Atomic layer deposition of high-$\kappa$ dielectric layers on Ge and III-V MOS channels

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Ge and III-V semiconductors are potential high performance channel materials for future CMOS devices. In this work, we have studied Atomic Layer Deposition (ALD) of high-$\kappa$ dielectric layers on Ge and GaAs substrates. We focus at the effect of the oxidant ($\text{H}_2\text{O}$, $\text{O}_3$, $\text{O}_2$, $\text{O}_2$ plasma) during gate stack formation. GeO$_2$, obtained by Ge oxidation in $\text{O}_2$ or $\text{O}_3$, is a promising passivation layer. The germanium oxide thickness can be scaled down below 1 nm, but such thin layers contain Ge in oxidation states lower than 4+. Still, electrical results indicate that small amounts of Ge in oxidation states lower than 4+ are not detrimental for device performance. Partial intermixing was observed for high-$\kappa$ dielectric and GeO$_2$ or GaAs$_x$O$_{1-x}$, suggesting possible correlations in the ALD growth mechanisms on Ge and GaAs substrates.

Introduction

Hafnium-based oxide, deposited by Atomic Layer Deposition (ALD), has recently been implemented as the gate dielectric in 45-nm CMOS devices (1). To further increase transistor performance, high-$\kappa$ gate dielectrics may be combined with Ge and III-V channel materials, which have higher intrinsic carrier mobilities as compared to Si. One of the issues when integrating high-$\kappa$ dielectrics with Ge or III-V channels is obtaining a low defect density at the Ge or III-V interface. Initial devices fabricated on Ge substrates with high-$\kappa$ dielectric layers aimed at preventing an interfacial oxide layer between the Ge channel and the high-$\kappa$ dielectric in order to minimize the equivalent oxide thickness (EOT) of the gate stack. However, the interface state densities of such devices were much too high for device applications (2). In subsequent work, passivating interfacial layers between the Ge channel and the high-$\kappa$ dielectric were developed, for example a thin (~0.6 nm) epitaxial Si passivation layer (3, 4), germanium oxynitride (5), and S-passivation (6, 7).

Passivation of Ge using germanium oxide has been discarded for some time because the limited stability of GeO$_2$ and its solubility in water complicate device fabrication. Yet, recent reports show good passivation using germanium oxide both on n- and p-type Ge. Applying GeO$_2$ prepared by electron-cyclotron-resonance plasma irradiation resulted in an interface trap density ($D_{it}$) of $\sim$6 $\times$ 10$^{10}$ cm$^{-2}$eV$^{-1}$ at the midgap, measured by the ac
conduction method (8). Thermal oxidation and photo-oxidation of Ge were also investigated (8, 9, 10, 11). Thermally grown GeO$_2$ was shown to be an effective passivation layer in combination with HfO$_2$ and Al$_2$O$_3$ deposited by ALD (12, 13, 14). In the oxide, Ge was present in the 4+ oxidation state, and densities of interface states as low as mid $10^{11}$ cm$^{-2}$eV$^{-1}$ were achieved. Ge/GeO$_x$/CeO$_2$ and Ge/GeO$_x$/CeO$_2$/HfO$_2$ gate stacks grown by molecular beam deposition exhibited similar interface state densities even though the GeO$_x$ layer contained Ge in low (2+ and 3+) oxidation states (15). On InGaAs, promising electrical properties in terms of leakage current and interface state density were claimed for Al$_2$O$_3$ layers deposited by ALD (16, 17). The reduction of Fermi-level pinning of the gate insulator/channel interface was ascribed to the removal of arsenic oxides during the ALD process.

ALD is often used to deposit high-$\kappa$ dielectrics on Ge and III-V materials as the electrical quality of the Ge or III-V interfaces can be controlled at the typically low deposition temperature (18). In this work, we investigate structures formed by ALD of Al$_2$O$_3$ and HfO$_2$ on Ge and GaAs substrates. We focus on the role of the oxidant (H$_2$O, O$_3$, O$_2$, O$_2$ plasma) during gate stack formation. We have characterized the oxide layer before and after ALD of the high-$\kappa$ dielectric on Ge and GaAs substrates. The oxidation states, oxide thickness, impurities, and possible intermixing with the high-$\kappa$ dielectric are investigated. The aim of this work is to understand how to control the attained interfacial oxide layer during gate stack formation, and finally to identify electrically active defects limiting the ultimate electrical performance.

