Full Paper

Investigating the Superoxide Formation and Stability in Mesoporous Carbon Perovskite Solar Cells with an Aminovaleric Acid Additive

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Perovskite solar cells have attracted a great deal of attention thanks to their high efficiency, ease of manufacturing, and potential low cost. However, the stability of these devices is considered their main drawback and needs to be addressed. Mesoporous carbon perovskite solar cells (m-CPSC), consisting of three mesoporous layers (TiO₂/ZrO₂/C) infiltrated with CH₃NH₃PbI₃ (MAPI) perovskite, have presented excellent lifetimes of more than 10 000 h when the additive NH₂(CH₂)₄CO₂HI (5- aminovaleric acid iodide; 5-AVAI) is used to modify the perovskite structure. Yet, the role of 5-AVAI in enhancing the stability has yet to be determined. Here, superoxide-mediated degradation of MAPI m-CPSC with and without the 5-AVAI additive is studied using the fluorescence probe dihydroethidium for superoxide detection. In situ X-ray diffractometry shows that aminovaleric acid methylammonium lead iodide (AVA-MAPI) perovskite infiltrated in mesoporous layers presents higher stability in an ambient environment under illumination, evidenced by a slower decrease of the MAPI/PbI₂ peak ratio. Superoxide yield measurements demonstrate that AVA-MAPI generates more superoxide than regular MAPI when deposited on glass but generates significantly less when infiltrated in mesoporous layers. It is believed that superoxide formation in m-CPSC is dependent on a combination of competitive factors including oxygen diffusion, sample morphology, grain size, and defect concentration.

1. Introduction

Halide perovskite solar cells (PSCs) have attracted great interest in the past decade thanks to their excellent optical properties, high carrier mobility, and diffusion length producing photovoltaic devices with relatively high power conversion efficiency (PCE).

Recent advances have led to impressive 25.2% PCE in 2019 giving merit for commercialization of this technology. However, the stability of PSCs needs to be addressed to realize the goal of mass production. PSCs can exhibit poor stability (of the order of days to weeks without encapsulation) due to degradation of the perovskite layer caused by moisture, oxygen, light, and combinations of these. In the case of methylammonium lead iodide (CH₃NH₃PbI₃, MAPI) perovskite, it has been shown that water hydrates MAPI leading to conversion into methylammonium iodide (MAI) and lead iodide (PbI₂). UV light can as well degrade perovskites through the photocatalytic effect of the mesoporous titanium dioxide layer in PSCs.

More recently, superoxide-mediated degradation has been identified as a significant contributor to degradation, aided by relatively quick diffusion of O₂ in the perovskite (=10⁻²–10⁻⁹ cm² s⁻¹, meaning that oxygen can completely diffuse into a 2 µm perovskite film in less than 20 s). Superoxide is formed when an excited electron, in the perovskite's conduction band, reduces molecular oxygen. Iodide vacancies, acting as trap states in the perovskite bandgap, have been shown to be energetically favorable sites for the formation of superoxide when occupied by a trapped electron. The formation of superoxide leads to the passivation of iodide vacancy trap states, reducing non-radiative recombination and therefore photobrightening the perovskite prior to superoxide-mediated degradation of the perovskite. Smaller grains tend to generate more superoxide than larger grains. The density of surface vacancies, which are well known to be more reactive than bulk
Figure 1. Superoxide formation in MAPI. If the conduction band minimum (CBM) of the electron transport layer (ETL) has a lower energy than the CBM of the perovskite, the electron can be injected in the ETL (left). In the opposite case, the photoelectron can react with molecular oxygen present in the air to form superoxide (right).

Figure 2. a) Schematic and b) band diagram of an m-CPSC. The photoactive area infiltrated with perovskite is highlighted in green.
assessed by monitoring the fluorescence of 2-hydroxyethidium, product of dihydroethidium and superoxide.[34] We compare the superoxide yield results to the degradation rate in air under illumination obtained with in situ X-ray diffractometry (XRD).

