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23.2% efficient low band gap perovskite solar cells with cyanogen management

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W. Hashini K. Perera,^{†a} Thomas Webb,^{†b} Yuliang Xu,^c Jingwei Zhu,^c Yundong Zhou,^d Gustavo F. Trindade,^d Mateus G. Masteghin,^a Steven P. Harvey,^e Sandra Jenatsch,^f Linjie Dai,^{gh} Sanjayan Sathasivam,^{ij} Thomas J. Macdonald,^k Steven J. Hinder,^l Yunlong Zhao,^{dm} Samuel D. Stranks,^{gh} Dewei Zhao,^c Wei Zhang,^a K. D. G. Imalka Jayawardena,^{a*} Saif A. Haque^{*b} and S. Ravi P. Silva^{a*}

Managing iodine formation is crucial for realising efficient and stable perovskite photovoltaics. Poly(3,4-ethylenedioxythiophene) polystyrene sulfonate (PEDOT:PSS) is a widely adopted hole transport material, particularly for perovskite solar cells (PSCs). However, improving the performance and stability of PEDOT:PSS based perovskite optoelectronics remains a key challenge. We show that amine-containing organic cations de-dope PEDOT:PSS, causing performance loss, which is partially recovered with thiocyanate additives. However, this comes at the expense of device stability due to cyanogen formation from thiocyanate-iodine interaction which is accelerated in the presence of moisture. To mitigate this degradation pathway, we incorporate an iodine reductant in lead-tin PSCs. The resulting devices show an improved power conversion efficiency of 23.2% which is among the highest reported for lead-tin PSCs, and a 66% enhancement for the T80 lifetime under maximum power point tracking in ambient conditions. These findings offer insights for designing next-generation hole extraction materials for more efficient and stable PSCs.

Introduction

Efficiency, stability, and scalability – all within a low-cost envelope remains the major challenge of perovskite solar cell (PSC) technologies. Independently verified power conversion efficiencies (PCEs) of 26.7% and 23.6% have been reported for single junction devices based on lead (Pb)¹ and lead-tin (Pb-Sn)² compositions respectively. The rapid development of Pb-Sn perovskite devices is particularly crucial towards realising all-perovskite multijunction cells which have now exceeded 30%¹ and are anticipated to compete with perovskite/silicon multijunction cells as a low-cost alternative.

In recent years, PSCs have also seen impressive progress in operational stability with reports of stability beyond 4500 hours for the regular architecture³ and over 1000 hours for the inverted architecture^{4–6}. The instability of PSCs is accelerated

through iodine formation which occurs when the perovskite absorber is subjected to stressors such as light, heat and moisture^{4,7–9}. Once formed, iodine rapidly degrades the perovskite absorber, transport layers, and metal electrodes⁹ and thereby, device performance^{10–12}. Therefore, further advances in PCE and long-term operational stability will ultimately depend on the ability to identify and mitigate iodine-induced degradation pathways in PSCs.

For inverted PSCs, the buried interface plays a crucial role in device stability^{10,11,13}. Poly(3,4-ethylenedioxythiophene) polystyrene sulfonate (PEDOT:PSS) is a flexible, water-based hole transport layer (HTL) used in a range of applications including light-emitting diodes (LEDs), biosensors, thermoelectrics and organic solar cells¹². Its low cost¹⁴, makes PEDOT:PSS a promising candidate for upscaling of PSCs for manufacturing. PEDOT:PSS also exhibits good charge transport

^a Advanced Technology Institute, School of Computer Science and Electronic Engineering, University of Surrey, Guildford, Surrey GU2 7XH, UK.

^b Department of Chemistry and Centre for Processable Electronics, Imperial College London, London W12 0BZ, UK.

^c College of Materials Science and Engineering and Engineering Research Centre of Alternative Energy Materials and Devices, Ministry of Education, Sichuan University, Chengdu 610065, China.

^d National Physical Laboratory, NICE-MSI, Teddington TW11 0LW, UK.

^e Materials Science Centre, National Renewable Energy Laboratory, Golden, Colorado 80401, United States.

^f Fluxim AG, Katharina-Sulzer-Platz 2, 8406 Winterthur, Switzerland.

^g Department of Chemical Engineering and Biotechnology, University of Cambridge, Cambridge CB3 0AS, UK.

^h Department of Physics, Cavendish Laboratory, University of Cambridge, Cambridge CB3 0HE, UK.

ⁱ School of Engineering, London South Bank University, London SE1 0AA, UK.

^j Department of Chemistry, University College London, 20 Gordon Street, London WC1H 0AJ, UK.

^k Department of Electronic & Electrical Engineering, University College London, London WC1E 7JE, UK.

^l The Surface Analysis Laboratory, Faculty of Engineering and Physical Sciences, University of Surrey, Guildford, Surrey GU2 7XH, UK.

^m Dyson School of Design Engineering, Imperial College London, London SW7 2BX, UK.

[†] These authors contributed equally.

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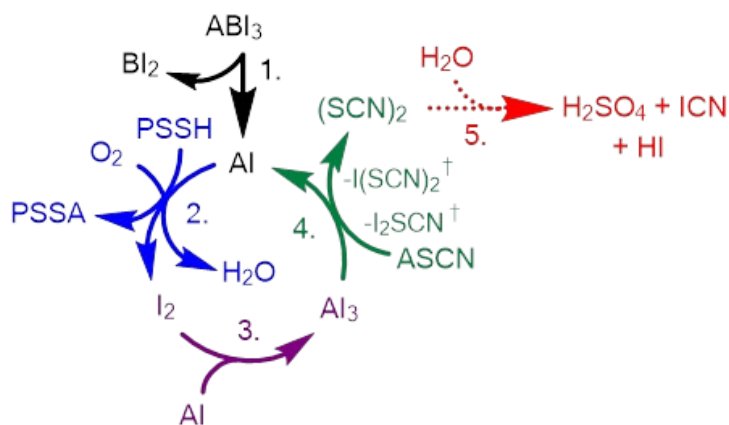


Fig. 1 Reaction scheme for interfacial chemistry occurring between PEDOT:PSS and perovskite. A denotes an organic cation, B a metal cation and PSS is polystyrenesulfonate anion. **Reaction 1:** breakdown of ABI_3 perovskite lattice into its constituent salts (Bi_2 , Al). **Reaction 2:** Oxidation of iodides (I^-) and reduction of $PEDOT^{n+}$ at the PEDOT:PSS interface leading to the formation of an organic cation-polystyrene sulfonate salt ($PSS-A$). **Reaction 3:** Reaction of I_2 within the perovskite to form corrosive triiodide species Al_3 . **Reaction 4:** Reduction of triiodide or iodide with thiocyanate salts via iodo-thiocyanate intermediates, reforming the organohalide and unstable thiocyanogen species. **Reaction 5:** Hydrolysis of thiocyanogen into cyanogen and iodine cyanide (ICN), sulphuric acid and iodic acid.

at film thicknesses >30 nm ensuring better tolerance to large-area manufacturing by reducing pinhole formation, a challenge faced with thin organic HTLs and self-assembled monolayers (SAMs)^{14–17}. PEDOT:PSS has also been the HTL of choice for single junction Sn^{18,19} and Pb-Sn^{2,20} PSCs, as well as in multijunction tandem devices with Pb-Sn sub-cells^{21–23}, mainly due to its favourable energy level alignment that allows efficient hole extraction²⁴.

PEDOT:PSS comprises of two coulombically bound ionomers which offers unique challenges concerning performance and stability. The dependence of the conducting poly(3,4-ethylenedioxythiophene) (PEDOT) polymer upon polystyrene sulfonate (PSS) for both stabilisation of charge and aqueous solubility strongly influences the doping and morphology characteristics and thereby its electronic properties²⁵. Consequently, PEDOT:PSS films are highly sensitive to factors such as pH, processing parameters, and undesirable de-doping. To this end, significant PCE and stability losses have been observed when PEDOT:PSS is used in PSCs based on Pb, Pb-Sn and Sn compositions compared to other HTLs^{26–28}, whose origins are yet to be fully understood.

Thiocyanate (SCN^-) salts are one of the most widely used additives in single and multijunction PSCs in both wide bandgap Pb, and Pb-Sn absorbers^{29–33}. However, the mechanism(s) by which SCN^- improves the performance of PSCs is/are still a topic of debate. An increase in perovskite grain size has been near-universally reported within the literature upon the direct addition of SCN^- salts to the precursor³⁴, while other studies have also suggested incorporation within the perovskite lattice, phase stabilisation, and complexation with the metal centre as well as a reduction of the trap density at the buried interface between PEDOT:PSS and the perovskite^{35,36}. While PSCs incorporating SCN^- have been reported to result in improved device stability, a majority of these have focused on the stability of devices in an inert environment (**Supplementary Table 1**) with devices tested under ambient and high humidity conditions showing poor stability relative to other HTLs³⁷. Therefore, understanding the origin and mechanism of this

instability will be a key step towards the commercialisation of perovskite photovoltaics based on PEDOT:PSS³⁸.

