

## RESEARCH ARTICLE

View Article Online  
View Journal | View IssueCite this: *Mater. Chem. Front.*,  
2018, 2, 1515Cs<sub>1-x</sub>Rb<sub>x</sub>SnI<sub>3</sub> light harvesting semiconductors  
for perovskite photovoltaics†Kenneth P. Marshall,<sup>a</sup> Shuxia Tao,<sup>b</sup> Marc Walker,<sup>c</sup> Daniel S. Cook,<sup>a</sup>  
James Lloyd-Hughes,<sup>b</sup> Silvia Varagnolo,<sup>a</sup> Anjana Wijesekera,<sup>a</sup>  
David Walker,<sup>b</sup> Richard I. Walton<sup>a</sup> and Ross A. Hatton<sup>\*a</sup>

We show that films of the 3-dimensional perovskite Cs<sub>1-x</sub>Rb<sub>x</sub>SnI<sub>3</sub> can be prepared from room temperature *N,N*-dimethylformamide solutions of RbI, CsI and SnCl<sub>2</sub> for  $x \leq 0.5$ , and that for  $x \leq 0.2$  film stability is sufficient for utility as the light harvesting layer in inverted photovoltaic (PV) devices. Electronic absorption and photoluminescence spectroscopy measurements supported by computational simulation, show that increasing  $x$  increases the band gap, due to distortion of the lattice of SnI<sub>6</sub> octahedra that occurs when Cs is substituted with Rb, although it also reduces the stability towards decomposition. When Cs<sub>0.8</sub>Rb<sub>0.2</sub>SnI<sub>3</sub> perovskite is incorporated into the model inverted PV device structure; ITO|perovskite|C<sub>60</sub>|bathocuproine|Al, an  $\sim 120$  mV increase in open-circuit is achieved which is shown to correlate with an increase in perovskite ionisation potential. However, for this low Rb loading the increase in band gap is very small ( $\sim 30$  meV) and so a significant increase in open circuit-voltage is achieved without reducing the range of wavelengths over which the perovskite can harvest light. The experimental findings presented are shown to agree well with the predictions of density functional theory (DFT) simulations of the stability and electronic structure, also performed as part of this study.

Received 9th April 2018,  
Accepted 4th May 2018

DOI: 10.1039/c8qm00159f

rsc.li/frontiers-materials

## Introduction

The black gamma (B- $\gamma$ ) polymorph of CsSnI<sub>3</sub> perovskite is a promising material for photovoltaic (PV) applications because it offers an ideal band gap for single junction PV devices ( $\sim 1.34$  eV<sup>1</sup>) together with a high electron and hole mobility (1.6 and 585 cm<sup>2</sup> V<sup>-1</sup> s<sup>-1</sup>, respectively)<sup>1,2</sup> and small exciton binding energy.<sup>3,4</sup> This wholly inorganic perovskite comprises a 3-dimensional structure of corner sharing SnI<sub>6</sub> octahedra with Cs cations surrounded by 8 SnI<sub>6</sub> octahedra, and adopts the orthorhombic B- $\gamma$  phase below 89 °C.<sup>2</sup> To date it has been used as the light harvesting semiconductor in PV devices based on mesoporous TiO<sub>2</sub><sup>5,6</sup> and planar<sup>7-9</sup> device architectures, demonstrating the potential to achieve very high short circuit current density ( $J_{sc}$ ) of  $> 20$  mA cm<sup>-2</sup> under one sun illumination<sup>6</sup> and high device fill factor (FF).<sup>8</sup> The primary factor limiting the power conversion efficiency of B- $\gamma$  CsSnI<sub>3</sub> based PVs is the low open-circuit voltage ( $V_{oc}$ ): to date the highest  $V_{oc}$  reported is 0.55 V,<sup>8</sup> achieved for an inverted device

architecture using phenyl-C<sub>61</sub>-butyric acid methylester (PCBM) as the electron transport layer (ETL), which is approximately half that attainable using the lead analogue CsPbI<sub>3</sub>.<sup>10</sup> When using C<sub>60</sub> in place of PCBM the highest  $V_{oc}$  reported is reduced to 0.36 V due to the larger electron affinity of the former.<sup>9</sup>

The  $V_{oc}$  in inverted perovskite PV devices depends strongly on a number of factors including the energetics at the perovskite-ETL interface<sup>7</sup> and the degree of crystallinity in the ETL.<sup>11</sup> Additionally, for PV devices using B- $\gamma$  CsSnI<sub>3</sub> as the light harvesting layer, we have shown that SnCl<sub>2</sub> is an effective additive for reducing the density of tin vacancy defects in the band gap and reducing the reverse saturation current,<sup>8</sup> both of which help to improve  $V_{oc}$ . However,  $V_{oc}$  is ultimately limited by the relatively small ionisation potential ( $I_p$ ) of B- $\gamma$  CsSnI<sub>3</sub> of  $\sim 4.9$  eV,<sup>9,12</sup> which is approximately half an electron volt smaller than that of lead halide perovskites.<sup>13</sup> Fortunately, similar to lead halide perovskites, B- $\gamma$  CsSnI<sub>3</sub> is amenable to substitution of iodide ions with bromide ions, which results in an increase in  $I_p$  and band gap which translates to an increase in  $V_{oc}$  in PV devices.<sup>14</sup> However, this benefit must be balanced against the inevitable reduction in short circuit current density ( $J_{sc}$ ) resulting from fewer photons having sufficient energy to excite electrons across the larger band gap. An alternative strategy for tuning the electronic structure of B- $\gamma$  CsSnI<sub>3</sub> is substitution of A-site Cs cation with the smaller Rb cation to form Cs<sub>1-x</sub>Rb<sub>x</sub>SnI<sub>3</sub>. Unlike the case of halide ion substitution, the orbitals of the A-site metal cation do not

<sup>a</sup> Department of Chemistry, University of Warwick, CV4 7AL, Coventry, UK.  
E-mail: Ross.Hatton@warwick.ac.uk

<sup>b</sup> Center for Computational Energy Research, Department of Applied Physics,  
Technische Universiteit Eindhoven, P.O. Box 513 5600 MB, Eindhoven,  
The Netherlands

<sup>c</sup> Department of Physics, University of Warwick, CV4 7AL, Coventry, UK

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8qm00159f



contribute directly to the conduction and valence band edges,<sup>4,15</sup> but still indirectly affect the band gap as a result of the distortion of the lattice of  $\text{SnI}_6$  octahedra *via* tilting that occurs when reducing the size of the A-site ion.<sup>16,17</sup>  $\text{RbSnI}_3$  has a tolerance factor of 0.840 which is very close to the value that allows for the likely formation of a 3D structure; 0.85,<sup>18</sup> which is consistent with the fact that it has only been reported in the one-dimensional yellow phase.<sup>19</sup>

Herein we report the facile room temperature preparation of thin films of 3-dimensional (3D)  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  and show how partial exchange of Cs with Rb can be used to increase  $I_p$  and band gap. Notably, for low levels of substitution it is possible to achieve a significant increase in  $I_p$  with only a very small increase in band gap which, in model inverted PV devices, results in a sizable increase in  $V_{oc}$  with no significant adverse impact on the light harvesting capability. The experimental findings presented are shown to agree well with the predictions of density functional theory (DFT) simulations of the stability and electronic structure, also performed as part of this study.

