

Light Harvesting Schemes for High Efficiency Thin Film Silicon Solar Cells

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ABSTRACT — In Thin Film Silicon (TF-Si) solar cells light harvesting schemes must guarantee an efficient light trapping in the thin absorber layers without decreasing the silicon layers quality and consecutively the p-i-n diodes electrical performance. TF-Si solar cells resilience to the substrate roughness is reported to be possibly improved through optimizations of the cell design and of the silicon deposition processes. By further tailoring the superstrate texture, amorphous silicon / microcrystalline silicon (a-Si:H/ μ c-Si:H) tandem solar cells with an initial efficiency up to 13.7 % and a stabilized efficiency up to 11.8 % are demonstrated on single-scale textured superstrates. An alternative approach combining large and smooth features nanoimprinted onto a transparent lacquer with small and sharp textures from as-grown LPCVD ZnO is then shown to have a high potential for further increasing TF-Si devices efficiency. First results demonstrate up to 14.1 % initial efficiency for a TF-Si tandem solar cell.

Index Terms — thin film, amorphous silicon, light trapping, photovoltaic cells, silicon, tandem cells.

I. INTRODUCTION

To further enhance the potential of Thin Film Silicon (TF-Si) photovoltaic technology, efficient light harvesting schemes are essential to maximize absorption in the thin absorber layers. Most importantly, all optical performance optimization must be realized without negatively impacting the electrical performances of the p-i-n junctions. Parasitic absorptions in the electrodes and doped layers have to be minimized while still ensuring sufficient conduction. In that respect, progress realized in our laboratory with the development of mixed phase doped silicon rich silicon oxide materials is detailed in section II, while the implementation of a high mobility transparent conductive oxide in combination with a transparent textured superstrate realized by nanoimprinting is detailed in section IV. In order to then efficiently trap light in the thin absorber layers, light scattering at rough interfaces is usually used and is typically introduced by utilizing surface-textured substrates. However, it was also reported that TF-Si layers deposited on highly textured substrates can exhibit locally low quality material regions or 2D nano-porous phases, which lead to deterioration of the electrical performance of TF-Si p-i-n junctions [1-7]. To maximize TF-

Si devices efficiency, one has therefore to find the optimum trade-off between high electrical and optical performances. This trade-off is first reported in section II to be possibly optimized by the development of cell design techniques and silicon deposition processes which improve TF-Si solar cell resilience to the substrate roughness, allowing retaining high electrical performance on rougher substrates [1, 5-6]. Then, the morphology of the superstrate texture can be adapted to provide an efficient light scattering without exhibiting too sharp features, so as to ensure high quality silicon films [7]. Best amorphous silicon / microcrystalline silicon (a-Si:H/ μ c-Si:H) micromorph tandem solar cells obtained so far on single-layer textured superstrates made of plasma treated ZnO grown by low-pressure chemical vapor deposition (LPCVD) are reported in section III.

Alternative approaches to the standard single-scale textured electrode superstrates are finally described in section IV. Conduction, transparency and texture requirements of TF-Si superstrates can be possibly decoupled via a multi-layer approach. A highly transparent TCO layer can ensure conduction with no restriction on its texture, while light scattering is achieved with a textured surface realized with a highly transparent medium. In that way, conduction – transparency optimization of the TCO is decoupled from the texture morphology optimization. Then, multi-scale texturing is reported: it was developed to provide efficient light incoupling and light scattering over the full spectral range while enabling morphologies adapted for the growth of high quality μ c-Si:H material and therefore for high electrical performance TF-Si devices [8]. Following such approach, a tandem device with an initial efficiency surpassing 14 % is reported.

II. CELL ARCHITECTURE AND PECVD PROCESS CONDITIONS FOR TEXTURED SUBSTRATES

Doped nano-structured silicon rich silicon oxide layers were synthesized by PECVD from a gas mixture of silane (SiH_4), hydrogen, of an oxygen precursor (CO_2 in this study)

