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Effects of Downstream Plasma Exposure on β-Ga_2O_3 Rectifiers

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The effects of downstream plasma exposure with O_2, N_2 or CF_4 discharges on Si-doped Ga_2O_3 Schottky diode forward and reverse current-voltage characteristics were investigated. The samples were exposed to discharges with rf power of 50 W plasma at a pressure of 400 mTorr and a fixed treatment time of 1 min to simulate dielectric layer removal, photoresist ashing or surface cleaning steps. Schottky contacts were deposited through a shadow mask after exposure to avoid any changes to the surface. A Schottky barrier height of 1.1 eV was obtained for the reference sample without plasma treatment, with an ideality factor of 1.0. The diodes exposed to CF_4 showed a 0.25 V shift from the I–V of the reference sample due to a Schottky barrier height lowering around 14%. The diodes showed a decrease of Schottky barrier height of 2.5 and 6.5% with O_2 or N_2 treatments, respectively. The effect of plasma exposure on the ideality factor of diodes treated with these plasmas was minimal; 0.2% for O_2 and N_2, 0.3% for CF_4, respectively. The reverse leakage currents were 1.2, 2.2 and 4.8 μA cm^{-2} for the diodes treated with O_2, and CF_4, and N_2 respectively. The effect of downstream plasma treatment on diode on-resistance and on-off ratio were also minimal. The changes observed are much less than caused by exposure to hydrogen-containing plasmas and indicate that downstream plasma stripping of films from Ga_2O_3 during device processing is a relatively benign approach.
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The ultra-wide bandgap semiconductor β-Ga_2O_3 has attracted increasing attention because of the prospects for use in next generation high-power electronics.[1–4] The high breakdown field of this material has enabled numerous demonstrations of vertical rectifiers with breakdown voltages in the kV range.[5–22] This is of interest for electric vehicle (EV) charging stations, power management in residential solar systems and battery energy storage systems. This high breakdown field of Ga_2O_3 enables greater voltage blocking capability and lower conduction losses due to a lower on-resistance reduction in comparison with Si-based devices.[1–4]

One important factor that needs to be considered when forming devices using β- Ga_2O_3 is the electronic behavior at the surface or interfaces. The surface termination, relaxation and surface energies for different faces of β-Ga_2O_3 have been reported by Bermudez.[23] However, it is not widely appreciated that the surface of β-Ga_2O_3 can be strongly affected by exposure to gaseous or plasma environments and the influence of changing conductivity and roles of surface states in oxidizing or reducing environments are not well-established.[24–37] This is despite the fact that Ga_2O_3 rectifiers are known to be sensitive detectors of hydrogen.[25,26] The (−201) Ga_2O_3 surface has been reported to be particularly sensitive to plasma-induced damage, leading to device performance degradation.[33]

In our study, we have employed truly downstream plasma exposures of Ga_2O_3 to CF_4, N_2 and O_2 to separate out chemical effects from physical damage due to ion bombardment and also to simulate processes like photoresist ashing, dielectric removal and surface cleaning that occur during device fabrication.[29,34] We find that only CF_4 exposure leads to significant changes under these conditions. This work complements previous studies where the Ga_2O_3 was exposed to these plasmas in immersive mode, where combined chemical and physical bombardment effects are present.[24,28,29]

Experimental

The starting samples consisted of vertical rectifier structures. The drift region of the material consisted of a 10 μm thick, lightly Si doped epitaxial layer grown by halide vapor phase epitaxy (HVPE) with carrier concentration of 3×10^{16} cm^{-3}, and this epitaxial layer was grown on a (001) surface orientation Sn-doped ($n = 10^{19}$ cm^{-3}) β-Ga_2O_3 single crystal (Novel Crystal Technology, Japan). The wafer surfaces were ultrasonically cleaned in acetone, methanol, and isopropyl alcohol prior to all experiments. The Fermi level is found to be relatively unpinned on the bulk β-Ga_2O_3 (001) substrate, suggesting the presence of lower density of oxygen vacancy states on its surface.[37]

