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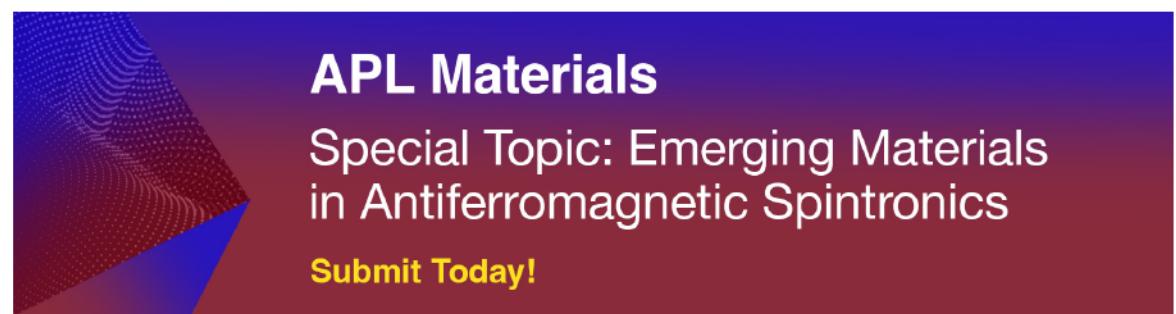
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ABSTRACT

We report the use of suboxide molecular-beam epitaxy (S-MBE) to grow β -Ga₂O₃ at a growth rate of \sim 1 μ m/h with control of the silicon doping concentration from 5×10^{16} to 10^{19} cm⁻³. In S-MBE, pre-oxidized gallium in the form of a molecular beam that is 99.98% Ga₂O, i.e., gallium suboxide, is supplied. Directly supplying Ga₂O to the growth surface bypasses the rate-limiting first step of the two-step reaction mechanism involved in the growth of β -Ga₂O₃ by conventional MBE. As a result, a growth rate of \sim 1 μ m/h is readily achieved at a relatively low growth temperature ($T_{\text{sub}} \approx 525$ °C), resulting in films with high structural perfection and smooth surfaces (rms roughness of <2 nm on \sim 1 μ m thick films). Silicon-containing oxide sources (SiO and SiO₂) producing an SiO suboxide molecular beam are used to dope the β -Ga₂O₃ layers. Temperature-dependent Hall effect measurements on a 1 μ m thick film with a mobile carrier concentration of 2.7×10^{17} cm⁻³ reveal a room-temperature mobility of 124 cm² V⁻¹ s⁻¹ that increases to 627 cm² V⁻¹ s⁻¹ at 76 K; the silicon dopants are found to exhibit an activation energy of 27 meV. We also demonstrate working metal–semiconductor field-effect transistors made from these silicon-doped β -Ga₂O₃ films grown by S-MBE at growth rates of \sim 1 μ m/h.

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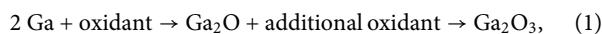
INTRODUCTION

With its very high bandgap, dopability, good mobility for electrons, and the availability of a large-diameter native substrate, β -Ga₂O₃ has emerged as a promising material for high-power

electronics.^{1,2} Although molecular-beam epitaxy (MBE) is the leading technique for the growth of most compound semiconductors, for the growth of β -Ga₂O₃, it has serious limitations, and metalorganic chemical vapor deposition (MOCVD) is currently the method

of choice.^{3,4} A direct comparison of the best electrical properties reported for the growth of β -Ga₂O₃ films by MBE^{5,6} and MOCVD³ at growth rates near current limits for each is shown in Table I, revealing the shortcomings of MBE.

The conventional MBE growth of β -Ga₂O₃ starts with metallic gallium, which undergoes a two-step reaction to form solid Ga₂O₃,^{7–9}



where the oxidant is either oxygen plasma or ozone. The suboxide produced in the first step, Ga₂O, is volatile at typical growth temperatures, and if there is insufficient oxidant present to oxidize it, it will desorb from the substrate surface diminishing the throughput of the second step. The first step is rate-limiting and even operating at the oxidant pressure limit that hot filaments in an MBE can withstand, i.e., $\sim 10^{-5}$ Torr, the growth rate of β -Ga₂O₃ is relatively slow, in the $\sim 0.1 \mu\text{m/h}$ range.^{2,5,6} It is possible to enhance this meager growth rate by catalytic means¹⁰ using a process referred to as metal-oxide catalyzed epitaxy (MOCATAXY)¹¹ or metal-exchange catalyzed molecular beam epitaxy (MEXCAT-MBE).¹² The metallic catalyst can be added indium, which can be incorporated into β -(Al_xGa_{1-x})₂O₃ films at the $\sim 1\%$ level¹¹ or through the addition of a dopant, e.g., tin.¹³ Another way to increase the growth rate is to bypass the first step of the two-step reaction mechanism and directly supply an incident molecular beam of Ga₂O, i.e., suboxide MBE (S-MBE),¹⁴ rather than by supplying a molecular beam of gallium.

Using S-MBE increases the growth rate by overcoming the kinetic limits imposed by the first reaction shown in Eq. (1), leading to an order of magnitude higher growth rate in the $\sim 1 \mu\text{m/h}$ range.¹⁴ There are a couple of options to produce a molecular beam of Ga₂O suboxide. One is to use solid Ga₂O₃ itself,^{15,16} and the other is to use a mixture of solid Ga₂O₃ and liquid gallium.¹⁷ Vapor pressure calculations show that the dominant species in the molecular beam for both methods is the suboxide Ga₂O.^{18,19} An advantage, however, of using the Ga₂O₃ + gallium mixture is the far lower temperature needed to produce the Ga₂O molecular beam.^{18,19} This allows more types of crucibles to be used to contain the mixture (e.g., Al₂O₃ and BeO crucibles), increases the Ga₂O flux that can be achieved (and with it the growth rate of the β -Ga₂O₃ film), and decreases contaminants coming from the hot crucible and effusion cell. For example, when solid Ga₂O₃ has been used as a source, even at source temperatures in the 1700 °C range, the maximum growth rate that has been

achieved is 0.14 $\mu\text{m/h}$ and the films have been contaminated with $\sim 5 \times 10^{18}$ iridium atoms cm^{-3} coming from the iridium crucible used to contain the very hot Ga₂O₃ source.^{20–22} In contrast, films of β -Ga₂O₃ produced using S-MBE using a Ga₂O₃ + gallium mixture result in growth rates exceeding 1 $\mu\text{m/h}$, excellent crystallinity, and smooth surfaces all at a relatively low growth temperature, T_{sub} .¹⁴ The disadvantage of S-MBE is that generating a Ga₂O molecular beam at low temperatures utilizes a source containing a mixture of Ga₂O₃ *powder* plus gallium metal.^{14,17} Unfortunately, the highest purity of Ga₂O₃ powder that we are able to obtain commercially is only 99.999% (5N), whereas 99.99999% (7N) pure gallium is the norm for conventional MBE. This lower purity raises questions about whether S-MBE can grow device-quality films. The direct way to answer this question is to make devices on β -Ga₂O₃ layers grown by S-MBE and see how they perform.

