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Solar cells based on n⁺-AZO/p-BaSi₂ heterojunction: advanced opto-electrical modelling and experimental demonstration

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We performed advanced opto-electrical simulations on thin-film BaSi₂ solar cells. First, absorption spectra of BaSi₂-pn homojunction solar cells on Si substrate were calculated based on flat and/or pyramidally-textured surfaces, wherein 20-nm-thick n⁺-BaSi₂ was the topmost electron transport layer. By changing the front surface structure from flat to texture, the reflectance decreased in the wavelength (λ) range 700 – 1200 nm and the photocurrent density (J_{ph}) delivered by the photogenerated carriers in the 500-nm thick p-BaSi₂ layer increased by 1.2 mA/cm². Simulations revealed that the key factor inhibiting light absorption in the p-BaSi₂ layer was parasitic absorption in the n⁺-BaSi₂ and in the c-Si substrate. To solve these optical issues, we propose a new device structure, Al-doped n⁺-ZnO (AZO, 50 nm)/i-ZnO (20 nm)/p-BaSi₂ (500 nm) heterojunction solar cell (HJSC). In this device structure, the parasitic absorption reduced drastically, and J_{ph} reached 30.23 mA/cm². Furthermore, by replacing the Si substrate with a glass substrate, the light trapping worked more effectively, and the absorber layer thickness required for J_{ph} to saturate was reduced to 1 μ m, yielding 32.06 mA/cm². Based on these simulation results, we manufactured n⁺-AZO/p-BaSi₂ HJSC. The internal quantum efficiency exceeded 30% at $\lambda = 600$ nm, meaning that we demonstrated the operation of n⁺-AZO/p-BaSi₂ HJSC for the first time. We investigated origins of small efficiencies compared to those simulated, and found that the passivation of defects in the p-BaSi₂ layer and the reduction of carrier recombination at the i-ZnO/p-BaSi₂ interface would significantly improve the solar cell performance.

I. INTRODUCTION

Thin-film solar cells (TFSCs), such as III-V [1,2], CIGS [3], perovskite [4-6], CdTe [7,8], and others [9,10] have been developed to achieve high power conversion efficiency (η). However, large-scale deployment of these solar cells equivalent to crystalline Si (c-Si) solar cells is not easy because the materials used in TFSCs contain non-abundant and/or toxic elements. Hence, there is a special need for alternative thin-film materials for photovoltaic (PV) applications. Under such circumstances, we have focused on semiconducting barium disilicide (BaSi_2) as a new material for TFSCs [11]. It has a suitable bandgap (E_g) of 1.3 eV, a large optical absorption coefficient of $3 \times 10^4 \text{ cm}^{-1}$ for a photon energy of 1.5 eV (more than 40 times larger than that of c-Si), and excellent minority-carrier properties [12–16]. Furthermore, it is composed of only safe, stable, and earth-abundant elements. BaSi_2 has a small lattice mismatch with Si(111), i.e., 0.1% and 1.1% along the b - and c -axes, respectively, allowing for epitaxial growth on an inexpensive Si substrate [17]. For these reasons, BaSi_2 is considered to be a promising material for terawatt-class power generation. We have achieved η values approaching 10% in p^+ - $\text{BaSi}_2/\text{n-Si}$ heterojunction solar cells (HJSCs) without any special treatment for passivation [18–20], wherein the depletion region stretched toward the n-Si side and therefore most of the photons were absorbed in the n-Si region. Our next target is BaSi_2 -pn homojunction solar cells (SCs), for which η is expected to exceed 25% according to calculations (Fig. 1) [21]. Furthermore, BaSi_2 PV technology could be applied to Si-based tandem solar cells by expanding its E_g via alloying BaSi_2 with C [22]. Currently, we focus on further improvement of the optical and transport properties of BaSi_2 films. For this purpose, we have used photoresponsivity as a measure to find optimum growth conditions for light absorber layers. In recent years, the photoresponsivity of BaSi_2 films has increased drastically due to elevated substrate temperatures during thin-film growth [23], three-step growth [24], lightly B-/As-doping [25,26] and H-passivation [27–30] to passivate Si vacancies (V_{Si}), reaching $\sim 1 \text{ A/W}$ at 800 nm under a bias voltage of 0.1 V between the front surface and back surface electrodes.

However, even with such high-quality BaSi₂ light absorber layers, the η of BaSi₂-pn SCs was quite small ($\eta = 0.28\%$) [31], and the reason for this remains unclear. We therefore suppose that the extraction of photogenerated carriers is somehow hindered in the solar cell device structures. In fact, our previous research showed that the external quantum efficiency (*EQE*) in the short wavelength region is as low as 40% [32]. The structural design of solar cells is very important in achieving high η . For Si solar cells, devices with various structures such as heterojunction with intrinsic thin-layer (HIT) solar cells [33-36], tunnel oxide passivated contact (TOPCon) solar cells [37-41], and passivated emitter and rear cell (PERC) solar cells [42] have been studied for years and their transport mechanisms unveiled [43,44]. In this study, we firstly aimed to clarify the factors that reduce the η in BaSi₂-pn SCs by advanced opto-electrical modelling. Simulation results are an important tool for solar cell design, as they provide, as in this study, detailed insight in reflection and parasitic absorption losses as well as recombination mechanisms which degrade BaSi₂-pn SCs performance. We therefore proposed a new device structure, Al-doped n⁺-ZnO (AZO)/p-BaSi₂ HJSC. With this device structure we address both previous optical and electrical shortcomings and show a pathway for reaching efficiencies well above 20%. In the last part of our contribution, we report on the first experimental demonstration of n⁺-AZO/p-BaSi₂ HJSC.

2. METHODS

2.1 Simulation Method

In this study, we performed optical simulations using the GENPRO4 [45] software and electrical simulation using ASA [46,47] version 7 (ASA-7), both developed at the Delft University of Technology. The complex refractive index ($n + ik$) of each layer was measured independently as a function of wavelength (λ) using spectroscopic ellipsometry (see Fig. 2) and was used as an input. Note that the extinction coefficient k of both p-BaSi₂ and p⁺-BaSi₂ materials was set to zero for wavelengths longer than 950 nm. This complies with the experimental evidence of

absence of photoresponsivity signal in such materials for $\lambda > 950$ nm [48]. By integrating the resulting absorption spectra of BaSi₂ light absorber layers over the AM1.5 spectrum range, the related photocurrent density J_{ph} values were obtained.

