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Advanced light management techniques for two-terminal hybrid tandem solar cells

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A B S T R A C T

Multi-junction solar cells are considered for various applications, as they tackle various loss mechanisms for single junction solar cells. These losses include thermalization and non-absorption below the band gap. In this work, a tandem configuration comprising copper-indium-gallium-di-selenide (CIGS) and hydrogenated amorphous silicon (a-Si:H) absorber layers is studied. Two main challenges are addressed in this work. Firstly, the natural roughness of CIGS is unfavorable for monolithically growing a high quality a-Si:H top cell. Some sharp textures in the CIGS induce shunts in the a-Si:H top junction, limiting the electrical performance of such a configuration. To smoothen this interface, the possibility of mechanically polishing the intermediate i-ZnO layer has been explored. The second challenge that is addressed, is the significant current mismatch in these tandem architectures. To enhance absorption in the current-limiting top cell, the ZnO:Al front electrode was textured by means of wet-etching the entire tandem stack. We demonstrated that one can manipulate the morphology of the random textures by varying the growth conditions of the ZnO:Al, leading to better light management in these devices.

1. Introduction

Thin-film technologies can have certain attractive characteristics in terms of weight, shapes, and possible translucence. In search for higher voltages and efficiencies, much research has focused on multi-junction thin-film photovoltaic devices. This is not limited to the field comprising III-V semiconductors, but also thin-film technology materials have been extensively researched. Combining a-Si:H with nanocrystalline silicon – the micromorph solar cells – has widely been investigated [1], but also work on (all silicon-based) triple-junction [2] and even quadruple junction solar cells are found in literature [3,4]. Likewise, multi-junction configurations are considered in the field of organic photovoltaics [5,6]. Beyond that, merging of different photovoltaic technologies in so-called hybrid multi-junctions is also being explored. Recent development are found in combining one junction of crystalline silicon with a junction of perovskite [7] or III-V semiconductors [8]. In addition, in the field of CIGS such stacks have been proposed, including combining this junction with various dye sensitized solar cells [9–11]. In this latter field, an additional incentive is to limit the use of the relatively scarce Indium. Using multi-junction devices, thinner CIGS layer can generate higher photo-conversion efficiencies, due to both higher voltages obtained in thinner absorber layers [12,13], as well as the better spectral utilization.

In terms of contact and interconnection design, multi-junction devices can be categorized in two configurations; two-terminal and four-terminal devices. The challenge for four-terminal multi-junction devices lies in the complex electrical components required for the integration of these cells in modules and complete systems. The challenge for two-terminal devices is to accomplish a design in which top cell and bottom cell are close to current matching while preserving a high value for the fill factor (FF). Consequently, two-terminal configurations face the biggest challenges in the design and fabrication of the cells.

This work focuses on two-terminal double-junction devices, where all layers are monolithically integrated. In these devices, two PV cells based on semiconductor absorber layers with different energetic bandgaps are stacked, with the cell based on the widest bandgap material facing the front side at which the irradiance is incident. In such a tandem cell the photons have - on average - less excess energy relative to the bandgaps, reducing thermalization losses, and thereby resulting in a more efficient utilization of the solar spectrum.

CIGS is a direct bandgap material with a high absorption coefficient till close to its band gap around 1.2 eV. This makes CIGS a suitable semiconductor material to function as a bottom cell in these devices. The amorphous silicon solar cell technology regards a wide bandgap range.
material (1.66 eV), is well-developed, and is processed at a temperature below 200 °C. These properties making a-Si:H a suitable candidate for use as top cell in this configuration.

The integration of these CIGS and amorphous silicon cells in one monolithic stack faces several challenges. First of all, in conventional single junction amorphous silicon and micromorph solar cells illumination through the p-doped layer is favorable, as these cells are constraint by the drift length of the holes. In this novel device structure, the a-Si:H top cell is to be illuminated through the n-doped layer to align it with the configuration of the CIGS. Secondly, it is challenging to grow high-quality a-Si:H layers on the rough CIGS. The relatively sharp texture results in a-Si:H with a high shunt density. In addition, the amorphous silicon cell is current limiting. Increasing the light absorption of the top cell can significantly improve the performance of the tandem cell.

In this work, we demonstrate two approaches to tackle the above limitations to improve the performance of a-Si:H/CIGS tandem device. First of all, the CIGS cell is smoothened by mechanical polishing the ZnO interface. The smoother CIGS half fabricate facilitates the growth of a shunt-free a-Si:H top cell, increasing the $V_{oc}$ and fill factor of the tandem cell. Very little related research to this methodology of interface smoothening can be found in literature. This concept has previously been demonstrated for micromorph solar cells where an intermediate i-ZnO layer [14] or nanocrystalline SiO$_2$:H [14–16] has been polished. An extensive study of polishing ZnO built the framework for this research [17]. Mechanically polishing layers of top of CIGS solar cells is unprecedented. Such smoothening approach induces additional challenges in terms of optimization, due to relative weak semiconductor-metal bonding.

