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High performance carbon-based printed perovskite solar cells with humidity assisted thermal treatment†

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We report humidity assisted thermal exposure (HTE) as a post-treatment method for carbon based printed perovskite solar cells (CPSCs). The method does not only improve the interfaces of different layers of the printed stack, but also provides a pathway to fabricate high performance CPSCs with low hysteresis along with high stability. The HTE treatment directly influences over the associated components in the stack and remarkably improves each photovoltaic parameter of the CPSCs as seen by several characterization schemes presented in this study. The average initial efficiency ($9.0\% \pm 0.2\%$) of the CPSCs of a batch was significantly improved to $13.1\% \pm 0.2\%$ i.e. as high as 45% when subjected to HTE treatment for a period of 200 hours. Furthermore, the highest average efficiency obtained from the same batch from reverse scanning was $13.8\% \pm 0.4\%$ with a CPSC attaining as high as 14.3% when exposed to the same thermo-humid environment for a period of 115 hours. Above all, the stability of the HTE treated CPSCs was also not compromised for over 350 hours under full-sun illumination stress testing at 40 °C. The results presented in this work provide an opportunity to adopt HTE treatment as a complementary step for the fabrication of high-performance carbon-based perovskite solar cells with low hysteresis accompanied by high durability and performance reproducibility.

Organic–Inorganic lead halide perovskite based solar cells (PSCs) have recently exhibited very high efficiencies (>22%) in their traditional device structures,¹ thus showing potential to compete with other commercialized photovoltaic (PV)

technologies, such as cadmium telluride (CdTe, 22.1%), multi-crystalline Si (c-Si, 21.3%) or copper indium gallium selenide (CIGS, 22.3%) solar cells, along with their rapid energy payback time.^{2,3} Nevertheless, poor long-term stability and costly materials (i.e. high purity Spiro-OMeTAD approximately costs 200–400 \$ per g) along with expensive manufacturing methods and high lead (Pb) wastage are the biggest challenges in the commercialization of these traditional PSCs technologies.^{4–7} Therefore, several alternative device structures have been explored by integrating these solution processed perovskite compounds to address the aforementioned key issues.^{8–12}

Amongst alternative device structures,^{11–13} carbon-based perovskite solar cells (CPSCs)^{12,14–20} look highly promising due to their low cost and abundantly available materials (TiO₂, ZrO₂, carbon black and graphite powders),^{14,20} cost-efficient scalable fabrication methods¹⁴ and the inherent high stability reported by several research groups.^{12,15,16,21} However, the high cell resistance of a thicker stack of their active materials (~12–14 μm)¹² along with uncontrolled and limited crystallization of the perovskite precursor solution in this thick porous stack are the limiting factors for significant enhancements in their solar-to-electrical conversion efficiencies.^{12,15} Therefore improvements in the photovoltaic performance of CPSCs may be realized if the aforementioned challenges could be addressed prudently.²²

Keeping this motive in mind, we report here humidity assisted thermal exposure (HTE) as a post-treatment method for these CPSCs, which does not only improve the interfaces of different layers of the printed stack, but also provides a pathway to fabricate high performance and highly stable CPSCs with low hysteresis.

In this work, we observed significant improvements in the photovoltaic parameters when a batch of CPSCs (7 devices) was first infiltrated with the traditional precursor solution (as reported before^{12,15,16}) containing lead iodide (PbI₂), methyl ammonium iodide (MAI) and 5-ammoniumvaleric acid (5-AVAI) in gamma butyrolactone (GBL), and then exposed to high humidity (70% ± 5%) at a constant temperature (40 °C) for up to 200 hours in an environmental chamber (Arctest 150 – Finland,²³ see also the ESI†).