A full area Ti/Au backside Ohmic contact was formed by e-beam evaporation and was annealed at 550 °C for 30 s under N_2 ambient. After backside Ohmic formation, the front of the sample was cleaned using HCl and then treated with ozone for 20 min to remove residual hydrocarbons. Previous reports have indicated that untreated substrates contain a significant amount of adsorbed carbon contaminations at the surface, which can be partly removed by annealing at 800 °C in UHV. In that uncleaned state, upward band bending of about 0.5 eV that increases with annealing is present, leading to an electron depletion layer at the near-surface region.[27] This effect could be removed either by annealing in oxygen at high temperatures or by chemical cleaning/etching of the surface prior to deposition of metal contacts.[27]

The plasma treatments were performed using a downstream PIE Scientific Tergeo Plasma Cleaner with O_2, N_2 or CF_4 discharges. The plasma was generated with a 13.56 MHz high frequency rf power supply with automatic impedance matching for the in situ plasma source. The RF power source could operate over the range 0–150 watt, and in our case we used 50 W to generate the plasma at a pressure of 400 mTorr, with a fixed treatment time of 1 min. The system can be operated either in immersion mode (samples are immersed in plasma) or downstream mode (samples are placed outside the plasma) and we used the latter in all cases. In the downstream mode, there is no physical bombardment of the sample surface by energetic ions that would occur in the immersion mode. However, there can still be chemically-induced changes to the near-surface in the downstream mode that affect properties such as Schottky barrier height. In addition to the ion bombardment effects in the immersion mode, there can be synergistic effects due to the combined ion and reactive neutral components. This is absent in the downstream mode.

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Next, the front of the samples was treated with ozone for 15 min, followed by cleaning with 1:10 diluted HCl, then was treated with ozone for another 15 min to remove surface contamination. the Schottky contacts were formed by e-beam evaporation. Ni/Au Schottky metallization was deposited through a shadow metal rather than photoresist patterning to avoid changing the chemistry of the Ga_2O_3 surface. Edge termination was not used, also in order to focus on the surface characteristics free of any edge effects.[9] The completed devices had circular contact diameters of 800 μm.

The current-voltage (I–V) characteristics were recorded at room temperature. Forward and reverse current measurements were recorded with a HP 4156 parameter analyzer. The forward direction was dominated by the thermionic emission (TE) current over most of the temperature range, while in the reverse direction, the thermionic field emission (TFE) and tunneling currents played an important role at high reverse bias. To extract the zero-bias equivalent barrier height (Φ_b) and ideality factor (n), we used the relationship for current density in TE theory and the linear portion of the forward bias characteristics through the correction factor $_{[eE/4\pi\varepsilon]}0^{.5}$, where E is the electric field at the Ni/Au/Ga_2O_3 interface and ε is the dielectric constant of the semiconductor,[37] i.e. $J = J_0 \exp(eV_A/nkT)$ [1-exp $(-eV_A/kT)$ where $J_0 = A^* m_{eff}/m_0 T^2 \exp(\Phi_B/kT)$, e is electronic charge and A^* is the Richardson constant (33.7 $A.cm^{-2}K^{-2}$) and V_A is the bias voltage applied. The values of barrier height were corrected for the image force (IF) lowering, as described elsewhere.[38] Capacitance-Voltage (C–V) characteristics were recorded with an Agilent 4284 A Precision LCR Meter to conform the carrier concentration in the epi layer.

The forward turn-on voltage, V_F, for a Schottky rectifier is given by

$$V_F = \frac{nkT}{e}\ln\left(\frac{J_F}{A^{**}T^2}\right) + n\Phi_B + R_{ON}\cdot J_F$$

where R_{ON} is the on-state resistance. We defined V_F as the bias at which the forward current density is 100 $A\cdot cm^{-2}$.The diode on/off ratio is another figure-of-merit and was measured when switching from 1 V forward to reverse biases up to 100 V.

