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Published in:
Journal of the Electrochemical Society

DOI:
10.1149/2.0671509jes

Published: 29/06/2015

Please cite the original version:
The Effect of Electrolyte Purification on the Performance and Long-Term Stability of Dye-Sensitized Solar Cells

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In this study the effect of the purification of electrolyte material on the performance and long-term stability of dye-sensitized solar cells is investigated. The combined effect of purifying all the electrolyte materials has been examined, as has the effect of purifying each compound to identify those compounds worth purifying and to eliminate unnecessary production steps on an industrial scale. Statistical methods were employed to draw statistically significant conclusions from the experimental results. No effect on the initial cell performance is found in this study. The purity of the electrolyte solvent (here methoxypropionitrile) is shown to have a remarkable effect on the cell lifetime: it could even double when the cell is properly purified. It is shown that exposing the cell to even relatively small amounts of UV light resulted in cell degradation through electrolyte bleaching during a 1000 hour aging test. Here the impurities in the electrolyte solvent lead to an almost doubled rate of electrolyte bleaching under UV light.

Material purification has been proposed to increase the stability of the cells. The use of highly pure materials is also regarded as an essential step in the preparation of most efficient cells. The reported data do not, however, include direct comparisons of purified and unpurified materials, which would be essential for assessing the importance of material purification in cell fabrication. From the industrial point of view, material purification increases the production costs of the cells. Thus the necessity of material purification should be studied carefully, just like any step that increases the complexity of the production. The objective of this study is to investigate if cell efficiency and stability increase as a result of the purification of the electrolyte material. The effect of the purification of all the electrolyte materials has been studied and the effect of each compound has been statistically compared and analyzed. The research continues in, where the effect of dye purification on cell efficiency and lifetime is studied. It is still quite common in the field to utilize only a few tests, mainly IV curve measurements in aging studies, or to test the cells only at the beginning and end of the aging. This is sufficient when the only purpose is to show that the cells remain stable but in order to gain a profound understanding of the aging mechanisms, a more detailed investigation is required. In this study, the cells are analyzed with a multitude of electrical, electrochemical, and visual tests: IV tests, electrochemical impedance measurements, cell photographing, quantum efficiency tests, and a combination of liquid chromatography and mass spectrometry. Automatic IV curve and impedance measurements are also performed continuously to monitor the trends in cell degradation.

In aging studies, the aging process is often accelerated by continuous, sunlight-simulating illumination, elevated temperatures or high air humidity. Still, a typical duration for an aging study is 1000 hours, which corresponds to approximately a year’s operation in sunlight in Northern European latitudes. The long duration makes it challenging to perform aging studies and treat the results because of the variability in test conditions, which is practically impossible to avoid, accumulate during the whole test period. One approach to this inherent challenge of aging tests is statistical treatment of the results. Statistical techniques can be used to compensate the differences in test conditions and evaluate the significance of the results. At the moment, the utilization of statistical tools is rare in the field of DSSC research. Here we present practical statistical methods (Pierce’s criterion, regression analysis, analysis of covariances, and one-way analysis of variances) applied to aging studies that we consider would bring significant improvements to the field. We suggest that these methods could be applied more generally in the investigations of dye solar cells.

Materials and Methods

Material purification.— The electrolyte components and the components of the dye solution were purified, especially in order to minimize the water absorbed in the materials. Thus, the material purification was performed under an inert atmosphere either using standard Schlenk techniques and an argon line or using a nitrogen-filled glovebox. Additionally, all the glassware was oven-dried before use.

The electrolyte consisted of iodine (I2), 1-methylbenzimidazole (NMBI), 1-propyl-3-methylimidazolium iodide (PMII), guanidinium thiocyanate (GuSCN) and 3-methoxypropionitrile (MPN). The iodine was not purified. The purification of the other materials required the use of purified solvents or reagents: tert-butyl methyl ether (TBME, Sigma-Aldrich), tetrahydrofuran (THF, Sigma-Aldrich), n-propyliodide and N-methylimidazole were stirred overnight at 60 °C and isolated by filtration in a glovebox. Additionally, all the glassware was oven-dried before use.

Statistical techniques can be used to compensate the differences in test conditions, which it is practically impossible to avoid, accumulate during the whole test period. One approach to this inherent challenge of aging tests is statistical treatment of the results. Statistical techniques can be used to compensate the differences in test conditions and evaluate the significance of the results. At the moment, the utilization of statistical tools is rare in the field of DSSC research. Here we present practical statistical methods (Pierce’s criterion, regression analysis, analysis of covariances, and one-way analysis of variances) applied to aging studies that we consider would bring significant improvements to the field. We suggest that these methods could be applied more generally in the investigations of dye solar cells.

