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Innovative PIN-type perovskite solar cells with 17% efficiency: processing and characterization†

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To develop high-performance tandem cells, it seems necessary to optimize semi-transparent PSC single junctions with a PIN-type architecture. In this article, the development of this PIN-type architecture by the optimization of the Hole Transport Layer (HTL) is proposed. Firstly a study of three HTLs with different surface properties such as PEDOT:PSS, PTAA and TFB was carried out, to understand their influence on the deposition conditions and on the final properties of the deposited perovskite layer. Indeed, the use of layers such as PTAA and TFB represents a real alternative, already mentioned as very promising in NIP and PIN type architectures. But unlike PEDOT:PSS, these hydrophobic layers disrupt the deposition of the perovskite layer. Once the material was chosen the HTL, here TFB, which has real originality, the optimization of the process was carried out. The latter based on the evaluation of the antisolvent ejection time and the drying time allowed a drastic increase in the PCE from 10% to 17%. Another important result of this study is the methodology used for this optimization. Indeed, the main marker used is the observation of defects in the perovskite layer at the macroscopic and microscopic scales. This optimization of the perovskite layer morphology leads to a significant increase in performance. This approach will be very useful to the community for a possible optimization of other devices.

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Introduction

Perovskite materials have been widely developed in the photovoltaic (PV) field for a few years.¹ These materials, processed at low temperatures, present exceptional optoelectronic properties.² A wide bandgap range (from 1.17 to 3.11 eV)³ combined with an excellent mobility⁴ and very good absorption properties⁵ make them the materials of choice for PV tandem architectures,^{6,7} in particular with silicon solar cells. The main objective is to exceed the Shockley-Queisser efficiency limit.⁸

Two different architectures can be used for perovskite solar cells (PSCs), depending on the order of the layer deposition, with a hole or an electron transport layer (HTL or ETL), underneath the perovskite. The NIP-type architecture is the most common because it was easier to process at the first stage of perovskite PV and led to the first PV yields with a TiO₂⁹ based ETL. The first PIN-type structures appeared for a second time with PEDOT:PSS¹⁰ as the HTL. Today, both architectures achieved high efficiencies in the range of 20–22%,^{11,12} but the NIP-type architecture has led to the record efficiencies for single

junction cells.¹¹ In comparison, somewhat lower efficiencies are consistently obtained for PIN-type architectures.¹³

In the case of a monolithic tandem application combining perovskite and silicon solar cells, the highest efficiencies are obtained with perovskite cells in the PIN-type architecture.¹⁴ The latter allows the reduction of parasitic absorption at the front side of the device, compared to the NIP-type architecture.¹⁵ A thin (despite absorbing) N-type interface within the PIN-type architecture (e.g. C₆₀ layer) appears indeed to be optically less detrimental than the thick highly absorbing HTL typically employed in the standard one (e.g. spiro-OMeTAD).¹⁴ To develop high-performance tandem cells, it therefore seems to be necessary to optimize semi-transparent PSC single junctions with the PIN-type architecture. This type of cell is gradually more studied, and the efficiencies increased accordingly up to 17%.^{16–21} The performances depend essentially on the materials used as active and interfacial layers. The main weakness of these architectures is the lower value of V_{oc} , due to the use of non-suitable N- and P-type interfacial layers. This prevents the optimal operation of the perovskite layer and favors the radiative recombination.²²

The current work focuses on the development of innovative PIN-type perovskite solar cells. For this, we propose to use mixed perovskites, which present several advantages over the most common form MAPbI₃ (with MA = methylammonium). The multiple cation/anion perovskites lead to higher yields, and better stability and reproducibility.^{23,24} In addition,

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controlling the stoichiometry in mixed perovskites (especially the iodide/bromide ratio) allows adjustment of the bandgap energy of the material in view of a tandem application.²⁵ An $FA_{1-y}Cs_yPb(I_{1-x}Br_x)_3$ (with FA = formamidinium) perovskite has recently been developed with a formulation specifically optimized for the development of low-temperature perovskite devices in the NIP-type architecture.²⁶ The optimum ratios were found to be 88% for I/Br and 94% for FA/Cs. This previous work suggested good potential of this formulation for both efficiency and stability. To improve the performances of these devices and in particular their V_{oc} , it is also proposed to study several HTLs with very different surface properties, PEDOT:PSS, PTAA and TFB, in order also to understand their influence on the deposition and on the properties of the perovskite layer deposited afterwards. Indeed, the use of layers such as PTAA and TFB represents a real alternative, already mentioned as very promising in both NIP- and PIN-type architectures.^{27–29} But unlike PEDOT:PSS, these hydrophobic layers disturb the deposition of the perovskite layer. Several approaches are proposed in the literature to improve the wettability of the perovskite on these hydrophobic layers,^{30,31} but it is preferable to preserve the nature of the materials, and quite possible to optimize the quality of the perovskite layer by solely varying the processing conditions. We propose here double optimization of the material and process to maximize the PCE of the devices. The main originality concerning the material concerns the use of TFB as the HTL. To vary the processing, a combined optimization of the antisolvent ejection time and the drying time before annealing was performed. These two steps permitted a drastic increase in the PCE from 10% to 17%. In addition, comprehensive elements are provided to broaden the experimental conclusions, and unravel new process/micro-structure/properties of the perovskite. This will be very useful to the community for a possible optimization of other devices.

Experimental part

The preparation of perovskite solutions, fabrication of solar devices and their photovoltaic characterization were carried out in a glove box containing less than 1 ppm oxygen and moisture.

Materials

All anhydrous solvents (dimethylformamide (DMF), dimethylsulfoxide (DMSO), chlorobenzene (CB), *o*-xylene, ethyl acetate (EA), diethyl ether (DE) and isopropanol (IPA)) were purchased from Sigma Aldrich, as were $PbBr_2$, CsI and PTAA. PbI_2 was purchased from Tokyo Chemical Industry and formamidinium iodide (FAI) from Greatcell Solar Materials. PEDOT:PSS and TFB were respectively purchased from Heraeus (ref. Clevios P-VPAI-4083) and American Dye Source (ref. ADS259BE). Np-SnO₂ (a dispersion of nanoparticles at 2.5 wt% in butanols (20.7 mg mL⁻¹)) was purchased from Avantama (ref. N30) and PC₆₀BM from Merck (ref. Lisicon PV-A600). Silver used for metallic contacts was purchased from Neyco, and the ITO coated substrates (square resistivity of 7 Ω/\square) from Visiontek (ref. ITOGLASS 07P).

Solar cell preparation

The perovskite precursor solution of $FACsPbBrI$ at 0.87 or 1.15 mol L⁻¹ (for lead) was made in a glovebox from the three following intermediate solutions: a perovskite precursor solution of $FAPbI_3$ at 0.87 or 1.15 mol L⁻¹ (lead) prepared with FAI:PbI₂ at a 1.05:1 molar ratio in DMF:DMSO at 4:1 in volume, stirred one night at 40 °C; a 0.87 or 1.15 mol L⁻¹ $PbBr_2$ solution in DMF:DMSO at 4:1 in volume, also stirred one night at 40 °C; and a 1.5 mol L⁻¹ CsI solution in DMSO, stirred at room-temperature. The three intermediate solutions were mixed for about 30 min before use (during which the final solution was cooled down to room temperature).

ITO coated substrates were sonicated sequentially in acetone, isopropanol and eventually deionized water. Before the first deposition layer, these substrates were treated with UV-ozone for 30 min. For PTAA as the HTL, the layer was deposited by spin-coating in a glovebox from a solution at 2 mg mL⁻¹ in chlorobenzene, filtered using a 0.45 μ m PTFE filter, and then dried at 110 °C for 10 min. For PEDOT:PSS as the HTL, the layer was deposited by spin-coating in air from a commercial solution, filtered using a 0.45 μ m GMF w/GMF filter, and then dried at 180 °C for 1 hour in an oven. For TFB as the HTL, a very thin layer, from a solution at 0.6 mg mL⁻¹ in *o*-xylene, was deposited by spin-coating in a glovebox and then dried at 150 °C for 10 min. In order to suppress possible macroscopic defaults, a surface treatment by DMF was applied onto PTAA and TFB right before the perovskite layer deposition. The previously described perovskite precursor solution, filtered using a PTFE filter of 0.45 μ m, is then spread and spin-coated, where XX ∈ {10, 20, 30, 40, 50, 60} s. 150 μ L of ethyl acetate are used as antisolvent and poured on the perovskite layer 5 s before the end of the spin-coating program ($t_{ASE} = XX - 5$ s). Afterwards, the resulting perovskite layer is dried at room temperature in a glovebox for 0 or 20 min (t_{DBA}), and finally annealed at 80 °C for 1 h. After the samples were cooled down, around 20 nm of PCBM are then deposited by spin-coating, from a 10 mg mL⁻¹ concentrated solution in chlorobenzene, filtered with a PTFE filter of 0.45 μ m. No drying or annealing was performed after PCBM deposition. 50 μ L of np-SnO₂ at 2.5 wt% in butanols, previously filtered with a PP w/GMF filter of 0.45 μ m, was finally deposited by dynamic spin-coating by means of an electronic pipette, leading to a thickness of about 20 nm. Dynamic spin-coating was chosen for minimizing the contact time between the solvent of np-SnO₂ (butanols) and the perovskite. No drying or annealing was performed after np-SnO₂ deposition. 100 nm of silver were deposited by thermal evaporation under vacuum to complete the setup, delimitating an active area of 0.34 cm².