In this work, we employ a combination of chemical, electronic and computational methods, to identify the origin of the poor performance and stability of PSCs utilising PEDOT:PSS as an HTL and the important role of thiocyanate additives on the performance and stability in these devices. We show that de-doping of PEDOT:PSS originates from the diffusion of amines from the perovskite photoactive layer (**Reactions 1-2: Fig. 1**). This results in the generation of highly corrosive iodine species and triiodide species which subsequently facilitates the degradation of device performance and stability (**Reaction 3: Fig. 1**). We show that the incorporation of amine-containing thiocyanates significantly impedes this diffusion process, thereby reducing the extent of de-doping at the PEDOT:PSS interface, and effectively scavenges any corrosive iodine species (**Reaction 4: Fig. 1**). This results in a $\sim 30\%$ performance improvement in lead and Pb-Sn PSCs. However, under high humidity conditions relevant to ISOS test protocols (relative humidity, RH = 65%), we identify that the thiocyanogen by-products of the neutralisation of I_2 with SCN^- are highly susceptible to hydrolysis, forming corrosive cyanogens (**Reaction 5: Fig. 1**). These compounds can readily compromise the ambient stability of the thiocyanate containing devices, particularly those susceptible to I_2 formation, contributing to the short device lifetimes typically observed for tin and Pb-Sn PSCs^{2,29,30}. By adding benzyl hydrazine chloride (BHC), an iodine reductant³¹, we demonstrate a champion PCE of 23.2%. Specifically, the use of BHC introduces a new chemical pathway that successfully competes with **Reaction 4 (Fig. 1)** and minimises cyanogen formation. Further, these BHC incorporated devices show a $\sim 66\%$ improvement in the T_{80} lifetime under maximum power point tracking (MPPT) in ambient conditions compared to devices without BHC. Our findings offer new chemical insights for the rational design of next-generation hole extraction materials and iodine management strategies to enhance efficiency and stability in PSCs.

Results and discussion

Effect of PEDOT:PSS and thiocyanate additives on device performance and stability

To evaluate the influence of the buried interface on device performance, we fabricated and tested devices with a methylammonium lead iodide (MAPI) absorber layer based on the inverted *p-i-n* architecture (Fig. 2a and b) consisting of indium tin oxide (ITO)/PTAA/MAPI/C60/bathocuproine (BCP)/Silver (Ag) (reference) and ITO/PEDOT:PSS/MAPI/C60/BCP/Ag. MAPI was chosen as a simple archetypal representation of the PEDOT:PSS/perovskite which circumvents complications including composition variation, phase segregation and the instability of tin precursors^{22,30,32}.

The current density vs voltage (*J-V*) characteristics of the champion devices (Fig. 2c) shows a significant loss for the open circuit voltage (V_{oc}) and short circuit current density (J_{sc}) are evident for devices based on PEDOT:PSS. For devices based on PTAA, we achieved a champion PCE of ~19%, while PEDOT:PSS devices consistently showed a lower PCE of 11.4%. The external quantum efficiency (EQE) spectra (Fig. 2d), also shows notable loss in photocurrent throughout the absorption range for the PEDOT:PSS device indicating either higher recombination within the bulk absorber and/or poor charge extraction. With the addition of thiocyanates (lead thiocyanate ($Pb(SCN)_2$) or guanidinium thiocyanate (GASCN)), we observed a significant PCE enhancement driven by improvements across all device parameters in PEDOT:PSS devices under optimised concentrations of the additives (Fig. 2c and d, Supplementary Figs. 1-3). Further, a difference in J_{sc} was observed between the solar simulator and EQE measurements for devices based on PEDOT:PSS, which is reduced upon the incorporation of SCN^- . We attribute this to the more dominant ion migration in these devices compared to those based on PTAA as discussed later. We note that the improvements in device performance following the addition of SCN^- was also observed for Pb-Sn perovskites (Supplementary Fig 4).

To identify the origins of the observed device losses, we carried out time and frequency domain characterisation of the device stacks. Using transient absorption spectroscopy (TAS), we obtained Shockley-Read-Hall recombination rate (k_1) constants of $1.22 \times 10^6 \text{ s}^{-1}$ for PEDOT:PSS/MAPI followed by a slower rate constant of $5.40 \times 10^5 \text{ s}^{-1}$ for PEDOT:PSS/MAPI+ SCN^- with PTAA/MAPI showing the slowest rate constant of $1.05 \times 10^5 \text{ s}^{-1}$ (Supplementary Fig. 5 and Supplementary Note 1). The smaller k_1 for PEDOT:PSS/MAPI+ SCN^- as compared to PEDOT:PSS/MAPI indicates a reduction in the trap density upon the addition of SCN^- . These results are further verified through transient photocurrent measurements (TPC, Supplementary Fig. 6 and Supplementary Note 2). Additionally, using intensity modulated photocurrent spectroscopy (IMPS), we observe a low-frequency peak (i.e. <100 Hz) for perovskite devices prepared on PEDOT:PSS, characteristic of increased ion migration³³ for perovskite devices prepared on PEDOT:PSS. These result points towards the buried PEDOT:PSS interface as a direct contributor to the observed performance losses.

We next studied the stability of these devices under ISOS-D-1 (dark, 65% RH) and ISOS-D-1I (dark, N_2 environment) test conditions³⁹. Under ISOS-D-1 conditions (Fig. 2e and Supplementary Figs. 7, 8), PEDOT:PSS devices showed a relative increase of ~20% in the PCE while an onset of degradation was observed for PTAA devices. However, PEDOT:PSS devices containing SCN^- showed an accelerated degradation with a T_{80} lifetime of ~250 h, significantly shorter than that of other device architectures investigated here. Along with changes in PCE, we also observed changes in the perovskite microstructure (Supplementary Figs. 9, 10). X-ray photoelectron spectroscopy (XPS) characterisation of half cells degraded under similar conditions indicates the presence of Pb(0) on fresh perovskites formed on PEDOT:PSS in the absence of SCN^- (Supplementary Figs. 11, 12), indicating the loss iodide from the absorber. In comparison, all three device architectures tested under ISOS-D1-I conditions demonstrated significantly improved stability over the same testing period (Fig. 2f). To assess whether the ISOS-D1 humidity-driven instability of devices containing SCN^- is universal to other HTLs, we fabricated and tested the stability of MAPI+ SCN^- devices based on PTAA and Me-4PACz modified with PFN-Br and alumina nanoparticles¹⁵ which are generally regarded as more stable alternatives compared to PEDOT:PSS. However, we observed a more rapid degradation for these devices, (Supplementary Figs. 13, 14), emphasising the key role of SCN^- additives in driving device degradation. The faster degradation observed for PTAA and Me-4PACz based PSCs compared to PEDOT:PSS based devices points towards the importance of thick HTLs in minimising degradation in SCN^- containing PSCs. This is attributed to the slower release of iodine to the perovskite bulk following the iodine formation at the buried interface or within the buried HTL, the mechanism of which is discussed below.

Chemistry at the buried interface

So far, we have shown that: (i) PEDOT:PSS/MAPI based solar cells exhibit relatively poor PCE as compared to devices based on other HTLs such as PTAA; (ii) addition of SCN^- salts to the MAPI active layer significantly improves the PCE of PEDOT:PSS based devices and (iii) the improvement in PCE as reported in (ii) comes at the expense of stability upon exposure to a humid environment.

To understand the origin of these observations, we first studied the changes to the oxidation state of PEDOT:PSS using UV-Visible absorbance spectroscopy where an increasing absorbance in the near IR region (> 1100 nm) corresponds to a greater degree of oxidation (Supplementary Fig. 15)^{27,40,41}. With the addition of MAI to PEDOT:PSS, we observed a decrease in absorbance in the NIR range and an increase at 850 nm (Fig. 3a) where the latter absorbance peak is associated with the de-doped form of PEDOT:PSS (Supplementary Fig. 16)⁴⁰. This is further confirmed by electron paramagnetic resonance (EPR) measurements which showed a decrease in the integrated spin signal intensity, resembling NaOH induced de-doping (Supplementary Fig. 17).

To identify the origins of the observed PEDOT de-doping, we evaluated changes to the absorbance spectra of PSS with the