Dye-sensitized solar cells (DSSC) have the potential to become a low-cost alternative to silicon solar cells in the future. They can be made out of sustainable materials, they have a wide application area, and they are on the threshold of large-scale production. They are also efficient: an 11.9% efficiency record for DSSCs has been achieved. At the moment one of the main challenges is the cell stability. DSSC stability for approximately 20 years in outdoor conditions is an essential requirement for the moment one of the main challenges is the cell stability. DSSC stability for approximately 20 years in outdoor conditions is an essential requirement.
After the purification the MPN was noticed to have changed color from slightly yellowish to fully transparent, which also indicates that some impurities were removed during the purification.

The test cells were dyed in purified cis-bis(isothiocyanato)-bis(2,20-bipyridyl-4,40-dicarboxylato)-ruthenium(II) tetra-8-8-diammonium (N719) dye, which was dissolved in tert-butanol (Sigma-Aldrich) and acetonitrile (Sigma-Aldrich) 1:1 solvent. The tert-butanol and acetonitrile were purified in a similar way to the isopropanol. Purified N719 was made from cis-bis(isothiocyanato)-bis(2,20-bipyridyl-4,40-dicarboxylato)-ruthenium(II) (N3) dye. N3 was prepared according to the procedure published in the literature. The preparation procedure for the N719 was the following: N3 dye that was dissolved in tert-butanol and water was purified by gel filtration on Sephadex LH20 using water as an eluent. The mean dark red-colored band was collected and titrated with 0.05-M HNO3 to pH 4.3. The resulting suspension was stirred for 3 h at room temperature and centrifuged. The precipitate was washed with water and centrifuged again. 2.5 eq of 40% solution of tert-butanol in water were added to the solid N719 thus obtained and the gel filtration and precipitation steps were repeated once more. Finally, the dye was dried in a vacuum. According to NMR spectroscopy (data not shown here), the spectrum of the self-prepared purified dye contained a smaller number of impurity peaks than the spectrum of commercially available dye (Solaronix), indicating that the self-prepared dye was purer.

### Table I. Compositions of electrolyte batches (A-F). It is indicated for each batch whether the electrolyte components were purified (P) or unpurified (U).

<table>
<thead>
<tr>
<th>Batch</th>
<th>I2</th>
<th>GuSCN</th>
<th>PMII</th>
<th>NMBI</th>
<th>MPN</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>P</td>
<td>P</td>
<td>P</td>
<td>P</td>
<td>U</td>
</tr>
<tr>
<td>B</td>
<td>P</td>
<td>P</td>
<td>P</td>
<td>U</td>
<td>P</td>
</tr>
<tr>
<td>C</td>
<td>P</td>
<td>P</td>
<td>U</td>
<td>P</td>
<td>P</td>
</tr>
<tr>
<td>D</td>
<td>P</td>
<td>U</td>
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<td>P</td>
<td>P</td>
</tr>
<tr>
<td>E</td>
<td>P</td>
<td>U</td>
<td>U</td>
<td>U</td>
<td>P</td>
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<tr>
<td>F</td>
<td>P</td>
<td>P</td>
<td>P</td>
<td>P</td>
<td>P</td>
</tr>
</tbody>
</table>

Aging tests and equipment.—. The test cells were aged for a total of 1000 hours under illumination. The aging tests consisted of three main parts: pre- and post-aging measurements, automatic measurements performed continuously during the aging, and manual measurements performed on a weekly basis during the aging.

First, the cells were subjected to light soaking at an illumination of approximately 0.5 Sun for 16 hours to stabilize the cell operation. Second, the following pre-aging tests were performed on the cells: current-voltage (IV) curve measurements and electrochemical impedance spectroscopy (EIS) at open circuit voltage ($V_{oc}$) in a solar simulator, IV curve measurements as a function of light intensity, and incident-photon-to-collected-electron efficiency (IPCE) tests.

The solar simulator was calibrated with an official calibration solar cell with a KG5 color glass filter (PV Measurements, Inc.) to a state equivalent to the 1000 Wm$^{-2}$ AM 1.5G (1 Sun) illumination spectrum in the visible range. The spectrum of the halogen lamps used in the solar simulator is shown in S.I. Fig. B1. The IV curves were measured with a Keithley 2420 SourceMeter with black tape masks attached on top of the cells. EIS at $V_{oc}$ under 1 Sun were performed with a Zahner Zennium potentiostat. In the impedance measurements, the frequency range $10^{-1}$-$10^{3}$ Hz was swept back and forth with 10 mV amplitude. The EIS data was fitted using the ZView2 program with an equivalent circuit presented in. The analysis of the EIS results is explained in more detail in the S.I. section C. An Autolab PGSTAT302 N potentiostat combined with a white LED light source built-in house was used for measuring the IV curves as a function of varying light intensity up to the equivalent of approximately 0.5 Sun. The IPCE tests were performed with a PV Measurements QEX7 Solar Cell Spectral Response Measurement System in DC mode with no bias light, and in the wavelength range 300–900 nm.

