A 3-in-1 doping process for interdigitated back contact solar cells exploiting the understanding of co-diffused dopant profiles by use of PECVD borosilicate glass in a phosphorus diffusion

Sebastian Gloger*, Axel Herguth, Josh Engelhardt, Giso Hahn and Barbara Terheiden

University of Konstanz, Department of Physics, Universitätsstr. 10, Konstanz, 78457, Germany

ABSTRACT

Boron and phosphorus doping of crystalline silicon using a borosilicate glass (BSG) layer from plasma enhanced chemical vapor deposition (PECVD) and phosphorus oxychloride diffusion, respectively, is investigated. More specifically, the simultaneous and interacting diffusion of both elements through the BSG layer into the silicon substrate is characterized in depth. We show that an overlying BSG layer does not prevent the formation of a phosphorus emitter in silicon substrates during phosphorus diffusion. In fact, a BSG layer can even enhance the uptake of phosphorus into a silicon substrate compared with a bare substrate.

From the understanding of the joint diffusion of boron and phosphorus through a BSG layer into a silicon substrate, a model is developed to illustrate the correlation of the concentration dependent diffusivities and the emerging diffusion profiles of boron and phosphorus. Here, the in diffusion of the dopants during diverse doping processes is reproduced by the use of known concentration dependences of the diffusivities in an integrated model. The simulated processes include a BSG drive in step in an inert and in a phosphorus containing atmosphere.

Based on these findings, a PECVD BSG/capping layer structure is developed, which forms three different n++, n+ and p+ doped regions during one single high temperature process. Such engineered structure can be used to produce back contact solar cells. Copyright © 2016 John Wiley & Sons, Ltd.

KEYWORDS
doping; co diffusion; boron; phosphorus; silicon; solar cell; borosilicate glass (BSG); interdigitated back contact (IBC)

*Correspondence
Sebastian Gloger, University of Konstanz, Department of Physics, Universitätsstr. 10, Konstanz, 78457, Germany.
E mail: sebastian.gloger@uni.konstanz.de

1. INTRODUCTION

About 90% of solar cells manufactured in 2014 were based on crystalline silicon [1,2]. Eighty percent of these cells use aluminum back surface fields (BSF [3]) [4]. However, their efficiency is limited to less than 20% because of the mediocre internal reflection and surface passivation of the aluminum contact [5]. The efficiency of solar cells can be increased up to 25% with back contact solar cell concepts [6] omitting the trade off between low shading and resistance losses. To realize such concepts, differently doped regions are needed on the rear side of a solar cell. These regions can conveniently be created in separate steps of deposition and in diffusion of dopants using plasma enhanced chemical vapor deposition (PECVD) silicate glass or other standard doping processes.

Recent research has focused on the formation process of PECVD silicate glass emitters [7-9], monitoring the quality of these PECVD emitters [10], co diffusion processes with PECVD emitters [11,12], contacting of these emitters [13,14], passivation through the same BSG layer [15], and the fabrication of solar cells using PECVD silicate glass [7,9,12,14,16,19]. Solar cell efficiencies of above 20% have been achieved with PECVD silicate glass processes for n type small area back junction IBC cell concepts [16], large area back junction [18,19], and front junction bifacial solar cell concepts [14]. Evidence for interacting diffusion phenomena have also been reported using a BSG layer and phosphorus oxychloride (POCl3) diffusion [15,17].

The objective of this study is therefore to reveal the interacting diffusion phenomena of boron and phosphorus...
in crystalline silicon. The co diffusion processes of boron and phosphorus through a BSG layer are investigated experimentally, and a model allowing for the prediction of co diffused dopant profiles is presented. Furthermore, we introduce an industrially feasible process to fabricate three differently doped regions, phosphorus $n^+$, $n^+$, and boron $p^+$ doped, using one single high temperature step for highly efficient interdigitated back contact silicon solar cells.

2. MODEL OF THE DIFFUSION OF DOPANTS IN SILICON

As a basis for the studies on dopant diffusion in crystalline silicon, using borosilicate glass, a model of the diffusion of dopants is introduced. It is developed as an integrated model to simulate the isolated in diffusion of boron and of phosphorus as well as the joint and interacting in diffusion of boron and phosphorus through a borosilicate layer into a silicon substrate.