The second approach is based on texturing of the ZnO:Al front electrode to enhance the generated photocurrent in the top sub-cell by forward scattering. Texturing of TCOs is a well-known strategy for light management in photovoltaic devices [18,19]. In this work, a wet-etching procedure is used for creating a textured front TCO surface. When a TCO is subject to an acid, the acid starts to react with the material, along its surface. Due to the columnar growth of TCOs, the acid will also penetrate partially along the grain boundaries and voids and etch the bulk of the material. Due to etching along the increased surface area at the grain boundaries, an inhomogeneous lateral etching speed is realized and a textured surface is realized. Likewise, when the acid penetrates and etches inside the voids, these develop into relatively larger craters. Wet-etching a TCO using this procedure, creates random textures that induce light scattering and improve light coupling. By varying the type of TCO, deposition parameters, and etchants, one can alter the type of textures.

2. Methodology

2.1. Fabricating bottom sub-cell

The half fabricate comprises the glass substrate, molybdenum back contact, CIGS absorber layer, CdS buffer layer and the intrinsic ZnO (glass/Mo/CIGS/CdS/i-ZnO). A 1 mm thick sodalime glass substrate was cleaned in five steps. Firstly, it was scrubbed clean using isopropanol on tissue, followed by immersion in 4 consecutive baths of 10 min; first an ultrasonic bath of acetone; secondly an ultrasonic bath of isopropanol, followed by two baths of HNO$_3$ (a 99% HNO$_3$ bath at room temperature, and a 69% HNO$_3$ bath at 100 °C). After these steps the substrates are extensively rinsed with de-ionized water (DI water).

A 420 nm layer of molybdenum was sputtered on this substrate in two pressure steps. First, a more porous layer to adhere well to the substrate was deposited, followed by a more dense layer to form a well-conducting back contact. A 1 μm CIGS was deposited using three-stage co-evaporation. The CIGS deposition, the sodium diffuses into the CIGS layer through natural diffusion from the substrate. The first stage was processed at 400 °C, stage II and III at 550 °C. No post-deposition treatments were performed. The 70 nm CdS was deposited by 6-min chemical bath deposition at 65 °C, using the precursors thiourea and CdSO$_4$, and NH$_3$ as a complexing agent.

Using these two pressures, polishing times were varied. After polishing, the slurry was removed from the surface by scrubbing with acetone and DI-water. These smoothened half fabricates have been characterized with an atomic force microscope and inspected with cross sectional scanning electron microscopy (SEM).

2.2. Polishing

The half fabricate (glass/Mo/CIGS/CdS/i-ZnO) with (natural) rough surface morphology was cut in samples with a size of 2.5 × 2.5 cm. The samples were mechanically polished using non-dry polishing fluid and a 200 mm diameter chemopolishing pad. The rotational speed both the mounting head and the table was set to 150 RPM, rotating in opposite directions. The applied downward forces were set to 10 N or 20 N, resulting in 16 kPa and 32 kPa pressure, respectively. Using these two pressures, polishing times were varied. After polishing, the slurry was removed from the surface by scrubbing with acetone and DI-water. These smoothened half fabricates have been characterized with an atomic force microscope and inspected with cross sectional scanning electron microscopy (SEM).

2.3. Fabrication of the top sub-cell

The top subcells were fabricated in a PE-CVD cluster tool having dedicated chambers for layers with different types of doping. Firstly, a 30 nm of phosphorus doped hydrogenated nanocrystalline SiO$_x$:H (p-nc-SiO$_x$:H) was deposited. This was followed by two nanocrystalline SiO$_x$:H layers of 7 nm and 14 nm in thickness with different doping concentrations. The first p-nc-SiO$_x$:H layer (with a narrower bandgap) serves as part of the tunnel recombination junction. The following layer was a wider bandgap p-layer to serve as hole-collector in the top subcell. After deposition of the amorphous silicon, another 30 nm n-type nanocrystalline n-nc-SiO$_x$:H was deposited, and the cell is completed by RF sputtering an ZnO:Al 250 nm front electrode. As a final step, a 300 nm Ag front grid was evaporated on the devices. The short circuit current of these devices was determined with an EQE, and $V_{oc}$ and FF with a triple A rated JV tester, calibrated with calibration diodes from the Fraunhofer ISE institute.

