Long Thermal Stability of Inverted Perovskite Photovoltaics Incorporating Fullerene-Based Diffusion Blocking Layer

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In this article, the stability of inverted (p-i-n) perovskite solar cells is studied under accelerated heat lifetime conditions (60 °C, 85 %, and N₂ atmosphere). By using a combination of buffer layer engineering, impedance spectroscopy, and other characterization techniques, the interaction of the perovskite active layer with the top Al metal electrode through diffusion mechanisms is proposed as the major thermal degradation pathway for planar inverted perovskite photovoltaics (PVs) under 85 °C heat conditions. It is shown that by using thick solution processed fullerene buffer layer the perovskite active layer can be isolated from the top metal electrode and improve the lifetime performance of the inverted perovskite photovoltaics at 85 °C. Finally, an optimized reliable solution processed inverted perovskite PV device using thick fullerene diffusion blocking layer with over 1000 h accelerated heat lifetime performance at 60 °C is presented.

1. Introduction

Perovskite-based solar cells have been among the most prominent photovoltaic (PV) technologies over the last couple of years offering a tremendous advancement in power conversion efficiency (PCE) from 3.8 to 22.7%.[1-5] Several characteristics such as high absorption coefficient,[3] tunable bandgap,[4,5] long carrier diffusion length,[6] high carrier mobility,[7] and low exciton binding energy[8,9] have allowed perovskite-based solar cells to reach such high values in PCE, even if casted from solution at low processing temperatures.[10]

Although important advancements have been made to improve PCE, the long-term stability of perovskite-based devices still remains a challenge however. Among others, important factors that affect the stability of perovskite solar cells are the moisture, heat, and light irradiation. These factors can deteriorate the performance of both the active layer and electrodes of the device. The organic–inorganic perovskite semiconductor materials are known to be intrinsically unstable to moisture[11,12] since they can form hydrated crystal phases, which mobilize the organic species, and decompose the crystal to lead iodide (PbI₂).[13] Decomposition of the perovskite active layer to PbI₂ can also happen in the presence of O₂ and light. It was suggested that superoxide ions (O₂⁻) can be formed upon irradiation of CH₃NH₃X₃ (X = Cl, Br, I) halides which, in turn, deprotonate the methylammonium cation (MA⁺) and decompose the perovskite structure.[13,14]

As a result, considerable research efforts have been dedicated toward the improvement of light and especially humidity stability of perovskite-based solar cells. It has been previously shown that under accelerated humidity conditions, there is a significant active layer degradation due to the decomposition of the perovskite crystal. Shielding the active layer from humidity, and therefore, improvement in device lifetime has been reported through the usage of hydrophobic layers such as thiols.[11] Fang et al. have also reported efficient moisture resistance by incorporating a cerium oxide (CeO₂) bilayer achieving stability of 200 h under ambient conditions.[15] In a recent work, Yang et al. have reported that the stability of perovskite solar cells under continuous illumination can also be improved by doping the [6,6]-phenyl-butyric acid methyl ester (PCBM) with graphene quantum dots (GQDs).[16] More recently, it has been reported that photooxidation of the perovskite active layer leads to the formation of charge trap centers which, in turn, may cause significant changes in charge density and charge extraction efficiency of the device.[17] On the other hand, by incorporating mixed-cation-based devices, such as formamidinium-cesium (FA₀.₈₃Cs₀.₁₇Pb₁₋ₓIₓBrₓ), the oxygen-induced degradation can be markedly minimized.[18] Research regarding the heat stability of such devices has been scarce however, especially at elevated temperatures such as 85 °C. The 85 °C is a particularly interesting temperature since, according to International Standards (IEC 61646 climatic chamber tests), it represents the elevated temperature of a roof during a hot summer day.[19] Although CH₃NH₃PbI₃ has been previously reported to be mostly intrinsically stable at elevated temperatures as high as 85 °C in the absence of O₂ and ambient conditions,[19,20]
the devices themselves are largely unstable. Zhao et al. have reported that 2,2',7,7'-tetakis[N,N-di(4-methoxyphenyl)amino]-9,9'-spirobifluorene (spiro-MeOTAD) exhibits a state change at elevated temperatures, which reduces the hole mobility. These issues can be tackled by using the thermally stable NiO and PCBM materials as the hole-transporting layer (HTL) and electron-transporting layer (ETL), respectively.[20] In another work where the active layer was damaged by subsequent electrode interaction was reported by Domanski et al. In their work, they suggested that temperatures as low as 70 °C can cause Au metal migration toward the active layer. The migration of Au can be blocked by incorporating a Cr interlayer between the Au metal electrode and HTL layer.[21] Using recent modeling tools it has also been proposed that blocking layers can reduce the voltage drop on perovskite active layer, providing reduced ionic motion and predicted improved stability.[22] In a more recent work, Bi et al. have reported improved heat stability by using an ETL consisting of N-doped graphene, PCBM, and carbon quantum dots (CQDs).[23]

