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High-voltage field effect transistors with wide-bandgap β -Ga₂O₃ nanomembranes

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Nanoscale semiconductor materials have been extensively investigated as the channel materials of transistors for energy-efficient low-power logic switches to enable scaling to smaller dimensions. On the opposite end of transistor applications is power electronics for which transistors capable of switching very high voltages are necessary. Miniaturization of energy-efficient power switches can enable the integration with various electronic systems and lead to substantial boosts in energy efficiency. Nanotechnology is yet to have an impact in this arena. In this work, it is demonstrated that nanomembranes of the wide-bandgap semiconductor gallium oxide can be used as channels of transistors capable of switching high voltages, and at the same time can be integrated on any platform. The findings mark a step towards using lessons learnt in nanomaterials and nanotechnology to address a challenge that yet remains untouched by the field. © 2014 AIP Publishing LLC. [<http://dx.doi.org/10.1063/1.4879800>]

High-voltage switching transistors are essential for a multitude of electronic applications that require the transport, conversion, conditioning, and distribution of electrical power.¹ These applications range from microscale on-chip power management for microcontrollers to heavy-duty electric motor drives in automotive applications. Power transistor switches are typically large devices.¹ Large length dimensions are necessary for sustaining high voltages ranging from 100s to 1000s of volts. For a constant electric field, sustaining a larger voltage requires a thicker layer. This is because the critical *electric fields* that can be sustained in semiconductors are limited by their bandgaps through electrical breakdown processes. Traditionally silicon power transistors have been the most widely used.² But the $E_g \sim 1.1$ eV bandgap and its associated critical breakdown field restricts the miniaturization, efficiency, and voltage handling capability of Si power transistors. These limitations have led to increased interest and adoption of power switching devices using wider bandgap semiconductors such as SiC ($E_g \sim 3.3$ eV) and GaN ($E_g \sim 3.4$ eV).³ The wider bandgaps and corresponding higher critical breakdown fields allow higher voltage handling capabilities, and a number of efficiency improvements result from scaling to smaller dimensions. Though wide-bandgap semiconductor devices are finding increased adoption, bulk single crystal substrates remain very expensive for SiC, or challenging for GaN.

A recent breakthrough in the fabrication of bulk single-crystal substrates of a new wide-bandgap semiconductor

β -Ga₂O₃ has attracted much attention in this regard.⁴ β -Ga₂O₃ has a bandgap of ~ 4.9 eV, significantly larger than GaN and SiC. