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Recent progress in Ga₂O₃ power devices

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Abstract

This is a review article on the current status and future prospects of the research and development on gallium oxide (Ga_2O_3) power devices. Ga_2O_3 possesses excellent material properties, in particular for power device applications. It is also attractive from an industrial viewpoint since large-size, high-quality wafers can be manufactured from a single-crystal bulk synthesized by melt–growth methods. These two features have drawn much attention to Ga_2O_3 as a new wide bandgap semiconductor following SiC and GaN. In this review, we describe the recent progress in the research and development on fundamental technologies of Ga_2O_3 devices, covering single-crystal bulk and wafer production, homoepitaxial thin film growth by molecular beam epitaxy and halide vapor phase epitaxy, as well as device processing and characterization of metal–semiconductor field-effect transistors, metal–oxide–semiconductor field-effect transistors and Schottky barrier diodes.

Keywords: gallium oxide, Ga_2O_3 , power devices, edge-defined film-fed growth, halide vapor phase epitaxy, molecular beam epitaxy

(Some figures may appear in colour only in the online journal)

1. Introduction

SiC and GaN power devices have attracted much attention as key components of the high-efficiency power conversion required for energy saving in the near future. Their device performance can far exceed that of the Si-based devices mainly used in current power electronics. However, SiC and GaN do not have to be the only candidates for next-generation power devices in the near future, because the materials and devices with the best performance and cost advantage should be chosen for society. Against this background, we have been proposing a new oxide compound semiconductor, gallium oxide (Ga₂O₃), as another promising candidate because of its excellent material properties and suitability for mass production.

The bandgap (E_g) and the expected Baliga's figure of merit (FOM) of Ga₂O₃ are much larger than those of SiC and GaN. These outstanding material properties will in principle enable Ga₂O₃ power devices with even higher breakdown

voltage (V_{br}) and efficiency than their SiC and GaN counterparts.

SiC wafers are produced from bulk ingots typically grown using a sublimation method at extremely high temperatures above 2000°C. Various growth methods are being used for GaN bulk and/or wafer production, such as the high nitrogen pressure method, the ammonothermal technique and hydride vapor phase epitaxy; however, it is an open question which technique will dominate the market. As mentioned above, SiC and GaN wafers are produced from bulks synthesized by growth methods other than melt growth, because no atmospheric melt growth methods, which are standard for the production of semiconductor single-crystal bulks to date, are applicable for both SiC and GaN. However, in the case of Ga₂O₃, a single-crystal bulk can be synthesized using several standard melt growth methods; therefore, this is a strong advantage for Ga₂O₃ over SiC and GaN, in particular for mass production.

This review article mainly summarizes the recent progress in research and development (R&D) activities on this



Figure 1. Atomic unit cell of β -Ga₂O₃.

emerging semiconductor Ga_2O_3 power device technology. Details about the above two attractive features of Ga_2O_3 will be introduced in the following sections. At present, the device technology is still at a primitive stage; however, the developed transistors and diodes with simple structures have verified its great potential for power devices.

2. Crystal structures of Ga₂O₃

The Ga₂O₃ single crystal is known to show five polymorphs, labeled α , β , γ , δ and ϵ [1]. Among the crystal structures, Ga_2O_3 mainly crystallizes in the monoclinic β -gallia structure with lattice constants of a = 12.2 Å, b = 3.0 Å and c = 5.8 Å as shown in figure 1. The β phase is the most stable structure, and the other phases are meta-stable ones. Note that β -Ga₂O₃ is the only crystal structure that can be grown from the melt. Most of the reported scientific studies have been on the crystal growth and material properties of β -Ga₂O₃. The second metastable polymorph is α -Ga₂O₃ with a corundum structure, which is often stabilized by low-temperature heteroepitaxial growth on sapphire substrates [2, 3]. The heteroepitaxial growth of γ -Ga₂O₃ thin films, which has a defective-spinel structure, has also been reported [4, 5]. Formation of the meta-stable crystalline phases is strongly dependent on the substrate lattice and growth temperature. Typically, high growth temperatures lead to the formation of β -Ga₂O₃ even in the case of heteroepitaxial growth on foreign substrates. In this review, we focus on β -Ga₂O₃.

