# PROCESS DEVELOPMENT OF SILICON HETEROJUNCTION INTERDIGITATED BACK-CONTACTED (SHJ-IBC) SOLAR CELLS BONDED TO GLASS

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ABSTRACT: In imec's i<sup>2</sup>-module concept, silicon heterojunction interdigitated back-contacted (SHJ-IBC) solar cells are fabricated on monocrystalline foils bonded to glass. The proposed technology allows for cell processing on thin wafers mechanically supported by the glass, increasing the yield of processing such thin wafers. A process sequence for SHJ-IBC cell fabrication that can be applied to bonded thin foils is described. We investigated and optimized individual process steps on thick wafers. Then the developed steps were integrated into a process flow to fabricate solar cells on wafers with different thicknesses and bonding agents. On wafers with a thickness of 190  $\mu$ m, functional cells with efficiencies of 22.6% and 21.7% were made on freestanding and silicone bonded wafers, respectively. On thin wafers of 57  $\mu$ m, our best SHJ-IBC cell on an EVA bonded wafer exhibits excellent V<sub>oc</sub> of 740 mV and efficiency of 20.0%, which demonstrates the high potential of the i<sup>2</sup>-module concept.

Keywords: Amorphous Silicon, Heterojunction, Silicon Solar Cell, interdigitated back-contacted, Superstrate Processing

#### 1 INTRODUCTION

The cost of silicon material accounts for approximately 61% of today's solar cell cost and 30% to 40% of the total c-Si solar module cost [1]. Hence, to reduce the silicon consumption, fabrication of solar cells on thin monocrystalline silicon foils is being studied by several groups [2-4].

Imec proposed the i<sup>2</sup>-module concept, which allows for large-scale module level processing of interdigitated back-contacted (IBC) solar cells on 40  $\mu$ m thin silicon foils bonded to glass [5]. The glass delivers mechanical support to the thin foil and thus significantly reduces the foil breakage during cells processing. The choice for an IBC device allows to process the junction and the contacts after bonding. Certain constraints are imposed to the cell process mainly due to the bonding and the glass. For that reason, an a-Si:H heterojunction process is chosen for junction formation as that allows to keep the processing temperature low.

In this contribution, we report the recent progress in the process development of silicon heterojunction IBC (SHJ-IBC) solar cells. An integration process is presented, which is fully compatible with thin foils bonded to glass. Functional cells were fabricated on both freestanding and bonded n-type float zone (FZ) wafers with different thicknesses. Two different bonding adhesives, i.e., silicone and Ethylene Vinyl Acetate (EVA), are evaluated at cell level. The factors limiting the cell performance are identified and the main losses of the cells are discussed in detail.

### 2 RESULTS AND DISCUSSION

### 2.1 Integration process

All cells, with an area of 3.97 cm<sup>2</sup>, were processed on 190  $\mu$ m and 57  $\mu$ m thick n-type FZ wafers (6 inch, <100>, 3 $\Omega$ -cm).

The wafers were exposed to Sulfuric acid/Ozone mixture (SOM) and following HF:HCl:H<sub>2</sub>O (2:5:93)

Front-side texturing
a-Si:H (i) front passivation + $SiN_x ARC$
Bonding to glass
a-Si:H (i/n+) rear passivation (= BSF)
SiO <sub>x</sub> /a-Si mask deposition
Litho of SiO <sub>x</sub> /a-Si mask
Wet etching of a-Si:H (i/n <sup>+</sup> )
a-Si:H (i/p+) repassivation (= emitter)
Lift-off of SiO <sub>x</sub> and a-Si (i/p <sup>+</sup> )
ITO + metallization + litho patterning
Dicing + anneal

Figure 1: Integration process flow of SHJ-IBC cells on bonded substrate.