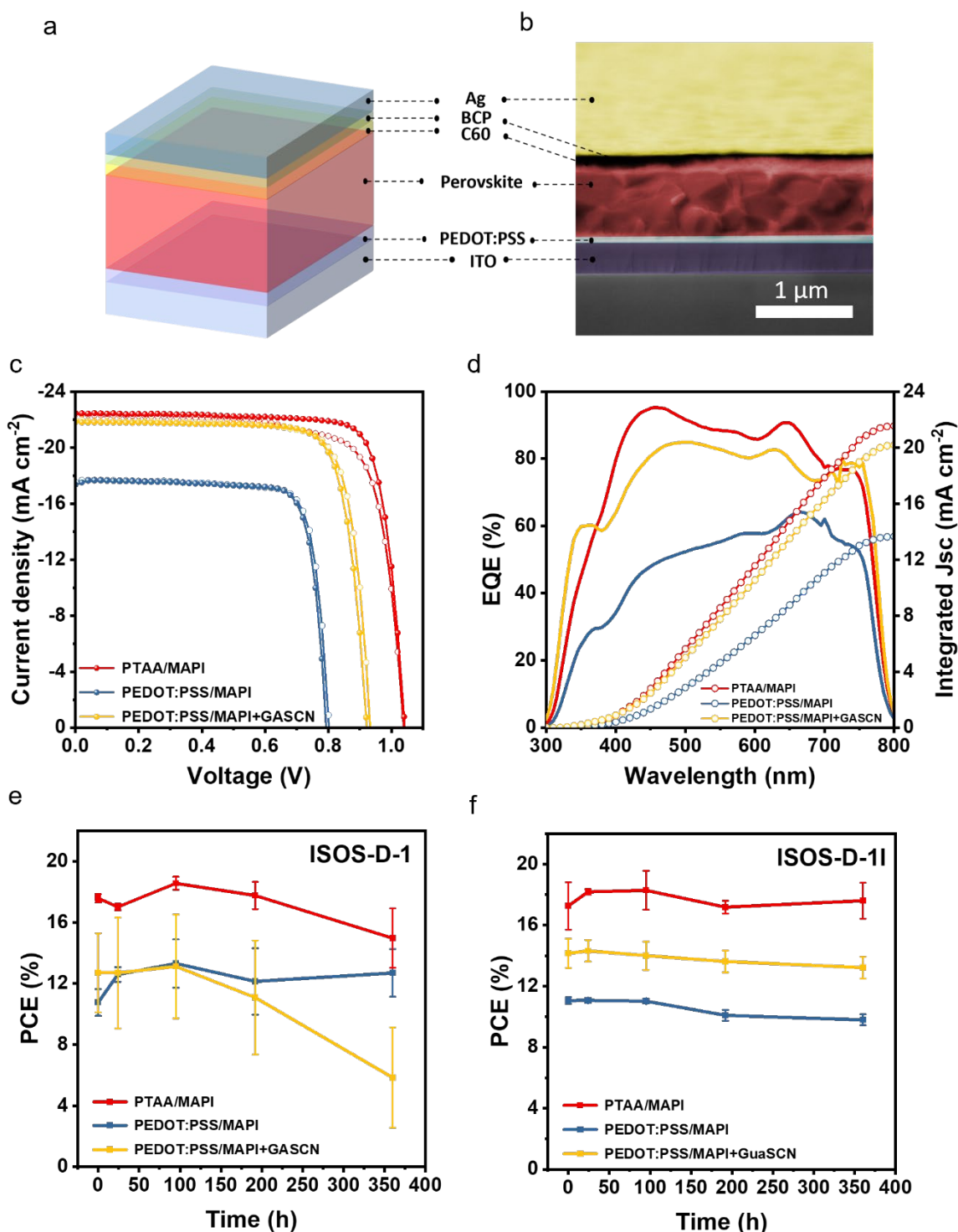


Fig. 2 Photovoltaic device performance and stability. (a) Device architecture of the PSCs. (b) A cross-sectional scanning electron microscope (SEM) image of the device, **acquired at a 45° tilt**. (c) J - V curves of the champion devices fabricated on PTAA and PEDOT:PSS with and without GASCN in MAPI under AM1.5G illumination. (d) Corresponding external quantum efficiency (EQE) spectra and integrated photocurrent densities (J_{sc}). (e) – (f) Stability testing of devices under (e) ISOS-D-1 (65% RH) and (f) ISOS-D-1I (N_2) conditions.

MAI which led to the emergence of a new absorbance feature at 366 nm (**Supplementary Fig. 18**). Mass spectrometry (MS) of the solution (**Fig. 3b**)²⁷ identifies the newly formed species as triiodide (I_3^-) which was further confirmed upon comparison of reference absorbance spectra (**Supplementary Fig. 18**). These findings suggest that the mechanism of PEDOT de-doping by the organohalide proceeds via the oxidation of iodide (I^-) anions and

the deactivation of PSS similar to NaOH induced de-doping (**Supplementary Fig. 19**)^{27, 28}.

To demonstrate that the I_3^- is formed from the interaction of the organohalide and PSS within a solid-state interface, substrates coated with MAI and PSS films were pressed together and pressure applied (**Supplementary Fig. 20**). Within hours, the films stored under ambient conditions took a yellow

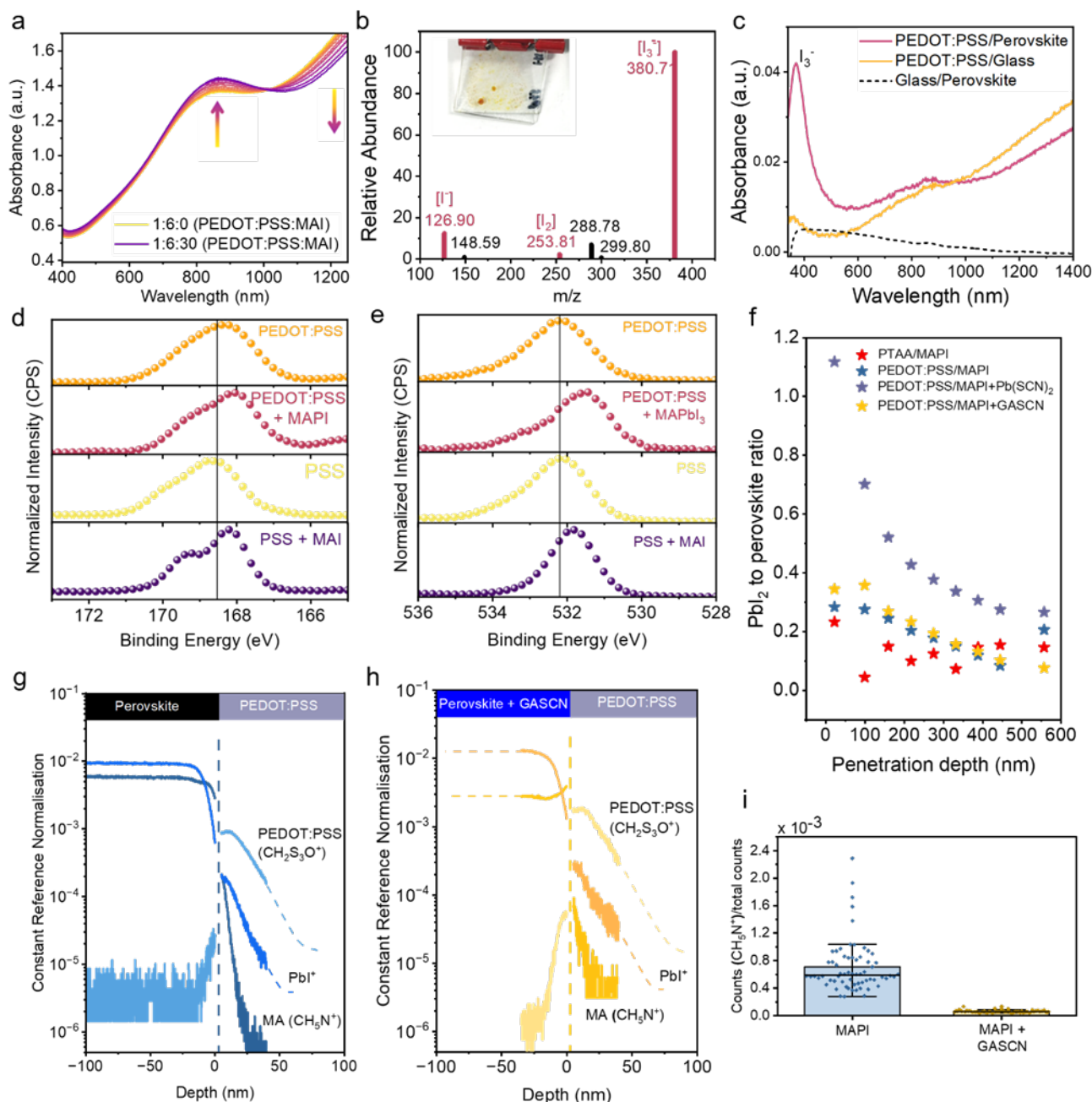


Fig. 3 Mechanism of de-doping and performance loss at the PEDOT:PSS/perovskite interface. (a) Absorbance spectra of PEDOT:PSS solution with increasing concentrations of methylammonium iodide (MAI). (b) Mass spectra collected of PSS solution combined with MAI revealing high concentrations of triiodide, (inset) photograph of PSS coated glass pressed against MAI coated glass. (c) Comparison of absorbance spectra of PEDOT:PSS films pressed against glass (yellow) or MAPI (pink). (d) – (e) X-ray photoelectron spectroscopy (XPS) of the (d) S 2p and (e) O 1s atomic orbitals. (f) Peak intensity ratios of PbI_2 to perovskite (110) as a function of penetration depth calculated from GIXRD data for grazing angles ranging from 0.25° to 2.5° . (g) – (i) Normalised ToF-SIMS depth profiles of (g) PEDOT:PSS/MAPI (h) PEDOT:PSS/MAPI + GASCN and (i) Total intensity analysis of MA^+ (CH_3NH_2^+) in PEDOT:PSS with and without GASCN.

appearance consistent with the formation of I_3^- , and confirmed via absorbance spectroscopy. Interestingly, under inert N_2 conditions, significantly less I_3^- is detected, indicating that (i) oxygen is needed for the oxidation of I^- and (ii) sufficient oxygen remains trapped or electrocatalytically bound to the surface to drive the mechanism even within an inert atmosphere⁴². Replacing the PSS with PEDOT:PSS also facilitated the formation of I_3^- along with de-doping of the PEDOT:PSS (Fig. 3h), thus highlighting the role of the

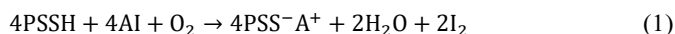
organohalide on PEDOT:PSS at the solid-state interface. Similarly, the formation of I_3^- and de-doping of the PEDOT transport material was also observed in a solid-state PEDOT:PSS/MAPI interface (Fig. 3c).