Results and Discussion

The I–V results before and after downstream plasma exposure are shown in Fig. 1 on both log (a) and linear (b) scales. Near-ideal Schottky characteristics with n values of 1.0 were obtained for the reference diodes, with a barrier height of 1.1 eV. We used I–V data rather than capacitance—voltage data because nonlinear doping profiles can affect the latter. From the forward I–V characteristic, as illustrated in Fig. 1, the diodes exposed to CF_4 showed a 0.25 V shift from the I–V of the reference sample due to a Schottky barrier height lowering around 14%, as shown in Table I. This can result from the change in barrier height from the relation above for V_F. Previous work has shown that under partially immersive CF_4 plasma exposure conditions, high concentrations of fluorine are incorporated into the near-surface region of Ga_2O_3 and that it remains in the material to temperatures beyond 400 °C.[24] It is clear that compensation of Si donors by F^- ions occurs in Ga_2O_3, leading to changes in effective barrier height.[24]

The diodes exposed to downstream N_2 and O_2 plasmas showed small decreases of Schottky barrier height of 2.5 and 6.5% with O_2 or N_2 treatments, respectively, as shown in Fig. 2. It has been previously reported that Ga_2O_3 displays electron accumulation when the surface is terminated by O–H groups,[27] resulting in downward band bending. After removal of the hydrogen by surface cleaning, the direction of this band bending is reversed and the surface displays electron depletion. This has been explained by the charge neutrality level, which is found to be 0.6 eV below the conduction band minimum.[27] This is supported by the determination of the charge state transition level for *H* interstitials, which do not disrupt the bonding on the Ga_2O_3 surface, but are captured by O lone-pairs

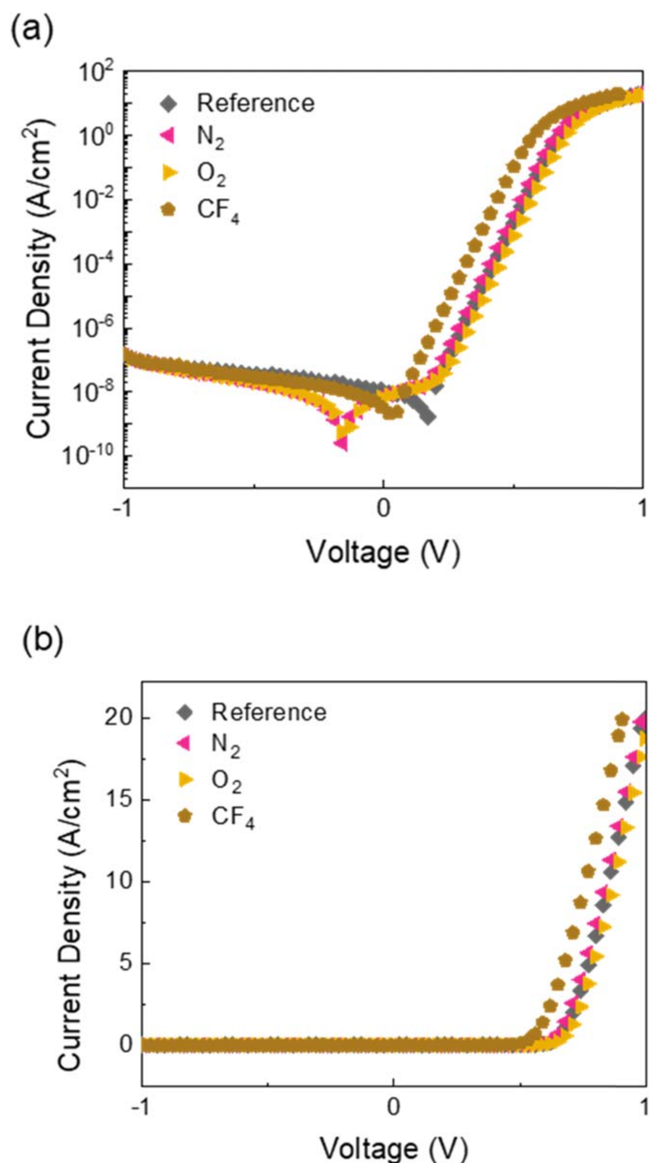

Figure 1. Forward I-V characteristic from −1 to 1 V for samples exposed to different plasmas shown in (a) log or (b) linear scale.

on the surface, forming shallow donors.[27,36,39,40] However, in our case, it is clear that the surface is sufficiently stable after downstream plasma exposures to O_2 that subsequent exposure to air does not change the barrier height to the level exhibited by O–H bonding. The effect of plasma exposure on the ideality factor of diodes treated with these plasmas was minimal; 0.2% for O_2 and N_2, 0.3% for CF_4, respectively, as also shown in Fig. 2.

