

## Perovskites

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## B-Site Co-Alloying with Germanium Improves the Efficiency and Stability of All-Inorganic Tin-Based Perovskite Nanocrystal Solar Cells

Maning Liu, Hannu Pasanen, Harri Ali-Löytty, Arto Hiltunen, Kimmo Lahtonen, Syeda Qudisia, Jan-Henrik Smått, Mika Valden, Nikolai V. Tkachenko, and Paola Vivo\*

**Abstract:** Colloidal lead-free perovskite nanocrystals have recently received extensive attention because of their facile synthesis, the outstanding size-tunable optoelectronic properties, and less or no toxicity in their commercial applications. Tin (Sn) has so far led to the most efficient lead-free solar cells, yet showing highly unstable characteristics in ambient conditions. Here, we propose the synthesis of all-inorganic mixture Sn-Ge perovskite nanocrystals, demonstrating the role of  $\text{Ge}^{2+}$  in stabilizing  $\text{Sn}^{2+}$  cation while enhancing the optical and photophysical properties. The partial replacement of Sn atoms by Ge atoms in the nanostructures effectively fills the high density of Sn vacancies, reducing the surface traps and leading to a longer excitonic lifetime and increased photoluminescence quantum yield. The resultant Sn-Ge nanocrystals-based devices show the highest efficiency of 4.9%, enhanced by nearly 60% compared to that of pure Sn nanocrystals-based devices.

## Introduction

Colloidal halide perovskite nanocrystals (PNCs), with their size-tunable narrow-band emission, higher photoluminescence quantum yields (PLQY) than the corresponding bulk crystals, and enhanced surface-to-volume ratio,<sup>[1]</sup> are promising candidates for the next-generation optoelectronics including PVs,<sup>[2,3]</sup> light-emitting diodes,<sup>[4,5]</sup> photodetectors,<sup>[6]</sup>

and lasers.<sup>[7]</sup> Because of the intrinsic instability in ambient conditions of organic cations (for example,  $\text{CH}_3\text{NH}_3^+$ ) at the A site of a typical  $\text{ABX}_3$  perovskite structure,<sup>[8]</sup> all-inorganic lead (Pb) based PNCs, such as  $\text{CsPbX}_3$  ( $\text{X} = \text{Cl}, \text{Br}, \text{or I}$ ), are preferred to pursue high stability while retaining the advanced optical properties, for example, the high PLQY.<sup>[9]</sup> However, toxic Pb is still the key constituent of the most efficient halide perovskite materials, thus impeding PNCs widespread commercialization.

Tin (Sn) is one of the most promising alternative candidates to replace Pb, being its closest analogue with similar structural and electronic properties.<sup>[10,11]</sup> As a result, the most efficient Pb-free perovskite solar cells (PSCs) to date always include Sn in the perovskite composition.<sup>[12–14]</sup> However, integrating Sn in spatially confined nanocrystals (NCs) has been rarely successful because of its high instability.  $\text{Sn}^{2+}$  (the B site in the perovskite structure) is rapidly oxidized to  $\text{Sn}^{4+}$  in ambient conditions.<sup>[15,16]</sup> Furthermore, the high density of surface vacancies<sup>[17]</sup> result in the low photoluminescence quantum yield (PLQY) of Sn-based PNCs. As another promising Pb-free candidate, germanium (Ge) is a defect-tolerant and nontoxic semiconductor still highlighted by the limited studies.<sup>[18,19]</sup> Among the synthesized materials,  $\text{CsGeI}_3$  has a direct band gap ( $E_g$ ) of 1.6 eV, which is suitable for PVs. When compared to  $\text{CsSnI}_3$ ,  $\text{CsGeI}_3$  does display, however, inferior optoelectronic properties owing to the lone-pair effect<sup>[20]</sup> and relatively larger  $E_g$ . Inspired by the heterovalent B-site co-alloying strategy for Pb and Sn, several reports have proposed to mix Sn with Ge for a new candidate of mixture lead-free perovskite.<sup>[21–26]</sup> By first-principles computations, Ju et al. predicted several potentially promising mixed Sn-Ge halide perovskites as light absorbers for solar cells.<sup>[27]</sup> Ma et al. presented a comprehensive theoretical study on the mixed vacancy-ordered inorganic double perovskite  $\text{Cs}_2\text{Sn}_{(1-x)}\text{Ge}_x\text{I}_6$ , indicating that the concentration of Ge doping severely influences the thermodynamic, electronic, and mechanical properties.<sup>[25]</sup> Ito et al. experimentally showed an improved power conversion efficiency (PCE) of 4.48% for  $\text{FA}_{0.75}\text{MA}_{0.25}\text{Sn}_{1-x}\text{Ge}_x\text{I}_3$ -based solar cells with a 5% Ge doping into the perovskite, compared with that (3.31%) of pure Sn-based ones.<sup>[28]</sup> Later on, Chen et al. reported the first synthesis of  $\text{CsSn}_{0.5}\text{Ge}_{0.5}\text{I}_3$  bulk crystals via a vapor method, showing a relatively high PCE of 7.11% but employing a complex, non-scalable, and expensive synthetic route.<sup>[29]</sup> However, there are no examples of mixed Sn-Ge perovskite nanocrystals.

Here, we pioneered the synthesis of all-inorganic Ge alloyed  $\text{CsSnI}_3$  perovskite nanocrystals (Sn-PNCs) via a facile

[\*] Dr. M. Liu, H. Pasanen, Dr. A. Hiltunen, Prof. N. V. Tkachenko, Assoc. Prof. P. Vivo  
Chemistry and Advanced Materials Group  
Faculty of Engineering and Natural Sciences, Tampere University  
P.O. Box 692, 33014 Tampere (Finland)  
E-mail: paola.vivo@tuni.fi

Dr. H. Ali-Löytty, Prof. M. Valden  
Surface Science Group  
Faculty of Engineering and Natural Sciences, Tampere University  
P.O. Box 692, 33014 Tampere (Finland)

Dr. K. Lahtonen  
Faculty of Engineering and Natural Sciences, Tampere University  
P.O. Box 692, 33014 Tampere (Finland)

S. Qudisia, Dr. J. H. Smått  
Laboratory of Molecular Science and Engineering  
Åbo Akademi University  
Porthansgatan 3–5, 20500 Turku (Finland)