Studying the mobility of β -Ga₂O₃ films and fabricating β -Ga₂O₃-based devices necessitate controlled doping. Silicon is the preferred dopant for β -Ga₂O₃ films as it not only yields films with highest mobility^{2,3,23–25} but has been demonstrated to provide abrupt and controlled doping over the 10^{16} – 10^{20} cm^{-3} range in β -Ga₂O₃ films grown by MOCVD.^{2,3,24–27} Silicon segregates less to the surface during the MBE growth than does tin,⁶ making it a superior dopant, and it has the advantage over germanium of a far lower diffusion coefficient in β -Ga₂O₃ at temperatures required for device processing.²⁸ The traditional MBE approach to dope with silicon is to create a silicon molecular beam by heating silicon to high temperatures in an MBE source. Unfortunately, in the high oxidant pressure ($\sim 10^{-5}$ Torr) used for the growth of β -Ga₂O₃ by MBE, the surface of the silicon doping source may also oxidize.²⁹ Whether the silicon surface gets covered over by a layer of SiO₂ or is able to desorb gaseous SiO at the same rate that the silicon source is oxidized and, thus, remain uncoated depends on the temperature of the silicon and the oxidant pressure. For the case that the oxidant pressure is fixed at 1×10^{-5} Torr, at high temperatures ($T_{\text{Si}} \gtrsim 830^\circ\text{C}$), the silicon surface is able to desorb all of the oxide that forms and remains clean, whereas at low temperature, ($T_{\text{Si}} \lesssim 830^\circ\text{C}$), the silicon surface gets coated by solid SiO₂.^{30–32} These changes with the oxidation of the silicon source result in huge changes in the flux of silicon or SiO emanating from the silicon source, thwarting controlled and reproducible doping at device-relevant concentrations.^{4,5,29}

This paper tackles the challenges of growing silicon-doped β -Ga₂O₃ thin films by taking advantage of the suboxide forms of both silicon and gallium. In this work, we demonstrate both controllable and reproducible silicon doping of β -Ga₂O₃ at device-relevant

TABLE I. Comparison of the highest reported electrical mobilities of β -Ga₂O₃ films grown at a relatively high rate by MBE vs MOCVD.

Mobility at room temperature ($\text{cm}^2 \text{ V}^{-1} \text{ s}^{-1}$)	Peak mobility ($\text{cm}^2 \text{ V}^{-1} \text{ s}^{-1}$)	Temperature of peak mobility (K)	Growth rate ($\mu\text{m/h}$)	Range of controlled doping (cm^{-3})	References
MBE ^a	129	390	97	10^{17} – 10^{20}	5
MBE ^b	136	168	165	4×10^{16} – 2×10^{19}	6
MOCVD	190	3425	53	2×10^{16} – 4×10^{19}	3

^aConventional plasma-assisted MBE.

^bMetal-oxide catalyzed epitaxy (MOCATAXY).

concentrations, i.e., the 5×10^{16} – 10^{19} cm $^{-3}$ range, with a growth rate of ~ 1 $\mu\text{m}/\text{h}$ by S-MBE. Having overcome the doping challenge, we then evaluate the mobility of the silicon-doped β -Ga₂O₃ layers and demonstrate working devices made by S-MBE at a growth rate of ~ 1 $\mu\text{m}/\text{h}$.

EXPERIMENTAL

All films were grown using a Veeco GEN10 MBE system equipped with retractable and differentially pumped effusion cells that can be exchanged without venting the entire MBE system. This facilitates refilling the Ga₂O₃ + gallium source. The film growth took place on (0001)-oriented sapphire substrates (Kyocera) or (010)-oriented iron-doped β -Ga₂O₃ single-crystal substrates (Novel Crystal Technology) with $10 \times 10 \times 0.5$ mm 3 dimensions held in substrate holders made of Haynes® 214® alloy. The backside of each substrate was coated with a ~ 200 nm thick platinum layer (on top of a ~ 20 nm thick titanium adhesion layer) to enable it to be radiatively heated by an SiC heating element to $T_{\text{sub}} = 525$ or 550 °C as measured by an optical pyrometer operating at a wavelength of 980 nm.

A detailed description of the thermodynamics and kinetics of the growth of Ga₂O₃ by suboxide MBE is given in prior publications.^{14,33} Thermodynamic calculations in the present work were performed using the Scientific Group Thermodata Europe (SGTE) Substance Database (SSUB5)³⁴ within the Thermo-Calc software.³⁵

To obtain a molecular beam of gallium suboxide, we shake the combination of Ga₂O₃ powder (Alfa Aesar, 99.999% purity) and molten metallic gallium (Alfa Aesar, 99.9999% purity) to produce a mixture with a molar fraction of oxygen of $x(\text{O}) = 0.4$. The Ga₂O₃ + gallium charge is contained within a beryllium oxide (BeO) crucible (Materion, 99.7% purity) in the MBE effusion cell. Our thermodynamic calculations estimate that such a mixture heated to the 750–1000 °C range of temperatures used in this study will produce a molecular beam that is 99.98% Ga₂O.¹⁴ Films were grown at a background pressure of distilled ozone ($\sim 80\%$ ozone with the balance being O₂) of $P_{\text{O}_3} = (1\text{--}5) \times 10^{-6}$ Torr. The growth rate of films grown on *c*-axis sapphire substrates was determined by measuring the thicknesses of the β -Ga₂O₃ films by x-ray reflectivity (XRR). For films grown on (010)-oriented iron-doped β -Ga₂O₃ single-crystal substrates, the growth rate was estimated by growing calibration films on *c*-axis sapphire substrates and assuming that the growth rate was the same on the (010)-oriented β -Ga₂O₃ substrates. The thickness of one sample (sample a) was measured by scanning transmission electron microscopy (STEM) and found to be $\sim 40\%$ thicker than this simple estimation. For two of the homoepitaxial β -Ga₂O₃ films (samples k and l), the growth rate was established by depth measurements made using a stylus profilometer on secondary-ion mass spectrometry (SIMS) craters.

To dope silicon into β -Ga₂O₃ films, we tried two silicon-containing oxide sources: SiO (Alfa Aesar, 99.99% purity) and SiO₂ (Kurt J. Lesker, 99.99% purity) contained within Al₂O₃ crucibles (McDanel, 99.8% purity). From vapor pressure calculations, both sources are expected to produce a molecular beam of suboxide SiO.¹⁹ This approach is an attempt to circumvent the challenges associated with going back and forth between the active and passive oxidation regimes of elemental silicon as the silicon source temperature is

changed, as unfortunately occurs in the growth of β -Ga₂O₃ by conventional MBE using a silicon source.^{5,29} Both SiO and SiO₂ have previously been used as doping sources for silicon in oxide MBE: SiO to dope β -Ga₂O₃ films^{36,37} and SiO₂ to dope α -Al₂O₃ films.³⁸

The SiO_x flux emanating from the SiO and SiO₂ doping sources was measured in two ways. At high SiO and SiO₂ source temperatures, where the SiO_x flux was sufficient to build up a multi-nanometer thick film in a reasonable time, its thickness was determined by XRR. XRR measurements were made on amorphous SiO_x films deposited on unheated (0001) Al₂O₃ or (100) MgO substrates using the calibration scheme described by Ref. 37. The silicon fluxes calculated from these XRR measurements were independent of the oxidant employed for background pressures of distilled ozone ranging from none (vacuum) up to at least 2.5×10^{-6} Torr. At lower SiO and SiO₂ source temperatures, the SiO_x flux was inferred from Hall effect measurements of silicon-doped β -Ga₂O₃ films. The Hall effect measurements were made using a van der Pauw geometry³⁹ with four ohmic contacts on 1 μm thick silicon-doped β -Ga₂O₃ films grown by S-MBE at a growth rate of ~ 1 $\mu\text{m}/\text{h}$ on iron-doped β -Ga₂O₃ (010) substrates that were $10 \times 10 \times 0.5$ mm 3 in size. For highly silicon-doped samples (greater than 1×10^{19} cm $^{-3}$), contacts to the films were made by soldering indium at the corners. In order to make ohmic contact to more lightly silicon-doped films, ~ 90 nm of n^+ epitaxial β -Ga₂O₃ was regrown on the four corners of the films using a shadow mask in an Agnitron Agilis 100 MOCVD system. The donor density (N_d) in these n^+ layers was greater than 1×10^{19} cm $^{-3}$ to ensure that the contacts are ohmic. The MOCVD reactor pressure was 50 Torr, and the substrate temperature was 630 °C. Triethylgallium (TEGa), oxygen (99.999%), and silane (25 ppm SiH₄ in argon) were used as precursors, with argon (99.999%) as the carrier gas. Then, indium was soldered on top of the regrown n^+ β -Ga₂O₃ contacts. A Nanometrics HL5500 Hall system was used on these samples to determine the sheet carrier concentration of mobile charges at room temperature. The silicon flux was estimated under the assumption that all silicon is activated at room temperature in the β -Ga₂O₃ film. The temperature-dependent transport properties of select samples (samples a, b, and d) were measured as a function of temperature using a vacuum cryostat with a closed-cycle helium compressor and an electromagnet with an applied magnetic field of 0.5 T for Hall-effect measurements.