Third, the cells were aged. The light soaking device built in-house that was used for aging the cells uses the same lamps as the solar simulator and thus gives illumination equivalent to 1 Sun in the visible range. Calibrated this way, the lamps emit approximately 20% of the UV in the AM 1.5G spectrum (the spectrum of the lamps is shown in S.I. Fig. B1). The temperatures of all the cells stayed below 40 °C during the light soaking, which was confirmed by weekly infrared photographs (Flir InfraCam) and by monitoring the temperature continuously during the aging with thermocouples. Because of the large number of cells in the aging, the illumination level in the light soaking device was not as uniform as in the solar simulator. To be able to compensate for the variations in the light intensity, the light intensities of each cell were measured manually every time the cells were moved or a bulb in the light soaking device was changed.

During the aging, automatic IV measurements and EIS measurements at different voltages between the short and open circuit voltages of the cells were performed with a Solar Cell Aging Test Unit (SCATU) built in-house using a Bio-Logic SP-150 potentiostat with similar EIS settings to those that were used in the Zahner potentiostat. SCATU takes one measurement every five minutes, cycling through all the cells, so the degradation rate of the cells is monitored continuously.

In addition to the automatic SCATU tests, manual tests were performed on a weekly basis. These tests included EIS in the dark (Zahner Zennium potentiostat) and cell photographing (the system built in-house described in which includes an Olympus E-620 camera). The EIS settings were the same as in the measurements performed under illumination but the tests were repeated at a voltage range from...
0 V to 0.7 V at intervals of 0.1 V. Cell photographing was used for the quantitative monitoring of the changes in the color of the electrolyte over the aging test period. As the yellow color of the electrolyte comes mainly from iodine, the changes in the blue RGB pixel values of the photographs of the cell electrolyte can be related to the changes in the iodine concentration in the electrolyte.\cite{28} Cell photographing is also suitable for monitoring dye desorption to the electrolyte because desorption increases the red RGB pixel values of the photographs of the electrolyte.\cite{24} A detailed description of the photographing method and the photographing system can be found in the literature.\cite{24}

Fourth, the post-aging tests were performed after an aging period of 1000 hours. These tests were otherwise the same as the pre-aging tests but one more test was added: liquid chromatography-mass spectrometry (LC-MS) measurements for a couple of cells in each cell group. Dye was extracted from one cell from each batch with 0.1 M NaOH and analyzed with LC-MS according to a procedure described elsewhere.\cite{25} MS was performed in order to gain further information about the structure of the observed dye degradation products. The LC-MS results of the cells were compared with the results of both the pure starting material (dye solution) and reference cells that had not been aged under illumination. The results were not compared to the pre-aging measurements of each cell because LC-MS is a destructive analytical method.

Statistical methods for analyzing the aging data.— The experimental results in this study were subjected to statistical testing in order to compensate for the effects of random errors and varying light intensity during the aging test. One-way analysis of variances (ANOVA) combined with paired comparison tests was utilized for checking if the differences between the cell groups in the averages of the measured parameters were significant.\cite{26,27} For the aged data, regression analysis was applied for line plots showing result parameters as a function of the light intensity during the aging test. Linear regression lines and confidence intervals were calculated with the Matlab software with the Matlab functions polyfit and polyconf. Analysis of covariance (ANCOVA)\cite{27} is a method in the middle ground of ANOVA and regression analysis, used for the comparison of groups with respect to a variable, regardless of the covariate. Here, the performance of different cell groups was compared despite of the varying light intensity during the aging test with ANCOVA.

Before ANOVA or ANCOVA were performed, the results had to be pre-treated by removing significant outliers from the untreated measurement data. The selected exclusion method was Peirce’s criterion. It has been described accurately in the literature.\cite{28} This easy method is designed in such a way that it can be applied to several suspicious data points and it adjusts the criterion for excluding a data point to the number of data points.\cite{28} Peirce’s criterion is designed for detecting random errors in data sets without taking any additional parameters into account but it seems to function also for the data that contains external dependencies, such as the light intensity dependency. Like all outlier detection methods, Peirce’s criterion must be combined with further inspection of the data and the cells in order to avoid false positive and negative detections.

Results and Discussion

IV measurement results.— Initially, all the groups performed similarly, as shown in Fig. 1. This was also confirmed by the 95% confidence level ANOVA test described in Statistical methods for analyzing the aging data section. Surprisingly, the result indicates that the purification of the electrolyte components did not affect the initial cell performance. There seems to be no need to purify electrolyte components in studies that do not include cell stability testing, even though this has been quite common practice in the field.

The efficiencies of the cells in this study are moderate (Fig. 1) because of the cell geometry used in this study, and the geometry does not affect the possible chemical reactions related to the impurities in the electrolyte. Thus we believe that these results apply also for more efficient cells. Here, the cell geometry was not optimized for cell efficiency but for the ease of photographing the electrolyte and to mimic a small piece of a larger module so that the results would also be meaningful for upscaled cells. Therefore, the cell dimensions are rather large, increasing, e.g., the series connected resistances.