Supporting information and the ORCID identification number(s) for the author(s) of this article can be found under:  
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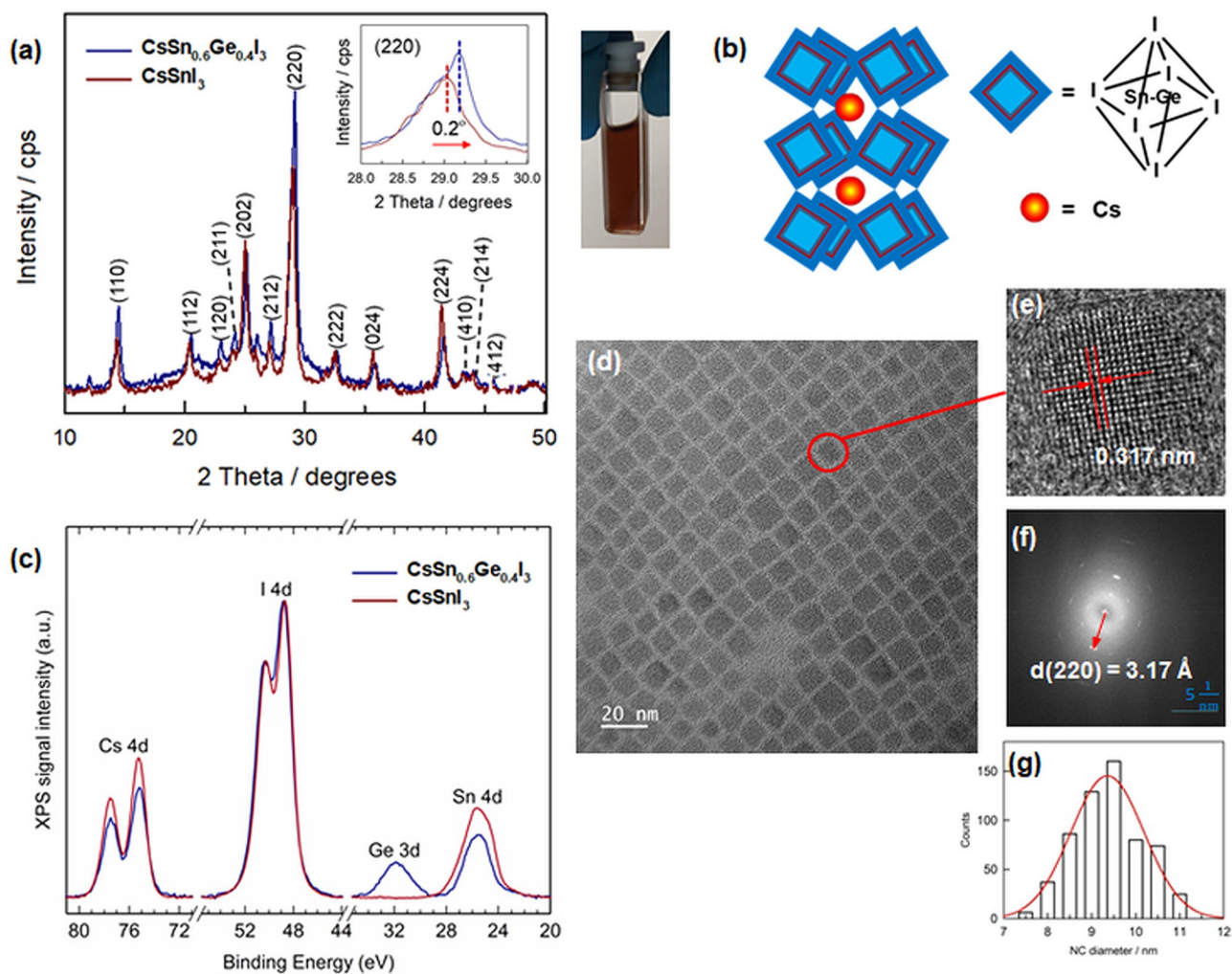
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and low-cost solution process. The novel  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  perovskite nanocrystals (SnGe-PNCs) exhibit significantly improved optical and photophysical properties compared to those of reference Sn-PNCs. The stability of as-formed  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  has also been effectively enhanced upon the introduction of Ge. The partial replacement of Sn atoms by Ge atoms in the nanostructure can effectively fill the high density of Sn vacancies and reduce the surface traps, resulting in a longer excitonic lifetime and improved PLQY. In this work, the PCE (4.9%) of the PSCs employing  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  nanocrystals is enhanced by nearly 60% compared to that (3.1%) of  $\text{CsSnI}_3$ -based devices.

## Results and Discussion

### Synthesis of $\text{CsSn}_x\text{Ge}_{1-x}\text{I}_3$ NCs

Based on the previous hot-injection methods to produce Sn-based PNCs,<sup>[30,31]</sup> we developed a modified reaction route for the synthesis of  $\text{CsSn}_x\text{Ge}_{1-x}\text{I}_3$  NCs. An equal molar ratio of  $\text{SnI}_2$  and  $\text{GeI}_2$  has been utilized to target the half replacement of Sn atoms by Ge atoms in the expected stoichiometry (namely,  $x = 0.5$ ). We found that the injection temperature is crucial to form stable SnGe-PNCs at least before the purification process. The temperature should be above 240 °C. After the injection of Cs-oleate into the precursor of Sn/Ge halide source, a sudden color change from transparent orange to dark brown occurred (see the appearance of  $\text{CsSn}_x\text{Ge}_{1-x}\text{I}_3$  NCs suspension in the side photo in Figure 1a) and no further color change was observed after 3 s reaction time, indicating that the formation of SnGe-PNCs features the extremely swift reaction kinetics, similarly as in the case of



**Figure 1.** a) X-ray diffraction patterns of the as-prepared  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs and reference  $\text{CsSnI}_3$  NCs. The side picture shows the appearance of  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs in suspension. b) Illustration of a unit cell of  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs in orthorhombic crystal structure. c) X-ray photoelectron spectroscopy (XPS) analysis of  $\text{CsSnI}_3$  NCs and  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs. d) Transmission electron microscopy (TEM) image of  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs. e) A high-resolution TEM (HRTEM) image of a single  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  nanocrystal. f) Fast Fourier transform (FFT) analysis of the HRTEM image. g) Narrow size distribution histogram of the particles.

conventional Pb-based PNCs.<sup>[32]</sup> The detailed synthetic route for SnGe-PNCs and the reference Sn-PNCs has been described in Supporting Information.

To examine the crystal structure of as-synthesized PNCs, X-ray diffraction (XRD) patterns for PNCs in film state have been measured. Figure 1a shows the comparison of XRD patterns for  $\text{CsSn}_x\text{Ge}_{1-x}\text{I}_3$  NCs and reference  $\text{CsSnI}_3$  NCs with the complete assignment of all the featured peaks corresponding to the halide perovskite structure.<sup>[31,33]</sup> The XRD scan of reference Sn-PNCs matches well with JCPDS reference 00-043-1162 from the ICDD database (Supporting Information, Figure S1a), indicating the formation of an orthorhombic crystallographic phase that belongs to the space group *Pnma* (62), which is consistent with previous literature findings for similar Sn-based PNCs.<sup>[31]</sup> It is noted that the XRD pattern of the newly synthesized SnGe-PNCs shows slight shifts in the peak positions compared to that of the reference, that is,  $14.2^\circ$  to  $14.5^\circ$  at (110) and  $29.0^\circ$  to  $29.2^\circ$  at (220) (see inset in Figure 1a), suggesting a slight change in the size of the unit cell dimension. This change is attributed to a partial exchange of the large Sn atoms with relatively small Ge atoms, resulting in the shrinkage of the unit cell and an upward shift in the  $2\theta$  angles. Therefore, we similarly assign the structure of as-formed SnGe-PNCs with orthorhombic crystallographic phase (Figure 1b) as in the case of the reference. The surface compositions of  $\text{CsSn}_x\text{Ge}_{1-x}\text{I}_3$  NCs and reference  $\text{CsSnI}_3$  NCs in film state were analyzed by X-ray photoelectron spectroscopy (XPS) measurements (Figure 1c). The concentration of iodine should be considered as a lower estimate because of the X-ray and electron beam induced desorption of I during XPS and EDS measurements.<sup>[34]</sup>  $\text{CsSn}_x\text{Ge}_{1-x}\text{I}_3$  NCs had similar surface concentrations for Sn and Ge that support the successful substitution of Sn with Ge (Supporting Information, Table S1). Only one chemical state was resolved on  $\text{CsSn}_x\text{Ge}_{1-x}\text{I}_3$  NCs for Cs (Cs  $4d_{5/2}$  at 75.2 eV), Sn (Sn  $4d_{5/2}$  at 25.2 eV), Ge (Ge  $3d_{5/2}$  at 31.6 eV), and I (I  $4d_{5/2}$  at 48.8 eV), which can be attributed to  $\text{Cs}^+$ ,  $\text{Sn}^{2+}$ ,  $\text{Ge}^{2+}$ , and  $\text{I}^-$ , respectively, in accordance with the  $\text{Cs}^+(\text{Sn}/\text{Ge})^{2+}\text{I}_3^{1-}$  perovskite structure.<sup>[29,35]</sup> The thorough XPS analysis confirms the stoichiometry of the PNCs films at the surface (that is, within the XPS information depth of 11 nm), to be  $\text{CsSnI}_3$  and  $\text{CsSn}_{0.5}\text{Ge}_{0.5}\text{I}_3$ , with minor deviations (Supporting Information, Table S1). To further verify the stoichiometry of PNCs in bulk, we conducted energy-dispersive X-ray spectroscopy (EDS) measurements for the PNCs films. The EDS layered images SnGe-PNCs (Supporting Information, Figure S2) for the compositional elements show the even elemental distribution on the surface of PNCs film. The EDS analysis elucidates the stoichiometry of SnGe-PNCs bulk film (Supporting Information, Table S1). Interestingly, the elemental ratio between Sn and Ge turns to 2:1, meaning that the actual replacement proportion of Sn atoms by Ge atoms is nearly 40% while considering the results of XPS data for the EDS analysis. We thus specify  $x=0.6$  in the stoichiometry, outlining  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  as the final composition. The Ge amount dependent synthesis and properties of SnGe-PNCs will be investigated in a separate work. It is noted that the ratio of Cs:I in the bulk case is less than the nominal value (for example, 1:3; Supporting Information, Table S1),