X-ray diffraction (XRD) and x-ray reflectivity (XRR) measurements were performed in a coplanar symmetric geometry using a PANalytical Empyrean system equipped with a copper anode and a hybrid incident-beam monochromator to provide Cu $K_{\alpha 1}$ radiation. Rocking curves were collected in a triple-axis configuration using a 220 Ge analyzer crystal. An Asylum Research Cypher ES atomic force microscope (AFM) was used to measure the surface roughness.

SIMS measurements were made either using a Hiden Analytical SIMS Plus Workstation or by EAG Laboratories. The Hiden Analytical SIMS Plus Workstation used an O₂⁺ ion source as the primary beam to profile the sample. The system features a MAXIM quadrupole mass analyzer to detect and analyze the emitted secondary ions at 30° to the probe axis. The O₂⁺ primary ion beam was oriented at 45° relative to the sample surface. The primary beam voltage and current for the analysis were 2 kV and 140 nA, respectively. The crater area, scan density, electronic gate, and oxygen flooding pressure were 400×400 μm^2 , 100 × 100 pixels 2 , 5% of the

raster area, and 4.0×10^{-6} Torr, respectively. The SIMS system base pressure was $\sim 5.0 \times 10^{-10}$ Torr.

The microstructure of one epitaxial β -Ga₂O₃ film (sample a) was examined using STEM. The sample was prepared in the cross-sectional geometry perpendicular to the [201] direction and thinned to electron transparency using a Thermo Fisher Helios G4 UX focused ion beam with a final milling step of 5 keV to minimize the damage of the top layer. Carbon and platinum protective layers were deposited on the sample prior to milling to minimize the ion-beam damage. STEM measurements were taken with an aberration-corrected Thermo Fisher Spectra 300 CFEG operated at 300 keV with a high-angle annular dark-field (HAADF) detector.

Metal–semiconductor field-effect transistors (MESFETs) were fabricated using annealed Ti/Al/Ni/Au ohmic contacts, a Ni/Au gate, and mesa isolation via reactive-ion etching,⁴⁰ with a channel length of 3 μ m and a gate length of 1 μ m.

RESULTS AND DISCUSSION

Utilizing a Ga₂O₃ + gallium two-phase mixture to produce a molecular beam of the suboxide Ga₂O, we first map out the growth rate of β -Ga₂O₃ on (0001) Al₂O₃ substrates as a function of suboxide flux under conditions yielding epitaxial films. In Fig. 1, we demonstrate that the growth rate of β -Ga₂O₃ is a function of substrate temperature, distilled ozone pressure, and Ga₂O flux. For a constant substrate temperature and background pressure of distilled ozone, the growth rate, at first, increases linearly with the Ga₂O flux and then plateaus. This behavior is seen in Fig. 1(a). For the growth conditions studied, the β -Ga₂O₃ films grown on (0001) Al₂O₃ were fully epitaxial by reflection high-energy electron diffraction and XRD with (201) orientation in agreement with prior reports of the growth of β -Ga₂O₃ on (0001) Al₂O₃ by pulsed-laser deposition,⁴¹ MOCVD,⁴² conventional MBE,^{7,8,37,43,44} and S-MBE.^{14–16} The linear region corresponds to oxidant-rich conditions, where all of the

Ga₂O supplied in the incident flux is oxidized to Ga₂O₃ during its residence time on the substrate surface. When the Ga₂O flux increases further—into the Ga₂O-rich regime—the excess Ga₂O, beyond what can be oxidized by the background pressure of the distilled ozone, is desorbed from the surface due to the volatility of Ga₂O at the substrate temperatures employed. This results in a plateau in the growth rate with increasing gallium suboxide Ga₂O flux. The growth rate in the plateau regime depends solely on the background pressure of the distilled ozone (and substrate temperature); it is an adsorption-controlled growth regime, where the insufficient ozone flux limits the adsorption of the excess Ga₂O flux. As the background ozone pressure is increased, more Ga₂O can be oxidized during its residence time and the growth rate is found to increase. This occurs by an extension of the oxidant-rich (linear) regime to a new plateau of the growth rate at a higher Ga₂O flux, beyond which there is insufficient ozone present to oxidize all of the Ga₂O to Ga₂O₃ during its residence time on the substrate surface. At a substrate temperature of 525 °C and a background distilled ozone pressure of $P_{O_3} = 5 \times 10^{-6}$ Torr, we achieve growth rates as high as 2.5 μ m/h on (0001) Al₂O₃. Similarly, when keeping the ozone pressure constant and varying the substrate temperature, the growth rate is seen [Fig. 1(b)] to decrease with increasing substrate temperature due to the decreased residence time of the Ga₂O on the substrate surface as the substrate temperature is increased. The oxidant-rich and adsorption-controlled growth regimes are still observed, just shifted due to the change in the residence time. These results agree with a kinetic model developed for S-MBE.^{14,33}

Having established the conditions for the epitaxial growth of β -Ga₂O₃ films at a rate of $\sim 1 \mu$ m/h, we next consider doping them with silicon. For this purpose, we tried both of the oxides of silicon, first SiO and then SiO₂, as source materials to dope β -Ga₂O₃. To calibrate the flux of the SiO molecular beam, XRR was used to determine the growth rate of amorphous SiO_x films. As XRR provides a measure of both the film thickness and the film density,⁴⁵ the

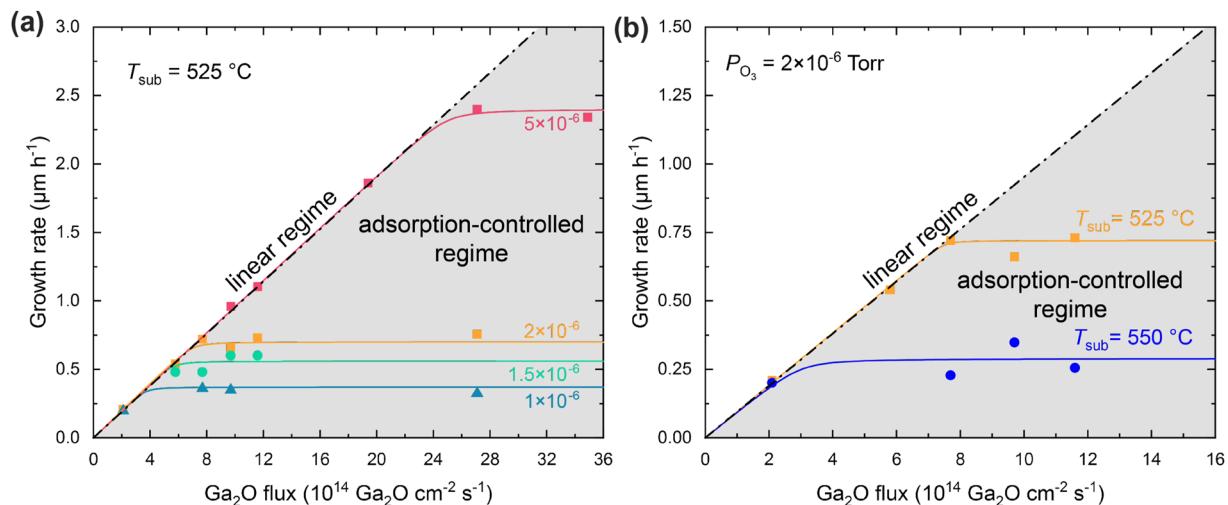


FIG. 1. Growth rate of β -Ga₂O₃ on (0001) Al₂O₃ substrates as a function of Ga₂O flux. (a) Growth rate at a constant substrate temperature ($T_{sub} = 525$ °C) at four different background pressures (in Torr) of distilled ozone showing the oxidant-rich (linear) regime and the Ga₂O-rich (adsorption-controlled) regime. (b) The effect of substrate temperature on the growth rate. The smooth curves indicate a fit to the experimental growth rates by the kinetic model described in Ref. 33.