Characterization

Photocurrent–voltage (*J*–*V*) curves were generated in a glovebox with a solar simulator (Newport Oriel SP94043), calibrated using a silicon reference solar cell at a light intensity of 100 mW cm⁻². A voltage sweep from 2.5 V to -0.2 V (reverse scan) with a step of 0.1 V and 20 ms of acquisition time for each data point was the parameter used for the *J*–*V* measurements. They were performed every minute for at least 15 min under



illumination for each cell to reach stabilization of their efficiency. All the UV-visible characterization studies were performed in air using a UV-visible spectrophotometer (UV-2600, Shimadzu, Japan) with an integration sphere. X-ray diffraction (XRD) patterns were collected in air using a Malvern PANalytical apparatus (Aeris version) equipped with an X-ray source ($\lambda(K_\alpha) = 0.178901$ nm), a divergence slit of $\frac{1}{2}$ °, an angle aperture of 0.04 rad and a 23 mm beam mask. Photoluminescence (PL) spectra were obtained with a fluorescence spectrophotometer (Hitachi F-4500 FL, Hitachi, Japan) with an excitation wavelength of 467 nm (corresponding to the highest intensity response) and a speed sweep of 1200 nm min $^{-1}$ (an excitation slit of 10.0 nm, an emission slit of 20.0 nm and a photomultiplier tube voltage of 400 V). A set of two filters was used to remove the 2θ harmonics and instrumental artifacts: a low-pass filter at 575 nm upstream and a high-pass filter at 600 nm downstream. For this experiment, the samples were only composed of the perovskite layer and the underlying HTL deposited on the glass/ITO substrate. The thicknesses were estimated with a profilometer (D500, KLA Tencor, USA) after scratching the upper layers, with the following parameters: a displacement speed of 0.05 mm s $^{-1}$, an equivalent strength of 1 mg, and the height range of 10 μ m. Work functions were acquired in air at room-temperature and in the dark using a Kelvin probe from KP Technology equipped with a digital control setup managed by Kelvin probe measurement system version DELTA software. During acquisitions, the oscillation frequency of the tip was set to 96 Hz, with an amplitude of 38 V, an electromotive force of ± 7 V, a trigger probe of 114 counts, and with an electrostatic force gradient of 300 N m $^{-1}$. For samples with perovskite, 300 consecutive measurements in one location were realized. Copper tape was used to guarantee the electrical contact between samples and the sample holder of the Kelvin probe. The tip work function was deduced by using a thermally evaporated gold layer of 100 nm. To establish the experimental standard deviation and the average work function of gold in air (attributed to -4.7 eV), 32 100 consecutive measurements were achieved at three different locations of the gold layer before and after samples' measurement campaigns. Images from optical microscopy were acquired with a Leica DMLM microscope and a Leica

DFC420 camera, in transmission for orange-like images and in reflection for blue-like images with a $\times 10$ magnification. Picture analysis was performed with ImageJ. Analyses to evaluate the microscopic pinholes' surface percentage were conducted with ImageJ software on 3 different pictures for each condition: pictures are set in the greyscale image at first, then a threshold is applied to settle and distinguish pinholes in the layer, and the binarization function is lastly applied to the image before extracting the surface area corresponding to the pinholes. SEM images were acquired using a Nova NanoSEM 630 apparatus equipped with secondary electron detectors, with a working distance of around 5 mm and a voltage of 7 kV.

Results and discussion

The present study focuses on the development of PIN solar cells, with a specific goal to upgrade the perovskite layer on an innovative and promising HTL. Fig. 1a presents the architecture of the studied cell. A mixed perovskite with two cations and two anions serves as the active layer. Its chemical structure $(\text{CH}(\text{NH}_2)_2)_{(1-y)}\text{Cs}_y\text{Pb}(\text{I}_{(1-x)}\text{Br}_x)_3$ will be simplified as (FACsPbIBr). This complex stoichiometry results from the iterative optimization from different studies to favor yields, stability and robustness. 23,24 For instance, as the energy bandgap depends on the molar ratios of the different species, the stoichiometry may be used as a means to customize the materials and match the desired absorption. This may help to reach a variety of specific needs. For tandem 25 applications, the iodine to bromine ratio is commonly modified to optimize the bandgap of the subcell to be combined with the silicon one. The addition of cesium (Cs) in the structure allows improvement of the stability of the crystalline structure of the brominated and iodinated perovskites. 33 The presence of Cs hinders the formation of the photo-inactive delta phase ($\delta\text{-FAPbI}_3$). 26,33 To facilitate the processing and improve the stability, 34 the perovskite structure without methylammonium may be formulated with formamidinium and cesium. The performance (higher than 12%) and stability recently obtained with these types of perovskite in NIP-type architectures are satisfactory. 26

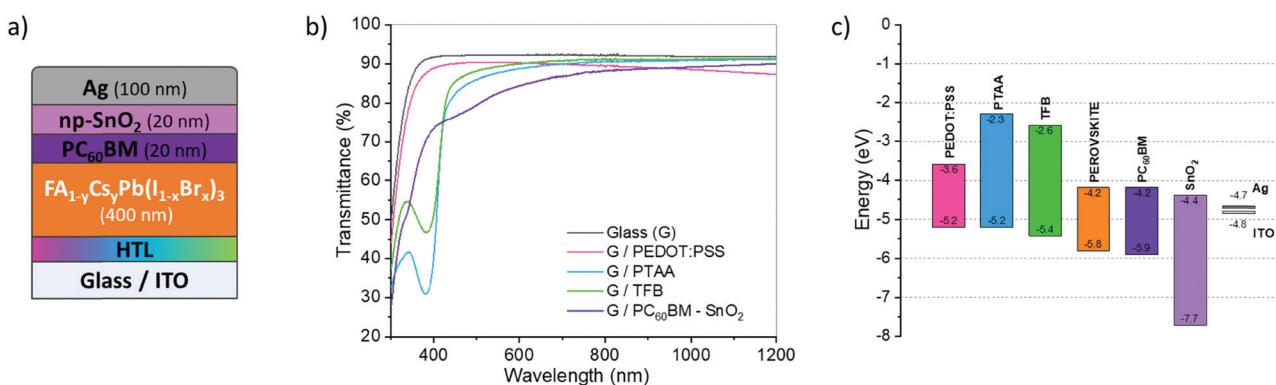


Fig. 1 Structure and properties of the solar cells developed in the present study. (a) PIN-type architecture on the ITO coated glass substrate; (b) optical properties in transmission of studied HTLs and ETL; and (c) energy levels of the constitutive materials (data issued from the literature).



The PCBM/BCP layer is intensively used for its efficiency in opaque PIN-type cells,^{22,35,36} and alternatives are also being investigated for tandem application.^{7,37} PCBM combines a good stability with a strong ability to extract electrons from the perovskite layer,^{38,39} Fig. 1c. It may safely be kept in the final architecture. In contrast, the BCP layer should be replaced by a more robust one. The substitution material ideally combines the roles of the buffer and hole blocking layer,³⁵ in addition to the ability to develop a good interface with the metal (low resistance and ohmic behavior). We decided to focus on SnO₂ that has already been identified among the best buffer layers and ETL.⁴⁰ Here a SnO₂ layer based on nanoparticles has been used. A preliminary study was thus carried out to stack PCBM and SnO₂ in order to optimize the two thicknesses with regard to the overall PCE (presented in Fig. S1, ESI[†]). This means that the best optical and physical properties compromise for a semi-transparent application.

Different HTLs were investigated including PEDOT:PSS, PTAA and TFB. For PIN-type architectures, PEDOT:PSS's hydrophilic character and its excellent wettability for the perovskite precursor solutions facilitate the processing of the active layer.^{41,42} Unfortunately, the resulting cells present low V_{oc} , due to their greater ability to form recombinant traps at the HTL/perovskite interface.⁴³ Other alternative materials have been tested, among which PTAA was considered a most promising candidate. It combines excellent optical properties and work function but its hydrophobicity limits the wettability of the perovskite layer. TFB was also considered because the latter presents an even better energy level alignment with perovskite, due to its lower HOMO level value close to 5.4 eV. This should induce an improvement of the V_{oc} , Fig. 1c. TFB also shows better electronic mobility than PTAA, while keeping similar and excellent optical properties,^{30,44} Fig. 1b. However, TFB is hydrophobic in

nature such as PTAA, and a dedicated protocol should be developed for these HTLs to obtain a homogeneous and controlled morphology of the perovskite layer.