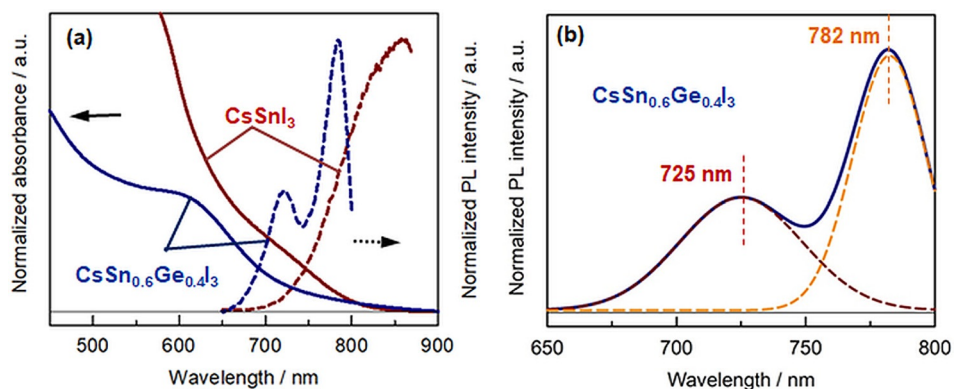
which can be attributed to the small ratio (1:4) of the injected volume (Cs-oleate) and that of  $\text{SnI}_2/\text{GeI}_2$  precursor, creating the halide-rich environment to form polyhedral  $[\text{SnI}_6]^{4-}$  as nucleus for the growth of nanocrystals.<sup>[36]</sup>

The morphology of as-synthesized  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs was investigated using transmission electron microscopy (TEM). Figure 1d reveals that the nanoparticles have a cubic shape with an average square diameter of about 9.4 nm for  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs, which is slightly larger than that (ca. 8.3 nm) of  $\text{CsSnI}_3$  NCs (Supporting Information, Figure S3a). Since the synthesis conditions were identical for both PNCs, this may suggest that small Ge atoms can potentially assist large Sn atoms to de-focus over a longer growth duration more effectively than pure Sn-based ones according to a kinetics model for the growth of Pb-based PNCs,<sup>[37]</sup> hence resulting in the bigger size of SnGe-PNCs. Based on the HRTEM image of a single  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  nanocube (Figure 1e), the lattice distance is 0.317 nm that corresponds to (220) facets, highly consistent with the calculated lattice spacing of 3.17 Å from the Fast Fourier Transform (FFT) analysis (Figure 1f) of the HRTEM image. This, together with the previous XRD data, further supports an orthorhombic crystal structure.<sup>[31]</sup> Narrow size distributions have been observed with full widths at half-maximum (FWHM) of 1.9 nm (Figure 1g) and 1.4 nm (Supporting Information, Figure S3b) for SnGe-PNCs and Sn-PNCs, respectively. This suggests that our modified hot-injection synthetic route can precisely control the nucleation and growth of nanocubes.

#### Optical Properties of $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$ NCs

The optical properties of as-synthesized  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs were investigated and compared to those of reference  $\text{CsSnI}_3$  NCs at room temperature in ambient conditions. The comparison of absorption and photoluminescence (PL) spectra of both PNCs in suspension is presented in Figure 2a. A clear blue shift has been observed for both absorption onset and PL peak from  $\text{CsSnI}_3$  NCs to  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs due to the wider  $E_g$  obtained upon the introduction of Ge in the Sn-based nanocrystal structure. The PL spectrum of  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs shows a main peak at around 780 nm (1.6 eV) with a FWHM of 65 nm, while a small PL sub-peak at around 725 nm has been detected that was not observed for the case of reference  $\text{CsSnI}_3$  NCs. We deconvoluted the emission spectrum with two contributed bands showing respective Gaussian peaks in Figure 2b, centered at 725 and 782 nm.<sup>[38]</sup> Interestingly, we have found that a small amount of SnGe-based nanorods formed together with the nanocubes. The nanorods have an average diameter of about 4.5 nm and a mean length of about 50 nm (see the corresponding TEM image in the Supporting Information, Figure S4a). This finding explains the origin of the 725 nm emission. The band gap of perovskite nanorods is closely related to their diameter or width.<sup>[39,40]</sup> The small size of the Sn-Ge nanorods (diameter of 4.5 nm) may induce a quantum confinement effect compared to the big SnGe-nanocubes (diameter of 9.4 nm), with a corresponding shift towards a higher band gap when the size of PNCs is less than their Bohr radius.<sup>[39]</sup> This



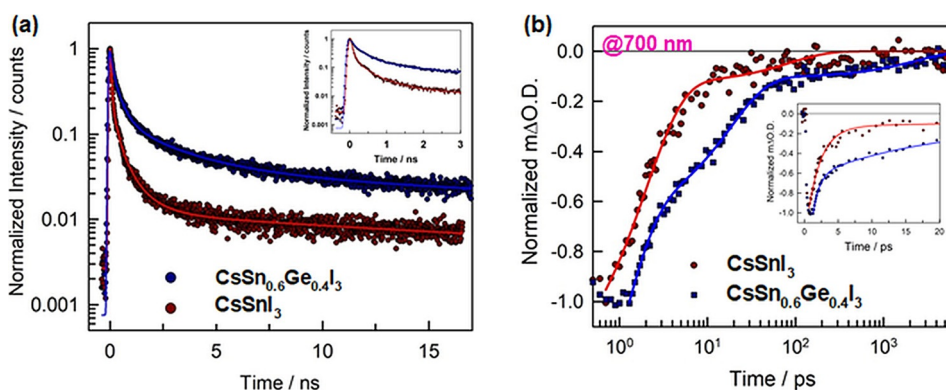


**Figure 2.** a) Absorption spectra (solid lines) and photoluminescence (PL) spectra (dashed lines) of CsSnI<sub>3</sub> NCs and CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs in suspension, respectively. b) Deconvolution of the PL profile of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs.

phenomenon has not been observed for reference Sn-PNCs, since in that case a change in the morphology has not been detected, as for the case of the mixed Sn<sup>2+</sup> and Ge<sup>2+</sup> cations. This point will be more thoroughly investigated in a future work. The absorption and PL spectra in film state for both PNCs (Supporting Information, Figure S4b) are nearly identical to those in suspensions, suggesting that no morphological change occurs during the formation of the PNC bulk film (see the inset photo of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs film in the Supporting Information, Figure S4b). The PL quantum yields (PLQYs) were sequentially measured as 0.95 % and 0.16 % for CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs and reference CsSnI<sub>3</sub> NCs, respectively. Although the absolute values of PLQYs are low compared to the lead analogues, a dramatic increase in the PLQY of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs has been achieved compared to that of the Ge-free counterpart. Since the dominant defects in Sn-based PNCs are interstitial, and Sn displays low defect formation energy in terms of the metal terminate mechanism,<sup>[41]</sup> the involvement of Ge effectively fills in Sn vacancies and reduces the number of traps, resulting in the enhanced PLQY.