2.1. Theory of diffusion (Fick’s law)

The diffusion of dopants like phosphorus (P) and boron (B) in silicon can be described by Fick’s first law of diffusion

$$ j_p = D_p \text{grad} \ c_p, \quad j_B = D_B \text{grad} \ c_B. \quad (1) $$

Here $D$ denotes the diffusivity and $j$ the particle current density of the respective dopant, which is opposing the gradient of its concentration $c$. The dopants are preserved according to the continuity equation

$$ \frac{\partial c_p}{\partial t} = \text{div} \ j_p, \quad \frac{\partial c_B}{\partial t} = \text{div} \ j_B. \quad (2) $$

In the case of boron, a source term accounts for a boron reservoir $r_B$ in the BSG layer.

The diffusivity of the dopants depends on the temperature, following an Arrhenius dependence as seen later on in the results section.

2.2. Determination of the diffusivity by Boltzmann–Matano analysis

The diffusivity $D$ of a dopant can be determined by means of Boltzmann Matano analysis from a diffused depth profile $c(x)$ of the dopant [20]. Therefore, the diffusivity of the dopant may depend exclusively on its concentration. To determine $D(c)$ the diffusion equation is rearranged [21–23].

The diffusivity $D$ of a specific concentration $c^*$ equals the inverse function $x(c)$ of the doping profile $c(x)$, integrated from the substrate concentration $c_0$ to the concentration $c^*$, multiplied with its negative differential at the position $c^*$, and divided by twice the diffusion duration $t$ [20],

$$ D(c^*) = \frac{1}{2t} \left( \frac{dx}{dc} \right)_c c^* \int_{c_0}^{c^*} x \ dx. \quad (3) $$

2.3. Simulation of diffusion processes

Solar cells and their manufacturing processes can be simulated using software like Sentaurus [24,25]. The diffusion processes in this study can already be described by a simpler model. Therefore, the diffusion and continuity equations of the dopants will be solved directly for the conditions as relevant within these investigations. In this way the effects to be investigated can be deduced clearly.

For the simulation of the in diffusion of boron and phosphorus only the diffusion of dopants is described with concentration dependent diffusivities in silicon. Diffusion of point defects and their interaction with dopants is not modeled independently. Furthermore, no interface effects like an oxidation of the silicon substrate and effects linked to it like the generation of point defects or segregation, which may lead to a formation of precipitates, are taken into account. Also the diffusion in the borosilicate glass layer is described by a constant diffusivity only, and the uptake of phosphorus from the diffusion furnace atmosphere is not modeled. Despite this simplification of the diffusion process, the model describes well the depth profiles and the observed effects of mutual decrease and enhancement of the diffusion of dopants. Diverse diffusion processes can be predicted with it.

To simulate the diffusion of dopants into a silicon substrate, the partial differential equations (1) and (2) are solved in one dimension for the conditions of the respective investigation. This is carried out numerically with the software FlexPDE [26] using the finite element method [27]. The following three processes are simulated with one collective model:

POCl$_3$ diffusion: POCl$_3$ diffusion is simulated as diffusion of phosphorus into a silicon substrate. Here a constant concentration of phosphorus is adopted at the surface of the substrate. There is no boron.

BSG diffusion in $N_2$ atmosphere: BSG diffusion in an inert nitrogen atmosphere is simulated as diffusion of boron from an overlying BSG layer into the silicon bulk. Here a certain concentration of boron in the BSG is used, which defines the surface concentration of boron in the silicon substrate. There is no phosphorus.

BSG diffusion in POCl$_3$ diffusion furnace: BSG diffusion in a POCl$_3$ diffusion furnace is simulated as concurrent diffusion of boron and phosphorus. Boron diffuses from a BSG layer into a silicon substrate while phosphorus diffuses through the borosilicate glass into the silicon substrate.
The change of the diffusivity of boron and phosphorus in crystalline silicon because of the respective other dopant [28–31] is taken into account.

The modeled sample structure consists of a silicon substrate with an overlying BSG of thickness $d_{BSG}$, see Figure 1. In the case of POCl$_3$ diffusion, the BSG thickness is assumed as zero. The two layers distinguish by different concentrations and diffusivities of the dopants. The formation of a phosphosilicate glass (PSG) [32,33] and the transport of phosphorus onto the substrate are not modeled separately. Instead the surface concentration of phosphorus is used as a parameter to describe the amount of phosphorus, which is available at the surface of the sample.