Optically, we investigated BaSi₂-pn SCs structure: ITO (80 nm) / a-Si (3 nm) / n⁺-BaSi₂ (20 nm) / p-BaSi₂ (500 nm) / p⁺-BaSi₂ (20 nm) / Si substrate (500 μ m) / Ag (100 nm), which is the same as the layered structure used for the first demonstration of BaSi₂-pn SCs [31]. In this work, next to simulating flat interfaces, we also introduced the typical 3D pyramidal texture at front and/or back surfaces of the Si substrate to investigate the anti-reflection and light trapping effects. For this purpose, an atomic force microscopy image of a pyramid-type texture obtained by chemical-etching of a Si substrate was used (see Fig. 2(c)). In addition, to increase light absorption in BaSi₂ light absorber layers, we investigated n⁺-AZO/p-BaSi₂ HJSCs, in which the electron transport layer (ETL) was changed from n⁺-BaSi₂ to more transparent n⁺-AZO and the substrate was also eventually changed from Si to glass. The ZnO-based ETL was set as AZO (50 nm) / ZnO (20 nm).

For the electrical modelling, the generation rate provided by GENPRO4 was combined with ASA-7 software, which solves the semiconductor equations including bulk and surficial recombination mechanisms [49] as well as advanced transport mechanisms such as non-local tunneling currents [50], band-to-band tunneling [51] and trap-assisted tunneling [52,53]. suitable for both single- and multi-junction devices.

2.2 Experimental Method

An ion-pumped molecular beam epitaxy (MBE) system equipped with an electron-beam gun for high-purity (10N) Si and standard Knudsen cells for low-purity (3N) Ba and 3N-B was used. We used B-implanted p-Si(111) substrates (resistivity $\rho = 1 - 4$ Ω cm) with a dose of 2.5×10^{14} cm⁻². A projected range was set at 50 nm so that the hole concentration p would be approximately 2×10^{19} cm⁻³ on the surface after activation annealing at 900 °C for 3 min.

Before growing the films, the Si substrates were cleaned by the standard RCA procedure, followed by thermal cleaning at 900 °C for 30 min in an ultra-high vacuum chamber to remove the protective oxide layer from the surface. For epitaxial growth of BaSi₂ films, we evaporated Ba onto the heated Si substrate at a substrate temperature of $T_s = 500$ °C to form a 5-nm-thick BaSi₂ template layer by reactive deposition epitaxy (RDE). This layer acts as a seed for the growth of subsequent layers [54]. In the RDE process, the Ba deposition rate was fixed at 1 nm/min. Next, we formed a stack of heavily B-doped p⁺-BaSi₂ (20 nm) / lightly B-doped p-BaSi₂ (500 nm) by MBE. The p of these layers was 2.0×10^{19} and 1×10^{17} cm⁻³, respectively, at room temperature (RT). Finally, a 3-nm-thick amorphous Si (a-Si) layer was deposited *in-situ* on the surface at 180 °C. The a-Si layer prevents oxidation of the BaSi₂ surface [55] and does not hinder carrier transport across the a-Si/BaSi₂ interface [56-58]. After exposing the sample to air to load it into another chamber for sputtering, 50-nm-thick ZnO and 300-nm-thick AZO with a diameter of 1 mm were fabricated on the front surface at 150 °C by sputtering and Al electrodes were used on the back side. The electron concentration of ZnO and AZO were $n \sim 10^{15}$ and 10^{20} cm⁻³, respectively, at RT.

Solar cell performance was evaluated using a mask with a 1-mm-diameter hole under AM 1.5 conditions at 25 °C. Photoresponse and reflectance spectra were evaluated using a lock-in technique with a xenon lamp (150 W) and a single monochromator with a focal-length of 25 cm (Bunko Keiki SM-1700A and RU-60N). The light intensity was calibrated using a pyroelectric sensor (Melles Griot 13PEM001/J).

3. RESULTS AND DISCUSSION

3.1 Effect of textured surfaces on the photocurrent of BaSi₂-pn SCs

As shown in Fig. 3, four types of solar cell structures were simulated, namely, “Flat/Flat” structure with flat surfaces on both sides, “Texture/Flat” structure with texture only on the front side, “Flat/Texture” structure with texture only on the back side, and “Texture/Texture”

structure with both sides textured. By changing the front surface structure from flat to textured (see Fig. 3(b)), the reflectance reduced in the λ range 700 – 1200 nm compared to that in Fig. 3(a), and the J_{ph} increased by 1.2 mA/cm². On the other hand, the light trapping effect by the rear texture structure was not pronounced as shown in Fig. 3(c,d). This is probably because almost all the transmitted light was absorbed by the Si substrate before bouncing back to the p-BaSi₂ layers. In addition, it was found that the key factor inhibiting light coupling in the p-BaSi₂ layer was the parasitic absorption in the n⁺-BaSi₂ ETL. This parasitic absorption is unavoidable when BaSi₂ is used for ETL as the sunny side, because BaSi₂ has higher light absorption coefficients than CIGS.¹²

3.2 Simulation of n⁺-AZO/p-BaSi₂ HJSCs

Extensively studied TFSCs, e.g. CIGS, CdTe are a kind of HJSCs which consist of p-type absorber layers and n-type window layers. The wide E_g of the window layer allows more photons to reach the absorber layer. Another advantage of a heterojunction solar cell is that the recombination in the wide E_g window layer is lower than that of a homojunction solar cell. Based on these arguments, we proposed n⁺-AZO/p-BaSi₂ HJSCs in which the n⁺-BaSi₂ ETL was replaced with n⁺-AZO layers. Figure 4 shows the schematic and the band diagram of n⁺-AZO/p-BaSi₂ HJSCs.

