybrid halide perovskites, demonstrating that fluoride perovskites have the same electronic mechanism as the halide perovskite.

In order to see how charge carriers such as electrons and holes behave after being generated by absorbing photons, we evaluated their effective masses, dielectric constants, and exciton binding energies by using the PBE functional. Such optical properties are the key quantities for the qualitative assessment of charge carrier mobility. As listed in Table 1, the Si-based compounds  $\text{ASiF}_3$  ( $A = \text{Na}$  and  $\text{K}$ ) have much lighter effective masses of electrons and holes than the Ge-based compounds  $\text{AGeF}_3$ . In accordance with the band gaps, this indicates that the former is more favourable for light absorbers than the latter. In particular,  $\text{NaSiF}_3$  exhibits the lightest effective masses of electrons and holes as  $0.26m_e$  and  $0.41m_e$  among the four fluoride perovskites under study, implying the highest mobilities of charge carriers. These values are slightly larger than those of  $\text{ABl}_3$  ( $A = \text{Cs}, \text{Rb}$  and  $B = \text{Pb}, \text{Ge}$ ),<sup>68,69</sup> but

comparable with those of tetragonal  $\text{KGeI}_3$ .<sup>70</sup> We note that the effective masses in  $\text{AGeF}_3$  are still small enough for applications as charge carrier conductors.

Also, the static dielectric constants  $\epsilon_s$  of Si-based fluoride perovskites, calculated using the DFPT approach while considering the atomic displacement effect, are smaller than those of the Ge-based ones (see Table 1). The calculated values range from 15.28 for  $\text{KSiF}_3$  to 21.10 for  $\text{NaGeF}_3$ , which are considerably larger than those of organic-inorganic hybrid MA-based perovskites (3–5).<sup>57,63</sup> This indicates that these fluoride perovskites are expected to have much smaller exciton binding energies than the hybrid chloride perovskites as the exciton binding energy is inversely proportional to the square of the static dielectric constant, which is favourable for solar cell performance.

Then, using the calculated effective masses and static dielectric constants, we determined the exciton binding energies  $E_b$  with eqn (2), which is a crucial quantity to estimate whether the electrons and holes recombine or behave as free charge carriers. The calculated exciton binding energies in fluoride perovskites  $\text{ABF}_3$  are provided in Table 1. We found that the  $E_b$  values are very low, below 12 meV, which can be attributed to very light effective masses and relatively large static dielectric constants. When compared with other halide perovskites, these values are comparable to those of all-inorganic Ge-based perovskites  $\text{AGeX}_3$  ( $A = \text{Cs}, \text{Rb}$  and  $X = \text{I}, \text{Br}, \text{Cl}$ ),<sup>69</sup> while distinctly lower than those of hybrid perovskite  $\text{MAPbX}_3$  (40–300 meV).<sup>57,63</sup> We also note that the calculated exciton binding energy has the similar variation tendency according to the selection of A- and B-site cations as the effective mass and band gap. To sum up the calculated band



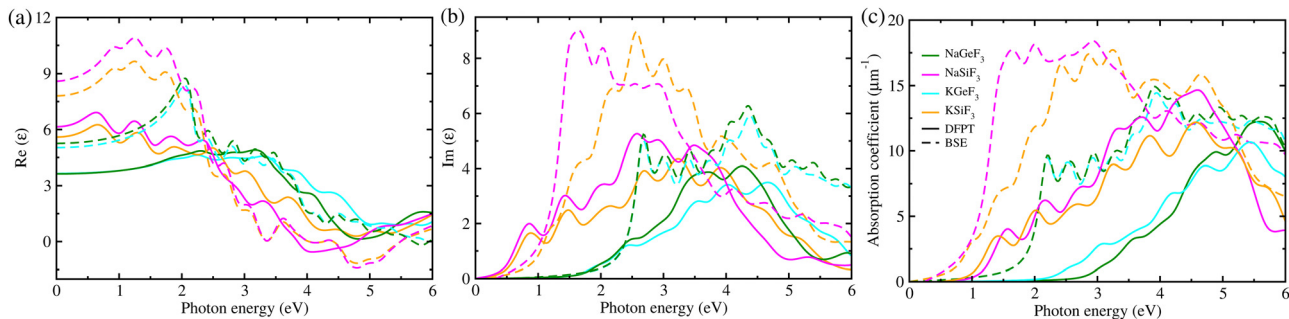


Fig. 3 (a) Real and (b) imaginary parts of frequency-dependent dielectric function  $\epsilon(\omega)$ , and (c) light-absorption coefficient calculated with DFPT (solid) and BSE/GW (dashed) methods in fluoride perovskites  $ABF_3$  ( $A = Na, K$  and  $B = Si, Ge$ ).