2.4. Diffusion of substitutional dopants

Dopants of group III and V of the periodic table, like boron and phosphorus, form covalent bonds with their surrounding silicon lattice atoms [34], and are almost exclusively built in substitutionally [21,35]. They diffuse mainly because of interaction with the two intrinsic point defects, vacancies, and interstitial silicon atoms [21,34,35]. Because their diffusion takes place through mobile dopant defect pairs, the diffusivity $D_A$ of a dopant depends on both, the concentration of the intrinsic point defect that governs its diffusion and the concentration of the dopant itself [22].

From the concentration $c_X$ of a point defect X in charge state $m$, the concentration of dopant defect pairs $c_{AX} = c_X$ can be developed. Using the sum of the current densities of the diffusing dopant defect pairs $D_{AX}$, the diffusivity $D_A$ of the dopant emerges as [34,35]

$$D_A = h \sum_m D_{AX} \cdot \left(\frac{n}{n_i}\right)^m$$

(4)

with the intrinsic diffusivities of the diffusing species $D_{AX} \cdot \nu_i$ and the field enhancement factor $h$.

2.5. Parameterization of the diffusivity of phosphorus

Under intrinsic conditions ($n < n_i$), the diffusion of phosphorus in silicon takes place through neutral silicon interstitials [22,35–37] and for high phosphorus concentrations through double negatively charged vacancies [22,36,38]. So for the diffusivity of phosphorus $D_p$ a simple model is used, which assumes a different diffusivity in the kink and in the tail region and whose parameters are chosen based on the phosphorus profiles shown in Figure 2. The diffusivity of phosphorus is depicted in Figure 3. For its parameterization according to equation (4) and [38] the following relations are used as presented in [39] ($c_{B}$: boron concentration; $c_{Pa(S)}$: electrically active phosphorus (surface) concentration; $c_{Peff}$: effective phosphorus net doping; $n_P$ constant):

$$D_{PK} = D_{PK0} \left(1 + \left(\frac{c_{Peff}}{n_P}\right)^2\right),$$

(5)

Diffusivity of phosphorus in kink region

$$D_{PT} = D_{PT0} \left(1 + \left(\frac{c_{PaS}}{n_P}\right)^3\right),$$

(6)

Diffusivity of phosphorus in tail region

$$D_{TE} = \left(D_{PT} - D_{PK}\right) \frac{1}{\exp\left(\frac{c_{PaS}}{n_P}\right) + 1}.$$

(7)

Diffusivity enhancement in tail region

![Figure 1](image1.png)

**Figure 1.** Sketch of the model of a BSG diffusion step in a POCl$_3$ diffusion furnace. Boron and phosphorus diffuse simultaneously into a silicon substrate.

![Figure 2](image2.png)

**Figure 2.** Phosphorus depth profiles experimentally produced by a POCl$_3$ diffusion (ECV measurement, symbols) and simulated according to section 2 (lines). The electrically active concentration of phosphorus $c_{Pa}$ is plotted against the depth in the substrate $x$ for a diffusion temperature $T$ of 806, 840, and 891 °C.
Furthermore, for a high concentration of effective boron net doping $c_{\text{Beff}} = c_{\text{B}}$, in p type silicon a proportional decrease of the diffusivity according to $(1 + c_{\text{Beff}}/n_i)^{-1}$ is taken into account [30,31], which is shown in Figure 4. So the overall diffusivity of phosphorus emerges as

$$D_P = (D_{PK} + D_{TE}(0)) \frac{1}{1 + \frac{c_{\text{Beff}}}{n_i}}.$$

The process temperature exceeds the glass transition and flow temperatures of borosilicate glass, which lie at about 600 °C [40] and 800 to 850 °C [41], respectively. Therefore, the diffusivity of phosphorus in the BSG layer $D_{\text{BSG}}$ is modeled with a higher value than in the crystalline silicon substrate. Values of $4 \times 10^{-10}$ to $3 \times 10^{-12}$ cm$^2$/s arise for it from the experimentally determined doping profiles.