To further study the nature of the interaction between PSS and organic cation salts at the interface of PEDOT:PSS/MAPI, we combined XPS with sputtering. For this, MAPI was spin coated on PEDOT:PSS (Sample 1) and compared with a model PSS/MAPI interface (Sample 2) prepared by pressing together films as

discussed previously. In sample 1, we observed a decrease in the binding energy of O 1s and S 2p orbitals, attributed to the sulfonate group ($-\text{SO}_3^-$) of PSS, relative to that of pristine PEDOT:PSS. An identical decrease in the electron binding energy of the O 1s and S 2p orbitals was observed in the spectra of sample 2 (Fig. 3d, e), further demonstrating the interaction of PSS and MAI as the origin of the chemical modification of PEDOT:PSS when in PSCs. Interestingly, while a similar decrease in binding energy of the $-\text{SO}_3^-$ group was observed upon exposure of PSS to NaOH (Supplementary Fig. 21), no shift was present upon the addition of HI, highlighting the importance of the cation and suggesting the formation PSS-organic cation salt. To obtain further evidence for the interaction of PEDOT:PSS and the perovskite, grazing incidence X-ray diffraction (GIXRD) was used to probe relative changes in the PbI_2 content as function of the perovskite absorber depth. For PEDOT:PSS/MAPI we observed a notable increase in PbI_2 closer to the buried interface (Fig. 3f and Supplementary Fig. 22) consistent with the diffusion of MA^+ near the buried interface into PEDOT:PSS. This agrees with the chemical insights discussed above. Notably, the formation of a localised organic-cation poor region around the interface was not observed for MAPI deposited on PTAA. This highlights the importance of PEDOT:PSS in enabling this change in composition. Moreover, the PbI_2 excess (organohalide deficiency) near the buried interface was reduced upon inclusion of GASCN, which also showed a more homogenous distribution of PbI_2 through the film thickness, a feature also evident from time-of-flight secondary ion mass-spectrometric (ToF-SIMS) measurements (Supplementary Fig. 23). A similar trend was observed for $\text{Pb}(\text{SCN})_2$. To verify the diffusion of MA^+ into PEDOT:PSS, we carried out time-of-flight secondary ion mass-spectrometry (ToF-SIMS) on PEDOT:PSS exposed by cleaving of PEDOT:PSS/MAPI samples prepared without or with the SCN^- salts (see methods for details on sample preparation). In agreement with previous results, the ToF-SIMS measurements show the tendency of MA^+ to diffuse within PEDOT:PSS in samples prepared with MAPI (Fig. 3g) which is reduced upon the addition of GASCN (Figure 3g – i and Supplementary Fig. 24).

Having shown that the MAI salt in the perovskite reacts with PSS, thereby de-doping the HTL and generating iodine (Fig. 1 and Equation (1)), we next demonstrate that this mechanism is ubiquitous amongst other organohalides (Supplementary Fig. 25). As before, using absorbance spectroscopy, we compared the changes in absorbance intensity of PEDOT:PSS at 1200 nm and 866 nm (corresponding to its higher and lower oxidation states respectively) upon the addition of MAI, FAI and GAI. From Fig. 4a and b, it is evident that increasing the number of amine groups on the organohalide salt leads to increased de-doping of PEDOT:PSS, which is further supported by EPR spectroscopy (Supplementary Fig. 26). We also observed aggregation of PEDOT following the deactivation of PSS with the organohalide (Supplementary Figs. 27-33), a consequence of the organohalides blocking PSS's role in producing micelles around PEDOT during deposition. Furthermore, we find that both the

degree of aggregation and de-doping ability increases with increasing amine groups ($\text{MA}^+ < \text{FA}^+ < \text{GA}^+$).



We employed Density Functional Theory (DFT) to investigate the thermodynamics of Equation (1) and the associated ionic binding energies of the salts involved. DFT predicts a favourable thermodynamic driving force (ΔG_r) for the forward reaction Equation (1), for all three cations investigated (Fig. 4c). Next, we evaluated the ionic binding energy $\Delta G_b^g(\text{A:PSS})$ between the PSS anion and organic cation A^+ . As expected, increasing the delocalisation of the cation across multiple amine groups decreased the strength of the ionic binding energy with the PSS anion (Supplementary Fig. 34)⁴³⁻⁴⁵. This is further observed in the electronic surface potential (ESP) maps of the A:PSS salts following structural optimisation (Fig. 4d - g) and by consideration of the total Mulliken charge across the PSS sulfonate anion ($-\text{SO}_3^-$) (Supplementary Fig. 35). Using XPS to compare the electron density on the S 2p orbital of the PSS sulfonate group in the absence, and presence of the organic cation salts, upon exposure to MA^+ , we observed an increase in the electron density on the $-\text{SO}_3^-$ which is further increased upon incorporation of FAI (Supplementary Fig. 36). We calculated the displacement energy $\Delta G_d(\text{PEDOT/A:PSS})$, defined as the difference in free energy for an isolated A^+ cation to displace a PEDOT trimer from PSS, forming a highly ionic A:PSS-salt and isolated PEDOT cations (see Methods). The $\Delta G_d(\text{PEDOT/A:PSS})$ values for MA^+ , FA^+ and GA^+ (Table 1) indicates the capability of these cations to displace and destabilise PEDOT when exposed to PEDOT:PSS, in agreement with the experimentally observed de-doping at the perovskite/PEDOT:PSS interface.

Table 1 Gas-phase DFT calculations for reaction, displacement and ionic binding energies (kJ mol^{-1}).

Organic Cation	ΔG_r (Eq.1)	ΔG_b (A: PSS)	ΔG_d (PEDOT/A:PSS)
MA^+	-125.4	-473.6	-194.5
FA^+	-129.6	-457.0	-177.9
GA^+	-120.6	-421.1	-142.0

When accounting for the factors which can dictate trends in de-doping, it is important to consider the binding energy $\Delta G_b(\text{A:I})$ of the organohalide salts (Table 1). In this case, the increased delocalisation of the charge reduces the ionic binding energy with the halide. This allows for the oxidation of the iodide anions which generates free organic cations to form a salt and deactivate the PSS (Equation (1)). Additionally, we note that charge delocalisation across multiple amine groups increases $\Delta G_b(\text{A:PSS})$ when a second neighbouring polystyrene group is introduced. In this case, the delocalisation of the charge enables bridging between the two sulfonate groups thereby allowing a single organic cation to interact strongly with adjoining PSS monomers (Supplementary Fig. 37). As such, the local stoichiometry and concentration effects between the organic cations and PSS will likely influence the co-ordination to PSS and its subsequent deactivation. From this comparison, we

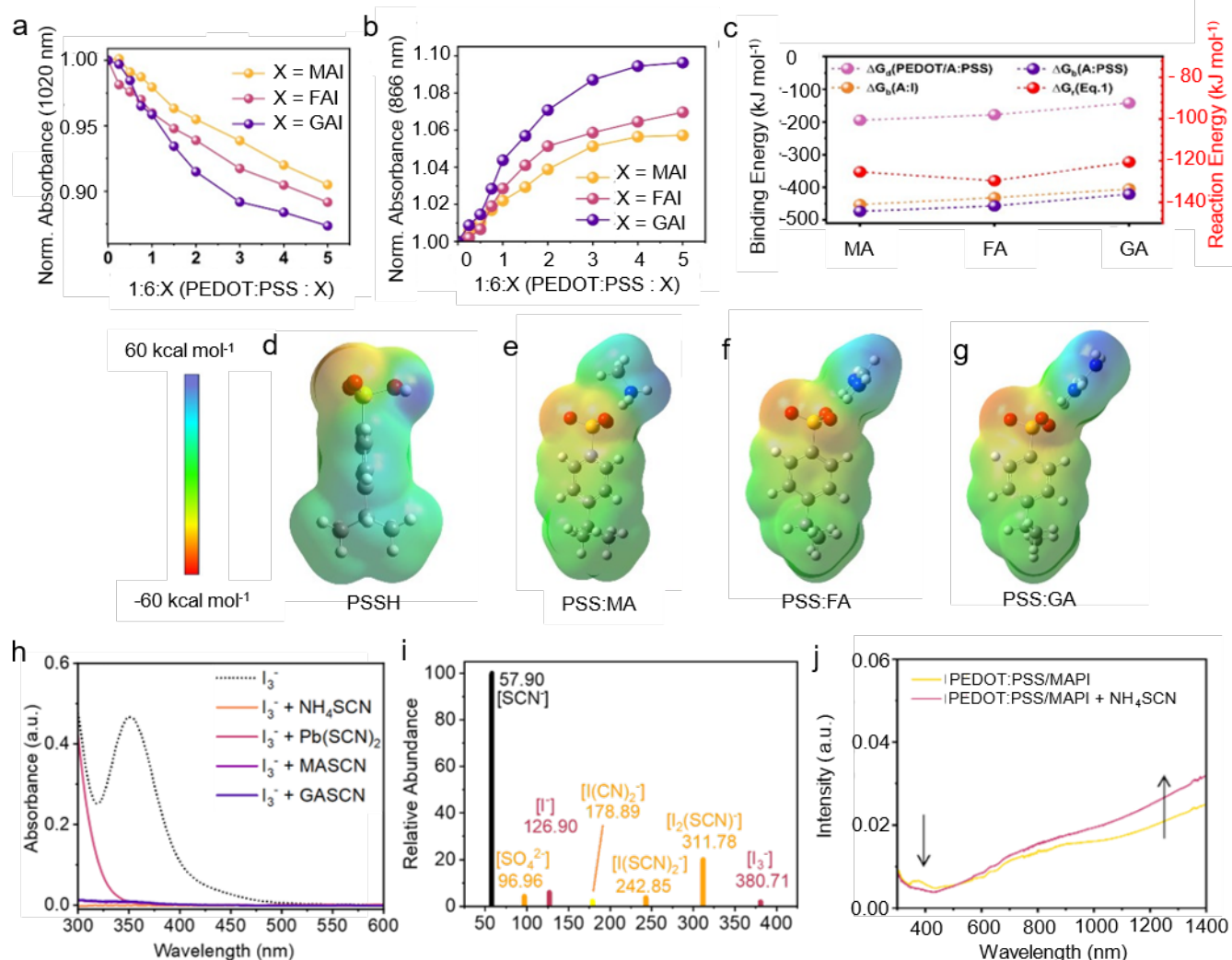


Fig. 4 The effect of cation choice on PEDOT:PSS de-doping. (a) – (b) Absorbance of PEDOT:PSS with various A site cations of (a) high oxidation states (1200 nm) and (b) lower oxidation states (866 nm). (c) DFT predictions of the displacement energy between PEDOT:PSS and A:PSS (PEDOT/A:PSS), ionic binding energy of the AI salt (A:I), ionic binding energy of the A:PSS salt (A:PSS), and reaction energy of equation 1 (Eq.1), calculated using a 6-311G(d,p) (H,C,N,O,S) + LANL2DZ (I) hybrid basis set (see methods). (d) – (g) Electronic surface potential of PSS-cation ionic pair when interacting with (d) H⁺, (e) MA⁺, (f) FA⁺, and (g) GA⁺. (h) Absorbance spectra showing neutralisation of a MAI₃ reference solution following addition of various thiocyanate salts. (i) Mass spectra of MAI₃ solution following addition of NH₄SCN. (j) Absorbance spectra of PEDOT:PSS films following pressing against MAPI perovskite films with and without NH₄SCN in an N₂ atmosphere for 7 days.

hypothesise that the observed trend in de-doping ability, from a thermodynamic viewpoint, is driven by a combination of the stability of the initial organohalide salt, the concentration of the organic cation available to displace PEDOT in PEDOT:PSS, and the binding strength of the PSS with the organic cation. However, additional factors such as the PSS micelle size, the inability of GA⁺ to occupy a 3D perovskite lattice position, mobility/diffusion of the organic cation, and defect density will also have a role in dictating de-doping of PEDOT:PSS.

