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Thermal stability improvement of metal oxide-based contacts for silicon heterojunction solar cells

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Abstract

Metal oxides are interesting materials for use as carrier-selective contacts for the fabrication of doping-free silicon solar cells. In particular, MoO_x and TiO_x have been successfully used as hole and electron selective contacts in silicon solar cells, respectively. However, it is of paramount importance that good thermal stability is achieved in such contacts. In our work, we combined i-a-Si:H/ MoO_x based hole contacts with electron contacts featuring i-a-Si:H/ TiO_x /low work function metal (ATOM) to fabricate doping-free cells, termed MolyATOM cells. We found that the thermal stability of the ATOM contact was improved when the i-a-Si:H was annealed (300 °C for 20 min in N_2) before depositing TiO_x (i.e. pre- TiO_x annealing), which reduces the hydrogen content in i-a-Si:H by about 27 %_{rel}, and thereby the H-related degradation of the ATOM contact characteristics. Moreover, it was found that reducing the thickness of the low-work function metal on top of the TiO_x enhanced the thermal stability of the ATOM contact. With these adaptations, the MolyATOM cell efficiency was improved by 3.5 %_{abs}, with the highest efficiency of 17.6 %. Moreover, the cells show improved thermal stability after the above-mentioned pre- TiO_x annealing, which is confirmed by annealing tests at cell level as well as damp-heat tests at module level. The insights of this study could be used to tailor other metal-oxide based electron or hole contacts.

1. Introduction

Photovoltaics (PV) is gaining more importance worldwide, and has become the largest newly installed renewable energy source with newly installed capacity in 2018 of more than 100 GWp [1]. Today crystalline silicon (c-Si)-based PV modules occupy a market share above 95% of the annually installed PV capacity [2]. Research in the field of c-Si solar cells is focused on getting higher efficiencies, combined with lower PV module prices [3]. Higher efficiencies can be achieved by minimising the total recombination current densities ($J_{0,\text{total}}$) in the solar cell, which is the sum of recombination current densities at the passivated surfaces ($J_{0,\text{passi}}$), at the metallized areas of the surfaces ($J_{0,\text{metal}}$) as well as in the bulk ($J_{0,\text{bulk}}$). Currently, homojunction solar cells, such as aluminum back-surface-field (Al-BSF), passivated emitter and rear cells (PERC), and passivated emitter and rear totally diffused (PERT) solar cells, are dominant in the market [4]. In these solar cells, metal-semiconductor (MS) contacts are made on the heavily doped Si regions. This results in high recombination current densities at the metalized areas ($J_{0,\text{metal}}$), which range from about a few hundred to a few thousand fA/cm^2 depending on the cell technology [5–8]. The high $J_{0,\text{total}}$ caused by a full-area rear MS contact in Al-BSF solar cells is addressed by implementing local contact structures in the PERC [9] and PERT [10] cells. This reduces the metallized area fraction and hence the contribution of $J_{0,\text{metal}}$ to $J_{0,\text{total}}$, thereby enabling improved efficiencies compared to Al-BSF cells. Since $J_{0,\text{passi}}$ below 20 fA/cm^2 can be readily achieved for both phosphorus-doped and boron-doped surfaces [11], $J_{0,\text{metal}}$ is the main factor limiting the achievable cell efficiency in these homojunction solar cells. Lowering the contribution of $J_{0,\text{metal}}$ further by reducing the metal contact area even more would lead to increased contact resistance (R_C) i.e. there is a trade-off between $J_{0,\text{total}}$ and R_C .

The current approach in the field of silicon photovoltaics to achieve both low $J_{0,\text{metal}}$ and low contact resistivity (ρ_c) is through the implementation of a passivating contact, whereby the passivating layer is part of the contact scheme. Representative examples of such passivating contacts are silicon heterojunctions (SHJ) based on hydrogenated amorphous or nano-crystalline silicon (a-Si:H or nc-Si:H) with a transparent conductive oxide (TCO) [12,13] as well as contacts based on poly-Si on an ultra-thin oxide (i.e. TOPCon or POLO) [14,15]. Both SHJ solar cells and the cells with poly-Si/oxide passivating contacts have demonstrated efficiencies above 25%, due to the excellent contact passivation and the resulting low $J_{0,\text{metal}}$, combined with the low ρ_c achieved in these solar cells [12,14,16]. Therefore, these contact schemes are intensively studied for industrial implementation [17].

These passivating contacts generally require the deposition of a doped layer (e.g. plasma-enhanced chemical vapor deposition (PECVD) of doped a-Si or low-pressure CVD (LPCVD) of *in situ* doped poly-Si) or a doping process (e.g. diffusion or implantation of dopants into intrinsic poly-Si). A novel approach for realizing passivating contacts that is of interest in the PV research community is the fabrication of cells without such doping processes or deposition of doped layers. By applying certain metal compounds, particularly transition metal oxides (TMOs), in place of such doped layers, carrier selectivity can be achieved at the contacts by virtue of their band structure, and their low or high work function with respect to c-Si. One of the benefits of this approach is that the manufacturing process could be simplified due to the absence of doping processes and critical pre- and post-diffusion cleaning steps. Moreover, optoelectrical losses occurring in the heavily doped regions can be avoided by replacing the doped layers with the aforementioned high bandgap metal compounds.

For doping-free hole contacts, several high work function materials were proposed, such as MoO_x , VO_x , WO_x , NiO_x , and CuI [18–21], however, only MoO_x with an intrinsic a-Si:H (i-a-Si:H) buffer layer on c-Si could achieve high solar cell efficiencies above 20 % [22–25]. Without the i-a-Si:H buffer layer, MoO_x can induce stronger upward band bending at the hole contact region of c-Si, which is beneficial for charge transport [26], however it also results in poor passivation quality which limits the cell efficiency [27]. Therefore, many studies were aimed at replacing only the p-type a-Si:H (p-a-Si:H) in the i/p-a-Si:H stack of SHJ cells with MoO_x . The high bandgap of MoO_x reduces the parasitic absorption at short wavelength region and its high work function induces upward band bending for efficient hole collection, in a similar way to p-a-Si:H in SHJ cells [24]. In this work, we too study MoO_x on i-a-Si:H as the hole contact.

For doping-free electron contacts, relatively many material options are available. Metal oxides (e.g. TiO_x) [28–31], metal nitrides (e.g. TaN) [32,33], metal fluorides (e.g. LiF) [34–36] or a combination of those layers with low work function metals (e.g. Ca or Yb) [37–44] have been studied. In theory, when a metal is directly in contact with c-Si or a-Si:H, a large number of surface states are created at the interface. As a result, the Fermi level of the metal ($E_{F,\text{metal}}$) is pinned at a certain energy level of semiconductor regardless of work function of metals [45,46]. This can lead to weak or undesired band bending of semiconductor, and result in high ρ_c . To

depin the $E_{F,\text{metal}}$ on c-Si or a-Si:H, a thin metal oxide can be introduced between the metal and the semiconductor (i.e. c-Si or a-Si:H) [47,48]. Previously, we have reported on the development of a unique electron contact stack featuring i-a-Si:H/TiO_x/low work function metal, in short, ATOM contact [39–41,48,49]. The i-a-Si:H layer ensures excellent contact passivation with a low $J_{0,\text{metal}}$ of less than 25 fA/cm², while the TiO_x layer is used to eliminate the Fermi level (E_F) pinning of the low work function metals (e.g. Ca or Yb) on the i-a-Si:H surface, which enables a low ρ_c of below 0.1 $\Omega\cdot\text{cm}^2$ to be attained [38–41]. In this work, we study the ATOM contact with Yb (Φ_{metal} 2.5 eV) as the low work function metal.