### 2.6. Parameterization of the diffusivity of boron

Boron diffuses in silicon together with neutral and positively charged silicon interstitials [34–36,42,43]. For the diffusivity of boron also a simple model is used. Its parameters are chosen based on the boron profiles shown in Figure 5. The diffusivity of boron is parameterized after equation (4) by the following relation according to [39] and shown in Figure 6.

$$D_{\text{Beff}} = D_B \left(1 - \frac{c_{\text{Beff}}}{n_i}\right)$$

Extrinsic diffusivity of boron (9)

For a high concentration of effective phosphorus net doping $c_{\text{Peff}} = c_{\text{P}}$, in n type silicon a proportional decrease of the diffusivity according to $(1 + c_{\text{Peff}}/n_i)^{-1}$ is taken into account [28,29], see Figure 4. So the overall diffusivity of boron emerges as

$$D_B = (D_{\text{Beff}} + D_{\text{TE}}(0)) \frac{1}{1 + \frac{n_i}{c_{\text{Peff}}}}.$$

The diffusivity of boron in the BSG layer is modeled with the same value as the diffusivity of phosphorus. It is distinctly higher than $D_B$ in silicon and therefore not limiting the diffusion process.

The boron and phosphorus concentrations are parameterized as presented in [39]. For the overall concentration...
of boron in the BSG layer at the start of the process a value of $5 \times 10^{21} \text{cm}^{-3}$ is used, which is estimated by inductively coupled plasma spectrometry.

2.7. Solving of the differential equations

The simulated depth profiles of the dopants phosphorus $c_P(x,t)$ and boron $c_B(x,t)$ follow with time the partial differential equations (1) and (2) taking into account the diffusivities $D_P$ from equation (8) and $D_B$ from equation (10) as well as the boundary conditions of the concentration $c_P$ and $c_B$ of the dopants for the respective process.

The values of the solubilities and the Arrhenius fits of the diffusivities ($D_{PK0}$, $D_{PT0}$, $n_P$, $D_{Bi}$) of the dopants phosphorus and boron in silicon are determined by investigations of the respective dopant profiles without counter doping and in agreement with literature values, see [39]. With this model the joint in diffusion of both dopants can be reproduced.

3. EXPERIMENTAL DETAILS

3.1. Sample preparation

(100) orientated, phosphorus doped n type and boron doped p type float zone (FZ) and Czochralski (Cz) silicon substrates with an area of $50 \times 50 \text{mm}^2$ are used to manufacture doping samples. The FZ silicon substrates have a thickness of $250 \mu\text{m}$, the Cz substrates $180 - 190\mu\text{m}$.

The saw damage at the surfaces of the Cz silicon samples is removed in a chemical polish (CP) [44-47] or in diluted sodium hydroxide solution (NaOH) [47]. Afterwards the samples are cleaned in diluted hydrochloric acid (HCl) and diluted fluoric acid (HF) and undergo a RCA cleaning [48-49]. The RCA cleaning consists of the steps SC1, HF, SC2, HF. Before and after these immersions the samples are rinsed in deionized water.

The samples receive a direct plasma enhanced chemical vapor deposition (PECVD) of a SiO$_x$N$_y$B$_z$:H layer, which will be denoted as borosilicate glass (BSG) in the following [50]. For this purpose an Oxford Instruments Plasmalab 100 System with silane (SiH$_4$), nitrous oxide (N$_2$O), diborane (B$_2$H$_6$), and hydrogen (H$_2$) as precursor gases is used. Diborane is diluted to 0.5% in hydrogen. The PECVD BSG layer has an average thickness of 27 or 73 nm and a boron concentration of $(5 \pm 1) \times 10^{21} \text{cm}^{-3}$, a silicon concentration of $(1.4 \pm 0.2) \times 10^{22} \text{cm}^{-3}$, and a refractive index of 1.5 at a wavelength of 633 nm.

Some samples receive a silicon nitride layer (SiN$_x$:H) [50] which is deposited in the same PECVD system, to mask the substrate against an in diffusion of phosphorus during a high temperature step in a phosphorus diffusion furnace. The SiN$_x$:H layer has a thickness of 70 nm and a refractive index of 2.0 at a wavelength of 633 nm.

The samples are heated in a Centrotherm horizontal tube furnace at atmospheric pressure to diffuse boron from the BSG layer into the silicon substrate. This diffusion step is carried out at temperatures of 850 - 1012 °C for 100 min in nitrogen atmosphere (N$_2$) or in an oxidation tube or in a phosphorus oxychloride diffusion tube (POCl$_3$).

3.2. Characterization

The thickness of the deposited layers is determined with a Woollam V VASE ellipsometer [51] in the wavelength range of 300 - 1000 nm. After the diffusion step, the borosilicate glass layer is dissolved in dilute hydrofluoric acid with a concentration of 0.5 to 25%. The concentration of boron, phosphorus, and silicon of the dissolved BSG layer is determined by inductively coupled plasma spectrometry.

