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Surface Passivation of Germanium with ALD Al$_2$O$_3$: Impact of Composition and Crystallinity of GeO$_x$ Interlayer

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Abstract: Germanium is an excellent material candidate for various applications, such as field effect transistors and radiation detectors/multijunction solar cells, due to its high carrier mobilities and narrow bandgap, respectively. However, the efficient passivation of germanium surfaces remains challenging. Recently, the most promising results have been achieved with atomic-layer-deposited (ALD) Al$_2$O$_3$, but the obtainable surface recombination velocity (SRV) has been very sensitive to the surface state prior to deposition. Based on X-ray photoelectron spectroscopy (XPS) and low-energy electron diffraction (LEED), we show here that the poor SRV obtained with the combination of HF and DIW surface cleaning and ALD Al$_2$O$_3$ results from a Ge suboxide interlayer (GeO$_x$, $x < 2$) with compromised quality. Nevertheless, our results also demonstrate that both the composition and crystallinity of this oxide layer can be improved with a combination of low-temperature heating and a 300-Langmuir controlled oxidation in an ultrahigh vacuum (LT-UHV treatment). This results in a reduction in the interface defect density ($D_{it}$), allowing us to reach SRVs as low as 10 cm/s. Being compatible with most device processes due to the low thermal budget, the LT-UHV treatment could be easily integrated into many future devices and applications.

Keywords: surface passivation; atomic layer deposition; aluminum oxide; ultrahigh vacuum; corona oxide characterization of semiconductors; quasi-steady-state microwave-detected photoconductance decay; germanium; X-ray photoelectron spectroscopy; low-energy electron diffraction

1. Introduction

Germanium (Ge) is used in various semiconductor devices due to its attractive electronic properties, such as high carrier mobility and a relatively small bandgap [1,2]. The high carrier mobility is particularly beneficial for metal–oxide–semiconductor field effect transistors (MOSFET) in alleviating the drain current saturation problem [3–5], whereas the small bandgap enabling efficient absorption in a wide wavelength range (200–1800 nm) [1,6,7] is crucial for numerous optoelectronic devices, including near-infrared detectors and thermophotovoltaic and multijunction solar cells [8–17]. Despite the many beneficial properties offered by Ge, utilizing its full potential in devices has turned out to be challenging. This is primarily due to the unfavorable surface properties of Ge, namely the difficulties in being able to achieve efficient surface passivation [1,18,19]. Even though Ge oxides can form a high-quality interface (with Ge) [18,20,21], they are known to be highly unstable and water-soluble [1,18,19], which practically prevents their use as surface passivation layers in recombination-sensitive devices. As a result, many surface passivation studies have been carried out with various externally deposited thin films, such as aluminum oxide (Al$_2$O$_3$), zirconium dioxide (ZrO$_2$), hafnium dioxide (HfO$_2$), amorphous silicon (a-Si), amorphous silicon carbide (a-SiC$_x$), and silicon nitride SiN$_x$ [1,18,22–32]. Similar to silicon (Si), one of the most promising candidates has turned out to be atomic-layer-deposited...
(ALD) Al₂O₃, which has recently been shown to significantly reduce the achievable surface recombination velocities (SRV) down to values as low as 2.7 cm/s [27–29].

The SRV of a Ge surface passivated with ALD Al₂O₃ has been shown to be very sensitive to the condition of the surface right before the deposition of the passivation layer [27]. Due to the extremely high etch rate of Ge in hydrogen peroxide solution [33,34], neither standard clean 1 nor standard clean 2 are applicable surface-cleaning treatments, narrowing the selection down to wet treatments such as HCl and HF dips. Surface cleaning performed with mere HCl without a following rinse in deionized water (DIW) (referred to as reference later on in the text) prior to the ALD of Al₂O₃ was seen to produce significantly better results than the combination of HF and DIW rinsing [27]. Although the latter treatment is speculated to result in a thin GeOₓ layer with low quality, determining the exact reason for the worse SRV needs further studies. In the case of Si, it is well known that a thin SiO₂ layer at the Si surface prior to the ALD of Al₂O₃ is of crucial importance for achieving good surface passivation and a low SRV [35,36]. Furthermore, the combination of low-temperature heating and controlled oxidation in an ultrahigh vacuum (LT-UHV treatment) of this thin SiO₂ layer prior to ALD has been shown to increase the negative fixed charge density (Qₜot) and to decrease the defect density (Dₓ) at the interface, thus resulting in improved surface passivation via both increased field effect and chemical passivation [37,38]. The question is whether a similar approach could be used in the case of Ge to, e.g., increase the degree of crystallinity and/or alter the composition of the thin GeOₓ layer, thus possibly improving the resulting surface passivation as well.