Thermal instability of doping-free electron and hole selective contacts based on TMOs has been widely reported in literature. For instance, it has been reported that annealing treatments on the TiO_x/Ti contact on c-Si causes oxygen migration from TiO_x to the Ti metal, leading to the conversion of TiO_x to semimetallic TiO. This leads to high ρ_c due to the reappearance of $E_{F,\text{metal}}$ pinning [50]. We have reported in our earlier work on the increase in ρ_c of the ATOM contact stack based on TiO_x/Yb/Ag on a-Si:H when a thermal treatment was applied [41].

Similarly, there have been many studies reporting on the degradation observed in hole selective contacts featuring i-a-Si:H/MoO_x. The degradation mechanism is apparently related to oxygen loss in the MoO_x layer due to reaction with hydrogen from the i-a-Si:H layer [27,51]. Changes in the stoichiometry of MoO_x directly affect the work function of MoO_x [52], and thus, the band bending at the c-Si surface and hole collection. One suggested a solution to this problem is to pre-anneal the i-a-Si:H layer at 250 °C in N₂ for 20 min before MoO_x deposition. With this approach, reduced degradation of MoO_x and increased cell efficiencies have been reported [27].

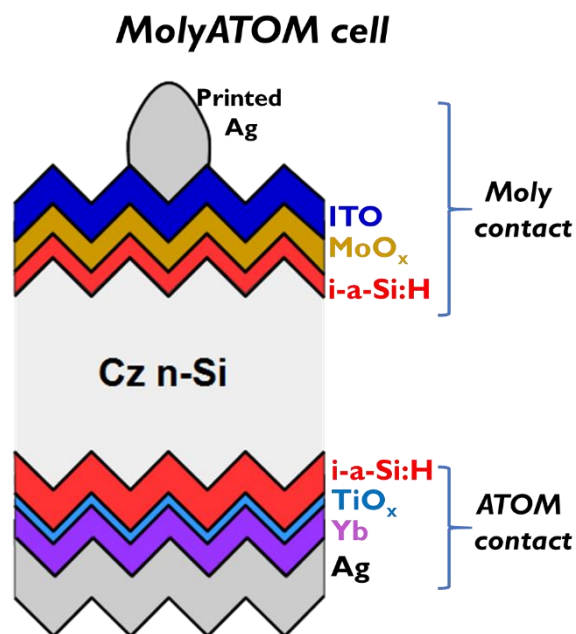


Figure 1. A schematic cross-section of the fully doping-free MolyATOM cell.

Building on the understanding gathered from the i-a-Si:H/MoO_x based hole contact, the first goal of this paper is to gain further insight into the role of hydrogen in the degradation observed in the ATOM electron contact based on i-a-Si:H/TiO_x/Yb/Ag. The impact of pre-annealing the i-a-Si:H layer on the thermal stability of the ATOM contact is investigated in terms of ρ_c . Additionally, it was found that the thickness of the low work function metal affected the thermal stability of the ATOM contact via an increase in ρ_c .

The second goal of this paper is to demonstrate, after determining the optimal pre-annealing conditions to ensure high thermal stability, fully doping-free cells, shortly called MolyATOM cells (Molybdenum oxide-based hole contact + ATOM electron contact, shown in Figure 1), whereby the doped layers on both the electron and hole contacts were replaced by TMOs. The thermal stability of the MolyATOM cells was evaluated at cell and module level.

2. Experimental methods

2.1. Effect of annealing i-a-Si:H on contact resistivity of ATOM contacts

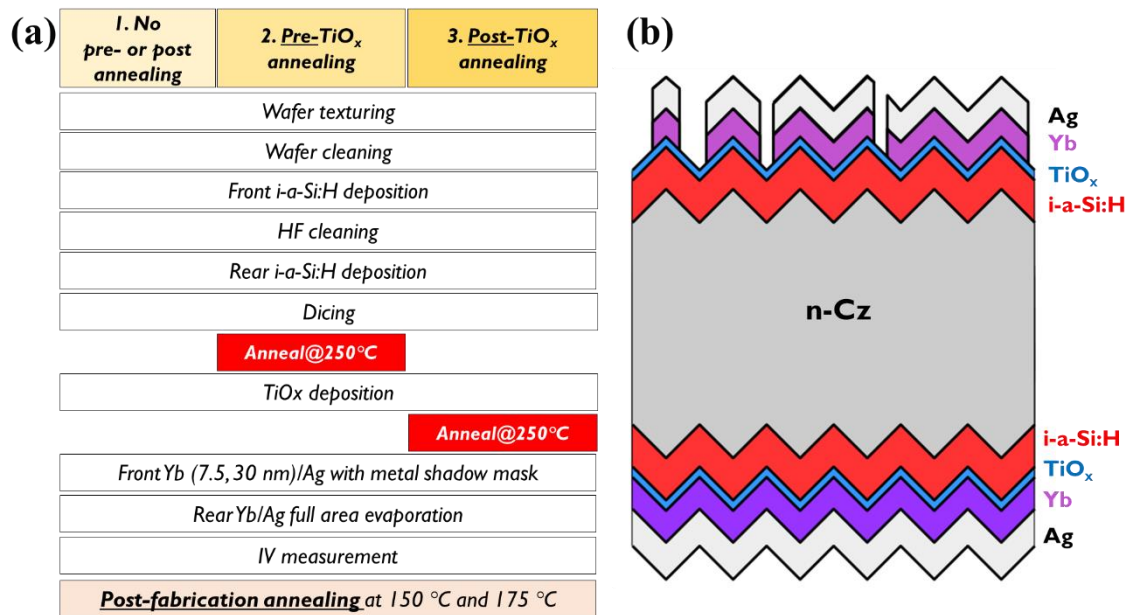


Figure 2. (a) Process flow for studying pre- and post-TiO_x annealing effect on ρ_c , and (b) a schematic sample structure for this study.

For evaluating the impact of annealing the i-a-Si layer before and after TiO_x deposition on the ρ_c of ATOM contacts, test samples were prepared using double-side textured (3 Ω ·cm, 180 μ m thickness after texturing process) n-type Cz-Si wafers. The process flow used to fabricate these test samples and their cross-sectional structure are depicted in Figure 2. After wafer cleaning using O₃/HCl/deionized water and HF/HCl solutions, 8 nm thick i-a-Si:H was deposited by PECVD on the front side. After HF cleaning, an i-a-Si:H layer of 8 nm was deposited on the rear side of all samples. TiO_x of 1 nm was deposited by atomic layer deposition (ALD) using Ti(OCH₃)₄ and H₂O as Ti precursor and oxidant, respectively. Two groups of the sample were

annealed at 250 °C for 20 min in N₂ using rapid thermal annealing (RTA) before and after TiO_x deposition, respectively, and one group was not annealed at all. Then, the wafers were diced to the size of 3 cm x 3 cm for ρ_c measurement. Two thicknesses of Yb (7.5 nm and 30 nm) with a fixed Ag thickness of 100 nm were evaporated without breaking vacuum on the front side of different samples through a metal shadow mask to form circular dots of different diameters. For the rear side, full area Yb of 30 nm and Ag of 100 nm were deposited using the same technique. At the end of the fabrication sequence, additional annealing treatments were done at 150°C and 175°C. The method for determining ρ_c using the test structure shown in Figure 2 (b) is explained in the elsewhere [38,41].

2.2. Solar cell fabrication

MolyATOM	Moly/in	Classical SHJ
Wafer texturing		
Wafer cleaning		
Front i-a-Si:H deposition		Front i/p-a-Si:H
HF cleaning		
Rear i-a-Si:H deposition	Rear i/n-a-Si:H deposition	
Pre-ann. @ 200, 250, 300 °C	Pre-ann. @ 300 °C	
TiO _x deposition		
Front vapor HF	HF cleaning	
MoO _x evaporation		
Front ITO	Front and rear ITO	
Front Ag screen print	Front and rear Ag screen print	
Sintering @ 160 °C		
Rear Yb (7.5 nm)/Ag evaporation		
Post-fabrication ann. @ 150 °C		

Figure 3. Process flow for the fabrication of different solar cells reported in this paper: MolyATOM, Moly/in and classical SHJ cells.