The sheet resistivity of the emitter is determined by means of four terminal sensing [52]. The depth profile of the net dopant concentration is determined by electrochemical capacitance voltage (ECV) profiling [22,53] with a WEP CVP21 wafer profiler. For selected samples the dopant profiles are determined by secondary ion mass spectrometry (SIMS) [22,54] using a Cameca IMS 4f spectrometer.

4. RESULTS

4.1. PECVD BSG emitter diffused-in in nitrogen atmosphere

The depth profile $c_B(x)$ of a boron emitter, which is produced by a borosilicate glass layer with a thickness of
$d_{BSG} = 73 \text{ nm on n type substrates}$, is shown in Figure 5 for diffusion temperatures of 850 to 1012 °C for 100 min. With rising temperature, the depth of the emitter increases. Its boron concentration continuously declines from the surface. The surface concentration of the boron emitter matches well the solubility of boron in silicon [55-58]. Therefore, the used borosilicate glass works as an unlimited diffusion source [59,60].

In Figure 6 the diffusivity of the dopant boron $D_B$ obtained by Boltzmann Matano analysis with equation (3) is plotted against its concentration $c_B$. It increases with rising temperature. Furthermore, it increases in the extrinsic area with rising boron concentration. It is fitted according to equation (9) with

$$D_B = 2.8 \exp \left( \frac{-3.5 \text{ eV}}{k_B T} \right) \left( 1 + \frac{c_B}{n_i} \right) \ cm^2 \ s^{-1} \ \ \ \ (11)$$

The determined diffusivity $D_B$ matches values presented in [55] and [61].

The profiles of the prepared emitters are well reproduced by the simulation, see Figure 5. Deviations emerge mainly because of a scattering of the produced emitters through process inaccuracies.

### 4.2. PECVD BSG emitter with SiNx:H capping layer

The BSG boron emitter profiles with and without SiNx:H capping correspond well with each other. The boron concentration is slightly reduced with a capping layer. When the diffusion step is carried out in a phosphorus diffusion tube, the phosphorus concentration in the substrate is less than $1 \times 10^{17} \text{ cm}^{-3}$.

Thus, PECVD BSG emitters can be produced with a silicon nitride capping layer. Also, the capping layer works as a diffusion barrier and prevents a compensating diffusion of phosphorus into the silicon substrate, see also [62-66].

### 4.3. PECVD BSG without SiNx:H capping layer in phosphorus diffusion

In this part of the investigation the in diffusion of phosphorus during a conventional POCl$_3$ diffusion into a silicon substrate coated with a PECVD BSG layer without a capping layer is determined. Here, silicon substrates with a BSG layer and unmasked substrates are exposed to a POCl$_3$ diffusion at a temperature of 832 °C and a duration of 44 min. With this process conventional phosphorus profiles arise independently of the presence of a BSG layer, revealing a high diffusivity of phosphorus in the BSG.

The phosphorus depth profile, which occurs with and without a PECVD BSG layer, is shown together with the respective simulated phosphorus profile in Figure 7. The depth profile exhibits the kink and tail region usually observed for a phosphorus diffusion. It is hardly affected by the BSG layer. Furthermore, only small differences occur with and without BSG layer, like a slight drop of the surface concentration and a slight reduction of phosphorus concentration for concentrations below $1 \times 10^{16} \text{ cm}^{-3}$. The sheet resistivity of the occurring phosphorus profile with BSG layer is about 10% higher than the resistivity without BSG layer.

The phosphorus profile with and without PECVD BSG layer is well matched by the simulation. Further, the simulation yields a completely compensated convex boron profile, which is accompanied by a low diffusivity of boron compared to phosphorus and a reduction of the boron diffusivity in the kink region of the phosphorus profile.