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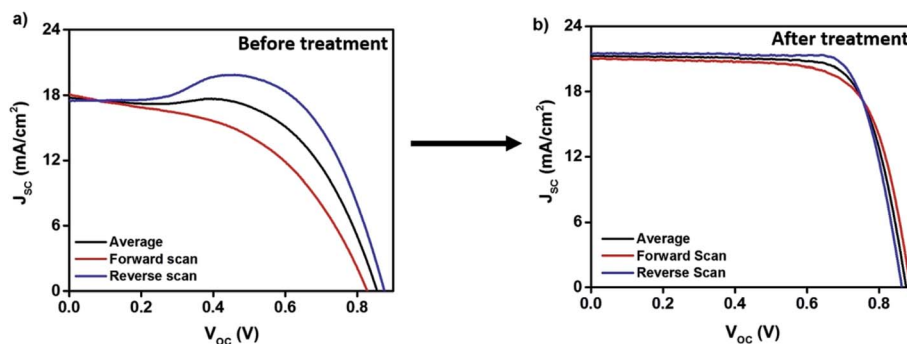


Fig. 1 (a) J - V curves of the champion CPSC (b) J - V curves of the same CPSC after HTE treatment for a period of 115 hours.

Table 1 Photovoltaic parameters of the champion CPSC as they appeared in Fig. 1a and b

Name	Scan	J_{SC} (mA cm^{-2})	V_{OC} (mV)	FF	η (%)	R_{CELL} (Ω)
Fresh	Forward	18.1	0.828	0.49	7.3	75.5
	Reverse	17.5	0.875	0.72	11.1	50.4
	Average	17.8	0.851	0.60	9.2	62.9
HTE treated (115 h)	Forward	21.0	0.887	0.71	13.3	28.2
	Reverse	21.5	0.863	0.77	14.3	29.5
	Average	21.2	0.875	0.74	13.8	28.8

Fig. 1a and b and Table 1 present the J - V curves and the photovoltaic parameters respectively of the initial and HTE treated CPSC (from a batch of 7 devices) that was exposed for a period of 115 hours to the aforementioned environmental conditions which resulted in drastic enhancements in the preliminary measured values of short circuit current density ($J_{SC} = 19\%$), fill factor (FF = 23%), and efficiency ($\eta = 50\%$) as well as a severe decrease in the overall cell resistance ($R_{CELL} = 54\%$) which is determined from the slope of the J - V curves. Above all, the hysteresis in the fabricated devices (which has been a critical issue in determining the reliable efficiencies of CPSCs) was also remarkably improved explaining the post-curing effects, which have not been observed in earlier reports of these CPSCs.^{19,22}

The detailed trends of these impressive changes in the photovoltaic parameters are also summarized in Fig. 2, where the fabricated batch (7 cells) was measured periodically up to 200 hours, after which slight signs of degradation appeared at the edges of some of the devices in this batch, indicating the extreme limits of exposure (Fig. 3a and b). Hence the HTE treatment was restricted to 115–120 hours for further batches of CPSCs fabricated for this study. Nevertheless, the overall photovoltaic performance was not compromised ($\eta = 13.1\% \pm 0.2\%$) and no major physical changes were observed, which was further verified by measuring the reflectance of the fabricated CPSCs (Fig. 3a–c).

More importantly, the repetitive J - V curves of some of these HTE treated CPSCs were also obtained by scanning them under full sunlight illumination to determine if the drastic improvements in their overall photovoltaic parameters was permanent or not. Fig. 4 presents ten consecutive J - V curves of a HTE treated CPSC which showed exceptionally high stability by

exhibiting no change in the overall conversion efficiency (13%) with a minimal compensation in the J_{SC} (1.5% increased) and V_{OC} (only 0.5% decrement) values, which were stabilized after five scans (Fig. 4).

Motivated by these promising findings, we further investigated both the fresh and HTE exposed CPSC device structures *via* isolating key components, which directly influence their overall photovoltaic performance. In this regard, the carbon back contact electrode was first examined by the four-probe sheet resistance measurement method (see the ESI†) since it directly contributes to the open circuit voltage (V_{OC}) and fill factor (FF), which were significantly improved during the HTE treatment. As expected, the sheet resistance ($R_{SH} = 8.5 \pm 0.10 \Omega \text{ Sq}^{-1}$) of the HTE treated CPSCs was impressively reduced (by 13%) compared to that of the fresh CPSCs before exposing them to HTE treatment (Table 2), thus attesting to the improvements in the overall conductivity of the porous carbon electrode. We confidently state that these improvements are mainly attributed due to an increase in the interconnectivity of carbon nanoparticles through a possible crystal growth of the perovskites (after their precursor infiltration and heating at 50°C), which may be enhanced within the porous stack during HTE treatment that consequently influenced the fill factor and open circuit voltage of the fabricated CPSCs (Fig. 2).