To improve the wettability of the perovskite on the highly hydrophobic organic HTL, prior surface treatment with DMF allows avoiding macroscopic holes (as observable in Fig. 2b). The use of an anti-solvent should also be necessary to obtain a good quality and covering layer.⁴⁵ In the case of a NIP architecture, the perovskite deposition is done on a rather hydrophilic layer (TiO₂, SnO₂) and chlorobenzene is generally used as an anti-solvent. In the case of deposition on hydrophobic layers (PTAA and TFB), the nature of the anti-solvent largely affects the quality of the deposited perovskite layer. As shown in Fig. S2 (ESI[†]) and Fig. 2a, the use of diethyl ether (DE) here leads to a macroscopically non-compliant perovskite layer, and the use of the common chlorobenzene (CB) leads to large holes of about 30 μ m without perovskite and the HTL beneath, indicating the dissolution of the latter, as also observed in the literature³¹ and confirmed by the test presented in Fig. S2 (ESI[†]). Although the use of other anti-solvents leads to more homogeneous deposits, microscopic observations show much smaller pinhole diameters (less than 15 μ m, and without any HTL dissolution) in the case of ethyl acetate (EA) and isopropanol (IPA). Such an effect of the nature of the anti-solvent has already been observed in the literature, but generally at a lower scale.^{46,47} IPA and EA appear here to be the best candidates. Our choice turned to EA because of literature results that show higher yields in single junctions⁴⁸ and tandem cells,¹⁵ in contrast to IPA which is known to easily solubilize the organic compound of the perovskite material.⁴⁹ EA is by the way reported to offer a better moisture resistance during perovskite fabrication.⁵⁰

Without changing the nature of the materials, the quality of the deposited layer may be largely altered by varying the processing conditions. For instance, adjusting the anti-solvent

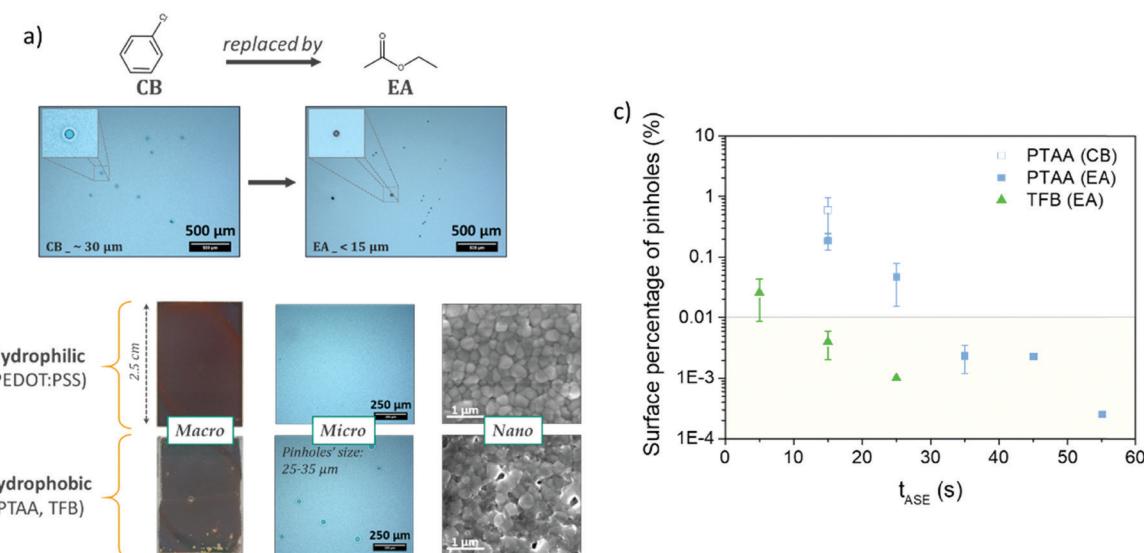


Fig. 2 (a) Impact of anti-solvent's nature (chlorobenzene vs. ethyl acetate) on the micro-scale pinholes in the perovskite layer (examples on the PTAA layer, $t_{ASE} = 15$ s); (b) impact of the HTL nature (hydrophilic vs. hydrophobic) on the perovskite layer at macro-, micro- and nano-scales (examples with EA, $t_{ASE} = 15$ s); and (c) surface pinhole proportion (determined by optical microscopy, see Fig. S3, ESI[†]) as a function of anti-solvent ejection time (t_{ASE}) and HTL nature.



ejection time (t_{ASE}) seems interesting:⁵¹⁻⁵⁴ the principle is presented in Fig. 3. The latter has hardly been used, mainly for the mono-cation perovskite (MAPbI_3). The authors concluded that the t_{ASE} influences the absorbance and crystallinity of the perovskite. The optimisation of the t_{ASE} can thereby significantly improve the performances of the solar cells, although in a phenomenological manner exclusive to each specific system. Y. Bai *et al.* revealed an optimum in the PCE yields as a function of the t_{ASE} for different perovskite formulations. They proposed a much longer optimal time (40 s⁵¹) than previous studies (10 s⁵³ or even 5 s⁵²). As presented in Fig. 2b, with an anti-solvent ejection time ≤ 15 s, micro-scale holes and nano-scale defects are visible in the case of hydrophobic HTLs like PTAA and TFB. Such defects are not observable in the case of the PEDOT:PSS-based HTL, confirming the role played by the nature of the HTL surface. As suggested in the literature, we varied the ejection time between 5 and 55 s and observed the obtained perovskite layers with an optical microscope (Fig. S4, ESI†). After image analysis (ImageJ), the relative number of micro-pinholes within the surface was calculated and plotted as a function of the anti-solvent ejection time, Fig. 2c. For the two hydrophobic layers studied (PTAA and TFB), this proportion of defects decreases significantly with the ejection time, together with the average diameter of the holes. This improvement is much faster with TFB, for which no microscopic defects remain after only 15 s, compared to 35 s with PTAA.

To better understand the effect of this anti-solvent ejection time, the perovskite layer deposited on the three HTLs was analyzed by X-ray diffraction and UV-Visible absorption spectroscopy. The X-ray diffraction pattern of the active layer for t_{ASE} is shown in Fig. S5 (ESI†). The diffraction peak located at 2θ close to 12.7° is commonly attributed to the PbI_2 phase.⁵⁵ The presence of bromine in the formulation can lead to PbX_2 , generally detected at 2θ close to $13.1-13.2^\circ$.^{22,56} The presence of PbI_2 was further confirmed by 25.7° and 38.9° peaks characteristic of the second- and third-order of the diffraction pattern. The other peaks could be attributed to the perovskite $\text{FA}_{1-y}\text{Cs}_y\text{Pb}(\text{I}_{1-x}\text{Br}_x)_3$. Various crystal orientations are observed, mainly $(00l)$ for which three orders of diffraction are detected at 14.1° (001), 28.4° (002) and 43.1° (003). Other orientations

can also be detected (see details in the ESI†). The solution of perovskite precursors was formulated as $\text{FA}_{0.94}\text{Cs}_{0.06}\text{Pb}(\text{I}_{0.88}\text{Br}_{0.12})_3$. The formed perovskite phase will be referred to as FACsPbIBr in the following. It has a cubic structure like FAPbI_3 ,^{3,57} with peaks slightly shifted towards larger angles. This is due to the reduction of the lattice plan associated with the presence of the smaller bromine atoms in the formulation, as compared to iodine. Finally, a small amount of the photo-inactive and undesirable $\delta\text{-FAPbI}_3$ phase is detected at 2θ of $11.7-11.8^\circ$.⁵⁸

As expected, all the tested t_{ASE} promote the nucleation and growth of the same few crystal phases (FACsPbIBr , PbI_2 and $\delta\text{-FAPbI}_3$), Fig. S5 (ESI†). From a quantitative standpoint the proportions of pertinent phases such as FACsPbIBr , PbI_2 and $\delta\text{-FAPbI}_3$ were evaluated with the anti-solvent ejection time t_{ASE} , Fig. 4. For the three HTLs studied, the proportion of the FACsPbIBr phase decreases progressively with an increase in t_{ASE} from a value of 15 s. Concerning the variations of PbI_2 , the proportion of this phase follows another trend with a minimum located respectively around 35 s, 15 s and 25 s for respectively PTAA, PEDOT:PSS and TFB. As this compound can have a negative effect on the functioning of the cell, it seems necessary to choose deposition conditions that limit it to a maximum. It is therefore possible to define an indicator corresponding to the ratio of FACsPbIBr and PbI_2 phases to highlight optimal conditions for the preparation of the perovskite layer. The latter is also maximal for t_{ASE} equal to 35 s, 15 s and 25 s, respectively, for PTAA, PEDOT:PSS and TFB. We note for these conditions that the proportion of the $\delta\text{-FAPbI}_3$ phase is negligible and thus should have little consequence on the functional properties of the perovskite layer.