All the cells degraded during the aging (Fig. 1) and the degradation was related to the light soaking because the reference cells kept in the dark during the aging stayed stable. Cells containing similar ingredients to those used in this study have been reported to have passed 1000-hour aging tests on a regular basis\cite{26,27} and even in our own studies using the same aging system.\cite{24} The reasons that led to the aging of the cells in this study are analyzed later in this section. The degradation is mainly related to reduced $I_{sc}$ and $FF$, while $V_{oc}$ decreased only moderately (30–40 mV). Interestingly, quite similar cells exposed to a shock test at 100 °C and under UV illumination showed a similar 36 mV decrease in $V_{oc}$\cite{15}.

On the evidence of Fig. 1, it seems that after the aging, group A and possibly also group E performed worse than the other groups. The cells in both groups contain unpurified MPN, while the other groups contain purified MPN. This hypothesis was tested by a 95% confidence level ANCOVA test with average light intensity during the aging as a covariate by dividing the cells into six groups (A–F). The test did not confirm reliable differences between the individual groups because the variations within the groups are large compared to the differences in the mean values of the groups, making it difficult to analyze the differences. This is also clear to the eye from Fig. 1. When the group size was increased by dividing the samples into only two groups, purified MPN (B, C, D, F) and unpurified MPN (A, E), ANCOVA resulted in differences in performance between the groups: the cells containing unpurified MPN had a lower $I_{sc}$, $FF$, and $\eta$ after the aging than the cells containing purified MPN. Therefore the effect of the purity of the MPN seems relevant to cell stability when the effects of the purity of the other electrolyte components are neglected.

One significant reason that we suspect accounts for the deviations in the post-aging cell efficiencies within the cell groups is the variation in the light intensity. The test cell series is so large that it was not possible to keep the variations in the light intensity between the cells negligibly small during the aging test. Therefore, the cells might have aged at different rates and the differences accumulated during the whole aging period.

This effect has to be dealt with in all the measurements performed on the cells. Additionally, there are some measurements that are

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{figure1.png}
\caption{Efficiencies, fill factors, short circuit currents and open circuit voltages of the cells with group standard deviations before and after the 1000-hour aging test. The green and red dots represent the two reference cells from groups E and F, respectively, which were not aged under illumination but kept in the dark.}
\end{figure}
of the MPN in the cell is larger than the volume of other electrolyte components. Fig. 2 shows that after 700 hours of aging, the η of the cells containing purified MPN has mainly decreased because of decrease in FF. For cells containing unpurified MPN, the degradation has proceeded further, also causing a decrease in I_{sc}.

Some cells have been discarded from the regression line calculations in Fig. 2. Peirce’s criterion, described in Statistical methods for analyzing the aging data section, was utilized as the main guideline in the cell rejection. Peirce’s criterion was applied for the I_{sc} values of the cells when the cells were divided into two groups (purified and unpurified MPN) and into six groups (A, B, C, D, E, and F). Cell C3 was spotted as an outlier in the two-group test and cells B3, C3, D3, and F3 were spotted as outliers in the six-group test. The aging data of cell C3 were consistent with the rest of the data of the same cell and showed that the cell had degraded very fast. It is possible that C3 had become contaminated during the cell assembly. Therefore, cell C3 was discarded as an outlier. On further inspection of the I_{sc} data of cells B3 and D3 it was noticed that the values at 700 hours were not consistent with the values before or after that time. The recorded light intensity for these cells is probably erroneous at 700 hours, which creates the discontinuities in I_{sc} and makes the data at that time invalid. Therefore, these cells were discarded from the regression line calculations. Peirce’s criterion spotted cell F3 as an outlier in group F when the cells were divided into six groups because F3 had a somewhat low current (0.94 in Fig. 2c) compared to the other three cells in group F (on average 1.00 in Fig. 2c). The difference is actually quite small compared to the deviation of all the cells with purified electrolyte solvent and the IV values of cell F3 fit this group well. Therefore, F3 was regarded as a false positive result of the Peirce’s criterion test and was not discarded. Peirce’s criterion seems to be a useful method in the analysis of aging data even if the efficiencies of the cells analyzed in Fig. 2 had a notable light intensity dependence.

After Peirce’s criterion had been applied to the SCATU IV data, the data were tested by 95% confidence level ANCOVA and paired comparison tests that have been described in Statistical methods for analyzing the aging data section. The tests resulted in the following conclusions: Groups A and E differed from the other groups (B-D and F, containing purified MPN) in I_{sc} and η but resembled each other. In F, groups A and E resembled each other but, additionally, group E resembled groups B and F, which had a slightly lower FF than groups C and D. The differences in V_{oc} between the groups were quite small and paired comparison tests for V_{oc} showed just that group A differed from group F in having a higher average value. The results indicate that groups A and E perform worse than the other groups in the IV tests. The difference is clear in η and I_{sc}. For FF, the differences between the groups are not as large as for η and I_{sc} but still, group A performs worse than the other groups.