In this work, we utilize surface science methods such as X-ray photoelectron spectroscopy (XPS) and low-energy electron diffraction (LEED) to study the surface of Ge in detail after the two different cleaning treatments: (i) mere HCl as a reference and (ii) the combination of HF and DIW rinsing. The results are compared to the degree of surface passivation, i.e., the SRV, Qₜot, and Dₓ, obtained for both treatments after the ALD of Al₂O₃. The SRV is defined from minority carrier lifetimes measured by quasi-steady-state microwave photoconductance decay (QSS-µPCD), while Qₜot and Dₓ are measured with the corona oxide characterization of semiconductors (COCOS) technique. Additionally, we apply an LT-UHV treatment for the HF-cleaned and DIW-rinsed samples, characterize its effect on both the sample surface prior to ALD and the resulting passivation quality (the SRV, the Qₜot, and the Dₓ) after ALD, and compare the results to earlier data reported for Si. Finally, the obtained results are discussed from a device perspective.

2. Experimental Details

The process flow of the conducted experiments is depicted in Figure 1. A batch of 189 ± 4 μm thick, single-side polished, 4-inch, n-type [100] Czochralski-grown (CZ) Ge wafers with 18–37 Ωcm base resistivity was used in these experiments. The first process step was surface cleaning. The reference samples were treated with 31.6% v/v HCl for 60 s without any DIW rinsing, whereas other samples were treated with 1% v/v HF for 90 s, followed by DIW rinsing for another 90 s. A high HCl concentration was selected due to its reported ability to efficiently remove carbon and oxide layers while still resulting in a smooth Ge surface with monochloride and dichloride termination [19]. For HF, the selected concentration and duration of the treatment were a compromise between the inefficient removal of impurities (e.g., oxygen, carbon, and copper) and resulting surface roughness enhancement [19]. For one set of the HF + DIW-treated samples, an additional low-temperature heating and controlled oxidation in an ultrahigh vacuum (LT-UHV treatment) was carried out immediately after the surface cleaning. For the LT-UHV treatment, the sample set was loaded into a UHV chamber and placed on top of a heater element using a cradle. The chamber was pumped down to a vacuum level of 10⁻⁸ mbar. The wafers were heated at 350 °C for 30 min, which was followed by a 300-Langmuir (L) oxidation at the same temperature. These parameters were selected based on earlier studies on Si [37,38]. After the treatment, the wafers were cooled down to room temperature under
UHV conditions. After unloading the sample set from the UHV chamber, it was rejoined with the other samples and the rest of the process was identical for all of them.

![Figure 1. The process flow of the conducted experiments.](image)

After surface cleaning (and optional LT-UHV treatment), the sample surfaces were characterized with XPS and LEED to obtain information on the oxidation state and crystallinity. Photoemission spectra were measured using a Thermo Scientific Nexsa XPS system with a monochromatized Al Kα radiation source and dual-mode charge compensation. High-resolution core-level spectra were collected with a 50 eV pass energy. The X-ray spot size was 200 µm. Next, a 23 nm thick Al₂O₃ layer was deposited on the samples through thermal ALD (Beneq TFS 500). The deposition was carried out at 200 °C with H₂O and trimethylaluminum (TMA) precursors. The number of cycles was 200. After ALD, to activate the Al₂O₃ passivation, the samples were annealed at 400 °C in N₂ for 30 min [27].

