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Diminished band discontinuity at the p/i interface of narrow-gap a-SiGe:H solar cell by hydrogenated amorphous silicon oxide buffer layer

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ABSTRACT

We optimized hydrogenated amorphous silicon oxide (a-SiO_x:H) as a buffer layer at the p/i interface of hydrogenated amorphous silicon germanium (a-SiGe:H) solar cells. This buffer layer was intended to effectively diminish the band discontinuity between the high-bandgap p-window layer and low-gap a-SiGe:H absorption layer, thereby enhancing the cell performance. The open-circuit voltage (V_{oc}) increased by 3.7% as the [CO₂]/[SiH₄] gas ratio increased from 0 to 0.06. In addition, the short-circuit current density (J_{sc}) gradually increased with the gas ratio. Using the optimized a-SiO_x:H buffer layer, a high performance of 9.6% was recorded for narrow-gap a-SiGe:H thin film solar cells. a-SiGe:H solar cells with optimized a-SiO_x:H buffer layers, used as middle subcells, are expected to enhance the total V_{oc} of triple-junction configuration solar cells.

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1. Introduction

Thin-film silicon solar cells have been the subject of extensive research for many decades as solar energy-harvesting devices [1–4]. In spite of suffering from light-induced performance degradation upon light soaking, known as the Staebler–Wronski effect [5], their competitive advantages such as low production cost, large-scale manufacturing, flexibility, light weight, and especially their potential applicability to building-integrated photovoltaic systems have continued to foster further improvements [6–8]. In particular, the recent world-record achievements of triple-junction solar cells with initial performance of over 16% have paved the way for the persistent pursuit of high-performance and cost-effective solar cells [9,10].

The main advantage of the triple-junction configuration is the use of three absorption layers with different bandgaps for capturing the full range of solar spectra and thus increasing the cell performance. In the triple-junction configuration, generally, hydrogenated amorphous silicon germanium (a-SiGe:H) solar cells play an important role as the middle subcells, in combination with

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of the a-SiGe:H middle subcells can improve the overall performance of the triple-junction configuration.

Consequently, this study aims to further enhance the cell parameters, especially V_{oc} , of narrow-gap a-SiGe:H solar cells. For this purpose, we examined a-SiO_x:H materials with various bandgaps as buffer layers at the p/i interface of a-SiGe:H solar cells. The effect of these buffer layers on the cell parameters was discussed in comparison with the conventional a-Si:H buffer layer.

2. Experiment

We used a cluster multi-chamber plasma-enhanced chemical vapor deposition system for the fabrication of a-SiGe:H thin-film solar cells. All cells were fabricated in a p-i-n superstate configuration on commercial Fluorine doped Tin oxide (FTO)-coated glass substrates with the random roughness in range of 5–20 nm. The window layers were double p-type layers, including hydrogenated microcrystalline silicon oxide (p- μ c-SiO_x:H) and hydrogenated amorphous silicon oxide (p- α -SiO_x:H) with optical bandgaps of 2.1 and 2.0 eV, respectively. The n-layers were hydrogenated microcrystalline silicon (n- μ c-Si:H). The absorption layers or intrinsic layer (i-layers) were a-SiGe:H layers with a thickness of 300 nm and an optical bandgap of 1.58 eV. In order to smooth the p/i interface, the intrinsic a-SiO_x:H buffer layers were inserted between the p-and i-layers.

Single a-SiO_x:H layers of thickness 50 nm were deposited on eagle glass. The deposition parameters of these layers are exhibited in Table 1. The electrical properties, including the dark and photoconductivity, of the a-SiO_x:H layers were measured with a paired aluminium planar electrode configuration using a semiconductor parameter analyser (EL420C). The thickness and absorption coefficient were inferred from fitting the ellipsometry spectra using the VASE J. A. Woollam system over a wavelength range of 240–1700 nm. Single a-SiGe solar cells were characterized under dark and illuminated conditions. A one-sun AM 1.5 solar tester with 100 mW/cm² light intensity was used for the illumination characterizations at 25 °C. The external quantum efficiency (EQE) model QEX7 was utilized.