2. Results and Discussion

Over 100 h under illumination at room temperature and in air, the m-CPSC infiltrated with the AVA-MAPI solution presents a much higher stability compared to the MAPI sample (10% PCE loss compared to 88%, respectively) (Figure 3). This shows the improved stability of AVA-MAPI compared to MAPI consistent with previous reports.[31,33]

The degradation rate in air and under illumination of MAPI and AVA-MAPI infiltrated in ZrO₂, TiO₂/ZrO₂, and TiO₂/ZrO₂/C was studied through the evolution of the ratio of the (110) peak of tetragonal MAPI and the (001) peak of trigonal PbI₂ peak under illumination (film side toward light source) over a period of 11 h (Figure 4 and Figure S2, Supporting Information). MAPI infiltrated in ZrO₂ presents the highest degradation rate with the perovskite being almost entirely converted to lead iodide after 11 h. The same architecture using AVA-MAPI is much more stable in comparison with an PbI₂/MAPI ratio of 0.5 after 11 h indicating a significant amount of non-degraded perovskite. TiO₂/ZrO₂/MAPI shows a similar degradation rate to ZrO₂/MAPI and is also fully degraded after 11 h. TiO₂/ZrO₂/AVA-MAPI is initially less stable than ZrO₂/AVA-MAPI and displays a similar initial decay to ZrO₂/MAPI but slows after the first 2 h. TiO₂/ZrO₂/C stacks present higher stabilities than other samples using the same perovskite, likely thanks to inhibition of moisture degradation by the carbon layer. In particular, TiO₂/ZrO₂/C/AVA-MAPI shows the lowest degradation rate demonstrating the higher stability of AVA-MAPI compared to MAPI. AVA-MAPI infiltrated in any of the architectures studied here is therefore more stable than MAPI. In particular, AVA-MAPI infiltrated in ZrO₂ is less degraded after 11 h than MAPI infiltrated in the full stack of TiO₂/ZrO₂/C. This is somewhat surprising as it is expected that the thick carbon layer offers significant protection to the perovskite film. We believe this highlights the stability benefit of the 5-AVAI additive in air.

The superoxide formation of MAPI and AVA-MAPI was investigated by monitoring the fluorescence spectra of 2-hydroxyethidium. When drop-casted onto glass substrates the MAPI solution formed a film with large crystals while smaller crystals were obtained with the AVA-MAPI solution (Figure 5).

Glass/AVA-MAPI absorbs more light than glass/MAPI partially owing to the slightly higher surface coverage (56% and 52%, respectively, 8% more). We note that absorptance is evident above 820 nm likely due to scattering. Whether the absorptance is corrected by shifting it so that the absorptance at 850 nm is zero, or not, does not significantly affect the corrected superoxide yield (Figures S15–S19, Supporting Information). The corrected superoxide yield \( I(t)/I(t = 0) \) at 660 nm shows that glass/AVA-MAPI absorbs more light than glass/MAPI on the timescale studied (Figure 6). Glass/AVA-MAPI generates more superoxide than glass/MAPI on the timescale studied, consistent with the recent work of Lin et al.[24] Although AVA-MAPI has been reported to have a lower defect concentration than MAPI,[24] the smaller size of the crystals of AVA-MAPI lead to an increase surface/volume ratio which may increase the overall defect concentration of AVA-MAPI compared to MAPI, and thus may explain the higher superoxide yield measured. The variations of the probe fluorescence without a sample present were recorded twice as a baseline for our experiments. We observed a slight decrease of the fluorescence intensity over time which may be explained by the photodegradation of 2-hydroxyethidium.
The observed behavior of these perovskites infiltrated in mesoporous layers is significantly different than on glass. AVA-MAPI absorbs more light than MAPI in ZrO$_2$, and TiO$_2$/ZrO$_2$ mesoporous architectures, likely due to 5-AVAI improving the perovskite infiltration (Figure 7) and thus more perovskite being present in the mesoporous scaffold. Again, we also studied the superoxide yield in the case where the absorbance is assumed zero below the bandgap energy (Figures S15–S19, Supporting Information). The correction factors $\beta$ for these samples are given in Figure S13, Supporting Information. Figure 8 shows the superoxide yield of the MAPI and AVA-MAPI perovskite solutions drop-casted on ZrO$_2$, TiO$_2$/ZrO$_2$, and TiO$_2$/ZrO$_2$/C. Similar trends are observed independently of the direction of illumination (i.e., glass side or film side) other than for TiO$_2$/ZrO$_2$/C samples. While superoxide formation is measured in these samples when illuminated through the glass, no or very little superoxide formation is observed when illuminated through the film side as the carbon top layer prevents the creation of excited carriers in the perovskite. TiO$_2$/ZrO$_2$/C samples illuminated through the film are thus ignored for the rest of the discussion. We also note that the fluorescence intensity increase is not linear when the samples are illuminated though the film and this was due to exhaustion of the dihydroethidium probe (Figure S4, Supporting Information). Independently of the direction of illumination, ZrO$_2$/MAPI generates more superoxide compared to the other samples studied. When infiltrated in TiO$_2$/ZrO$_2$, the superoxide yield of MAPI decreases likely due to the possibility of charge injection into the mesoporous TiO$_2$. Little difference is observed between the superoxide yield of MAPI infiltrated in TiO$_2$/ZrO$_2$/C and TiO$_2$/ZrO$_2$ when illuminated through the glass. The carbon layer is non-selective and therefore can extract holes and electrons. Lower electron concentration in the perovskite reduces the chance of superoxide formation, however one might expect lower hole concentration to result in reduced recombination with electrons, increasing the excited electron lifetime, therefore increasing the chance of superoxide formation (Figure 8a). AVA-MAPI infiltrated in ZrO$_2$ generates the lowest superoxide of all the samples studied independently of the illumination direction. TiO$_2$/ZrO$_2$/AVA-MAPI presents a higher superoxide yield than ZrO$_2$/AVA-MAPI and similar to TiO$_2$/ZrO$_2$/MAPI. This is quite surprising as a reduction in the superoxide yield may be expected due to the presence of the mesoporous TiO$_2$ layer as observed with MAPI, we attribute this, at least partly, to changes in morphology as discussed below. Repeat measurements showed that AVA-MAPI generates...
less superoxide than MAPI when infiltrated in mesoporous TiO₂ (Figure S14b, Supporting Information). When illuminated through the glass substrate, TiO₂/ZrO₂/C/AVA-MAPI generates slightly less superoxide than in TiO₂/ZrO₂ but more than in ZrO₂ only (Figure 9). We note that given the difference in lamp power used during the superoxide measurement (1.5 mW cm⁻²) and the XRD experiment (19 mW cm⁻²), the results obtained in Figure 8 over 55 min are consistent with the ones obtained in Figure 4 during the first few minutes: AVA-MAPI samples present the lowest superoxide yield/highest stability, TiO₂/ZrO₂/MAPI is slightly more stable than ZrO₂/MAPI while TiO₂/ZrO₂/AVA-MAPI is less stable than ZrO₂/AVA-MAPI. The only difference with the superoxide experiment is observed for TiO₂/ZrO₂/C architectures which are more stable than the other stacks and which can be explained by the presence of the carbon layer which prevents moisture infiltration and moisture-mediated degradation of the perovskite due to its hydrophobic properties.