## Results and discussion

The tin halides  $\text{SnI}_2$ ,  $\text{SnCl}_2$  and  $\text{SnF}_2$  are typically used as a source of excess Sn during the preparation of B- $\gamma$   $\text{CsSnI}_3$  films to minimise the density of tin vacancy defects.<sup>5,7,8,12</sup> In the first part of this study  $\text{SnI}_2$  was used as the source of excess Sn to avoid the possibility that Cl or F might be incorporated into the lattice of  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$ , which would complicate the interpretation. Films of  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  were prepared with Rb content  $x = 0, 0.2, 0.5, 0.8$  and 1, with a thickness of  $\sim 50$  nm on ITO glass and  $\sim 80$  nm on plain glass. All films were deposited under nitrogen by spin casting from solutions of *N,N*-dimethylformamide (DMF) at room temperature. Films prepared with  $x = 0, 0.2$  and 0.5 (Fig. S1, ESI<sup>†</sup>) were an intense dark colour typical of 3D tin halide perovskite, whilst films with  $x = 0.8$  and 1 were orange/yellow in colour indicating that they did not form 3D perovskite phases.

Fig. 1 shows the X-ray diffraction (XRD) patterns for  $\sim 80$  nm thick  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  films supported on glass at room temperature, along with simulated patterns for B- $\gamma$   $\text{CsSnI}_3$  and  $\text{RbSnI}_3$ . It is evident from these patterns that for  $x = 0$  and 0.2 the crystal structure is similar to that of B- $\gamma$   $\text{CsSnI}_3$ . For  $x = 0.5$  the films were very unstable, with the colour evolving from a deep red to yellow in the few minutes taken to load the sample into the X-ray diffractometer.

Lattice parameter fitting; ESI<sup>†</sup> (Fig. S2), shows there is no significant or systematic change in lattice parameters between B- $\gamma$   $\text{CsSnI}_3$ ,  $\text{Cs}_{0.9}\text{Rb}_{0.1}\text{SnI}_3$  and  $\text{Cs}_{0.8}\text{Rb}_{0.2}\text{SnI}_3$ , entirely consistent with the computational simulations of Jung *et al.*<sup>16</sup> and DFT simulations performed as part of this study (ESI<sup>†</sup>, Fig. S3 and S4), both of which predict only a small change in lattice parameters between the 3D perovskite B- $\gamma$   $\text{CsSnI}_3$  and the hypothetical 3D  $\text{RbSnI}_3$ , due to a much higher degree of tilting between the corner sharing  $\text{SnI}_6$  octahedra in the latter. Simulation of the X-ray diffraction pattern based on a crystallographic information file of  $\text{CsSnI}_3$  in which 20% of Cs atoms have been replaced with Rb

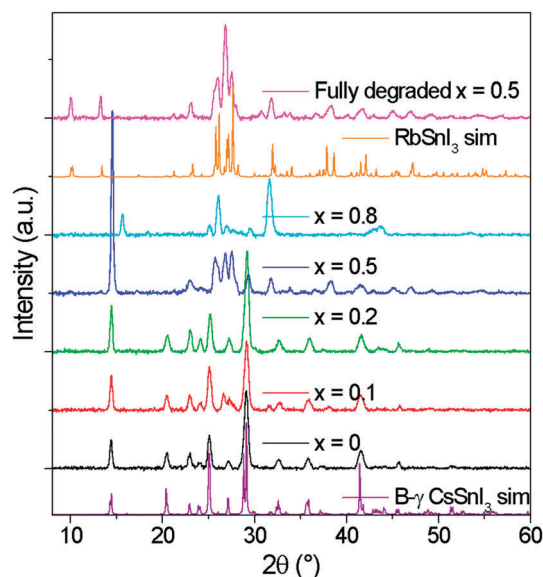


Fig. 1 XRD patterns of  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  where  $x = 0, 0.1, 0.2, 0.5, 0.8$  with simulated patterns of B- $\gamma$   $\text{CsSnI}_3$ . All films were deposited from DMF solutions of RbI, CsI and  $\text{SnI}_2$  at room temperature.

atoms shows that there should be some difference in the peak intensities (Fig. 2): the simulation predicts that the (101) and (020) reflections should have increased intensity relative to the (202) and (040) reflections when Cs is partially substituted with Rb. Comparing the measured patterns of  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  there is no significant trend in the relative ratios of the (101) + (020):(202) + (040) intensities, although both samples incorporating Rb have higher relative intensities of (101) and (020) compared with (202). Additionally, before the sample degraded the  $\text{Cs}_{0.5}\text{Rb}_{0.5}\text{SnI}_3$  sample has a very intense peak at  $14.5^\circ$  which is assigned to (101) and (020) Miller planes for perovskite material. By comparison, the fully degraded  $\text{Cs}_{0.5}\text{Rb}_{0.5}\text{SnI}_3$  has peaks at  $10.0^\circ$  and  $13.3^\circ$ , characteristic of the yellow phase.<sup>2,19</sup> It should be noted that preferred orientation effects may also modify the

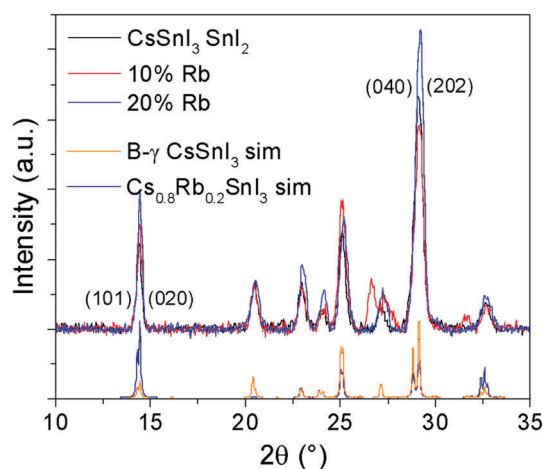


Fig. 2 XRD patterns of  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  where  $x = 0, 0.1, 0.2$ , with simulated patterns of B- $\gamma$   $\text{CsSnI}_3$ <sup>2</sup> and of  $\text{CsSnI}_3$  in which 20% of Cs atoms are replaced by Rb.