The optical properties were also investigated by UV-Visible spectroscopy. The raw data obtained are presented in Fig. S6 (ESI†) and the bandgaps, calculated from the Tauc plot method, in Fig. 4c. The optical properties are not much altered by t_{ASE} : negligible variations in the optical bandgap and in absorbance magnitude. The bandgaps are however slightly different depending on the HTL, respectively, 1.58 eV, 1.60 eV and 1.61 eV for TFB, PEDOT:PSS and PTAA.

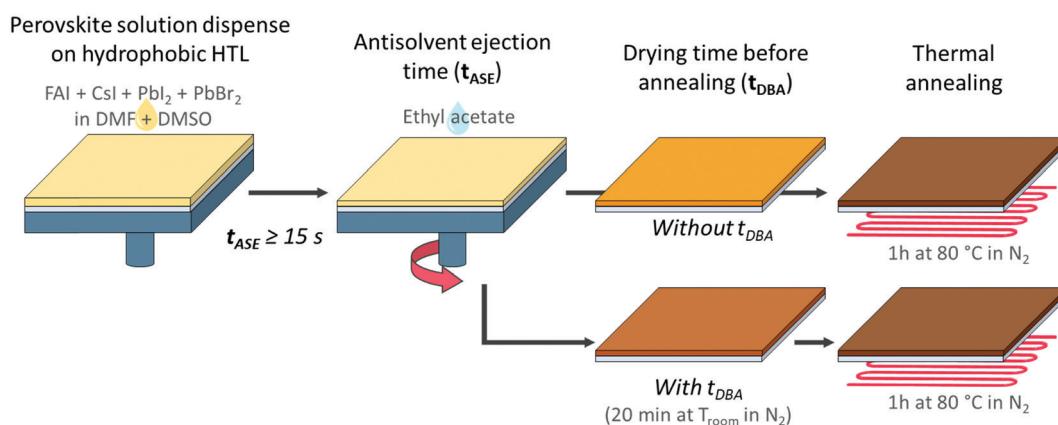


Fig. 3 Illustration of the deposition process for the perovskite layer on an HTL with the two studied parameters: the anti-solvent ejection time (t_{ASE}) and the drying time before annealing (t_{DBA}).



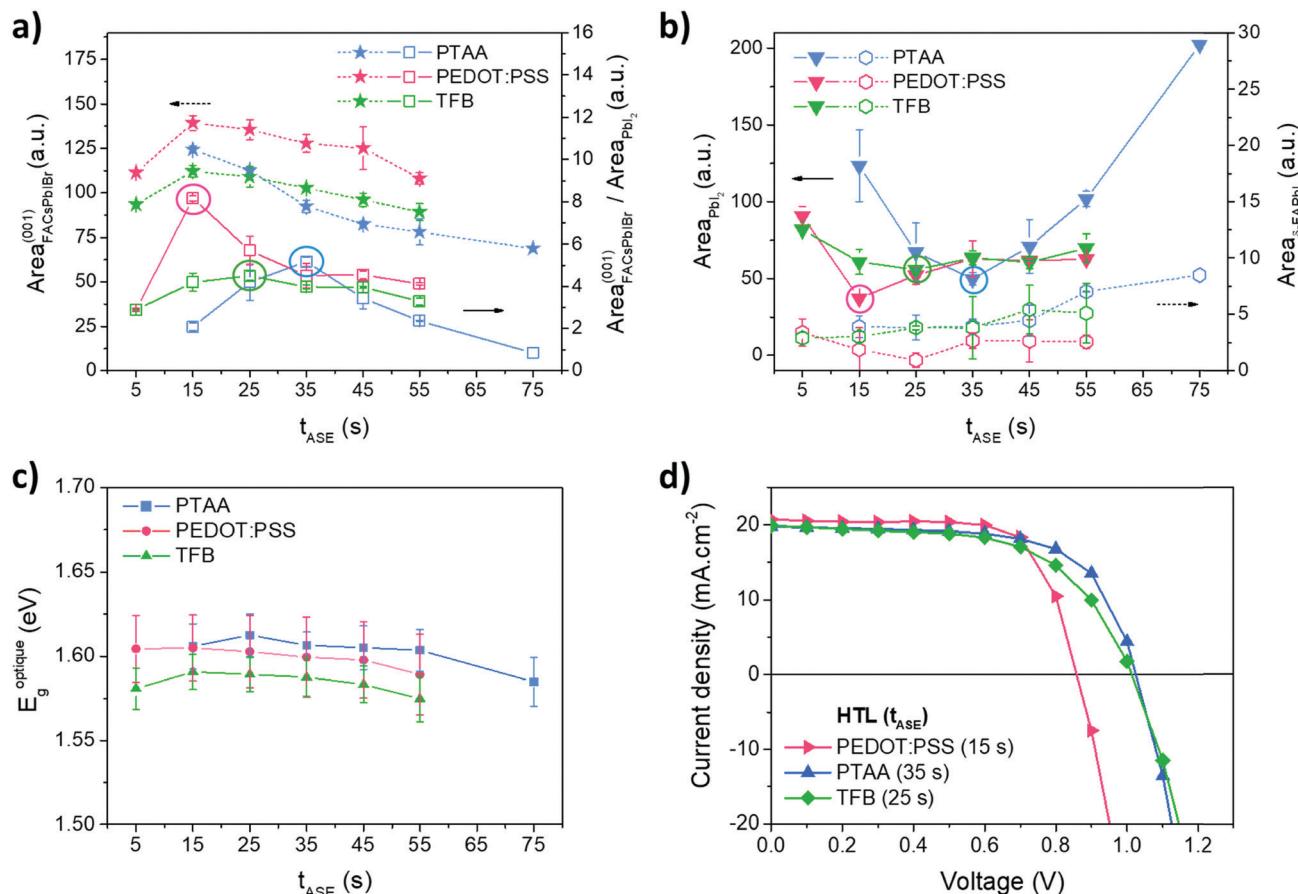


Fig. 4 For the three studied HTLs PTAA (blue), PEDOT:PSS (pink), and TFB (green): (a) and (b) extracted areas of the main diffraction peaks attributed to the FACsPbI_3 (along the orientation (001)), PbI_2 and $\delta\text{-FAPbI}_3$ phases, and the areas ratio of the FACsPbI_3 (along the orientation (001)) and PbI_2 phases as a function of t_{ASE} ; (c) optical bandgaps determined by the Tauc plot method as a function of t_{ASE} ; and (d) J - V measurements in reverse scan of solar cells based on PEDOT:PSS, PTAA and TFB with respectively a t_{ASE} of 15 s, 35 s and 25 s.

The optimal t_{ASE} determined by XRD analyses has been tested in solar cells: the corresponding J - V measurements and PV parameters are shown in Fig. 4d and Table 1. Efficiencies of around 12% are obtained for cells based on PEDOT:PSS and PTAA. The TFB leads to somewhat lower efficiencies, closer to 11%. The FF values are rather good, around 60% for PTAA and TFB and above 70% for PEDOT:PSS. The values of J_{sc} are similar regardless of the HTL studied and close to 20 mA cm^{-2} which is in agreement with the literature. The most interesting result

concerns the values of V_{oc} . As expected and demonstrated in the literature,⁵⁹ this is below 1 V for PEDOT:PSS, while an increase towards 1 V is obtained for the other two HTLs. The work function of PEDOT:PSS probably does not match well with that of the perovskite absorber.⁶⁰ However these values remain a little low compared to those obtained in the literature,^{61–63} and it seems that our systems can still be improved.