The observed superoxide yield difference in ZrO₂ and TiO₂/ZrO₂ can be partially explained by the morphology of the samples (Figure 10). All samples present capping layers with similar morphologies, with the presence of larger (>10 µm) and smaller (<10 µm) grains (Table 1). In the case of AVA-MAPI, the smaller grains form a continuous and smooth capping layer on top of the mesoporous layers. The larger grains are a few µm thick and appear to be located inside the capping layer formed by the smaller grains and not sitting on top of it (Figure S6, Supporting Information). The smaller crystals in MAPI samples are bigger and do not form a smooth layer leaving many pinholes. Since AVA-MAPI forms smaller grains with ergo a higher defect concentration, we could expect the superoxide yield to be higher than in MAPI samples (as observed on glass). However, we believe, pinholes observed in MAPI samples allow increased oxygen diffusion into the samples while the compact capping layer formed in AVA-MAPI samples hinders oxygen diffusion. This may explain the higher superoxide yield observed in MAPI compared to AVA-MAPI in both ZrO₂ and TiO₂/ZrO₂ architectures. The higher superoxide yield measured in TiO₂/ZrO₂/AVA-MAPI compared to ZrO₂/AVA-MAPI could be explained by the increased surface roughness of TiO₂/ZrO₂/AVA-MAPI (especially for the observed dark spot) resulting in increased surface area to react with oxygen thus counterbalancing the reduction in the concentration of excited state electrons from charge injection.

Figure 9. Comparison of the superoxide yield of MAPI and AVA-MAPI samples illuminated through a) the glass and b) the film after 55 min (see Tables S1 and S2, Supporting Information for values).

Figure 10. Top-down SEM images of the capping layer of MAPI and AVA-MAPI drop-casted in ZrO₂ with inserts showing a photograph of the sample. The TiO₂/ZrO₂/AVA-MAPI sample presented lighter and darker areas as indicated by the black arrows.
Perovskite grain size in TiO2/ZrO2 calculated from 24 measurements for each image.

| Perovskite grain size in TiO2/ZrO2 [µm] | Perovskite grain size in TiO2/ZrO2 [µm] |
|----------------------------------------|----------------------------------------|
| Large | Small | Large | Small |
| AVA-MAPI | 17.6 ± 3.0 | 12.1 ± 0.3 | 21.9 ± 7.8 | 1.6 ± 0.4 |
| MAPI | 15.2 ± 3.4 | 7.6 ± 2.4 | 11.8 ± 2.0 | 7.0 ± 1.3 |

See Figure S7, Supporting Information for more details.