Three types of solar cells were prepared, namely 1) MolyATOM cells (with i-a-Si:H/MoO_x as hole contact and i-a-Si:H/TiO_x/Yb/Ag as ATOM electron contact), 2) Moly/in cells (with i-a-Si:H/MoO_x as hole contact and i/n-a-Si:H stack as electron contact), and 3) classical SHJ cells (with i/p-a-Si:H and i/n-a-Si:H stacks as hole and electron contacts, respectively). The respective process flows for the fabrication of these solar cells is given in Figure 3. Double-side textured (3 Ω·cm, 180 μm thickness after texturing process) n-type Cz-Si wafers were used.

After cleaning the textured wafers, an i-a-Si:H layer was deposited on the front for MolyATOM and Moly/in cells while an i/p-a-Si:H stack was deposited on the same side for classical SHJ cells. After HF cleaning, i-a-Si:H was deposited on the rear for the MolyATOM cells while an i/n-a-Si:H stack was deposited for the Moly/in and classical SHJ cells.

Afterward, for the 1st study about pre-annealing conditions, 3 groups of wafers from the MolyATOM split were pre-annealed using rapid thermal annealing (RTA) at 200 °C, 250 °C and 300 °C, respectively, for 20 min in N₂ atmosphere, while a fourth group was not annealed at all. For the 2nd study about the comparison of MolyATOM with classical SHJ cells, one group of MolyATOM cells was pre-annealed at 300 °C. One group of wafers from the Moly/in split was also pre-annealed at 300 °C under the same conditions, while the other Moly/in wafers were not. For the classical SHJ wafers, no pre-annealing was carried out.

On the rear-side of the MolyATOM wafers, a 1 nm thick TiO_x layer was deposited using ALD. Then, a vapor HF process or an HF cleaning was performed on the wafers of the MolyATOM and the Moly/in splits, respectively. Vapor HF allows one-sided native oxide removal on the front-side of the MolyATOM wafers without affecting the TiO_x on the rear-side. After a MoO_x evaporation of 6 nm on the front-side for the MolyATOM and Moly/in splits, ITO deposition and Ag screen-printing were done for all cell types on the front-side. For the rear-side, ITO sputtering and Ag screen-printing were done only for the Moly/in and classical SHJ splits. Sintering was then carried out on all cells including the MolyATOM devices using a belt furnace in N₂ at 160 °C to increase the conductivity of the printed Ag grid on the front-side and to recover the sputtering damage from ITO deposition. For the MolyATOM split, an Yb/Ag stack was thermally evaporated on the rear-side without breaking vacuum.

Illuminated IV characteristics were measured with an aperture opening of the same size as the cell area of 4 x 4 cm² at standard 1 sun conditions (AM 1.5G, 1000 W/m² at 25 °C). External quantum efficiency (EQE) and reflectance (R) were measured using an illumination spot size of 1.5 x 1.5 cm². Therefore, the EQE and reflectance results include the shading and reflection effects of the front metal grid. A post-fabrication anneal was performed on part of the cells at 150 °C for varied durations.

2.3 Evaluation of hydrogen content in i-a-Si:H

The evolution of the hydrogen content in i-a-Si:H after annealing at 200 °C, 250 °C and 300 °C was investigated using elastic recoil detection (ERD) on a 30 nm thick a-Si:H layer deposited using PECVD on mirror polished 730 μm thick Cz wafers.

2.4. Mini-module fabrication

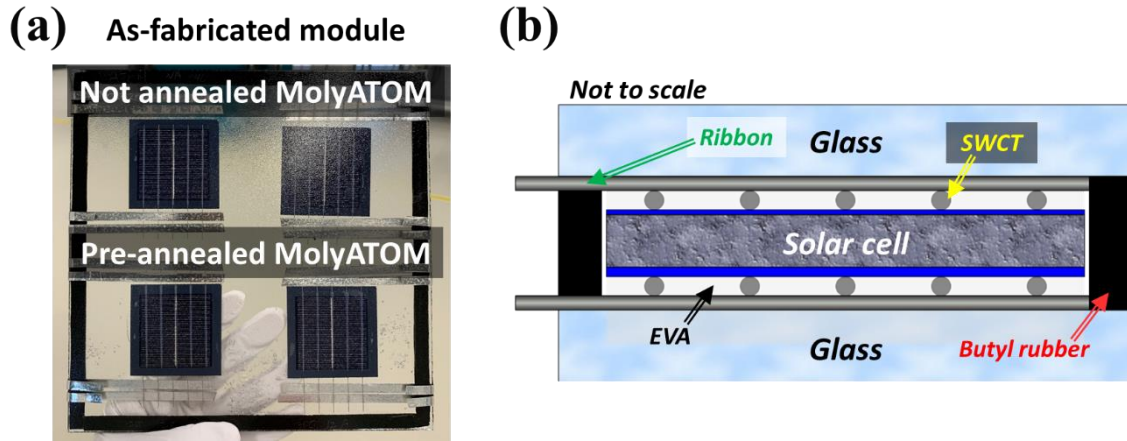


Figure 4. (a) As-fabricated state the mini-module and (b) a schematic cross-sectional image of the mini-module.

A mini-module (glass/glass) was fabricated using the MolyATOM cells with an Yb thickness of 7.5 nm with and without pre-anneal treatment at 300 °C. The four cells in the mini-module shown in Figure 4 (a) are not interconnected. An UV-blocking ethylene vinyl acetate (EVA) sheet, a smart-wire connection technology (SWCT) foil (from Meyer Burger) and glass were used. A solder-coated ribbon was used to connect the smart wires and bring the electrical contacts of each individual cell to outside of the mini-module, so that each one can be measured individually. A butyl rubber tape was applied for hermetic sealing at the edges. The lamination process was carried out at 150°C for about 20 min. During the course of the damp heat test (85 °C/85 % humidity/1000 hour), illuminated IV curves (AM 1.5, 1000 W/m² at 25 °C) were measured after every 250 hours. An aperture area of 4×4 cm² was used for these measurements by using an opaque black tape around the edge of the active cell area. The same module structure and test method were reported in reference [20].

3. Improving the thermal stability of the ATOM contact

As noted earlier, deterioration in ρ_c of the ATOM contact, based on TiO_x/Yb/Ag on a-Si:H, by about an order of magnitude upon annealing at 150 °C or 180°C has been reported previously [41]. Thus, finding a way to improve the thermal stability of the ATOM contact is necessary. It is postulated that the mechanism behind this thermal degradation is related to the reduction of TiO_x, which could occur due to the reaction between H in a-Si:H and O in TiO_x, or due to O migration towards the low work function metal on the other side. In addition, TiO_x may undergo microstructural changes upon annealing.

For these reasons, annealing treatments before (labeled pre-TiO_x anneal) and after TiO_x deposition (post-TiO_x anneal) were tested as possible means to improve the thermal stability. The idea is that a pre-TiO_x anneal could dehydrogenate the i-a-Si:H layer and therefore mitigate the reaction between H in i-a-Si:H and O in TiO_x. A post-TiO_x anneal, on the other hand, could stabilize the microstructure of TiO_x before deposition of the low work function metal, such that

O migration to the interface with the low work function metal or the a-Si:H can be inhibited, thereby minimizing thermal degradation.

As a method for evaluating thermal stability, the change in ρ_c of the ATOM contact was evaluated after post-fabrication annealing consecutively at 150 °C and 175 °C for 10 min each. In addition, different Yb thicknesses of 7.5 nm and 30 nm were tested to investigate the impact on thermal stability of the ATOM contact in terms of ρ_c . Since loss of hydrogen from a-Si:H could affect the passivation quality, the evolution of the implied V_{OC} (iV_{OC}) and the $J_{0,total}$ of cell precursors was also monitored as a function of the pre-annealing temperature.