3. Material properties of β -Ga₂O₃

Some of the basic material parameters of β -Ga₂O₃ have not been fully studied and understood. There is great variability in

the reported E_g values, between 4.5 and 4.9 eV [6–9]. The *n*type doping technology is relatively easy and mostly established. Sn and Si atoms are known to be shallow donors with small activation energies in Ga_2O_3 . Electron densities (*n*) can be controlled in the wide range of 10^{15} - 10^{19} cm⁻³ [7, 8, 10, 11]. In contrast, there has been no report on effective hole conduction in Ga_2O_3 . We consider that it is difficult to find acceptor atoms with a small activation energy since Ga₂O₃ is a wide bandgap oxide semiconductor. Furthermore, a very low hole mobility (μ) prohibiting effective ptype conductivity in Ga₂O₃ was expected from the firstprinciples calculation of the Ga_2O_3 band structure [12–14]. Even in the worst case, tight localization of holes at specific sites in a Ga_2O_3 bulk was also predicted [15]. The thermal conductivity of Ga₂O₃ strongly depends on the crystal orientation due to its asymmetric crystal structure as shown in figure 1. The highest thermal conductivity of $0.27 \text{ W} (\text{cm K})^{-1}$ can be obtained in the [010] direction, which is more than twice as large as that in the [100] direction [16]. However, even the [010] thermal conductivity of Ga₂O₃ is still much smaller than those of other semiconductors.

4. Advantages of Ga_2O_3 power devices from the material point of view

Here, we discuss why there is a real possibility that Ga₂O₃based transistors and diodes can have excellent power device characteristics such as high V_{br}, high power, and low loss due to its material properties. Table 1 summarizes the physical properties of β -Ga₂O₃ and other major semiconductors for power devices. The Ga₂O₃ bulk electron μ of 300 cm² Vs⁻¹ was estimated on the basis of the experimental data for the typical *n* range of 10^{15} - 10^{16} cm⁻³ for the drift layers of vertical power devices [11, 17]. From the interpolation of the relationships among the E_g and breakdown electric fields (E_{br}) of the other semiconductors, the E_{br} of Ga_2O_3 is expected to have a very large value of about 8 MV cm⁻¹ [17]. The high $E_{\rm br}$ is the most attractive property of Ga₂O₃, because the Baliga's FOM, which is the basic parameter to show how suitable a material is for power devices, is proportional to the cube of $E_{\rm br}$, but is only linearly proportional to μ . The fundamental limits of on-resistances (R_{on}) as a function of V_{br} for Ga₂O₃ and representative semiconductors are plotted in figure 2. It suggests that the conduction loss of Ga_2O_3 devices can be one order of magnitude lower than those of SiC and GaN devices at the same $V_{\rm br}$. Therefore, we consider that Ga_2O_3 will best show its potential in unipolar devices.

5. Single-crystal Ga₂O₃ bulks and wafers

In general, large-diameter single-crystal wafers are required to mass produce vertical devices that are favorable for high-voltage and high-current power devices. Single-crystal Ga₂O₃ bulks have been synthesized by several melt growth methods such as the Czochralski [18, 19], the floating-zone (FZ)

Table 1. Material properties of major semiconductors and β -Ga₂O₃.

	Si	GaAs	4H-SiC	GaN	Diamond	β -Ga ₂ O ₃
Bandgap $E_{\rm g}$ (eV)	1.1	1.4	3.3	3.4	5.5	4.5-4.9
Electron mobility μ (cm ² Vs ⁻¹)	1,400	8,000	1,000	1,200	2,000	300
Breakdown field $E_{\rm br}$ (MV cm ⁻¹)	0.3	0.4	2.5	3.3	10	8
Relative dielectric constant ϵ	11.8	12.9	9.7	9.0	5.5	10
Baliga's FOM $\epsilon \mu E_{\rm br}^3$	1	15	340	870	24,664	3,444
Thermal conductivity (W (cm K) $^{-1}$)	1.5	0.55	2.7	2.1	10	0.27 [010]
						0.11 [100]



Figure 2. Theoretical unipolar performance limits of R_{on} as a function of V_{br} for β -Ga₂O₃ and other major semiconductors power devices. Reproduced with permission from [17]. Copyright 2012, AIP Publishing LLC.