**Experimental details**

Ge(100) or Ge-on-Si substrates were cleaned in HF solution. The subsequent surface preparation, germanium oxidation, and high-$\kappa$ deposition were performed in a Polygon® cluster (19) without air exposure between these process steps. Germanium oxidations in O$_2$ were explored in a Polygon 8200 cluster, equipped with an EPSILON™ reactor for thermal treatments and a PULSAR® 2000 reactor for ALD. The samples were first annealed in H$_2$ at 650°C for 10 minutes to desorb O or C that can be present after HF clean (20). Oxidations were subsequently performed in the same reactor in O$_2$ at 700 Torr at temperatures between 300 and 450°C. HfO$_2$ and Al$_2$O$_3$ layers were deposited on the GeO$_2$ layers at 300°C in the ALD reactor from HfCl$_4$ or Al(CH$_3$)$_3$ (trimethylaluminium or TMA) with H$_2$O. Germanium oxidizations in O$_3$ were performed in a PULSAR® 3000 reactor, with subsequent H$_2$O or O$_3$ based ALD of HfO$_2$ or Al$_2$O$_3$ in the same reactor. For this end, 15% O$_3$ was generated in an O$_2$/N$_2$ mixture and 0.5 slm of this flow was further diluted with 2 slm N$_2$ in the ALD reactor. The pressure in the ALD reactor was ~1 Torr.

GaAs substrates were cleaned before ALD in HCl or (NH$_4$)$_2$S solutions. Immediately after the clean, the wafers were blown dry with a N$_2$ gun. Depositions were also performed on the native oxide on GaAs. Thermal ALD of HfO$_2$ and Al$_2$O$_3$ were performed using HfCl$_4$ or TMA with H$_2$O in an ASM PULSAR® 2000 reactor at 300°C (19). Plasma Enhanced ALD (PEALD) of Al$_2$O$_3$ using TMA and O$_2$ plasma was performed in an Oxford Instruments FlexAl® reactor (21) at 300°C.

Angle Resolved X-ray Photoelectron Spectroscopy (AR-XPS) was carried out in a Theta300 system (Thermo Instruments) in parallel angle resolved mode using monochromatized Al K$_\alpha$ radiation. Oxide thicknesses were also verified with
spectroscopic ellipsometry (SE). The areal density of Hf (Hf/nm$^2$) on GeO$_2$ was measured by Rutherford Backscattering (RBS) in a RBS400 endstation with a 1 MeV He$^+$ beam. The areal density of Al (Al/nm$^2$) on GaAs was measured by Total X-ray Reflection Fluorescence (TXRF) in a TEI-Atomika 8300W system. Time of Flight Secondary Ion Mass Spectroscopy (TOFSIMS) was used for depth profiling, using a dual ion beam set up with a 500eV Xe$^+$ ion beam.

Electrical measurements were performed on MOS capacitors with Pt gates. 50 nm thick Pt dots of different sizes were deposited by thermal evaporation using a shadow mask. The electrical characteristics were measured with a Keithley K4200 semiconductor parameter analyzer and a HP4284 LCR meter. The EOT was estimated by fitting the capacitance-voltage (CV) curves using the Hauser routine, including quantum mechanical corrections (22).

**Oxidation of Ge in O$_2$ and O$_3$**

Ge substrates were oxidized in either O$_2$ or O$_3$. The oxidation was carried out at a relatively low temperature range (225 – 450°C) in order to avoid decomposition of GeO$_2$ and desorption of GeO. For the O$_2$ case, the pressure was set at 700 Torr of O$_2$ in order to achieve a sufficiently high oxidation rate. The oxidation of Ge by O$_3$ was performed in the ALD reactor at a nominal O$_3$ partial pressure of 0.04 Torr (not taking into account decomposition by transport from the O$_3$ generator to the reactor and thermal decomposition, as described below).

The oxidation rate in O$_3$ and O$_2$ are compared in Figure 1. For O$_2$, the oxidation rate increases with temperature in the temperature range 300-450°C and follows the Deal-Grove model, as described previously (23). When comparing oxidation in O$_3$ and O$_2$ at 300°C, we observe that O$_3$ oxidation proceeds slightly faster than O$_2$ oxidation even though the partial pressure of O$_3$ was much lower. O$_3$ is more reactive than O$_2$ because O$_3$ dissociates at the Ge surface to produce atomic oxygen thus enhancing oxidation (24):

$$\text{O}_3 \ (\text{surface}) \rightarrow \text{O}_2 + \text{O}$$

Still, the oxidation rate in O$_3$ is relatively slow in the studied temperature range (225 - 370°C), likely because of the low pressure of O$_3$.

O$_3$ oxidation shows a maximum rate at ~300°C: the oxidation rate increases when increasing the temperature from 225 to 300°C, and then again decreases slightly when further increasing the temperature to 370°C (Figure 1). The oxidation rate of Ge by atomic oxygen increases with temperature. The reduced oxidation rate at 370°C cannot be associated with GeO desorption as GeO is known to desorb only at temperatures higher than 420°C (25). Moreover, the decrease in oxidation rate was also observed for oxidation of Si in O$_3$ in the same conditions, and SiO desorption occurs at an even higher temperature range. Another possible explanation for the lower oxidation rate at 370°C is a lower concentration of O$_3$. Indeed, the lifetime of O$_3$ decreases with temperature (24). Before O$_3$ arrives at the Ge or Si surface, O$_3$ can thermally decompose by collisions with other gas molecules (M):

$$\text{O}_3 + \text{M} \leftrightarrow \text{O}_2 + \text{O} + \text{M}$$
\[ O_3 + O \rightarrow O_2 + O_2 \]

Also, \( O_3 \) can be lost at the walls of the reactor

\[ 2 O_3 \rightarrow 3 O_2 \]

Which destruction reactions of \( O_3 \) are predominant will to a large extent depend on the geometry of the reactor. As the oxidations were performed in an ALD reactor where the distance between the wafer and the top wall is only a few mm, we expect a large contribution of wall destruction of \( O_3 \) especially at higher temperature. Indeed, the SiO\(_2\) layer obtained at 370°C showed a decrease in thickness in the direction of the gas flow. The oxide layers grown at 225 and 300°C demonstrated good thickness uniformity (1 – 4% standard deviation) both for Si and Ge.