In this work, we have performed accelerated heat lifetime tests at 85 °C under inert (N2) atmosphere in the dark, using p-i-n perovskite-based solar cells with the following structure: indium tin oxide (ITO)/poly(3,4-ethylenedioxythiophene) polystyrene sulfonate(PEDOT:PSS)/CH$_3$NH$_3$PbI$_3$/PC[70]BM/AZO/Al, following the perovskite formulation reported by Liang et al.[24] [6,6]-Phenyl-C$_70$-butyric acid methyl ester (PC[70]BM) is a widely used material in p-i-n perovskite solar cells that not only functions as an ETL but also passivates charge trap sites present on the surface and grain boundaries of the perovskite films.[25] Aluminum-doped zinc oxide (AZO) was also incorporated since it has been previously reported to increase the electron selectivity of such device structures.[16] Thus, both PC[70]BM and AZO ETLS play an essential role on the device operation of p-i-n perovskite PVs. In this report, two different thicknesses of PC[70]BM were used for heat lifetime performance tests. These include CH$_3$NH$_3$PbI$_3$-based devices with PC[70]BM layers of ≈70 nm (thin fullerene-based devices) and ≈200 nm (thick fullerene-based devices) thickness. We show that thick PC[70]BM yields improved heat stability over the thin PC[70]BM counterparts at the expense of reduced PCE. We have shown that PC[70]BM layer isolation is the key for achieving improved stability. Finally, we also show a perovskite formulation based on CH$_3$NH$_3$PbI$_{3-x}$Cl$_x$ that is optimized to function efficiently using a thick PC[70]BM ETL while still maintaining good PCE. This developed device retained very good thermal stability at 60 °C for 1000 h.

2. Results and Discussion

2.1. CH$_3$NH$_3$PbI$_3$ Film Characterization

CH$_3$NH$_3$PbI$_3$ (MAPbI$_3$) has been previously reported to be mostly intrinsically stable at 85 °C under N$_2$ atmosphere by Conings et al.[18] In order to confirm previous observations, ITO/PEDOT:PSS/CH$_3$NH$_3$PbI$_3$ films were annealed at 85 °C on a hotplate inside a N$_2$-filled glove box in the dark for 96 h. The X-ray diffraction (XRD) patterns of fresh and thermally annealed films are shown in Figure 1. Similar to what has been previously reported in the literature,[17] the XRD patterns show no traces of PbI$_2$, judging by the absence of diffraction peaks at 12.7° and 38.7°. Although there is a peak at 52.4° which has been previously attributed to PbI$_2$ phase, the size of this peak does not change upon thermal treatment. This suggest that CH$_3$NH$_3$PbI$_3$ is, for the most part, intrinsically stable at temperatures to 85 °C under N$_2$, which is in agreement with previous studies.[19] The UV–vis absorption spectra of fresh and aged perovskite films also remain quite unchanged and are similar to the ones reported by Conings et al. which further points toward the intrinsic stability of the perovskite film (Figure S1, Supporting Information).

2.2. Heat Stability and Characterization of CH$_3$NH$_3$PbI$_3$-Based Devices

As it was previously mentioned, we tested two different sets of p-i-n solar cells incorporating different fullerene thicknesses, that is, PC[70]BM with ≈70 nm (thin fullerene-based devices) and PC[70] BM ≈200 nm (thick fullerene-based devices) film thickness. The devices were encapsulated using a UV-curable encapsulation epoxy and a small glass slide in order to avoid any possible ingress of moisture and then placed on a hotplate at 85 °C in a N$_2$-filled glove box under dark. The devices were taken outside of the glove box to measure the PV device performance parameters and undergo the characterization process. The devices were then placed back again inside the glove box to continue the aging process. This procedure was repeated at 24 h intervals for up to 168 h. The deterioration of the average normalized photovoltaic parameters from eight devices from each device set was monitored over the course of 168 h at 24 h intervals, and is shown in Figure 2. We note that the experimental lifetime behavior of the solar cells under study reported in Figure 2 was repeatedly observed in four independent experimental runs under the same conditions.

From Figure 2 it can be seen that there are striking differences between the two sets of devices. The thin fullerene-based device exhibits a very sharp drop to all of their photovoltaic parameters only just after 24 h of heating. The drop in photovoltaic parameters seems to stabilize at the 72 h mark. However, the device...
PCE has already dropped well below 20% by that point. On the other hand, thick fullerene-based device retains almost 100% of the PCE even after 168 h of heating. Although there is a noticeable drop of the $V_{oc}$ at $\approx$80% at the first 24 h, there is an equally increase in $J_{sc}$ (115%), and especially FF (140%), for this device. Both the drop in $V_{oc}$ and increase in $J_{sc}$ and FF are stabilized after 24 h, allowing the thick fullerene-based device to preserve its high PCE ($\approx$100%) even after 168 h of heating.