### 4.4. Boron diffusion from a PECVD BSG without capping in a POCl$_3$ diffusion furnace

The in diffusion of the dopants during a PECVD BSG diffusion in a POCl$_3$ diffusion furnace is determined. Here silicon substrates with a BSG layer and uncoated silicon substrates undergo a high temperature step of 920 °C for 100 min in a POCl$_3$ diffusion furnace. As process gas, nitrogen is used without addition of POCl$_3$. Yet the inner walls of quartz tubes of the POCl$_3$ diffusion furnace are coated with PSG by previous POCl$_3$ diffusions and serve as a phosphorus source. Therefore, the BSG drive in process yields a phosphorus profile, which exhibits a completely compensated boron concentration. The high concentration of dopants leads to a mutual reduction of the diffusivity and hence to a convex boron profile.
The phosphorus concentration of a PECVD BSG layer is determined to \( (2.5 \pm 0.5) \times 10^{21} \text{ cm}^{-3} \) after a diffusion step in a POCl\(_3\) diffusion furnace, with its inner walls coated with PSG by previous POCl\(_3\) diffusions. The depth profiles of the phosphorus and boron concentration with and without PECVD BSG layer are shown in Figure 8 together with the respective simulated depth profiles.

A phosphorus profile can be determined in case of a diffusion with and without BSG layer. In the case of samples without BSG layer the phosphorus profile has a much higher sheet resistivity and a much lower surface concentration than a phosphorus profile arising at this temperature with an addition of POCl\(_3\). The concentration of the phosphorus profile emerging without BSG is much lower than the one arising with BSG. It does not exhibit a kink region.

In the case of samples with BSG, the phosphorus profile has a sheet resistivity much lower and a surface concentration much higher than without BSG. Also there is a completely compensated boron concentration besides the phosphorus concentration in the substrate with BSG layer. The boron concentration is much lower than the phosphorus concentration and proceeds convex.

The simulation matches well the profiles of the phosphorus and boron concentration with PECVD borosilicate glass.

The reduction of the concentration of the phosphorus profile compared to a POCl\(_3\) diffusion at this temperature without BSG and the steep concentration decrease of the phosphorus profile close to the surface are consequences of the high boron concentration in this region, which decreases the phosphorus diffusivity, see also [30,31]. The high boron concentration close to the surface forms at the beginning of the diffusion step, when it is not yet over compensated by phosphorus. Likewise the high phosphorus concentration leads to a decrease of the boron diffusivity, see also [28,29]. The decrease of the boron diffusivity results in a convex boron profile, which is much lower than in the case of an in diffusion without phosphorus.

The boron concentration decreases steeply. This corresponds to a low diffusivity. Hence the phosphorus concentration, which decreases the boron diffusivity, is present early and the diffusivity of phosphorus in the BSG layer is high or phosphorus is taken up during heating up already, see also subsection 5.2. The diffusivity in the BSG becomes an important quantity for the profile of a BSG drive in step \( (T > 900 \, ^\circ\text{C}) \) in a POCl\(_3\) diffusion furnace. In contrast, the diffusivity of phosphorus in the BSG layer does not limit the formation of a phosphorus profile during a standard POCl\(_3\) diffusion \( (T < 850 \, ^\circ\text{C}) \), see Figure 7. This arises as a result of the reduction of the phosphorus diffusivity by the non compensated boron concentration at the beginning of the drive in step.

The simulated boron profile shows a concentration decrease with lower slope in the phosphorus tail region compared to the experimental boron profile and so a higher diffusivity of boron. This is seen as a result of the used parameterization of the boron diffusivity. A more detailed model of the boron diffusivity in general [61] and in this case of an overcompensated emitter push effect [28,29] taking into account the different diffusing species [38] is expected to correct the inaccuracy.

5. DISCUSSION

5.1. Phosphorus uptake by borosilicate glass

The use of a borosilicate glass layer on a silicon substrate during a POCl\(_3\) diffusion or a high temperature drive in step in a POCl\(_3\) diffusion furnace without any POCl\(_3\) flow leads to a simultaneous in diffusion of boron and phosphorus into the silicon substrate. This happens because of the boron concentration in the BSG layer, resulting in a low glass transition and flow temperature of the BSG [40,41] and with that in a high diffusivity of phosphorus. Therefore, a BSG layer does not work as a diffusion barrier for phosphorus, unlike silicate glass without boron [62,63], which has a much higher glass transition and flow temperature [40,41,67]. Thus phosphorus can diffuse from POCl\(_3\) present in the diffusion furnace through the BSG layer into the silicon substrate.