![Graph showing oxide thickness as a function of oxidation time](image)

Figure 1. Germanium oxide thickness as a function of oxidation time for oxidation of Ge substrates in \( O_3 \) and \( O_2 \) at several temperatures, determined by ellipsometry. Trend lines are indicated as a guide to the eye.

The oxidation states of Ge in the germanium oxide layers grown in \( O_2 \) or \( O_3 \) were investigated by means of XPS. Note that XPS measurements were performed ex-situ, but air exposure between germanium oxidation and loading the samples in the XPS tool was limited to less than ~15 minutes. In order to determine the oxidation states of Ge in the oxide, the Ge3d spectrum was fitted using binding energy shifts of 0.8, 1.8, 2.6, and 3.4 eV for Ge3d in the 1+, 2+, 3+, and 4+ oxidation states (26). As described in reference (15), precise determination of the fraction of Ge in different oxidation states is complicated by the uncertainty in the XPS fitting and interpretation. The XPS fits should therefore be used as a qualitative guide to the key oxidation states, not for quantitative estimation of the ratios of Ge oxidation states. The fraction of Ge\(^{4+}\) (at\% Ge\(^{4+}\) / (at\% Ge\(^{1+}\) + at\% Ge\(^{2+}\) + at\% Ge\(^{3+}\) + at\% Ge\(^{4+}\) ) is plotted as a function of the GeO\(_x\) thickness in Figure 2, while a summary of all oxidation states is shown in TABLE 1.

The fraction of Ge\(^{4+}\) in the germanium oxide layers grown in both \( O_3 \) and \( O_2 \) depends clearly on the thickness of the germanium oxide layer, and less on the oxidation conditions (temperature, \( O_2 \), \( O_3 \), etc.) (Figure 2). For all germanium oxide layers thicker than ~1 nm, obtained either in \( O_2 \) or \( O_3 \), Ge\(^{4+}\) is the predominant oxidation state. The
highest fraction of Ge\(^{4+}\) was observed for the thickest germanium oxide layers grown in O\(_2\) at 450°C. The 1 nm germanium oxide grown at 300°C in O\(_3\) also contained Ge mainly in the form of Ge\(^{4+}\), only a small fraction of Ge\(^{2+}\) was present.

![Graph](https://via.placeholder.com/150)

**Figure 2.** Fraction of Ge\(^{4+}\) \((\text{at}\% \text{Ge}^{4+}/(\text{at}\% \text{Ge}^{1+} + \text{at}\% \text{Ge}^{2+} + \text{at}\% \text{Ge}^{3+} + \text{at}\% \text{Ge}^{4+})\) as a function of the germanium oxide thickness (XPS) for different oxidation conditions (see TABLE 1).

**TABLE 1.** Summary of the Ge oxidation states in GeO\(_x\) layers obtained by thermal oxidation in O\(_2\) or O\(_3\), as determined by XPS.