Representative devices from each device configuration were chosen for characterization. The representative devices yielded the following photovoltaic parameters: thin fullerene-based device ($V_{oc} = 0.91$ V, $J_{sc} = 14.75$ mA cm$^{-2}$, FF = 79.2%, PCE = 10.64%), thick fullerene-based device ($V_{oc} = 0.87$ V, $J_{sc} = 10.57$ mA cm$^{-2}$, FF = 40.8%, PCE = 3.75%). These measurements indicated a considerable decrease in FF when we increase the thickness of PC[70]BM, due to the limited electrical conductivity of the material.\(^4\) The purpose of this study was to identify the major degradation pathway of the CH$_3$NH$_3$PbI$_3$ formulation-based devices under accelerated heat conditions and to evaluate the effect of different fullerene thickness on the device lifetime performance. For the above reasons achieving the highest possible PCE for each device set was not within the targets of this experimental plan.

Figure 3 shows the illuminated $J$–$V$ characteristic of fresh and aged devices after 96 h of heating. The 96 h mark was chosen as the point of degradation where all the characterization studies were performed on the examined samples. The reason for this choice was that at the 96 h mark the decline of the photovoltaic parameters has been substantially reduced. As it can be seen from Figure 3, the shape of the $J$–$V$ curve for thin fullerene-based devices is severely distorted after 96 h of heating, which is consistent with the severe drop on photovoltaic parameters, as evident from Figure 2. Furthermore, significant signs of hysteresis start to appear within the device performance characteristics, which were absent from the fresh devices. Hysteresis in perovskite solar cells has been previously attributed, among other phenomena, to non-steady-state capacitive currents resulting from electrode polarization.\(^26\) Thus, the manifestation of hysteresis in the $J$–$V$ plot could point to electrode degradation in the device, which prevents the efficient extraction of carriers. On the other hand,
the thick fullerene-based devices do not exhibit any shape deterioration of the J–V plot or any hysteresis behavior after degradation. This correlates well with the stable performance of the thick fullerene-based devices, where the initial values of the photovoltaic parameters are mostly retained. Furthermore, the J–V plots of thick fullerene-based devices show an S-shaped profile growth prior to the heating test, which completely disappears after 96 h of annealing. An S-shape in the J–V curves has been previously attributed to poor charge extraction, which in our devices can be related with the large thickness of the PC[70]BM layer.\(^{27}\) The absence of the S-shape after 96 h heating could be a result of the light soaking effect from the constant light illumination during the characterization of the devices. It has been previously reported that this phenomenon is a result of the trap-assisted recombination of excitons,\(^{28}\) which can be strengthened due to the increase in trap density in the thick PC[70]BM film. The different \(J_{sc}\) behavior from both the lifetime plots and illuminated J–V characteristics was also observed when we performed photocurrent testing (PCT) measurements (Figure S2, Supporting Information). PCT is a very useful technique for degradation studies that have been previously used in organic photovoltaics (OPVs) as well\(^{29}\) for the visualization of how the current intensity is distributed inside a device. As it was expected, the device incorporating thin PC[70]BM layer exhibited a severe loss to current intensity, whereas the device with thick PC[70]BM layer retained its current intensity to a good degree.

Complementary to the previous characterization techniques, impedance spectroscopy was also used. Impedance spectroscopy is a very powerful technique that can be used to provide insight on the physical processes occurring inside the device.\(^{31}\) The impedance spectroscopy data were analyzed and represented using Nyquist as well as Mott–Schottky plots.

Figure 4a shows the standard shape of two frequency responses for perovskite solar cells (a high- and low-frequency feature). The feature at high frequencies has been previously attributed to charge transport resistance \((R_{tr})\) of the HTLs and ETLs as well as their interface with the perovskite active layer.\(^{30,31}\) The low-frequency feature has been attributed to the recombination resistance \((R_{rec})\) and ionic diffusion.\(^{30–32}\) From the results in Figure 4a, a significant decrease in \(R_{rec}\) is observed between fresh and aged thin fullerene-based devices as well as an increase in \(R_{tr}\). The increase to \(R_{tr}\) denotes that the ETL/HTL or their interface with the perovskite has been altered, making carrier movement inside the device more difficult. This in return results in a decrease of \(R_{rec}\) due to more frequent charge recombination events. The combination of increasing \(R_{tr}\) and decreasing \(R_{rec}\) is in good agreement with the decreased FF (≈40%) noted for thin fullerene-based devices, highlighting the difficulty for carrier extraction. On the other hand, the \(R_{tr}\) for thick fullerene-based devices remain relatively unchanged, whereas a slight increase of the \(R_{rec}\) was observed which could be tied with the normalized FF increase.

Mott–Schottky analysis is often used for the differentiation of processes occurring at the active layer with the ones occurring at the interfaces and outer contacts of the device.\(^{33}\) The capacitive plateau at negative voltages of Figure 4b provides information on the dielectric constant, which is an intrinsic property of the bulk material. A decrease in \(V_{oc}\) is observed (point of intersection of the slope with the x-axis) for both thin and thick fullerene-based devices, which denotes that the energy equilibration at the contacts has shifted.\(^{33}\) More importantly, \(V_{bi}\) for thin ETLs-based devices has dropped by ≈0.5 V whereas for thick fullerene-based devices only by ≈0.25 V. This is in accordance with the more rapid decrease of normalized \(V_{oc}\) for the thin ETLs-based devices compared to the thick fullerene-based devices, where the \(V_{oc}\) drop is a lot less apparent and more stable.