Finally, both the injection-dependent effective minority carrier lifetime (τ_{eff}) and the Ge-Al₂O₃ interfacial properties (namely, the thin-film charge (Q_{tot}) and the interface defect density (D_{it})) were measured using a Semilab PV2000A semiconductor characterization tool [39]. The τ_{eff}, which reflects the overall recombination activity of a sample, was measured with the quasi-steady-state microwave-detected photoconductance decay technique (QSS-µPCD) as an average over five points around the center of the sample [40]. The obtained τ_{eff} was used to extract the maximum SRV at an injection level of 7 × 10^{14} cm⁻³ by assuming negligible bulk recombination, which was likely a valid assumption since high-quality substrates were used. Ge-Al₂O₃ interfacial properties, namely Q_{tot} and D_{it}, were assessed with the corona oxide characterization of semiconductors (COCOS) method [41].

### 3. Results and Discussion

To compare the Ge surface state after mere HCl (reference) and combined HF + DIW cleaning, Figure 2a presents the Ge 3d spectra measured with XPS from both samples. The Ge 3d spectra characterize the direct bonds between Ge and oxygen atoms. A difference between the samples is seen at binding energies in the range of 31.0–32.5 eV. The shoulder in the Ge 3d spectrum of the HF + DIW-cleaned sample indicates the presence of a larger...
amount of Ge suboxides (GeO_x with x < 2, i.e., a Ge atom has fewer than four direct oxygen bonds). This is in good agreement with the current literature, which suggests that GeO_2 is readily water-soluble, whereas GeO_x (with x < 2) is much more resistant to chemicals. Even after an HF dip, small amounts of GeO_x were found, although a DIW rinse carried out due to safety concerns is more likely to cause additional oxide formation [42]. Similarly to here, when comparing HBr and HF cleaning treatments prior to their PECVD a:Si:H Ge surface passivation, Berghuis et al. also reported some remains of GeO_x on the Ge surface after HF cleaning [29].

To study the Ge surface oxygen content more closely, we also measured the oxygen O 1s XPS spectra for both of the samples. The obtained results are presented in Figure 2 b). As expected, both samples similarly show the Ge Auger peak at a binding energy of 535 eV [43]. However, more interestingly, a clear oxygen peak at around 531 eV is seen only for the HF + DIW-cleaned sample and not for the reference. This is in agreement with the Ge 3d spectra presented in Figure 2a, and further proves the presence of an oxide layer at the surface of the HF + DIW-cleaned sample. The impact of this oxide layer on the surface crystallinity was qualitatively studied with the LEED patterns presented in Figure 2c). The reference sample provides sharper diffraction spots and, thus, a more crystalline surface than the HF + DIW-cleaned sample.

These findings complement the previous results, where the combination of HF and DIW rinsing prior to the ALD of Al_2O_3 was seen to lead to a worse SRV than mere HCl [27]. In contrast to the case of Si—where the combination of HF and DIW rinsing removes oxides efficiently and leaves the surface H-terminated, preventing subsequent oxidation [27]—in the case of Ge, there is clearly an oxide layer remaining on the surface after the treatment. While for Si, the presence of a thin chemical oxide layer has been shown to improve ALD Al_2O_3 passivation [35], for Ge, this seems detrimental. In the case of Ge, the presence of a considerable amount of poor quality GeO_x (x < 2) at the Ge/Al_2O_3 interface could (i) reduce the amount of negative fixed charge obtainable by the Al_2O_3 passivation layer,
decreasing the field effect, and/or (ii) increase the defect density (compared to better-quality GeO$_2$), deteriorating the chemical passivation. The COCOS results presented (Figure 4) and discussed later on shed more light on this [18,44–49].