<table>
<thead>
<tr>
<th>Oxidation conditions</th>
<th>Ge(^{1+}) (at%)</th>
<th>Ge(^{2+}) (at%)</th>
<th>Ge(^{3+}) (at%)</th>
<th>Ge(^{4+}) (at%)</th>
<th>GeO(_x) thickness (nm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>O(_2) 700 Torr 300°C 120 min</td>
<td>22%</td>
<td>0%</td>
<td>20%</td>
<td>57%</td>
<td>0.88</td>
</tr>
<tr>
<td>O(_2) 700 Torr 300°C 120 min</td>
<td>15%</td>
<td>1%</td>
<td>7%</td>
<td>77%</td>
<td>0.71</td>
</tr>
<tr>
<td>O(_2) 700 Torr 350°C 60 min</td>
<td>7%</td>
<td>7%</td>
<td>0%</td>
<td>86%</td>
<td>1.10</td>
</tr>
<tr>
<td>O(_2) 700 Torr 350°C 120 min</td>
<td>12%</td>
<td>3%</td>
<td>15%</td>
<td>70%</td>
<td>1.35</td>
</tr>
<tr>
<td>O(_2) 700 Torr 400°C 30 min</td>
<td>12%</td>
<td>3%</td>
<td>16%</td>
<td>69%</td>
<td>1.41</td>
</tr>
<tr>
<td>O(_2) 700 Torr 400°C 60 min</td>
<td>10%</td>
<td>0%</td>
<td>12%</td>
<td>77%</td>
<td>1.77</td>
</tr>
<tr>
<td>O(_2) 700 Torr 400°C 60 min</td>
<td>5%</td>
<td>3%</td>
<td>3%</td>
<td>90%</td>
<td>1.30</td>
</tr>
<tr>
<td>O(_2) 700 Torr 400°C 120 min</td>
<td>7%</td>
<td>0%</td>
<td>11%</td>
<td>82%</td>
<td>2.33</td>
</tr>
<tr>
<td>O(_2) 700 Torr 450°C 5 min</td>
<td>3%</td>
<td>6%</td>
<td>0%</td>
<td>91%</td>
<td>1.30</td>
</tr>
<tr>
<td>O(_2) 700 Torr 450°C 18 min</td>
<td>0%</td>
<td>0%</td>
<td>0%</td>
<td>100%</td>
<td>2.50</td>
</tr>
<tr>
<td>O(_2) 700 Torr 450°C 5 min</td>
<td>0%</td>
<td>8%</td>
<td>6%</td>
<td>86%</td>
<td>1.33</td>
</tr>
<tr>
<td>O(_2) 700 Torr 450°C 30 min</td>
<td>0%</td>
<td>0%</td>
<td>4%</td>
<td>96%</td>
<td>2.72</td>
</tr>
<tr>
<td>O(_2) 700 Torr 450°C 60 min</td>
<td>0%</td>
<td>0%</td>
<td>8%</td>
<td>92%</td>
<td>4.20</td>
</tr>
<tr>
<td>O(_2) 0.06 Torr 400°C 5 min</td>
<td>0%</td>
<td>25%</td>
<td>43%</td>
<td>32%</td>
<td>0.37</td>
</tr>
<tr>
<td>O(_2) 12 Torr 400°C 5 min</td>
<td>8%</td>
<td>6%</td>
<td>32%</td>
<td>55%</td>
<td>0.70</td>
</tr>
<tr>
<td>O(_2) 0.06 Torr 500°C 5 min</td>
<td>0%</td>
<td>20%</td>
<td>48%</td>
<td>32%</td>
<td>0.50</td>
</tr>
<tr>
<td>O(_2) 12 Torr 500°C 5 min</td>
<td>0%</td>
<td>12%</td>
<td>21%</td>
<td>67%</td>
<td>0.55</td>
</tr>
<tr>
<td>O(_3) 225°C 30 min</td>
<td>8%</td>
<td>12%</td>
<td>6%</td>
<td>73%</td>
<td>0.66</td>
</tr>
<tr>
<td>O(_3) 300°C 30 min</td>
<td>0%</td>
<td>13%</td>
<td>0%</td>
<td>87%</td>
<td>1.03</td>
</tr>
</tbody>
</table>

Layers thinner than ~1 nm contain a higher fraction of lower oxidation states. Which oxidation states are present depends on the oxidation conditions. Thin layers grown in 700 Torr O\(_2\) at 300°C contain, next to Ge\(^{4+}\), about 10-20 % of Ge in the 1+ and 3+ oxidation states, while almost no Ge\(^{2+}\) is observed. On the other hand, thin germanium oxide layers grown at lower O\(_2\) pressure (0.06 – 12 Torr) and higher temperature (400-500°C) contain Ge\(^{2+}\) and Ge\(^{3+}\) and almost no Ge\(^{1+}\). The presence of Ge\(^{3+}\) in the thin
oxides grown at higher temperature is surprising considering that desorption of GeO can occur at temperatures higher than 420°C (25). On the other hand, no Ge$^{2+}$ is found in the samples grown at lower temperature. This may be related to the much longer oxidation time due to the low oxidation rate of O$_2$. The 0.7 nm germanium oxide layer grown in O$_3$ at 225°C contains predominantly Ge$^{4+}$, while small fractions of Ge$^{1+}$, Ge$^{2+}$ as well as Ge$^{3+}$ are observed.

**ALD of HfO$_2$ and Al$_2$O$_3$ on GeO$_2$ and Ge substrates**

O$_3$ and H$_2$O based ALD of both HfO$_2$ and Al$_2$O$_3$ was performed on thermally grown GeO$_2$ layers, and on HF cleaned Ge substrates for comparison. Air exposure of GeO$_2$ was avoided by performing the oxidation and ALD in clustered reactors. The ALD was performed at relatively low temperature (300°C or lower) to avoid GeO desorption. Still, the ALD has an impact on the thickness of the GeO$_2$ layer, the abruptness of the Ge/GeO$_2$/high-$\kappa$ interfaces, and the predominant oxidation states of Ge.

**Effect of ALD precursors on the germanium oxide interfacial layer thickness**

In order to investigate the effect of the ALD precursors and process conditions on the thickness of the germanium oxide interfacial layer, we considered ALD on 1 nm GeO$_2$ layers (obtained by 20 minutes O$_3$ oxidation at 300°C) and on HF cleaned Ge substrates (0.2 – 0.3 nm GeO$_x$ remains present on HF cleaned Ge substrates). The GeO$_2$ thickness was measured by means of XPS, and the results are summarized in Figure 3.