### 2.3. Buffer Layer Device Engineering Experimental Methods

In order to try and isolate the effect of heat in each layer of the device, buffer layer engineering methods were used. This is a powerful technique, which was also previously used in OPVs,\(^{28}\) allowing us to study the effect of the PV parameters in a fabricated device using aged films at 85 °C for 96 h, similar to the conditions used for device characterization. To achieve our goal, semifinished device structures (four device structures from each device set) were initially fabricated using aged ITO/PEDOT:PSS/CH\(_3\)NH\(_2\)PbI\(_3\) and aged ITO/PEDOT:PSS/CH\(_3\)NH\(_2\)PbI\(_3\)/PC[70]BM/AlZO device structures at 85 °C for 96 h. The best performing photovoltaic parameters of the aged device structures completed with fresh PC[70]BM/AlZO and Al, respectively, were compared with the values of fresh (not-aged) reference devices. The results are shown in Table 1.

From Table 1 aged device structures (ITO/PEDOT:PSS/CH\(_3\)NH\(_2\)PbI\(_3\) and aged ITO/PEDOT:PSS/CH\(_3\)NH\(_2\)PbI\(_3\)/PC[70]BM/AlZO) at 85 °C for 96 h with fresh PC[70]BM/AlZO/Al and fresh Al retain their photovoltaic performance with minor losses compared to the fresh reference devices. This further
highlights the intrinsic stability of MAPbI₃ and the important role that the top electrode interaction with the perovskite active layer plays for the stability of the device. If we compare the device structures which were fabricated using aged MAPbI₃ with the aged devices of Figure 3, aging the MAPbI₃ does not play a significant role in the degradation of the device since at 96 h the PCE drops below ~20% of its initial value, whereas the device structures (with aged active layer and fresh PC[70]0.17PB([0.83]Br0.17)3 formulation. [21] As already mentioned above, the isolation of these two layers is of utmost importance. Al and other commonly used electrode metals (Au, Ag, and Ca) have been previously reported to penetrate into fullerene layers even in the process of evaporation. This increases the probability of such metal and the perovskite layer to come in direct contact with each other chemically reacting. [35,36] This chemical reaction between the metal electrodes and perovskite leads to a change in the perovskite film color from dark brown to yellow due to the decomposition to PbI₂. [23] Figure 5 shows devices presented within this paper: (a) bottom side of fresh device stack, (b) bottom side of thermally aged device stack of thin PC[70]BM-based devices, (c) bottom side of thermally aged devices stack of thick PC[70]BM-based devices, (d) top side of fresh device stack, (e) top side of thermally aged device stack of thick PC[70]BM-based devices, and (f) top side of thermally aged devices stack of thick PC[70]BM-based devices.

From Figure 5b we can see that the perovskite film in thin PC[70]BM-based devices exhibited a color change from dark brown to yellow, indicating decomposition to PbI₂, upon thermally aging the whole device stack at 85 °C and N₂ similar to previously reported observations. [23] On the contrary, the perovskite retains its color at a respectable degree for devices incorporating thick PC[70]BM (Figure 5c). Furthermore, metallic-like spots start to appear in the thin PC[70]BM-based devices located at the bottom side of the device which were absent from both the fresh and thick PC[70]BM-based devices. Since the ITO/PEDOT:PSS/CH₃NH₃PbI₃ film has shown nondetectable signs of decomposition from the XRD and absorption spectra measurements, the color change from dark brown to yellow upon aging alongside the Al electrode could be a result of Al diffusion toward the active layer. The top side of the thick PC[70]BM device also shows some change in the color of Al electrode where it became less shiny and whiter (Figure 5e). This is an indication of Al corrosion due to diffusion of I⁻ as previously reported. [23] Both the fresh (Figure 5d) and aged thick fullerene-based devices (Figure 5f) do not show any apparent change in the Al electrode.

Recently reported time-of-flight secondary ion mass spectrometry (ToF-SIMS) measurements have shown metal diffusion of air stable metal such as Au in a Cs-containing FA₀.8₃M A₀.1₇Pb[I₀.8₃Br₀.1₇]₁₃ formulation. [21] As already mentioned above,

Table 1. Photovoltaic parameters for fresh reference and buffer layer engineered devices.

| Device type                                      | Vₜ [V] | Jₜ [mA cm⁻²] | FF [%] | PCE [%] |
|-------------------------------------------------|--------|---------------|--------|---------|
| Fresh reference device                           | 0.91   | 14.75         | 79.2   | 10.64   |
| Aged ITO/PEDOT:PSS/CH₃NH₃PbI₃ and fresh PC[70]BM/AZO/Al | 0.89   | 15.13         | 67.3   | 9.03    |
| Aged ITO/PEDOT:PSS/CH₃NH₃PbI₃/PC[70]BM/AZO and fresh Al | 0.84   | 16.43         | 73.2   | 10.10   |