[7, 20, 21] and the edge-defined film-fed growth (EFG) [22] methods. In contrast, SiC and GaN bulk crystals are grown using methods such as sublimation, vapor phase epitaxy and high-pressure synthesis other than melt growth methods [23– 25]. We propose that EFG has an advantage against the other melt growth methods regarding the bulk size, which is a key factor for high-volume production of large-size single-crystal Ga_2O_3 wafers. Figure 3 shows a photograph of a 4 inch diameter single-crystal Ga2O3 wafer produced from an EFGgrown Ga₂O₃ bulk. The crystal quality of Ga₂O₃ wafers prepared from EFG bulks has already been very good, with a full-width at half-maximum of the x-ray diffraction rocking curve (XRC) as narrow as 17 arcsec and a dislocation density on the order of 10^3 – 10^4 cm⁻² as characterized by surface etch pits. The wafer surface was atomically flat and smooth after chemical-mechanical polishing (CMP) with a small rootmean-square (rms) surface roughness of 0.11 nm. The good material workability of Ga₂O₃ is another important feature. Undoped Ga_2O_3 bulks show *n*-type conductivity due to unintentional Si incorporation from the Ga₂O₃ powder source. Thus, at present, compensation doping with deep acceptors such as Mg and Fe is required for the production of semiinsulating bulk crystals and wafers.

a203 p 0a203 p 0a203 p 0a203 p 0a20	3 p.00203 p.00203 p
$\beta\text{-}\text{Ga}_2\text{O}_3\ \beta\text{-}\text{Ga}_2\text{O}_3\ \beta\text{-}\text{Ga}_2\text{O}_3\ \beta\text{-}\text{Ga}_2\text{O}_3\ \beta\text{-}\text{Ga}_2\text{O}_3\ \beta\text{-}$	$Ga_2O_3 \beta - Ga_2O_3 \beta - Ga_2$
a2O3 β-Ga2O3 β-Ga2O3 β-Ga2O3 β-Ga2O	β -Ga ₂ O ₃ β -Ga ₂ O ₃ β
β-Ga2O3 β-Ga2O3 β-Ga2O3 β-Ga2O3 β-	Ga2O3 B-Ga2O3 B-Ga
$a_2O_3 \beta - Ga_2O_3 \beta - Ga_2O_3 \beta - Ga_2O_3 \beta - Ga_2O_3$	3 β-Ga2O3 β-Ga2O3
β-Ga2O3 β-Ga2O3 β-Ga2O3 β-Ga2O3 β-	Ga2O3 B-Ga2O3 B-Ga
a2O3 β-Ga2O3 β-Ga2O3 β-Ga2O3 β-Ga2	O ₃ β-Ga ₂ O ₃ β-Ga ₂ O ₃
β-Ga2Q3 β-Ga2O3 β-Ga2O3 β-Ga2O3 β-Ga2O3 β-	Ga2O3 B-Ga2O3 B-Ga
a ₂ O ₃ β-6a ₂ O ₃ β-Ga ₂ O ₃ β-Ga ₂ O ₃ β-Ga ₂ O	0 ₃ β-Ga ₂ O ₃ β-Ga ₂ O ₃ β
β-Ga2O β-Ga2O3 β-Ga2O3 β-Ga2O3 β-	Ga2O3 B-Ga2O3 B-Ga
a2O3 β-G2O3 β-Ga2O3 β-Ga2O3 β-Ga2O	$\beta_3 \beta - Ga_2O_7 \beta - Ga_2O_3$
β-Ga2O3 β-Ga2O3 β-Ga2O3 β-Ga2O3 β-	Ga2O3 B Ga2O3 B-Ga
a2O3 β-Ga2O3 β-Ga2O3 β-Ga2O3 β-Ga2O	$\beta_3 \beta - Ga_2O_3 \beta - Ga_2O_3$
β-Ga2O3 β-Ga2O3 β-Ga2O3 β-Ga2O3 β-	Ga2O3 B-Ga2O3 B-Ga
a2O3 β-usimm	un hara hara hara hara hara hara hara har
β-Ga ₂ O mm 1 2 3 4 5 6 7	8 9 10 11 1
a ₂ O ₃ β-Ga ₂ O ₃ β-Ga ₂ O ₃ β-Ga ₂ O ₃ β-Ga ₂ O	ο ₃ β-Ga ₂ O ₃ β-Ga ₂ O ₃

Figure 3. Photograph of 4 inch diameter single-crystal Ga₂O₃ wafer.