### Photophysical Properties of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs

We then turn to assess the PL dynamics of as-synthesized nanocrystals by conducting the time-correlated single photon counting (TCSPC) measurements in the pico- and nano-second regime. The PL decays of nanocubes in suspension are presented in Figure 3a. Apparently, the emission decay of CsSnI<sub>3</sub> NCs is faster than that of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs. Poisson statistics of defects distribution in PNCs have been employed to model decays and fit the experimental data (see the detail of the Poisson model in Supporting Information).<sup>[42,43]</sup> It was found that two types of defects need to be assumed to obtain a reasonable data fit. The fitting results are shown in the Supporting Information, Table S2, where  $\tau$  is the quenching time constant in a nanocrystal having exactly one defect and  $c$  is the relative defect concentration, or the average number of defects per nanocrystal. The observation of short PL lifetime for CsSnI<sub>3</sub> NCs makes a good agreement with previously reported fast decays for Sn-based nanocrystals within several hundreds of picoseconds.<sup>[30,31]</sup> In contrast to the reference, the quenching time constants of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs have been extended by a factor of 4 to 490 and 4200 ps, respectively. At the same time the relative defect concentrations are also



**Figure 3.** a) Time-correlated single photon counting (TCSPC) decays of CsSnI<sub>3</sub> NCs and CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs in suspension (the inset shows magnified decays in early timescale), excited at 405 nm. Solid lines present the fitting results by Poisson statistics model. c) Ultrafast transient absorption (TA) decays of the encapsulated CsSnI<sub>3</sub> NCs and CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs films excited upon a 500 nm pump light with an excitation energy intensity of 10 μJ cm<sup>-2</sup>. Solid lines show the fitting results with a bi-exponential function.

lower for  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs,  $c_1 = 2.0$ ,  $c_2 = 1.7$ , vs.  $c_1 = 2.3$ ,  $c_2 = 3.3$  of  $\text{CsSnI}_3$  NCs (Table S2). This supports our hypothesis of Ge filling the Sn vacancies, which has the two side effects of 1) lowering the density of defects and 2) the less detrimental effect of the defects left, that is, slower quenching. Furthermore, it is noted that the PL decay lifetimes of both PNCs in film (Supporting Information, Figure S5) state are similar to those of their suspensions, suggesting that the defect distribution for both PNCs is less influenced by their variable phases.

To further investigate the influence of Ge replacement on the charge carrier dynamics and the nonradiative processes, we also carried out the ultrafast transient absorption (TA) spectroscopy on the encapsulated  $\text{CsSnI}_3$  NCs and  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs films on glass, respectively. The TA spectra of the  $\text{CsSnI}_3$  NCs film excited upon a 500 nm pump light after different time delays are shown in the Supporting Information, Figure S6a. The overall TA spectra are dominated by one distinct bleach signal with a peak at 700 nm, which are corresponding to the first (660–800 nm) excitonic absorption band, in a good agreement with previously reported assignments for  $\text{CsSnI}_3$  QDs.<sup>[44]</sup> It is noted that there is a positive signal appearing with a peak at 600 nm, likely attributed to the overlap of the bleach with the photoinduced absorption (PIA) between 570 and 650 nm. The comparative TA spectra of  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs film measured upon the identical condition as for the case of reference Sn-PNCs film are shown in the Supporting Information, Figure S6b. Interestingly, a second bleached band appears at 750 nm in addition to the main bleached band below 700 nm. We conclude that the second bleach arises from the small amount of nanorods mixed into the sample. After assigning the origins in the TA spectra of both PNCs films, we now turn to assess the TA kinetics. A comparison of TA decays for  $\text{CsSnI}_3$  NCs and  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs films, monitored at 770 nm in early time-scale are shown in the Supporting Information, Figure S6c. It is clearly observed that both TA profiles instantly rise within the instrument time (ca. 0.2 ps), corresponding to the formation of hot carriers after excitation well above the band gap (500 nm pump light). The hot carriers then relax to the band edge within a cooling time of about 2.5 ps for both films, indicating that the partial replacement of Ge atoms in nanostructure does not make obvious effect on the hot carrier kinetics. We then compared the TA decays of both films monitored at 700 nm (Figure 3b) to evaluate the dynamics of bleaching recovery. The TA decay curves can be well fitted using a bi-exponential function:  $\Delta\text{O.D.} = A_1 \times \exp(-t/\tau_1) + A_2 \times \exp(-t/\tau_2)$ , which provides two exponential decay constants for both films (summarized in the Supporting Information, Table S3). As discussed earlier, Ge atoms can effectively

fill the high density of Sn vacancies, reducing the shallow traps and leading to a much longer average lifetime ( $\tau_{\text{avg}} = 259.4$  ps) compared to that of reference  $\text{CsSnI}_3$  NCs film ( $\tau_{\text{avg}} = 10.1$  ps), which is highly consistent with the trend of TRPL data in the Supporting Information, Table S2. This further suggests that, compared to reference  $\text{CsSnI}_3$  NCs, the  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs possess the extended excited-state lifetime that allows efficient charge transfer, which is beneficial for the device performance.

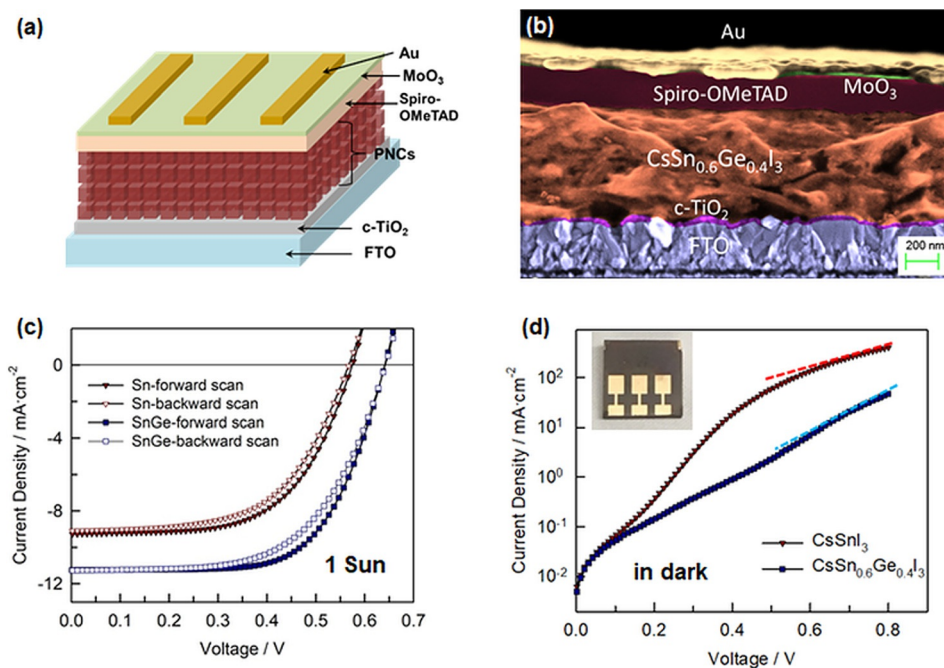
### Enhanced Performance of $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$ NCs-based Solar Cells