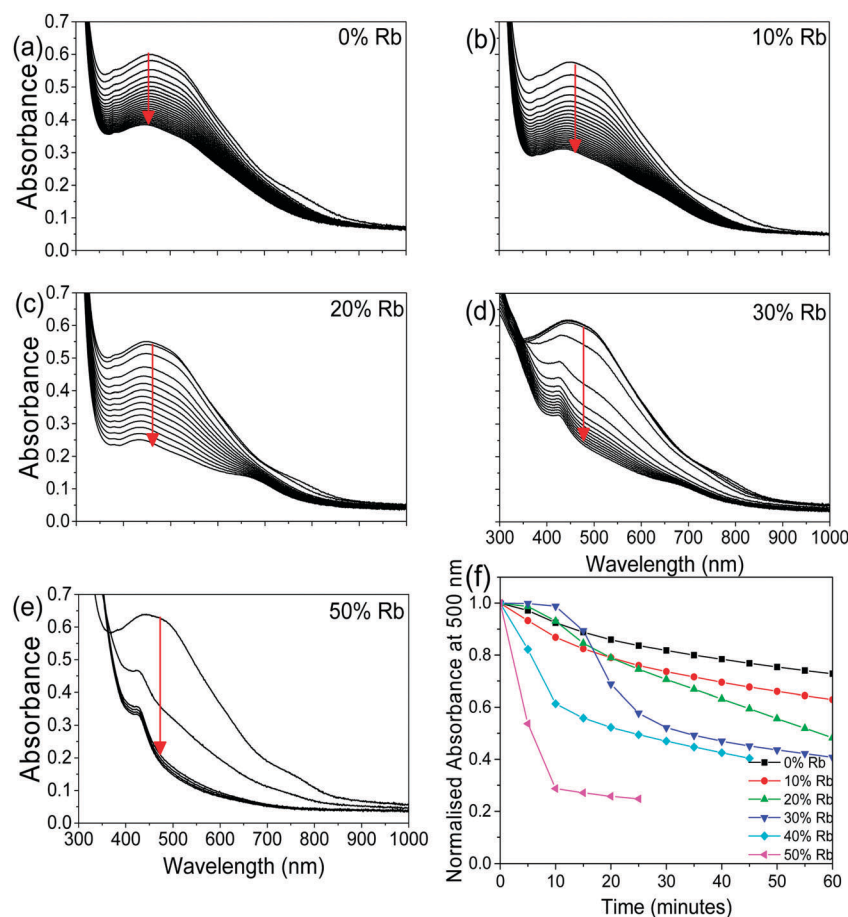


Fig. 3 (a–e) UV/vis/NIR spectra as a function of time in air (measurements made every 5 minutes) for  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3:\text{SnI}_2$  with  $x = 0, 0.1, 0.2, 0.3$ , and  $0.5$ . (f) Evolution of normalised absorbance at a wavelength of 500 nm for data shown in (a–e) as a function of time in ambient air. Data for  $x = 40$  are given in ESI,† Fig. S6.

relative intensities of reflections, indeed we have previously shown that B- $\gamma$   $\text{CsSnI}_3$  films deposited in the same way as used in the current study can exhibit substrate specific preferred crystallite orientation.<sup>9</sup> Since the degree of orientation may also conceivably be affected by the compositions of the samples this analysis cannot be quantified.

To determine how the stability of the 3D perovskite films in ambient air depends on Rb content, the evolution of the absorption spectrum of  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  films with a thickness of  $\sim 50$  nm on glass substrates was monitored as a function of time in ambient air. It is evident from Fig. 3 that incorporation of Rb into the lattice destabilises the film towards oxidation in air since there is a faster degradation of the absorption spectra with time with increasing Rb content, although for low Rb content (10%) this effect is relatively small. The observed reduction in stability with increasing substitution of Cs with Rb is consistent with the results of DFT calculation (Fig. 4), which show that the formation energy of the yellow phase of  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  from the 3D perovskite is slightly exothermic and becomes increasingly exothermic with increasing Rb content, facilitating the formation of the yellow phase. DFT simulations also show that the yellow phase of  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  will spontaneously convert to  $(\text{Cs}_{1-x}\text{Rb}_x)_2\text{SnI}_6$  upon exposure to  $\text{O}_2$  in air due to the

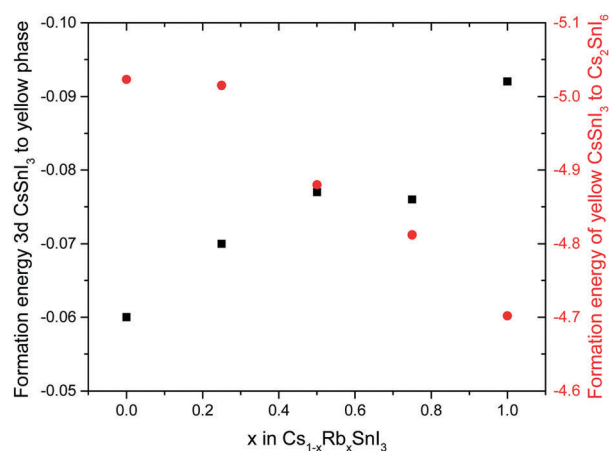


Fig. 4 Calculated formation energy of yellow  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  from 3D  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  perovskites (black squares) and formation energy of  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_6$  from yellow  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  (red circles). Both energy scales are in electron volts (eV).

large negative formation energies of the oxidised products. The reduction in stability with increasing Rb content is attributed to the increased octahedral tilting that results from substitution of Cs by Rb, leading to increased strain. Direct evidence for



increased octahedral tilting of the lattice upon incorporation of Rb, based on measurement of the optical band gap and DFT simulation, is discussed below. Corroborating evidence for a distortion of the Sn–I–Sn bonds with increasing Rb content is also provided by core level photoelectron spectroscopy; ESI,† Fig. S5(a), which shows a continuous increase in the binding energy of the Sn3d peaks with increasing Rb substitution. Notably, scanning electron microscopy images of films with  $x = 0$  and  $x = 0.2$  and  $x = 0.5$  (ESI,† Fig. S7) show that the film porosity significantly increases with increasing Rb content, which may also partially account for the differences in film stability, since more porous films have an increased surface area to volume ratio.