6. Epitaxial growth of Ga₂O₃ thin films

For the epitaxial growth of Ga_2O_3 thin films, there have been many reports on molecular beam epitaxy (MBE) [26–31], halide vapor phase epitaxy (HVPE) [32–35], metal–organic chemical vapor deposition [36–39] and mist chemical vapor deposition [2, 40]. In this section, the current status of our MBE and HVPE technologies for Ga_2O_3 thin film growth will be introduced.

6.1. MBE

At the time of writing this article, scientific studies on MBE growth have been the most advanced among the epitaxial growth techniques of Ga_2O_3 thin films. In general, normal gas-source MBE machines equipped with a turbo molecular pump are used for Ga_2O_3 growth. There are two types of oxygen sources used for the MBE growth of oxide semiconductors; one is oxygen radicals, generated by an radio-frequency (RF)-plasma cell [26–29], and the other is ozone [30, 31]. Our work on the ozone MBE growth of undoped and Sn-doped Ga_2O_3 films on native Ga_2O_3 (010) substrates will be described in the following [31].



Figure 4. (a) Surface AFM images and (b) their rms roughnesses of Ga_2O_3 epitaxial films grown at various temperatures by ozone MBE. Reproduced with permission from [31]. Copyright 2014, Elsevier.

The specification of our ozone MBE machine used for Ga_2O_3 thin film growth is as follows. Ga and Sn (*n*-type dopant) are, respectively, supplied from Ga metal with six nines purity and four nines SnO₂ powder heated in conventional Knudsen cells. The oxygen source is an ozone (5%)–oxygen (95%) gas mixture generated by an ozone generator. One of the noteworthy advantages of ozone over RF-plasma is its high growth rate, reaching a few μ m h⁻¹.

Figure 4(a) shows the surface morphologies of Ga_2O_3 epitaxial films grown at various temperatures, which were observed by atomic force microscopy (AFM). The rms surface roughnesses of the Ga_2O_3 epitaxial films are plotted as a function of growth temperature (T_g) in figure 4(b). As shown in figure 4(a), step bunching along the [100] direction is observed for T_g of higher than 700°C, and the surface becomes more rough with increasing T_g . A different type of



Figure 5. N_d - N_a depth profiles in Sn-doped Ga₂O₃ epitaxial films grown at 540, 570, and 600°C. Reproduced with permission from [31]. Copyright 2014, Elsevier.

rough surface due to three-dimensional growth is shown for $T_{\rm g} = 500^{\circ}$ C. These experimental data indicate that 550–650°C is the optimum window of $T_{\rm g}$ for obtaining a smooth surface.

Next, the electrical properties of the ozone–MBE Ga₂O₃ films were characterized. The dependence of effective carrier concentration (N_d – N_a) on T_g was characterized by using electrochemical capacitance–voltage (C–V) measurements. Figure 5 plots the N_d – N_a depth profiles in three Sn-doped Ga₂O₃ epitaxial films grown at $T_g = 540$, 570 and 600°C. Uniform distributions of the doped Sn atoms along the growth direction were observed for the samples grown at $T_g = 540^{\circ}$ C and 570°C. On the other hand, a delay in the Sn doping at the initial stage was confirmed for the sample grown at $T_g = 600^{\circ}$ C, which was likely to be due to Sn segregation. Therefore, the optimum T_g for Sn-doped *n*-Ga₂O₃ films was determined to be 550–570°C from the overlap of the growth conditions for the best structural and electrical properties.

6.2. HVPE

MBE is a very useful tool for scientific research, because we can obtain high-quality, high-purity epitaxial films by omitting the possibility of the incorporation of unintentional impurities. However, the productivity of MBE, which requires an ultra-high-vacuum atmosphere, is open to question. We chose HVPE from the several epitaxial growth methods as the first candidate for Ga_2O_3 thin film growth for future mass production of Ga_2O_3 epitaxial wafers. Ga_2O_3 HVPE R&D is still at a primitive stage; however, we have already established several important technologies [33, 34].