It was further decided to focus the rest of the study on a single HTL. TFB was chosen for its promising character

Table 1 PV parameters corresponding to J - V measurements in reverse scan presented in Fig. 4d and 5c. Values in brackets correspond to the champion cell for each condition. The anti-solvent ejection time is indicated in parentheses

HTL	V_{oc} (V)	J_{sc} (mA cm^{-2})	FF (%)	PCE (%)	R_s ($\Omega \text{ cm}^2$)	R_{oc} ($\Omega \text{ cm}^2$)	R_{sc} ($\Omega \text{ cm}^2$)
PEDOT:PSS ^a (15 s)	0.86 ± 0.02 (0.85)	18.8 ± 1.3 (20.4)	73 ± 5 (76)	11.7 ± 0.8 (13.2)	3.5 ± 0.7 (3.2)	8.8 ± 1.3 (7.4)	11583 ± 37291 (179 025)
PTAA ^a (35 s)	1.00 ± 0.03 (1.03)	19.8 ± 0.8 (19.5)	60 ± 8 (67)	12.0 ± 1.9 (13.4)	2.7 ± 0.2 (2.5)	9.1 ± 1.8 (7.4)	489 ± 245 (514)
TFB ^a (25 s)	0.91 ± 0.05 (1.00)	20.2 ± 0.6 (19.8)	56 ± 7 (61)	10.4 ± 1.6 (12.1)	3.6 ± 0.7 (3.2)	11.8 ± 3.0 (9.2)	446 ± 247 (589)
TFB ^b (15 s)	1.09 ± 0.01 (1.08)	20.5 ± 0.5 (21.2)	64 ± 3 (66)	14.3 ± 0.8 (15.3)	2.6 ± 0.2 (2.3)	9.9 ± 2.6 (6.0)	1122 ± 456 (1370)

^a Perovskite concentration = 0.87 mol L^{-1} . ^b Perovskite concentration = 1.15 mol L^{-1} .

according to the recent results obtained in the literature.^{51,61,64} It presents notably better properties compared to PAA such as its mobility (10^{-2} cm² V⁻¹ s⁻¹ vs. 4×10^{-3} cm² V⁻¹ s⁻¹, according to data from suppliers) and energy levels that more suitable for the architectures tested here.

The optimization of the anti-solvent and its ejection time could not entirely solve the identified issue of pinholes within the perovskite layer, particularly on the microscale and nanoscale (Fig. 2b). In addition, the number of pinholes is not entirely reproducible, meaning that the process could not be totally controlled. It was decided to improve the process by increasing at first the concentration of the perovskite precursor (1.15 mol L⁻¹ instead of 0.87 mol L⁻¹), leading to more reproducible and efficient devices. This also induces an increase in the thickness of the active layer from 250 nm to 400 nm. A proportional increase in absorption is observed below 750 nm and the effect on the optical properties of the devices is negligible for wavelengths targeted for tandem applications (above 800 nm), as shown in Fig. S7 (ESI†). A summary of the corresponding results is presented in Fig. 5. The amount of the crystalline FACsPbI_3 phase within the active layer increased for all the t_{ASE} tested, Fig. 5a. This proportion seems maximal between 15 s and 25 s of t_{ASE} . According to Fig. 5b, the plot of our previously defined indicator thus allows defining an optimal ejection time slightly different from the one previously determined, *i.e.* close to 15 s. Even if this indicator is slightly lower for this new concentration, the performance in solar cells is better presented in Fig. 5c, indicating a more important effect of the quality and thickness of the deposited perovskite layer than those of its microstructure. These tests in solar cells enable validation of this first optimization phase. As shown in Fig. 5e, we note a further improvement of the V_{oc} after increasing the perovskite precursor concentration and an increase in efficiency from 10% to 14%, Table 1, probably explained by an improvement of the perovskite microstructure.

In addition to the anti-solvent, an annealing procedure after the perovskite deposition permits the full conversion of the intermediate perovskite precursors into the desired perovskite and often improves the properties of the perovskite layer, and

the overall performance of the cells.^{65,66} This approach consists in annealing the perovskite layer under milder conditions to avoid a too fast evaporation of the solvents and to obtain a more homogeneous layer at the nanoscale. In the literature, this type of approach has already been encountered essentially for the perovskite of form MAPbI_3 .⁶⁷⁻⁶⁹ In some studies, this is also carried out with a solvent (or solvent mixtures) to assist the perovskite crystallization during the annealing.^{65,66,70,71} The nature of the anti-solvent is important, and ethyl acetate (EA) is well-suited for this optimization^{67,68} for mono- and mixed-cation perovskites.⁶⁹ The annealing procedure is usually performed in a single step, typically by setting the temperature at 100 °C. The conversion of mixed perovskite has also been reported to possibly occur at room-temperature due to the addition of Cs^+ in the perovskite formulation.^{69,72} However, several successive steps including lower intermediate temperature seem to be helpful to obtain higher quality of the perovskite and thus higher device efficiency.^{73,74} This approach can therefore be particularly applicable to our case study. Here, in the search for the optimisation of the perovskite deposition procedure, we have therefore introduced a drying time before annealing (t_{DBA} , Fig. 3) at room temperature without solvent in order to limit the number of study parameters and to have a greater control of this new process. As illustrated in Fig. S8 (ESI†), we noticed that 20 min t_{DBA} may help to enhance the PCE of our devices. Between 0 and 15 min t_{DBA} , a progressive enhancement in device performances can be observed. After 15–20 min, a plateau is then reached, hence the choice of 20 min was extended in the cases of other t_{ASE} . The enhancement magnitude, however, seems dependent on t_{ASE} . As a result, the present study deals with the coupled effects of t_{ASE} and t_{DBA} upon cell efficiencies and perovskite properties. The goal is thus to optimize the PSCs, and share a way to do so.

The morphological properties of the perovskite layer were studied by scanning electron microscopy (SEM) as a function of t_{ASE} duration, as presented in Fig. 6a1, b1, c1 and d. For $t_{\text{DBA}} = 0$ min, the perovskite morphology is characterized by nanoscaled pinholes and grain size ranging between 200 and 300 nm. It appears that the grains are slightly larger after 5 s t_{ASE} . This is supported by statistical analysis (Fig. 6d), where the

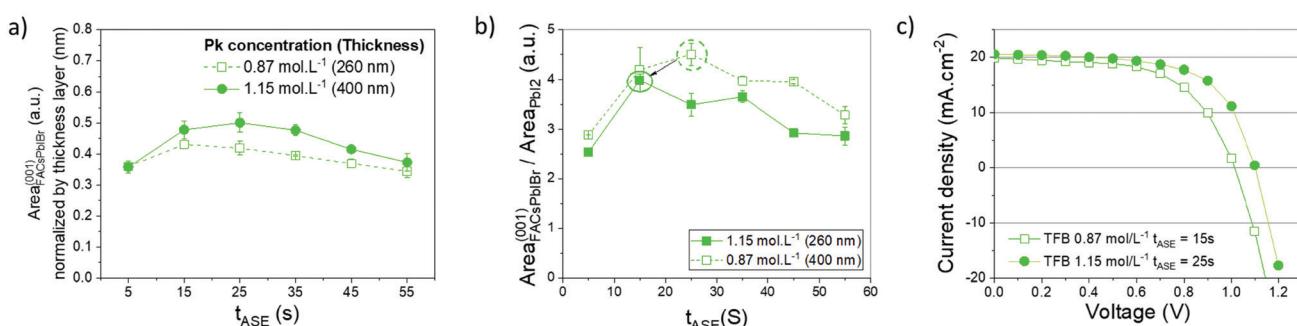


Fig. 5 Effect of perovskite concentration, 0.87 mol L⁻¹ (open symbols dashed lines) vs. 1.15 mol L⁻¹ (closed symbols continuous lines), on: (a) extracted areas of the diffraction peaks related to the FACsPbI_3 phase according to orientation (001) normalized with the thickness of the perovskite layer as a function of t_{ASE} ; (b) ratio of FACsPbI_3 and PbI_2 phases as a function of t_{ASE} ; and (c) J – V measurements in reverse scan obtained for the 2 concentrations with their respective optimal t_{ASE} .



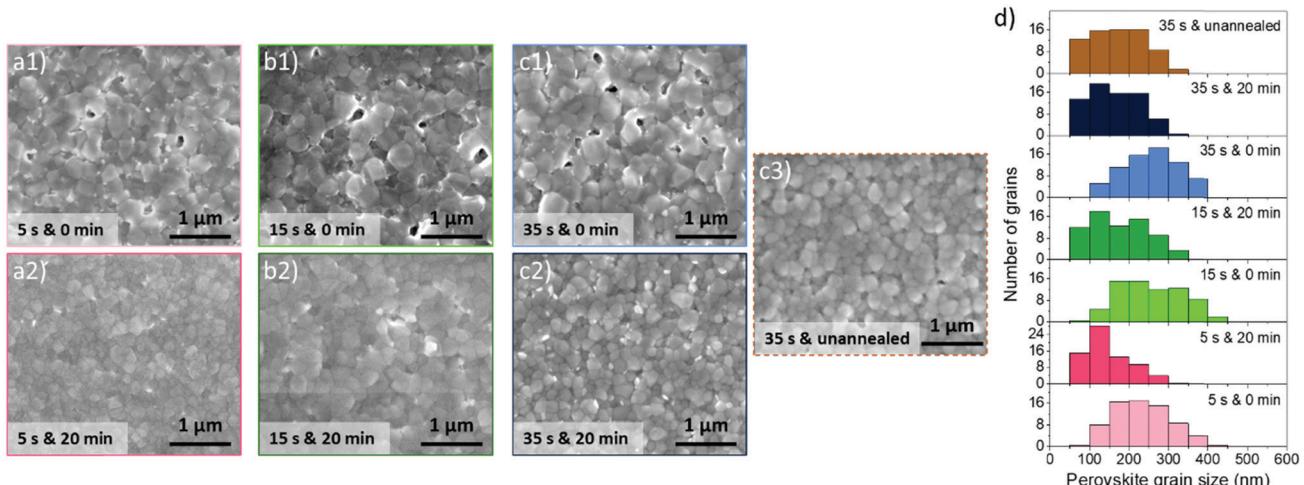


Fig. 6 (a)–(c) SEM observations of perovskite layers (performed with a perovskite concentration of 1.15 mol L^{-1}) for different t_{ASE} & t_{DBA} ; and (d) corresponding grain size distributions.

averages of grain size distributions are respectively 250–300 nm (15 s and 35 s) and 200–250 nm (5 s).