XRR spectra of smooth SiO_x films enable the silicon flux to be calculated as a function of temperature of the SiO source, T_{SiO} . From the XRR spectra, the density of the amorphous SiO_x films deposited by the SiO source was about 1.8 g/cm^3 , comparable to that reported in prior studies of vacuum-deposited SiO_x films deposited under similar conditions.^{37,46} To evaluate the stability of the SiO source and the associated reproducibility of growing silicon-doped $\beta\text{-Ga}_2\text{O}_3$ films over a range of desired conditions, we grew amorphous SiO_x calibration films at oxidant pressures ranging from vacuum to $P_{\text{O}_3} = 5 \times 10^{-6}$ Torr. While the SiO source behaved well at the high values of T_{SiO} used for the growth of the XRR samples, at the lower values of T_{SiO} relevant to doping $\beta\text{-Ga}_2\text{O}_3$ films with silicon concentrations in the $10^{17}\text{--}10^{19} \text{ cm}^{-3}$ range, SIMS measurements revealed a major challenge with using SiO as a source. At a lower T_{SiO} , the flux is not reproducible. Not only did doping at the same value of T_{SiO} vary considerably from growth to growth (as seen in Hall measurements), but often, the doping concentrations measured in SIMS stacks would not follow an Arrhenius relationship as T_{SiO} was varied. Presumably, this is due to the surface of the SiO forming an SiO_2 crust in the presence of ozone, causing the flux to plummet. This is consistent with the results reported by Ardenghi *et al.*³⁷ for SiO used in conjunction with an oxygen plasma in the growth of silicon-doped $\beta\text{-Ga}_2\text{O}_3$ by plasma-assisted MBE (PAMBE). For this reason, we found SiO to be unsuitable as a controlled and reproducible doping source and moved on to evaluating SiO_2 for this purpose.

In calibrating the SiO_2 source, XRR measurements of amorphous SiO_x on (0001) Al_2O_3 substrates revealed that a well-behaved and reproducible silicon flux emanates from it, despite changing the oxidant pressure from vacuum to $P_{\text{O}_3} = 2.5 \times 10^{-6}$ Torr. From the XRR spectra, the density of the amorphous SiO_x films deposited by the SiO_2 source was about 2.1 g/cm^3 as tabulated for each sample in the *supplementary material* (Table S1). Having established that the SiO_2 source performed well when its temperature, T_{SiO_2} , was high, we moved on to evaluating its stability at lower T_{SiO_2} relevant to

doping $\beta\text{-Ga}_2\text{O}_3$ films. For these lower fluxes of the silicon-containing dopant species (mainly SiO from vapor pressure calculations¹⁹), Hall measurements were performed on silicon-doped homoepitaxial $\beta\text{-Ga}_2\text{O}_3$ films in both the oxidant-rich and adsorption-controlled regimes. The silicon flux is determined from the Hall measurements assuming full activation at room temperature, an assumption that may underestimate the silicon flux. On the other hand, mobile carriers due to the unintentional silicon contamination that often occurs at the interface between the $\beta\text{-Ga}_2\text{O}_3$ film and the underlying substrate^{14,25,47–49} could lead to the Hall measurements overestimating the silicon flux. Tables II and S1 (*supplementary material*) list the growth conditions of the samples grown using an SiO_2 source. Table II lists the samples investigated by Hall measurements (samples a–l), while Table S1 lists the samples measured by XRR (samples o–v).

Figure 2 shows an Arrhenius plot of the silicon flux calculated from both the XRR and Hall data as a function of T_{SiO_2} . A clear Arrhenius behavior with an activation energy of about 5 eV is seen between the silicon flux and $1/T_{\text{SiO}_2}$. To assess our ability to control silicon doping over the $10^{16}\text{--}10^{19} \text{ cm}^{-3}$ range using the SiO_2 sub-oxide doping source, we attempted to grow two $7 \mu\text{m}$ thick films for SIMS analysis, one in the oxidant-rich regime and the other in the adsorption-controlled regime. Each film starts off with a $1 \mu\text{m}$ thick undoped $\beta\text{-Ga}_2\text{O}_3$ buffer layer followed by alternating $0.5 \mu\text{m}$ thick layers of silicon-doped and undoped $\beta\text{-Ga}_2\text{O}_3$ as shown by the schematics in Figs. 3(a) and 3(c) for the SIMS stacks grown in the oxidant-rich and adsorption-controlled regimes, respectively. To keep the growth conditions similar and yet explore the two growth regimes to see whether they affect the dopant incorporation or the doping profile in some way, both SIMS stacks were grown at the same $T_{\text{sub}} = 525^\circ\text{C}$ and the same $P_{\text{O}_3} = 2.5 \times 10^{-6}$ Torr. Accessing the oxidant-rich regime vs the adsorption-controlled regimes was achieved by using different Ga_2O fluxes for these two SIMS stacks: $6 \times 10^{14} \text{ molecules cm}^{-2} \text{ s}^{-1}$ for the oxidant-rich regime and 1×10^{15}

TABLE II. Growth parameters and electrical characteristics of the $\beta\text{-Ga}_2\text{O}_3$ films measured by the Hall effect and SIMS.^a

Name	Mobility ($\text{cm}^2 \text{ V}^{-1} \text{ s}^{-1}$)	Carrier density (cm^{-3})	SiO_2 source temperature ($^\circ\text{C}$)	Thickness (nm)	Ga_2O flux ($\text{molecules cm}^{-2} \text{ s}^{-1}$)
Sample a	124	2.7×10^{17}	1286	1000	1×10^{15}
Sample b	119	2.9×10^{17}	1134	1000	6×10^{14}
Sample c	111	3.5×10^{17}	1115	1000	6×10^{14}
Sample d	129	3.5×10^{17}	1119	1000	1×10^{15}
Sample e	104	9.8×10^{17}	1187	1000	6×10^{14}
Sample f	86	1.3×10^{18}	1225	1000	1×10^{15}
Sample g	98	1.5×10^{18}	1171	1000	6×10^{14}
Sample h	91	2.4×10^{18}	1176	1000	1×10^{15}
Sample i	75	5.5×10^{18}	1262	1000	1×10^{15}
Sample j	79	1.1×10^{19}	1286	1000	1×10^{15}
Sample k	62	3.0×10^{19}	1319	1000	1×10^{15}
Sample l	68	4.4×10^{19}	1350	1000	1×10^{15}
Sample m	Figure 3(a)	6500	6×10^{14}
Sample n	Figure 3(c)	9000	1×10^{15}

^a All samples were grown at $T_{\text{sub}} = 525^\circ\text{C}$ in $P_{\text{O}_3} = 2.5 \times 10^{-6}$ Torr on iron-doped (010) $\beta\text{-Ga}_2\text{O}_3$ substrates and doped with an SiO_2 source.

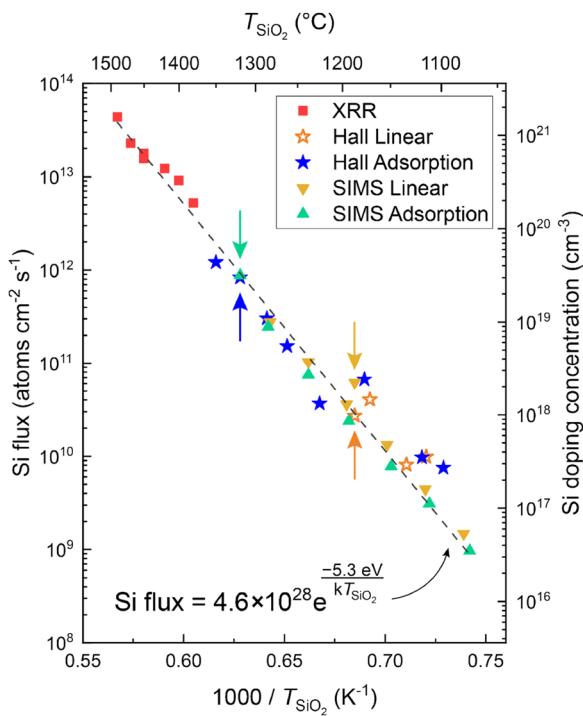


FIG. 2. Silicon flux incorporated into β -Ga₂O₃ films as measured by XRR (red squares), SIMS (triangles), and the Hall effect (stars) as a function of the temperature of the SiO₂ doping source (T_{SiO_2}). The linear fit shown is calculated based on the XRR and SIMS data points, where T_{SiO_2} is in kelvin. Two of the samples on which the Hall effect was measured were also measured by SIMS. For those two samples, the calculated silicon flux values are indicated by the vertical arrows, showing that the fraction of silicon that is electrically active is lower when the β -Ga₂O₃ is less highly doped. The vertical axis on the right side of the figure gives the silicon doping concentration that the silicon flux would produce in a film of β -Ga₂O₃ grown at 1 $\mu\text{m}/\text{h}$.

molecules $\text{cm}^{-2} \text{s}^{-1}$ for the adsorption-controlled regime [Fig. 1(a)]. The growth rates in these two regimes for the growth conditions used differ (0.86 vs 1.3 $\mu\text{m}/\text{h}$), requiring different T_{SiO_2} values. During the growth of each undoped layer, the temperature of the SiO₂ source is increased to provide the silicon flux targeted for the next silicon-doped layer.