2.4. Texturing of the front electrode

In this work, the ZnO:Al front electrode has been wet-etched to create a front texturing. Regarding the experimental design, it is crucial that there is a full area coverage of ZnO:Al, as the 250 nm of ZnO:Al (after etching) serves as a protective layer, preventing any water and acid ingress to the cells underneath. The deposition pressures have been varied between 1.5 μbar, 2.5 μbar, and 3.5 μbar and deposition temperatures between 100 °C, 125 °C, and 150 °C. Higher temperatures are unsuitable in this architecture, considering that ZnO:Al is the final layer to be processed in these devices and higher temperatures would induce undesired diffusion of cadmium. In all cases etching was done in a 0.5% HCl bath for 45 s. After etching, the sample was rinsed for at least 3 min in DI water. The initially deposited thicknesses of ZnO:Al were optimized as such that a 250 nm layer of textured ZnO:Al remained after etching. As the etching rate was dependent on the deposition parameters, the initial thicknesses of the ZnO:Al was varied.

3. Results

3.1. Polishing

The deposition of the 500 nm i-ZnO layer was performed in 10 consecutive passes of the carrier by the sputtering target. This
procedure has an effect on the growth of the crystal structure as shown in Fig. 1 below. The crystal structure is still showing continuous columnar growth as is expected for state-of-the-art ZnO layer. However, an interesting artifact is observed induced by the movement of the carrier relative to the sputtering target. Fig. 1 shows that a wiggle structure is created.

During the mechanical polishing procedure, it is important to secure that the i-ZnO layer is not fully removed. A fully removed i-ZnO would risk a partly or full removal of the underlying CdS layer as well. A damaged CdS layer would deteriorate the cell performance. Thus, the removal rate is an essential variable. The removal rate (RR) was estimated using the modified Preston equation (Eq. (1)), compiled by Luo et al. [20]:

\[ RR = K(P - P_{th})v + R_c. \]  

In this equation the removal rate is expressed in terms of pressure \( P \), linear velocity \( v \) and the following constants; the Preston constant \( K \), a pressure constant \( P_{th} \), and a removal constant \( R_c \). Gupta et al. [17] determined empirically that for polishing i-ZnO these constants should be to be \( K = 2.69 \times 10^{-10} \text{Pa}^{-1}, P_{th} = 1.235 \times 10^4 \text{Pa} \) and \( R_c = 9 \text{Å/min} \), respectively. Using these values the initial removal rate was been estimated to be \( RR = 209 \text{nm/min} \).

After polishing, the thickness was determined using SEM. The removal rates have been verified using 500 nm thick i-ZnO layer on Asahi-VU substrates. In Fig. 2 the cross-sectional SEM images of the unpolished and polished ZnO are displayed. In addition, an isometric overview of the polished sample is shown in Fig. 2(c). The dashed line in Fig. 2(b) indicates the very smooth interface and surface morphology of the polished interface. Some pinholes are observed in the isometric SEM image. These pinholes are assumed to reflect the pre-existing voids in the i-ZnO layer. The removal rate was determined to be \( 204 \pm 30.3 \text{nm/min} \), and thus consistent with removal rates reported in literature [17].

The mechanical polishing process was repeated on a CIGS bottom cells (glass/Mo/CIGS/CdS/i-ZnO). The surface morphology of these (un)polished interfaces were characterized using an atomic force microscope and the obtained RMS roughness is tabulated in Fig. 3 below. The decaying trends are correlated to the applied pressure and polishing time. Polishing for 60 s with 20 N downward force resulted in an impressive low RMS roughness of 2.80 nm. From experience, longer polishing times than 60 s are inadvisable to be used. Longer polishing times induced some unwelcome edge effects such as peeling of the CIGS layer.

Looking at the external parameters (Table 1 and Fig. 5a) one can see that especially the \( V_{oc} \) strongly increased after polishing. We think that this enhancement is partially induced by the thinner ZnO layer, and partially due to the better growth conditions for the amorphous silicon. As the \( V_{oc} \) of the polished device is very close the summation of the \( V_{oc} \) of the single junctions (800–850 mV and 610–630 mV for a-Si:H and CIGS, respectively) and no significant impact is observed in the fill factor, we conclude that the electrical performance is not significantly affected and we successfully polished an intermediate interface in these devices.

Regarding the spectral response of these solar cells (Fig. 5b), it is observed that polishing the interface significantly increases the interference. This significantly influences the spectral response of the bottom cell. However, as this layer is not current-limiting, this does not reflect in the device performance.

3.2. Textured ZnO:Al

The front electrode ZnO:Al was fabricated in the cluster-tool, with a stationary substrate holder. Hence, after the incubation phase of this columnar growth of this layer, a homogenous growth is expected. Fig. 6
shows a scatter plot of the measured RMS roughness versus the measured correlation length of the various surface textures of the several varied parameters. In this figure, the contours indicate the aspect ratio of the surface texture, which is defined as the ratio between the RMS roughness and the correlation length. The higher the aspect ratio, the larger the forward scattering in the visible spectral range can be expected.