The coupled influence of t_{ASE} and t_{DBA} has also been studied, and the corresponding SEM images of the resulting perovskite layer morphology are depicted in Fig. 6a2, b2 and c2. As soon as 20 min t_{DBA} is applied to develop the perovskite layer, the morphology changes significantly whatever t_{ASE} is used. We observe that the additional drying step leads to more homogeneous surfaces, without nano-scaled pinholes, also noticeable when no annealing was performed after the drying step, Fig. 6c. This seems to be confirmed by the grain size distributions, which appear to be somewhat wider and show a decrease in the average grain size of around 150 to 250 nm. The introduction of this step leads therefore to perovskite layers with a much more homogeneous and smoother quality from a morphological point of view, which can undoubtedly have positive repercussions on the PV performances of the manufactured cells. Thus the manner in which the perovskite layer crystallizes during the annealing step determines strongly its nanoscale morphology. Therefore, delaying the annealing slows down perovskite layer crystallization, which then allows a better nanoscale morphology. It is believed that crystallization at room temperature affects the diffusion magnitude of perovskite precursors within the “fresh” undried thin perovskite layer (right after the spin-coating step) by slowing it down as well. Perhaps, the mechanism would be similar to the one that occurs when a surfactant is added in the perovskite solution.⁷⁵

As mentioned in the introduction, the main novelty of this work is the combined study of the effect of the t_{ASE} and t_{DBA} parameters on the PV performance of cells. The results relating to PV performances are presented in Fig. 7. Without t_{DBA} , the best performances were reached with the lowest t_{ASE} : *i.e.*, 5 s and 15 s (dotted lines in Fig. 7). Average PCEs around 14–15% are obtained for these two conditions. Increasing t_{ASE} reduces the performances to about 12%. These variations essentially result from the FF and V_{oc} parameters, Fig. 7 and Table 2. A FF above 60% is measured for t_{ASE} of 5 s and 15 s, whereas it is

reduced for longer t_{ASE} . These observations are compatible with an increase in the series resistance R_s as a function of the t_{ASE} , indicating lower effective charge transport through the whole device and likely more specifically between the active layer and the interfacial layers, Table 2. V_{oc} is also larger for shorter t_{ASE} and reaches up to 1.09 V, while delaying the ejection of the anti-solvent leads to a V_{oc} reduction of about 0.10 V, Table 2. This may thus also result from an increase in non-radiative recombinations,^{76,77} as shown in Fig. S9 (ESI†).

The addition of 20 min t_{DBA} leads to a clear improvement of the PCE, except for the t_{ASE} of 5 s. For a t_{ASE} of 15 s, the average PCE is indeed shifted from $14.3 \pm 0.8\%$ to $15.9 \pm 0.9\%$ and for a t_{ASE} of 35 s, it is shifted from $11.9 \pm 1.1\%$ to $15.7 \pm 0.6\%$. The gain obtained by the variation of this processing parameter seems to be more significant than that obtained by the variation of the t_{ASE} . These variations in efficiency are explained by variations in FF and V_{oc} , as shown in Fig. 7, where the FF reaches up to 72% for t_{ASE} of 15 s and 35 s and the V_{oc} tends towards 1.1 V. This is a small change for 5s and 15s t_{ASE} , but the increase is larger for 35s t_{ASE} . The J_{sc} seems little impacted. We also note an improvement in the R_s at 15 s/20 min. Thus, the two conditions 15 s/20 min and 35 s/20 min appear very relevant and equivalent in average PCE, between 15.5 and 16%. The champion cell achieved 17.1% for the 15 s/20 min condition (out of 6 cells in total), due to a high FF (77.1%). The best cell at 35 s/20 min is 16.6%, due to a slightly lower FF (71.8%). However, the latter has the highest V_{oc} : 1.13 V against 1.10 mV at best for 15 s/20 min. Consequently, it seems that an improvement in perovskite morphology by an optimization of processing conditions (ejection time and dripping time) induces an improvement in solar cell performance in particular the FF.

The last tested condition, namely 35 s/unannealed, highlights that the V_{oc} improvement is clearly brought by the drying step at room-temperature, Fig. 7c. However, very poor FF and J_{sc} were obtained under this condition. This indicates that standard thermal annealing is still required to achieve reasonable



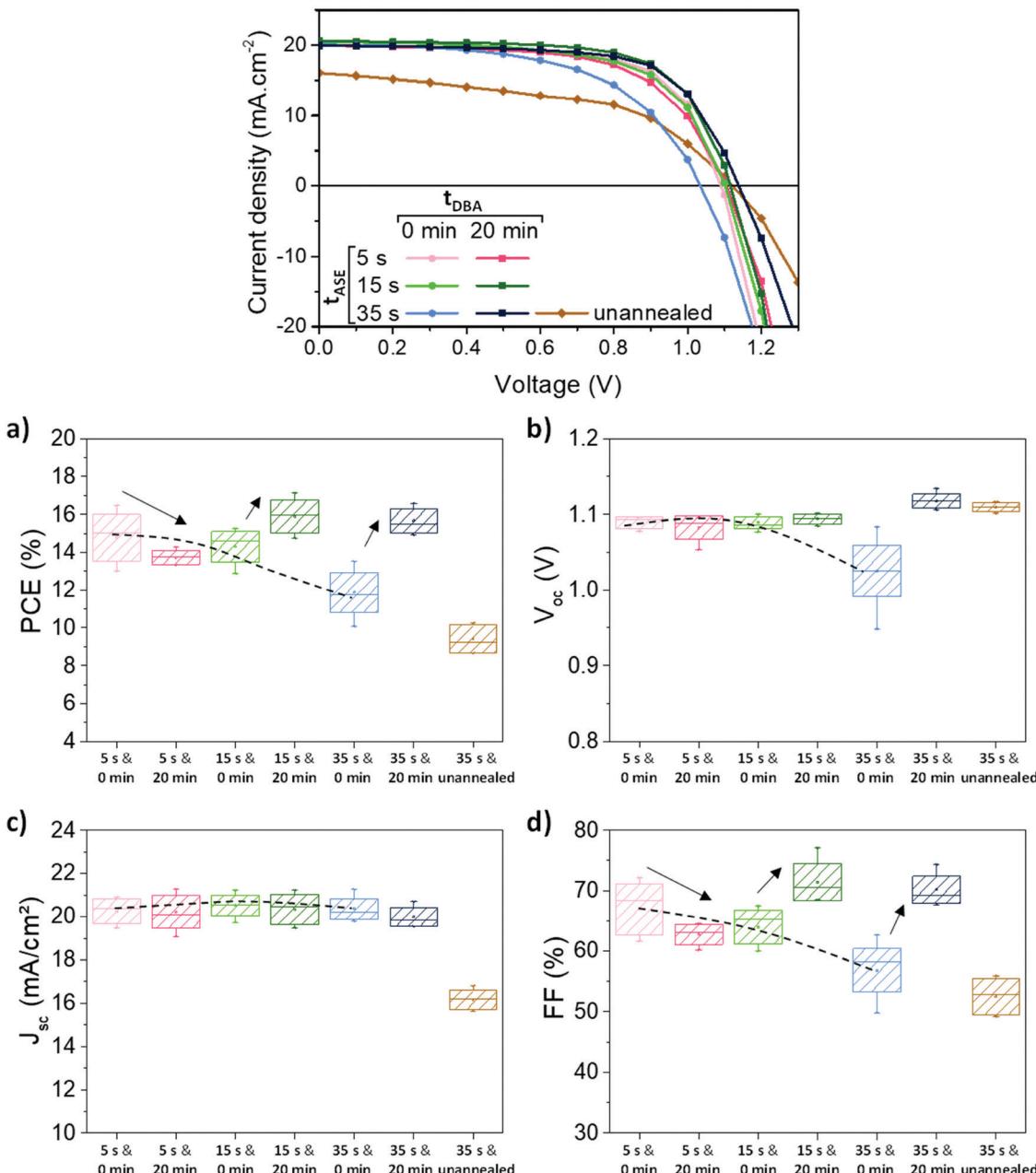


Fig. 7 Influence of the processing parameters of the perovskite layer on the device performances. The two durations of t_{ASE} and t_{DBA} are varied. (a) Representative J - V measurements in reverse scan, with an active area of 0.34 cm^2 ; means (points), medians (lines), standard deviations (boxes) and minimum–maximum (extrema) of the PV parameters (statistics on 6 to 15 cells); (b) PCE, (c) V_{oc} , (d) J_{sc} and (e) FF. The dotted lines indicate the trend as a function of t_{ASE} , while the effect of t_{DBA} is represented by arrows.

perovskite conversion and extraction/transport of the electronic charges within the device.