3.1. Hydrogen content of i-a-Si:H after annealing at different temperatures

Pre-annealing of i-a-Si:H at 250 °C for 20 min during the fabrication of solar cells with i-a-Si:H/MoO_x based hole contacts was reported to result in higher thermal stability and cell efficiency [27]. While we may expect a similar effect for our ATOM electron contact, the optimal conditions for this pre-annealing treatment would depend on the properties of the as-grown a-Si:H layer and the details of its interaction with TiO_x. Thus, in our work, various pre-annealing temperatures were tested.

Annealing of i-a-Si:H, prior to capping it with metal layers, leads to effusion of hydrogen and concomitant reduction in its hydrogen content. ERD measurements can be used to determine the hydrogen content in a-Si:H films. The concentration of hydrogen in a 30 nm thick i-a-Si:H film was determined in the as-deposited state and then after annealing at different temperatures of 200 °C, 250 °C and 300 °C for 20 min, and the results are summarised in Table 1.

Table 1. The hydrogen concentration in a-Si:H of 30 nm thickness after annealing at different temperatures for 20 min, determined using ERD measurements.

Annealing temperature	No annealing	200 °C	250 °C	300 °C
H (at %)	15.7 ± 2	14.3 ± 1	14.0 ± 1	11.4 ± 1
Si (at %)	84.3 ± 2	85.7 ± 2	86.0 ± 2	88.6 ± 2
(H / H_{non-annealed}) -1 (%rel)	-	- 9	- 11	- 27

The as-deposited film has a high H concentration of approximately 15.7 %, and annealing the film results in a loss of hydrogen at all investigated temperatures. This proves that indeed H effuses out of the a-Si:H film upon annealing at these temperatures. The higher the temperature, the greater the loss of H from the film. While annealing at 200 °C and 250 °C results in a loss of H of around 10%_{rel}, a sharp reduction in H of 27%_{rel} is observed when annealing at 300 °C.

While a 30 nm thick i-a-Si:H film was used for this study, the phenomenon is expected to be similar for an 8 nm thick film.

3.2. Effect of pre- and post-TiO_x annealing on thermal stability of the ATOM contact as monitored via ρ_c

To study the impact of annealing the i-a-Si:H layer on the ρ_c of the ATOM contact, a fixed pre-annealing temperature of 250 °C for 20 min was used (see Figure 2 (a)). The ρ_c results of the ATOM contact are summarised in Figure 5. To minimize series resistance losses, ρ_c should be as low as possible. However, an upper limit for ρ_c of 0.1 $\Omega\cdot\text{cm}^2$ was set, at which significant V_{OC} losses are expected due to non-negligible potential drop at the contact [53,54].

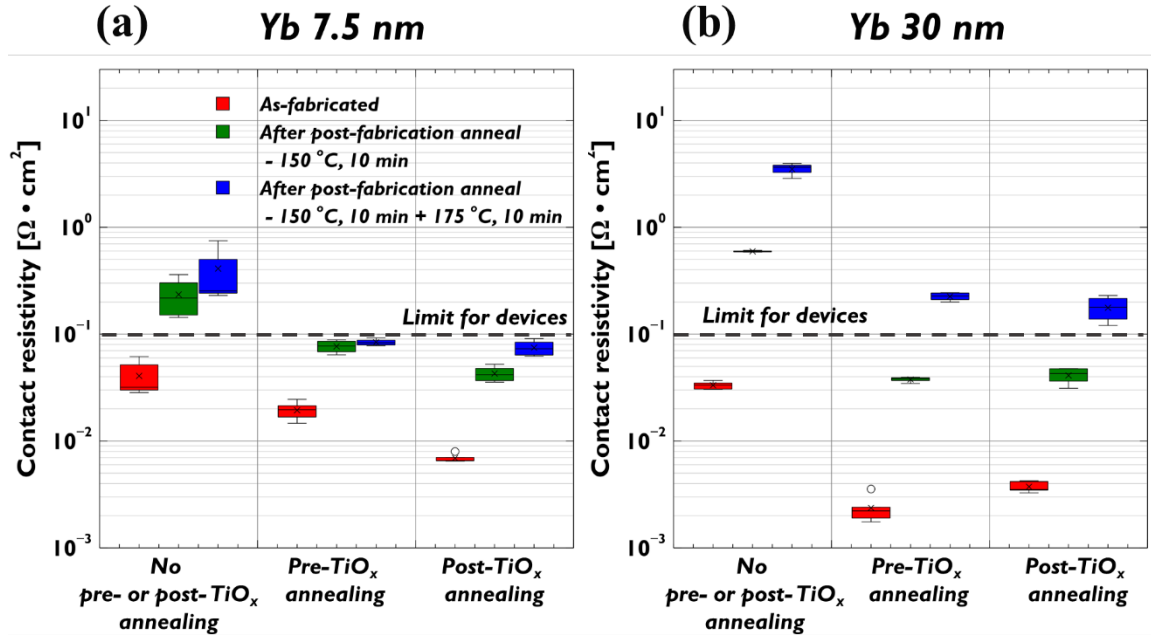


Figure 5. Evolution of contact resistivity (ρ_c) after sequential post-fabrication anneals at 150 °C and 175 °C for 10 min each, for the three anneal groups shown in Figure 2 (i.e. no pre- or post-TiO_x annealing, pre- and post-TiO_x annealing). Pre- and post-TiO_x anneals were done at 250 °C. Results for ATOM contact with Yb thickness of (a) 7.5 nm and (b) 30 nm, respectively, are plotted. Each bar contains five data points. The targeted ρ_c for device implementation is below 0.1 $\Omega\cdot\text{cm}^2$ (indicated by the dashed line).

The first observed trend is that the ρ_c of the ATOM contact increases after post-fabrication annealing treatments at 150 °C and 175 °C for all investigated conditions. However, the extent of the degradation is much lower for the pre- and post-TiO_x anneal splits compared to the non-annealed case without neither pre- or post-TiO_x annealing. Analogous to studies on degradation of the ρ_c of i-a-Si:H/MoO_x hole contacts [27], it appears that annealing the i-a-Si:H layer before capping it with thick metal layers is indeed also beneficial for the ATOM contact in ensuring higher thermal stability. As discussed in Section 3.1, annealing reduces the H content of the i-a-Si:H layer, which mitigates the reaction between the H in i-a-Si:H and O in TiO_x. In this

respect, to maximize O concentration in TiO_x by avoiding thermal treatment of TiO_x as much as possible, the pre- TiO_x annealing route would be more suitable than the post- TiO_x annealing.

Comparing Figure 5 (a) with Figure 5 (b), a second trend that can be noticed is that for non-annealed TiO_x , thicker Yb causes a stronger degradation in ρ_c after post-fabrication annealing. In other words, thicker Yb seems to reduce TiO_x more aggressively. According to Fu et al., oxygen transport to the interface from TiO_2 to metal was dependent on the overlying metal work function [55]. They explained that the lower work function of a metal with respect to metal oxides promoted out-diffusion of oxygen from the metal oxides. A similar phenomenon was also found by Allen et al. whereby TiO_x was reduced by Ca (Φ 2.9 eV) and a TiO phase was detected by electron energy-loss spectroscopy (EELS) [37]. Since Yb has a work function of 2.5 eV, such oxygen migration at the interface between TiO_x and Yb can be expected to be greater compared to the interface between TiO_x and Ca or Al [37,55]. However, this phenomenon might be less severe for the case of thin Yb, whose work function could be increased by the alloying of Ag and Yb [39,41]. Thus, it is clear that using a thin Yb layer is beneficial in minimizing the thermal degradation of the ATOM contact.

The third trend that can be seen from Figure 5 is that both pre- and post- TiO_x annealing treatments lower the ρ_c compared to that of non-annealed samples before post-fabrication anneal treatments. We believe this is mainly related to structural and/or compositional changes to the i-a-Si:H layer. Dehydrogenation would lead to increased defects in a-Si:H, which could in turn increase trap-assisted tunneling through the i-a-Si:H.