solution for cleaning. Tetramethyl ammonium hydroxide (TMAH) based solution was used for thinning down and texturing of the wafers. As shown in Figure 1, the textured front side underwent PECVD a-Si:H passivation and SiN<sub>x</sub> antireflection coating, followed by bonding to glass using EVA or silicone encapsulant. We have reported previously that plasma-silicone interaction during a-Si:H deposition has a detrimental impact on passivation quality [6]. Hence, silicone bonded wafers were treated with Ar plasma to form a more densely crosslinked layer by local modification of the exposed silicone surface [6]. Then the depositions of a-Si:H (i/n+)and a stack of SiO<sub>x</sub>/a-Si:H were performed. This SiO<sub>x</sub> layer is a sacrificial mask for subsequent lift-off process and the top a-Si:H layer is used to protect SiOx during HF dip prior to a-Si:H (i/p+) deposition. Photolithography and wet etching were used to define the back surface field (BSF) area. The emitter was passivated by a-Si:H (i/p+) and patterned by self-alignment lift-off process, which was carried out in dilute HF. A stack of indiumdoped tin-oxide (ITO) and Cu was deposited to form the rear side metal contact. Thermal annealing was applied to the finished cells in order to improve the contact behaviour. To investigate the influence of the presence of bongding agents and glass on the cell performance, identical cells were also fabricated on freestanding wafers without bonding to glass.



Figure 2: The cross-sections (not drawn to scale) of the bonded SHJ-IBC solar cells are depicted in the diagram.

## 2.2 Device results

The cell results are shown in Table I. For the best cells before anneal, we observe an improvement of  $\Delta \eta = 0.5\%$  on freestanding cells with respect to silicone bonded cells. This is due to an absolute increase  $\Delta V_{oc} = 8$  mV and  $\Delta FF = 0.7\%$ , but more significantly due to an increase  $\Delta J_{sc} = 1.1$  mA/cm<sup>2</sup>. Figure 3 compares the external quantum efficiency (EQE) of cells on freestanding wafers with that on silicone and EVA bonded wafers. The difference in EQE suggests that glass and silicone at the front side lead to additional reflectance and absorption losses for a wide range of wavelengths. Nevertheless, a high J<sub>sc</sub> of 41.2 mA/cm<sup>2</sup> was still achieved on silicone bonded cells.

 Table I: Best Cell results on freestanding and bonded wafers

Anneal	J <sub>sc</sub> (mA/ cm <sup>2</sup> )	V <sub>oc</sub> (mV)	FF (%)	η (%)
before	42.3	735	69.7	21.5
after	41.8	730	74.8	22.6
before	41.2	727	70.4	21.0
after	40.8	734	73.1	21.7
before	40.7	718	68.6	20.0
after	39.9	724	71.6	20.7
before	38.8	738	67.5	19.0
after	38.6	740	71.7	20.0
	before after before after before after before after	Anneal $3sc(mA/cm^2)$ before42.3 afterafter41.8before41.2 afterafter40.8before40.7 afterafter39.9before38.8 afterafter38.6	Anneal $3se \ (mA/) \ (mV)$ $mA/$ $(mV)$ $max$ $(mV)$ before42.3after41.8730before41.2 $727$ after40.8734before40.7718after39.9724before38.8after38.6740	Anneal $3se$ $7se$ $7se$ $11$ $(mA/)$ $(mV)$ $(\%)$ before $42.3$ $735$ $69.7$ after $41.8$ $730$ $74.8$ before $41.2$ $727$ $70.4$ after $40.8$ $734$ $73.1$ before $40.7$ $718$ $68.6$ after $39.9$ $724$ $71.6$ before $38.8$ $738$ $67.5$ after $38.6$ $740$ $71.7$

For the cells on EVA bonded thick wafers, the best  $J_{sc}$  is 40.7 mA/cm<sup>2</sup> that is 0.5 mA/cm<sup>2</sup> lower than that of silicone bonded cells. As shown in Figure 3, this result can be attributed to the light absorption loss at short wavelength due to the UV blocker contained in EVA. And for the EVA bonded cells on 57 µm thin wafer, a  $J_{sc}$  of 38.8 mA/cm<sup>2</sup> was achieved, which is further limited by

the light absorption loss at long wavelength.