No further oxidation of Ge occurs during H$_2$O based ALD, both for HfCl$_4$/H$_2$O and TMA/H$_2$O ALD, as reported previously (27). On thermally grown GeO$_2$, the GeO$_2$ thickness remains ~ 1 nm after the H$_2$O based ALD. Also the thin GeO$_x$ layer present after HF clean does not grow further during H$_2$O based ALD.

On the other hand, the Ge substrate oxidizes during O$_3$ based ALD. This is most noticeable for deposition on HF cleaned Ge. A 0.7 nm germanium oxide layer is formed when using the TMA/O$_3$ ALD. An even thicker germanium oxide layer, 1.7 nm, is formed during HfCl$_4$/O$_3$ ALD. First, the differences in germanium oxide thickness for the two processes can be related to differences in O$_3$ pulse times. In the HfCl$_4$/O$_3$ ALD, O$_3$ pulse times of at least 10 seconds are required to achieve a uniform HfO$_2$ layer (Figure 4a). On the other hand, O$_3$ pulse times of about 2 orders of magnitude shorter (< 500 ms) are used for TMA/O$_3$ process (Figure 4b), resulting in substantially less pronounced oxidation of the Ge substrate during the ALD. Second, the GeO$_2$ thickness after HfCl$_4$/O$_3$ ALD is even larger than for oxidation of Ge in O$_3$ at the same temperature, indicating a catalytic effect of Hf species at the Ge surface on the oxidation.

On 1 nm GeO$_2$, we observe a similar difference between the TMA/O$_3$ and HfCl$_4$/O$_3$ ALD. The GeO$_2$ thickness is not affected by the TMA/O$_3$ process, while for the HfCl$_4$/O$_3$ ALD the GeO$_2$ thickness increases to 1.7 nm, the same value as for deposition on HF cleaned Ge. Thus, the GeO$_2$ thickness is independent on the starting surface for HfCl$_4$/O$_3$ ALD due to the excessive re-growth of GeO$_2$ during ALD. The re-growth during HfCl$_4$/O$_3$ ALD can be decreased by decreasing the ALD temperature from 300 to 225°C, where only 1.2 nm GeO$_2$ is observed. Decreasing the deposition temperature for TMA/O$_3$ from 300 to 225°C does not result in a significant thickness decrease.
Figure 3. GeO$_2$ thickness determined by XPS after ALD of 2 nm HfO$_2$ or Al$_2$O$_3$ (a) on 1 nm GeO$_2$ (20 minutes O$_3$ at 300°C) (b) on HF cleaned Ge.

Figure 4. Effect of O$_3$ pulse time on thickness and uniformity (49 points on 300 mm Si wafer); (a) 100 cycles HfCl$_4$/O$_3$ ALD at 300°C; (b) 100 cycles TMA/O$_3$ ALD at 300°C.

Effect of ALD precursors on the abruptness of Ge/GeO$_x$/high-$\kappa$ interfaces

The quality of the interfaces between Ge/GeO$_x$/high-$\kappa$ and the extent of intermixing of the GeO$_2$ and high-$\kappa$ layer also depend on the ALD process parameters, as demonstrated by TOFSIMS depth profiling. We report Ge$^+$ profiles for different ALD layers in Figure 5, similar trends were observed for the GeO$^+$ profiles.

For HfO$_2$ and Al$_2$O$_3$ ALD on 1 nm GeO$_2$, we observe partial intermixing of GeO$_2$ with the high-$\kappa$ dielectric, both for GeO$_2$ grown in O$_2$ (12) and in O$_3$ (Figure 5a, Figure 5b). Still, the amount of GeO$_x$ in the high-$\kappa$ layers is small (note the exponential scale in the TOFSIMS Ge$^+$ profiles). For example, for Al$_2$O$_3$, the intensity of the Ge$^+$ signal is diminished by one order of magnitude at a distance of 1 nm from the GeO$_2$ interface, indicating about 3 at% of Ge. This may explain why this small degree of intermixing was not observed previously by means of AR-XPS (12), while TOFSIMS is sensitive enough to detect it. Still, such small concentrations of Ge in the high-$\kappa$ layer can affect the
electrical properties. Note that TOFSIMS cannot determine the oxidation state of Ge in the high-κ layer, but most likely it is present in an oxidized state and not as Ge\(^0\), as TOFSIMS demonstrated previously that HfCl\(_4\)/H\(_2\)O ALD on HF cleaned Ge substrates results in an abrupt Ge/HfO\(_2\) transition (27). First principles calculations demonstrated that the formation of Ge–O bonds or Hf–O–Ge bonds at or near the interface does not introduce interface states to the Ge energy band-gap (28, 29). On the other hand, it was suggested that such bonds could be responsible for hysteresis often observed in CV curves (23).