The improved photophysical properties of  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs compared to the case of reference  $\text{CsSnI}_3$  NCs has a direct effect on the performance of corresponding PSCs. To confirm this, we incorporated both types of nanocubes in standard *n-i-p* photovoltaic architectures. Owing to the reported high charge diffusion coefficient ( $> 1.3 \text{ cm}^2 \text{ s}^{-1}$ ) for typical bulk  $\text{CsSnI}_3$  crystals, a planar structure has been adopted for this type of perovskite.<sup>[45,46]</sup> The full PSC structure is FTO/c-TiO<sub>2</sub>/PNCs/spiro-OMeTAD/MoO<sub>3</sub>/Au, as illustrated in Figure 4a. The devices consisted of circa 550 nm thick PNCs layer (based on the cross-sectional SEM image of a typical  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs-based solar cell, Figure 4b), fabricated in a nitrogen-filled glovebox. The current density (*J*)-voltage (*V*) curves of the best-performing (champion) devices based on SnGe-PNCs and reference Sn-PNCs, recorded under standard 1 Sun condition ( $100 \text{ mW cm}^{-2}$  AM 1.5 G illumination) and in dark condition, are depicted in Figure 4c,d, respectively. Interestingly, the *J-V* curves of both PNCs-based devices display a negligible hysteresis effect between forward and backward scans (Figure 4c). This suggests that ion migration under an electric field has a nearly negligible influence on the photocurrent flow within the perovskite nanocrystals layer. This, in turn, may lead to a more stable power output of the corresponding PNCs devices compared to that of bulk crystals-based devices, such as the popular tri-cation perovskite solar cells in *n-i-p* structure.<sup>[2]</sup> The averaged photovoltaic performances (16 devices for each structure) along with the standard deviation, together with the photovoltaic parameters of the champion cells, are summarized in Table 1. The high reproducibility of our results is demonstrated by the small standard deviations. The  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs lead to PSCs with a remarkable 58 % performance enhancement (average PCE = 4.1 %) compared to devices based on reference  $\text{CsSnI}_3$  NCs (Table 1) in this work, resulting in the highest PCE of 4.9 % for the champion cell (shown in the inset photo in Figure 4d).

**Table 1:** Summary of PNCs based solar cells performance.<sup>[a]</sup>

PNCs	$J_{\text{sc}}$ [mA cm <sup>-2</sup> ]	$V_{\text{oc}}$ [V]	FF	PCE [%]	$R_{\text{s}}$ [Ω cm <sup>2</sup> ]	$R_{\text{sh}}$ [kΩ cm <sup>2</sup> ]
Sn-PNCs	8.6 ± 0.8 (9.4) <sup>[b]</sup>	0.54 ± 0.04 (0.58)	0.54 ± 0.03 (0.57)	2.7 ± 0.4 (3.1)	23.3 ± 1.5 (21.8)	3.5 ± 0.3 (3.8)
SnGe-PNCs	10.9 ± 0.9 (11.8)	0.61 ± 0.03 (0.65)	0.60 ± 0.03 (0.64)	4.1 ± 0.8 (4.9)	13.4 ± 1.1 (12.3)	6.3 ± 0.4 (6.7)

[a] The data are from the forward scans. [b] Values in the brackets refer to the photovoltaic parameters of the champion cells.



**Figure 4.** a) Architecture of *n-i-p* PNCs based solar cells in planar structure studied in this work. b) Cross-sectional SEM image of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs-based solar cell, observed at 3 kV. c), d) *J-V* curves of the champion Sn-PNCs and SnGe-PNCs based solar cells (c) under 1 sun illumination (forward and backward scans with a scan rate of 10 mV s<sup>-1</sup>) and (d) in dark condition (forward scan) while dash line shows the fitting to extract series resistance. The inset picture shows the appearance of a typical CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs-based solar cell.

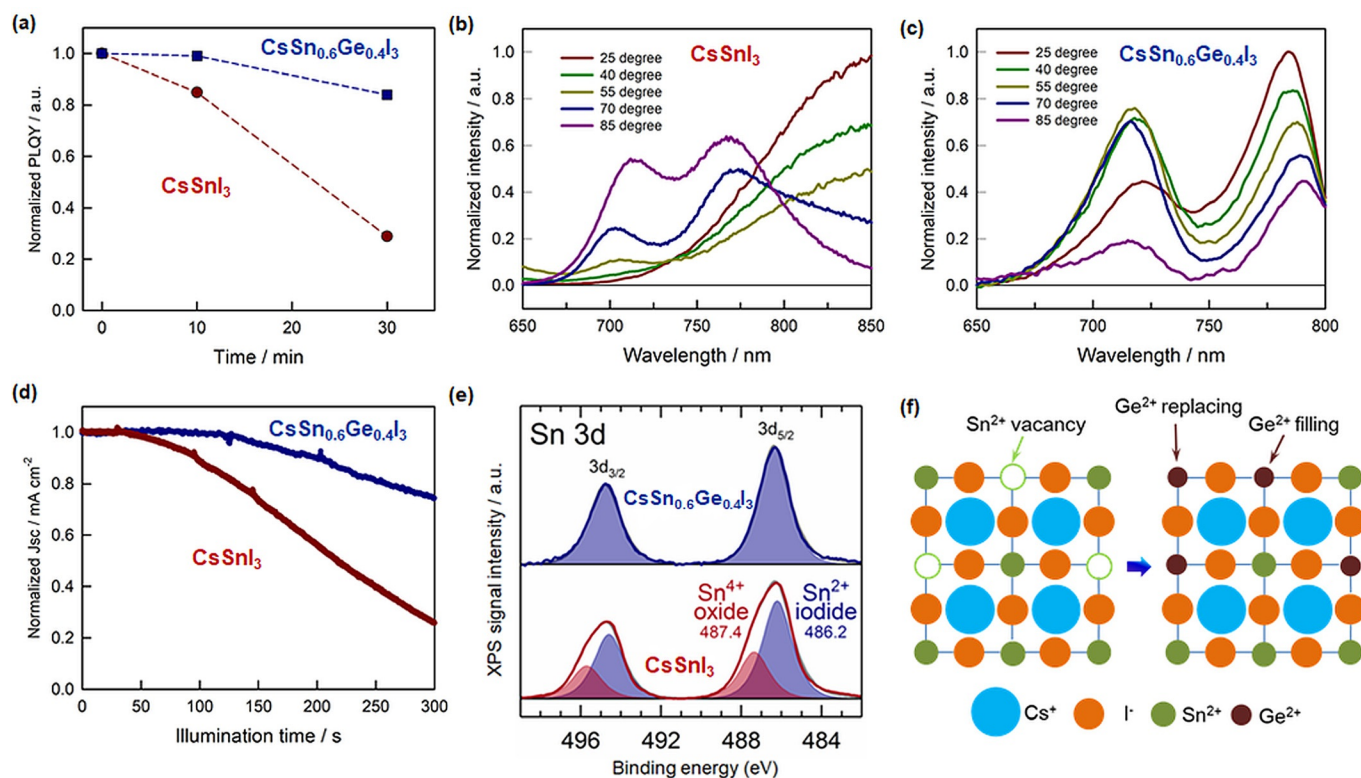
The improvement in the PCE is clearly determined by the increase in all device parameters, that is,  $J_{SC}$ ,  $V_{OC}$ , and FF. We have extracted the series ( $R_S$ ) and shunt ( $R_{SH}$ ) resistances of the devices, according to the 1-diode model,<sup>[47]</sup> from the *J-V* curves scanned in dark with the axis of current density plotted in logarithmic scale (Figure 4c and Table 1). Both  $R_{SH}$  and  $R_S$  affect the FF, with lower  $R_S$  and higher  $R_{SH}$  being required to increase the FF. The extracted resistances in Table 1 confirm this trend since CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs-based cells have lower  $R_S$  and higher  $R_{SH}$  (nearly twice) than that of the CsSnI<sub>3</sub> NCs-based cells. In addition, the excited-state lifetime (ca. 4 ns) of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs clarified from the TCSPC and TA measurements, which is nearly three times longer than that (ca. 1.5 ns) of CsSnI<sub>3</sub> NCs, in turn leads to a more effective charge separation at the interface between PNCs film and charge-transporting layers (TiO<sub>2</sub> or spiro-OMeTAD).<sup>[48]</sup> This can be one key reason for the enhanced  $J_{SC}$  and FF in CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs-based cells. The above-demonstrated involvement of Ge in reducing the Sn vacancies (that function as trapping sites) suggests that CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs-based active layer possesses much fewer defect sites compared to the CsSnI<sub>3</sub> NCs film, leading to the hindering of charge recombination within PNCs film beneficial to the  $V_{OC}$ .<sup>[49,50]</sup> Furthermore, we investigated the surface morphology of PNCs films by taking SEM topographic imaging for CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs (in Figure S7a) and CsSnI<sub>3</sub> NCs (Supporting Information, Figure S7b). It is noted that the surface of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs film is much smoother and denser (see inset image in the Supporting Information, Figure S7a) than that of CsSnI<sub>3</sub> film. Probably, upon the involvement of Ge, Sn phase aggregation