Overall, for the case of 7.5 nm thick Yb, the ρ_c values of the pre- and post- TiO_x annealed samples were below the targetted threshold of $0.1 \Omega\cdot\text{cm}^2$ after successive post-fabrication annealing treatment at 150 °C and 175 °C while the non-annealed ATOM contact was already above the limit after the post-fabrication annealing even at 150 °C. In conclusion, pre- TiO_x annealing in combination with thin Yb should improve the thermal stability of ATOM contacts in terms of ρ_c .

3.3. Impact of pre- TiO_x annealing on iV_{OC} and $J_{0,total}$ of MolyATOM cell pre-cursors

Hydrogen in a-Si:H plays an important role in passivating the c-Si/a-Si interface. Thus, dehydrogenation of the i-a-Si:H layer during pre- TiO_x annealing can potentially affect the passivation quality of the i-a-Si:H layer. This was verified by measuring the effective lifetime of MolyATOM cell pre-cursors (1) after pre- TiO_x annealing, (2) after TiO_x deposition and (3) after MoO_x deposition, from which implied V_{OC} (iV_{OC}) and $J_{0,total}$ can be determined.

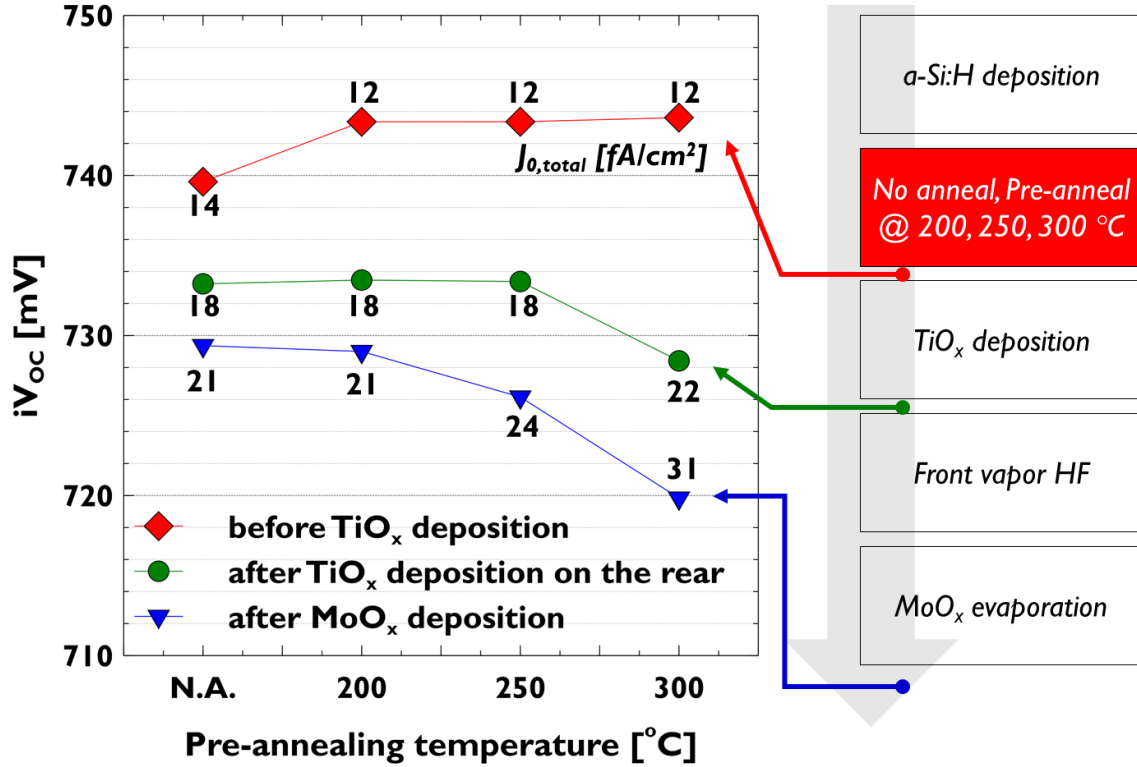


Figure 6. Progress in iV_{OC} of pre-annealed or not-annealed samples at different process steps. The numbers in the figure are the $J_{0,total}$ values of samples based on a J_{SC} of 36 mA/cm^2 .

The evolution of iV_{OC} and $J_{0,total}$ of the cell pre-cursors at different steps as a function of pre-TiO_x annealing temperature is shown in Figure 6. The passivation quality of annealed i-a-Si:H is similar to that of non-annealed i-a-Si:H, and high $iV_{OC} > 740 \text{ mV}$ and low $J_{0,total}$ of 12 fA/cm^2 are obtained for the cell pre-cursors with annealed i-a-Si:H. However, the deposition of TiO_x results in a decrease of iV_{OC} of all splits, with the sample pre-annealed at $300 \text{ }^\circ\text{C}$ showing the strongest reduction in iV_{OC} . Similarly, the subsequent deposition of MoO_x on the opposite side of the wafers leads to a further decrease in iV_{OC} of all splits. In addition, a decreasing trend in iV_{OC} with pre-TiO_x anneal temperature develops. Thus, we can conclude that pre-annealing treatment negatively impacts on the passivation quality after TMO depositions. Nevertheless, the iV_{OC} of the worst sample is still 720 mV , which is promising for device implementation. Thus, the developed pre-TiO_x annealing step was integrated into the device fabrication flow (see Figure 3).

4. MolyATOM cells and comparison to classical SHJ cells

4.1. Impact of pre-TiO_x annealing temperature on MolyATOM cell parameters

In this section, we studied the impact of the pre-TiO_x annealing temperature on the device parameters of the MolyATOM cell, particularly the fill factor (FF) and V_{OC} , which depend on the ρ_c and $J_{0,total}$, respectively.

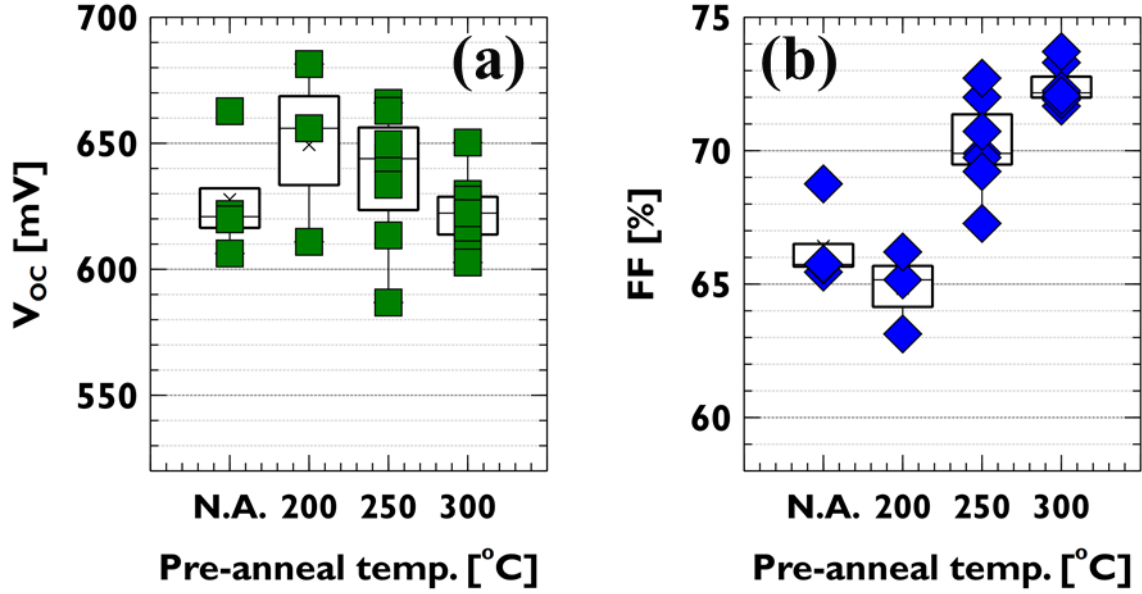


Figure 7. (a) V_{oc} and (b) FF of MolyATOM cells as a function of pre- TiO_x annealing temperature.