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degradation due to diffusion of $I^-$ is often a result of the intrinsic decomposition of the perovskite layer and introduction of iodine vacancies in the process. Even though decomposition of the perovskite layer was not detectable, in the process of characterization via XRD (from the introduction of PbI$_2$ peaks) and reduction of the absorption spectra, it can still be present. Although the lack of changes in absorption and XRD results reported within this paper indicating the lack of degradation in MAPI, it is well known that MAPI and cell degradation can start long before those characterization methods will detect changes.

The results reported in the buffer layer engineering section, showing that aging the semifinished ITO/PEDOT:PSS/CH$_3$NH$_3$PbI$_3$ and ITO/PEDOT:PSS/CH$_3$NH$_3$PbI$_3$/PC$_{60}$BM/AZO device stacks does not play a detrimental role to the PV parameters of the fabricated device. In a recent work, Akbulatov et al. have shown that similar device structures of ITO/PEDOT:PSS/CH$_3$NH$_3$PbI$_3$/PC$_{60}$BM/Ag can easily degrade under open-circuit conditions, constant illumination at 70 mW cm$^{-2}$, and 60 °C thermal aging even in inert conditions. Through ToF SIMS and theoretical calculations it was shown that the degradation had occurred through liberation of volatile methylammonium iodide (MAI) and formation of AgI in the Ag electrode. It was also reported through ToF SIMS that the Ag electrode showed some sign of penetration inside the PC$_{60}$BM upon aging. The experimental results reported in this paper show that buffer layer engineering using thick fullerene-electron transporting/passivation layers can be used to block diffusion and improve the thermal stability of inverted perovskite PVs.

### 2.4. Reducing the Efficiency–Stability Gap of Perovskite Photovoltaics

Using a 200 nm thick PC$_{70}$BM ETL we were able to isolate the CH$_3$NH$_3$PbI$_3$ active layer from the metal electrode, a procedure that effectively improves the lifetime performance of device upon heating at 85 °C from 24 h to over 168 h. This improved stability however is attained at the cost of strong PCE reduction due to the thick PC$_{70}$BM fullerene-based buffer layer used for the isolation of the perovskite active layer from the top metal electrode. Furthermore, despite the positive effects of the toluene CH$_3$NH$_3$PbI$_3$ washing step in efficiency this processing step can introduce some reliability issues on the reproducibility of the device performance. To reduce the efficiency–stability–reliability gap of perovskite photovoltaics, we have tested the behavior of the proposed diffusion blocking layer to devices that are optimized to work effectively with thick PC$_{70}$BM. These devices were based on a highly reliable perovskite formulation of CH$_3$NH$_3$PbI$_x$Cl$_{1-x}$. Typically, devices based on CH$_3$NH$_3$PbI$_x$Cl$_{1-x}$produce perovskite films with high roughness ($R_{ms} \approx 12.45$ nm) compared to the previously used CH$_3$NH$_3$PbI$_3$ formulation that produces films with relatively low roughness ($R_{ms} \approx 4.21$ nm), as shown in the atomic force microscopy (AFM) height images in Figure S3 (Supporting Information).

![Figure 6](image-url) Normalized a) $V_{oc}$, b) $J_{sc}$, c) FF, and d) PCE of CH$_3$NH$_3$PbI$_3$ and CH$_3$NH$_3$PbI$_{x}$Cl$_{1-x}$-based devices over 1000 h of heating at 60 °C and N$_2$ atmosphere.
Information). The thicker PC[70]BM is therefore necessary to ensure uniform coverage of the rough perovskite films and to improve device reliability.

Using CH$_3$NH$_3$PbI$_3$-Cl$_x$ and thick (200 nm) diffusion based PC70BM blocking layer we have managed to achieve devices with $V_{oc} = 0.88$ V, $J_{sc} = 12.99$ mA cm$^{-2}$, FF = 62.3%, PCE = 7.11% (average), and a champion device with $V_{oc} = 0.90$ V, $J_{sc} = 15.79$ mA cm$^{-2}$, FF = 66.8%, and PCE = 9.48% (Figure S4, Supporting Information). The heat stability of these devices was also tested and compared with the efficient CH$_3$NH$_3$PbI$_3$-based devices using optimized for efficiency thin (70 nm) fullerene ETL. The lifetime profile of the average normalized PV parameters from eight devices of each device set is shown in Figure 6. We note that the experimental lifetime behavior of the solar cells under study reported in Figure 6 was repeatedly observed in three independent experimental runs under the same conditions.