The photoluminescence (PL) peak of ZrO2/AVA-MAPI is slightly narrower (full-width-at-half-maximum ≈ 44 nm) than TiO2/ZrO2/AVA-MAPI (≈ 50 nm) indicating a lower trap states concentration in the former (Figure S8, Supporting Information). 5-AVAI has been previously shown to slow down the movement of perovskite iodide ions in the TiO2/ZrO2 architecture.[35] This may then limit trap state passivation (e.g., iodide ions passivating iodide vacancies) leaving more trap state to react with oxygen to form superoxide. This is consistent with the slower photobrightening observed in TiO2/ZrO2/AVA-MAPI samples compared to ZrO2/AVA-MAPI (Figure 11). This needs to be put in perspective with passivation of trap states (and thus photobrightening) by superoxide and moisture. The photobrightening observed in Figure 11 is the result of ion migration, superoxide, and moisture. Ignoring the contribution of moisture, the slower photobrightening of the TiO2/ZrO2/AVA-MAPI sample indicates that passivation of trap states by superoxide is not enough to balance the slow ion migration. In both samples, large grains emit much less and photobrighten much slower than the small grains. Photodarkening of the small grains is observed in both samples and hints that the greater photobrightening in these grains is not due to higher enhanced trap state passivation by ion migration but rather trap state passivation by superoxide as the superoxide then degrades the perovskite. This is consistent with the higher defect concentration of small grains compared to large ones.

To further assess superoxide formation with respect to illumination direction, we studied further eight samples, namely MAPI and AVA-MAPI infiltrated in TiO2/ZrO2 (two samples of each, note new samples were used for each illumination direction) (Figure 12). Most excited charge carriers are localized within a few hundred nm in the film with respect to the illumination direction.[36] The superoxide yield does not change for MAPI depending on the illumination direction as oxygen can easily diffuse in the film and react with the excited charges. Conversely, less superoxide is formed when illuminating through the glass layer of AVA-MAPI than when illuminating from the film since the seemingly dense perovskite capping layer limits the rate of oxygen diffusion into the film. We believe this provides strong evidence for the increased oxygen diffusion in the MAPI samples and highlights the importance of the morphology in prevention of oxygen diffusion and hence superoxide formation. The processes and factors affecting superoxide formation are summarized in Figure 13.

3. Conclusion

We have studied the role of 5-AVAI in improving the stability of MAPI perovskite in m-CPSG. Changes in the perovskite and lead iodide XRD peak intensities under illumination showed higher degradation rates for MAPI samples than AVA-MAPI samples. Using dihydroethidium as a probe, the superoxide formation in MAPI and AVA-MAPI has been measured. AVA-MAPI deposited on glass presented a higher superoxide yield than MAPI. However, when drop-casted on ZrO2, TiO2/ZrO2, and TiO2/ZrO2/C mesoporous architectures, AVA-MAPI presented significantly lower formation of superoxide than MAPI. We have shown that superoxide formation in these samples is the result of a combination of multiple factors including oxygen diffusion, grain morphology, and defect concentration. AVA-MAPI samples have smaller grains with higher surface/volume ratio and therefore more surface defects which are counterbalanced by an overall lower defect concentration through the binding of AVA to iodide vacancies, and poor oxygen diffusion into the film due to the compact perovskite layer, resulting in an overall lower superoxide yield. MAPI samples have larger grains with lower surface/volume ratio and thus less surface defects which are balanced by higher defect concentration compared to AVA-MAPI and the increased oxygen diffusion in the film.
Infiltrated in mesoporous architectures

- Infiltrated lower defect concentration from passivated trap states due to 5-AVAI
- Higher defect concentration due to smaller crystals with higher surface/volume ratio
- Overall high defect concentration

Accessibility

TiO2/ZrO2/AVA-MAPi compared to ZrO2/AVA-MAPi which we believe is due to the very porous perovskite capping layer. Lower defect concentration measured when MAPI is infiltrated in TiO2/ZrO2 architecture compared to the ZrO2 architecture by the improved charge injection into the former architecture. Conversely, we observed a higher superoxide yield in TiO2/ZrO2/AVA-MAPi compared to ZrO2/AVA-MAPi which we believe is due to the higher perovskite porosity observed in the former architecture. All these factors are heavily influenced by the sample morphology and thus better understanding and control of morphology is required to help to lower superoxide formation and thus increase the device stability.