Figures 5(a,b) show the absorption spectra of the n⁺-AZO/p-BaSi₂ HJSCs with (a) Flat/Flat and (b) Texture/Flat surfaces deployed on Si substrate. The parasitic absorption in the ETL was significantly reduced, and the J_{ph} increased by more than 10 mA/cm² compared with those of the BaSi₂-pn SCs in Fig. 3. The J_{ph} reached 30.23 mA/cm² using the front textured surface in Fig. 5(b). Next, in order to reduce the absorption in the Si substrate and to make light-trapping effect work well, we simulated n⁺-AZO/p-BaSi₂ HJSCs in which the Si substrate was replaced by a glass substrate. The formation of high-photoresponsivity BaSi₂ films on glass was already demonstrated [59]. A 100-nm thick ITO layer was used as a back contact on top of the

glass substrate and 100-nm-thick Ag was set as a reflection film on the rear side of the glass. Figures 5(c,d) show the absorption spectra of the n^+ -AZO/p-BaSi₂ HJSCs on glass. The J_{ph} is further increased by about 1 – 2 mA/cm² and reached a maximum of 32.06 mA/cm² in Fig. 5(d). We attribute this increase to the enhanced light absorption in the p-BaSi₂ layer owing to the light reflected back to the p-BaSi₂ layer because of the Ag film.

In Fig. 6 we summarize the J_{ph} as a function of p-BaSi₂ thickness (t_{p-BaSi_2}) in the range of 0.02 – 10 μ m. For BaSi₂-pn SCs, J_{ph} saturated when t_{p-BaSi_2} reached 3 μ m, resulting in $J_{ph} \sim 20$ mA/cm². By replacing n^+ -BaSi₂ ETL with n^+ -AZO ETL, the saturated J_{ph} increased drastically to more than 30 mA/cm². Furthermore, by changing the surface from flat/flat to texture/flat, it increased by more than 2 mA/cm². By replacing the Si substrate with glass, the saturated J_{ph} did not change, but the light absorption became more efficient for small t_{p-BaSi_2} (< 3 μ m). The t_{p-BaSi_2} necessary for J_{ph} to saturate was reduced to 1 μ m.

Next, using ASA-7, we performed electrical simulations of this novel solar cell configuration. We simulated the ideal case with no defect properties. Input electrical parameters were shown in Table 1. Figure 7 shows the current density versus voltage (J - V) characteristics of three types of BaSi₂ solar cells. The short-circuit current density (J_{SC}) of n^+ -AZO/p-BaSi₂ HJSCs is much larger than that of BaSi₂-pn SCs due to the more efficient light absorption. Furthermore, a large open circuit voltage (V_{OC}) well over 0.8 V results from n^+ -AZO/p-BaSi₂ HJSCs. We can therefore state that the novel n^+ -AZO/p-BaSi₂ HJSCs is a device structure in which the excellent optical properties of BaSi₂ as a light absorber layer can be fully utilized.

3.3 Experimental demonstration of n^+ -AZO/p-BaSi₂ HJSCs

Based on these simulated results, we fabricated the novel n^+ -AZO/p-BaSi₂ HJSCs. The device structure is the same as that shown in Fig. 4. Figure 8 shows the measured J - V characteristics under AM1.5 illumination and the internal quantum efficiency (IQE) spectrum for n^+ -AZO/p-BaSi₂ HJSCs. It shows $\eta = 0.04\%$, $J_{SC} = 3.7$ mA/cm², and $V_{OC} = 50$ mV. IQE exceeded 30% at

$\lambda = 600$ nm. This efficiency is almost the same as that previously obtained for BaSi₂-pn SCs [31]. This is the first demonstration of this novel n⁺-AZO/p-BaSi₂ HJSCs.

3.4 Investigation of the defect properties existing in the SCs

In Fig. 9(a) we compare the experimental and the simulated J - V characteristics, noting that the η of the fabricated solar cell is much smaller than that expected from the simulations. To explain the experimental results, we performed electrical simulation modelling different defect profiles. As density of states (DOS) distribution, we used conduction band tail (CBT), valence band tail (VBT), and localized states within the forbidden band gap, as shown in Fig. 9(b). The standard model of the DOS distribution consists of a parabolic conduction band (CB) and a parabolic valence band (VB), an exponentially decaying CBT and VBT. Localized states in the forbidden band gap consist of a donor-like state, the so-called DB⁺⁰, and an acceptor-like state, so-called DB⁰⁻, and are therefore represented by two energy levels E^{+0} and E^{0-} in the band diagram, respectively [60]. These energy levels are called the transition energy levels. The transition energy levels are separated from each other by a correlation energy U_{corr} .

In these electrical simulations, we first attempted to find the key parameters significantly affecting the solar cell performance and then we fine-tuned the values. We assumed that the i-ZnO layers would electrically contact the p-BaSi₂ layers despite the presence of the a-Si layer in between. This is because the a-Si layer is very thin and is likely to be damaged during the sputtering of i-ZnO layers. Therefore, the values of E_g and electron affinity ($q\chi$) of a-Si was set to be the same as those of BaSi₂. The parameters used for each layer which reproduced well the experimental result are summarized in Table 2. The most influential parameter of all was found to be the localized state density in the p-BaSi₂ light absorber layer ($N_{\text{T,BaSi}_2}$). Figure 10(a-f) shows the dependence of $N_{\text{T,BaSi}_2}$ on the J - V characteristic, DOS, J_{sc} , V_{oc} , η , and fill factor (FF), respectively. The defect level in the p-BaSi₂ layer is located at 0.55 eV from the CBM as shown in Fig. 10(b). As the $N_{\text{T,BaSi}_2}$ was increased from 10^{12} to 10^{18} cm⁻³,