So far, we have discussed the chemical reactions at the PEDOT:PSS/perovskite interface which are responsible for de-doping of the HTL and leading to the formation of I₃⁻. We next investigated the role of SCN⁻ salts in mitigating this issue. We first directly added thiocyanate salts to MAI₃ solutions and observed a colour change from yellow to colourless consistent with the reduction of I₃⁻ to I⁻ (Fig. 4h and Supplementary Fig.

38). Notably, in the case of Pb(SCN)₂, the reaction with MAI₃ formed a bright yellow precipitate attributed to PbI₂ (Supplementary Fig. 38). Cyclic voltammetry (CV) was used to compare the redox potentials of I₂ and NH₄SCN using tetrabutylammonium hexafluorophosphate [nBu₄PF₆] an anhydrous and degassed acetonitrile electrolyte (Supplementary Fig. 39). The collected voltammograms revealed similar redox potentials between the SCN⁻ and I₂^{46–48} revealed similar redox potentials, making the reaction susceptible to additional parameters including the concentrations of [I⁻]:[I₃⁻]:[I₂] and most notably, the concentration of trace water (the importance of this is discussed shortly). Indeed, upon addition of SCN⁻ ions to I₂ or vice versa, we observed the formation of I⁻ of a second redox peak consistent with the introduction of I⁻ into the equilibrium and subsequent formation of I₃⁻. To further confirm the reduction

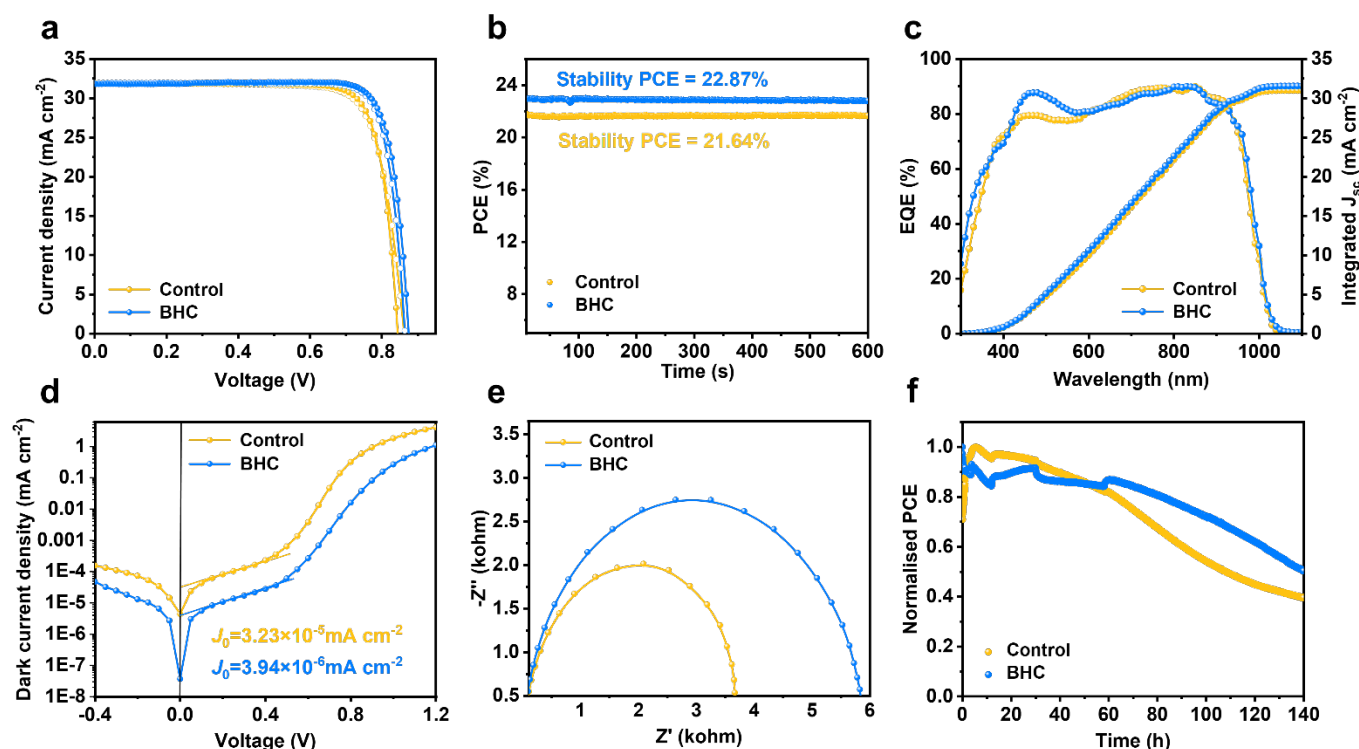
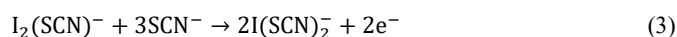
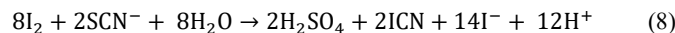


Fig. 5 Device performance and characterisations of Pb-Sn PSCs with and without BHC. (a) J - V curves of the champion Pb-Sn PSCs. (b) Steady-state efficiency outputs. (c) Corresponding EQE spectra and integrated J_{sc} . (d) Dark J - V curves. (e) EIS spectra. (f) Continuous MPP tracking of encapsulated devices under a 100 mW cm^{-2} multicolour LED solar simulator in ambient air.

of I_2 , as well as to identify by-products, MS analysis was performed on MAI_3 following the addition of NH_4SCN (Fig. 4i). Two additional small peaks in the mass spectra with fragment masses of 242.85 a.u. and 311.78 a.u. are identified as $[I(SCN)_2]^-$ and $[I_2(SCN)]^-$ and correspond to the intermediates of iodine reduction/ SCN^- oxidation as given in Equations (2)–(4)⁴⁹.



Mass spectra also revealed additional peaks with masses of 96.96 and 178.88 a.u. corresponding to SO_4^{2-} and $[ICN(CN)]^-$, respectively. This is consistent with the hydrolysis of thiocyanogen $(SCN)_2$, the oxidation product of thiocyanate. Under this mechanism, upon contact with ambient moisture, hygroscopic $(SCN)_2$ species undergo hydrolysis forming ICN, HCN and H_2SO_4 as indicated in Equations (5) – (8) (Reaction 5, Fig. 1). Additional mass spectra were collected using anhydrous isopropyl alcohol dried over molecular sieves where identical hydrolysis products were recorded, demonstrating the high degree of moisture sensitivity of the reactions following the brief exposure to ambient conditions during sample loading (Supplementary Fig. 40).



A combination of 1H and ^{13}C nuclear magnetic resonance (NMR) techniques (Supplementary Fig. 41) were employed to further study the nature of the decomposition products. Upon addition of I_2 to a suspension of $Pb(SCN)_2$ in $CDCl_3$, we observed a downfield shift in the H_2O trace peak from 1.59 ppm to 1.62 ppm consistent with acidification and the formation of the hydronium ion (H_3O^+). Likewise, attenuated total reflection Fourier Transform Infrared (ATR-FTIR) spectroscopic measurements on NH_4SCN before and after I_2 exposure shows the formation of a new peak around 3500 cm^{-1} , consistent with H_2SO_4 evolution (Supplementary Fig. 42). ^{13}C NMR on the same $Pb(SCN)_2$ solution before and the addition of I_2 (Supplementary Fig. 41) showed no clear peaks in the former owing to the insolubility of the SCN^- salt in the solvent. Upon addition of I_2 to the solution, we observed a new peak at 30.95 ppm, consistent with the formation of ICN. However, we note this peak was not observed when NMR samples were prepared using extra dry solvents and carefully desiccated inorganic salts whereby thiocyanogen compounds do not undergo hydrolysis and subsequent decomposition.

We next investigated the impact of ambient moisture in facilitating the reduction of I_3^- via oxidation and subsequent hydrolysis of thiocyanate anions. To do this, changes in the CN bond were measured using ATR-FTIR following the reaction of NH_4SCN with MAI_3 in solid-liquid and solid-solid phases under ambient conditions (Supplementary Fig. 43). In each case, the C-N stretch of SCN was shifted to lower frequencies. These

results are as expected owing to the highly hygroscopic nature of SCN^- salts. Similarly, confocal Raman spectroscopy probing NH_4SCN before and after mixing with solid MAI_3 revealed a shift and broadening of the low-frequency S-C bend, demonstrating the availability of sufficient ambient moisture for SCN^- to react with I_3^- (Supplementary Fig. 44).