can be avoided, supporting the possibility of suppressed interfacial charge recombination.

#### Improved Stability of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs

Finally, the effect of Ge on the stability of the corresponding PNCs is thoroughly studied. Figure 5a shows the comparison between the time-dependent PLQYs of CsSnI<sub>3</sub> NCs and CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs in suspension in air (25 °C and 50 % RH). While the PLQY of CsSnI<sub>3</sub> NCs swiftly decreased to below 30 % of its initial value after 30 min (Supporting Information, Figure S8a), that of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs was retained to above 80 % of the original PLQY in the same time span (Supporting Information, Figure S8b). This suggests that upon the protection of Ge atoms in the Sn-based perovskite against the oxidation, the optical properties of PNCs can be effectively maintained. Figure S9 presents the comparison of XRD patterns from unencapsulated reference CsSnI<sub>3</sub> NCs (Supporting Information, Figure S9a) and CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs (Supporting Information, Figure S9b) films exposed for 10, 20, and 30 min to ambient atmosphere. The relative intensity of the main XRD peak can sustain above 75 % of its initial value for the case of SnGe-PNCs after 30 min exposure while that of Sn-PNCs has dropped below 45 % of its corresponding initial intensity, consistent with the enhancement of optical stability upon the introduction of Ge. Moreover, we measured the temperature-dependent PL spectrum (Figure 5b,c) for both PNCs under ambient conditions to clarify the thermal stability influenced by the involvement of Ge in the perovskite structure. Interestingly,





**Figure 5.** a) Time-dependent PLQYs of  $\text{CsSnI}_3$  NCs and  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs in suspension under ambient condition. Temperature dependent PL spectrum of b)  $\text{CsSnI}_3$  NCs and c)  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs in suspension under ambient condition. d)  $J_{\text{SC}}$  evolution of typical unencapsulated  $\text{CsSnI}_3$  NCs and  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs-based PSCs in continuous operation under 1 sun illumination in air. The sampling time is 100 ms. e) XPS analysis of Sn 3d transition for  $\text{CsSnI}_3$  NCs and  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs after 10 min exposure to air atmosphere. f) Scheme of the Ge filling effect in the Sn vacancies and the replacement of Sn atoms by Ge atoms in the nanocrystal structure.

upon temperatures above 55 °C, the PL spectrum of  $\text{CsSnI}_3$  NCs exhibited a clear decrease in the PL intensity with a distinct blue-shift and an additional sub-peak at about 710 nm, indicating a sign of the phase transition or degradation, that is, the formation of  $\text{Sn}^{4+}$ . On the contrary, the PL spectrum of  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs retained its shape feature when temperature increased, although the peak PL intensity at 782 nm (corresponding to nanocubes) decreased, as commonly observed for other types of perovskite nanocrystals.<sup>[51,52]</sup> It is interestingly noted that the sub-peak PL intensity at 725 nm (corresponding to nanorods) effectively increased up to 70 °C and then decreased with the increasing temperature (85 °C), indicating that some temperature-dependent morphological transition from nanocubes to nanorods may occur which will be investigated in detail in a separate work. The overall study of PL stability suggests that Ge atoms can effectively stabilize the nanocrystal structures upon the thermal treatment. We then studied the stability of the corresponding photovoltaic devices by measuring the change in the short circuit current density ( $J_{\text{SC}}$ ) under continuous 1-sun illumination in air. Figure 5d compares the decrease in  $J_{\text{SC}}$  upon unencapsulated cells illumination for both  $\text{CsSnI}_3$  NCs- and  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs-based devices. As predicted,  $J_{\text{SC}}$  of pristine Sn-PNCs cells decreased by more than 75 % of its initial value after 300 s of continuous illumination. On the other hand, SnGe-PNCs based devices can still retain more than 80 % of initial  $J_{\text{SC}}$ , which is

consistent with the variation trend of the time-dependent PLQYs of both PNCs. Hence,  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs exhibit significantly improved ambient, thermal, and photostability compared to  $\text{CsSnI}_3$  NCs.

To investigate the underlying reason for the improved stability of  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs, we can further refer to the XPS analysis in Figure 5e. Clearly, only one chemical state containing Sn could be extracted for  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs (Sn 3d<sub>5/2</sub> at 486.2 eV) after 10 mins exposure to air, that is, during the sample preparation before the XPS measurement. In contrast, within the same time frame, a second chemical state was clearly observed for  $\text{CsSnI}_3$  NCs related to Sn 3d transition: a second pair of doublet peaks (Sn 3d<sub>5/2</sub> at 487.4 eV) is found in addition to the  $\text{Sn}^{2+}$  iodide 3d<sub>5/2</sub> at 486.2 eV. The high binding energy component corresponds to  $\text{Sn}^{4+}$  oxide.<sup>[53–55]</sup> This was also confirmed by the O 1s metal oxide component at 530.7 eV, which has been observed only for the case of pure Sn-PNCs. Furthermore, we have conducted the XPS analysis on  $\text{CsSn}_{0.6}\text{Ge}_{0.4}\text{I}_3$  NCs after 24 hours of exposure to air (Supporting Information, Figure S10). Upon the long-time exposure to air, Sn 3d/4d peaks are shifted to higher binding energies, indicating the oxidation of  $\text{Sn}^{2+}$  cations into  $\text{Sn}^{4+}$  state. Yet, the Ge 3d peak has shown no obvious change with the fixation of most  $\text{Ge}^{2+}$  cations in the perovskite structure. Interestingly, the fast formation of  $\text{Ge}^{4+}$  oxide that was recently observed by Chen et al. on bulk perovskite with a similar elemental composition (bulk