An atmospheric-pressure horizontal hot-wall HVPE system as schematically shown in figure 6 is used for Ga_2O_3 growth [33]. Currently, we use EFG-grown Ga_2O_3 (001)



Figure 6. Schematic diagram of the in-house-built atmosphericpressure HVPE system for Ga₂O₃ growth.

substrates for the HVPE growth. Based on thermodynamic analysis, GaCl and O₂ were determined to be appropriate precursors for Ga₂O₃ growth [33]. In the HVPE system, GaCl is generated by the reaction between high-purity Ga metal and Cl₂ gas at 850°C in an upstream region of the reactor. The GaCl and O_2 are separately introduced into a growth zone using N₂ as a carrier gas, and the Ga₂O₃ substrate is placed on a quartz glass susceptor. The SiCl₄ dopant gas is simultaneously supplied in case of *n*-type Ga₂O₃ growth. The growth rate has been controlled up to as high as about 20 μ m h⁻¹ without degradation of the material quality of the Ga₂O₃ epitaxial film [33]. Unintentionally doped (UID) Ga₂O₃ thin films grown by HVPE possess excellent structural and electrical properties, evidenced by an XRC peak as narrow as the one from the EFG bulk single crystal and an extremely low residual carrier concentration of less than $1\times 10^{13}~{\rm cm}^{-3}$ [34]. The *n*-type conductivity of Ga_2O_3 films can be controlled by the Si doping density in the wide range on the order of $10^{15} \sim 10^{18}$ cm⁻³. Details of the material and electrical properties of HVPE-grown n-Ga2O3 thin films will be reported elsewhere.

7. Ga₂O₃ transistors

7.1. Ga₂O₃ MESFETs

Transistor action for single-crystal Ga₂O₃ devices was first demonstrated using simple metal–semiconductor field-effect transistor (MESFET) structures [17]. By ozone MBE, a Sndoped *n*-type Ga₂O₃ layer with a thickness of 300 nm was grown on a Mg-doped semi-insulating Ga₂O₃ (010) substrate. The Sn doping concentration was set at 7×10^{17} cm⁻³. The activation energy of Sn in Ga₂O₃ was estimated to be about 60–80 meV; therefore, about half of the Sn dopants were activated in the MBE-grown Ga₂O₃ layer at room temperature (RT). Figure 7(a) shows a cross-sectional schematic of the *n*-Ga₂O₃ MESFET. We employed a circular FET pattern as



Figure 7. (a) Cross-sectional schematic illustration and (b) optical microscope micrograph of Ga₂O₃ MESFET. Reproduced with permission from [17]. Copyright 2012, AIP Publishing LLC.

shown in figure 7(b). In the first process of ohmic contact formation, a reactive ion etching (RIE) treatment was performed using a gas mixture of BCl₃ and Ar followed by evaporation of Ti(20 nm)/Au(230 nm) and lift off. We found that the RIE process is effective in improving the ohmic contacts. Finally, Schottky gates were fabricated by Pt (15 nm)/Ti(5 nm)/Au(250 nm) deposition and lift off. Surface dielectric passivation was not performed for the device. The gate length, source–drain spacing and diameter of the inner circular drain electrode were 4, 20 and 200 μ m, respectively.

Figures 8(a) and (b) show the dc output and transfer characteristics of the Ga₂O₃ MESFET, respectively. The drain current density (I_d) was effectively modulated by a gate voltage (V_g), and the device exhibited a perfect pinch off. The maximum I_d was 26 mA mm⁻¹ at a drain voltage (V_d) of 40 V and a V_g of +2 V. The three-terminal destructive breakdown in the off-state, which resulted in permanent degradation of the gate electrode, happened at a V_d of over 250 V. The maximum transconductance (g_m) was 2.3 mS mm⁻¹ for $V_d = 40$ V. The off-state drain leakage current was as small as 5 μ A mm⁻¹, and the on/off I_d ratio reached about four orders of magnitude.

The Ga_2O_3 MESFET showed excellent device characteristics as the first FET demonstration. However, the devices also exhibited two clear issues. The most serious problem in the device characteristics was the high contact resistance of source/drain (S/D) electrodes. In addition, the I_d on–off ratio was limited by the small leakage current through the unpassivated Ga_2O_3 surface.