We now consider the implications of this fundamental chemistry on device performance. The presence of I_3^- has previously been linked to the formation of shallow traps, while de-doping of PEDOT:PSS leads to reduced conductivity and the formation of a charge extraction barrier which degrades PSC performance (Supplementary Figs. 45–47)^{31,50,51}. We attribute the improved device PCE in thiocyanate containing devices to the neutralisation and the reduction of I_3^- traps. Indeed, PEDOT:PSS / MAPI interfaces prepared with NH_4SCN added to the perovskite precursor resulted in less I_3^- and reduced de-doping of the PEDOT:PSS, as compared to those without SCN^- (Fig. 4j and Supplementary Fig. 48). While the reduction in I_3^- is undoubtedly beneficial to the performance and longevity of the resulting cells, we note this stability improving mechanism (I_3^- scavenging via thiocyanates, Reaction 4, Fig. 1) is likely to be offset by the rapid hydrolysis of the formed thiocyanogen (Reaction 5, Figure 1), particularly under humid conditions or at high thiocyanate loadings. Indeed, the corrosive nature of cyanogen and iodocyanogens explains the poor stability of thiocyanate containing devices under moisture or high relative humidity conditions (Figure 2e-f).

Effect of iodine reductant on stability and efficiency in Pb-Sn cells

Next, we sought to investigate if the incorporation of an iodine reducing agent can improve the performance and stability of Pb-Sn mixed PSCs by switching off Reaction 4 (Fig. 1). To this end, we incorporated benzyl hydrazine chloride (Supplementary Note 4) (BHC), an iodine reductant⁴⁴, into the Pb-Sn precursor and fabricated devices with the architecture of glass/ITO/PEDOT:PSS/perovskite/C60/BCP/Cu. Examples for other reductants and scavengers are discussed in (9) and exemplars are given in Supplementary Table 2. The incorporation of BHC resulted in devices with a champion PCE of 23.18%, J_{sc} of 31.84 mA cm^{-2} , V_{oc} of 0.875 V and a fill factor (FF) of 83.23% (Fig. 5a, Table 2 and Supplementary Fig. 49). In comparison, the control device showed a lower PCE of 21.86%, with a J_{sc} of 31.53 mA cm^{-2} , V_{oc} of 0.852 V and a FF of 81.41%. The enhancement in the FF particularly is ascribed to improved perovskite bulk quality as a result of BHC reducing iodine to iodide. The steady-state efficiencies of the BHC device and the control (without BHC) are 22.87% and 21.64% (Fig. 5b) at a bias of 0.728 V and 0.710 V respectively. The higher stabilised power output for BHC device agrees well with the $J-V$ measurements. Fig. 5c shows the corresponding EQE spectra of the devices. The EQE-integrated J_{sc} over the entire solar spectrum was improved from 30.98 to 31.56 mA cm^{-2} upon the incorporation of BHC, in accordance with the values from $J-V$ measurement. This significant enhancement of integrated J_{sc} originates from the notable increase in the spectral response over the 415–600 nm wavelength range. To gain more insights into the reduced non-radiative recombination in BHC devices, we measured the dark

$J-V$ curves of these devices (Fig. 5d). The dark saturation current density (J_0) of BHC device was decreased from $3.23 \times 10^{-5} \text{ mA cm}^{-2}$ (for control device) to $3.94 \times 10^{-6} \text{ mA cm}^{-2}$, indicating a reduced leakage current and a smaller barrier for charge transfer. The electrochemical impedance spectroscopy (EIS) results (Fig. 5e) showed increased recombination resistance in BHC incorporated devices, demonstrating the suppression of non-radiative recombination, consistent with the smaller J_0 obtained from the dark $J-V$ curve and suppression of I_3^- induced charge trapping. To examine the impact of BHC incorporation on stability, we then conducted MPP tracking to monitor the change of operational stability in ambient air. The encapsulated BHC device maintained 80% of its peak efficiency (i.e. T_{S80} lifetime) after continuous 1-sun illumination for 83 h, while the control device showed rapid degradation (Fig. 5f). We attribute this to the iodine scavenging effects of BHC which reduces carrier recombination and improves device efficiency. However, degradation can be observed which indicates that alternate strategies are required, some of which we highlight in Supplementary Notes 3, 4 and Supplementary Figs. 50, 51. We note that a majority of testing reported in literature for Pb-Sn devices utilise a nitrogen atmosphere, thereby masking the crucial role of a strong encapsulant and moisture removal for the stability of this technology. We also carried out topographical, structural and optical analysis to identify any changes to the absorber following the addition of BHC (Supplementary Figs 50–52). The addition of BHC did not result in a noticeable change to the root mean square roughness of the sample (which was ~49 nm), grain sizes as well as the crystalline structure nor the optical properties.

Table 2 Summary of photovoltaic parameters of champion Pb-Sn PSCs with and without BHC.

Condition	Scan direction	PCE (%)	FF (%)	V_{oc} (V)	J_{sc} (mA cm^{-2})
Control	Reverse	21.86	81.41	0.852	31.53
	Forward	21.34	79.14	0.858	31.43
BHC	Reverse	23.18	83.23	0.875	31.84
	Forward	22.89	82.93	0.864	31.93

Conclusions

In conclusion, we identify that the properties of PEDOT:PSS are significantly modified upon depositing a layer of perovskite owing to undesirable chemical interactions at the interface. More specifically, the nature of these interactions corresponds to the diffusion of the organohalide cations from the perovskite into the underlying PEDOT:PSS layer. This leads to the deactivation of PSS and subsequent de-doping of PEDOT:PSS while simultaneously forming trap states in the perovskite absorber, thereby resulting in device performance loss. The above process further results in the oxidation of iodide anions to form triiodide, proceeding first via the formation of iodine. The addition of thiocyanates reduces the device performance losses by reducing or preventing triiodide formation. However, when stored under ambient conditions, the reaction of the SCN^-

oxidation product (SCN)₂ with moisture results in the formation of highly corrosive ICN and HCN, leading to a trade-off between minimising device losses and compromising ambient stability. We show that iodine reducing additives can mitigate these losses and improve device efficiency. Using Pb-Sn single junction PSCs, an architecture that to date has heavily relied on the use of PEDOT:PSS as the HTL, we show an improvement in device efficiency upon adding BHC with a champion efficiency of 23.2% being realised, which is among the highest reported for Pb-Sn mixed PSCs. In parallel, we also observe an improvement of T₈₀ lifetime by 66% under maximum power point tracking in ambient conditions. Based on our findings, we offer insights for designing next-generation hole extraction materials to enhance efficiency and stability in large-area PSCs.

Experimental

Materials

Formamidinium iodide (FAI, ≥99.99%), methylammonium iodide (MAI, >99.99%) and guanidinium iodide (GAI) were purchased from GreatCell Solar Materials, Australia. Lead(II) iodide (PbI₂, 99.99%) and benzylhydrazine hydrochloride (BHC) were purchased from Tokyo Chemical Industry Co., Ltd. (TCI, Japan). Poly(3,4-ethylenedioxythiophene) polystyrene sulfonate (PEDOT:PSS, CLEVIOS™ P VP AI 4083) was purchased from Heraeus, Germany. Poly[(9,9-bis(3'-(N,N-dimethyl)-N-ethylammonium)-propyl)-2,7-fluorene]-alt-2,7-(9,9-dioctylfluorene)]dibromide (PFN-Br) was purchased from 1-material, Canada. [4-(3,6-dimethyl-9H-carbazol-9-yl)butyl]phosphonic acid (Me-4PACz) was purchased from Dyenamo, Sweden. Poly(triaryl amine) (PTAA), lead(II) thiocyanate (Pb(SCN)₂, 99.5% trace metals basis), guanidine thiocyanate (GASCN, for molecular biology, ≥99%), aluminium oxide (Al₂O₃, 20% (v/v) in 2-propanol), tin(II) iodide (SnI₂, 99.999%), tin(II) fluoride (SnF₂, 99%), ethylenediammonium diiodide (EDAI₂, 98%) and ammonium thiocyanate (NH₄SCN), and solvents dimethyl sulfoxide (DMSO, anhydrous, ≥99.9%), N,N-dimethylformamide (DMF, anhydrous, 99.8%), toluene (anhydrous, 99.8%) and chlorobenzene (CB, 99.9%) were purchased from Sigma Aldrich, UK. Solvents diethyl ether (99.5%, Extra Dry, Acros), methanol (HPLC grade, ≥99.9%), ethanol (anhydrous, >99.5%) and 2-propanol (anhydrous, 99.5%, Acros) were purchased from Fisher Scientific, UK. Caesium iodide (CsI, 99%) was purchased from Alfa Aesar. C₆₀ was purchased from Ossila, UK or Nano-C. Bathocuproine (BCP, sublimed grade >99.5%) was purchased from Ossila, UK or Jilin OLED Company, China. Copper (Cu) was purchased from Zhongnuoxincai Co., Ltd.

Substrate cleaning and preparation

ITO coated glass substrates purchased from South China Science and Technology Ltd and from Youxun, China (20 mm x 20 mm with a thickness of 1.1 mm and a sheet resistance of 15 Ω/□) were first cleaned by sonicating in a 2% (v/v) Hellmanex in water solution for 30 minutes at 35–40 °C. The substrates were then rinsed with deionised water and sonicated in water for further

30 minutes. Thereafter, they were sequentially cleaned in acetone, 2-propanol and methanol in an ultrasonic bath at ~25 °C for 15 min each and blow dried with N₂. Prior to coating with PEDOT:PSS and PTAA, the substrates were treated with an oxygen plasma for 5 min at 200 W using a Diener Zepto system. Prior to coating with Me-4PACz, the substrates were subjected to a UV-O₃ pre-treatment for 10 min using a Jelight UV-O3 Cleaner Model 24.

For Pb-Sn devices (Sichuan University): The pre-patterned ITO substrates were cleaned by sonicating sequentially in deionised water and anhydrous ethanol for 15 min each, and then dried under N₂ stream and treated by UV-Ozone for 15 min.