The incorporation of F- in Ga_2O_3 under immersive plasma conditions was previously found to lead to an increase of the barrier height.[28] However, in this current work with remote plasmas, the data shows a small decrease of the barrier height. Thus it is clear that the incorporation of F is assisted by the ion bombardment present in immersive plasmas. This is plausible given the need for the F to diffuse enough to produce the compensation effect of Si donors by F- ions. The ion bombardment can produce enough near-surface point defects to assist this diffusion.In the case of downstream plasma, the F compensation effect is absent and the changes are due to subtle near-surface changes in the chemistry of the upper layers

The reverse leakage currents were 1.2, 2.2 and 4.8 $\mu A\ cm^{-2}$ for the diodes treated with O_2, and CF_4, and N_2 respectively, as shown in Fig. 3. The respective reverse breakdown voltages (V_B) are shown

Table I. Summary of diode characteristics as a function of plasma treatments.

	Schottky barrier height (eV)	Ideality factor	R_{ON} (mΩ.cm^2)	V_B (V)	AFM FWHM (nm)	Ga/O ratio
Reference	1.1	1.00	28.6	135	3.5	0.40
N_2 plasma	1.04	1.02	29.6	105	3.5	0.40
O_2 plasma	1.08	1.01	29.1	130	3.6	0.40
CF_4 plasma	0.96	1.04	30.5	105	3.5	0.42

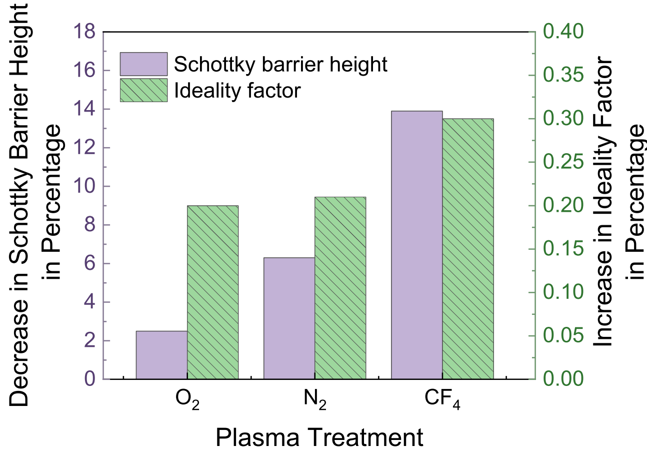

Figure 2. Percentage change of Schottky barrier height and ideality factor for the Ga_2O_3 diodes exposed to O_2, N_2 or CF_4 plasma.

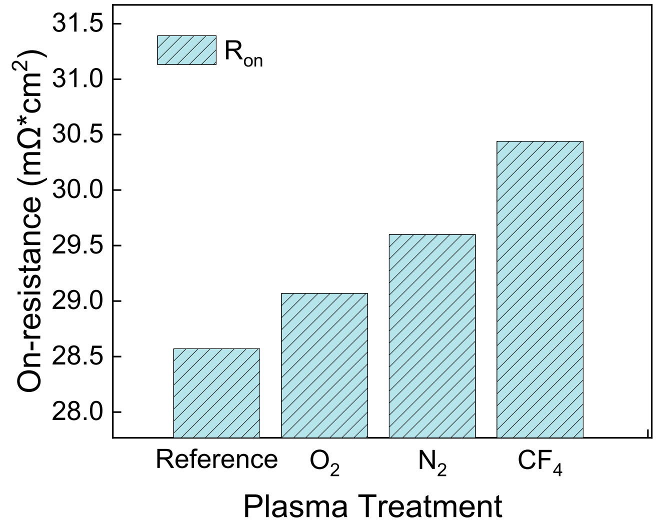

Figure 4. On-resistance for samples exposed to different plasmas.