For Al\(_2\)O\(_3\) ALD, the amount of intermixing is independent of the oxidant precursor (H\(_2\)O or O\(_3\), Figure 5a). In contrast, for HfO\(_2\) ALD the extent of intermixing depends on the oxidant: the layers intermix more with O\(_3\) than H\(_2\)O based ALD (Figure 5b). For the O\(_3\) based process, the intermixing decreases slightly when decreasing the ALD temperature from 300 to 225°C. The presence of GeO\(_x\) in the high-κ layer indicates that reactions other than the conventional ligand exchange reactions must contribute during ALD on GeO\(_2\), as suggested previously for HfCl\(_4\)/H\(_2\)O ALD on 0.3 nm GeO\(_x\) by considering the first reaction cycle (27).

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Not only the thickness but also the oxidation states of Ge in the interfacial germanium oxide layer can be affected by ALD of the high-κ dielectric. We have analyzed this effect for Al\(_2\)O\(_3\) ALD on GeO\(_2\) by investigating the XPS Ge3d spectra. Unfortunately, a similar analysis cannot be performed for HfO\(_2\) ALD as the Ge3d peak is located at a binding energy very close to the O2s peak from HfO\(_2\) (32.5 eV). The quantitative determination of the oxidation states of Ge is complicated by the presence of the high-κ layer (partial intermixing as described above and potential charge effects). Therefore, instead of deconvolution of the Ge3d signal, we compare the apparent binding energy shift of the germanium oxide 3d peak (Figure 6). The smaller the binding energy shift, the higher the fraction of low oxidation states in the oxide; the larger the binding energy shift, the higher the fraction of high oxidation states in the oxide.
For Al₂O₃ ALD on GeO₂ layers with thicknesses larger than 1 nm, the main oxidation state of Ge remains Ge⁴⁺, indicated by a binding energy shift of the oxidized Ge3d peak of ~3 eV (Figure 6). Thus, contributions of Ge in oxidation states lower than 4+ are small. The binding energy shift decreases when decreasing the thickness of germanium oxide below 1 nm, suggesting the occurrence of Ge in oxidation states of 4+ and lower. This observation is similar to that made for the GeO₂ layers without high-κ dielectrics on top (Figure 2). Thus, when scaling down the thickness of germanium oxide for EOT reduction, lower oxidation states may occur in germanium oxide.

![Figure 6. XPS binding energy shift of the Ge3d oxide peak (relative to the substrate peak) as a function of the germanium oxide thickness for GeO₂ layers and for GeO₂ covered with 2 nm of Al₂O₃. Trend lines are indicates as a guide to the eye. Binding energy shifts of for Ge3d in the 2+, 3+, and 4+ oxidation states (26) are indicated as a guides to oxidation states that can be present.](image)

Electrical characteristics of devices with GeO₂ interfacial layers

Capacitance-Voltage characteristics

Capacitance-Voltage characteristics for Pt gated capacitors with 4-nm thick high-κ dielectrics (H₂O or O₃ based ALD) are compared in Figure 7. The CV characteristics of the capacitors shown here behave nicely. Frequency dispersion in accumulation at higher frequencies is observed due to series resistance as undoped Ge-on-Si wafers were used.

The measured EOT of the capacitors is summarized in Table 2. The determined EOT values are in excellent agreement with EOT estimations based on the thickness of the high-κ (4 nm) and GeO₂ (XPS), assuming a κ-value of 6 for GeO₂ (30), 20 for HfO₂ (31) and 9 for Al₂O₃ (32).

The lowest EOT, 1.5 nm, was obtained for 4 nm HfCl₄/H₂O ALD deposited on 1 nm GeO₂. As shown above, the HfCl₄/H₂O ALD has the advantage of no additional oxidation of Ge during ALD. A higher EOT, 2.0 nm, is obtained for HfCl₄/O₃ ALD, due to the presence of a thicker GeO₂ layer (1.7 nm). XPS indicated that decreasing the ALD temperature for the HfCl₄/O₃ process results in a thinner GeO₂ layer. However, devices with HfCl₄/O₃ deposited at 225°C demonstrated a high leakage current, making these films unsuitable for gate dielectric applications. TOFSIMS indicated that the Cl-content was more than 1 order of magnitude higher for O₃ or H₂O based ALD at 300°C.
Figure 7. Capacitance-Voltage characteristics as a function of frequency (a) 4 nm HfO$_2$ (HfCl$_4$/H$_2$O) on 1 nm GeO$_2$ (20 min O$_3$ at 300°C); (b) 4 nm HfO$_2$ (HfCl$_4$/O$_3$) on 1 nm GeO$_2$ (20 min O$_3$ at 300°C); (c) 4 nm Al$_2$O$_3$ (TMA/O$_3$) on HF cleaned Ge; (d) 4 nm Al$_2$O$_3$ (TMA/O$_3$) on 1 nm GeO$_2$ (20 min O$_3$ at 300°C).