4. Experimental Section

Sample Manufacturing: MAPI precursor solutions were prepared from a 1:1 molar solution of lead iodide (PbI2, TCI) and MAI (Dyesol) in gamma-butyrolactone (Sigma-Aldrich). The AVA-MAPi solution was prepared as above for MAPI with the addition of 4% 5-AVAI, (Dyesol). In order to understand how each layer of the mesoporous structure affects the stability of the devices, a range of combinations of mesoporous layers were studied: ZrO2 only, TiO2, and ZrO2, and the full stack of TiO2, ZrO2, and C, all deposited on FTO glass (Figure 14).

The samples were prepared by screen-printing commercially available pastes as described earlier.[28, 29] The perovskite solutions were drop-casted on the different architectures, left at room temperature for 10 min, and then annealed at 50 °C for 1 h.

Power Conversion Efficiency Measurement: The current–voltage (J–V) curves of complete c-TiO2/TiO2/ZrO2/C devices were recorded (Oriel Sol3A solar simulator) under simulated AM1.5 sun illumination. The stability of these devices was measured in air. Between each measurement, the cells were kept at open circuit and under illumination (Oriel Sol3A solar simulator) under simulated AM1.5 sun illumination.

The samples were illuminated at 1.5 mW cm−2 with a Sylvania FVT 12V CY6,35 halogen lamp (Figure 15a) filtered with a 530 nm high pass filter and a near infrared filter to avoid degradation of the probe (Figure S1a, Supporting Information).
The fluorescence spectra of 2-hydroxyethidium were measured with a HORIBA FluoroMax-4 fluorometer ($\lambda_{\text{exc}} = 520 \text{ nm}$). The intensity ratio at 660 nm ($I(t)/I(t=0)$) was used to monitor the superoxide yield (note that although this yield was used to compare samples, these are not absolute yields of superoxide formation). The superoxide yields are corrected for absorbance using
\[
\beta \left( \frac{I(t)}{I(0)} \right)_{\text{corr}} = \frac{I(t)}{I(0)} + \frac{I(t) - I(0)}{\beta} \cdot \frac{I(0)}{I(0)}
\]
where $\beta$ is a correction factor calculated from the total number of photons absorbed (and hence, the total number of excited charge carriers) by the sample
\[
\beta = \int I_{\text{lamp}}(\lambda) \cdot A(\lambda) \cdot d\lambda
\]
with $I_{\text{lamp}}(\lambda)$ the intensity of the lamp and $A(\lambda)$ the absorbance of the sample at wavelength $\lambda$, and $S$ the area of the sample. The corrected superoxide yield is simply referred as $I(t)/I(t=0)$ in the main text. Superoxide formation was studied with illumination through the glass substrate and through the film (Figure 16b).

In Situ XRD: XRD measurements were performed using a Bruker D8. In situ degradation was performed by illuminating the samples at $\approx 19 \text{ mW cm}^{-2}$ with a LED Lenser M7R torch ($\approx 19 \text{ mW cm}^{-2}$—about $13 \times$ higher than for the superoxide experiment—Figure 16c and Figure S1b, Supporting Information). The compounds weight ratio evolution was then determined through the intensity ratio of the (110) tetragonal MAPI peak and the (001) PbI$_2$ peak using the relation
\[
I_{\text{MAPI}}/I_{\text{PbI}_2} \propto \omega_{\text{MAPI}}/\omega_{\text{PbI}_2}
\]
where $I_{\text{MAPI}}/I_{\text{PbI}_2}$ is the intensity of the peak associated with MAPI (PbI$_2$) and $\omega_{\text{MAPI}}/\omega_{\text{PbI}_2}$ is its respective weight.

Data Analysis: Data were processed with Python$^{[18]}$ with the numpy,$^{[40]}$ scipy,$^{[41]}$ scikit-image,$^{[42]}$ and OpenCV$^{[43]}$ packages, and figures were generated using the matplotlib package.$^{[44]}$ Surface coverage of grains was calculated using image thresholding and grain sizes were measured using the ImageJ software.$^{[45]}$

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

5-ammonium valeric acid iodide, AVA-MAPI, dihydroethidium, fluorescence, in situ x-ray diffractometry

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Figure 16. a) Superoxide setup. An external illumination source filtered with a 530 nm high pass and NIR filters is used to generate excited states in the sample. The sample is placed in a cuvette with 10 mL of dihydroethidium probe solution. The fluorescence of the probe solution is periodically measured using a fluorometer coupled with an automatic shutter. b) Schematic of the superoxide experiments during which samples were illuminated through the glass or through the film (i.e., the perovskite infiltration side), and c) XRD setup where a torch is used to illuminates the sample while the XRD pattern of the later is periodically measured.