and of phosphine (PH_3) or tri-methyl boron ($\text{B}(\text{CH}_3)_3$) respectively for n-type and p-type doping. Optimization of the PECVD processes permitted a clear phase separation in the developed layers, with the creation of silicon nano-filaments elongated in the growth direction, embedded in a silicon oxide matrix [9], as shown in the energy-filtered TEM image of Figure 1. For the prepared mixed-phase material, a high n-type doping ratio $\text{PH}_3/(\text{PH}_3+\text{SiH}_4)$ of up to 5 % could be realized without impacting the crystalline volume fraction or the nano-structure of the silicon rich phase. For such n-type films, layers with a refractive index down to 1.7 at a wavelength of 600 nm were successfully implemented in cells. For p-type doping of the developed silicon rich silicon oxide materials, optical and electrical performances are detailed in [10]. The p-type doping ratio $\text{B}(\text{CH}_3)_3/(\text{B}(\text{CH}_3)_3+\text{SiH}_4)$ is here limited to about 0.5 %. A higher $\text{B}(\text{CH}_3)_3$ concentration is then shown to affect the crystalline volume fraction of the Si phase [10], consecutively reducing the layer conductivity. In comparison to n-type doping, p-type layers with a minimum refractive index of about 2.5 can therefore be developed with sufficient transverse conductivity to be integrated in cell. For both doping types and both a-Si:H and $\mu\text{-Si:H}$ junctions, the optimum oxide layers integrated in the cells detailed in the following correspond to layers with a maximum oxygen content (i.e. maximum CO_2/SiH_4 gas flux ratio) while keeping a sufficient transverse conductivity to not impact the cell electrical performance.

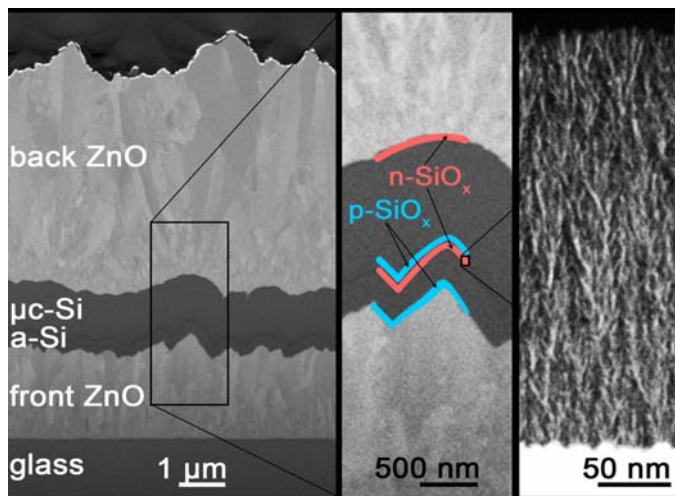


Fig. 1. Left: SEM image of a FIB cross section of an a-Si:H/ $\mu\text{-Si:H}$ tandem cell with ZnO front and back electrodes. Middle: zoom in the SEM image to illustrate the implementation of p-type and n-type doped silicon rich silicon oxide layers in both top and bottom cell. Right: EFTEM image of a n-type doped silicon rich silicon oxide layer exhibiting a semiconducting silicon filament nano-structure embedded a highly transparent silicon oxide matrix: white contrast regions correspond to silicon rich phase, dark contrast region to silicon oxide rich phase (right, [9]).

The implementations of both n-type and p-type silicon rich silicon oxide layers in thin film silicon solar cells permit to:

- Improve the light management [9-14]: transparent intermediate reflector layer in micromorph solar cells and reduced parasitic absorption (for instance highly transparent window layer).
- Increase electrical performance and cell resilience to the substrate texture [1, 5-6]: electrical performance gain scales with substrate roughness, and up to 6 % and 20 % gains were demonstrated respectively for a-Si:H and $\mu\text{-Si:H}$ p-i-n junctions [1].

Silicon oxide doped layers therefore not only improve the cell optical performance but were demonstrated to also mitigate the impact of the local low quality material regions onto the cell electrical properties [1, 15]. The implementations of these layers both for a-Si:H and $\mu\text{-Si:H}$ p-i-n junctions thus permits the integration of rougher superstrates: these doped layers are integrated both in the top and bottom cells of the tandem cells presented in next sections (configuration as shown in Figure 1).

In addition, it was recently demonstrated that plasma-enhanced chemical vapor deposition (PECVD) process conditions of silicon layers have to be optimized not only to provide a high bulk material quality but also to reduce the density and/or to increase the quality of the local defective material regions which typically grow on high angle V-shaped valleys of the substrate texture [1, 5]. Some PECVD processes can lead to films with low bulk defect density, but with strongly defectives localized areas near cracks. It must therefore be emphasized that for a given plasma process, the global material quality and the resulting cell electrical properties will strongly depend on the substrate texture. Optimum plasma conditions will therefore depend on the exact morphology on which the film is grown.