Figures 3(b) and 3(d) are the resulting SIMS measurements depicting the concentration of elements of interest—silicon, iron, aluminum, and beryllium—with respect to the sputtered depth. The SIMS data show clear and well-defined steps, demonstrating controllable silicon-doping from 5×10^{16} atoms cm^{-3} to 1×10^{19} atoms cm^{-3} . The steepness of the step edges of the silicon profile differs between the oxidant-rich and adsorption-controlled regimes, with the oxidant-rich regime consistently showing sharper steps. The underlying reason could be the increased rms roughness of the $\sim 9 \mu\text{m}$ thick SIMS stack grown in the adsorption-controlled regime (53 nm) in comparison with the $\sim 6.5 \mu\text{m}$ thick SIMS stack grown in the oxidant-rich regime (11 nm). We also note that when the silicon shutter is open, the silicon profile is also observed to be flatter for the oxidant-rich regime [Fig. 3(b)], whereas it has a

slight upward slope (indicative of silicon riding the surface) for the adsorption-controlled regime [Fig. 3(d)].

Despite calibrating the silicon incorporation by XRR and Hall measurements, the silicon concentration measured by SIMS in Figs. 3(b) and 3(d) does not precisely match the incorporation targeted in our SIMS stack design in Figs. 3(a) and 3(c). At high silicon-doping ($T_{\text{SiO}_2} > 1187^\circ\text{C}$), the match is good, but at lower T_{SiO_2} , the concentration observed by SIMS is lower than the predicted silicon incorporation by up to a factor of 5. One explanation of this discrepancy is that the Hall data overestimate the silicon concentration at low doping because of the unintentional silicon contamination at the substrate interface^{14,25,47–49} giving rise to a concentration of mobile carriers beyond that provided by the silicon flux during growth. To assess this hypothesis, additional SIMS measurements were conducted on two Hall samples (samples k and e), one in each growth regime. If this hypothesis were true, then the concentration of silicon measured by SIMS should be less than the mobile carrier concentration measured on the same samples by the Hall effect. The results are shown in Fig. 2, where the vertical arrows point to the silicon fluxes measured by SIMS and Hall effect on the same sample. The data do not support the hypothesis. Rather, the data show that the silicon concentration measured by SIMS is as large as or larger than the concentration measured by the Hall effect. The data show that at high doping, almost all of the silicon is electrically active, $\sim 90\%$ at $3 \times 10^{19} \text{ cm}^{-3}$ doping. At lower doping, however, the fraction of silicon dopants that produce mobile carriers is far lower, with $\sim 40\%$ activation of silicon at $2 \times 10^{18} \text{ cm}^{-3}$ doping. This could be from a background of compensating acceptor states in our films, but the concentration involving $\sim 10^{18} \text{ cm}^{-3}$ of acceptors or traps is so high that it is inconsistent with the relatively high mobilities seen in other films as we describe below.

The concentration of silicon incorporated into the β -Ga₂O₃ films as a function of T_{SiO_2} in the SIMS stacks shown in Figs. 3(b) and 3(d) was used to calculate the silicon flux at doping-relevant temperatures and is also plotted in Fig. 2. From these SIMS values and the XRR values, a more accurate fit to the Arrhenius behavior between the silicon flux and $1/T_{\text{SiO}_2}$ can be obtained. Unlike the Hall measurement, which only probes mobile carriers, the SIMS and XRR data measure all of the silicon incorporated. The resulting best fit has an activation energy of 5.3 eV/molecule. Although some scatter exists, particularly between Hall and SIMS values, the results in Fig. 2 show a linear trend that extends over 5 orders of magnitude, establishing that SiO₂ is a well-behaved doping source for the growth of β -Ga₂O₃ by suboxide MBE. We attribute the more stable behavior of the SiO₂ doping source over an SiO or silicon doping source to the fact that SiO₂ is fully oxidized and free of the active/passive oxidation issues^{30–32} that plague SiO³⁷ and silicon^{5,29} sources when used at the high oxidant pressures involved in the growth of β -Ga₂O₃ by MBE.

In addition to measuring the concentration of silicon, the concentrations of iron, aluminum, and beryllium were also measured by SIMS. This is because these elements are the major impurities in our Ga₂O₃ powder or they are the major constituents of the BeO and Al₂O₃ crucibles used to contain the Ga₂O₃ + gallium mixture and SiO₂ sources, respectively. A composition analysis on the 99.999% Ga₂O₃ powder reveals aluminum, boron, sodium, and iron to be the only elements present at above the ppm level. A broad screening of these elements and more by SIMS analysis (not shown) indicated