7.2. Depletion-mode Ga_2O_3 MOSFET version 1: MBE-grown Sn-doped Ga_2O_3 channel

To overcome the drawbacks of the MESFETs, we next fabricated depletion-mode Ga_2O_3 MOSFETs [41]. Figure 9 shows a schematic cross section of the Ga_2O_3 MOSFET. The Sn-doped Ga₂O₃ layer with a thickness of 300 nm was grown at 560°C on an Fe-doped semi-insulating Ga₂O₃ (010) substrate by ozone MBE. The device process started with multiple Si-ion (Si^+) implantations to the regions under the S/D electrodes to form a 150 nm deep box profile with $Si = 5 \times 10^{19} \text{ cm}^{-3}$, followed by activation annealing at 925°C for 30 min in an N2 gas atmosphere. Then, a Ti/Au metal stack was deposited on the implanted regions and annealed again at 470°C for 1 min in an N2 gas atmosphere. This metal annealing process is effective to further decrease contact resistance, probably due to the reaction between the metal and Ga₂O₃ at the interface. We obtained a specific contact resistance as low as $8.1 \times 10^{-6} \ \Omega \ cm^2$ for the annealed contacts. Further details of the Si⁺ implantation doping in Ga₂O₃ and its application to the fabrication of lowresistance ohmic contacts are reported elsewhere [42]. A 20 nm thick Al₂O₃ gate dielectric and passivation film was formed on the Ga₂O₃ layer at 250°C by plasma atomic layer deposition. Note that the conduction band offset at the Al₂O₃- Ga_2O_3 interface was estimated to be about 1.5 eV [43]. The gate metal was formed with a Pt-Ti-Au stack on top of the Al_2O_3 film. The gate length and width were 2 μ m and 500 μ m, respectively. The spacing between the S/D Si⁺implanted regions was 20 μ m.

Figure 10(a) shows the dc output characteristics of the Ga₂O₃ MOSFET at RT. As in the case of the MESFET, the I_d was effectively modulated by V_g with good saturation and sharp pinch-off characteristics. Device self-heating caused by the poor thermal conductivity of Ga₂O₃ gave rise to the negative output conductance with increasing V_d . The maximum I_d was 39 mA mm⁻¹ at $V_g = +4$ V. The three-terminal off-state V_{br} was as high as 370 V at $V_g = -20$ V. The I_d - V_g characteristic of the MOSFET at $V_d = 25$ V is shown in figure 10(b). The I_d on–off ratio was extremely high, exceeding ten orders of magnitude with the measured off-state leakage reaching the lower limit of the measurement



Figure 8. (a) Dc I-V and (b) transfer characteristics of Ga_2O_3 MESFET. Reproduced with permission from [17]. Copyright 2012, AIP Publishing LLC.

instrument. These device characteristics were much superior to those of the Ga_2O_3 MESFETs.

We also investigated the performance of the Ga_2O_3 MOSFET at high-temperature operation. The device characteristics evolved smoothly with increasing device operating temperature. No kink or abrupt change that might be indicative of breakdown events and/or permanent degradation was observed, suggesting stable device operation in the whole temperature range from 25 to 250°C. The MOSFET maintained a high I_d on–off ratio of about four orders of magnitude even at 250°C. Furthermore, the device characteristics were almost entirely recovered after operation at 250°C and cool down to RT.



Figure 9. Schematic cross section of Ga₂O₃ MOSFET structure with an MBE-grown Sn-doped channel layer. Reproduced with permission from [41]. Copyright 2013, AIP Publishing LLC.

7.3. Depletion-mode Ga_2O_3 MOSFET version 2: Si⁺ - implanted Ga_2O_3 channel

The Ga₂O₃ MOSFETs with a Sn-doped n-Ga₂O₃ channel suffered from reproducibility of the doping profile, because the suppression of the delay of Sn incorporation into the Ga₂O₃ at the initial stage of the MBE growth requires the precise control of T_g in the narrow temperature window of 550–570°C. This issue led to both poor channel thickness control and non-uniform in-plane *n*. We addressed the problem in [44] by applying Si⁺implantation doping to not only the S/D electrodes but also the channel layer for reliable doping.

The device structure of the Ga₂O₃ MOSFET presented in this subsection was mostly the same as that of the Sn-dopedchannel MOSFET shown in figure 9, except that the channel was formed by Si⁺ implantation doping to a UID Ga₂O₃ layer grown by MBE. The thickness of the MBE-grown UID Ga₂O₃ layer was 300 nm, which was the same as the Sndoped channel. The implanted Si atom densities in the channel layer and the S/D regions were 3×10^{17} and $5\times10^{19}~\text{cm}^{-3},$ respectively. Multiple implantations were performed to obtain box profiles with depths of 300 nm for the channel and 150 nm for the S/D regions. Implant activation annealing was performed once at 925°C for 30 min in N_2 atmosphere for both the channel and S/D regions. After the implantation and activation process, all fabrication procedures were identical to those employed for the Sn-dopedchannel Ga₂O₃ MOSFETs.