3. Results and discussion

The main purpose of this work was to optimize the a-SiO_x:H buffer layer at the p/i interface of narrow-gap a-SiGe:H solar cells to significantly limit the band discontinuity between high-bandgap p-type window layers and narrow-gap intrinsic a-SiGe:H layers, which is expected to enhance the cell parameters, resulting in improved cell performance. In the first step, we examined the optoelectronic properties of the single a-SiO_x:H buffer layers with respect to modifying the $[CO_2]/[SiH_4]$ gas flow ratios during preparation. Fig. 1 shows the optoelectronic properties of 50 nm single a-SiO_x:H layers as a function of the $[CO_2]/[SiH_4]$ gas flow ratios, including the transmittance (1a), optical bandgap inferred by Tauc's plot (1 b), dark conductivity (σ_d), photo-conductivity (σ_p), and photosensitive value (1c), which is defined as ratio of the photo-to-

Table 1

Deposition parameters of a-SiO_x:H layers.

Parameters	values
[H ₂]/[SiH ₄] flow ratio	5
[CO ₂]/[SiH ₄] flow ratio	0.0–0.6
Power density	42.55 mW/cm ²
Working Pressure	300 mTorr
Deposition temperature	160 °C
Thickness	50 nm



Fig. 1. a) Transmittance; b) Plot of $(\alpha.E)^{1/2}$ versus photon energy – E (the linear fitting, dash line, for determination of the Tauc optical bandgap); and c) Dark – and Photo – conductivity (the inset is the photosensitive which is determined by photo to dark conductivity ratio) of single a-SiO_x:H layers.

dark-conductivities. The transmittance and optical bandgap of the films gradually increased with the gas ratios, whereas the electronic properties, such as σ_d , and σ_p , substantially decreased. It is well-known that the incorporation of oxygen elements into silicon thin-film materials can enhance the optical bandgap and therefore increase the transmittance [19-22]. In addition, this incorporation can cause high defect densities, reducing carrier transport, and as a result decreasing the conductivity [20,23]. The photosensitive value has been frequently used as one of the parameters to estimate the quality of a single silicon layer [24-26]. As seen in Fig. 1c, the photosensitive values gradually decreased with the gas ratios and especially decreased by one to two orders of magnitude at gas ratios greater than 0.06. The decrease in the photosensitive values suggested that the high oxygen concentration via the increased CO₂ flow rate detrimentally affected the guality of the a-SiO_x:H films, i.e., increased the defect density in the films.

In the second step, we fabricated single a-SiGe:H solar cells with various the $[CO_2]/[SiH_4]$ gas flow ratios of the a-SiO_x:H buffer at the p/i interface, as shown in Fig. 2. We examined the effect of the buffer layers on the cell parameters such as Jsc, FF, and Voc. These parameters were extracted from the illuminated J-V curves, as shown in Fig. 3a. In addition, Fig. 3b summarizes the variation of these parameters with the gas ratios. First, as can be seen in Fig. 3b, the J_{sc} value regularly increased with the gas ratios. This increase is consistent with that in the transmittance and optical bandgap of the buffer layers as shown in Fig. 1. This means that the increase in Isc could be due to the broadened optical bandgap of the buffers, resulting in reducing parasitic absorption and therefore enhanced transmittance, especially at short wavelengths, into the absorption a-SiGe:H layers. To demonstrate the enhanced short-wavelength absorption of the absorption layers, we examined the EQE measurements, as shown in Fig. 4. The results indicated the clear increase in the short wavelength of EQE with the gas ratios.

Second, the FF values in Fig. 3b decreased slightly from 0.66 to 0.64 as the $[CO_2]/[SiH_4]$ gas flow ratio increased from 0.0 to 0.06, and dropped significantly when the ratio was higher than 0.16. The decrease in FF can be due to the reduced conductivity of the a-SiOx:H buffer layers with gas ratios, which can cause considerable increases in the series resistance (R_s) value of the device. Based on the dark I–V characteristics [27], we calculated the R_s value of the a-SiGe:H cells, as shown in Fig. 5a. In addition, Fig. 5b summarizes

the R_s values of the cells as a function of the gas ratio. It can be seen that the R_s value gradually increases from 8 Ω to 14 Ω with the gas ratio. It is well-known that the FF value is proportional to the shunt resistance and inversely proportional to the series resistance of a cell device [28,29]. It can be inferred that the reduction in conductivity of the buffer layers with the gas ratio could result in an increase in R_s and therefore a reduction in the FF value.