Furthermore, a BSG layer promotes an in diffusion of phosphorus into a silicon substrate during a high temperature step in a coated POCl\(_3\) diffusion furnace. This takes place by an enhanced phosphorus uptake from the
atmosphere of the diffusion furnace compared to a bare silicon substrate. Without the addition of oxygen, more phosphorus is taken up in a borosilicate glass layer than on a bare silicon substrate, as oxygen is needed for the formation of a PSG layer according to the reaction \[22\]

\[
4 \text{POCl}_3 + 3 \text{O}_2 \rightarrow 2 \text{P}_2\text{O}_5 + 6 \text{Cl}_2.
\]

In case of a BSG layer on the silicon substrate, oxygen can be provided from the BSG. Furthermore, a borosilicate or silicate glass layer exhibits compressive stress [68]. The stress is reduced with increasing phosphorus and boron fraction [69,70]. This also is assumed as a reason for the uptake of phosphorus in a borosilicate and silicate glass layer.

The used simple model of the diffusion of the two dopants, phosphorus and boron, matches well the results determined experimentally for the different diffusion steps investigated exemplarily. Nonetheless, the use of a more detailed model of phosphorus [38,71 73] and boron diffusion [61,74] in the silicon bulk [36,76 78] but also of the BSG layer and the uptake of the dopants from the atmosphere of the diffusion furnace [33,60,79,80] is expected to lead to an even better match of simulation and experiment.

5.2. Diffusivity of the dopants in the borosilicate glass layer

The diffusivity of the dopants in the borosilicate glass layer has a large effect on their depth profiles generated in the silicon substrate. To illustrate this, the simulated depth profiles of phosphorus and boron are shown respectively in Figures 9 and 10 for different values of the diffusivity of the dopants \(D_{\text{BSG}}\) and \(D_{\text{P in BSG}} = D_{\text{B in BSG}}\).

Here, the in diffusion of boron from a PECVD BSG layer in a coated POCl\(_3\) diffusion furnace is simulated according to subsection 4.4.

The simulated depth profiles of phosphorus are shown in Figure 9. For a low diffusivity of the dopants \(D_{\text{BSG}}\) no phosphorus enters the substrate. The BSG layer blocks the diffusion completely. Starting at a certain diffusivity, a depth profile of phosphorus arises in the substrate: According to Figure 9 at \(D_{\text{BSG}} \geq 10^{-16} \text{cm}^2/\text{s}\) at a BSG layer thickness \(d_{\text{BSG}} = 27 \text{nm}\). With rising diffusivity a depth profile with increasing concentration of phosphorus develops. Figure 9 shows that from a certain high value of the diffusivity, \(D_{\text{BSG}} \geq 10^{-14} \text{cm}^2/\text{s}\), the diffusion process is no longer limited by the diffusivity in the BSG layer. For a further rise of the diffusivity \(D_{\text{BSG}}\) the phosphorus profile shows no heavier doping.

The simulated depth profile of boron is shown in Figure 10. For a low diffusivity of the dopants \(D_{\text{BSG}}\) no phosphorus enters the substrate. The BSG layer blocks the diffusion completely. Starting at a certain diffusivity, a depth profile of phosphorus arises in the substrate: According to Figure 9 at \(D_{\text{BSG}} \geq 10^{-16} \text{cm}^2/\text{s}\) at a BSG layer thickness \(d_{\text{BSG}} = 27 \text{nm}\). With rising diffusivity a depth profile with increasing concentration of phosphorus develops. Figure 9 shows that from a certain high value of the diffusivity, \(D_{\text{BSG}} \geq 10^{-14} \text{cm}^2/\text{s}\), the diffusion process is no longer limited by the diffusivity in the BSG layer. For a further rise of the diffusivity \(D_{\text{BSG}}\) the phosphorus profile shows no heavier doping.

The simulated depth profile of boron is shown in Figure 10. For a low diffusivity of the dopants \(D_{\text{BSG}}\) a boron profile of shallow depth and low surface concentration emerges, as the diffusion of boron into the silicon substrate is limited by the low diffusivity in the BSG layer, in Figure 10 for \(D_{\text{BSG}} = 10^{-20} \text{cm}^2/\text{s}\). For increasing diffusivity heavier doped depth profiles of boron with higher surface concentrations arise, in Figure 10 for \(D_{\text{BSG}} = 10^{-19}\) to \(10^{-16} \text{cm}^2/\text{s}\). With a higher diffusivity the diffusion of phosphorus into the silicon substrate leads to the emitter push effect, resulting in much deeper boron profiles, in Figure 10 for \(D_{\text{BSG}} \geq 10^{-15} \text{cm}^2/\text{s}\). For a further rising diffusivity, the increasing phosphorus concentration leads to a reduction of the boron diffusivity close to the surface of the silicon substrate, yielding a convex depth profile of boron, in Figure 10 for \(D_{\text{BSG}} \geq 10^{-13} \text{cm}^2/\text{s}\). With increasing diffusivity, the high phosphorus concentration is already
present early during the BSG drive in process, reducing the diffusivity of boron in the silicon substrate already at an early stage, and thus resulting in shallow depth profiles of boron. In Figure 10 this is the case for $D_{\text{BSG}} = 10^{-13}$ cm$^2$/s to $10^{-8}$ cm$^2$/s.