Fig. 5 shows the photoluminescence spectra for encapsulated  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  films with increasing Rb content, from which it is evident that the band gap increases with increasing  $x$  from  $\sim 1.34$  eV to  $\sim 1.50$  eV. Films with  $x = 0.5$  were found to be unstable even with encapsulation under nitrogen and so the spectrum shown in Fig. 5 is a sample before complete degradation. The increase in band gap with increasing Rb substitution is consistent with the electronic structure calculations performed as part of this study using the DFT-1/2 method: Fig. 6. The DFT-1/2 method has the advantage of improved accuracy in band gap calculation by introducing a half-electron/half-hole occupation, and has been successfully applied for the accurate prediction of band gaps of several metal halide perovskites including  $\text{CsSnI}_3$ .<sup>20</sup> The smaller size of Rb compared with Cs increases tilting of the  $\text{SnI}_6$  octahedra, which reduces Sn–I orbital overlap leading to a larger band gap.<sup>16,17</sup> The increase in band gap is also compelling evidence that phase separation between domains of  $\text{RbSnI}_3$  and  $\text{B-}\gamma$   $\text{CsSnI}_3$  does not occur, since  $\text{RbSnI}_3$  is predicted to have a much larger band gap than  $\text{B-}\gamma$   $\text{CsSnI}_3$  (Fig. 6 and ref. 16) and so the photoluminescence spectrum of a film with phase separated domains of  $\text{RbSnI}_3$  and  $\text{B-}\gamma$   $\text{CsSnI}_3$  would still have a significant peak at 1.34 eV. It is evident from Fig. 6 that for  $x = 0.2$  the measured band gap is significantly smaller than predicted. This disparity is attributed to the relatively small structural

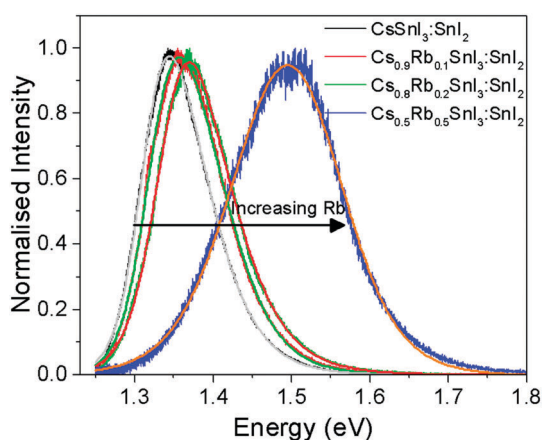


Fig. 5 Photoluminescence spectra of  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$ , where  $x = 0, 0.1, 0.2$ , and  $0.5$ , including fits for each data set, with maxima at  $1.34 \pm 0.06$  eV,  $1.36 \pm 0.08$  eV,  $1.37 \pm 0.05$  eV and  $1.5 \pm 0.11$  eV, respectively. Errors given are full peak width at half maximum.

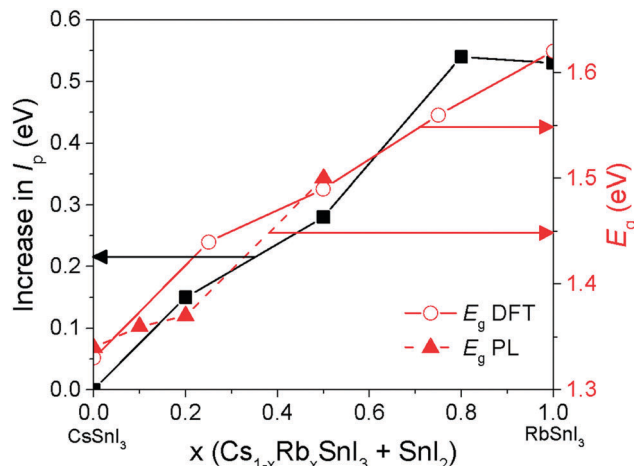


Fig. 6 Evolution of band gap ( $E_g$ ) and increase in  $I_p$  for  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  with increasing Rb substitution ( $x$ ). Band gap measured using photoluminescence spectroscopy (filled red triangles) and predicted using DFT-1/2 (open red circles). Increase in  $I_p$  measured using UPS (filled black squares). The connecting lines are included to guide the eye. The uncertainty in the DFT calculations is approximately  $\pm 0.1$  eV.

model used in the DFT calculations which will overestimate the distortion of the lattice and the octahedral tilting for small  $x$ ; ESI,† Fig. S3 and S4, giving rise to an overestimate of the band gap. However, the simulation correctly predicts the trend of increasing band gap with increasing Rb content, and for larger  $x$  the simulation and experiment are quantitatively in close agreement.

Further evidence of the uniform inclusion of Rb into the perovskite lattice is provided by the Cs:Rb elemental ratio estimated from the XPS peak intensities: ESI,† Table S1. The Cs:Rb ratios for those compositions that result in a stable perovskite structure (*i.e.*  $\text{Rb} \leq 50\%$  substitution) are in close agreement with the ratio used in the preparative solution. Since 95% of the XPS signal originates from the top  $\sim 8$  nm of the perovskite film,<sup>8</sup> this can only be the case if the Rb is uniformly incorporated into the perovskite lattice. Taken together with the photoluminescence spectroscopy, valence band photoelectron spectroscopy and prediction of simulation, this provides compelling evidence for the inclusion of Rb in to the perovskite lattice.

The potential of these materials as the light harvester in PV devices was tested in the model inverted device architecture:  $\text{ITO}|\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3:10 \text{ mol}\% \text{ SnI}_2|\text{C}_{60}|\text{BCP}|\text{Al}$ , for  $x = 0, 0.2$ , or  $0.5$ . Whilst using  $\text{C}_{60}$  as the ETL is known to give a lower  $V_{oc}$  than can be achieved using PCBM, due to its lower lying lowest unoccupied molecular orbital (LUMO),<sup>21</sup> it was used in the first instance because  $\text{C}_{60}$  can be deposited in a very controlled and highly reproducible way by vacuum deposition, rendering it well suited to this fundamental study. We have recently shown that  $\text{B-}\gamma$   $\text{CsSnI}_3$  PV devices with an inverted planar device architecture exhibit the best efficiency and stability when not using a hole-transport layer.<sup>8,9</sup> For this reason we have used this simplified device architecture as a test bed for these new perovskite materials that are closely related to  $\text{B-}\gamma$   $\text{CsSnI}_3$ .