It is observed that the devices which are based on CH$_3$NH$_3$PbI$_3$-Cl$_x$ with thick PC[70]BM show impressive heat stability for up to 1000 h. It has been reported that CH$_3$NH$_3$PbI$_3$-Cl$_x$ exhibits better intrinsic stability over other perovskite formulations such as CH$_3$NH$_3$PbCl$_x$ and CH$_3$NH$_3$PbI$_3$.[39] It is important to note that based on the experimental conditions used within this paper we have not observed any major intrinsic degradation of the CH$_3$NH$_3$PbI$_3$ from the XRD and UV–vis. Furthermore, we have shown using buffer layer device engineering experimental methods (please see Section 2.3) that using thermally aged CH$_3$NH$_3$PbI$_3$ and fresh top electrode yields almost no change to the efficiency parameters of the device compared to a fresh-reference device. Therefore, we believe that the improved stability of the CH$_3$NH$_3$PbI$_3$-Cl$_x$-based devices tested is mainly due to the incorporation of the thick PC[70]BM buffer layer that can be used to eliminate top electrode thermally activated degradation mechanisms of inverted perovskite photovoltaics.

3. Conclusion

To conclude we have proposed that the interaction of the Al metal electrode with the perovskite active layer through diffusion mechanisms is the main thermal degradation mechanism for inverted perovskite solar cell. Using a 200 nm thick PC[70]BM ETL we were able to reduce the effect of diffusion mechanisms and isolate the CH$_3$NH$_3$PbI$_3$ active layer from the metal electrode, a procedure that effectively improves the lifetime device performance upon accelerating lifetime heating at 85 °C from 24 h to over 168 h. This improved stability however is attained at the cost of reduced PCE for the CH$_3$NH$_3$PbI$_3$-based perovskite solar cells due to the thick PC[70]BM fullerene buffer layer used for the isolation of the CH$_3$NH$_3$PbI$_3$ active layer from the top Al metal electrode. To reduce the efficiency–stability gap of perovskite photovoltaics, we have tested the proposed diffusion blocking layer to highly reliable CH$_3$NH$_3$PbI$_3$-Cl$_x$-based devices that are optimized to work effectively with thick PC[70]BM. We have presented a hysteresis-free CH$_3$NH$_3$PbI$_3$-Cl$_x$ inverted perovskite PVs with thick PC[70]BM diffusion blocking layer that exhibit PCE in the range of 10% and thermal stability for over 1000 h at 60 °C.

4. Experimental Section

CH$_3$NH$_3$PbI$_3$-Cl$_x$-Based Device Fabrication: The commercial I201 perovskite precursor ink (nitrogen processing) by Ossila Ltd. was used. The precursor was heated at 70 °C inside a N$_2$ glove box for 2 h and then left to cool down to room temperature. ITO-patterned glass substrates (sheet resistance = 4.12 $\Omega$ sq$^{-1}$) were cleaned using an ultrasonic bath for 10 min in acetone followed by 10 min in isopropanol. The PEDOT:PSS films were prepared using the AI4083 solution from Heraeus. PEDOT:PSS was filtered using 0.22 µm polyvinylidene difluoride (PVDF) filters before coating. The PEDOT:PSS films were coated using spin-coating at 6000 rpm for 30 s followed by annealing at 150 °C for 10 min. The perovskite films were prepared inside a N$_2$ glove box using spin-coating at 4000 rpm for 30 s followed by annealing at 80 °C for 2 h. PC[70]BM (50 mg mL$^{-1}$) was then coated using spin-coating at 1000 rpm for 30 s. The AZO solution was prepared using a commercial solution (N-20X) from Avantama Ltd. AZO films were coated using spin-coating at 1000 rpm for 30 s. The Al metal was finally thermally evaporated achieving a thickness of 100 nm. The devices were then encapsulated using a small glass slide and a UV-curable encapsulation epoxy from Ossila Ltd.

CH$_3$NH$_3$PbI$_3$-Based Device Fabrication: The perovskite solution was prepared using a mixture (1:1 w) of Pbl$_3$ from Alfa Aesar and MAI from GreatCell Solar. The mixture was dissolved in a mixed solvent (7:3) of γ-butyrolactone and dimethyl sulfoxide (DMSO). The solution was stirred at 60 °C for 1 h. The preparation of PEDOT:PSS films is identical to the procedure followed for the fabrication of CH$_3$NH$_3$PbI$_3$-Cl$_x$-based devices. The perovskite solution was left to cool at room temperature inside the glove box followed by filtering using a 0.22 µm PVDF filter. The perovskite films were coated using a three-step spin-coating process: first step 500 rpm for 5 s, second step 1000 rpm for 45 s, and third step 5000 rpm for 45 s. During the third step, after the first 20 s of the duration of the step, 0.5 mL toluene was dropped onto the spinning substrate to achieve rapid crystallization of the films. The resulting perovskite films were annealed at 100 °C for 10 min. The PC[70]BM film was coated using a solution of 20 and 50 mg mL$^{-1}$ to achieve fullerene thickness of 70 and 200 nm, respectively, at 1000 rpm for 30 s. The AZO and Al coating was identical to the CH$_3$NH$_3$PbI$_3$-Cl$_x$-based formulation.