The excellent  $V_{oc}$  of 734 mV and 740 mV were obtained on the silicone bonded and EVA bonded cells after anneal, respectively, similar to the  $V_{oc}$  of cells without bonding agent. It indicates that the impact of bonding agents on the surface passivation quality has been prevented. Such achievement has been made possible by several adaptations in the process:

- The new encapsulants silicone and EVA were selected to replace the previously used PV 6100
   [7], because these alternative bonding agents were proven to be more resistant to chemicals and plasma treatments.
- An optimized curing/outgassing step reduces outgassing of bonding agents during a-Si:H deposition.
- The Ar plasma improves crosslinking of the silicone, and hence reduces the plasma-silicone interaction.
- The SiO<sub>x</sub> hard mask (~680 nm) also covers the bonding agents during rear side process and thus further reduces the interaction of plasma /chemicals and bonding agents.
- Optimum a-Si:H deposition conditions were chosen for surface passivation in presence of different bonding agents.



**Figure 3:** External quantum efficiency of a cell on freestanding wafer compared to that on bonded wafers. The thicknesses of the wafers are 190 μm.



Figure 4: Light J-V curve of the best SHJ-IBC cells after anneal on freestanding and bonded wafers.

Anneal	Cell parameters				FF loss analysis		
	$R_s$ ( $\Omega$ .cm <sup>2</sup> )	$rac{ m R_{sh}}{ m (\Omega.cm^2)}$	FF (%)	pFF (%)	∆FF <sub>Rs</sub> (% at	$\Delta FF_{Rsh}$ psolute)	
before	2.8	2.2×10 <sup>3</sup>	68.0	84.1	13.5	0.6	
after	1.6	2.1×10 <sup>3</sup>	73.8	82.5	7.9	0.6	

Table II: FF loss analysis of a cell on a freestanding wafer before and after anneal

For all cells before annealing, the efficiencies are strongly limited by the relatively low FF ( $\leq 70.4\%$ ). Thermal anneals of the finished cells at temperatures below 200°C were performed under air ambient, which yielded a significant improvement in FF. For cells on freestanding wafers, the highest FF of 74.8% has been achieved after annealing at 150°C. A detailed FF loss analysis is shown in Table II [8]. The series resistance Rs was determined by the Bowden method at different light intensities [9]. The shunt resistance R<sub>sh</sub> was determined by a linear fit of the dark I-V curve. The pseudo fill factor (pFF) was extracted from Suns-Voc measurements. The absolute FF loss  $\Delta$ FF<sub>Rs</sub> was calculated to be 13.5%, while  $\Delta FF_{Rsh}$  was only 0.6%. This reveals that the drop of the measured FF was primarily caused by series resistance. After annealing,  $R_s$  reduced significantly from 2.8  $\Omega$ .cm<sup>2</sup> to 1.6  $\Omega$ .cm<sup>2</sup>, where corresponding  $\Delta$ FF<sub>Rs</sub> reduced from 13.5% to 7.9%.

# 3 CONCLUSION AND OUTLOOK

We presented an integration process for module-level processing of SHJ-IBC solar cells, which is compatible with thin silicon foils bonded to glass. Individual process steps have been investigated and optimized. To demonstrate the developed process, highly efficient cells on 190  $\mu$ m and 57  $\mu$ m thick wafers were fabricated. Issues limiting the cell performance were identified. Current development is the introduction of thin epitaxially grown wafers instead of thin FZ wafers [10]. Next steps will be directed towards improvement of fill factor and proces simplifications, including investigations and integration of lithography-free emitter patterning approach, dry etching of a-Si:H (already demonstrated in [11]) and selective depositon of a-Si:H to replace lift-off.

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