Finally, the V_{0c} showed a low value of 0.78 V with the conventional a-Si:H buffer layer, with $[CO_2]/[SiH_4] = 0$. The V_{oc} gradually increased to 0.82 V at the ratio of 0.06 and abruptly decreased to 0.79 V at the ratios higher than 0.06. We used the amorphous semiconductor analysis (ASA) simulation program to simulate the p/i interface with various a-SiO_x:H buffer layers. The band diagram schematics of the cells are shown in Fig. 6. It can be observed that a significantly high potential barrier may be formed at the p/i interface of a cell without buffer layers. A conventional a-Si:H buffer layer can significantly reduce the band discontinuity with the ilayer, but considerably formed a band-offset with the p-layer. In addition, with high bandgap of the a-SiO_x:H buffer layer at the gas ratio of 0.6, the considerably high band-offset was formed with the i-layer, but strongly reduced at the p-side layer. In all these cases, the hole carrier transport can be significantly hindered by high potential carriers, resulting in detrimental effects on Voc and FF. Consequently, it is believed that the intermediate bandgap of the a-SiO_x:H buffer layer at the gas ratio of 0.06 can balance the bandoffset at both interfaces, as shown in Fig. 6, resulting in an enhancement in Voc. D. Lundszien el al [14,16]. Investigated complete effects of a-Si:H buffer layers at the interfaces on narrow-gap a-SiGe:H solar cells. It showed clearly that a-Si:H buffer layers can improve significantly the cell parameters such as Voc and FF, resulting in cell efficiency-enhancement. However, with a tendency to further widening the band gap of the window layers in order to enhance the light into the absorption layers [30-32], the a-Si:H buffer layers with the limited band gap seem to be a non-optimal choice for balancing the band discontinuity at the p/i interface. Consequently, tuneable-bandgap buffer layers such as a-SiC:H [18,33] or a-SiO_x:H [20,23] layers can play a potential role in the band-offset challenge of narrow-gap a-SiGe:H solar cells.

In conclusion, a high conversion $E_{\rm ff}$ of 9.6% was found with the a-SiO_x:H buffer with a [CO₂]/[SiH₄] gas flow ratio of 0.06. The cell parameters, including J_{sc} and particularly V_{oc}, were impressively



Fig. 2. Single a-SiGe:H cell structure with various the [CO₂]/[SiH₄] gas ratios of a-SiO_x:H buffer layers.



Fig. 3. a) Illuminated J-V curve characteristics of a-SiGe:H solar cells; and b) variation of cell parameters such as V_{oc} , J_{sc} , FF, and $E_{\rm ff}$ as a function of the gas ratios.

increased at this ratio, compared with those of the conventional a-Si:H buffer layer. The gradual increase in J_{sc} was attributed to the broadened optical bandgap of the a-SiO_x:H buffer layer owing to changes in the $[CO_2]/[SiH_4]$ gas ratios. The enhancement in V_{oc} could be due to the reduction in the band-offset at the p/i interface with the intermediate bandgap of a-SiO_x:H at the gas ratio of 0.06, whereas the gradual decrease in FF may be due to the increase in the R_s value of the cell device.



Fig. 4. External quantum efficiencies of a-SiGe:H solar cells in short wavelength.



Fig. 5. a) Plot of dV/dJ vs 1/J for a-SiGe:H solar cells from measured dark J(V) data. From the linear fitting, the intercept is series resistance (R_s); and b) variation of R_s as a function of the gas ratios.



Fig. 6. The band diagrams of a-SiGe:H solar cells, extracted from the ASA simulation, with various the gas ratios of a-SiO_x:H buffer layers.

4. Conclusion

We optimized the a-SiO_x:H buffer layers at the p/i interface for narrow-gap a-SiGe:H solar cells. The a-SiO_x:H buffer layer with a $[CO_2]/[SiH_4]$ gas ratio of 0.06 showed increases in V_{oc} and J_{sc} compared with the conventional a-Si:H buffer layer. A high cell efficiency of 9.6% was obtained with the a-SiO_x:H buffer at this ratio. The a-SiO_x:H buffer was credited with significantly diminishing the band-offset between the high-bandgap p-layer and narrow-gap intrinsic a-SiGe:H layer. With optimized a-SiO_x:H buffer layers, the V_{oc} values of a-SiGe:H solar cells are expected to be further improved. Therefore, the performance of the triple-junction configuration solar cells.

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