Table 2. Illuminated IV parameters of different MolyATOM cells (average and best cell values).

Pre-anneal temp. (# of sample)		J_{sc} [mA/cm ²]	V_{oc} [mV]	FF [%]	Efficiency [%]	R_s [$\Omega \cdot cm^2$]
No pre-anneal (4)	Average	33.7 ± 0.2	628 ± 24	66.4 ± 1.6	14.1 ± 1.0	3.3 ± 0.2
	Best	34.0	663	68.8	15.5	2.8
200 °C (3)	Average	33.8 ± 0.7	649 ± 36	64.8 ± 1.6	14.3 ± 1.4	3.7 ± 0.2
	Best	34.2	681	66.2	15.4	3.6
250 °C (7)	Average	33.5 ± 0.5	637 ± 28	70.2 ± 1.8	15.0 ± 1.2	2.5 ± 0.5
	Best	33.7	663	72.7	16.2	2
300 °C (7)	Average	33.2 ± 0.4	623 ± 15	72.4 ± 0.8	15.0 ± 0.7	1.9 ± 0.2
	Best	33.9	650	73.7	16.3	1.7

As shown in Figure 7 and Table 2, the V_{oc} of the MolyATOM cells showed a weak dependence on pre-annealing temperature, as expected from the results of Figure 6. The non-annealed cells show a somewhat lower average V_{oc} compared to the cells with pre- TiO_x anneal, due to a less efficient hydrogen chemical passivation. Despite the different hydrogenation levels of the i-a-Si:H films, the passivation quality achieved at the device level is quite similar for all investigated conditions of pre- TiO_x annealing. Nevertheless, the absolute V_{oc} values are much lower compared to the iV_{oc} values of Figure 6, indicating that the metallization process following MoO_x deposition (see Figure 3) affects the passivation quality even further. More detailed comparison will be discussed in section 4.3.

On the other hand, a general trend in FF of the MolyATOM cells demonstrates a strong dependence with pre-annealing temperature, which again correlates well with the contact resistivity results of Section 3. Less hydrogen content in i-a-Si:H after the pre-annealing

treatment has a positive effect for the carrier collection, likely due to increased defects in the i-a-Si:H layer, which aid trap-assisted conduction of charge carriers.

4.2. Dependence of thermal stability on pre-annealing temperature at cell level

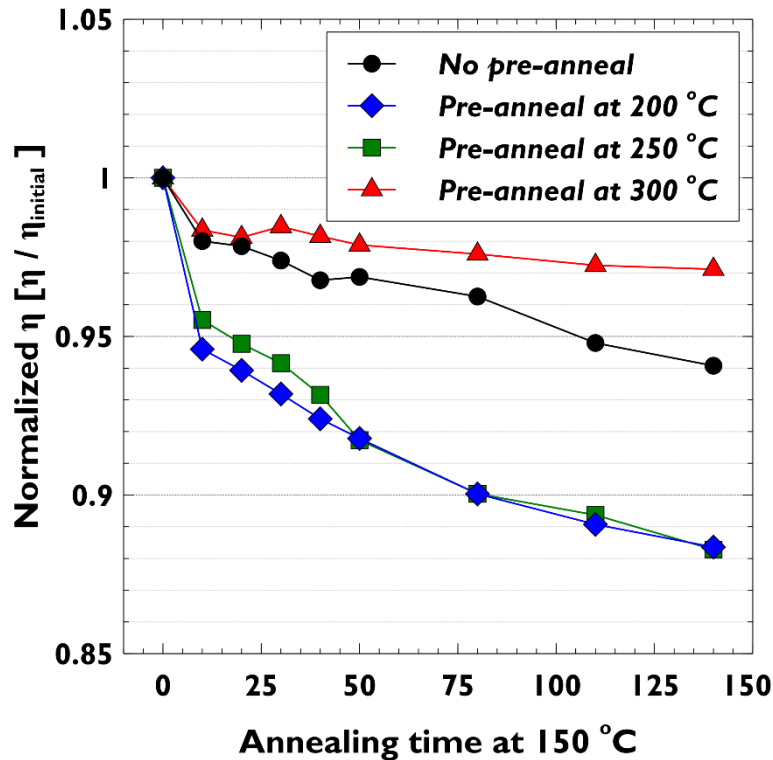


Figure 8. Efficiency progress as a function of annealing time at 150 °C in N_2 atmosphere.

Since the pre-annealing of the i-a-Si:H layer will affect both the $ATOM$ contact and the MoO_x based hole contact, the thermal stability was evaluated at device level using post-fabrication annealing at 150 °C for up to 140 min (see Figure 8). Cells with a pre- TiO_x annealing at 300 °C remained relatively stable, while the other cells showed a continuous decrease in efficiency with increasing annealing time. The not pre-annealed Moly $ATOM$ cell looks stable due to their low initial efficiency of 13.4 %. The initial efficiency of other cells was in a range of 14.6 % - 15 %. We extracted a reduction rate in the efficiency of the cells using the last four data points where degradation occurred linearly. The cell with pre- TiO_x annealing at 300 °C showed a reduction rate less than one-third of that of the cell without pre-annealing.

Based on the above results, the pre- TiO_x annealing temperature for Moly $ATOM$ device integration was chosen to be 300 °C for the study in section 4.3 in which we compare Moly $ATOM$ cells with other cell varieties.

4.3. Confirmation of improved thermal stability of Moly $ATOM$ cells at module level

The thermal stability of Moly $ATOM$ cells with and without pre-annealing at 300 °C was compared at cell and module level. To check the thermal stability at cell level, post-fabrication

annealing was done at 150 °C using the MolyATOM cells as shown in Figure 8. In addition, thermal stability of the MolyATOM cells at module level was investigated through the standard damp-heat testing (1000 hour, 85 % humidity, 85 °C). The MolyATOM cells with and without pre-annealing treatment at 300°C, shown in Figure 7, were used for fabricating a module. Performance of the mini-module was regularly measured every 250 hour during damp-heat test.

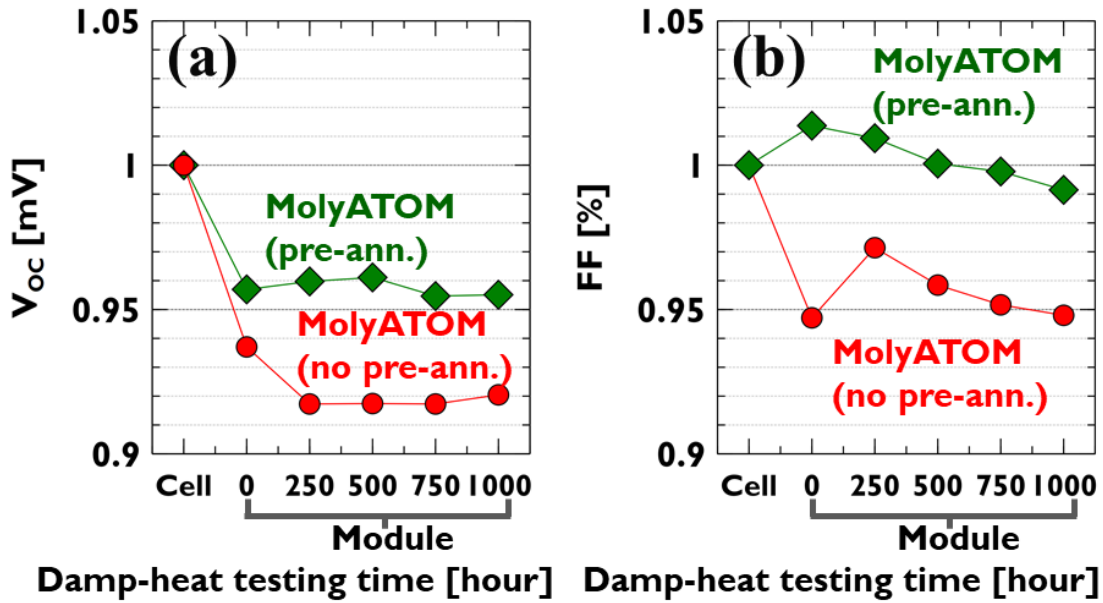


Figure 9. Thermal stability of MolyATOM cells at a module level. Progress in (a) V_{OC} and (b) FF during the damp-heat test.