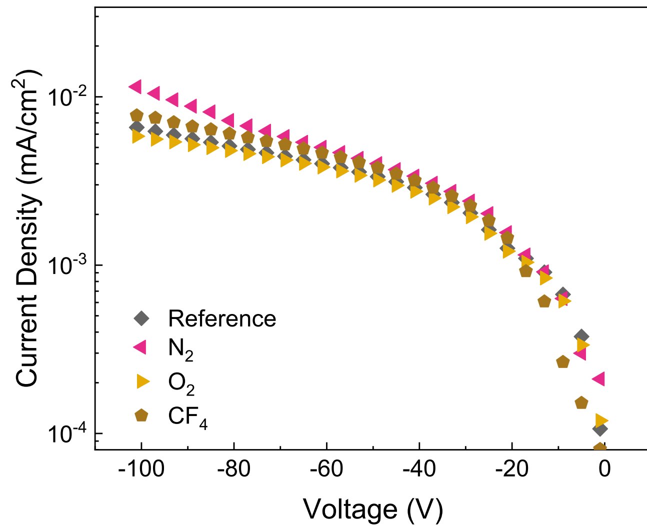

Figure 3. Reverse I–V characteristic from 0 to −100V for samples exposed to different plasmas.

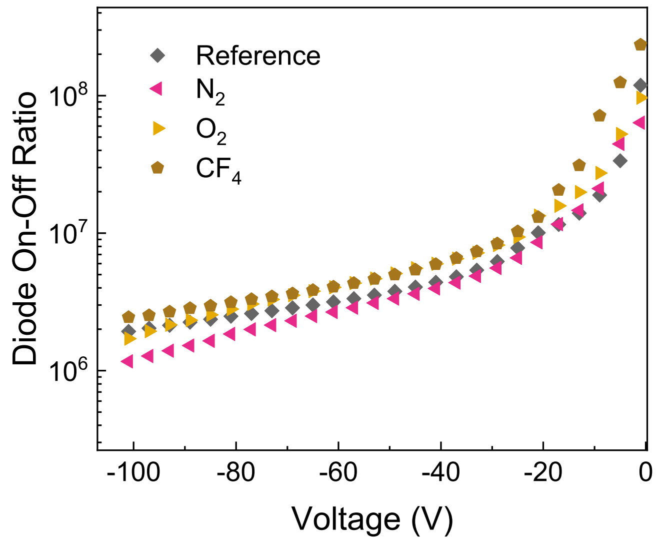

Figure 5. On/off ratio for samples exposed to different plasmas.

in Table I, with only minor changes as a result of the downstream plasma exposure. As expected, this indicates that the number of midgap states leading to recombination or tunneling currents is not significantly changed as a result of the downstream plasma exposure or else we would observe increases in the reverse current density. The absence of ion bombardment is clearly the main factor. Notice also in Table I that the surface roughness as measured by Atomic Force Microscopy (AFM) over an area of 5 x 5 μm^2, did not show any significant change. The near-surface Ga/O ratio averaged over a depth of ~100 Å by X-ray Photoelectron Spectroscopy did not change, as expected. It would be difficult to correlate these measurements to the change in electrical properties, since the latter are dominated by the first few layers of the surface.