A high diffusivity of the dopants in the PECVD BSG layer $D_{\text{BSG}}$ provides a high concentration of boron at the surface of the silicon substrate, see Figure 10. Consequently, $D_{\text{BSG}}$ can be determined from the surface concentration of the dopants. For the dopant profiles $D_{\text{BSG}}$ is estimated from $10^{-13}$ to $10^{-10}$ cm$^2$/s, independent of the phosphorus amount of the BSG layer [40]. This diffusivity allows for a sufficient transport of the dopants to the surface of the substrate to work as an infinite doping source in the case of the boron profile, see Figure 5, and the phosphorus profile, see Figure 7.

5.3. Back contact solar cell

The simultaneous creation of differently doped regions in a silicon substrate [80] can be used for the production of a back contact solar cell [81] with a rear side $p$ $n$ junction [82,83]. Through the simultaneous in diffusion of phosphorus and boron [16] all necessary doped regions can be realized during only one high temperature step.

To demonstrate this, $n$ type silicon substrates with a full area BSG layer ($d_{\text{BSG}} = 27$ nm) and a structured silicon nitride layer ($d_{\text{SiN}} = 70$ nm) as a diffusion barrier for phosphorus on their rear side are prepared. During a sequential diffusion (a boron drive in step with $T > 900$ °C followed by a POCl$_3$ diffusion step with $T < 850$ °C) in a POCl$_3$ diffusion furnace a $p^+$ emitter doped positively with boron forms in the areas with SiN$_x$:H layer, whereas in the unmasked areas a $n^{++}$ BSF [3,84] heavily doped with phosphorus, and on the front side of the substrate a $n^+$ front surface field (FSF) [85] forms. The doping structure of the silicon substrate produced like this is shown in Figure 11.

The sheet resistivities and the doping profiles of the different regions correspond to those of identically processed full area silicon substrates. Also the results hold essentially for $p$ and $n$ type substrates as well as for invertedly masked structures. Average sheet resistivities of $39 \Omega/\square$ for the $p^+$ region, $19 \Omega/\square$ for the $n^{++}$ region and $48 \Omega/\square$ for the $n^+$ region were obtained.

The dopant profiles of the three regions can be further optimized by applying a co diffusion step (a boron drive in step with a POCl$_3$ flow starting at a certain time of the step) in a coated POCl$_3$ diffusion furnace. Using the developed model of the simultaneous diffusion of boron and phosphorus, an adjustment of the BSG layer thickness and the process temperature and duration can be carried out to obtain desired profiles for the three regions.

6. CONCLUSION

This study reveals the interacting diffusion phenomena of boron and phosphorus in silicon. Co diffusion processes of boron and phosphorus through a BSG layer are investigated experimentally. Key findings are: An overlying BSG layer does not prevent the formation of a phosphorus emitter in silicon substrates during phosphorus diffusion, and it can even enhance the uptake of phosphorus into a silicon substrate compared to a bare substrate.

Based on these findings and known models of the diffusivities of boron and phosphorus a model allowing for the prediction of co diffused dopant profiles is presented. Furthermore, a potentially industrially feasible process to fabricate three diverse phosphorus $n^{++}$, $n^+$, and boron $p^+$ doped regions during one single high temperature step is introduced for the production of highly efficient interdigitated back contact silicon solar cells.

ACKNOWLEDGEMENTS

Part of this work was financially supported by the German Federal Ministry for the Environment, Nature Conservation and Nuclear Safety (FKZ 0325581).

REFERENCES


32. Wagner H, Dastgheib Shirazi A, Chen R, Dunham ST, Kessler M, Alternatt PP. Improving the predictive power of modeling the emitter diffusion by fully including the phosphosilicate glass (PSG) layer. Proceedings IEEE PVSC Seattle. 2011; 2957. DOI:10.1109/PVSC.2011.6186566