Therefore, the structure of the perovskite crystals within the CPSC stack was also examined *via* X-ray diffraction (XRD) measurements. Fig. 5a shows the XRD patterns of a CPSC collected at the initial state and up to 115 hours of HTE treatment which did not induce any growth of PbI_2 .²⁴ During HTE treatment, the perovskite structure remained unchanged at the atomic level: the d spacing remained constant, indicating no release of stress or strain and the FWHM of the diffraction



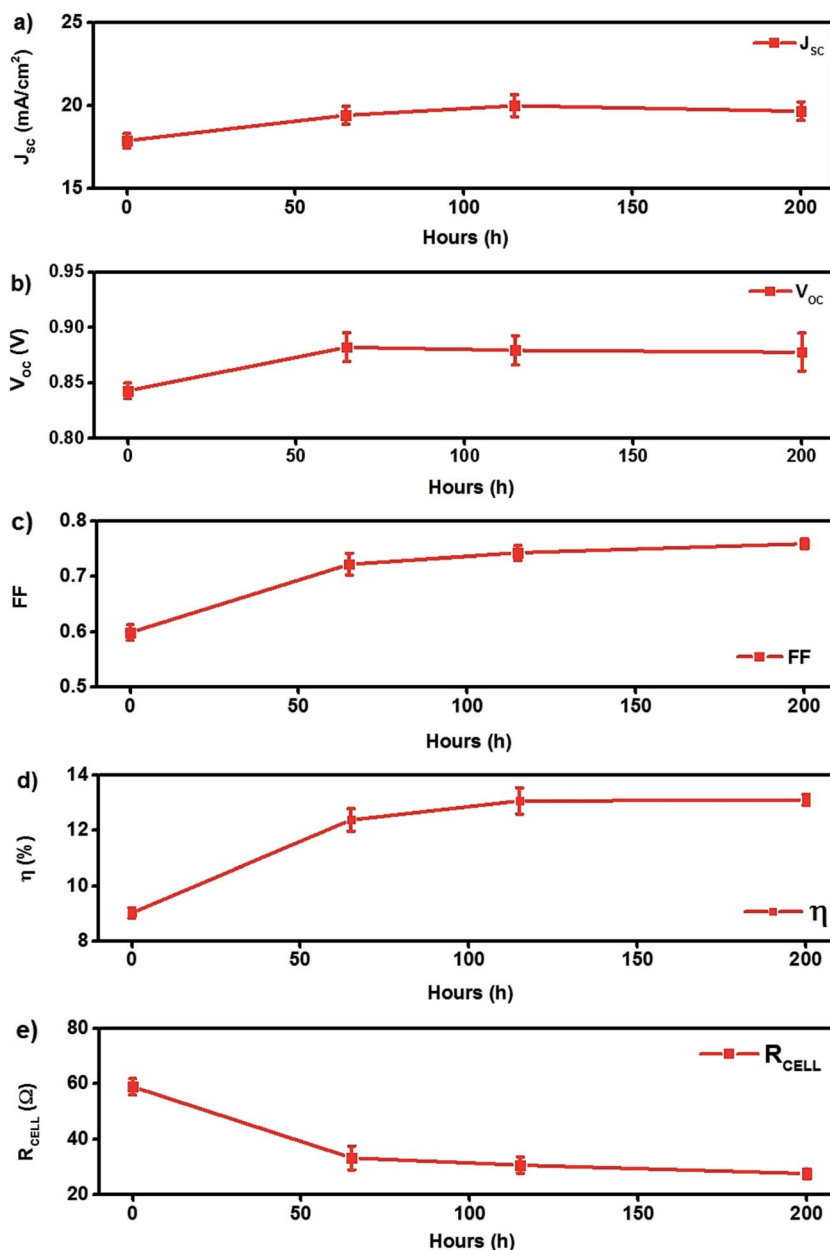


Fig. 2 Photovoltaic parameters of a batch (7 cells) of CPSCs exposed in the environmental chamber for the HTE treatment. The values represent the average of forward- and reverse-scanned J - V curves (see also the ESI†).