The XPS spectra and the LEED pattern for HF + DIW cleaned sample shown in Figure 2 demonstrated that there is a thin GeO$_x$ suboxide (x < 2) layer remaining on the sample after the cleaning. In the case of silicon with a similarly chemically grown thin SiO$_x$ layer at the wafer surface, the combination of low-temperature heating and controlled oxidation in an ultrahigh vacuum (LT-UHV treatment) before ALD Al$_2$O$_3$ surface passivation has been shown to improve both the oxide and the interface quality [37,38]. Here, we studied whether the quality of the thin GeO$_x$ layer, namely the composition and the crystallinity, could be similarly improved using LT-UHV treatment. Figure 3a,b show the effect of LT-UHV treatment on the Ge 3d and O 1s XPS spectra obtained for the HF + DIW-cleaned sample. The LT-UHV treatment was seen to slightly increase the overall oxygen content (the peak height at ~531 eV in O 1s spectra increased) and to shift the germanium oxide towards a higher oxidation state (the shoulder in the Ge 3d spectrum moved to 31.5–33.5 eV); in other words, GeO$_x$ after the LT-UHV treatment resembled GeO$_2$ more than GeO$_x$ (before the LT-UHV treatment) [50]. Furthermore, the LEED pattern comparison presented in Figure 3c shows sharper diffraction spots after the LT-UHV treatment, which indicates an increased degree of crystallinity at the oxidized surface. This matches well with the case of Si, where some of the SiO$_x$ was converted to SiO$_2$ followed by an increased degree of crystallinity [37,38]. It is worth noting that the LT-UHV treatment parameters used here were selected based on earlier studies on Si [37,38]. Keeping in mind the different oxidation kinetics for Ge, there could possibly be room for further improvement by optimizing the parameters specifically for Ge. Nevertheless, in terms of Ge surface passivation, both the oxide shift towards higher oxidation states and the increased degree of crystallinity pave the way for the reduced density of surface defects [18,45–49].

Figure 3. (a) Ge 3d and (b) O 1s XPS spectra for HF + DIW-cleaned samples with and without the LT-UHV treatment. (c) LEED patterns for same samples.
Finally, we studied the impact of increased Ge oxide quality and improved crystallinity obtained with the LT-UHV treatment for the HF + DIW-cleaned sample on ALD Al₂O₃ surface passivation. We characterized the surface passivation by measuring the effective minority carrier lifetime (τ_eff) and used that to determine the maximum surface recombination velocity. Figure 4a presents the injection-dependent τ_eff for the samples that experienced HCl cleaning (reference), HF + DIW cleaning, and HF + DIW cleaning followed by the LT-UHV treatment. Without the LT-UHV treatment, the HF + DIW sample has approximately a 200 μs lower lifetime across the whole injection level range when compared to the reference. This translates to a 42% higher maximum SRV at 7 × 10¹⁴ cm⁻³⁻¹ injection for the HF + DIW-cleaned sample. Similarly, Berghuis et al. reported earlier that a more efficient GeO₂ removal is beneficial for Ge surface passivation with PECVD a:Si:H [29]. Most notably, the addition of LT-UHV treatment improved the lifetime to the same level as in the reference sample and, thus, reduce the SRV to below 10 cm/s. Looking more closely at the injection level dependency, the reference sample was seen to have a slightly higher lifetime in the low-injection region (1250 μs vs. 1200 μs at an injection of 2 × 10¹⁴ cm⁻³⁻¹), whereas in the high-injection region, the LT-UHV-treated sample was seen to perform better (500 μs vs. 600 μs at an injection of 2 × 10¹⁵ cm⁻³⁻¹). The slightly smaller injection-level dependency of the LT-UHV-treated sample indicates that it should have improved chemical passivation quality (i.e., lower D_H) compared to the reference.

To study the reasons behind the improved surface passivation in more detail, we performed COCOS measurements on all three different sample types to determine the Q_tot and D_H. Figure 4b presents Q_tot, the figure of merit for field effect passivation, and Figure 4c presents D_H, the figure of merit for chemical passivation. The reference sample showed the highest thin-film charge in this comparison (Q_tot = −2.3 × 10¹² cm⁻²) followed by the HF + DIW (Q_tot = −1.9 × 10¹² cm⁻²) and HF + DIW with LT-UHV treatment (Q_tot = −1.5 × 10¹² cm⁻²) samples, respectively. Interestingly, the charge of the HF + DIW-cleaned sample decreased after the LT-UHV treatment, which was the opposite effect to the case of Si, where the LT-UHV treatment was shown to increase the Q_tot [37,38]. On the other hand, the D_H behaved in a completely opposite manner. The lowest D_H was achieved with the LT-UHV-treated sample (D_H = 2.3 × 10¹² cm⁻²) followed by the HF + DIW-cleaned sample (D_H = 3.7 × 10¹² cm⁻²), while the reference sample showed the worst value (D_H = 4 × 10¹² cm⁻²). Thus the LT-UHV treatment clearly reduced the D_H, which is in line with the surface science results and also agrees well with the LT-UHV treatment results obtained for Si [37,38]. It is interesting that even though the overall passivation after the LT-UHV treatment matched the reference, the passivation mechanisms differ from each other. The passivation of the reference samples relied on a strong field effect, while in the case of the LT-UHV-treated sample, the passivation was more balanced with improved chemical passivation.