The effects of downstream plasma treatment on diode on-resistance were also minimal as shown in Fig. 4, where the increases are <10% for all the different plasma chemistries. Since

$$R_{diode} = R_{drift} + R_{sub} + R_{contact}$$

where specific on-state resistance of a unipolar diode is a sum of the drift region resistance R_{drfit}, the contact resistance $R_{contact}$ and the substrate resistance. R_{sub},[1] this means that the latter hasn't changed much during exposure to the fluorine, oxygen or nitrogen- based discharges. A similar conclusion can be drawn from the diode on/off ratio data in Fig. 5, where the changes are small (<10%). The on-off ratio is another figure of merit in that having high on-current and low

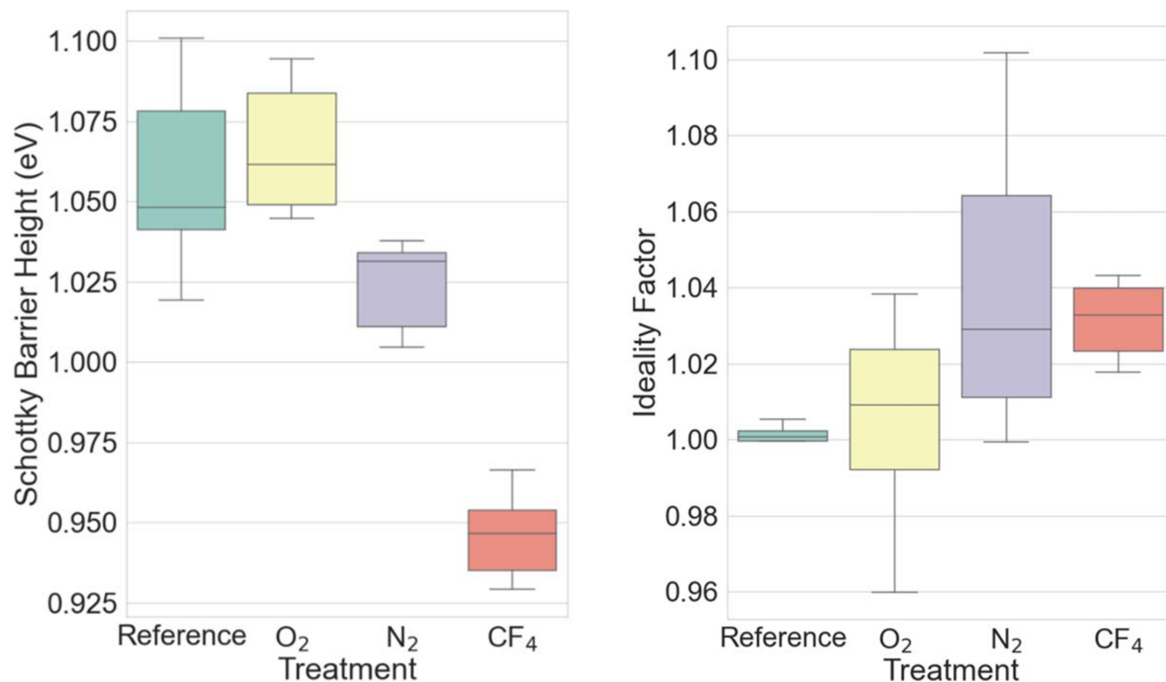

Figure 6. Schottky barrier height and ideality factor after different plasma treatments plotted as the mean of 6 different measurements (n = 6).