CsSn<sub>0.5</sub>Ge<sub>0.5</sub>I<sub>3</sub>) but prepared using a melt-crystallization method has not been observed in our case.<sup>[29]</sup> Furthermore, the enhanced stability of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs is in agreement with two factors demonstrated in the previous simulation results: 1) the Goldschmidt tolerance factor (*t*) has been reported for the mixed Sn-Ge halide perovskite in comparison to Ge-free counterpart,<sup>[56]</sup> that is, *t* (MASn<sub>0.5</sub>Ge<sub>0.5</sub>I<sub>3</sub>) = 0.90, and *t* (MASnI<sub>3</sub>) = 0.84, indicating that the involvement of Ge atoms in Sn-based perovskite structure is beneficial for the structural stability; 2) the activation barrier (0.23 eV) towards water or oxygen penetration of mixed Sn-Ge halide perovskite (RbSn<sub>0.5</sub>Ge<sub>0.5</sub>I<sub>3</sub>) has been reported as remarkably higher than that of pure Sn perovskite (0.17 eV for MASnI<sub>3</sub>)<sup>[56]</sup> or even of Pb-based perovskite (0.09 eV for MAPbI<sub>3</sub>)<sup>[27]</sup> suggesting that the terminated Ge atoms can effectively protect the inner Sn atoms, making them less susceptible to oxidization upon the replacement and filling effect from Ge atoms (see the proposal in Figure 5 f).

## Conclusion

We have shown the first-ever synthesis of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> nanocrystals with a narrow size distribution. The PLQY of Sn-Ge based NCs is close to 1 %, which has enhanced nearly one order magnitude than that of conventional pure Sn-based PNCs. The photophysical properties of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs are dramatically enhanced compared to those of pure Sn-PNCs, mainly attributed to the prolonged excited-state lifetime, as confirmed by both transient photoluminescence and transient absorption measurements. We assign the improvement of both optical and photophysical properties of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs to the reduced number of Sn vacancies upon their filling with Ge in the nanostructure. Vacancies act as trapping sites and alternative nonradiative recombination pathways. The overall improved charge transfer dynamics for CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs system, that is, efficient charge separation and suppressed charge recombination, is the key reason for the significant enhancement of solar cells performance when CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs are used as the light absorber instead of CsSnI<sub>3</sub> NCs. The highest PCE of CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs-based solar cells in this work, 4.9 %, is one of the highest values reported for all-inorganic lead-free PNCs solar cells. The introduction of Ge is highly beneficial for ambient, thermal, and photostability, due to the effective protection of Ge<sup>2+</sup> against Sn<sup>2+</sup> oxidization. We believe that, upon optimization (for example, by identifying the most suitable selective contacts matching the CsSn<sub>0.6</sub>Ge<sub>0.4</sub>I<sub>3</sub> NCs energy levels), the solar cells based on SnGe-PNCs can outperform current generation of lead-free PNCs based solar cells with the additional benefit of outstanding stability, hence bringing PNCs-photovoltaics a step closer to their commercialization.

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## Conflict of interest

The authors declare no conflict of interest.

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- [1] S. Khalifin, Y. Bekenstein, *Nanoscale* **2019**, *11*, 8665–8679.
- [2] R. Yang, R. Li, Y. Cao, Y. Wei, Y. Miao, W. L. Tan, X. Jiao, H. Chen, L. Zhang, Q. Chen, H. Zhang, W. Zou, Y. Wang, M. Yang, C. Yi, N. Wang, F. Gao, C. R. McNeill, T. Qin, J. Wang, W. Huang, *Adv. Mater.* **2018**, *30*, 1804771.
- [3] N. J. Jeon, H. Na, E. H. Jung, T.-Y. Yang, Y. G. Lee, G. Kim, H.-W. Shin, S. Il Seok, J. Lee, J. Seo, *Nat. Energy* **2018**, *3*, 682–689.
- [4] D. Nguyen Minh, J. Kim, J. Hyon, J. Hyun Sim, H. H. Sowli, C. Seo, J. Nam, S. Eom, S. Suk, S. Lee, E. Kim, Y. Kang, *Chem. Mater.* **2017**, *29*, 5713–5719.
- [5] J. Song, J. Li, X. Li, L. Xu, Y. Dong, H. Zeng, *Adv. Mater.* **2015**, *27*, 7162–7167.
- [6] Z. Liu, J. Hu, H. Jiao, L. Li, G. Zheng, Y. Chen, Y. Huang, Q. Zhang, C. Shen, Q. Chen, H. Zhou, *Adv. Mater.* **2017**, *29*, 1606774.
- [7] H. Cho, C. Wolf, J. S. Kim, H. J. Yun, J. S. Bae, H. Kim, J.-M. Heo, S. Ahn, T.-W. Lee, *Adv. Mater.* **2017**, *29*, 1700579.
- [8] Z. Ning, X. Gong, R. Comin, G. Walters, F. Fan, O. Voznyy, E. Yassitepe, A. Buin, S. Hoogland, E. H. Sargent, *Nature* **2015**, *523*, 324–328.
- [9] M. Gong, R. Sakidja, R. Goul, D. Ewing, M. Casper, A. Stramel, A. Elliot, J. Z. Wu, *ACS Nano* **2019**, *13*, 1772–1783.
- [10] W. Ke, C. C. Stoumpos, M. G. Kanatzidis, *Adv. Mater.* **2019**, *31*, 1803230.
- [11] Y. Yan, T. Pullerits, K. Zheng, Z. Liang, *ACS Energy Lett.* **2020**, *5*, 2052–2086.
- [12] X. Jiang, F. Wang, Q. Wei, H. Li, Y. Shang, W. Zhou, C. Wang, P. Cheng, Q. Chen, L. Chen, Z. Ning, *Nat. Commun.* **2020**, *11*, 1245.
- [13] Y. Wang, J. Tu, T. Li, C. Tao, X. Deng, Z. Li, *J. Mater. Chem. A* **2019**, *7*, 7683–7690.
- [14] F. Hao, C. C. Stoumpos, D. H. Cao, R. P. H. Chang, M. G. Kanatzidis, *Nat. Photonics* **2014**, *8*, 489–494.
- [15] F. Hao, H. Tan, Z. Jin, L. Ding, *Sci. Bull.* **2020**, *65*, 786–790.
- [16] E. W. G. Diau, E. Jokar, M. Rameez, *ACS Energy Lett.* **2019**, *4*, 1930–1937.
- [17] T.-B. Song, T. Yokoyama, S. Aramaki, M. G. Kanatzidis, *ACS Energy Lett.* **2017**, *2*, 897–903.
- [18] T. Krishnamoorthy, H. Ding, C. Yan, W. L. Leong, T. Baikie, Z. Zhang, M. Sherburne, S. Li, M. Asta, N. Mathews, S. G. Mhaisalkar, *J. Mater. Chem. A* **2015**, *3*, 23829–23832.
- [19] C. C. Stoumpos, L. Frazer, D. J. Clark, Y. S. Kim, S. H. Rhim, A. J. Freeman, J. B. Ketterson, J. I. Jang, M. G. Kanatzidis, *J. Am. Chem. Soc.* **2015**, *137*, 6804–6819.
- [20] Y. Dong, Y. Zhao, S. Zhang, Y. Dai, L. Liu, Y. Li, Q. Chen, *J. Mater. Chem. A* **2018**, *6*, 21729–21746.
- [21] T. Minemoto, Y. Kawano, T. Nishimura, Q. Shen, K. Yoshino, S. Iikubo, S. Hayase, J. Chantana, *Sol. Energy Mater. Sol. Cells* **2020**, *206*, 110268.