gaps and charge transport properties, we emphasize that the Si-based fluoride perovskites  $ASiF_3$  ( $A = Na, K$ ) are promising light-absorber materials for PSCs because of their suitable band gaps (1.2–1.4 eV) and low exciton binding energies (8.5–11.2 meV).

Subsequently, we considered the light-absorption coefficients  $\alpha(\omega)$  as a function of frequency  $\omega$ , which were calculated using the frequency-dependent dielectric functions  $\epsilon(\omega)$  with eqn (1). Fig. 3 displays the real and imaginary parts of  $\epsilon(\omega)$ , calculated with DFPT (solid) and BSE/GW (dashed) approaches, and the  $\alpha(\omega)$  value for four fluoride perovskites. The BSE/GW calculations provided higher peak values of  $\epsilon(\omega)$  compared with the DFPT calculations. With reference to  $\alpha(\omega)$ , the highest peaks were estimated to range from 11 (15) to 15 (18)  $\mu\text{m}^{-1}$  for DFPT (BSE/GW) calculations, which are lower than  $\sim 20 \mu\text{m}^{-1}$  in  $\text{MAPbX}_3$ <sup>57,63</sup> but higher than 2–4  $\mu\text{m}^{-1}$  in  $\text{AGeX}_3$  ( $A = Cs, Rb$  and  $X = I, Br, Cl$ )<sup>69</sup>, indicating that the fluoride perovskites conceived in this work can exhibit reasonable light harvesting capability. In accordance with the band gaps, the Si-based fluoride perovskites  $ASiF_3$  ( $A = Na, K$ ) show an absorption onset at a lower photon energy than the Ge-based ones. Moreover, we found that the light absorption of the Si-based perovskites spans the whole visible light spectrum, which is clearly beneficial for enhancing the efficiency of solar cells. Such findings provide strong evidence that the Si-based fluorides are a promising absorber for solar cell applications.

Using the calculated band gaps and light-absorption coefficients as input, we estimated the SLME values for PSCs employing fluoride perovskites as a light absorber, providing the maximum efficiency for a single-junction solar cell. Fig. 4 shows the calculated SLME as a function of absorber thickness in fluoride perovskites  $ABF_3$  ( $A = Na, K$  and  $B = Si, Ge$ ). We found that the SLMEs become saturated to nearly constant over a layer thickness of  $\sim 1 \mu\text{m}$  for the four fluoride perovskites. The resulting SLME values were found to be 34% and 13% for Si- and Ge-based fluoride perovskites, respectively. Therefore, the solar cells using Si-based fluoride perovskites  $ASiF_3$  ( $A = Na, K$ ) as a light-absorber are sufficiently competitive compared with other PSCs and the conventional Si-based solar cells. Meanwhile, the calculated SLME ( $\sim 34\%$ ) of Si-based fluoride perovskites is larger than the theoretical calculations ( $\sim 30\%$ ) of chalcogenide perovskites  $AZrS_3$  ( $A = Ca, Sr, \text{ and } Ba$ ),<sup>59</sup>



Fig. 4 Spectroscopic limited maximum efficiency (SLME) as a function of thickness in lead-free all-inorganic fluoride perovskites  $ABF_3$  ( $A = Na, K$  and  $B = Si, Ge$ ).