Upon reducing the deposition pressure (star – triangle – square) an increasing spread of values for the correlation length is observed. We explain this relation by the fact that a highly energetic precursor can induce defects and vacancies sub-surface in the bulk of the growing layer. At higher deposition pressures, the precursor atoms and ions experience more collisions with the argon atoms on their path towards the surface. Consequently, the atoms arrive at the surface with less energy, leading to layers with a higher density.

During the wet-etching process, denser films result in a more homogeneous lateral etching, due to dominance of etching along the surface rather than along the grain boundaries. As the films deposited at lower deposition pressures contain more voids, which the acid can penetrate along the grain boundaries, leading to more bulk etching and as a consequence larger features and correlation lengths.

In addition, Fig. 6 reveals the dependence on the substrate temperature during the ZnO:Al deposition. It shows that the higher the substrate temperature, the less the correlation length is affected by etching. We believe that deposition at higher substrate temperatures leads to higher mobility of the precursor species at the surface of the film leading to denser films. Denser films means that the role of bulk etching versus surface etching is reduced and consequently its effect on the correlation length.

As a result of these different effects the different processing conditions lead to three different regimes of surface morphology. One regime with high RMS roughness & large correlation length (I), one regime with high RMS roughness and smaller correlation lengths (II) leading to the highest aspect ratio. The last regime corresponds to most of the dense films that are clustered in Fig. 6 in the parameter space with
small correlation lengths and limited RMS roughness (regime III).

These propositions are supported by Fig. 7. The SEM images show four surfaces of wet-etched ZnO:Al, where the ZnO:Al was deposited at various substrate temperatures and pressures. The surface morphology in Fig. 7(a) corresponds to ZnO:Al deposited at both the lowest temperature and pressure (regime I). The dark spots observed in the ZnO:Al (a) are an indication of the voidy nature of this film. Images 7(b) and 7(c) visually show comparable surface morphology, with significant less dark spots. The SEM image depicted in Fig. 7(d), supports that combining a high temperature with high deposition pressure (regime II) leads to a more compact film and more surficial etching. This is revealed by the textures in this image being significantly more alike oriented when compared to the 7(a)-(c).

The best light trapping is expected for textured ZnO:Al with the combination of highest RMS roughness and a correlation length close to the thickness of the absorber layer of the top cell (250–300 nm), favoring the ZnO:Al deposited at a pressure of 1.5 µbar at 150 °C. Fig. 8(a) shows that all textures significantly improve the $J_{sc}$ of the devices. Additionally, one see that there is some spread in the $V_{oc}$ between the different textures. However, this is considered to be due to processing conditions, supported by the fact that also the ‘as grown’ reference device also has a fairly limited electrical performance.

For describing the light management qualities of these different textures it is more valuable to focus on overall device reflection (which can reduced by better light incoupling) and reduced coherence (indicating better light scattering). Fig. 8(b) shows how three of the textures (from the different regimes) affect the solar cell reflectance. Data of several textures is also summarized in Table 2 below. This shows that it can be concluded that all textures will significantly improve light incoupling (4 – 5% absolute reduction in averaged solar cell reflectance) as well as decrease the coherence of the light. Furthermore, the texture fabricated from the AZO deposited at 1.5 µbar at 150 °C (from regime II) outperforms the other textures both in terms of having the lowest average reflection as well as the smallest standard deviation of the reflected light, indicating that this texture functions best to reduce the coherence of the light.

Fig. 9 shows the EQE and 1-R (where R is reflectance) of tandem a-Si:H/CIGS tandem cells. One tandem cell has the optimized textured
working hybrid tandem devices have been presented. The RMS intermediate i-ZnO layer in hybrid a-Si:H/CIGS tandem solar cells optimizing the top cell architecture. We expect to further improve current matching by further increased the current of the top sub-cell with 7.9% relatively to the cell performance, eliminating the destructive interference dip observed in spectral ranges. The unpolished tandem structure. We attribute this increase to both reducing the thickness of the i-ZnO and improved growth conditions for ultrathin CIGS layers. However, as currently the top sub-cell is current-averaged solar cell reference. It has been found in 11.59 ± 2.61% reflection over a wavelength range of 300–1000 nm. As texturing our devices did not deteriorate the electrical and optical performance of our devices, we conclude that the front electrode can serve as a barrier against the acid treatment and water rinsing, protecting the cells underneath.

The demonstrated procedures are not solely applicable in architectures considering these type of the absorber layers, but with slight alterations many different types of multi-junction solar cells can benefit from these techniques. Therefore, this work can be considered as a fundamental study that can contribute to work towards high quality monolithic two-terminal hybrid tandem solar cells.

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