- [22] S. Zou, Y. Liu, J. Li, C. Liu, R. Feng, F. Jiang, Y. Li, J. Song, H. Zeng, M. Hong, X. Chen, *J. Am. Chem. Soc.* **2017**, *139*, 11443–11450.
- [23] F. Qian, M. Hu, J. Gong, C. Ge, Y. Zhou, J. Guo, M. Chen, Z. Ge, N. P. Padture, Y. Zhou, J. Feng, *J. Phys. Chem. C* **2020**, *124*, 11749–11753.
- [24] C. H. Ng, K. Nishimura, N. Ito, K. Hamada, D. Hirotsu, Z. Wang, F. Yang, S. Iikubo, Q. Shen, K. Yoshino, T. Minemoto, S. Hayase, *Nano Energy* **2019**, *58*, 130–137.
- [25] X.-X. Ma, Z.-S. Li, *Phys. Status Solidi* **2019**, *256*, 1800427.
- [26] R. Ali, G. J. Hou, Z. G. Zhu, Q. B. Yan, Q. R. Zheng, G. Su, *J. Mater. Chem. A* **2018**, *6*, 9220–9227.
- [27] M.-G. Ju, J. Dai, L. Ma, X. C. Zeng, *J. Am. Chem. Soc.* **2017**, *139*, 8038–8043.
- [28] N. Ito, M. A. Kamarudin, D. Hirotsu, Y. Zhang, Q. Shen, Y. Ogomi, S. Iikubo, T. Minemoto, K. Yoshino, S. Hayase, *J. Phys. Chem. Lett.* **2018**, *9*, 1682–1688.
- [29] M. Chen, M.-G. Ju, H. F. Garces, A. D. Carl, L. K. Ono, Z. Hawash, Y. Zhang, T. Shen, Y. Qi, R. L. Grimm, D. Pacifici, X. C. Zeng, Y. Zhou, N. P. Padture, *Nat. Commun.* **2019**, *10*, 16.
- [30] D. S. Dolzhenkov, C. Wang, Y. Xu, M. G. Kanatzidis, E. A. Weiss, *Chem. Mater.* **2017**, *29*, 7901–7907.
- [31] T. C. Jellicoe, J. M. Richter, H. F. J. Glass, M. Tabachnyk, R. Brady, S. E. Dutton, A. Rao, R. H. Friend, D. Credgington, N. C. Greenham, M. L. Böhm, *J. Am. Chem. Soc.* **2016**, *138*, 2941–2944.
- [32] I. Lignos, S. Stavrakis, G. Nedelcu, L. Protesescu, A. J. Demello, M. V. Kovalenko, *Nano Lett.* **2016**, *16*, 1869–1877.
- [33] Y. Zhou, H. F. Garces, B. S. Senturk, A. L. Ortiz, N. P. Padture, *Mater. Lett.* **2013**, *110*, 127–129.
- [34] Z. Dang, J. Shamsi, Q. A. Akkerman, M. Imran, G. Bertoni, R. Brescia, L. Manna, *ACS Omega* **2017**, *2*, 5660–5665.
- [35] C. J. P. Alexander, V. Naumkin, A. Kraut-Vass, S. W. Gaarenstroom, *NIST X-ray Photoelectron Spectrosc. Database, NIST Stand. Ref. Database 20, Version 4.1* **2000**, Accessed: 2018-2002-08.
- [36] X. He, Y. Qiu, S. Yang, *Adv. Mater.* **2017**, *29*, 1700775.
- [37] M. Koolyk, D. Amgar, S. Aharon, L. Etgar, *Nanoscale* **2016**, *8*, 6403–6409.
- [38] Y. Zhang, J. Yin, M. R. Parida, G. H. Ahmed, J. Pan, O. M. Bakr, J.-L. Brédas, O. F. Mohammed, *J. Phys. Chem. Lett.* **2017**, *8*, 3173–3177.
- [39] Y. Zhang, J. Liu, Z. Wang, Y. Xue, Q. Ou, L. Polavarapu, J. Zheng, X. Qi, Q. Bao, *Chem. Commun.* **2016**, *52*, 13637–13655.
- [40] A. B. Wong, M. Lai, S. W. Eaton, Y. Yu, E. Lin, L. Dou, A. Fu, P. Yang, *Nano Lett.* **2015**, *15*, 5519–5524.
- [41] J. Shamsi, A. S. Urban, M. Imran, L. De Trizio, L. Manna, *Chem. Rev.* **2019**, *119*, 3296–3348.
- [42] R. Koole, B. Luigjes, M. Tachiya, R. Pool, T. J. H. Vlugt, C. de Mello Donegá, A. Meijerink, D. Vanmaekelbergh, *J. Phys. Chem. C* **2007**, *111*, 11208–11215.
- [43] N. V. Tkachenko, *ChemPhotoChem* **2018**, *2*, 112–120.
- [44] F. Liu, C. Ding, Y. Zhang, T. S. Ripolles, T. Kamisaka, T. Toyoda, S. Hayase, T. Minemoto, K. Yoshino, S. Dai, M. Yanagida, H. Noguchi, Q. Shen, *J. Am. Chem. Soc.* **2017**, *139*, 16708–16719.
- [45] J. Zhang, D. Bai, Z. Jin, H. Bian, K. Wang, J. Sun, Q. Wang, S. F. Liu, *Adv. Energy Mater.* **2018**, *8*, 1870067.
- [46] M. Liu, M. Endo, A. Shimazaki, A. Wakamiya, Y. Tachibana, *ACS Appl. Energy Mater.* **2018**, *1*, 3722–3732.
- [47] S. Yoo, B. Domercq, B. Kippelen, *J. Appl. Phys.* **2005**, *97*, 103706.
- [48] S. Makuta, M. Liu, M. Endo, H. Nishimura, A. Wakamiya, Y. Tachibana, *Chem. Commun.* **2016**, *52*, 673–676.
- [49] W. Tress, M. Yavari, K. Domanski, P. Yadav, B. Niesen, J. P. Correa Baena, A. Hagfeldt, M. Graetzel, *Energy Environ. Sci.* **2018**, *11*, 151–165.
- [50] M. Liu, M. Endo, A. Shimazaki, A. Wakamiya, Y. Tachibana, *J. Photopolym. Sci. Technol.* **2018**, *31*, 633–642.
- [51] S. B. Naghadeh, S. Sarang, A. Brewer, A. Allen, Y. H. Chiu, Y. J. Hsu, J. Y. Wu, S. Ghosh, J. Z. Zhang, *J. Chem. Phys.* **2019**, *151*, 154705.
- [52] P. Liang, P. Zhang, A. Pan, K. Yan, Y. Zhu, M. Yang, L. He, *ACS Appl. Mater. Interfaces* **2019**, *11*, 22786–22793.
- [53] M. A. Kamarudin, D. Hirotsu, Z. Wang, K. Hamada, K. Nishimura, Q. Shen, T. Toyoda, S. Iikubo, T. Minemoto, K. Yoshino, S. Hayase, *J. Phys. Chem. Lett.* **2019**, *10*, 5277–5283.
- [54] T. M. Koh, T. Krishnamoorthy, N. Yantara, C. Shi, W. L. Leong, P. P. Boix, A. C. Grimsdale, S. G. Matheskar, N. Mathews, *J. Mater. Chem. A* **2015**, *3*, 14996–15000.
- [55] S. J. Lee, S. S. Shin, Y. C. Kim, D. Kim, T. K. Ahn, J. H. Noh, J. Seo, S. Il Seok, *J. Am. Chem. Soc.* **2016**, *138*, 3974–3977.
- [56] S. Nagane, D. Ghosh, R. L. Z. Hoyer, B. Zhao, S. Ahmad, A. B. Walker, M. S. Islam, S. Ogale, A. Sadhanala, *J. Phys. Chem. C* **2018**, *122*, 5940–5947.

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