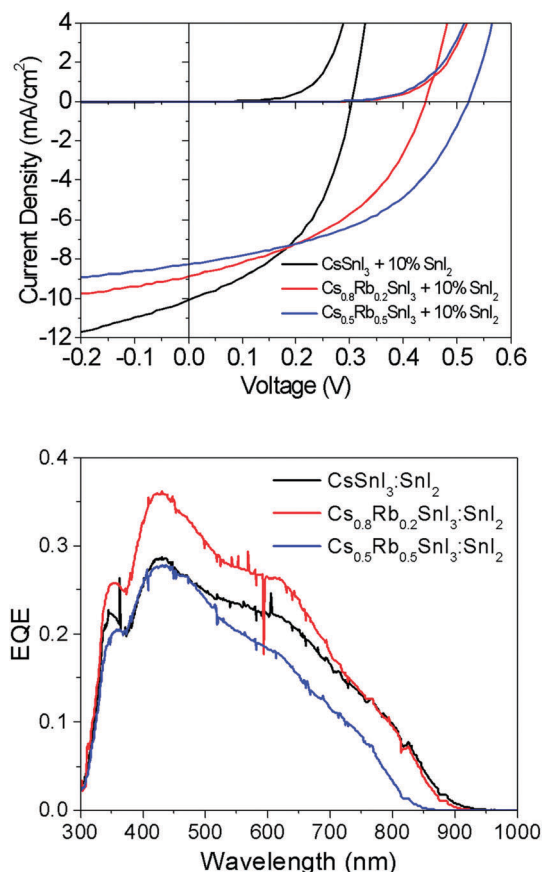


Fig. 7 (upper) Representative current–voltage and corresponding external quantum efficiency (EQE) characteristics for devices made using  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  + 10 mol%  $\text{SnI}_2$  for  $x = 0, 0.2$ , and  $0.5$  (lower) corresponding EQE spectra.

Fig. 7 shows representative current–voltage ( $J$ – $V$ ) characteristics in the dark and under 1 sun simulated illumination. The full data set is given in ESI†, Table S2. Most striking is the large increase in open-circuit voltage ( $V_{\text{oc}}$ ) with increasing Rb content, which increases by  $\sim 50\%$  from  $0.31$  V to  $0.48$  V when the Rb content is increased from  $x = 0$  to  $x = 0.5$ . Given that Rb inclusion into the B- $\gamma$   $\text{CsSnI}_3$  lattice increases the band gap, the simplest explanation for the increase in  $V_{\text{oc}}$  is a commensurate increase in the perovskite  $I_p$ , since for this device architecture the maximum  $V_{\text{oc}}$  is expected to scale with the energy difference between the valence band edge in the perovskite and the LUMO level of the fullerene ETL. To verify this hypothesis the change in  $I_p$  of  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  samples with increasing  $x$  was measured using ultra-violet photoelectron spectroscopy: Fig. 6 and ESI†, Fig. S8. From these measurements it is not possible to determine the absolute  $I_p$  in each case, because the excess  $\text{SnI}_2$  used during the preparation of the perovskite films is accumulated at the film surface<sup>8</sup> where it inevitably modifies the surface potential contribution to the measured  $I_p$ .

However, the direction and magnitude of the change in  $I_p$  can be deduced on the assumption that the excess  $\text{SnI}_2$  is distributed in a similar way in all of the perovskite films, giving rise to a comparable perturbation of the surface potential contribution to the  $I_p$  measurement for all of the samples.

Fig. 6 shows that substitution of Cs with Rb lowers the energy of the valence band edge with respect to the vacuum level (*i.e.* increases  $I_p$ ) and the magnitude scales with increasing Rb content. Indeed, the magnitude of the change in  $I_p$  with increasing Rb content correlates closely with the increase in device  $V_{\text{oc}}$ .

The change in band gap with Rb inclusion into the perovskite lattice is also evident from the change in device external quantum efficiency (EQE) spectra: Fig. 7(lower). It is estimated from the EQE spectrum for  $x = 0$  and  $0.2$  (ESI†, Fig. S9(a)) that the band gap is increased by  $\sim 20$  meV for  $x = 0.2$  which is in close agreement with the photoluminescence measurements (Fig. 5). Given that the magnitude of this increase is comparable to the thermal energy of an electron at room temperature it is barely significant, and so for  $x = 0.2$  the energy of both the valence and conduction band edges must be decreased by approximately the same amount, enabling a significant increase in  $V_{\text{oc}}$  of  $\sim 120$  meV with only a very small increase in band gap. For  $x = 0.5$  the increase in band gap compared with  $x = 0$  becomes significant, increasing by  $\sim 160$  meV from  $\sim 1.34$  to  $\sim 1.5$  eV. For  $x = 0.5$  the change in energy of the low energy edge in the EQE spectrum is, again, in close agreement with the photoluminescence measurements. The reduction in device  $J_{\text{sc}}$  with increasing  $x$  can be partially explained by the differences in perovskite film coverage, which reduces from  $\sim 98\%$  for  $x = 0$ , to  $\sim 94\%$  for  $x = 0.2$  and  $\sim 84\%$  for  $x = 0.5$  (Fig. S7, ESI†). For  $x = 0.5$  the significant reduction in band gap is also a plausible reason for the reduction in  $J_{\text{sc}}$ , since fewer long wavelength photons can be harvested.

To test the generality of this result, PV devices were also fabricated using perovskite films prepared using  $\text{SnCl}_2$  as the source of excess Sn and a PCBM ETL in place of  $\text{C}_{60}$  (Fig. 8), since we have previously shown that B- $\gamma$   $\text{CsSnI}_3$  devices using  $\text{SnCl}_2$  in conjunction with  $\text{PC}_{61}\text{BM}$  achieves substantially higher fill factor and  $V_{\text{oc}}$ , which is most pronounced after a few days storage in an inert atmosphere.<sup>8</sup>