Characterization Techniques: The thickness of the fullerene layers was measured using a Veeco Dektak 150 profilometer. XRD measurements were performed using PANalytical X’tal’s x pert pro MPD powder diffractometer (40 kV, 40 mA) using Cu Kα radiation (λ = 1.5418 Å). Optical measurements for the absorption spectra were performed with a Schimadzu UV-2700 UV–vis spectrophotometer. The AFM measurements were performed using Easy Scan-2 Nanosurf in tapping mode (phase contrast mode) and the roughness was extracted using the Gwydion software. The current density–voltage (J–V) characteristics were characterized with a Botest LIV Functionality Test System. Both forward and reverse scan directions were measured with 10 mV voltage steps and a time step of 40 ms. For illumination, a calibrated Newport Solar simulator equipped with a Xe lamp was used, providing an AM1.5G spectrum at 100 mW cm$^{-2}$ as measured by a certified oriel 91150 V calibration cell. A custom-made shadow mask was attached to each device prior to measurements to accurately define the corresponding device area. The photocurrent measurements were performed using a Botest PCT 1 test system. Impedance spectroscopy was performed using an Autolab PGSTAT 302N equipped with FRA32M module. To extract the Nyquist plots, the devices were illuminated using a red LED at 625 nm and 100 mW cm$^{-2}$. A small AC perturbation voltage of 10 mV was applied and the current output was measured using a frequency range of 1 MHz to 1 Hz. The steady-state DC bias was kept at 0 V. The C–V measurements for the Mott–Schottky plots were performed under dark using a steady frequency of 5 kHz throughout a voltage range of −1 to 1 V.
Supporting Information

Supporting Information is available from the Wiley Online Library or from the author.

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Conflict of Interest

The authors declare no conflict of interest.