During module fabrication, additional heat treatment due to the lamination process lowered the V_{OC} of both MolyATOM cells compared to the as-fabricated values (Figure 9 (a)). However, during the damp heat testing, the V_{OC} of the non-pre-annealed MolyATOM cell degraded further while that of the pre-annealed MolyATOM cell remained stable throughout the testing duration.

The FF of the pre-annealed MolyATOM cell during the damp heat testing (Figure 9 (b)) showed a similar evolution as that of the FF of the cell during the thermal stability test at cell level, which is a noticeable reduction after annealing at 150 °C for 10 min (Figure 8). Starting at DH 0 h, there is a gradual decrease in FF for the pre-annealed MolyATOM cell. The cell without pre-anneal though showed an anomalous behavior, with an increase in FF at DH 250 h, followed then by a gradual decrease in FF till DH 1000 h. Based on these results, we did not observe a significant difference in the thermal stability in terms of FF of the MolyATOM cells due to pre-annealing (DH 0 h vs DH 1000 h). Nevertheless, considering all the thermal stability results together, we can conclude that pre-annealing the i-a-Si:H during cell processing is beneficial for the thermal stability of MolyATOM cells. More generally, reducing the hydrogen content in the thin film contact stacks is helpful when using metals oxides. After the damp-heat test, the final efficiency was decreased by about 2.4 %_{rel} compared to that measured at DH 0 h. It meant

that the stability of MolyATOM cell at the module was passing the criteria of IEC61215 ($\eta < 5 \%_{\text{rel}}$).

4.3. Relative performance of MolyATOM cells compared to other cell types

To study relative performance, several types of solar cells were newly prepared. In Figure 10 and Table 3, MolyATOM cell parameters are compared with classical SHJ- and Moly/in cells. For the Moly/in cells, 2 splits of cells with and without pre-annealing treatment at 300 °C are also shown. For MolyATOM cells, only pre-annealed cells were prepared. In addition, the published results of cells having i/p-a-Si:H based hole contacts and ATOM electron contacts (i.e. ip/ATOM cells) have been included in Figure 10 [41]. Note that a different Ti precursor for TiO_x and thicker Yb of 40 nm were used for these ip/ATOM cells [41]. The other ATOM contacts had 7.5 nm thick Yb.

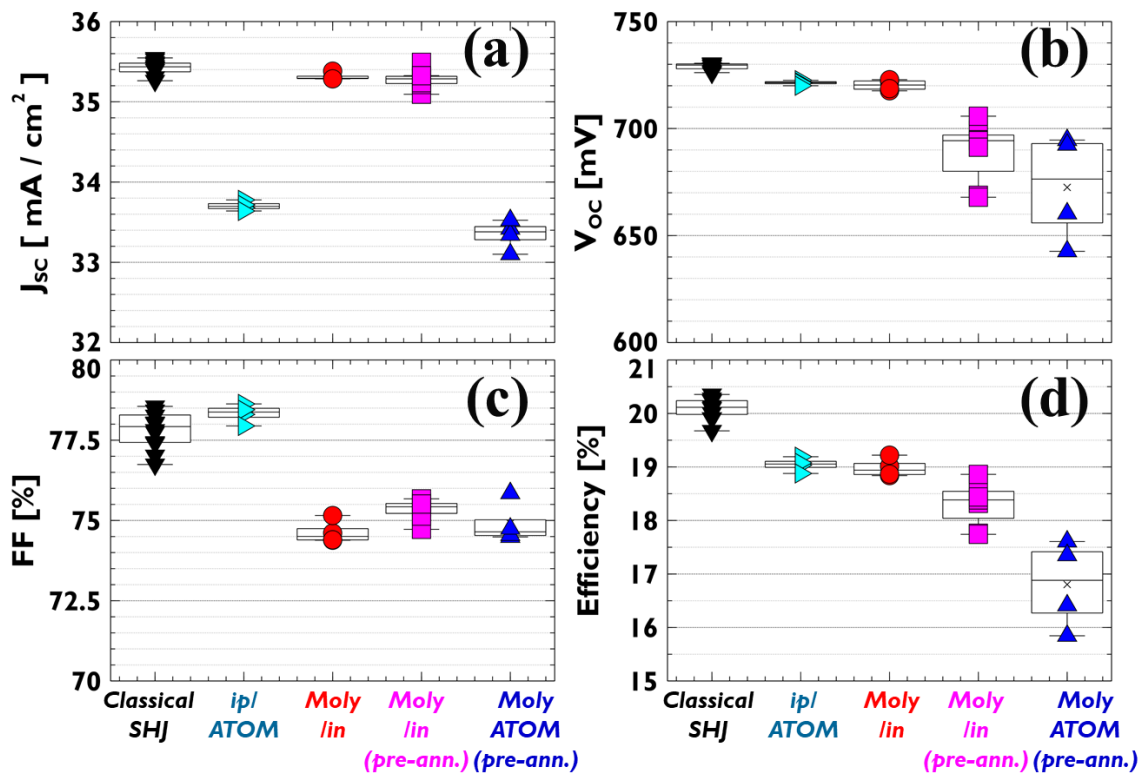


Figure 10. Performance comparison between classical SHJ, partial-doping free (Moly/in) and fully doping-free cells (MolyATOM). (a) J_{sc} , (b) V_{oc} , (c) FF and (d) efficiency. ip/ATOM data is from [41].

Table 3. The average and the best cell results of each solar cell group shown in Figure 10. Data for ip/ATOM is taken from [41].

Solar cell structure (# of sample)		J_{SC} [mA/cm ²]	V_{OC} [mV]	FF [%]	Efficiency [%]	R_s [$\Omega \cdot \text{cm}^2$]
Classical SHJ (12)	Average	35.5 ± 0.1	729 ± 2	77.7 ± 0.6	20.1 ± 0.2	1.4 ± 0.1
	Best	35.5	730	78.6	20.4	1.3
ip/ATOM (4) [41]	Average	33.7 ± 0.1	721 ± 1	78.3 ± 0.3	19.0 ± 0.1	1.0 ± 0.0
	Best	33.8	723	78.6	19.2	1.0
Moly/in (4)	Average	35.3 ± 0.0	720 ± 3	74.6 ± 0.4	19.0 ± 0.2	1.9 ± 0.1
	Best	35.4	723	75.2	19.2	1.8
Moly/in (Pre-ann.) (7)	Average	35.3 ± 0.1	689 ± 15	75.3 ± 0.3	18.3 ± 0.4	1.4 ± 0.1
	Best	35.5	706	75.3	18.9	1.4
MolyATOM (pre-ann.) (4)	Average	33.3 ± 0.2	673 ± 25	74.9 ± 0.6	16.8 ± 0.8	1.4 ± 0.0
	Best	33.4	695	75.8	17.6	1.3

The J_{SC} of the cells is drastically affected by the electron contact structure on the rear-side of the cells. A conventional i/n-a-Si:H/ITO/Ag electron contact at the rear-side of classical SHJ and Moly/in cells resulted in higher J_{SC} compared to that of MolyATOM and ip/ATOM cells. Moreover, Moly/in cells do not exhibit a higher J_{SC} compared to the classical SHJ cells. To understand these differences, the EQE and reflectance of the different cell types were compared in Figure 11.