All the device characteristics of the Si⁺-implantedchannel MOSFETs were almost the same as or slightly improved over those of the Sn-doped-channel Ga₂O₃ MOS-FETs. Figures 11(a) and (b) show the dc output and breakdown characteristics of the Ga₂O₃ MOSFET with a gate length of 4 μ m at RT. The maximum I_d was 65 mA mm⁻¹ at

+6 V

+2 V

-2 V

 $V_{g} =$



Figure 10. (a) Dc output and (b) $I_d - V_g$ characteristics of Ga₂O₃ MOSFETs with an MBE-grown Sn-doped channel layer. Reproduced with permission from [41]. Copyright 2013, AIP Publishing LLC.

 $V_{\rm g} = +6$ V. The three-terminal off-state $V_{\rm br}$ was as high as 415 V at $V_{\rm g} = -30$ V. The off-state $I_{\rm d}$ remained below the detection limit prior to device breakdown. At $V_d = 30$ V, a high I_d on-off ratio of about ten orders of magnitude was achieved, and the maximum $g_{\rm m}$ was 3.6 mS mm⁻¹. Figure 12 plots the temperature dependence of the transfer characteristics at $V_d = 30 \text{ V}$ of the MOSFET. The presented devices with the Si⁺-implanted channel also revealed stable operation up to 250°C as did the devices with the MBE-grown Sndoped channel. The MOSFETs maintained a low drain



80

60

40

= 4 μm, *W*_a = 500 μm

Figure 11. (a) Dc output and (b) breakdown characteristics of Ga₂O₃ MOSFET with a Si-doped channel layer by ion implantation. Reproduced with permission from [44]. Copyright 2013, IEEE.

leakage current of less than 5 μ A mm⁻¹ and a high I_d on–off ratio of about four orders of magnitude at 250°C.

8. Ga₂O₃ Schottky barrier diodes (SBDs)

8.1. Ga₂O₃ SBDs fabricated on UID Ga₂O₃ native substrates

For the purpose of evaluating the basic device performance of Ga₂O₃ SBDs, we first fabricated simple SBD structures on a



Figure 12. Temperature dependence of the transfer characteristics at $V_d = 30$ V of Ga₂O₃ MOSFET with a Si-implanted channel. Reproduced with permission from [44]. Copyright 2013, IEEE.

UID Ga₂O₃ (010) substrate with a thickness of 600 μ m, which were produced from an FZ-grown bulk crystal [45]. The substrates showed *n*-type conductivity due to the unintentionally incorporated Si in the Ga₂O₃ powder source. The 100 μ m diameter circular Schottky anode electrodes were fabricated on the front side of the substrate with a Pt–Ti–Au stack. Next, BCl₃/Ar RIE treatment was performed on the whole back area of the substrate, and the cathode electrode of Ti/Au was evaporated onto it. These were the same procedures employed for the MESFET to fabricate S/D ohmic contacts [17].

Note that n was uniform along the thickness of the substrate but showed in-plane variation from 3×10^{16} - 1×10^{17} cm⁻³ as evaluated by *C*-*V* measurements. The inplane distribution of n was caused by a non-uniform Si atom density, which was attributed to the characteristic of the Ga₂O₃ FZ growth. Here, we introduce device characteristics of the two selected SBDs having different n of 3×10^{16} cm⁻³ and 5×10^{16} cm⁻³, which were fabricated at different locations on the same substrate. The forward current densityvoltage (J-V) characteristics of the SBDs are shown in figures 13(a) and (b). The near-unity ideality factors of 1.04-1.06 were estimated for both devices, indicating the high crystal quality of the FZ Ga₂O₃ substrate and good Schottky interface property. A Schottky barrier height of 1.3-1.5 eV was extracted for the Pt-Ga₂O₃ interface from the semi logarithmic-plotted J-V and $1/C^2-V$ characteristics. The R_{on} of the Ga₂O₃ SBDs, which were determined from the slopes of the linear regions in figure 13(a), were relatively high at 7.85 and $4.30 \text{ m}\Omega \cdot \text{cm}^2$ because of the low conductivity of the bulk substrate. Figure 13(b) shows the reverse J-V characteristics of the Ga₂O₃ SBDs. The reverse $V_{\rm br}$ values were 150 and 115 V for $n = 3 \times 10^{16}$ and 5×10^{16} cm⁻³,



Figure 13. (a) Forward and (b) reverse J-V characteristics of two different Ga₂O₃ SBDs. They were fabricated at different locations on the same substrate with $n = 3 \times 10^{16}$ and 5×10^{16} cm⁻³. Reproduced with permission from [45]. Copyright 2013, IEEE.

respectively, which were reasonably high for the n and the simple device structure.