There is a significant difference between the regression line calculation and the ANCOVA tests: the first method is used for estimating the size of the difference between the groups and the latter is used for assessing whether any difference between the groups exists. Thus it is not odd that ANCOVA concluded with a high 95% confidence level that groups A and E behaved differently from the other groups, even if the size of the difference could not be estimated with such a high confidence level (70% confidence intervals of regression lines).

**Analysis of internal series connected resistances.**—The previous section showed that a great performance-limiting factor for the cells after 700 hours of aging is FF. Series connected resistances are the factors affecting FF: the series resistance, R_s, the resistance at the counter electrode/electrolyte interface, R_{CE}, and the steady state value of the diffusion impedance in the electrolyte at the counter electrode, R_{D}. These resistances are easy to investigate by EIS under illumination at V_{oc}.

EIS at V_{oc} was analyzed in its initial state and at 700 aging hours. Examples of the measurement results of the aged cells A6 (unpurified MPN) and F1 (purified MPN) and the fits to the equivalent circuit model are shown in S.I. Fig. E1. All the cells were analyzed but the only differences were found between cells containing unpurified MPN
The same post-aging data as shown in Fig. 3 are plotted cell-wise as a function of the average light intensity during the aging in Fig. 4. These graphs illustrate more clearly the differences between the two groups. First, the cells with an unpurified electrolyte solvent degraded regardless of the light intensity applied. Second, the standard deviations in this group are much higher than in the cells containing a purified solvent. On the basis of Fig. 4, \( R_{CE} \) and \( R_D \) seem to be relatively independent of the light intensity applied, at least in this intensity range. However, \( C_{CE} \) seems to decrease as the light intensity increases in the case of the cells containing unpurified MPN.

Analysis of photoelectrode performance with impedance measurements performed in the dark.— EIS in the dark as a function of voltage was analyzed at the moment of 700 aging hours to investigate possible changes at the photoelectrode. Only groups A (unpurified MPN) and F (purified MPN) were analyzed because the IV results presented in IV measurement results section showed that the purity of other cell components than the electrolyte solvent did not affect the cell stability. First, the equivalent circuit model of EIS (see S.I. Section C) was fitted to the measurement results. An example of typical data and the fits of both a fresh and an aged cell are shown in S.I. Fig. F1. The data are only shown for one cell (A6) as the data were very similar in all the cells.

The average fitted equivalent circuit model parameters, resistance, and capacitance at the photoelectrode/electrolyte interface, \( R_{PE} \) and \( C_{PE} \), and electron lifetime in the photoelectrode, \( \tau \), are shown in Fig. 5. There are no significant differences in the results of the two cell groups before and after the aging. This fact suggests that the purification of the electrolyte solvent did not affect the initial photoelectrode performance or photoelectrode degradation during the aging. \( R_{PE} \) decreased and \( C_{PE} \) increased slightly during the aging in both groups. A decrease in \( R_{PE} \) indicates that the recombination from TiO\(_2\) to the electrolyte has increased and thus the cell performance has decreased. The large standard deviations at 0.3 V and 0.4 V in \( C_{PE} \) arise from the fact that \( \log(C_{PE}(V)) \) is not linear but has a steep slope in this range. Thus the slight individual differences between the cells result in large differences in the values of \( C_{PE} \). Interestingly, \( \tau \) in Fig. 5c has not decreased but stayed rather constant during the aging and even increased to some extent at higher voltages.

Analysis of short circuit currents.— The \( I_s \) of the cells decreased during the aging, as Fig. 1 shows. The reasons for the decrease can be investigated by IPCE tests at low light intensity and IV measurements as a function of light intensity. \( I_s \) at IV tests performed under illumination of 1 Sun might be diffusion-limited, whereas in IPCE tests that are performed at low light intensity without bias light, the current is not diffusion-limited. Hence, the possible changes in the photocurrent-generating capability of the dye can be separated from the possible changes in the limiting current density, \( I_{lim} \), of the cell. As for the changes in \( I_{lim} \), they can be quantitatively estimated by IV measurements performed as a function of the light intensity.

IPCE was measured from one test cell in each cell group. The results for the aged and reference cells are shown in Fig. 6 and in S.I. Fig. G1, respectively. The IPCE curve is slightly lower for the cells with unpurified MPN. Actually the differences are so small that they do not significantly affect \( I_s \). This conclusion was checked by calculating the short circuit currents of the cells as

\[
I_{SC, TiO_2} = A q_l \int \Phi(\lambda) \eta_{IPCE}(\lambda, \lambda) d\lambda.
\]

where \( A \) is the cell area, \( q_l \) is the elementary charge, \( \lambda \) is the wavelength, \( \Phi(\lambda) \) is the flux of the halogen lamps used in this study, and \( \eta_{IPCE} \) is the IPCE efficiency of the cell,\(^3\) and comparing \( I_{SC, TiO_2} \) of

![Figure 3](image-url)
Figure 4. Counter electrode resistance and capacitance, $R_{CE}$ and $C_{CE}$, and diffusion resistance of the electrolyte, $R_D$, of the cells determined from EIS measured at $V_{oc}$ after 700 hours of aging as a function of the average light intensity during the aging.