TABLE 2. Summary of EOT for capacitors with HfO$_2$ and Al$_2$O$_3$ dielectrics.

<table>
<thead>
<tr>
<th>Starting surface</th>
<th>ALD</th>
<th>Temperature (°C)</th>
<th>Estimated EOT (nm)</th>
<th>Measured EOT (nm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>GeO$_2$ 20 min O$_3$</td>
<td>HfCl$_4$/O$_3$</td>
<td>300</td>
<td>1.9</td>
<td>2.0</td>
</tr>
<tr>
<td>GeO$_2$ 20 min O$_3$</td>
<td>HfCl$_4$/H$_2$O</td>
<td>300</td>
<td>1.4</td>
<td>1.5</td>
</tr>
<tr>
<td>HF cleaned Ge</td>
<td>TMA/O$_3$</td>
<td>300</td>
<td>2.2</td>
<td>2.4</td>
</tr>
<tr>
<td>GeO$_2$ 20 sec O$_3$</td>
<td>TMA/O$_3$</td>
<td>300</td>
<td>2.3</td>
<td>2.4</td>
</tr>
<tr>
<td>GeO$_2$ 20 min O$_3$</td>
<td>TMA/O$_3$</td>
<td>300</td>
<td>2.3</td>
<td>2.6</td>
</tr>
</tbody>
</table>

The TMA/O$_3$ process demonstrates the feasibility of obtaining a thin germanium oxide interfacial layer while keeping good CV characteristics. The EOT slightly decreases when the Ge oxidation time before ALD is reduced, due to a thinner germanium oxide layer (XPS). The CV characteristics are well behaved even for TMA/O$_3$ ALD directly on HF cleaned Ge (and starting the ALD with a TMA pulse) (Figure 7c). In this case, the interfacial oxide layer is only 0.7 nm, with Ge in oxidation states 4+ and lower. The CV characteristics are much better than those observed previously for H$_2$O based ALD on HF cleaned Ge, where large frequency dispersion and even flat CV characteristics were reported (2). As compared to the capacitors with ALD on 1 nm GeO$_2$ (Figure 7a, b, d), the CV curves are slightly more stretched out. Still, it is possible to achieve reasonable passivation with a thin germanium oxide interfacial layer where Ge oxidation states lower that 4+ are present.
Interface state densities of $1 - 3 \times 10^{12} \text{ cm}^{-2} \text{eV}^{-1}$ were obtained for all devices shown in Figure 7 even without post deposition treatments, using the low-frequency or quasistatic method (33). Further electrical characterization is necessary in order to investigate the relation between interface state density and the Ge oxidation states, germanium oxide thickness or intermixing. Whether or not scaling down the thickness of GeO$_2$ underneath HfO$_2$ is feasible will be the subject of further investigation. Theoretical calculations indicate that the formation of a Ge–Hf bonds at the interface, likely present if Hf is located in the sub-oxide interfacial layer (GeO$_x$ with $x < 2$), results in the formation of a defect level in the upper part of the Ge energy band-gap, degrading the electrical properties of metal-oxide-semiconductor devices (28).

HfCl$_4$/H$_2$O ALD on 1 nm GeO$_2$ (Ge substrates) results in a similar leakage current as on 1 nm SiO$_2$ (Si substrates), as shown in Figure 8 that compares the leakage current – EOT behavior of the Ge devices with that of similar devices on Si substrates. The HfCl$_4$/O$_3$ ALD on Ge gives a higher EOT due to the thicker GeO$_2$ layer. Still, the leakage current for HfCl$_4$/O$_3$ and HfCl$_4$/H$_2$O ALD is similar. Under gate injection, transport through the high-$\kappa$ layer limits the current, and is expected to depend mainly on the physical thickness of the high-$\kappa$ layer. The Al$_2$O$_3$ stacks also give a comparable leakage current at an even higher EOT due to the lower $\kappa$-value of Al$_2$O$_3$.

Figure 8. Leakage current as a function of EOT for Pt gated capacitors with 4 nm high-$\kappa$ dielectrics on Ge substrates, and 2-5 nm of HfO$_2$ ALD on Si substrates (HfCl$_4$/H$_2$O on 1 nm SiO$_2$).

The hysteresis in the CV characteristics is frequency dependent and varies from 70 mV to 250 mV for HfO$_2$ layers with and without pre-oxidation, as well as the Al$_2$O$_3$ layer without pre-oxidation. For the Al$_2$O$_3$ layer with O$_3$ pre-oxidation, the observed hysteresis varies less, from 70 mV to 100 mV. The hysteresis values are significantly lower than those reported previously for transistors with Ge/GeO$_2$/HfO$_2$/TiN gate stacks, where process steps at higher temperatures were included (12).