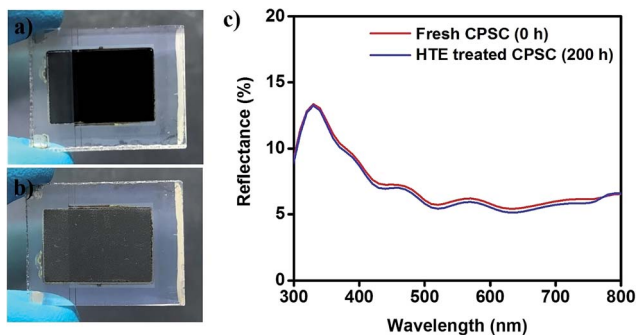


Fig. 3 (a) Front and (b) back view and (c) initial and aged reflectance spectra of a HTE treated CPSC for a period of 200 h.

peaks varied only slightly. The perovskite crystal structure was also cross-checked from FTIR spectra (Fig. 6) which also remained almost unaltered during the aging process, suggesting that the HTE treatment did not have any effect on the perovskite chemical configuration. On the other hand, an increase in the integrated intensity of the strongest perovskite reflection at 14.1° (ref. 15 and 16) was observed, most likely due to a slight increase in crystallite sizes which supports the idea of reorganization of the crystallites within the porous matrix. Hence, the effect of the HTE on the morphology and appearance of the perovskite layer was correspondingly analysed at the edge of the carbon electrode of CPSCs by scanning electron microscopy (Fig. 7 a). The micrographs revealed a steady perovskite



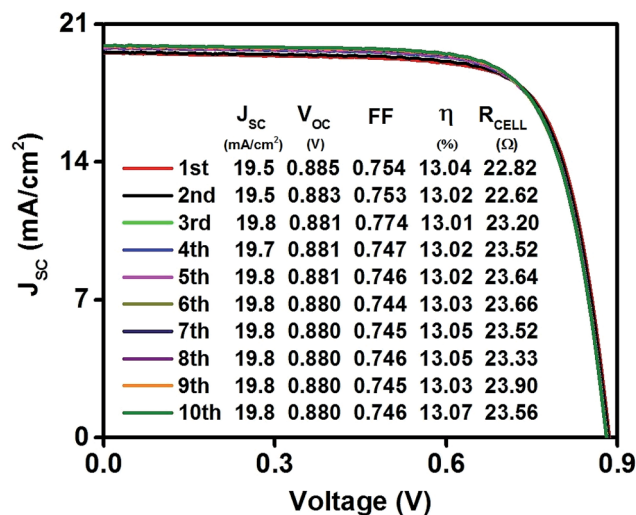


Fig. 4 J - V curves obtained from 1–10 times scanning of a high performance HTE treated CPSC for a period of 115 hours. The values represent the average of forward- and reverse-scanned J - V curves.

Table 2 Sheet resistances of the carbon electrodes

Electrode	Sheet resistance (Ω Sq ⁻¹)	No of electrodes
Fresh carbon back electrode (without perovskite precursor infiltration)	14.4 ± 0.1	3
Fresh carbon back electrode after perovskite infiltration and crystallization at 50 °C	9.8 ± 0.4	3
Carbon back electrode after HTE treatment (115 h)	8.5 ± 0.1	3

crystal growth (Fig. 7d, e and h) in the HTE-treated devices compared to the fresh ones (Fig. 7b) and consequently reduced the reduced overall porosity (Fig. 7g and h) which was well evident in the fresh devices (Fig. 7b–d) thus providing another