Solution preparation

PEDOT:PSS: CLEVIOS™ P VP AI 4083 was used as received. Prior to deposition, the dispersion was allowed to warm up to room temperature for 1 hour.

PTAA: A 2 mg/mL solution of PTAA was prepared in toluene and left to stir overnight inside the glovebox.

Me-4PACz: A 1 mmol L⁻¹ solution of Me-4PACz was prepared in ethanol and left to stir overnight inside the glovebox.

PFN-Br: A 0.5 mg mL⁻¹ solution of PFN-Br was prepared in methanol and left to stir overnight inside the glovebox.

Al₂O₃: Al₂O₃ 20% (v/v) in 2-propanol, Sigma Aldrich was diluted in 2-propanol to form a 1:1000 Al₂O₃ dispersion.

MAPI: PbI₂ (461 mg) and MAI (159 mg) were weighed into one vial. To this, DMF (595 μL) and DMSO (71 μL) were added and left to stir overnight in the glovebox at room temperature. For MAPI solutions containing Pb(SCN)₂ or GASCN, the thiocyanate was weighed into a separate vial and the MAPI solution was transferred to the powder in the vial and allowed to stir 1 hour before deposition. For all MAPI solutions, the precursor was stirred at 65 °C for 15 min prior to deposition and maintained at 65 °C during deposition.

PEDOT:PSS with organohalides: MAI, FAI or GAI was added to PEDOT:PSS (filtered through a 0.45 μm PTFE syringe filter) to form dispersions with 1:0.25, 1:0.5, 1:0.75, 1:1, 1:1.5, 1:2, 1:3, 1:4 and 1:5 PSS:organohalide molar ratios.

For Pb-Sn devices: NH₄SCN (2.74 mg), SnF₂ (14.10 mg), CsI (46.77 mg), MAI (85.84 mg), FAI (185.73 mg), SnI₂ (335.27 mg) and PbI₂ (414.91 mg) were dissolved in 1 ml mixed solvent of DMF and DMSO (DMF:DMSO = 3:1, v/v) to obtain the FA_{0.6}MA_{0.3}Cs_{0.1}Pb_{0.5}Sn_{0.5}I₃ 1.8 M precursor. For the BHC doping precursor, 2.1 mg (0.75 mol%) of BHC was added to 1 mL perovskite precursor. The post-treatment solution was prepared by dissolving 0.5 mg of EDAI₂ in 1 mL mixed solvent of toluene and IPA (toluene: IPA = 1:1, v/v). For the Pb-Sn devices fabricated at Surrey, 4 mg of GASCN was used instead of NH₄SCN.

Device fabrication

For PEDOT:PSS substrates, PEDOT:PSS was deposited through a 0.45 μm PTFE syringe filter and spun at 5000 rpm for 40 s. The films were then annealed at 150 °C for 30 min. For PTAA samples, the substrate was coated with 100 μL of the solution, spun at 6000 rpm for 33 s and then annealed at 100 °C for 10 min. For Me-4PACz substrates, 50 μL of the solution was spread

on the substrate and after a waiting time of 5 s, it was spun at 3000 rpm for 30 s. The films were then annealed at 100 °C for 10 min. For PTAA substrates, PFN-Br was used as a surface compatibiliser, while for Me-4PACz coated substrates, either PFN-Br or Al₂O₃ nanoparticles was used as previously reported¹⁵. For PFN-Br modified samples, 30 µL of the PFN-Br solution was dispensed at 5 s from the spin start time on a substrate spinning at 5000 rpm for 30 s. For Al₂O₃ modified samples, 40 µL of the solution was spin coated at 4000 rpm for 30 s. The perovskite was deposited by spin coating 35 µL of the MAPI at 4000 rpm for 30 s while 600 µL of diethylether, the antisolvent, was dropped at 7 s from the spin start time. The films were annealed, first at 65 °C for 2 min and then at 100 °C for 10 min. C60 (20 nm), BCP (7 nm) and Ag (100 nm) were thermally evaporated in an Angstrom EvoVac system.

For organohalide incorporated PEDOT:PSS samples, the same deposition procedure as PEDOT:PSS was followed using 75 µL of the dispersion.

For Pb-Sn devices (Sichuan University), PEDOT:PSS was spin coated on the ITO substrate at 4000 rpm for 50 s and annealed at 160 °C for 20 min. Thereafter, the substrates were transferred to the glovebox until they cooled to room temperature. 65 µL of the perovskite precursor was deposited on PEDOT:PSS using the two-step process of 1000 rpm for 10 s and 4000 rpm for 50 s. 600 µL of CB was dropped onto the substrate during the second spin-coating step. Then, the as-prepared perovskite films were annealed for 10 min at 100 °C. For the post-treatment, 100 µL of EDAl₂ solution was spin-coated onto the perovskite films at 4000 rpm for 30 s, followed by annealing at 100 °C for 5 min. Then, the substrates were transferred into a vacuum chamber for evaporating 20 nm of C₆₀ and 5 nm of BCP at a vacuum pressure of 3×10⁻⁴ Pa. Finally, 100 nm of Cu was evaporated with a mask.

For Pb-Sn devices (University of Surrey), PEDOT:PSS was spin coated on the ITO substrate at 5000 rpm for 40 s and annealed at 150 °C for 30 min in air followed by 140 °C for 45 min inside the glove box. 50 µL of the perovskite precursor was deposited on PEDOT:PSS using the two-step process of 1000 rpm for 10 s and 4000 rpm for 50 s. 200 µL of CB was dropped onto the substrate during the second spin-coating step. Then, the as-prepared perovskite films were annealed for 10 min at 100 °C. Then, the substrates were transferred into a vacuum chamber for evaporating 20 nm of C₆₀ and 7 nm of BCP at a vacuum pressure of less than 4×10⁻⁴ Pa. Finally, 100 nm of Ag was evaporated with a mask.

Device characterisation

Current (*I*) - Voltage (*V*) characteristics of the devices were evaluated using an Enlitech SS-X100R (Class AAA) solar simulator with a Keithley 2450 source measure unit with a scan rate of 0.2 V s⁻¹. The calibration of the simulator was carried out using a KG-5 filtered Si diode. A mask with 0.09 cm² aperture area was used to define the active area of the device. All devices were measured without any encapsulation in a N₂ glovebox at a light intensity of 100 mW cm⁻² (AM1.5G). The light intensity was calibrated using a silicon reference cell (KG5 filter, Fraunhofer ISE). No preconditioning of the cells was carried out.

For Pb-Sn devices (Sichuan University): *J-V* curves were obtained by using a Keysight Technologies B2901A source meter with a solar simulator (SS-F5, Enlitech) with a scan rate of 0.08 V s⁻¹, dwell time of 100 ms and scan range of -0.1 to 0.9 V under simulated AM 1.5 G solar illumination at 100 mW cm⁻². The light intensity was calibrated using a silicon reference cell (SRC-00331, Enlitech). The devices were measured using a black shadow mask with an aperture area of 0.0576 cm². No preconditioning of the cells was carried out.

For Pb-Sn devices (University of Surrey): *J-V* curves were obtained by using a Keysight Technologies B2901A source meter with a solar simulator (SS-F5, Enlitech) with a scan rate of 0.08 V s⁻¹, dwell time of 100 ms and scan range of -0.1 to 1 V under simulated AM 1.5 G solar illumination at 100 mW cm⁻². The light intensity was calibrated using a silicon reference cell (Newport). The devices were measured using a shadow mask with an aperture area of 0.09 cm². Measurements were carried out under ambient conditions without any encapsulation at a temperature of ~25 °C and a relative humidity of 30-35%. No preconditioning of the cells was carried out.

EQE measurements of the MAPI devices were carried out using a Bentham PVE300 system. All measurements were carried out under ambient conditions. Reflectance measurements were carried out using an integrating sphere incorporated to the same system. For Pb-Sn devices, EQE measurement was taken with an Enlitech QE-R system. All devices were measured without any encapsulation under ambient conditions at a temperature of ~25 °C and a relative humidity of 30-35%.

TPC and IMPS measurements were conducted using Fluxim's PAIOS all-in-one test setup with a pixel area of 4.5 mm².

EIS was measured using CHI600E. The Nyquist plots of the EIS were measured at the frequency range of 1 MHz to 1 Hz with a bias voltage of 0.8 V and amplitude 5 mV.

Stability testing: No encapsulation was used for stability testing. For ISOS-D-1 testing, samples were stored in the dark under ambient conditions with a relative humidity of ~65%. For ISOS-D11 testing, samples were stored in the dark in a N₂ glove box. For Pb-Sn devices: MPP tracking was performed with a multicolour LED solar simulator (Guangzhou Cryscos Equipment Co. Ltd) under AM1.5G, 100 mW cm⁻².

Thin film characterisation

SEM images were acquired using a TESCAN MIRA2 SEM under an accelerating voltage of 5 kV or a ThermoFischer Scientific Apreo under an accelerating voltage of 2 kV. Grain size analysis was carried out using ImageJ software.

GIXRD measurements were taken using a Panalytical X'pert Pro diffractometer using a GI stage with a Cu Kα1 X-ray source driven at 40 kV.

XPS spectra were obtained on a Thermo Fisher Scientific Instruments K-Alpha+ spectrometer consisting of a monochromated Al Kα X-ray source (hν = 1486.6 eV) with a spot size of ~400 µm radius. A pass energy of 200 eV was used for acquisition of the survey spectra. For obtaining high resolution core level spectra, a pass energy of 50 eV was used for all elements. For correction of possible charging effects that can occur during acquisition, the obtained spectra were charge

referenced against the C1s peak (285 eV). Fitting of the spectra was carried out using the manufacturers' Advantage software.