Effect of ALD precursors on the band gap of GeO$_2$

Internal photoemission indicates that the oxidant used in the HfO$_2$ ALD affects the band gap of the GeO$_2$ layer. For HfCl$_4$/O$_3$ on GeO$_2$, the band gap measured for GeO$_2$ is close to the value reported for stoichiometric GeO$_2$ (5.5 eV). This was also demonstrated
previously for HfCl₄/O₃ at 375°C on Ge substrates (34) and was confirmed by our measurements for HfCl₄/O₃ at 300°C on GeO₂. On the other hand, the band gap of GeO₂ layer is reduced to 4.3 eV when HfO₂ is deposited on GeO₂ by means of H₂O based ALD (35), even for GeO₂ layers with a thickness of more than 2 nm. It was suggested that hydroxyl groups are incorporated in the GeO₂ layer during H₂O based ALD, and that the incorporation of OH in GeO₂ lowers the value of the band gap. Thus, whereas the H₂O based process has the advantage of no additional Ge oxidation during ALD, the O₃ based process has the advantage of preserving the value of the GeO₂ band gap.

ALD on GaAs substrates

The oxidant precursor also plays an important role in the formation of the interfacial layer on GaAs substrates. For the thermal process, XPS indicates no interfacial layer on GaAs for both HfO₂ and Al₂O₃ ALD (Figure 9 for Al₂O₃ ALD and reference 36 for HfO₂ ALD). The As₃d oxide peak at binding energy of 43 – 46 eV decreases as a function of the Al₂O₃ thickness, and for Al₂O₃ layers of ~2 nm (60-70 Al/nm²), no As₂O₅ or As₂O₃ were detected. So, the AsOₓ-GaOₓ layers that were present before ALD are removed during H₂O based ALD at 300°C, at least for the HfCl₄/H₂O and TMA/H₂O ALD. Thinning of the initial oxide has also been reported previously for Al₂O₃ ALD (TMA/H₂O at 300°C) on GaAs and InGaAs (16, 17). On the other hand, the interfacial oxide thickness increased during PEALD using TMA and O₂ plasma (Figure 9). The XPS As₃d and Ga₃d spectra indicate that As₂O₅, As₂O₃, as well as Ga₂O₃ are present.

As and Ga partly intermix with the Al₂O₃ layers (Figure 10), similar as for ALD on GeO₂ substrates. Both the H₂O and O₂ plasma based Al₂O₃ ALD show a similar extent of intermixing. On the other hand, the HfCl₄/H₂O ALD of HfO₂ on GaAs shows more abrupt interfaces, again in correspondence with ALD on GeO₂ (see above). These trends are observed for deposition on both native oxide and HCl cleaned GaAs. Although the reaction mechanisms accounting for incorporation of GeO₂, GaOₓ and/or AsOₓ are unclear, these results suggest that there are correlations in the reaction mechanisms on Ge and GaAs substrates.

![Figure 9](image_url)  
Figure 9. XPS AsOₓ thickness as a function of Al-content for thermal and PEALD Al₂O₃ on GaAs with different surface preparations.
Preliminary device measurements on GaAs/high-κ/Pt capacitors indicate that the oxide interfacial layer has an impact on the device characteristics: for the thermal ALD process, less frequency dispersion, less hysteresis and higher capacitance are observed, probably linked to the lower content of As$_2$O$_3$, As$_2$O$_5$ and Ga$_2$O$_3$ (37). Nevertheless, all interfaces (using both ALD and PEALD) are still Fermi level pinned, as discussed in (38) for Al$_2$O$_3$ deposited by thermal ALD, and in (37) for both thermal and plasma enhanced ALD. Together with the investigation of the ALD precursors, the effect of surface preparation before the ALD and post-deposition treatments should be further investigated.

**Conclusions**

The ALD precursors play an important role in the characteristics of both Ge and III-V MOS devices. Passivation of interface defects using thermally grown GeO$_2$ is a promising approach. We demonstrated that the thickness, oxidation states of Ge, and the amount of intermixing of germanium oxide with the high-κ dielectric depend on the ALD precursors and process conditions.

H$_2$O based ALD has the advantage of no additional Ge oxidation during ALD but internal photoemission indicates a low band gap of germanium oxide (4.3 eV), supposedly by hydroxyl incorporation during H$_2$O based ALD. On the other hand, HfCl$_4$/O$_3$ ALD at 300°C results in a rather thick (1.7 nm) GeO$_2$ interfacial layer. Oxidation of Ge can be minimized in ALD processes with shorter O$_3$ pulse times, for example TMA/O$_3$, and by reducing the ALD temperature. As such, the germanium oxide thickness was scaled down below 1 nm. However, such thin germanium oxide layers contain Ge in oxidation states lower than 4+. Still, electrical results indicate that small amounts of Ge in oxidation states lower than 4+ are not detrimental for device performance.

Partial intermixing was observed for high-κ ALD on both GeO$_2$ and GaAsO$_x$, suggesting possible correlations in the ALD growth mechanisms on Ge and GaAs substrates. Obtaining good device characteristics with ALD high-κ dielectrics on GaAs substrates remains nevertheless challenging. Together with the investigation of the
influence of the applied ALD precursors, the effect of surface preparation before the ALD and post-deposition treatments should be further investigated.

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References