It is evident from the data in Fig. 8 that the correlation between  $V_{\text{oc}}$  and the Rb content for devices using  $\text{SnCl}_2$  as the source of excess Sn is consistent with that observed for devices using  $\text{SnI}_2$ . For freshly made devices, the magnitude of the increase is in very close agreement (Table S3, ESI†): using  $\text{SnI}_2$  with  $x = 0.2$  and  $0.5$  the increase in  $V_{\text{oc}}$  is  $\sim 120$  mV and  $\sim 170$  mV respectively. Using  $\text{SnCl}_2$  with  $x = 0.2$  and  $0.5$  the increase in  $V_{\text{oc}}$  is  $\sim 130$  mV and  $\sim 160$  mV respectively. Interestingly, Fig. 8 and Table S3 (ESI†) show that the difference in  $V_{\text{oc}}$  for devices with and without Rb incorporated into the perovskite lattice reduces from  $\sim 130$  mV to  $\sim 90$  mV after 12 days storage in an inert atmosphere due to an increase in the  $V_{\text{oc}}$  of devices using  $\text{CsSnI}_3$ , which does not occur in devices incorporating Rb. The reason why the  $V_{\text{oc}}$  in devices with Rb incorporated into the perovskite lattice does not also exhibit an increase in  $V_{\text{oc}}$  with storage time is not yet understood. However, it is important to note that for the case of  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_3$  prepared using  $\text{SnCl}_2$  as the source of excess Sn, the material system is complicated by the possible presence of  $\text{Cs}_{1-x}\text{Rb}_x\text{SnI}_{3-y}\text{Cl}_y$  and/or phase separated  $\text{RbSnCl}_3$ . Whilst the difference in  $J_{\text{sc}}$  for devices with  $0\%$  and  $20\%$  Rb substitution is small, consistent with the very small difference in band gap, the  $J_{\text{sc}}$  (Table S3, ESI†) for  $x = 0.5$  Rb



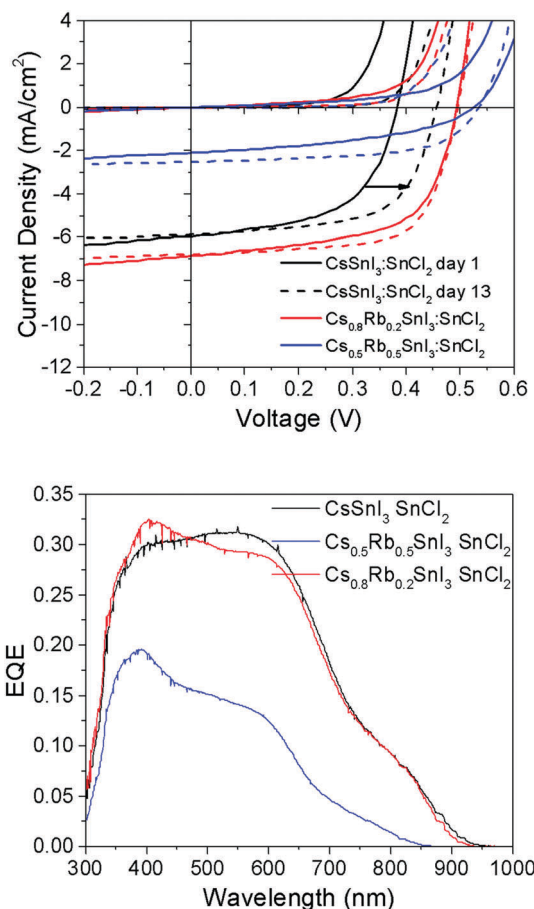


Fig. 8 (upper)  $J$ - $V$  characteristics of PV devices with the structure ITO|Cs<sub>1-x</sub>Rb<sub>x</sub>SnI<sub>3</sub> + 10 mol% SnCl<sub>2</sub>|PC<sub>61</sub>BM|BCP|Al ( $x = 0, 0.2$  or  $0.5$ ) tested in the dark and under 1 sun simulated solar illumination immediately after fabrication (solid lines) and after 12 days storage under nitrogen ( $<1$  ppm O<sub>2</sub> and H<sub>2</sub>O) (dashed lines). Full data set given in ESI,† Table S3. (lower) Corresponding EQE spectra.

substitution is greatly reduced. The latter is attributed to partial decomposition of the perovskite film *in situ* in the device, since even with encapsulation under nitrogen films with  $x = 0.5$  are unstable.

## Conclusions

In summary we have shown that the 3-dimensional perovskite Cs<sub>1-x</sub>Rb<sub>x</sub>SnI<sub>3</sub> can be prepared from room temperature solutions of RbI, CsI and SnCl<sub>2</sub> for  $x$  up to 0.5. Experiments and DFT simulations show that increasing Rb content simultaneously increases the band gap and reduces film stability, due to distortion of the lattice that occurs when Cs is partially substituted by the much smaller Rb cation at the A-site. We have shown that for small  $x$  ( $\leq 0.2$ ) film stability is sufficient for utility as the light harvesting layer in inverted PV devices. When used in conjunction with the model electron transport layer; C<sub>60</sub>, a 120 meV increase in  $V_{oc}$  was achieved for  $x = 0.2$ , which correlates closely with an increase in the perovskite ionisation potential. At the same time the increase in band gap for  $x = 0.2$

is only  $\sim 30$  meV and so the light harvesting capability of the perovskite film is not significantly degraded. Consequently, if effective strategies can be developed for the passivation of this type of perovskite, similar to that needed for lead perovskites, then inclusion of Rb into the B- $\gamma$  CsSnI<sub>3</sub> perovskite offers an effective means of increasing the ionisation potential, and thus device  $V_{oc}$ , without compromising light harvesting ability. Furthermore, we envisage that there is considerable scope for combining this strategy with halide ion substitution, which represents a fertile area for further research.

## Experimental

### Materials

CsI (Sigma-Aldrich 99.9%), SnI<sub>2</sub> (Alfa Aesar, 99.999%), SnCl<sub>2</sub> (Sigma Aldrich, 99.99%), RbI (Alfa Aesar 99.8%), PC<sub>61</sub>BM (Solenne, 99.5%), C<sub>60</sub> (nano-c, 99.9%), bathocuproine (BCP) (Alfa Aesar, 98%), *N,N*-dimethylformamide (DMF) (VWR, anhydrous, 99.8%), chlorobenzene (Sigma-Aldrich, anhydrous, 99.8%), acetone (Sigma-Aldrich, GPR, 99%), propan-2-ol (Sigma-Aldrich, GPR, 99.8%), deionised H<sub>2</sub>O (purite dispenser,  $>10$  M $\Omega$ ). CsI and non-anhydrous solvents were stored in air. All other chemicals were stored in a nitrogen filled glove box ( $<5$  ppm O<sub>2</sub> and  $<1$  ppm H<sub>2</sub>O).

### B- $\gamma$ CsSnI<sub>3</sub> films

In a dry nitrogen filled glovebox CsI, RbI, SnI<sub>2</sub> and tin(II) halide were mixed together in  $(1 - x):x:1:0.1$  molar ratio. To this mixture *N,N*-dimethylformamide (DMF) was added to make an 8 wt% solution (total mass of solids), which was stirred overnight before use. To deposit films, two drops of solution were cast onto a substrate spinning at 4000 rpm for 60 seconds. The B- $\gamma$  phase forms immediately upon solvent evaporation. For Cs<sub>0.5</sub>Rb<sub>0.5</sub>SnI<sub>3</sub>/SnI<sub>2</sub>, the film took several seconds to darken in colour after spin coating.