To better understand the effect of the processing conditions on cell performance, the perovskite layer has been further characterized. As mentioned previously, all the tested t_{ASE} promote the nucleation and growth of the same few crystal phases (FACsPbIBr, PbI₂ and δ -FAPbI₃), Fig. 8a (top graph) and Fig. S9 (ESI[†]). From a quantitative standpoint, the ratio of perovskite phases relative to PbI₂ was evaluated with the surface of the corresponding diffraction peaks, Fig. 8b. The maximum is

reached for a t_{ASE} close to 15 s. At this value, the width at the mid-height of the diffraction peak at 14.1° according to orientation (001) is also minimal. For the other orientations, the largest variations are observed for the (111) orientation, Fig. S10 (ESI[†]). The ratio of orientations (111) and (001) is minimal for a t_{ASE} of 15 s, confirming the strong (001) orientation at this stage of t_{ASE} . This could partially explain the evolution of the V_{oc} with t_{ASE} .³⁶

Concerning the combined influence of t_{ASE} and t_{DBA} , in addition to the same phases already observed, two new crystal phases φ_1 and φ_2 are detected when t_{DBA} is applied, Fig. 8a

Table 2 Averages and standard deviations of PV parameters corresponding to J – V measurements presented in Fig. 7. Values in brackets correspond to the champion cell for each condition

Conditions	V_{oc} (V)	J_{sc} (mA cm^{-2})	FF (%)	PCE (%)	R_s ($\Omega \text{ cm}^2$)	R_{oc} ($\Omega \text{ cm}^2$)	R_{sc} ($\Omega \text{ cm}^2$)
5 s & 0 min	1.089 ± 0.008 (1.093)	20.2 ± 0.6 (20.9)	67.0 ± 4.2 (72.1)	14.8 ± 1.2 (16.5)	2.47 ± 0.24 (2.24)	8.6 ± 2.7 (5.6)	934 ± 404 (1644)
5 s & 20 min	1.083 ± 0.015 (1.095)	20.2 ± 0.8 (20.8)	62.8 ± 1.7 (62.8)	13.7 ± 0.4 (14.3)	2.39 ± 0.55 (1.98)	9.7 ± 2.5 (8.3)	841 ± 436 (1062)
15 s & 0 min	1.090 ± 0.008 (1.084)	20.5 ± 0.5 (21.2)	64.0 ± 2.8 (66.4)	14.3 ± 0.8 (15.3)	2.56 ± 0.16 (2.32)	9.9 ± 2.6 (6.0)	1122 ± 456 (1370)
15 s & 20 min	1.094 ± 0.006 (1.097)	20.3 ± 0.7 (20.2)	71.4 ± 3.0 (77.1)	15.9 ± 0.9 (17.1)	2.32 ± 0.37 (2.30)	6.9 ± 1.5 (5.1)	1609 ± 183 (1788)
35 s & 0 min	1.025 ± 0.034 (1.084)	20.4 ± 0.5 (19.9)	56.8 ± 3.6 (62.7)	11.9 ± 1.1 (13.5)	3.65 ± 1.06 (2.57)	11.2 ± 1.6 (7.1)	741 ± 381 (1097)
35 s & 20 min	1.117 ± 0.009 (1.117)	20.0 ± 0.4 (20.7)	70.2 ± 2.2 (71.8)	15.7 ± 0.6 (16.6)	3.17 ± 0.60 (2.74)	7.7 ± 1.2 (7.4)	1480 ± 466 (1720)
35 s & unannealed	1.109 ± 0.006 (1.110)	16.1 ± 0.4 (16.8)	52.5 ± 3.1 (54.9)	9.4 ± 0.7 (10.2)	4.19 ± 0.93 (3.46)	15.0 ± 2.2 (13.4)	326 ± 29 (340)

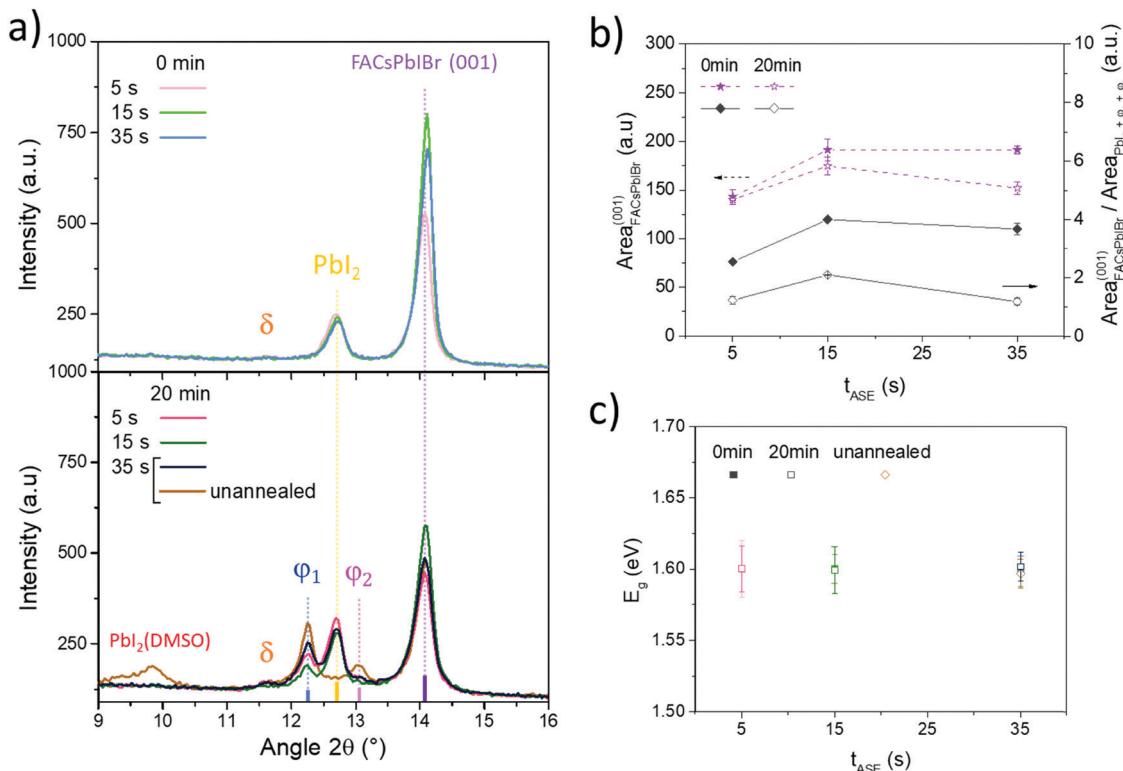


Fig. 8 (a) Zoom of X-ray diffraction patterns of perovskite layers for 2θ between 10° and 16° for different conditions of t_{ASE} (for $t_{DBA} = 0$ min on top and $t_{DBA} = 20$ min at the bottom); (b) areas of the main diffraction peaks relating to the FACsPbIBr phase and of the ratio of the FACsPbIBr phases compared to PbI_2 , φ_1 and φ_2 phases as a function of the t_{ASE} for t_{DBA} of 0 and 20 min; and (c) optical bandgap of the perovskite layers for the different tested conditions (calculated using the Tauc plot method).

(bottom graph). Their diffraction peaks appear respectively at 12.2 – 12.3° and 13.1° . According to the literature, these phases seem to correspond to the phases denoted 6H and 4H of the study by Gratia *et al.*,⁷⁸ which concerns the intermediate phases possibly observed in mixed perovskites during its conversion process.⁷⁸ The latter is crystalline intermediates between the δ - FAPbI_3 and perovskite phases. The peak at 13.1° , *i.e.* the φ_2 phase, can also correspond to the PbX_2 phase, located between 13.1° and 13.2° .²² However, the absorbance spectra obtained on the same samples show optical transitions similar to the intermediate phases reported by Gratia *et al.*⁷⁸ This therefore seems to confirm the hypothesis of the formation of crystalline intermediates induced by the application of t_{DBA} . However, very little work concerns the study of these two phases, in particular

their optoelectronic properties and therefore their impact on the PV performance of cells produced from these perovskites.