the J_{SC} , V_{OC} , and η started to sharply decrease especially when $N_{T,BaSi_2} > 10^{13} \text{ cm}^{-3}$. Properties of defects in BaSi₂ have been investigated by various methods such as Raman spectroscopy, photoluminescence, deep-level transient spectroscopy, positron annihilation spectroscopy, and electron paramagnetic resonance [23,61–65]. It was reported that most of the defects are related to V_{Si} . A $N_{T,BaSi_2}$ value of $8.8 \times 10^{16} \text{ cm}^{-3}$ which is reproduced well the experimental result as shown in Fig. 10(b), is much higher than that obtained by experiment in case of BaSi₂ epitaxial films on Si(111)($\sim 10^{16} \text{ cm}^{-3}$) [23]. In the solar cell structure, however, the lightly B-doped p-BaSi₂ absorber layer was grown on the defective heavily B-doped p⁺-BaSi₂ layer. Therefore, the $N_{T,BaSi_2}$ in lower quality p-BaSi₂ absorber layer is likely to increase compared to that directly grown on a Si(111) substrate. In our previous work [66], a few tens of nanometer diameter B clusters were found to form in heavily B-doped p-BaSi₂ films, and therefore defects such as dislocations due to B clusters are present in the p-BaSi₂ absorber layers. Such a deep defect level can thus be attributed to the defects coming from B clusters in the heavily B-doped p⁺-BaSi₂ underlayer. We therefore speculate that reducing the B concentration in the p⁺-BaSi₂ layer might be effective to reduce the defect density in the p-BaSi₂ absorber layer. Regarding the V_{OC} , however, a small value of $V_{OC} = 0.05 \text{ V}$ cannot be explained by $N_{T,BaSi_2}$ alone. As shown in Fig. 10(a), the reduction of V_{OC} was limited down to 0.6 V even at $N_{T,BaSi_2} = 10^{18} \text{ cm}^{-3}$, which is still far from the experimentally obtained V_{OC} . We thus consider another factor, the shunt resistance (R_{SH}). Figure 11(a) shows the depth profiles of electron and hole concentrations under illumination in n⁺-AZO/p-BaSi₂ HJCS with $N_{T,BaSi_2} = 8.8 \times 10^{16} \text{ cm}^{-3}$ at a forward bias voltage (V_{bias}) of 0.7 V, close to V_{OC} . The black-colored J - V plot in Fig. 11(b) corresponds to this solar cell. Due to large CB and VB discontinuities at the i-ZnO/p-BaSi₂ interface, electrons and holes accumulate at the interface, leading to the recombination of these carriers. To reflect this effect, we introduced R_{SH} in the equivalent circuit (Fig. 11(b)), and set R_{SH} to be in the range between 10^3 and $10^5 \Omega$. V_{OC} decreased with decreasing R_{SH} as shown in Fig. 11(b) and approached the experimental result. Finally, we summarize the J - V characteristics of n⁺-AZO/p-BaSi₂ HJSCs

with defect properties in Fig. 12(a). All the parameters are summarized in Table 2. By reflecting the introduced defect characteristics, the simulated J - V characteristics perfectly matched the experimental result in Fig. 12(a). The complete DOS distribution of the p-BaSi₂ absorber layer is shown in Fig. 12(b).

It is noted that we need to suppress the carrier recombination to achieve a larger V_{OC} as discussed above. Theoretical analysis of the effect of band offsets at window/absorber interfaces has been made on CIS solar cells [67]. Band offsets are caused by the difference in $q\chi$ between the BaSi₂ absorber and the window layer. The point is to insert an interlayer to separate electrons from holes spatially. We therefore turn our attention to materials such as Zn_{1-x}Ge_xO [68] whose $q\chi$ is close to that of BaSi₂ (3.2 eV) [69], but still tunable, likewise E_g , from 3.3 to 5.9 eV and 2.5 to 4.3 eV, respectively, by changing the Ge content x . Figures 13(a-c) and (a'-c') show the band diagrams and carrier concentration profiles of n⁺-AZO($q\chi = 4.3$ eV)/i-ZnO($q\chi = 4.3$ eV)/p-BaSi₂($q\chi = 3.2$ eV) and n⁺-AZO($q\chi = 4.3$ eV)/i-Zn_{1-x}Ge_xO($q\chi = 3.2$ eV)/p-BaSi₂($q\chi = 3.2$ eV) HJSCs, respectively, under illumination. The band diagrams were evaluated at $V_{bias} = 0$ V and 0.7 V, with the latter value close to that of the V_{OC} . In the case of ZnO interlayer, the accumulation of electrons and holes occurs at the i-ZnO/p-BaSi₂ interface (see Fig. 13(c)). This is caused by the difference in $q\chi$ between ZnO and BaSi₂. In contrast, such carrier accumulation is completely suppressed by inserting the Zn_{1-x}Ge_xO ($q\chi = 3.2$ eV) interlayers in Fig. 13(c'). In Table 3, we summarize the J_{SC} , V_{OC} , FF , and η of BaSi₂-pn SCs and BaSi₂ HJSCs with/without Zn_{1-x}Ge_xO interlayers. Such effect has actually been demonstrated in CIS solar cells. Insertion of a CdS interlayer at the interface of n⁺-AZO/p-CIS solar cells resulted in spatial separation of photogenerated carriers and improved the conversion efficiency [70]. On the basis of these discussions, we can state that the insertion of interlayers with their $q\chi$'s close to that of BaSi₂ between n⁺-AZO and p-BaSi₂ is a very effective means to decrease the carrier recombination and thus to improve V_{OC} .

4. CONCLUSION

We investigated device structures of BaSi₂ solar cells by advanced opto-electrical device modelling by means of GENPRO4 and ASA-7 software. The absorption spectra of BaSi₂-pn SCs on Si substrates showed that the front pyramid texture was effective in increasing the J_{ph} of the 500-nm-thick p-BaSi₂ absorber layer by 1.2 mA/cm². On the other hand, the rear-texture did not yield additional optical performance. This is mainly because the Si substrate absorbs the transmitted light. Moreover, we found that the key factor inhibiting light absorption in the p-BaSi₂ layer was parasitic absorption in the 20-nm-thick n⁺-BaSi₂ ETL. To prevent the parasitic absorption, we proposed a new device structure, n⁺-AZO/p-BaSi₂ HJSCs. By using a wide band gap ETL, the parasitic absorption reduced drastically, and the J_{ph} reached 30.23 mA/cm². Furthermore, by replacing Si substrate with glass substrate, the required absorber layer thickness was reduced to 1 μm. Based on these simulated results, we demonstrated the first operation of n⁺-AZO/p-BaSi₂ HJSCs experimentally. The J - V characteristics showed $\eta = 0.04\%$, $J_{SC} = 3.7$ mA/cm², and $V_{OC} = 50$ mV. IQE exceeded 30% at $\lambda = 600$ nm. The defects-based electrical model well reproduced the experimentally obtained J - V characteristics. According to the simulation results, the defect level of the p-BaSi₂ absorber layer was located at 0.55 eV from the CBM and its density was 8.8×10^{16} cm⁻³, probably caused by the defective p⁺-BaSi₂ underlayer. We also found that the carriers recombination at the heterointerface caused by carrier accumulation due to large band offsets decreased the V_{OC} significantly. An interlayer material such as Zn_{1-x}Ge_xO with its $q\chi$ close to that of BaSi₂ placed in between the n⁺-AZO and the p-BaSi₂ suppresses such carrier accumulation, drastically improving the V_{OC} .