The cells with purified and unpurified MPN. The results are shown in S.I. Table H1.

The original peaks in Fig. 6 did not move during the aging but an extra peak appeared at 400 nm for all the cells aged under illumination. The most likely reason for this peak is the bleached electrolyte: e.g., iodine in the electrolyte absorbs less light at shorter wavelengths, allowing the dye to absorb more light in this wavelength area. It is also noteworthy that the new peak does not appear in the IPCE curves of the reference cells E1 and F2, which were stored in the dark and apparently were not bleached. Thus, the electrolyte bleaching seems to be a photoinduced effect, which is in accordance with the literature.

The post-aging $I_{sc, TiO_2}$ calculated with Eq. 1 were compared to the actual $I_{sc}$ measured with the solar simulator to evaluate the effect of PE degradation on the decrease in $I_{sc}$. This is made possible by

Figure 5. Average photoelectrode resistance $R_{PE}$, photoelectrode capacitance $C_{PE}$, and electron lifetime $\tau$ shown as a function of impedance measurement voltage before and after the aging. The results for group A are marked in red and the results for group F in green. The “o” markers are for the pre-aging and the “x” markers for the post-aging results. The results show the standard deviations for the values.

Figure 6. Results of incident photon to collected electron efficiency (IPCE) measurements in absolute and relative values as a function of light wavelength. One cell has been measured for each group. The solid lines mark the pre-aging measurements and the dashed lines mark the post-aging (1000 h) measurements.
the fact that IPCE was measured at a low light intensity, where the current limitations do not occur, but the IV measurements were made at 1 Sun. Apart from one cell that remained very stable, all the aged cells had lower $I_{\text{lim}}$ than the IPCE results would have led us to assume: $I_{\text{lim},T1O_2}$ had decreased by 2-18%, whereas $I_{\text{lim}}$ had decreased by 35-80% (see S.I. Table H1). This indicates that mere PE degradation does not explain the decrease in $I_{\text{lim}}$ of the cells, but there are also current limitations in the cells.

The IPCE results and visual inspection showed that the electrolyte in most of the cells had bleached during the aging, indicating a loss of charge-transporting $I_3$ ions. Severe loss of $I_3$ can also lead to a decrease in $I_{\text{lim}}$, which was investigated by measuring $I_{\text{lim}}$ as a function of the light intensity (see S.I. Section J). The tests were performed before the aging and after 1000 hours of aging for one cell in each cell group. For some cells, $I_{\text{lim}}$ decreased during the aging test. The results for these cells are shown and explained in S.I. Section J. As a result, the IV curve measurements as a function of the light intensity suggest that unpurified MPN exposes the cells to a greater decrease in $I_{\text{lim}}$ of the cells, but there are also current limitations in the cells.

IPCE and $I_{\text{lim}}$ tests were performed for one cell in each cell group in order to limit the total measurement time. Therefore, it was not reasonable to perform statistical testing. ANOVA or ANCOVA, for these tests. Instead, these tests were used to support the more thorough tests: IPCE shows that the changes of the photodetectors observed in EIS are not the main reason for the reduced efficiency of the cells and $I_{\text{lim}}$ tests prove that the electrolyte bleaching observed in IPCE and visually (see Investigation of electrolyte bleaching section) is strong enough to limit the currents of the cells at 1 Sun.

Liquid chromatography-mass spectroscopy measurements.— LC-MS measurements supported the conclusions drawn from IPCE and EIS in the dark tests: all the aged cells were found to have degraded to some extent but no significant differences between the cells in terms of the amount and types of degradation products were observed. N719 was found to degrade by ligand substitution reactions between the dye and solar cell components such as solvent and electrolyte additives. The ruthenium complexes present in the extract were identified on the basis of their m/z ratio, fragmentation pattern, and comparison with data presented in the literature as $[\text{RuL}_2(\text{NCS})_2(\text{H}_2\text{O})]$, $[\text{RuL}_2(\text{NCS})(\text{MPN})]$, $[\text{RuL}_2(\text{NCS})(\text{NMBI})]$, $[\text{RuL}_2(\text{NCS})(\text{SCN})]$, and N719 or $[\text{RuL}_2(\text{NCS})_3]$. The amount of these degradation products could be determined only roughly by LC-MS analysis. Sample UV-chromatograms from one reference cell kept in the dark, a cell aged under illumination, and the dye used in the cells are presented in Fig. 7. The corresponding m/z ratios of the compounds that were identified are listed in Table II.