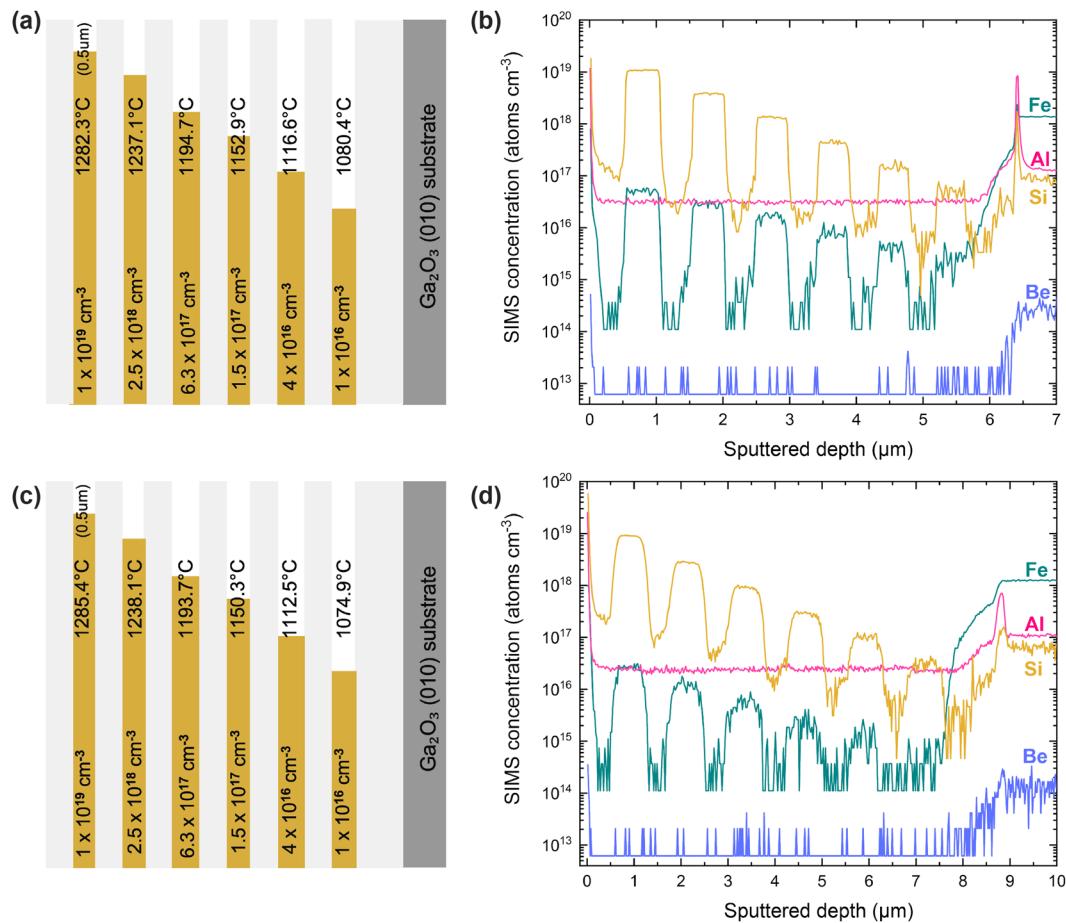


FIG. 3. SIMS profiles showing the concentrations of silicon, iron, aluminum, and beryllium as a function of depth of silicon-doped β -Ga₂O₃ films grown at $T_{\text{sub}} = 525$ °C and a background oxidant pressure of 2.5×10^{-6} Torr of distilled ozone. (a) Schematic of the targeted doping profile, including T_{SiO_2} for each layer, and (b) the measured SIMS profile of the silicon-doped β -Ga₂O₃ films grown in the oxidant-rich (linear) regime at a growth rate of $0.86 \mu\text{m}/\text{h}$ at a Ga₂O flux of $6 \times 10^{14} \text{ molecules cm}^{-2} \text{ s}^{-1}$. (c) Schematic of the targeted doping profile, including T_{SiO_2} for each layer, and (d) the measured SIMS profile of the silicon-doped β -Ga₂O₃ films grown in the Ga₂O-rich (adsorption-controlled) regime at a growth rate of $1.3 \mu\text{m}/\text{h}$ at a Ga₂O flux of $1 \times 10^{15} \text{ molecules cm}^{-2} \text{ s}^{-1}$.

that the major contaminants in our undoped β -Ga₂O₃ films grown by S-MBE are aluminum, silicon, and iron; all impurities that are not isoelectronic with gallium have concentrations below 10^{16} cm^{-3} .¹⁴ The low concentration of beryllium in the SIMS in Figs. 3(b) and 3(d) demonstrate that BeO is a suitable crucible for the Ga₂O₃ + gallium mixture. Of concern, however, are the high iron levels seen in Figs. 3(b) and 3(d). The steps in iron contamination seen by SIMS track the temperature changes and the shutter opening of the SiO₂ source. This indicates that the iron contamination is coming from either (1) the impurities in the SiO₂ source material, with its 99.99% purity; (2) the 99.8% pure Al₂O₃ crucible used to contain the SiO₂; or (3) the iron impurities from the iron-doped β -Ga₂O₃ substrate that ride the growth front⁵⁰ and are preferentially incorporated in the presence of silicon dopants.

When SiO₂ is heated to a temperature range of 1075–1490 °C, as is done in this work, the stable polymorph varies as is described

in Fig. S1 of the [supplementary material](#). Over this temperature range, the silicon-containing species with the highest vapor pressure is the suboxide SiO, followed by SiO₂. According to our thermodynamic calculations (Fig. S1), the SiO makes up 61% of the silicon-containing species in the gas phase at 1075 °C and 99.3% at 1490 °C. Experimentally, the dominant silicon-containing species observed in the gas phase when SiO₂ is evaporated is SiO.^{51–53} Furthermore, the activation of energy of the measured vapor pressure of SiO, averaged from three studies (all with $f < 4 \times 10^{-3}$, where f is a parameter characterizing the effusion cell and, at this small magnitude, is consistent with measurements of the equilibrium vapor pressure), is 5.29 ± 0.21 eV.^{51–53} This agrees well with the activation energy of 5.3 eV from the fit in Fig. 2.

Having established that SiO₂ is a well-behaved doping source, 1 μm thick films of silicon-doped β -Ga₂O₃ are grown in both the oxidant-rich and adsorption-controlled regimes at $\sim 1 \mu\text{m}/\text{h}$.

Figure 4(a) shows the mobility deduced from the Hall measurements of silicon-doped β -Ga₂O₃ films grown by S-MBE (red stars) together with the best results from the literature of silicon-doped β -Ga₂O₃ films grown by other leading techniques.^{3,5,24,27,54,55} From the mobility comparison, it is evident that β -Ga₂O₃ grown by S-MBE with a β -SiO₂ doping source is on a par with other leading techniques. Figure 4(b) clarifies which S-MBE films shown in Fig. 4(a) were grown in the oxidant-rich regime and which were grown in the adsorption-controlled regime by S-MBE. Interestingly, there is no clear difference between the mobility at room

temperature of silicon-doped β -Ga₂O₃ grown by S-MBE in the oxidant-rich or adsorption-controlled regime.

The sample with the highest mobility at room temperature is sample a, a 1 μ m thick film grown by S-MBE with the SiO₂ doping source in the adsorption-controlled regime. At room temperature, it has a mobility of 124 $\text{cm}^2 \text{V}^{-1} \text{s}^{-1}$. The temperature-dependence of this mobility measured by the Hall effect is shown in Fig. 5. The mobility peaks at 76 K at a mobility of 627 $\text{cm}^2 \text{V}^{-1} \text{s}^{-1}$. This value is significantly higher than all prior reports of β -Ga₂O₃ grown by MBE, i.e., higher than PAMBE⁵ and MOCATAXY.⁶ Using the

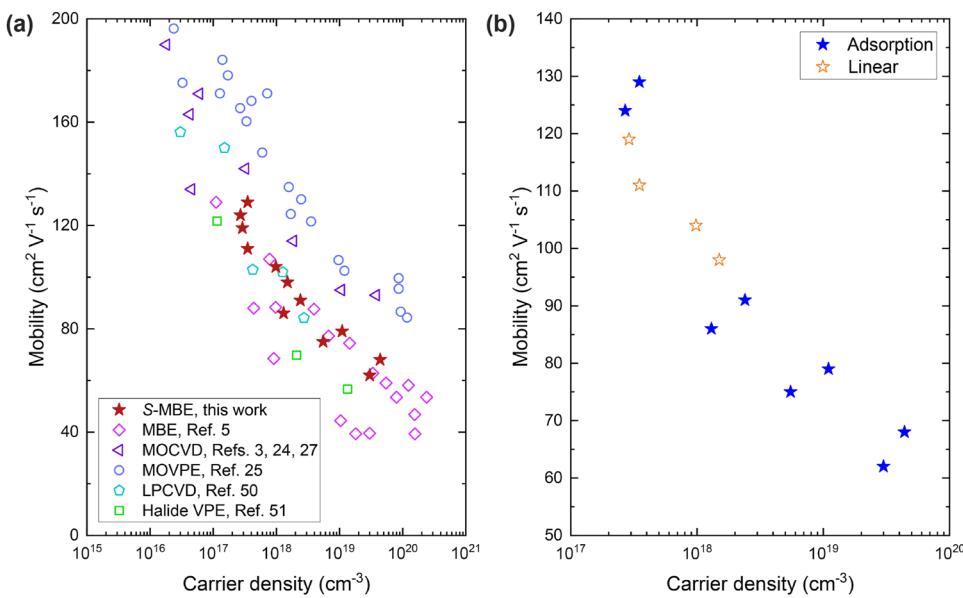


FIG. 4. (a) Comparison of the electron mobility as a function of electron density measured at room temperature by the Hall effect on the silicon-doped β -Ga₂O₃ films grown in this study at $\sim 1 \mu\text{m}/\text{h}$ by S-MBE to leading reports of silicon-doped β -Ga₂O₃ films by other techniques from the literature. The S-MBE results are indicated by the red stars. (b) A close-up of the S-MBE results [the results indicated by the red stars in (a)] showing which films were grown in the oxidant-rich regime (hollow stars) and which films were grown in the adsorption-controlled regime (solid stars).