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Table. 1: Input electrical parameters of each layer used for ideal simulation.

	n ⁺ -AZO	ZnO	a-Si	n-BaSi ₂	p-BaSi ₂	p ⁺ -BaSi ₂
E_g [eV]	3.3	3.3	1.3	1.3	1.3	1.3
$q\chi$ [eV]	3.5	3.5	3.2	3.2	3.2	3.2
N_c [cm ⁻³]	3.0×10^{18}	3.0×10^{18}	2.0×10^{20}	2.6×10^{19}	2.6×10^{19}	2.6×10^{19}
N_v [cm ⁻³]	1.7×10^{19}	1.7×10^{19}	2.0×10^{20}	2.0×10^{19}	2.0×10^{19}	2.0×10^{19}
ϵ	9	9	11.9	14	14	14
n/p [cm ⁻³]	1.0×10^{20}	1.0×10^{17}	9.7×10^{15}	1.0×10^{19}	1.0×10^{17}	2.0×10^{19}
μ_e [cm ² V ⁻¹ s ⁻¹]	50	50	5	500	800	500
μ_h [cm ² V ⁻¹ s ⁻¹]	31	31	1	30	100	30

Table 2: Input electrical and defective parameters used for each layer which reproduced well the experimental result.

	n ⁺ -AZO	ZnO	a-Si	p-BaSi ₂	p ⁺ -BaSi ₂
E_g [eV]	3.3	3.3	1.3	1.3	1.3
$q\chi$ [eV]	3.5	3.5	3.2	3.2	3.2
N_C [cm ⁻³]	3.0×10^{18}	3.0×10^{18}	2.0×10^{20}	2.6×10^{19}	2.6×10^{19}
N_V [cm ⁻³]	1.7×10^{19}	1.7×10^{19}	2.0×10^{20}	2.0×10^{19}	2.0×10^{19}
ϵ	9	9	11.9	14	14
n/p [cm ⁻³]	1.0×10^{17}	1.0×10^{17}	9.7×10^{15}	8.0×10^{16}	1.0×10^{18}
μ_e [cm ² V ⁻¹ s ⁻¹]	50	50	5	800	500
μ_h [cm ² V ⁻¹ s ⁻¹]	31	31	1	77	30
Hole capture rate in VBT [m ⁻³ s ⁻¹]	1.0×10^{-19}	1.0×10^{-19}	1.0×10^{-17}	1.0×10^{-15}	1.0×10^{-15}
Electron capture rate in VBT [m ⁻³ s ⁻¹]	5.0×10^{-17}	5.0×10^{-17}	1.0×10^{-15}	1.0×10^{-15}	1.0×10^{-15}
DOS in VBT [m ⁻³ eV ⁻¹]	1.0×10^{25}	1.0×10^{25}	1.0×10^{25}	1.0×10^{25}	1.2×10^{24}
Slope of the VBT [eV]	0.01	0.01	0.044	0.02	0.1
Hole capture rate in CBT [m ⁻³ s ⁻¹]	1.0×10^{-19}	1.0×10^{-19}	1.0×10^{-17}	1.0×10^{-15}	1.0×10^{-15}
Electron capture rate in CBT [m ⁻³ s ⁻¹]	1.0×10^{-19}	1.0×10^{-19}	1.0×10^{-15}	1.0×10^{-15}	1.0×10^{-15}
DOS in CBT [m ⁻³ eV ⁻¹]	1.0×10^{25}	1.0×10^{25}	1.0×10^{25}	1.0×10^{25}	1.2×10^{24}
Slope of the CBT [eV]	0.01	0.01	0.03	0.02	0.1
U_{corr} [eV]	0.2	0.2	0	0.1	0
Electron capture rate of neutral DB [m ⁻³ s ⁻¹]	1.0×10^{-17}	1.0×10^{-17}	1.0×10^{-10}	1.0×10^{-11}	1.0×10^{-11}
Electron capture rate of positive DB [m ⁻³ s ⁻¹]	1.0×10^{-17}	1.0×10^{-17}	1.0×10^{-10}	1.0×10^{-11}	1.0×10^{-11}
Hole capture rate of neutral DB [m ⁻³ s ⁻¹]	1.0×10^{-17}	1.0×10^{-17}	1.0×10^{-10}	1.0×10^{-11}	1.0×10^{-11}
Hole capture rate of negative DB [m ⁻³ s ⁻¹]	1.0×10^{-17}	1.0×10^{-17}	1.0×10^{-10}	1.0×10^{-11}	1.0×10^{-11}
Defect density N_T [cm ⁻³]	1.0×10^{17}	1.0×10^{15}	1.0×10^{14}	8.8×10^{16}	3.5×10^{17}
Energy relative to midgap [eV]	0.1	0.1	0.0	0.1	-0.2

Table 3: Progress in the modelling-based optimization of external parameters of different solar cells based on p-BaSi₂ absorber material. *Ultimate* stands for n⁺-Zn_{1-x}Ge_xO/i-Zn_{1-x}Ge_xO/p-BaSi₂ HJSC on glass.