The implementation of silicon oxide doped layers combined with PECVD processing conditions which can provide a dense absorber layer with reduced local defective regions creation recently permitted to reach an unprecedented $\mu\text{-Si:H}$ single junction solar cell efficiency of 10.9 % (for a cell area of $0.5 \times 0.5 \text{ mm}^2$), as realized in two different reactors in our laboratory [5-6].

II. MICROMORPH TANDEM CELL ON LPCVD ZnO

TF-Si micromorph tandem solar cells were first optimized for single-layer textured electrodes. ZnO grown by LPCVD provides as-grown pyramidal features exhibiting excellent light scattering properties. Even in the case of optimized TF-Si tandem solar cells using doped layers and PECVD processes as described in section II, sharp features of LPCVD ZnO texture are still detrimental to the cell electrical properties, mainly for ZnO layers with a thickness above 1.2 μm , i.e. for substrate feature sizes with peak to peak height above 300 nm and mean angle of surface incidence above 35

degrees. For larger feature sizes (thicker ZnO layers), a plasma treatment of the electrode surface can smoothen the most pronounced V-shaped valleys of the texture and allow to recover high silicon layers quality (reduced local nano-porous regions) and high cell electrical properties, at the possible price of reduced light in-coupling and scattering [7]. An optimum ZnO surface could therefore be determined for our a-Si:H/ μ c-Si:H tandem solar cell design: it has a typical thickness of 2 to 3 μ m and a soft plasma treatment of the surface [7]. Its typical surface texture is shown in Figure 2.

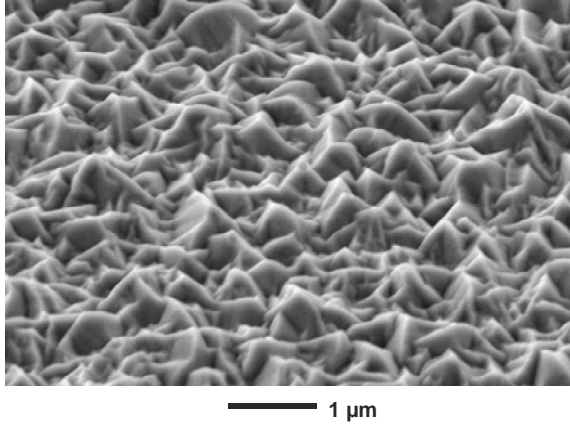


Fig. 2. SEM image of a tilted top surface of an optimum LPCVD ZnO front electrode: $\sim 2.5 \mu$ m thick layer with a soft plasma treatment.

Tandem cells were optimized on such substrate: they implement a 250 nm thick a-Si:H top cell, a 50 nm thick silicon rich silicon oxide intermediate reflecting layer (IRL), a 1.4 μ m thick μ c-Si:H bottom cell, a LPCVD ZnO back electrode and a dielectric white back reflector. All Si layers were deposited in KAI reactor types. The external quantum efficiency (EQE) and the corresponding short circuit current densities (J_{SC}) obtained with these devices in the top and bottom cells are shown in Figure 3 and described in Table I.

To further increase the light in-coupling, micrometric random square based pyramids were imprinted on a lacquer disposed at the air/glass interface. The application of this anti-reflective coating (ARC) described in [16] is shown to yield 5.8 % and 2.5 % relative J_{SC} increase respectively in the top and bottom cells (~ 4 % gain in total current). The pyramids morphology permits to reduce primary reflection losses via double rebound of the incoming light at the glass surface. In addition, possible anti-escaping effect and slight light diffusion result in a small improvement also in light trapping in the device, as described in [16]. Overall, combining the optimized ZnO superstrate and cell design with the micrometric texture at the air/glass interface permits to achieve a high total J_{SC} of 27 mA/cm^2 in the developed tandem device, even for a bottom cell thickness of 1.4 μ m. Most importantly, high optical performance is realized while maintaining high electrical performance, as

detailed in Table I. A high initial efficiency of 13.7 % is consecutively realized for a reasonably thick device (total Si layers thickness $< 1.7 \mu$ m). This solar cell current matching condition was not optimum: the tandem cell is in a top limitation case after 1000 hours of light induced degradation under a class A 1 sun illumination at 50 $^{\circ}\text{C}$. Still, a stabilized efficiency of 11.8 % was achieved with such device, demonstrating that stabilized efficiency approaching 12 % can certainly be realized with the use of single-layer LPCVD ZnO front and back electrodes and with a reasonable thickness of Si layers deposited in KAI reactors.