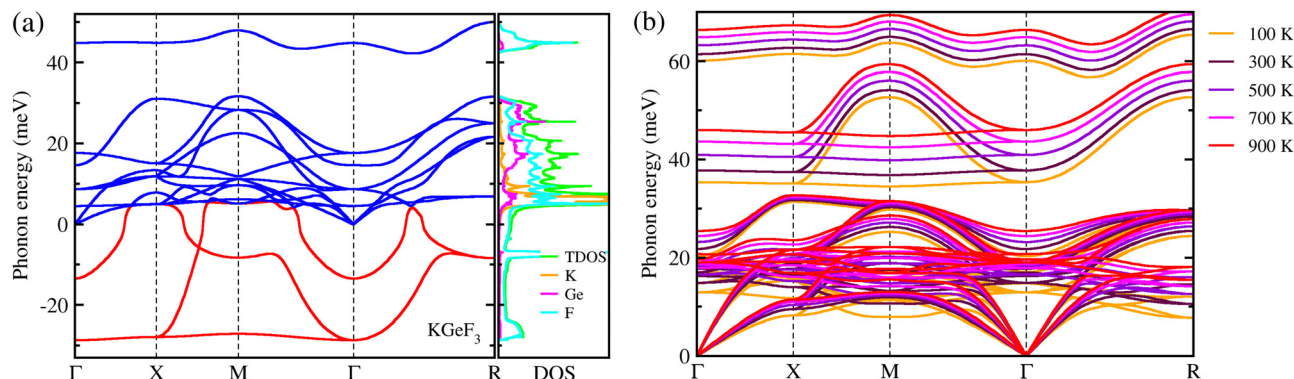
indicating that the Si-based fluoride perovskite is superior to the chalcogenide perovskite for solar cell applications.

Finally, we estimated the mechanical, chemical and thermodynamical stabilities of the fluoride perovskites based on the analysis of their elastic constants, and cohesive and formation energies. The elastic stiffness and compliance constants were obtained from the DFPT calculations to investigate the mechanical stability of the fluoride perovskites  $ABF_3$  ( $A = Na, K$  and  $B = Si, Ge$ ). As listed in Table 2, for all the fluoride perovskites, the three independent elastic stiffness constants of  $C_{11}$ ,  $C_{12}$  and  $C_{44}$  satisfy the well-known Born stability criteria for cubic crystals,<sup>80</sup> expressed as  $C_{11} > 0$ ,  $C_{44} > 0$ ,  $C_{11} - C_{12} > 0$  and  $C_{11} + 2C_{12} > 0$ , therefore implying their mechanical stability at zero pressure. Furthermore, it was found that all the fluoride perovskites  $ABF_3$  are ductile materials in accordance with the Pugh criteria<sup>62</sup> because the calculated values of Pugh's ratio  $B/G$  and Poisson's ratio  $\nu$  are larger than the threshold values of 1.75 and 0.26, respectively. On the other hand, the cohesive energies  $\Delta E_c$  were found to be negative for all the fluoride perovskites, but the formation energy  $\Delta E_f$  was estimated to be negative only for  $\text{KGeF}_3$ , thereby indicating that only the  $\text{KGeF}_3$  exhibits a good chemical stability against the



**Table 2** Elastic stiffness constant ( $C_{ij}$ ), compliance constant ( $S_{ij}$ ), bulk modulus ( $B$ ), shear modulus ( $G$ ), Young's modulus ( $E$ ), Pugh's ratio ( $B/G$ ), Poisson's ratio ( $\nu$ ), cohesive energy ( $\Delta E_c$ ) and formation energy ( $\Delta E_f$ ) per atom in all-inorganic lead-free fluoride perovskites  $ABF_3$  ( $A = Na, K$  and  $B = Si, Ge$ )

Compound	$C_{ij}$ (GPa)			$S_{ij}$ (TPa <sup>-1</sup> )			Elastic moduli (GPa)			Ratio		$\Delta E$ (eV per atom)	
	$C_{11}$	$C_{12}$	$C_{44}$	$S_{11}$	$S_{12}$	$S_{44}$	$B$	$G$	$E$	$B/G$	$\nu$	$\Delta E_c$	$\Delta E_f$
NaSiF <sub>3</sub>	173.65	63.45	35.96	0.72	-0.19	2.78	100.14	22.64	63.17	4.42	0.39	-4.25	0.28
NaGeF <sub>3</sub>	177.62	51.34	34.62	0.65	-0.14	2.88	93.54	23.86	65.97	3.92	0.38	-4.19	0.13
KSiF <sub>3</sub>	164.85	62.85	43.48	0.77	-0.21	2.29	96.87	24.17	66.95	4.00	0.38	-4.33	0.13
KGeF <sub>3</sub>	168.29	50.92	40.05	0.69	-0.16	2.49	89.94	24.67	67.82	3.64	0.37	-4.29	-0.04