	BaSi ₂ -pn on Si	n ⁺ -AZO/p-BaSi ₂ on Si		n ⁺ -AZO/p-BaSi ₂ on Glass		<i>Ultimate</i>
	Flat/Flat	Flat/Flat	Tex./Flat	Flat/Flat	Tex./Flat	Tex./Flat
d_{BaSi_2} [μm]	0.5	0.5	0.5	0.5	0.5	0.5
J_{SC} [mA/cm^2]	11.2	27.4	30.5	27.9	30.5	32.9
V_{OC} [V]	0.80	0.84	0.84	0.77	0.77	0.84
FF [-]	0.85	0.85	0.86	0.78	0.80	0.86
η [%]	7.6	19.6	22.0	16.7	18.9	23.9

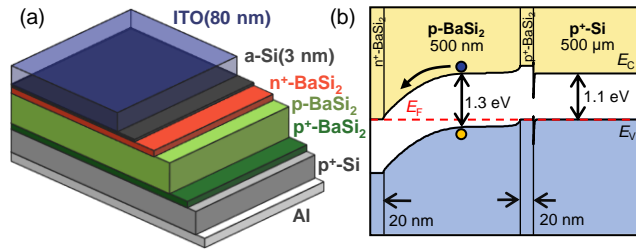


FIGURE 1 (a) Schematic image and (b) band alignment of BaSi₂-pn homojunction solar cells simulated by ASA-7. p-BaSi₂ is used as a light absorber, and photogenerated electrons flow to the n⁺-BaSi₂ side.

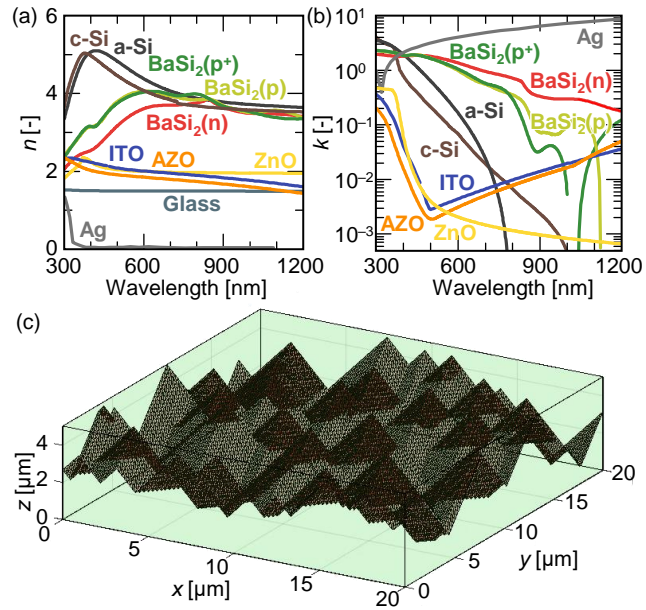


FIGURE 2 (a) Refractive index $n(\lambda)$ and (b) extinction coefficient $k(\lambda)$ of layers used for optical simulations. These data were measured by spectroscopic ellipsometry. (c) 3D pyramidal texture measured by AFM.

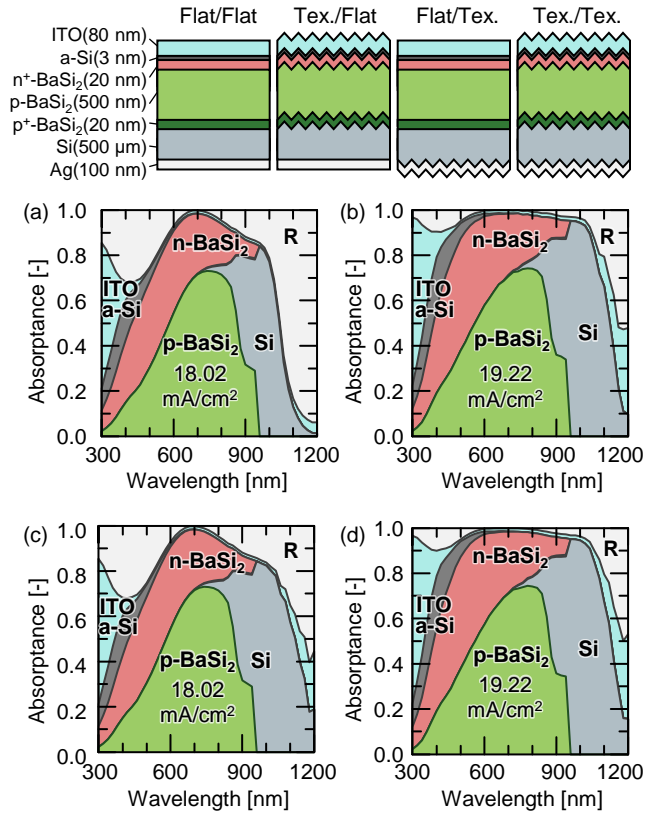


FIGURE 3 Reflection and absorption spectra of BaSi₂ homojunction solar cells with (a) Flat/Flat, (b) Texture/Flat, (c) Flat/Texture, (d) Texture/Texture surfaces. The photocurrent density J_{ph} generated in the p-BaSi₂ absorber layer is shown.

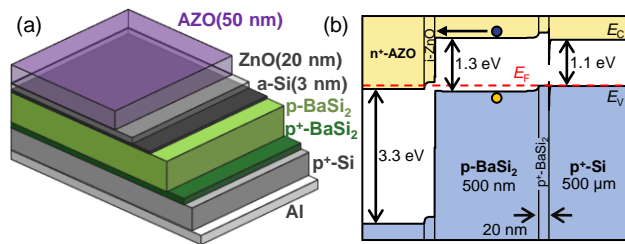


FIGURE 4 (a) Schematic and (b) band diagram of an n⁺-AZO/p-BaSi₂ heterojunction solar cell in equilibrium.