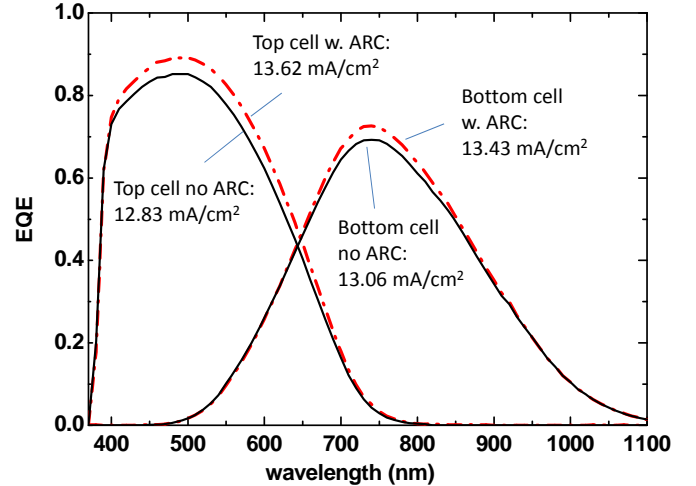


Fig. 3. External Quantum Efficiency (EQE) curves of a 250 nm thick a-Si:H top cell and a 1.3 μ m thick μ c-Si:H bottom cell implemented in a micromorph tandem solar cell deposited on a LPCVD ZnO single scale textured substrate and integrating a 60 nm thick silicon oxide intermediate reflecting layer.

TABLE I. Solar cell performance of 1 cm^2 a-Si:H/ μ c-Si:H TF-Si tandem solar cells implementing LPCVD ZnO front and back electrodes, 250 nm a-Si:H top cell, 50 nm IRL and a 1.4 μ m μ c-Si:H bottom cell.

	$J_{SC \text{ top}}$ (mA/cm^2)	$J_{SC \text{ bottom}}$ (mA/cm^2)	V_{OC} (V)	FF (%)	Eff. (%)
Initial state no ARC	12.85	13.06	1.395	72.5	13
Initial state w. ARC	13.6	13.4	1.395	73.5	13.7
Stabilized state after 1000 h LID, w. ARC	13.05	13.3	1.36	66.4	11.8

III. MULTI-SCALE SUPERSTRATES

Textures employed in TF-Si solar cells to provide an efficient light-trapping are generally dictated by the underlying growth and etching kinetics of the textured transparent conductive oxide (TCO) and/or glass. We recently

developed transparent electrodes with arbitrary surface morphologies by using ultraviolet nano-imprinting lithography (UV-NIL) to replicate an arbitrary structure made from a transparent or opaque arbitrary master material [17-18]. The textured nanoimprinted lacquer can then be combined with a thin and highly transparent hydrogen doped indium oxide $\text{In}_2\text{O}_3\text{:H}$ (IOH) TCO layer [18-19]. The efficient light scattering properties of the textured transparent lacquer combined with the high transparency and conductivity of IOH were shown to result in a high total current of 26 mA/cm^2 in a micromorph cell using a thin $\mu\text{-Si:H}$ bottom cell ($1.1 \mu\text{m}$) and no AR coating [18]. This approach shows a high potential as the texture is free from morphological growth constraints, while the employed TCO guarantees a high transparency [18-19]. These developments can then be combined with LPCVD ZnO electrodes to build nano-scale to micro-scale multi-textured superstrates. The objectives following such approach are to provide efficient light in-coupling and light scattering over the full spectral range with different texture scales [20-21] while enabling morphologies adapted for $\mu\text{-Si:H}$ cell growth [8]. Large and smooth LPCVD ZnO pyramids were replicated by UV-NIL and combined with a thin and highly transparent IOH layer to provide conduction. Finally a thin non-intentionally doped LPCVD ZnO layer was deposited to add small and sharp pyramidal features to the superstrate morphology. The obtained surface morphology is shown in Figure 4.

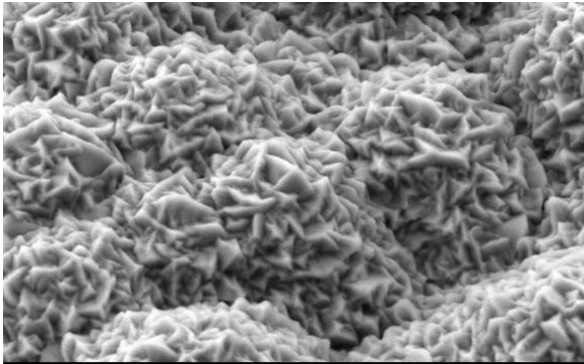


Fig. 4. Top: SEM image of a tilted top surface of a multi-scale superstrate implementing large understructures covered with small and sharp features of a thin LPCVD ZnO layer.