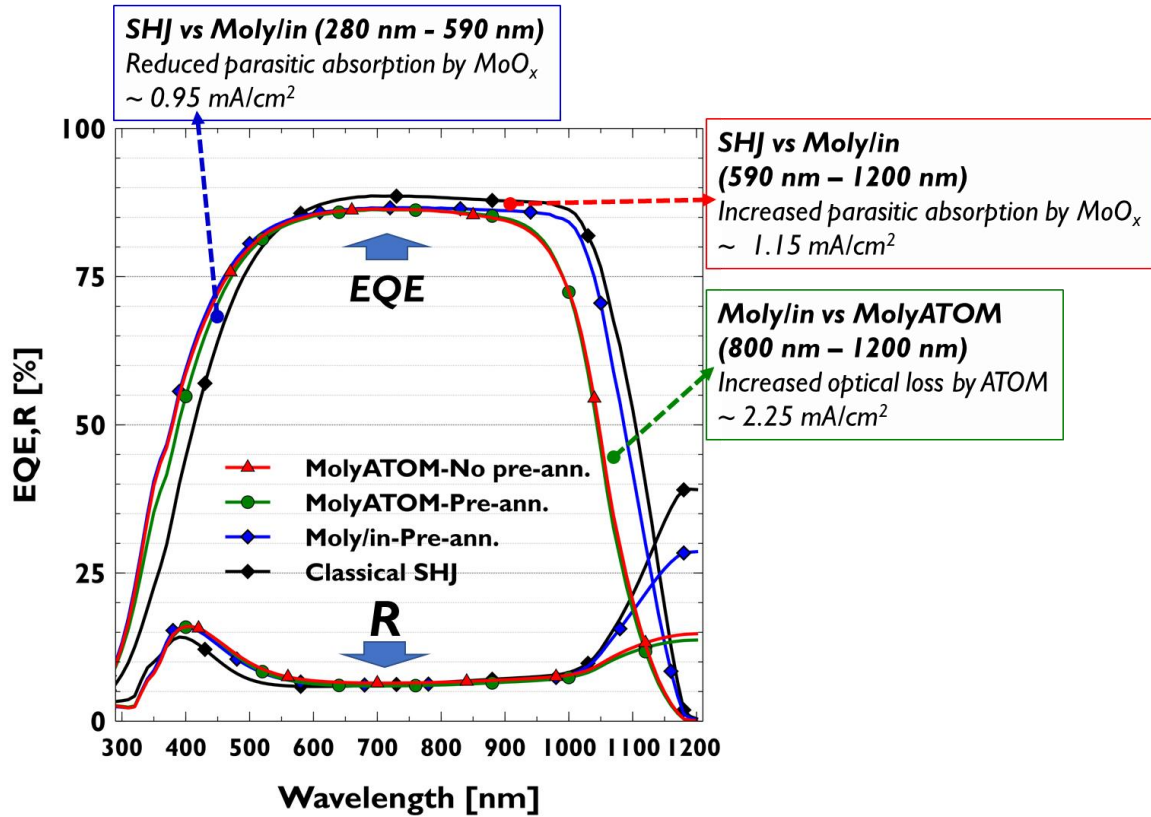


Figure 11. EQE and reflectance comparison between different cell structures; MolyATOM cell with/without pre-annealing treatment, Moly/in- and a classical SHJ cell.

Figure 11 shows that the cells with MoO_x based hole contacts exhibited higher EQE in the short wavelength region compared to the classical SHJ cells containing p-a-Si:H. However, this did not result in a higher J_{SC} for the Moly/in cells due to a lower EQE in the middle and long-wavelength regions compared to the classical SHJ cells, which is the result of parasitic absorption in MoO_x , as reported in [20]. A pre-annealing treatment does not change the EQE of the Moly/in cell (data is not shown here). Thus, transparent MoO_x needs to be developed for getting higher J_{SC} in Moly/in cells relative to classical SHJ cells. For the MolyATOM cells, lower EQE is observed at long wavelengths, which coincides with lower reflectance compared to the Moly/in or classical SHJ cells. This is expected to be mainly due to the proximity of the metal to the c-Si in the ATOM contact structure, which results in the excitation of surface plasmon polaritons, causing parasitic infrared light absorption losses. For the Moly/in and classical SHJ cells, on the other hand, ITO acts as a buffer layer.

As shown in Figure 10 (b), classical SHJ cells showed the highest V_{OC} among the test groups. When doping-free contacts were applied separately, V_{OC} of ip/ATOM and Moly/in were above 720 mV on average. For Moly/in cells, the pre-annealing condition used might not be the optimal due to the fact that pre-annealed cells showed lower V_{OC} . Different contact structure likely needs different thermal treatment condition. When combining MoO_x -based hole and ATOM electron contacts into the same device, an additional V_{OC} drop was observed. It is

believed that this reduction in V_{OC} might be caused due to integration difficulties when combining the two different doping-free contacts. For example, to deposit TiO_x , 300 mm diameter carrier wafers with a 200 mm round inner pocket, had to be used. During this loading of wafers on such carriers, the i-a-Si:H passivation layer on the front side of MolyATOM wafers touch the surface of carrier wafers, which may lead to contamination or scratches. A second possibility could be that the vapor HF, utilized in the case of MolyATOM cells, might not be sufficient to clean the surface of the wafers.

In Figure 10 (c), the average FF of ip/ATOM cells of about 78.3 % is higher compared to that of the classical SHJ cells which is 77.8 %. However, when replacing the p-a-Si:H layer with MoO_x , a reduction in FF of about 3.2 %_{abs} with respect to classical SHJ cell was observed. The FF was not further decreased for the MolyATOM cells thanks to the pre-annealing treatment in the cell process. The pre-annealing treatment of MolyATOM cells and Moly/in cells effectively reached the average R_S of $1.4 \Omega \cdot cm^2$ and $1.4 \Omega \cdot cm^2$, respectively, which is at a similar level to that in classical SHJ cells ($R_S \sim 1.4 \Omega \cdot cm^2$). This gain in FF and R_S due to pre-annealing treatment mostly come from the hole-contact side because ρ_c of the ATOM contact is lower than the limit of $0.1 \Omega \cdot cm^2$.

The champion MolyATOM cell achieved an efficiency of 17.6 % with J_{sc} of 33.4 mA/cm^2 , V_{OC} of 695 mV and FF of 75.8 %, for an active cell area of 16 cm^2 . So far, SHJ cells with doped a-Si:H layers at both electron and hole contacts outperform the MolyATOM cells. Overall, the gap in performance between MolyATOM and classical SHJ cells was observed and needs to be reduced. Thus, for the MolyATOM cells, good optical rear reflectance and stronger band bending at the ATOM electron contact, as well as lower R_S at the MoO_x based hole contact with higher transmittance and higher workfunction need to be achieved to attain similar or better performance compared to classical SHJ cells.

5. Conclusion

We have investigated the impact of annealing i-a-Si:H on the thermal stability of an electron contact featuring i-a-Si:H/ TiO_x /low work function metal (ATOM). Like MoO_x /i-a-Si:H based hole contacts, annealing the i-a-Si:H layer before metal oxide deposition (i.e. pre-annealing treatment) was helpful to increase the thermal stability of the ATOM contact. Doping-free cells, called MolyATOM cells, were prepared using both MoO_x /i-a-Si:H based hole contacts and ATOM based electron contacts. The champion cell fabricated with pre-annealing treatment at $300 \text{ }^\circ\text{C}$ for 20 min achieved a J_{SC} of 33.4 mA/cm^2 , a V_{OC} of 695 mV, a FF of 75.8 % and an efficiency of 17.6 % with a cell area of 16 cm^2 . High efficiencies and improved thermal stability of pre-annealed MolyATOM cells were confirmed at the cell as well as module level. It is believed that hydrogen-related degradation was mitigated by the pre-annealing treatment. The hydrogen content in i-a-Si:H was decreased from 15.7 % to 11.4 % after pre-annealing treatment at 300°C according to elastic recoil detection measurements. The insights of this study will also be useful for developing another novel stable and highly performing metal-oxide based electron or hole contacts.

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