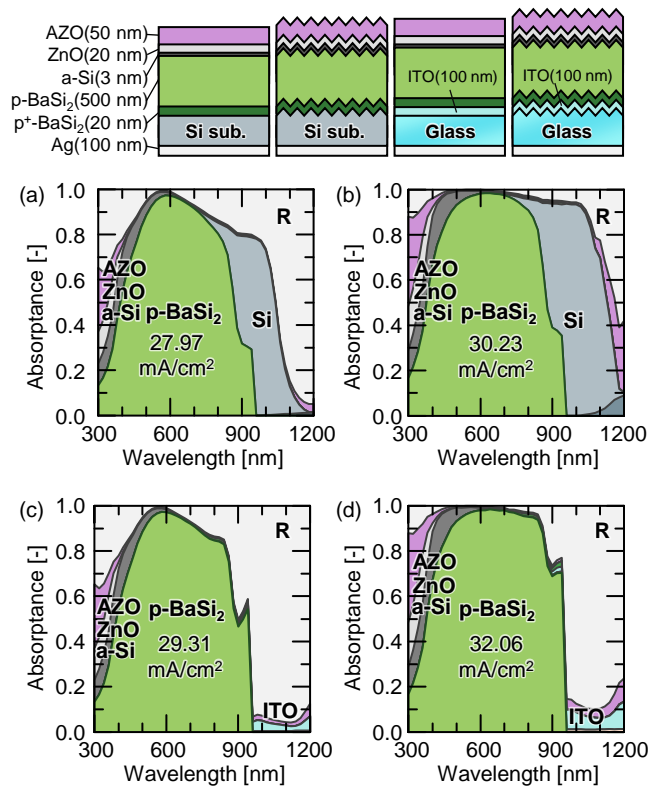


FIGURE 5 Absorption spectra of AZO/BaSi₂ heterojunction solar cells with (a)-(c) Flat/Flat and (b)-(d) Texture/Texture surfaces on Si or glass substrates. The photocurrent density J_{ph} generated in the p-BaSi₂ absorber layer is shown.

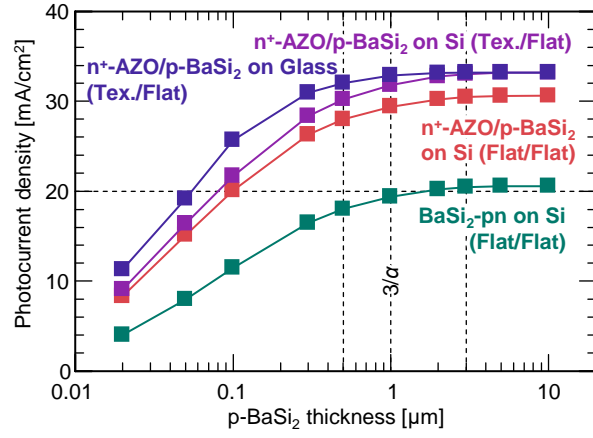


FIGURE 6 Dependence of J_{ph} on the thickness of p-BaSi₂ light absorber layer in four types of BaSi₂ solar cells (on Si substrate: flat BaSi₂-pn, flat n⁺-AZO/p-BaSi₂, textured n⁺-AZO/p-BaSi₂; on glass substrate: textured n⁺-AZO/p-BaSi₂).

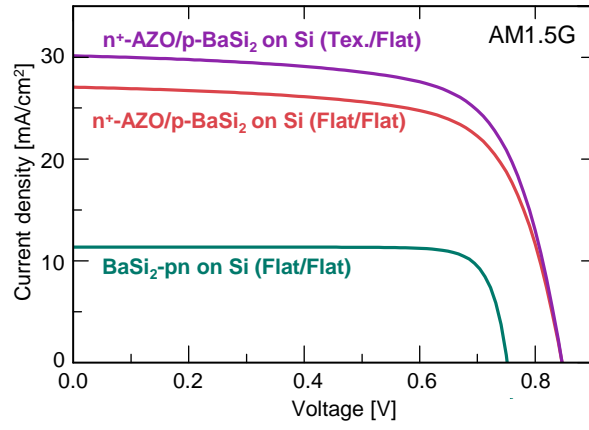


FIGURE 7 (a) J - V characteristics of three types of BaSi_2 solar cells (on Si substrate: flat BaSi_2 -pn, flat $\text{n}^+\text{-AZO/p-BaSi}_2$, textured $\text{n}^+\text{-AZO/p-BaSi}_2$).

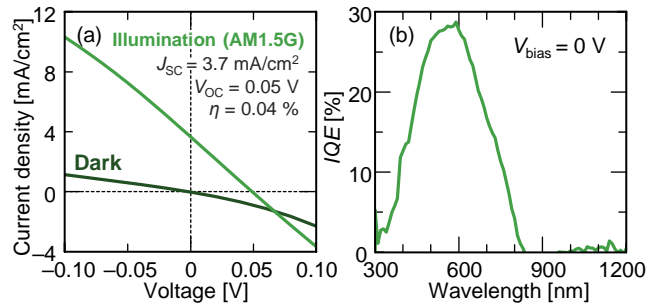


FIGURE 8 (a) J - V characteristics and (b) IQE spectrum of n^+ -AZO/ p -BaSi₂ HJSCs. The solar cell operation was demonstrated for the first time.

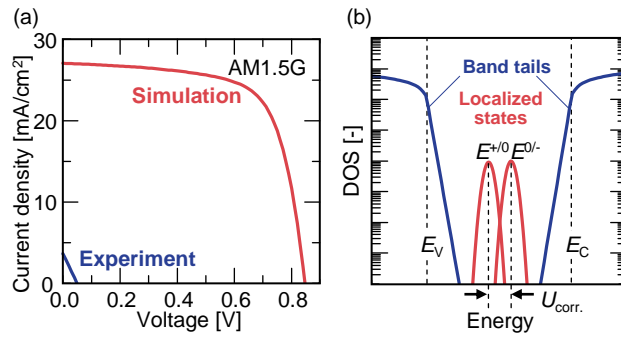


FIGURE 9 (a) Comparison of J - V characteristics between experimental and simulated flat n^+ -AZO/ p -BaSi₂ HJSCs; (b) Density of States distribution in p -BaSi₂ absorber.