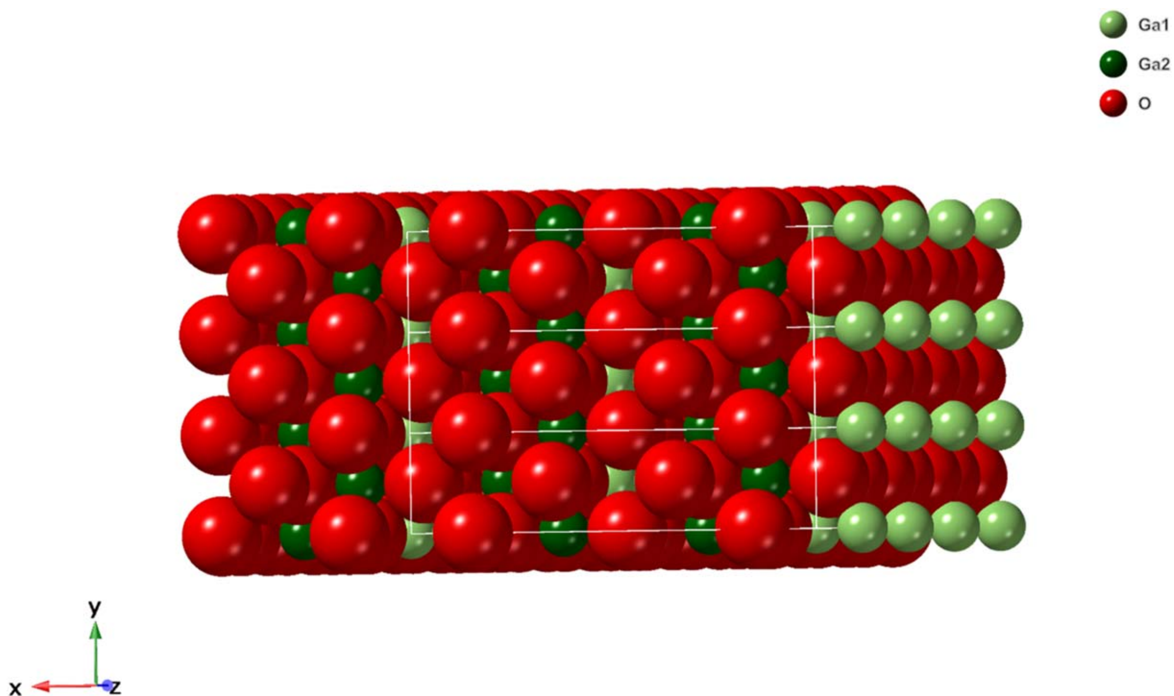

Figure 7. Schematic of β Ga_2O_3 crystal structure with the (001) direction being vertical to the top surface.[37]

leakage current in reverse bias is desirable. This was $>10^5$ for all devices measure, independent of plasma exposure.

Finally, to give some idea of variability across samples, the range of barrier heights and ideality factors measured from 6 different devices on each chip are shown in Fig. 6. Within the experimental error, only samples exposed to CF_4 show significant changes in these two parameters.

The crystal structure of β Ga_2O_3 consists of double octahedral chains, running parallel to the crystallographic *y*-axis.[41] The chains are cross-linked by tetrahedral GaO_4 groups. Figure 7 shows the crystal structure for the (001) orientation used here,[37] where the surface consists mostly of oxygen termination. In terms of intrinsic defects in n-type Ga_2O_3, the reported high formation energy of oxygen vacancies (V_O) suggests they are deep donors that are not

ionized in n-type material and this will not contribute to the conductivity.[28,36] The Ga vacancy (V_{Ga}) is expected to be a triple acceptor, while the oxygen interstitial (Oi) is neutral, and the gallium interstitial (Gai) are in the 3 + charge state.[28,36] The V_{Ga} concentration increases with oxygen partial pressure, leading to a conductivity compensation. What is clear from our data is that this O-terminated surface is stable against air and O_2 exposure, but there are small changes with fluorine exposure that are smaller than in the case of performing the CF_4 exposure in an immersive plasma environment. The ion bombardment in that case enhance the chemical effects of the fluorine.

Summary and Conclusions

There are numerous cases during Ga_2O_3 device fabrication where plasmas are used in resist stripping, surface cleaning or dielectric layer removal. Our results show that purely downstream plasmas produce only small changes in surface properties. There are still some unanswered questions as to what effect the crystal orientation has, since the surface termination is different in different crystal directions and it is known that the bulk properties of Ga_2O_3 are highly anisotropic. Similarly, what is the effect of polytype, since the α polytype is promising due to its even larger bandgap than for β Ga_2O_3. Lastly, what effect will alloying with Al and In have when those $(Al_xGa_{1-x})_2O_3$ and $(In_xGa_{1-x})_2O_3$ alloys are used in device structures ? It would be expected that the Al-based alloys will be more sensitive to both oxygen and fluorine adsorption effects.

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Data Availability Statement

All data that support the findings of this study are included within the article (and any supplementary files)

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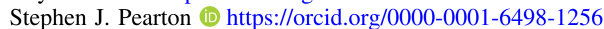
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