**Fig. 5** Phonon dispersion curves calculated at 0 K (a) and finite temperatures ranging from 100 to 900 K with a step of 100 K (b) for  $KGeF_3$ .

chemical decomposition into its chemical reactants. It should be noted that although  $ASiF_3$  perovskites have suitable band gaps and low exciton binding energy, they show poor chemical stability, which means that these materials are unstable and can be decomposed quickly. In order to address the thermodynamical stability of the mechanically and chemically stable  $KGeF_3$ , we calculated its phonon dispersions at 0 K and finite temperatures. As shown in Fig. 5(a), anharmonic phonon modes with negative phonon energies were observed in the dispersion curves at 0 K, which are so-called soft phonon modes, a typical characteristic of halide and oxide perovskites in the cubic phase. The atom-projected phonon DOS reveals that the soft phonon modes are mainly attributed to the atomic vibrations of F atoms with a small contribution from K and Ge atoms. However, the anharmonic phonon modes were all renormalized by considering the finite temperature effects, demonstrating that the cubic phase of  $KGeF_3$  is dynamically stable (Fig. 5(b)). To sum up, only  $KGeF_3$  is stable in mechanic, chemical and thermodynamic aspects among the four lead-free fluoride perovskites, together with a moderate SLME of  $\sim 13\%$ , making it a suitable and promising light-absorber for nontoxic, long-term stable and high-performance solar cells.

## 4 Conclusions

In this work, we have systematically investigated the structural, electronic and optical properties, and mechanical and chemical stabilities of all-inorganic lead-free fluoride perovskites  $ABF_3$  ( $A = Na, K$  and  $B = Si, Ge$ ) using first-principles calculations. From the assessment of geometric factors such as Goldschmidt tolerance and octahedral factors, we confirmed that  $ABF_3$  could

stabilize in the cubic perovskite structure. Our calculations illustrated that their material properties, such as lattice constants, band gaps, effective masses of electrons and holes, exciton binding energies, exhibit a similar increasing tendency according to the increase in ionic radii of A- and B-site cations. We emphasize that the band gaps and exciton binding energies of the Si-based perovskites  $ASiF_3$  were calculated to be 1.24 eV and 8.50 meV for  $A = Na$ , and 1.37 eV and 11.19 meV for  $A = Si$ , respectively, which are suitable for absorbing visible light and transferring charge carriers. Moreover, the calculated spectroscopic limited maximum efficiency for the Si-based fluoride perovskites reaches  $\sim 34\%$  with a layer thickness over 1  $\mu m$ , indicating a competitiveness of solar cells. Through the analysis of calculated elastic constants, formation energies and phonon dispersion curves, we revealed that all the fluoride perovskites  $ABF_3$  ( $A = Na, K$  and  $B = Si, Ge$ ) are mechanically stable in the cubic phase, whereas only  $KGeF_3$  is chemically stable, not decomposing back into its precursor materials due to a negative formation energy, and also thermodynamically stable. Finally, we concluded that  $KGeF_3$  is the most suitable and promising light-absorber for solar cells due to its moderate material properties combined with good mechanical, chemical and thermodynamic stabilities among the four compounds, highlighting the potentiality of using all-inorganic lead-free fluoride perovskites for nontoxic and high-performance solar cells.

## Author contributions

Chol-Jin Park and Un-Gi Jong developed the original project, performed the calculations and drafted the first manuscript. Chung-Jin Kang, Yun-Sim Kim, and Yun-Hyok Kye assisted in



DFT calculations and the post-processing of calculation results, and contributed to useful discussions. Chol-Jun Yu supervised the work. All authors reviewed the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work is supported as part of the basic research project "Design of New Energy Materials" (no. 2021-12) funded by the State Commission of Science and Technology, DPR Korea. Computations have been performed on the HP Blade System C7000 managed by the Faculty of Materials Science, Kim Il Sung University.

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