### Device fabrication

Indium tin oxide (ITO) coated glass slides (Thin Films Devices Inc.  $15 \pm 3 \Omega \text{ sq}^{-1}$ ) were held in vertical slide holders and ultrasonically agitated in an acetone bath, followed by a high purity water bath with a few drops of surfactant, followed by high purity deionized water only bath, and finally an isopropanol bath. After this, the slides were suspended in hot acetone vapour for 10 seconds before UV/O<sub>3</sub> treatment for 15 minutes.

Immediately after UV/O<sub>3</sub> treatment the slides were transferred into a dry nitrogen filled glovebox for CsSnI<sub>3</sub> film deposition, followed by deposition of a PC<sub>61</sub>BM film from 15 mg ml<sup>-1</sup> chlorobenzene solution using a spin speed of 1500 rpm, or C<sub>60</sub> which was deposited by thermal evaporation at a rate of  $0.4 \text{ \AA s}^{-1}$  to a thickness of 40 nm. This was followed by thermal evaporation of 6 nm at  $0.5 \text{ \AA s}^{-1}$  of bathocuproine (BCP) and then 50 nm of Al at  $1 \text{ \AA s}^{-1}$ . Thermal evaporation was performed at a pressure of  $\leq 1 \times 10^{-5}$  mbar (with substrate rotation). The Al electrode was deposited through a shadow mask to make six devices per slide, each with an area of 6 mm<sup>2</sup>.



## PV device testing

Device testing was performed in the same glove box as used for device fabrication or, as for the stability testing, using a solar simulator outside the glove box. Current density–voltage ( $J$ – $V$ ) curves were measured using a Keithley 2400 source-meter under AM1.5G solar illumination at 100 mW cm<sup>−2</sup> (1 sun), scanned from −1 V to +1 V at 0.1 V s<sup>−1</sup>. External quantum efficiency (EQE) measurements were carried out using a Sciencetech SF150 xenon arc lamp and a PTI monochromator, with the monochromatic light intensity calibrated using a Si photodiode (Newport 818-UV). The incoming monochromatic light was chopped at 180 Hz. For signal measurement a Stanford Research Systems SR 830 lock-in amplifier was used.

$J$ – $V$  and EQE measurements were made using custom LabVIEW programs.

## X-ray diffraction (XRD)

XRD was performed on thin films of CsSnI<sub>3</sub> prepared from 16 wt% (total solids) DMF solution deposited onto a glass substrate (13 × 13 mm<sup>2</sup>) spinning at 4000 rpm for 60 seconds. A thicker film than used in the devices was used here to obtain sufficient signal for the time dependent studies. Measurements were made on a Panalytical X'Pert Pro MRD equipped with a hybrid monochromator (4-bounce Ge (220) monochromator and mirror combined) to give pure Cu K<sub>α</sub> radiation and a solid state Pixel detector. The sample was kept under a flow of gas (dry nitrogen, dry air, humid nitrogen or ambient air) using an Anton Paar DHS 1100 stage with a graphite dome. Simulated diffraction patterns were calculated using the program Mercury 3.1<sup>22</sup> using CIFs from the Inorganic Crystal Structure Database (ICSD).

Powder X-ray diffraction patterns were analysed using the TOPAS software<sup>23</sup> with the Pawley method used to fit the measured profile and refine lattice parameters.

## X-ray/ultraviolet photoelectron spectroscopy

XPS was performed on film on gold coated glass substrates using a Kratos AXIS Ultra DLD. Samples were transferred using an inert atmosphere transfer arm from the nitrogen filled glovebox to the spectrometer ultra-high vacuum system and so were not exposed to the ambient environment. XPS measurements were carried out in a UHV system with a base pressure of 5 × 10<sup>−11</sup> mbar. The sample was excited with X-rays from a monochromated Al K<sub>α</sub> source ( $h\nu$  = 1486.7 eV) with the photoelectrons being detected at a 90° take-off angle. Curve fitting was performed using the CasaXPS package, incorporating Voigt (mixed Gaussian–Lorentzian) line shapes and a Shirley background.

UPS was performed in the same vacuum system as for XPS using a He 1 $\alpha$  source at 21.22 eV.

## Electronic absorption spectroscopy

UV/Vis/NIR spectra were measured for optically thin films of CsSnI<sub>3</sub> on glass substrates. Experiments from the same set were performed on the same day. Measurements were made using Agilent Technologies Cary 60 spectrometers.

## Computational methods and structural models

The initial structure optimisations were performed using DFT within the local density approximation (LDA)<sup>24</sup> as implemented in the Vienna *ab initio* simulation package (VASP).<sup>25</sup> The exchange–correlation (XC) functional is used as parameterized by Perdew and Zunger.<sup>26</sup> The outermost s, p, and d (in the case of Sn) electrons are treated as valence electrons whose interactions with the remaining ions is modelled by pseudopotentials generated within the projector-augmented wave (PAW) method.<sup>27</sup> ESI,† Fig. S10 shows the crystal structures and unit cells used in the DFT calculations. Unit cells with 12, 20, and 36 atoms are used for the case of orthorhombic and yellow phase CsSnI<sub>3</sub>, and face centered cubic (FCC) Cs<sub>2</sub>SnI<sub>6</sub> crystal structures, respectively. During the optimisation, the positions of the atoms, and the shape and volume of the unit cell are all allowed to relax. An energy cutoff of 500 eV and 6 × 4 × 6, 4 × 8 × 2, and 6 × 6 × 6  $k$ -point meshes (for orthorhombic, yellow phase, and FCC structures, respectively) are used to achieve energy and force convergence of 0.1 meV and 2 meV Å<sup>−1</sup>, respectively.