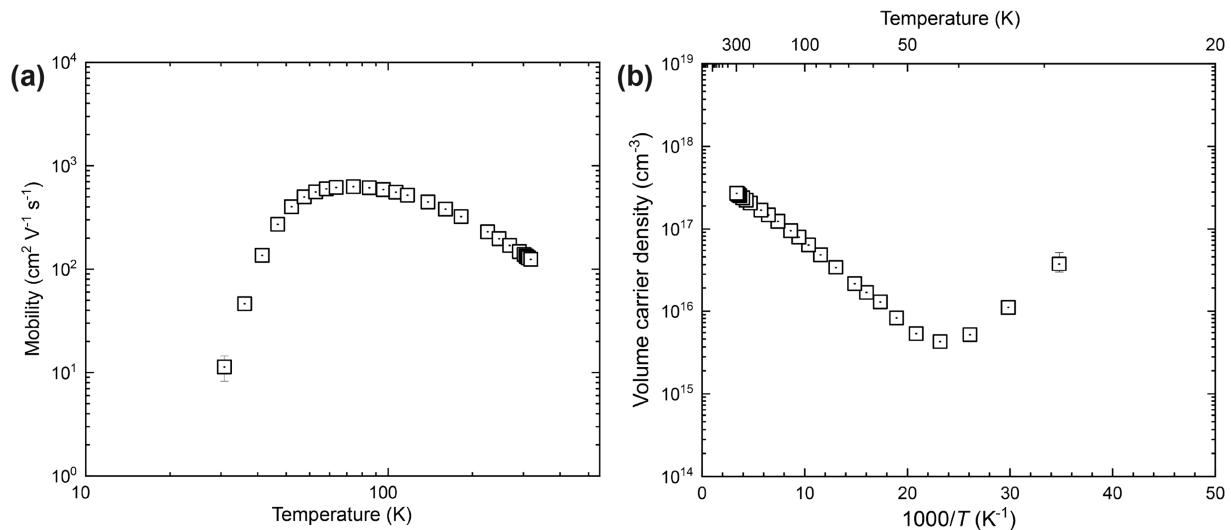


FIG. 5. Temperature-dependent Hall effect measurements made on a 1 μ m thick film (sample a) grown by S-MBE at a growth rate of $\sim 1 \mu\text{m}/\text{h}$ with a mobile carrier concentration of $2.7 \times 10^{17} \text{ cm}^{-3}$ and a room-temperature mobility of 124 $\text{cm}^2 \text{V}^{-1} \text{s}^{-1}$. The mobility of the charge carriers (electrons) (a) and their density (b) are shown as a function of temperature. The mobility peaks at 76 K at a mobility of 627 $\text{cm}^2 \text{V}^{-1} \text{s}^{-1}$.

temperature-dependent carrier density data for this sample as measured by the Hall effect, an activation energy of 27 meV and a compensating acceptor density (N_a) of $4 \times 10^{15} \text{ cm}^{-3}$ were determined by fitting the temperature-dependent Hall carrier density using the charge neutrality equation.^{56,57} This activation energy for silicon from our SiO_2 doping source is comparable to that of silicon in $\beta\text{-Ga}_2\text{O}_3$ grown by conventional MBE (with an elemental silicon doping source)^{5,57} and MOCVD.^{3,23,25}

The results of additional structural characterization measurements on this same high-mobility sample (Sample a) are shown in Fig. 6. The film is unremarkable when analyzed by XRD; single-phase epitaxial $\beta\text{-Ga}_2\text{O}_3$ is seen. Rocking curves of the 020 $\beta\text{-Ga}_2\text{O}_3$ peak reveal full widths at half maxima (FWHMs) of 43 and 122 arcsec along two perpendicular in-plane directions. These values are on a par with the rocking curves of the bare iron-doped $\beta\text{-Ga}_2\text{O}_3$ (010) substrates. AFM reveals an rms roughness of 1.8 nm for this 1 μm thick film grown at 0.86 $\mu\text{m}/\text{h}$. High-angle annular dark-field STEM (HAADF-STEM) imaging reveals a uniform, single-crystalline $\beta\text{-Ga}_2\text{O}_3$ film with high structural quality. The thickness of the epilayer was measured to be $\sim 1.4 \mu\text{m}$, where the interface between the film and substrate is indicated by the black dashed line. The slight contrast at the interface likely originates from contamination at the surface of the air-exposed substrate or from point defects as has been observed in other homoepitaxial $\beta\text{-Ga}_2\text{O}_3$ films.⁵⁸ Dislocations were

not observed at the film/substrate interface or in the epilayer. The variation in contrast at the surface of the film is a result of the TEM sample being prepared as a wedge that is thinner at the surface. We thus demonstrate that using S-MBE for both the semiconductor and the dopant tackles some of the major challenges associated with the conventional MBE growth of $\beta\text{-Ga}_2\text{O}_3$.

Having demonstrated that S-MBE appears to be a viable method for the growth of $\beta\text{-Ga}_2\text{O}_3$ with good structural and electrical properties at record growth rates (for MBE), we return to the question of whether it can produce device-quality materials. As an initial test, we prepared a simple MESFET on a 65 nm thick homoepitaxial $\beta\text{-Ga}_2\text{O}_3$ layer grown by S-MBE at a growth rate of $\sim 1 \mu\text{m}/\text{h}$ with the structure shown in Fig. 7(a). For this test device, the SiO_2 doping source was used to dope the epilayer. The MESFET results are shown in Figs. 7(b) and 7(c); although these are initial results, the ability of S-MBE to produce device-quality materials is evident. Device contacts were ohmic, and the channel showed good modulation, with a peak transconductance (G_m) of 23.4 mS/mm, threshold voltage $V_{th} \sim -6 \text{ V}$, and $V_{off} = -13 \text{ V}$. The relatively low on/off ratio can be attributed to the leakage through the parasitic channel formed by unintentional silicon contamination at the interface between the $\beta\text{-Ga}_2\text{O}_3$ film and the underlying substrate.^{14,25,47-49} These MESFETs perform comparably to devices with similar architectures,^{40,59} indicating that the