Keywords

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[1] A. Kojima, K. Teshima, Y. Shirai, T. Miyasaka, J. Am. Chem. Soc. 2009, 131, 6050.
[2] H. Shen, T. Duong, Y. Wu, J. Peng, D. Jacobs, N. Wu, K. Weber, T. White, K. Catchpole, Sci. Technol. Adv. Mater. 2018, 19, 55.
[3] Y. Liu, M. Bag, L. A. Renna, Z. A. Page, P. Kim, T. Emrick, D. Venkataraman, T. P. Russel, Adv. Energy Mater. 2015, 6, 1501606.
[4] A. Savva, I. Burgues-Ceballos, S. A. Choulis, Adv. Energy Mater. 2016, 6, 1600285.
[5] G. E. Eperon, S. D. Strank, C. Menelaou, M. B. Johnston, L. M. Herz, H. J. Snaith, Energy Environ. Sci. 2014, 7, 982.
[6] J. H. Noh, S. H. Im, J. H. Heo, T. N. Mandal, S. Seong, Nano Lett. 2013, 13, 1764.
[7] C. Wehrenfenning, G. E. Eperon, M. B. Johnston, H. J. Snaith, L. M. Herz, Adv. Energy Mater. 2014, 26, 1584.
[8] H. J. Snaith, J. Phys. Chem. Lett. 2013, 4, 3623.
[9] C. Stoumpos, C. D. Malliakas, M. G. Kanatzidis, Inorg. Chem. 2013, 52, 9019.
[10] H. Choi, C. K. Mai, H. Kim, S. Song, G. C. Bazan, J. K. Kim, A. J. Heeger, Nat. Commun. 2015, 6, 8348.
[11] J. Cao, J. Yin, S. Yuan, Y. Zhao, J. Li, N. Zheng, Nanoscale 2015, 7, 9443.
[12] I. C. Smith, E. T. Hoke, D. Solis-Ibarr, M. D. McGhee, H. I. Karunadasa, Angew. Chem. 2014, 126, 11414.
[13] T. Leitjnes, K. Bush, R. Chechaaroen, R. Beal, A. Bowring, M. D. McGehee, J. Mater. Chem. A 2017, 5, 11483.
[14] N. Aristidou, I. Sanchez-Molina, T. Chothuangchutchaval, M. Brown, L. Martinez, T. Rath, S. A. Haque, Angew. Chem., Int. Ed. 2015, 54, 8208.
[15] R. Fang, S. Wu, W. Chen, Z. Liu, S. Zhang, R. Chen, Y. Yue, L. Dong, Y. Cheng, L. Han, W. Chen, ACS Nano 2018, 12, 2403.
[16] Z. Yang, J. Xie, V. Arivazhagan, K. Xiao, Y. Qiang, K. Huang, M. Hu, C. Cui, X. Yu, D. Yang, Nano Energy 2017, 40, 345.
[17] J. A. Pearson, G. E. Eperon, P. E. Hopkinson, S. N. Habirseutinger, J. T. Wang, H. J. Snaith, N. C. Greenham, Adv. Energy Mater. 2016, 6, 1600014.
[18] M. Alsari, A. J. Pearson, J. T. Wang, Z. Wang, A. Montisci, N. C. Greenham, J. H. Snaith, S. Liliu, R. H. Friend, Sci. Rep. 2018, 8, 5977.
[19] B. Conings, J. Drijikoningen, N. Gauquelin, A. Babayigit, J. D’Haen, L. D’Ollieslaeger, A. Ethisrajan, J. Verbeeck, J. Manca, E. Mosconi, F. De Angelis, H. Boyen, Adv. Energy Mater. 2015, 5, 1500477.
[20] X. Zhao, H. Kim, J. Seo, N. Par, ACS Appl. Mater. Interfaces 2017, 9, 7148.
[21] K. Domanski, J. Correa-Baena, N. Mine, M. Nazeeruddin, A. Abate, M. Saliba, W. Tress, A. Hagfeldt, M. Gratzel, ACS Nano 2016, 10, 6306.
[22] N. Tessler, Y. Vaynzof, Appl. Energy Mater. 2018, 1, 676.
[23] E. Bi, H. Chen, F. Xie, Y. Wu, W. Chen, Y. Su, A. Islam, M. Gratzel, X. Yang, L. Han, Nat. Commun. 2017, 8, 15330.
[24] P. W. Liang, C. Chueh, S. T. Williamsa, A. Jen, Adv. Energy Mater. 2015, 5, 1402321.
[25] Y. Fang, C. Bi, D. Wang, J. Huang, ACS Energy Lett. 2017, 2, 782.
[26] B. Chen, M. Yang, S. Priya, K. Zhu, J. Phys. Chem. Lett. 2016, 7, 905.
[27] M. Glatthaar, M. Riede, N. Keegan, K. Sylvesset-Hivid, B. Zimmermann, N. Nigemann, A. Hinsch, A. Gombert, Sol. Energy Mater. Sol. Cells 2007, 91, 390.
[28] S. Shao, M. Abdu-Aguey, L. Qiu, L. Lai, J. Liu, S. Adjokatse, F. Jahanhi, M. E. Kamminga, H. Brink, T. M. Palstra, B. Kooi, J. C. Hummelen, M. A. Loi, Energy Environ. Sci. 2016, 9, 2444.
[29] F. Herrersmidt, A. Savva, E. Georgiou, S. M. Tuladhar, J. R. Durrant, I. McCulloch, D. D. C. Bradley, C. J. Brabec, J. Nelson, S. A. Choulis, Appl. Energy Mater. Interfaces 2017, 9, 14136.
[30] M. Bag, L. A. Renna, R. Y. Adhikari, S. Karak, F. Liu, P. M. Lahti, T. P. Russel, M. T. Tuominen, D. Venkataraman, J. Am. Chem. Soc. 2015, 137, 13130.
[31] F. Galatopoulos, A. Savva, I. T. Papadas, S. A. Choulis, J. Am. Chem. Soc. 2017, 5, 076702.
[32] A. Guerrero, G. Garcia-Belmonte, I. Mora-Sero, J. Bisquert, Y. Kang, J. Jacobson, J. Correa-Baen, A. Hagfeldt, J. Phys. Chem. C 2016, 120, 8023.
[33] A. Guerrero, J. Yu, C. Aranda, Y. Kang, G. Garcia-Belmonte, H. Zhou, J. Bisquert, Y. Yang, ACS Nano 2016, 10, 218.
[34] H. Back, G. Kim, J. Kim, T. Kim, H. Kang, J. Lee, S. Lee, K. Lee, Energy Environ. Sci. 2016, 9, 1258.
[35] J. You, L. Meng, T. Song, T. Guo, Y. Yang, Y. Michael, W. Chang, Z. Hong, H. Chen, H. Zhou, Q. Chen, Y. Liu, N. De Marco, Y. Yan, Nat. Nanotechnol. 2016, 11, 76.
[36] G. Zhang, S. Hawks, C. Ngo, L. Schelhas, T. Scholes, H. Kang, J. Aguirre, S. Tolbert, B. Schwartz, ACS Appl. Mater. Interfaces 2015, 7, 25247.
[37] K. Brinkmann, J. Zhao, N. Pourdavoud, T. Becker, T. Hu, S. Oltbo, K. Meerholz, L. Hoffmann, T. Gahlmann, R. Heiderhoff, M. F. Osajca, N. A. Luechinger, D. Rottger, Y. Chen, B. Cheng, R. Heiderhoff, Nat. Commun. 2017, 8, 13938.
[38] A. F. Akbulatov, L. A. Frolova, M. P. Griffin, I. R. Gearba, A. Dolocan, D. A. Vanden Bout, S. Tsarev, E. A. Katz, A. F. Shchetkov, K. J. Stevenson, P. A. Troshin, Adv. Energy Mater. 2017, 7, 1700476.
[39] D. Wang, Z. Liu, Z. Zhou, H. Zhu, Y. Zhou, C. Huang, Z. Wang, H. Xu, Y. Jin, B. Fan, S. Pang, G. Cui, Chem. Mater. 2014, 26, 7145.