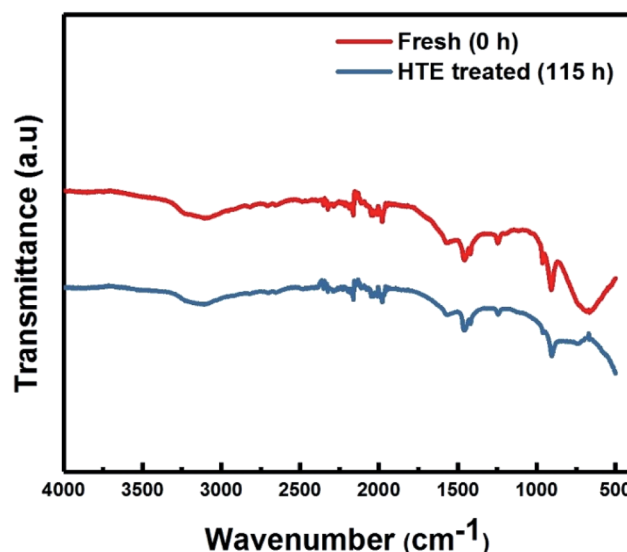


Fig. 6 FTIR spectra of fresh and HTE treated (115 h) CPSCs.

evidence for crystallite growth within the porous stack of the fabricated devices.

In addition to this, we also explored steady-state and time-resolved photoluminescence (PL, Fig. 8) to understand the influence of HTE on the emission characteristics of the perovskite layer. Upon exciting the perovskite samples at 450 nm, PL spectra corresponding to the band-to-band recombination appeared around 780 nm (inset; Fig. 8), in agreement with the intrinsic bandgap of $\text{CH}_3\text{NH}_3\text{PbI}_3$.²⁵ The injection of electrons from the conduction band (CB) of the perovskite into that of TiO_2 was further investigated by probing the decay of emission using the time-correlated single photon counting (TCSPC) technique. Fig. 6 shows the time-resolved photoluminescence (TRPL) spectra acquired from various perovskite devices measured at λ_{max} upon excitation at 408 nm from the FTO side. With increasing duration of HTE of the perovskite devices, PL quenching increased continuously, establishing that electron injection from the CB of the perovskite into that of TiO_2 becomes relatively more efficient when devices are subjected to

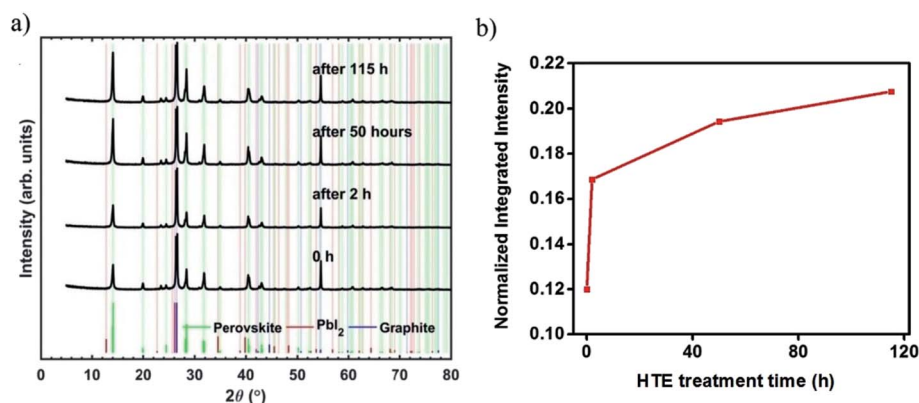


Fig. 5 (a) XRD patterns of a CPSC obtained via periodical measurements from 0–115 hours (b) normalized integral peak intensity of the same CPSC monitored from 0–115 hours.