The major degradation product in the aged cells is $[\text{RuL}_2(\text{NCS})(\text{NMBI})]$ (peak 4, Fig. 7). This compound has previously been found to work as a sensitizer, but is approximately two times less efficient than N719.41 This compound was not present in the reference cells kept in the dark, which indicates that $[\text{RuL}_2(\text{NCS})(\text{NMBI})]$ is a photo-induced degradation product. $[\text{RuL}_2(\text{NCS})(\text{MPN})]$ and $[\text{RuL}_2(\text{NCS})(\text{H}_2\text{O})]$ are degradation products that appeared both in the cells aged under illumination and in the reference cells kept in the dark but were not present in the fresh dye. The effectiveness of these products as dyes has not been studied. The solvent substitution product $[\text{RuL}_2(\text{NCS})(\text{MPN})]$ seems to be a minor degradation product. The product is not stable in the extraction solution and may therefore be more prominent in the intact solar cell.

There was no significant difference between the cells containing unpurified and purified solvent, even though the unpurified solvent probably contains more water than the purified one. This is especially interesting when the minor degradation product $[\text{RuL}_2(\text{NCS})(\text{H}_2\text{O})]$ is considered. The LC-MS result suggests that the amount of $[\text{RuL}_2(\text{NCS})(\text{H}_2\text{O})]$ in the cell is actually not dependent on the amount of water in the cell. It is possible that the formation of $[\text{RuL}_2(\text{NCS})(\text{H}_2\text{O})]$ competes with the other degradation reactions, and that a higher amount of the water ligand would thus be present in cells with a different electrolyte composition containing the same amount of water. As a final remark, in the fresh

<table>
<thead>
<tr>
<th>Peak</th>
<th>Retention time (min)</th>
<th>m/z</th>
<th>Ruthenium complex</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>8.0</td>
<td>663.0</td>
<td>$[\text{RuL}_2(\text{NCS})(\text{H}_2\text{O})]^{+}$</td>
<td>Minor impurity formed in the cell even in dark.</td>
</tr>
<tr>
<td>2</td>
<td>9.9</td>
<td>731.0</td>
<td>$[\text{RuL}_2(\text{NCS})(\text{MPN})]^{+}$</td>
<td>Minor impurity formed in the cell even in dark.</td>
</tr>
<tr>
<td>3</td>
<td>11.6</td>
<td>705.0</td>
<td>$[\text{RuL}_2(\text{NCS})(\text{SCN}) - \text{H}]^{-}$</td>
<td>Impurity present in the dye used to assemble the cells.</td>
</tr>
<tr>
<td>4</td>
<td>11.8</td>
<td>778.0</td>
<td>$[\text{RuL}_2(\text{NCS})(\text{NMBI})]^{+}$</td>
<td>Major, photo-induced impurity.</td>
</tr>
<tr>
<td>5</td>
<td>12.3</td>
<td>705.0</td>
<td>$[\text{RuL}_2(\text{NCS})_2]^{+}$ or N3 (fully protonated N719)</td>
<td>Minor impurity formed in the cell even in dark.</td>
</tr>
</tbody>
</table>

Figure 7. Sample UV-chromatograms from one of the cells aged under illumination (top), one of the reference cells stored in the dark (middle), and the dye used to assemble the cells (bottom) (see Table II for peak identifications).
dye, [RuL2(NCS)(SCN)] was not detected by LC-MS analysis. In solar cells with the type of electrolyte composition used in this study, water probably reacts with N719 at a slower rate than NMBI. In this work these degradation products were detected in cells aged at a temperature below 40 °C. In this study and suggested to be caused by oxidative species generated under the illumination. The illumination spectrum of the light soaking lamps is shown in S.I. Fig. B1. It indicates that the amount of UV in the lamp illumination is just one fifth of the UV illumination in the AM 1.5G spectrum. Even small amounts of UV light might, however, be harmful in the long term if UV light triggers side reactions leading to irreversible electrolyte degradation, as has been suggested. In this case, the damage caused by the UV light would simply be linearly related to the amount of UV in the illumination instead of having some threshold value below which no damage would be caused.

One suggestion for the reason for the different bleaching rates of the cells containing purified and unpurified MPN is that unpurified MPN contains more impurities that, together with UV or visible light, trigger the bleaching reaction. MPN could contain, e.g., water, which would react with the TiO2 when exposed to UV light. In this case, the bleaching could be retarded but not completely prevented by applying a UV filter, as there are no perfect UV filters. This is suggested also by our other study. Therefore, the effect of the purity of the electrolyte on bleaching is important for the overall cell stability.