When used as front electrode for a micromorph cell with respective top cell, intermediate reflective layer (IRL) and bottom cell thicknesses of 260 nm , 50 nm and $2.4 \mu\text{m}$, a very high total current density of 28.3 mA/cm^2 was reached. Similar micromorph solar cells were deposited directly on an IOH – thin LPCVD ZnO stack, and on a flat IOH layer directly on a flat glass.

The thin LPCVD ZnO layer is first shown to allow for a clear current gain in the top cell with almost 2 mA/cm^2 gain with respect to the flat IOH electrode case. The small

sharp pyramidal features are confirmed to provide an efficient light in-coupling and trapping for the top cell, while still allowing for the growth of high quality $\mu\text{-Si:H}$ bottom cell, so that only 10 mV of V_{OC} is lost in comparison to the electrically ideal flat electrode case. However, current is not sufficient in the bottom cell. This limitation is then solved by adding the large and smooth features understructure in the multi-scale approach: high top cell current is conserved thanks to the small and sharp features of the thin LPCVD ZnO layer, while the bottom cell light trapping is boosted by the large features understructure, resulting in up to 1.4 mA/cm^2 current gain in the bottom cell in comparison to the IOH – thin LPCVD ZnO electrode. Here again, a high V_{OC} of 1.41 V is conserved (20 mV loss in comparison to the flat electrode electrical ideal case), demonstrating that the two structures employed to realize an efficient light trapping in both sub-cells are adapted for the growth of high quality silicon layers. Overall, an initial efficiency of 14.1% is demonstrated for a thin film silicon tandem solar cell following the presented multi-scale approach.

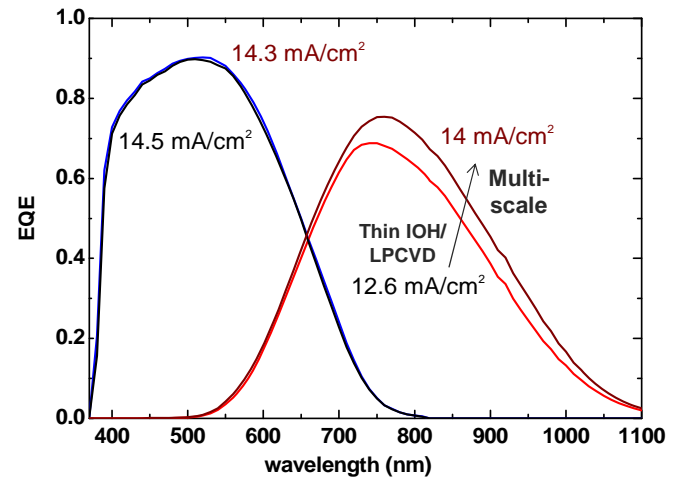


Fig. 5. External Quantum Efficiency (EQE) curves of micromorph tandem solar cell deposited on a IOH – thin LPCVD ZnO superstrate and on a multi-scale superstrate.

TABLE II. Initial state solar cell performance of 1 cm^2 a-Si:H/ $\mu\text{-Si:H}$ TF-Si tandem solar cells on various substrates (all cells have an ARC on the glass as described in [16]).

	$J_{\text{SC top}}$ (mA/cm^2)	$J_{\text{SC bottom}}$ (mA/cm^2)	V_{OC} (V)	FF (%)	Eff. (%)
Flat IOH	12.6	11.0	1.432	71.8	11.3
IOH + thin LPCVD ZnO	14.5	12.6	1.42	75	13.4
Nanoimprinted under- structure + IOH + thin LPCVD ZnO: multi-scale	14.3	14	1.41	71.5	14.1

V. CONCLUSION

High efficiency thin film silicon solar cells could be developed by optimizing the trade-off between optical and electrical performances. Advanced nanomaterials (silicon rich silicon oxide mixed phase layers) and PECVD processes optimized for dense $\mu\text{-Si:H}$ with a high bulk quality were shown to allow for up to 13.7 % initial efficiency and 11.8 % stabilized efficiency for TF-Si tandem solar cells deposited on an optimum LPCVD ZnO electrode. Advanced superstrate concept comprising large and smooth understructure guaranteeing light trapping for the bottom cell and small and sharp surface features leading to high light in-coupling and light trapping in the top cell was shown to have a high potential to realize an efficient light trapping while conserving high electrical performance. First results demonstrate a very high efficiency tandem solar cell with an initial efficiency of 14.1 % based on this concept.

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