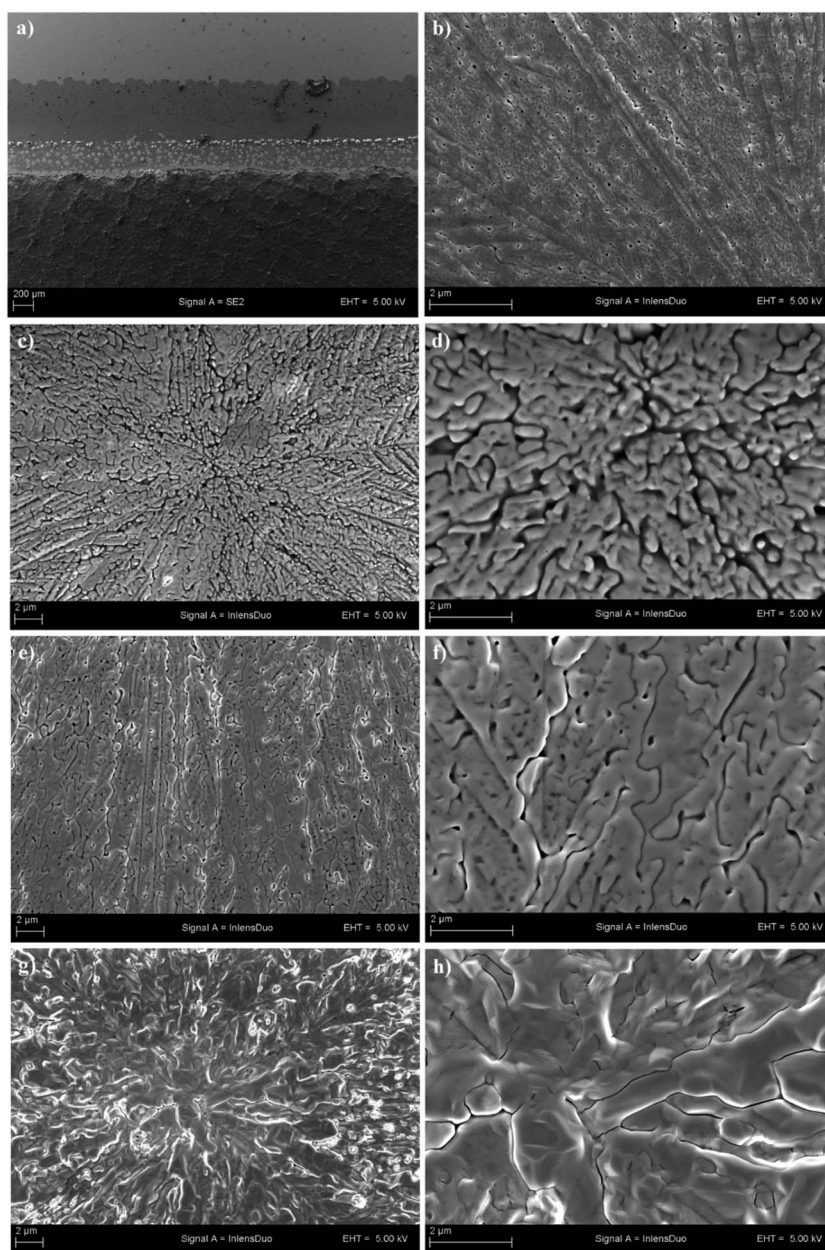


Fig. 7 SEM images: (a) top view of the carbon edge showing perovskite layer formation, (b) fresh perovskite layer at the carbon electrode edge, (c) 2 hour aged perovskite layer at the carbon electrode edge, (d) close view of the 2 hour aged perovskite layer, (e) 65 hour aged perovskite layer at the carbon electrode edge, (f) close view of the 65 hour aged perovskite layer at the carbon electrode edge, (g) 200 hour aged perovskite layer at the carbon electrode edge, and (h) close view of the 200 hour aged perovskite layer.

HTE.²⁶ Probably, in addition to improving the crystallinity, HTE makes the interface between mesoporous TiO₂ and the perovskite layer more favorable for electron injection, which eventually increases the J_{SC} .

The evidence of the perovskite crystal growth under humidity was first reported by You *et al.* in one of the traditional thin film geometry based PSCs where the thermal annealing of the perovskite precursor film was performed under ambient air (humidity = 35% ± 5%).²⁷ Similar enhancements (as presented here in this work) were also observed in their photovoltaic parameters, which include high FF (>80%) and high J_{SC} (~20

mA cm⁻²) values.²⁷ Nevertheless, due to the thin stack (~1 μm) in any type of combination for traditional PSCs (such as that presented in ref. 11), the devices do not survive in a highly humid environment over longer periods of time as due to its hygroscopic nature, the CH₃NH₃I rapidly decomposes within the perovskite crystals and affects the overall device performance.²⁷ Contrary to this, the 12–14 times thicker and robust device architecture in the case of CPSCs makes it possible to expose the devices to much higher humidity for longer periods of time. We propose that the thick (10–12 μm) hydrophobic carbon electrode of CPSCs, during the HTE treatment, works as