8.2. Ga_2O_3 SBDs fabricated on HVPE-grown n⁻- Ga_2O_3 drift layers

Recently, we started development on full-scale Ga_2O_3 SBDs with the advancement of the HVPE technology [33, 34]. The Ga_2O_3 SBDs were fabricated on epitaxial wafers with HVPE-



Figure 14. Schematic cross section of Ga_2O_3 SBD structure with an HVPE-grown n^- - Ga_2O_3 drift layer.

grown Si-doped n^{-} -Ga₂O₃ drift layers on n^{+} -Ga₂O₃ (001) substrates. SiCl₄ was simultaneously supplied during the growth as an *n*-type dopant gas. The growth rate of the Ga₂O₃ epitaxial layers was set at 10 μ m h⁻¹. After the HVPE growth, the surfaces of the epitaxial layers were rough with many pits; therefore, the CMP process was performed to flatten the surfaces. The thickness of the post-CMP drift layer was about 7 μ m.

A schematic cross section of the SBD structure is shown in figure 14. The depth profiles of $N_d - N_a$ extracted by the $d(1/C^2)/dV$ method in two *n*⁻-Ga₂O₃ epitaxial layers grown with different SiCl₄ flow rates were almost constant at 1.4×10^{16} cm⁻³ and 2.0×10^{16} cm⁻³, respectively. Figure 15(a) shows the forward J-V characteristics of the SBDs. From linear fits to the slopes within the range of J =100–200 A cm⁻², the specific R_{on} were estimated to be $3.0 \text{ m}\Omega \cdot \text{cm}^2$ for the device with $N_d - N_a = 1.4 \times 10^{16} \text{ cm}^{-3}$ in the drift layer and $2.4 \text{ m}\Omega \cdot \text{cm}^2$ for the one with $N_{\rm d}$ - $N_{\rm a} = 2.0 \times 10^{16} {\rm cm}^{-3}$. Note that the $R_{\rm on}$ values included a substrate resistance as large as $1.0 \text{ m}\Omega \cdot \text{cm}^2$. The ideality factors of the SBDs were 1.03-1.07, indicating that the Pt/ HVPE-grown Ga₂O₃ interface followed the ideal thermionic emission theory. The reverse J-V characteristics shown in figure 15(b) reveal a high $V_{\rm br}$ of around $-500 \,\rm V$ for both SBDs. Note that hard breakdown events happened with catastrophic damage in both devices due to electric-field concentration at the edge of the anode electrodes. Therefore, implementation of edge termination designs such as a field



Figure 15. (a) Forward and (b) reverse J-V characteristics of Ga₂O₃ SBDs with an HVPE-grown n^{-} -Ga₂O₃ drift layer.

plate and/or a guard ring to the presented device structure should lead to further improvement in $V_{\rm br}$.

9. Conclusion

This paper has attempted to give a review of the present status of R&D on new wide bandgap semiconductor Ga_2O_3 power devices. Ga_2O_3 bulk single crystals can be synthesized by several melt growth methods, and large-size, high-quality Ga_2O_3 wafers produced from the EFG bulk have already been available up to the 4 inch diameter size. Among several epitaxial growth techniques, MBE and HVPE succeeded in producing device-quality Ga_2O_3 epitaxial wafers. Significant progress has been made in the development of Ga_2O_3 transistors and diodes in the last few years, and the device characteristics demonstrated the great potential of Ga_2O_3 power devices for future high-power and high-voltage applications. We consider that vertical transistors and diodes operating at the voltage range of higher than 3 kV will be the main target for Ga_2O_3 power devices in future R&D. However, Ga_2O_3 devices are not limited to the power device applications and thus should also be developed for several different application fields such as high-frequency and extreme environment electronics. Further efforts and progress in R&D on Ga_2O_3 devices should pave the way for nextgeneration semiconductor electronics.

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