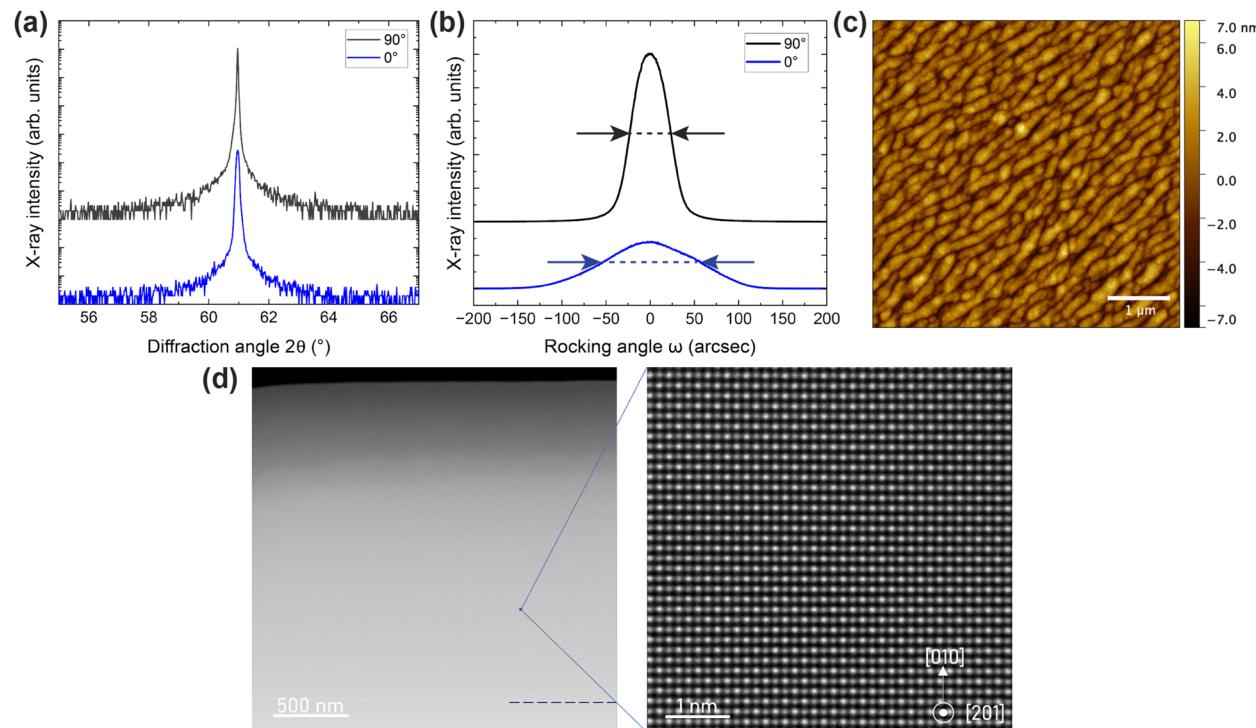


FIG. 6. Structural characterization of the same film shown in Fig. 5 by (a) θ -2 θ XRD scan in the vicinity of the 020 $\beta\text{-Ga}_2\text{O}_3$ peak where the scans along $\phi = 0^\circ$ and $\phi = 90^\circ$ are offset for clarity. (b) Rocking curve of the 020 $\beta\text{-Ga}_2\text{O}_3$ peak showing a FWHM of 122 arcsec (blue) and 43 arcsec (gray) along $\phi = 0^\circ$ and $\phi = 90^\circ$, respectively; the scans are offset for clarity. (c) AFM scan of the same film revealing an rms roughness of 1.8 nm. (d) HAADF-STEM images of the same film along the [201] zone axis, where the dashed line indicates the interface between the $\beta\text{-Ga}_2\text{O}_3$ (010) substrate and the grown $\beta\text{-Ga}_2\text{O}_3$ film.

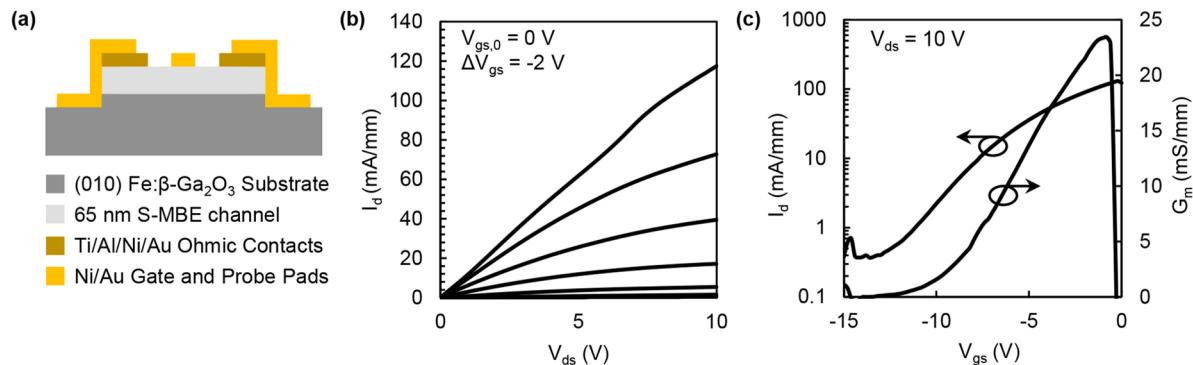


FIG. 7. (a) Schematic of the annealed-ohmic MESFET fabricated using a silicon-doped β -Ga₂O₃ film grown at $\sim 1 \mu\text{m}/\text{h}$ by S-MBE with $N_d = 10^{18} \text{ cm}^{-3}$. (b) Output curves at $V_{gs} = 0, -2, -4, -6, -8$, and -10 V and (c) transfer curve and transconductance of the MESFET.

channel material grown by S-MBE is, indeed, suitable for device fabrication.

CONCLUSIONS

We have overcome the kinetic limitation to the slow growth rate of β -Ga₂O₃ by conventional MBE and demonstrate a growth rate of epitaxial β -Ga₂O₃ as high as $2.5 \mu\text{m}/\text{h}$ on (0001) Al₂O₃ substrates by S-MBE. By using a doping source in which silicon is in its most oxidized state, SiO₂, we avoid the oxidation issues of previously used silicon^{4,5,29} and SiO dopant sources³⁷ when used at high oxidant pressures. This enables controllable and reproducible silicon-doping of β -Ga₂O₃ in the $5 \times 10^{16} \text{ cm}^{-3}$ – 10^{19} cm^{-3} range for the high growth rates used. The doped films grown at $\sim 1 \mu\text{m}/\text{h}$ exhibit mobilities at room temperature rivaling leading techniques and mobilities, at low temperatures, that are the highest achieved to date by MBE, $627 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ at 76 K. While these characteristics are still inferior to the electrical properties of β -Ga₂O₃ grown by MOCVD,^{2–4,23–25} S-MBE is emerging as a viable technique for the growth of electronic-grade β -Ga₂O₃ at rates enabling the intensive investigation of thick vertical heterostructures.

SUPPLEMENTARY MATERIAL

See the [supplementary material](#) for calculated partial pressures and additional sample data.

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AUTHOR DECLARATIONS

Conflict of Interest

The authors K.A., F.V.E.H., S.-L.S., P.V., Z.-K.L., and D.G.S. have been granted U.S. Patent No. 11,462,402 (4 October 2022) with the title “Suboxide Molecular-Beam Epitaxy and Related Structures.”

Author Contributions

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Investigation (supporting); Visualization (equal). **M. K. Indika Senevirathna**: Formal analysis (supporting); Investigation (supporting); Visualization (equal). **Selena Coyer**: Data curation (supporting); Formal analysis (supporting); Investigation (supporting); Visualization (equal). **Shun-Li Shang**: Data curation (supporting); Formal analysis (supporting); Investigation (supporting); Visualization (equal). **Jacob Steele**: Formal analysis (equal); Investigation (supporting); Methodology (equal); Visualization (equal). **Patrick Vogt**: Formal analysis (equal); Investigation (supporting); Methodology (equal); Visualization (supporting). **Nicholas A. Parker**: Formal analysis (supporting); Investigation (supporting); Visualization (supporting); Writing – original draft (supporting); Writing – review & editing (supporting). **Yorick A. Birkholzer**: Investigation (supporting); Writing – original draft (supporting); Writing – review & editing (supporting). **Jonathan P. McCandless**: Funding acquisition (equal); Investigation (supporting); Supervision (supporting); Writing – original draft (supporting); Writing – review & editing (supporting). **Debdeep Jena**: Funding acquisition (equal); Investigation (supporting); Supervision (supporting); Writing – review & editing (supporting). **Huili G. Xing**: Data curation (supporting); Funding acquisition (equal); Supervision (supporting); Writing – review & editing (supporting). **Zi-Kui Liu**: Data curation (supporting); Investigation (supporting); Supervision (supporting). **Michael D. Williams**: Investigation (supporting); Supervision (supporting). **Andrew J. Green**: Funding acquisition (equal); Investigation (supporting); Supervision (supporting). **Kelson Chabak**: Data curation (equal); Funding acquisition (equal); Investigation (supporting); Supervision (supporting); Visualization (equal); Writing – review & editing (equal). **David A. Muller**: Funding acquisition (equal); Resources (supporting); Supervision (supporting); Visualization (supporting); Writing – review & editing (supporting). **Adam T. Neal**: Data curation (equal); Funding acquisition (equal); Investigation (supporting); Supervision (supporting); Visualization (equal); Writing – review & editing (equal). **Shin Mou**: Funding acquisition (equal); Supervision (supporting); Writing – review & editing (equal). **Michael O. Thompson**: Conceptualization (lead); Funding acquisition (equal); Investigation (equal); Project administration (lead); Resources (lead); Supervision (supporting); Validation (equal); Writing – original draft (equal); Writing – review & editing (equal). **Hari P. Nair**: Data curation (equal); Funding acquisition (equal); Investigation (equal); Supervision (supporting); Visualization (equal); Writing – original draft (supporting); Writing – review & editing (equal). **Darrell G. Schloss**: Conceptualization (lead); Funding acquisition (lead); Investigation (equal); Project administration (lead); Resources (lead); Supervision (lead); Validation (equal); Visualization (supporting); Writing – original draft (equal); Writing – review & editing (lead).

DATA AVAILABILITY

The data that support the findings of this study are available within the article and are openly available at <https://doi.org/10.34863/zsda-pa72>.

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