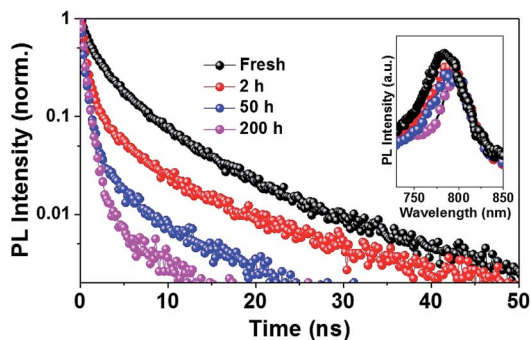


Fig. 8 Photoluminescence of the CPSC after HTE treatment from 2–200 hours. Time-resolved photoluminescence measured at λ_{max} upon excitation at 408 nm. The inset shows the steady-state photoluminescence recorded upon excitation at 450 nm.

a ‘mesh’ *i.e.* strongly blocks the rapid intrusion of large water droplets which may be produced in this ‘thermo-humid’ environment and only allows the gaseous state of the moisture to first pass through it and then proceed through more dense mesoporous ZrO_2 to assist the irreversible perovskite crystal growth both in the TiO_2 scaffold as well as in the whole porous stack without changing its chemical structure.

Finally, the remaining task in this work was to test the long-term stability of the HTE-treated CPSCs. In this regard, one more batch (6 cells) of the HTE exposed CPSCs was subjected to full sun illumination at 40 °C. Fig. 9a–e present the results of the photovoltaic parameters of the subjected CPSCs for a period of 350 hours of this ongoing stability test which retained ~92%