UPS was taken on a Thermo Scientific Theta Probe system using He-I excitation (21.22 eV). A -9 V bias was applied during measurement to the samples prepared on ITO substrates. The binding energy scale was calibrated to the Fermi edge of clean Au at 0 eV.

For TAS measurements, all samples were given glass-lid encapsulation. The output of a Ti:sapphire amplifier system (Spectra Physics Solstice Ace) operating at 1 kHz and generating ~100 fs pulses was used as femtosecond laser source. **For measurements, a 400 nm pump with fluence of ~2.3 $\mu\text{J cm}^{-2}$ (incident from the perovskite layer), corresponding to an injected carrier density of $1.06 \times 10^{17} \text{ cm}^{-3}$ was used.** The 400 nm pump pulses were created by sending the 800-nm fundamental beam of the Solstice Ace through a second harmonic generating (SHG) beta barium borate (BBO) crystal of 1 mm thickness (Eksma Optics). The pump was blocked by a chopper wheel rotating at 500 Hz. The broadband white light was provided by the Disco (Leukos Laser, STM-2-UV) and the pump-probe decay was controlled electronically. The white light was split into two identical beams (probe and reference) by a 50/50 beamsplitter. The reference beam passing through the sample did not interact with the pump, which allows for correcting for any shot-to-shot fluctuations in the probe that would otherwise greatly increase the structured noise in the experiments. Based on this arrangement, small signals with $\Delta T/T \sim 10^{-5}$ could be measured. The transmitted probe and reference pulses were collected with a silicon dual-line array detector (Hamamatsu S8381-1024Q, spectrograph: Andor Shamrock SR-303i-B) driven and read out by a custom-built board (Stresing Entwicklungsbüro). The recombination rate constants were calculated by fitting the decay characteristics using the general recombination rate equation⁵²:

$$-\frac{dn}{dt} = k_1 n + k_2 n^2 + k_3 n^3$$

where k_1 is the Shockley–Read–Hall monomolecular recombination rate (corresponding to traps), k_2 is the radiative bimolecular recombination rate, and k_3 is the Auger recombination rate. In this study, we focus on low carrier density region where k_1 dominates the recombination process and extract the k_1 rate constant.

ToF-SIMS (NREL): An ION-TOF ToF-SIMS V spectrometer was utilised for depth profiling using methods covered in detail in previous reports⁵³. Analysis was completed utilising a 3-lens 30 keV BiMn primary ion gun. High mass resolution depth profiles were completed with a 30 KeV Bi₃⁺ primary ion beam, (0.8pA pulsed beam current), a 50x50 μm area was analysed with a 128:128 primary beam raster. Sputter depth profiling was accomplished with 600 eV Cesium ion beam (2.6 nA sputter current) with a raster of 150 x 150 μm^2 .

ToF-SIMS (NPL): Samples were prepared following the configuration of glass/ITO/PEDOT:PSS/MAPI (with or without SCN⁻ additives)/polymethylmethacrylate (PMMA)/Gorilla 2-part epoxy (5 min) /glass. The glass/ITO used was 15 mm x 15 mm while a cover glass slips cut to the dimensions of

25 mm x 25 mm was used on the top. PMMA 950k A6 from Microchem was coated on the perovskite to realise a 200 nm thick film (inside a N₂ glove box) and was left to cure overnight. The Gorilla epoxy was then carefully coated on the samples and the glass cover slips were carefully placed on top and left to cure for at least 24 hours inside a N₂ glove box. The samples were then subjected to impact by dropping from ~1 m.

UV-VIS spectra of PEDOT:PSS + organohalide thin films were acquired using a Cary 5000 spectrophotometer while UV-VIS spectra of PEDOT:PSS dispersions incorporating organohalides were obtained using a Shimadzu 2600 spectrophotometer with integrating sphere accessory.

EPR spectroscopy was collected on a Bruker Magnettech ESR 5000 at a temperature of 29 °C, averaged over 3 sweeps using a Microwave power of 4 MW.

¹H and ¹³C NMR were collected on a Bruker 400 MHz NMR spectrometer using CDCl₃ as a deuterated solvent. Data was collected using top spin data acquisition software. Data chemical shift was corrected to the 7.26 ppm peak of CDCl₃ using Mestrenova software.

Attenuated total reflectance Fourier transform infrared spectroscopy (ATR-FTIR) was measured using a Carey 630 FTIR benchtop spectrometer using a Diamond ATR configuration with a KBr window.

MS measurements were acquired by direct injection using Atmospheric Pressure Chemical Ionisation (APCI) ion source connected to the Thermo Fisher Q-Exactive Orbitrap mass spectrometer using capillary temperature of 320 °C and discharge voltage was set between 3.5 and 4.0 kV. The mass spectrometer operated in positive and negative ionisation mode.

Cyclic voltammetry was collected using an electrolyte of 0.15 M n-butylammonium hexafluorophosphate in 50 ml anhydrous acetonitrile (99.8+%), dried over molecular sieves. CV data was collected on a Gamry 1010E potentiostat using a AgNO₃ reference electrode (connected through a ceramic luggin capillary), Pt counter and Pt working electrode (2 mm diameter). Samples were measured at a scan rate of 100 mV s⁻¹. To the electrolyte solution, 2 mM of sample was dissolved before the solution was degassed in N₂. In experiments where -SCN was added to I₂ or vice versa, 2 scan cycles were collected before the secondary compound was added, degassed, and remeasured.

Optical images of PEDOT:PSS incorporating organohalides was obtained using a Keyence VHX-5000 digital microscope. DFT and TD-DFT calculations were processed using Gaussian 16 Software. For each calculation the geometry was optimised using a hybrid 6-311G(d,p) [H,C,N,O,S], LANL2DZ [I] basis set and unrestricted either unrestricted B3LYP and ω B97XD functionals with a Grimme's dispersion correction modifier (GD3).

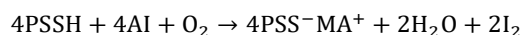
Calculation of oscillator strength of PEDOT 12-monomer chain: To calculate the oscillator strength of the PEDOT under different oxidation states, first the PEDOT geometry was optimised at between 0 and 50 % oxidation (PEDOT0 – PEDOT6) using uB3LYP with Grimme's D2 dispersion or ω B97XD functionals. From the calculated ground state geometries, the energies for different

spin multiplicities were calculated allowing for the lowest energy spin multiplicity to be determined. From this lowest energy predicted spin multiplicity, TD-DFT was used to predict electronic transitions and give a prediction of the oscillator strength across visible wavelengths.

Calculation of Binding/Dissociation/Reaction Free Energies: To calculate the gas-phase ion binding energies, the molecular geometries were first optimised before performing frequency calculations to obtain the zero-point energy corrected free energies at a default temperature of 298.15 K.

$$\Delta G_g^o = \Delta G_g(\text{PEDOT}:I_3) + \Delta G_g(A:\text{PSS}) - [\Delta G_g(\text{PEDOT}:\text{PSS}) + \Delta G_g(A:I_3)]$$

For this work, only gas phase calculations were performed owing to the inability of SMD models to accurately describe the hydration environment within PSS micelles. This is particularly important owing to the effects of polar mediums to screen the ionic charges. The predicted thermodynamic quantities were calculated using the equations below. In this work, a PSS monomer was used with a PEDOT trimer PEDOT_3^{+1} :



$$\Delta G_r = \left(\frac{1}{4}\right)[4\Delta G(A:\text{SS}_1) + 2\Delta G(\text{H}_2\text{O}) + 2\Delta G(\text{I}_2) - [4\Delta G(\text{SS}_1\text{H}) + \Delta G(\text{O}_2) + 4\Delta G(\text{AI})]]$$

$$\Delta G_b(\text{PSS}:A) = [\Delta G(\text{PSS}:A) - [\Delta G(\text{PSS}_1) + \Delta G(A)]]$$

$$G_a(\text{PSS}:\text{PEDOT}/A) = [\Delta G(\text{SS}_1:A) + \Delta G(\text{PEDOT}_3^{+1}) - [\Delta G(\text{PEDOT}:\text{SS}_1) + \Delta G(A^{+1})]]$$

$$\Delta G_b(A:I) = [\Delta G(A:I) - [\Delta G(A^+) + \Delta G(I^-)]]$$

Author contributions

Conceptualisation: W. H. K. P., T. W., K. D. G. I. J., S. A. H., S. R. P. S.

Methodology: W. H. K. P., T. W., Y. X., D. W. Z., K. D. G. I. J., S. A. H., S. R. P. S.

Investigation: W. H. K. P., T. W., Y. X., J. Z., Y. ZHOU, G. F. T., M. G. M., S. P. H., S. J., L. D., S. S., T. J. M., S. J. H., Y. ZHAO, K. D. G. I. J.

Funding acquisition: S. P. H., S. J., T. J. M., Y. ZHAO, S. D. S., D. W. Z., W. Z., K. D. G. I. J., S. A. H., S. R. P. S.

Supervision: T. J. M., S. D. S., G. F. T., D. W. Z., W. Z., K. D. G. I. J., S. A. H., S. R. P. S.

Writing – original draft: W. H. K. P., T. W., Y. X., K. D. G. I. J., S. A. H., S. R. P. S.

Writing – review & editing: W. H. K. P., T. W., Y. X., J. Z., Y. ZHOU, G. F. T., M. G. M., S. P. H., S. J., L. D., S. S., T. J. M., S. J. H., Y. ZHAO, S. D. S., D. W. Z., W. Z., K. D. G. I. J., S. A. H., S. R. P. S.

Conflicts of interest

S. D. S. is a co-founder of Swift Solar Inc. S. J. is an employee at Fluxim AG.

Data availability

Data for this article are available at University of Surrey Open Research Repository at <https://openresearch.surrey.ac.uk/esploro/>.

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