Evolution of iodine concentration in the electrolyte.— A difference in the bleaching rates of the cells containing unpurified and purified MPN was demonstrated in the previous section. The remaining questions are first, why cells have passed 1000-hour aging tests performed with the same light soaking device before but degraded in this study and second, whether the bleaching is caused by UV or visible light. These subjects are studied further in this section by estimating the evolution of the iodine concentration in the electrolyte in the cells of this study and comparing them to another study. The cell photographing results were used for estimating the iodine concentration as a function of the cumulative UV dose shown in Fig. 9. The cumulative UV dose is the product of the aging hours and the UV light level in proportion to the amount of UV light in AM1.5G illumination. A 0.03M concentration of iodine has been marked in Fig. 9 as representing the level below which current limitations probably appear in the cells.
The red and green lines in Fig. 9 represent the cells of this study. In this kind of plot the drastic difference in the bleaching rates of the cells containing purified and unpurified MPN is even more clearly visible than in Fig. 8: the cells with unpurified MPN cross the 0.03M line after roughly 20 UV1Sunh and the cells with purified MPN just after 45 UV1Sunh. Converted to the actual aging hours, these values are 100 hours and 225 hours, respectively. They match to the beginning of the performance degradation in the SCATU results quite well (IV measurement results section), which supports the hypothesis that electrolyte bleaching is the main reason for the degradation. The iodine concentration in both cell groups seems to saturate to the level that corresponds to the blue pixel value of the bare glass substrate in Fig. 9. Therefore, it is more probable that the saturation of the red and green curves in Fig. 9 illustrates a drop in the resolution power of the photographing technique when the electrolyte is nearly colorless than that it indicates the actual saturation of the bleaching.

A straightforward thought arising from Fig. 9 is whether the cell performance would remain stable for a longer time if more iodine was added to the electrolyte. Then the slope of the bleaching would still be the same but the starting point would be higher. This is also suggested by Mastroianni et al. The experimental confirmation of this question was not within the scope of this study, but a comparison was performed that was based on the aging data presented by Asghar et al. (the blue lines in Fig. 9). The most significant difference between the cells of Asghar et al. and the cells aged in this study is that the cells of Asghar et al. contained commercial Dyesol HSE-EL electrolyte. This is the same electrolyte that has been used in cells that have passed 1000-hour aging tests without a UV filter and using the exactly same lamps as in this study. The two blue curves in Fig. 9 are otherwise identical but have a different initial concentration of iodine because the commercial electrolyte used in article24 was estimated to contain 0.10-0.15 M iodine on the basis of the color of the electrolyte. The line showing a 0.1 M initial concentration crosses the 0.03 M limit just after 155 UV1Sunh and the line showing a 0.15 M initial concentration does not reach the 0.03 M level at all in 200 UV1Sunh. This indicates that the cells would probably have passed a 1000-hour aging test conducted with the same light soaking device (0.2 UV1Sun) as that was used in this study. By comparing the slopes of the lines in Fig. 9 it is clear that the cells containing purified MPN are not less stable than the cells of Asghar et al. - actually, the slopes are remarkably similar before the 85 UV1Sunh point. Therefore it can be concluded that the cells in this study are as stable as expected but failed to pass the aging test because they did not contain that much iodine.

Asghar et al. aged the cells under lamps that emitted 80% of the UV light in AM1.5G irradiation but the same amount of visible light as in this study. Because the bleaching rate in Fig. 9 is plotted as a function of the cumulative UV dose and the bleaching rates of the cells containing purified MPN and the cells prepared by Asghar et al. are the same, it is likely that the electrolyte bleaching was caused mainly by UV rather than visible light in this study (see investigation of electrolyte bleaching section).

Additionally, Fig. 9 illustrates the fact that with the bleaching rates of this study and article24 the cells would only survive under solar irradiation for a few dozen days without a UV filter. Based on the bleaching rate of the cells containing purified electrolyte solvent calculated in this study, similar cells would probably survive a 1000 hour aging test with a UV filter that would pass roughly 4.5% of the UV irradiation. This drastic difference might explain why controversial mechanisms in the cells containing purified and unpurified electrolyte solvent, but the degradation rate is clearly faster in the latter cells. The electrolytes degraded massively during the aging under illumination through bleaching. The bleaching rate of the cells containing unpurified electrolyte solvent was almost double that of the cells containing purified solvent. The photoelectrodes of the cells degraded only marginally. Despite that, we interestingly found dye degradation products that have previously been reported only at elevated temperatures in cells aged under UV and visual irradiation below 40°C. Some degradation products seem to form only under illumination, because they were not detected from cells kept in the dark at room temperature.

On the evidence of this study, UV-induced iodine bleaching seems to be directly proportional to the cumulative dosage of UV. Bleaching is most probably a combined effect of impurities and exposure to UV irradiation (approximately 20% of the UV in AM1.5G irradiation) during the aging test. UV is a significant factor in cell stability testing and therefore, the amount of UV in the illumination of aging lamps should be reported in cell stability studies. While the UV-induced degradation can be alleviated by using a UV filter, the filters are not perfect and preventing UV-related degradation, e.g., as shown here by purifying the solvent, is important.

Acknowledgments

The authors thank Prof. Leif Kronberg (Åbo Akademi) for his expertise in LC-MS measurements and Dr. Eeva Vilkkumaa (Aalto University) for her assistance in the statistical analysis. This study was partially funded by Tekes - the Finnish Funding Agency for Technology and Innovation (project KesMPV, 2928/31/2010) and partially by the Multidisciplinary Institute of Digitalization and Energy (MIDE) of Aalto University (project FerroPV, 751026). S. R. thanks the Fortum Foundation for financial support and K. M. is grateful for a post-doctoral project (253643) funded by the Academy of Finland.