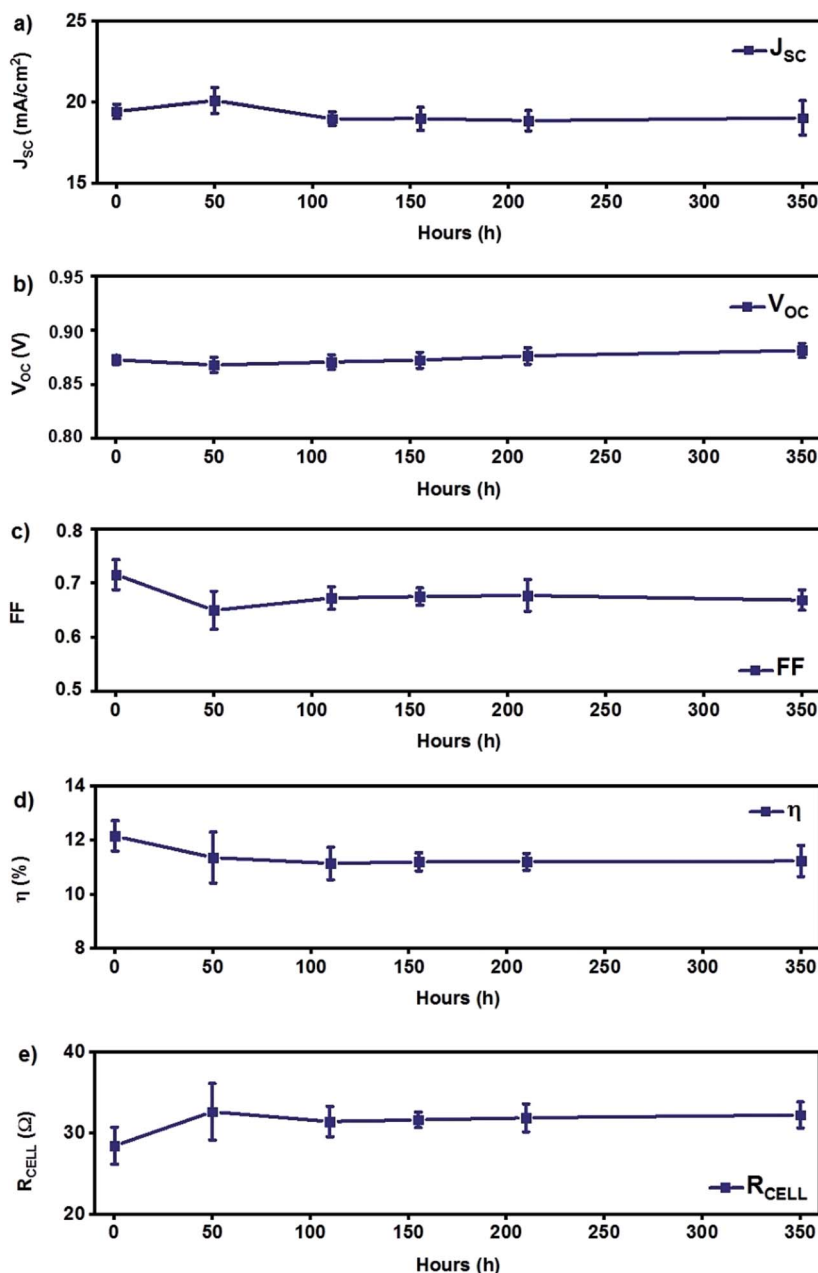


Fig. 9 Stability data of a batch of CPSCs (6 cells) along with their standard deviations for a period of 350 hours.



of the average initial efficiency thus endorsing the high performance durability and calling for an opportunity to apply this interesting method also over large area carbon-based perovskite solar modules to further enhance their performance which is now crossing 10%.^{22,28} The initial slight decrease and increase in the FF and R_{CELL} values, respectively, is an expected result due to the additional contacts (fabricated with copper tapes and silver paint and further protected with the slowly drying epoxy) that were slowly stabilized and consequently improved and retained the V_{OC} and the conversion efficiencies respectively (see also the statistical data in the ESI†).

In recent CPSC developments, efforts have mostly been made to formulate new precursor solutions of perovskites by introducing new solvents and materials which have shown good performances.^{19,20} However, the importance of reducing the overall cell resistance of CPSCs has been hardly discussed²² and has not been highlighted before this report. Further improvements in reducing the cell resistance in future work may also be realized by fabricating more conductive carbon back electrodes (*i.e.* replacing large graphite particles with low cost and conductive multi-walled carbon nanotubes (MWCNTs) or single walled carbon nanotubes (SWCNTs)). An additional HTE treatment such as the one presented in this study should potentially influence the overall photovoltaic performance by decreasing further the sheet resistance of these highly conductive carbon back electrodes within the CPSC device configuration.

To sum up, we successfully demonstrated here humidity assisted thermal exposure (HTE) as a post-treatment scheme to obtain high performance and durable carbon-based printed perovskite solar cells. It was shown in this work that traditional heating (at 50 °C) after perovskite precursor infiltration into their highly porous stack is not enough to generate sufficient perovskite crystal growth and also does not effectively improve the interfaces of the porous stack of the CPSCs. The additional HTE treatment directly influences over the associated components in the stack and remarkably improves each photovoltaic parameter of the CPSCs which were further certified by several characterization schemes presented in this study. With HTE, the average initial efficiencies ($9.0\% \pm 0.2\%$, calculated from forward and reverse scans of the J - V curves) of the fabricated devices were greatly improved to $13.1\% \pm 0.2\%$ *i.e.* as high as 45% after exposing one batch of solar cells for a period of 200 hours (Fig. 2, see also the ESI†). The highest average efficiency obtained from the same batch from the reverse scanning of the fabricated CPSCs was $13.8\% \pm 0.4\%$ with a CPSC attaining as high as 14.3% (Fig. 1 and Table 1) after 115 hours of exposure in an automated weather chamber. Moreover, the long-term photovoltaic performance of the HTE treated CPSCs has also not been compromised for greater than 350 hours under stressful conditions in the stability test. The results presented in this work are very encouraging for the adoption of HTE treatment as an additional step in the standard CPSC fabrication procedure to obtain high performance carbon-based perovskite solar cells with low hysteresis accompanied by high durability and reliable performance reproducibility.

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