Here, when t_{DBA} is applied, φ_1 and φ_2 phases are formed during the drying step, together with the improvement in the perovskite layer's morphology. Both may thus have a role to play in the assessed enhancements in PV performances and/or be interconnected. Regarding the final thermal annealing step, it allows the elimination of the remaining $\text{PbI}_2(\text{DMSO})$ complexes undoubtedly⁷⁹ responsible for the limited FF and J_{sc} values previously described for devices elaborated with the condition of 35 s/unannealed.

Furthermore, it is important to note that the angular positions of all diffraction peaks relative to the perovskite remain unchanged after application of t_{DBA} , indicating conservation of



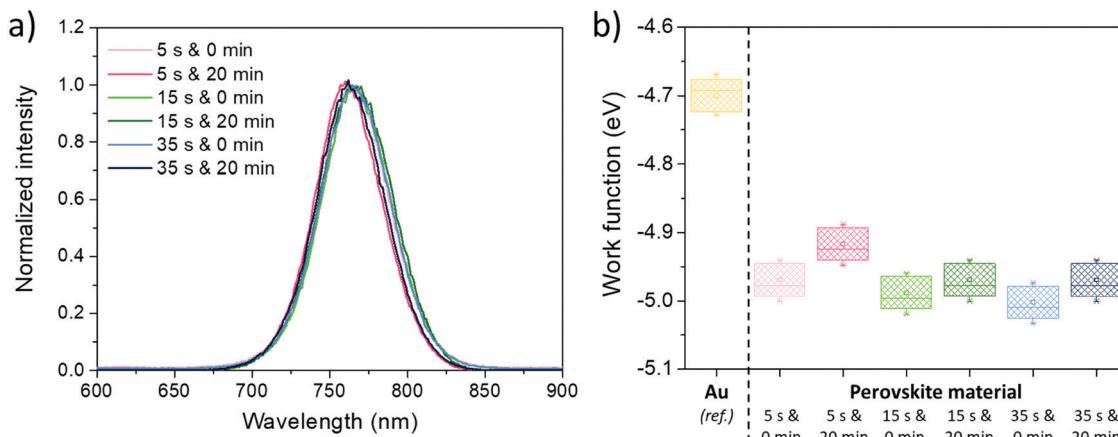


Fig. 9 (a) PL spectra for the different tested t_{ASE} and t_{DBA} , and (b) work functions of perovskite layers developed according to different conditions of t_{ASE} and t_{DBA} .

crystal lattice parameters. This is supported by UV-Visible spectroscopy analyses, which after exploitation of the Tauc plots show a constancy of the bandgap whatever the t_{ASE} and t_{DBA} conditions, Fig. 8c. In parallel, as shown in Fig. 8b (dotted lines), there is a loss of the FACsPbIBr phase in favor of the formation of other crystalline phases (PbI_2 , φ_1 and φ_2). This trend seems to be confirmed for each t_{ASE} , Fig. 8b. This also seems to correspond to a loss of absorption as shown in Fig. S11 (ESI[†]). As shown in Fig. S10 (ESI[†]), the FACsPbIBr phase is further disoriented as t_{ASE} increases for a t_{DBA} of 20 min. At the same time, widening of the width at half the height of the main diffraction peak of this phase is also noted, indicating a reduction of the crystal quality. Given the changes in efficiencies granted by the FF and V_{oc} improvements with t_{DBA} , these changes appear to be of lesser importance or at least the formation of these new phases (φ_1 and φ_2) seems to compensate for the decreased proportion of the perovskite phase and its lower quality.

Regarding the photoluminescence spectroscopy results, shown in Fig. 9a, if t_{ASE} seems to have no effect on the peak position, a blue-shift can be typically observed when t_{DBA} is applied. Due to the constancy of the optical bandgap observed previously, it can therefore be said that this cannot be explained by a chemical modification of the perovskite but by a probable improvement in the properties of the electronic defects within the perovskite layer. Indeed, the energy states of the electronic traps can be located in shallower states within the bandgap and this, coupled with a lower defect density, can lead to a spectral blue-shift in the PL position.⁸⁰ This could explain the performance gains observed only for 15 s and 35 s t_{ASE} . In addition, the results of the measurements carried out using a Kelvin probe are presented in Fig. 9b. In fact, the addition of t_{DBA} seems to systematically decrease the work function (Fermi level) of the perovskite layer by about 0.1 eV for the three t_{ASE} tested. It is therefore conceivable that the energy positions of valence and conduction bands are also modified. These changes in energetic properties are likely the overall result of all the previously described modifications brought by the room temperature additional drying step.

To check the stability of these phases, an ageing study has been performed (in a glovebox, and under dark and room temperature conditions), Fig. S12 (ESI[†]). Overall, the PV parameters are stable for at least 450 h. We however note a slight decrease of these parameters, probably because of the insufficient barrier level of the N-type layer regarding the diffusion of the silver electrode and perovskite compounds.⁸¹ This can be overcome by adding an ITO layer between SnO_2 and Ag layers, which can act as a barrier against diffusion, Fig. S12 (ESI[†]). Therefore, the perovskite layer and its crystalline phases are intrinsically stable over time. Finally, the hysteresis phenomenon of these solar cells has been evaluated before and after aging, Fig. S13 (ESI[†]). It is relatively stable over time and the hysteresis index⁸² was found to be close to 10%. Thus, the optimization performed seems to be sustainable over time and generates a low hysteresis phenomenon.

Conclusions

In this paper, the development of innovative perovskite cells with PIN architecture has been achieved through a material/process optimization approach. Thanks to a previous study carried out in the NIP architecture, the formulation of the perovskite used as an active layer was optimized. Then, the first optimization phase developed in this article concerned the perovskite deposition process according to the HTL beneath, thanks to the change of anti-solvent and the optimization of the anti-solvent ejection time. The interest in the PTAA and TFB layers was immediately highlighted, characterized by their potential for higher V_{oc} . The study was then refocused on TFB as the HTL due to its more suitable intrinsic properties and lower explored investigations in PV literature. After an optimization of the concentration of the perovskite solution, the double optimization of the anti-solvent ejection time (t_{ASE}) and the drying time before annealing (t_{DBA}) was performed, allowing improvement in the PCE of our devices from 10% to 17% with an optimized V_{oc} of about 1.1V. One of the important results of this study is the methodology



used for this optimization. Indeed, the main marker used is the observation of defects in the perovskite layer at the macroscopic and microscopic scales. This optimization of the perovskite layer morphology leads to a significant increase in performance. The additional gain in performance is then obtained by optimizing the nanostructure and in particular by defining markers resulting mainly from XRD analyses such as perovskite/ PbI_2 phase ratios. The formation of the φ_1 and φ_2 phases seems to have a very positive effect on the nanostructure of the perovskite and thus on its optoelectronic properties.

Abbreviations

t_{DBA}	Drying time before annealing
t_{ASE}	Anti-solvent ejection time
ETL	Electron transport layer (n)
FA	FormAmidinium
FF	Fill factor
GMF	Glass microFiber
HTL	Hole transport layer (p)
IPA	IsoPropyl alcohol
ITO	Indium tin oxide
$J-V$	Current density-voltage
J_{sc}	Short-circuit current density
MA	MethylAmmonium
MAPI	MAPbI_3
PCE	Power conversion efficiency
PEDOT:PSS	Poly(3,4-ethyleneDiOxyThiophene): poly(styrenesulfonate)
PL	PhotoLuminescence
PSC	Perovskite solar cell
PTAA	Poly(triaryl amine)
PTFE	Polytetrafluoroethylene
PV	Photovoltaic
R_{oc}	Resistance at open circuit condition
R_s	Series resistance
R_{sc}	Resistance at short circuit condition
SEM	Scanning electron microscope
TFB	Poly[(9,9-diOctylfluorenyl-2,7-diyl)-co-(4,4'-(<i>N</i> - <i>p</i> -butylphenyl))diphenylamine)]
V_{oc}	Open-circuit voltage
XRD	X-Ray diffraction
NIP	Negative-intrinsic-positive solar cell architecture
PIN	Positive-intrinsic-negative solar cell architecture
spiro-OMeTAD	2,2',7,7'-Tetrakis[<i>N,N</i> -di(4-methoxyphenyl)amino]- 9,9'-spirobifluorene

Conflicts of interest

There are no conflicts to declare.

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