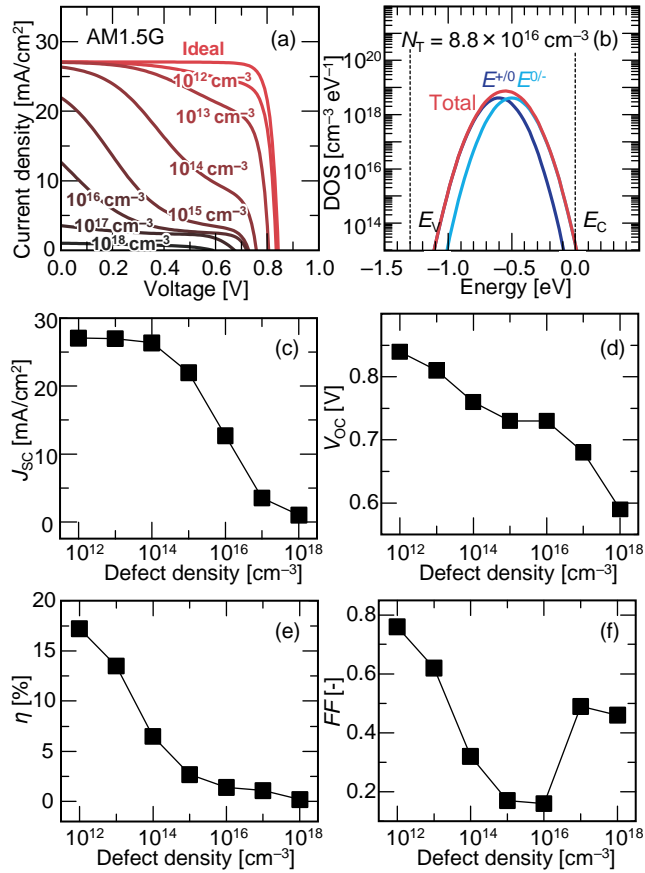


FIGURE 10 The effect of the defect density N_T in p-BaSi₂ light absorber. (a) J - V characteristics of n⁺-AZO/p-BaSi₂ HJSCs as function of defect level in p-BaSi₂ absorber and (b) DOS distribution; (c-f) dependence of the defect density in p-BaSi₂ on J_{sc} , V_{oc} , η , and fill factor (FF), respectively.

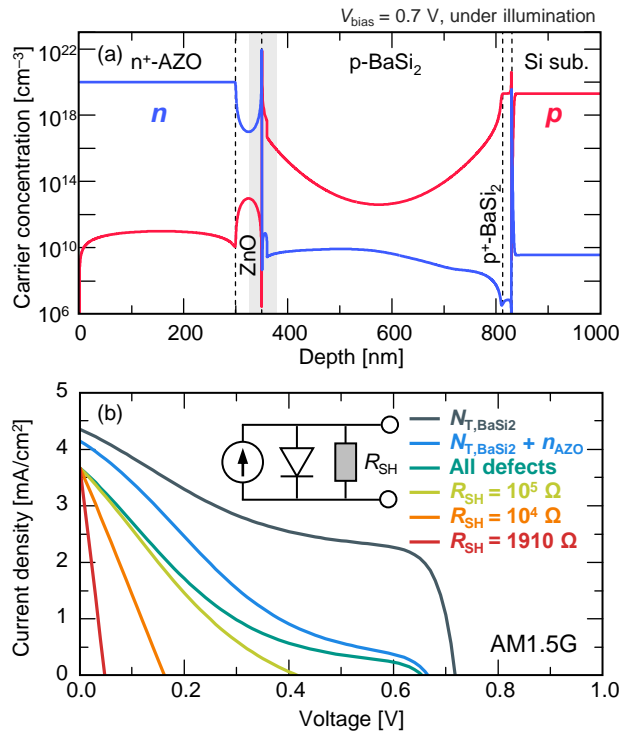


FIGURE 11 (a) The depth profile of carrier concentration in n^+ -AZO/p-BaSi₂ HJSCs under a forward bias voltage of 0.7 V under illumination and (b) calculated J - V characteristics at $N_{\text{T,BaSi}_2} = 8.8 \times 10^{16} \text{ cm}^{-3}$ and $n_{\text{AZO}} = 10^{17} \text{ cm}^{-3}$. By introducing shunt resistance, J - V profiles fitted to that obtained in experiments.

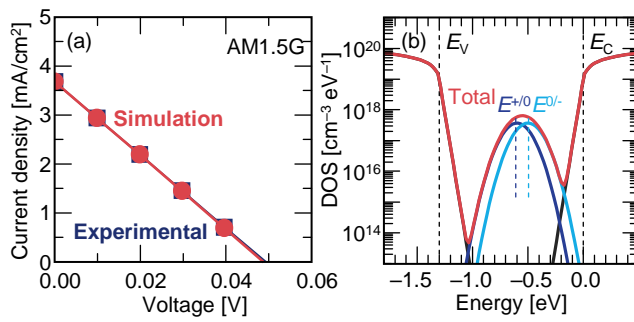


FIGURE 12 (a) J - V characteristics of experimental and simulated flat n^+ -AZO/p-BaSi₂ HJSCs and (b) profiles of tailored DOS and defect levels to reproduce the experimental J - V characteristics in (a).

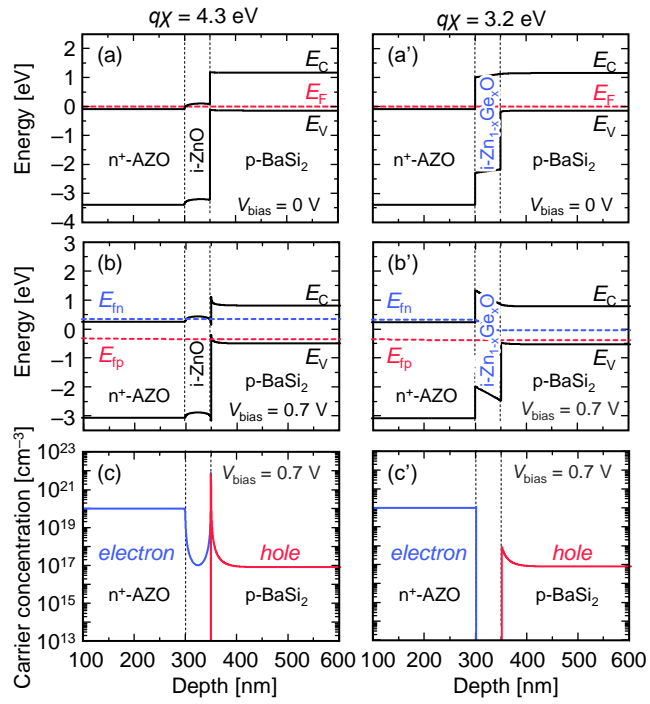


FIGURE 13 Band diagrams of n^+ -AZO/ p -BaSi₂ HJSCs with an inter layer of $q\chi =$ (a,b) 4.3 and (a',b') 3.2 eV at $V_{\text{bias}} = 0$ and 0.7 V under illumination. (c,c') Depth profiles of electron and hole concentrations at $V_{\text{bias}} = 0.7$ V .