Finally, it is important to consider what the obtained results mean from a Ge device perspective. Surface recombination is detrimental to the performance of many types of Ge devices and, hence, efficient passivation is a crucial part of the device-manufacturing process. For instance, in state-of-the-art photodiodes, surface recombination affects both the dark current and the sensitivity, which are important figures of merit [51]. Fortunately, the surface recombination characteristics achieved in this work are enough for high-performance devices [52]. Furthermore, the low thermal budget of the LT-UHV treatment makes it compatible with most device processes. Interestingly, very high absorption over the whole wavelength range determined by the Ge band gap has been achieved by nanostructuring the Ge surface [6,7,53,54]. This nanostructure, also known as black Ge, multiplies the surface area and, thus, requires more efficient passivation methods than planar counterparts. A recent study showed that, even with the surface passivation scheme combining HCl cleaning with subsequent ALD Al₂O₃ and improved by an additional intermediate H₂O₂-based etchback, the achievable minimum SRV of black Ge still seems to be heavily dependent on the surface area [55], indicating the need for improvements. One possible route towards enhanced passivation for the nanostructured surfaces could be...
applying the LT-UHV treatment seen here to reduce the \( D_{\text{it}} \). With such a large surface area, even a small reduction in \( D_{\text{it}} \) could lead to a significant upgrade.

**Figure 4.** (a) Injection-dependent \( \tau_{\text{eff}} \) of ALD-Al\(_2\)O\(_3\)-coated samples with different combinations of pretreatments (Reference (HCl), HF + DIW, and HF + DIW + LT-UHV treatment). All samples were post-deposition-annealed for 30 min at 400 °C in N\(_2\) ambient. The value labelled for each curve represents the corresponding SRV extracted at \( 7 \times 10^{14} \) cm\(^{-3}\) injection level. Extracted (b) \( Q_{\text{tot}} \) and (c) \( D_{\text{it}} \) values measured using COCOS.

**4. Conclusions**

The findings of this work suggest that Ge surface cleaning with the combination of HF and DIW rinsing prior to ALD resulted in a poor-quality GeO\(_x\) layer, which reduced the effectivity of a subsequent Al\(_2\)O\(_3\) surface passivation. Nevertheless, the combination of low-temperature heating and controlled oxidation in an ultrahigh vacuum (LT-UHV treatment) improved the GeO\(_x\) quality by shifting its composition towards higher oxidation states and by increasing the degree of its crystallinity. As a result, the SRV after a subsequent ALD Al\(_2\)O\(_3\) passivation was reduced by 32%. The reason for this improvement was the reduction in \( D_{\text{it}} \) from \( 3.7 \times 10^{12} \) cm\(^{-2}\) to \( 2.3 \times 10^{12} \) cm\(^{-2}\). Such a decrease in \( D_{\text{it}} \) might provide much-needed assistance for the surface passivation of Ge nanostructures in the future.


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Abbreviations

Manuscript abbreviations and their definitions.

ALD  atomic layer deposition
COCOS  corona oxide characterization of semiconductors
D_{it}  interface defect density
DIW  deionized water
QSS-μPCD  quasi-steady-state microwave-detected photoconductance decay
Q_{fr}  fixed charge density
LEED  low-energy electron diffraction
LT-UHV  low-temperature heating and controlled oxidation in an ultrahigh vacuum
SRV  surface recombination velocity
XPS  X-ray photoelectron spectroscopy
\tau_{eff}  effective minority carrier lifetime

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