DFT–LDA underestimates the lattice parameters of the orthorhombic CsSnI<sub>3</sub> by about 3%. The deviations of the band gaps (compared to those with experimental lattice constants) are in the range of Cs<sub>1−x</sub>Rb<sub>x</sub>SnI<sub>3</sub> about 250 meV to 350 meV when using LDA-optimized lattice constants. To be consistent, all the electronic structure calculations were performed with corrected lattice parameters by expanding the lattice parameters proportionally (to match experimental volume of the cells) while keeping the LDA-optimized shape of the cells. This procedure has shown to keep the *ab initio* aspects of the approach without compromising accuracy. The subsequent electronic structure calculations were performed using the DFT-1/2 method. The DFT-1/2 method stems from Slater's proposal of an approximation for the excitation energy, a transition state method,<sup>28,29</sup> to reduce the band gap inaccuracy by introducing a half-electron/half-hole occupation. Ferreira *et al.*<sup>30</sup> extended the method to modern DFT and particularly to solid-state systems. Recently, Tao *et al.*<sup>20</sup> has reported the successful application of the DFT-1/2 method in predicting accurate band gaps of several metal halide perovskites including CsSnI<sub>3</sub> with a calculated band gap of 1.33 eV. Fortunately, the computational effort is the same as for standard DFT, with a straightforward inclusion of spin–orbit coupling when coupled with VASP. In this work, the DFT-1/2 method with the same setting is used to looking at band gap evolution of Cs<sub>1−x</sub>Rb<sub>x</sub>SnI<sub>3</sub> by substituting Cs by Rb. The DFT simulations in Fig. 4 assume the following reaction pathway: 2Cs<sub>1−x</sub>Rb<sub>x</sub>SnI<sub>3</sub> + O<sub>2</sub> → SnO<sub>2</sub> + (Cs<sub>1−x</sub>Rb<sub>x</sub>)<sub>2</sub>SnI<sub>6</sub>.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The authors would like to thank the United Kingdom Engineering and Physical Sciences Research Council (EPSRC) for funding



(Grant number: EP/L505110/1 & EP/N009096/1). The authors also thank Adam Crocker and Edward Griffin (University of Warwick) who assisted with the collection of the photoluminescence measurements. All data supporting this study are provided as ESI,<sup>†</sup> accompanying this paper.

## References

- 1 B. Wu, Y. Zhou, G. Xing, Q. Xu, H. F. Garces, A. Solanki, T. W. Goh, N. P. Padture and T. C. Sum, *Adv. Funct. Mater.*, 2017, **27**, 1604818.
- 2 I. Chung, J.-H. Song, J. Im, J. Androulakis, C. D. Malliakas, H. Li, A. J. Freeman, J. T. Kenney and M. G. Kanatzidis, *J. Am. Chem. Soc.*, 2012, **134**, 8579–8587.
- 3 Z. Chen, C. Yu, K. Shum, J. J. Wang, W. Pfenninger, N. Vockic, J. Midgley and J. T. Kenney, *J. Lumin.*, 2012, **132**, 345–349.
- 4 L.-Y. Huang and W. R. L. Lambrecht, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2013, **88**, 165203.
- 5 M. H. Kumar, S. Dharani, W. L. Leong, P. P. Boix, R. R. Prabhakar, T. Baikie, C. Shi, H. Ding, R. Ramesh, M. Asta, M. Graetzel, S. G. Mhaisalkar and N. Mathews, *Adv. Mater.*, 2014, **26**, 7122–7127.
- 6 T.-B. Song, T. Yokoyama, S. Aramaki and M. G. Kanatzidis, *ACS Energy Lett.*, 2017, **2**, 897–903.
- 7 K. P. Marshall, R. I. Walton and R. A. Hatton, *J. Mater. Chem. A*, 2015, **3**, 11631–11640.
- 8 K. P. Marshall, M. Walker, R. I. Walton and R. A. Hatton, *Nat. Energy*, 2016, **1**, 16178.
- 9 K. P. Marshall, M. Walker, R. I. Walton and R. A. Hatton, *J. Mater. Chem. A*, 2017, **5**, 21836–21845.
- 10 Q. Wang, X. Zheng, Y. Deng, J. Zhao, Z. Chen and J. Huang, *Joule*, 2017, **1**, 371–382.
- 11 Y. Yan, *Nat. Energy*, 2016, **1**, 15007.
- 12 I. Chung, B. Lee, J. He, R. P. H. Chang and M. G. Kanatzidis, *Nature*, 2012, **485**, 486–489.
- 13 J. Endres, D. A. Egger, M. Kulbak, R. A. Kerner, L. Zhao, S. H. Silver, G. Hodes, B. P. Rand, D. Cahen, L. Kronik and A. Kahn, *J. Phys. Chem. Lett.*, 2016, **7**, 2722–2729.
- 14 D. Sabba, H. K. Mulmudi, R. R. Prabhakar, T. Krishnamoorthy, T. Baikie, P. P. Boix, S. Mhainsalkar and N. Mathews, *J. Phys. Chem. C*, 2015, **119**, 1763–1767.
- 15 L. Lang, J.-H. Yang, H.-R. Liu, H. J. Xiang and X. G. Gong, *Phys. Lett. A*, 2014, **378**, 290–293.
- 16 Y. K. Jung, J. H. Lee, A. Walsh and A. Soon, *Chem. Mater.*, 2017, **29**, 3181–3188.
- 17 R. Prasanna, A. Gold-Parker, T. Leijtens, B. Conings, A. Babayigit, H. G. Boyen, M. F. Toney and M. D. McGehee, *J. Am. Chem. Soc.*, 2017, **139**, 11117–11124.
- 18 C. Li, X. Lu, W. Ding, L. Feng, Y. Gao and Z. Guo, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2008, **64**, 702–707.
- 19 G. Thiele and B. R. Serr, *Z. Kristallogr.*, 1995, **210**, 64.
- 20 S. X. Tao, X. Cao and P. A. Bobbert, *Sci. Rep.*, 2017, **7**, 14386.
- 21 B. W. Larson, J. B. Whitaker, X.-B. Wang, A. A. Popov, G. Rumbles, N. Kopidakis, S. H. Strauss and O. V. Boltalina, *J. Phys. Chem. C*, 2013, **117**, 14958–14964.
- 22 C. F. Macrae, I. J. Bruno, J. A. Chisholm, P. R. Edgington, P. McCabe, E. Pidcock, L. Rodriguez-Monge, R. Taylor, J. Van De Streek and P. A. Wood, *J. Appl. Crystallogr.*, 2008, **41**, 466–470.
- 23 A. A. Coelho, *J. Appl. Crystallogr.*, 2000, **33**, 899–908.
- 24 W. Kohn and L. J. Sham, *Phys. Rev.*, 1965, **140**, A1133.
- 25 G. Kresse and J. Furthmüller, *Comput. Mater. Sci.*, 1996, **6**, 15–50.
- 26 J. P. Perdew and A. Zunger, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1981, **23**, 5048–5079.
- 27 G. Kresse and D. Joubert, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1999, **59**, 1758–1775.
- 28 J. C. Slater, *Advances in Quantum Chemistry*, Academic Press, 1972, vol. 6, pp. 1–92.
- 29 J. C. Slater and K. H. Johnson, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1972, **5**, 844–853.
- 30 L. G. Ferreira, M. Marques and L. K. Teles, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2008, **78**, 125116.

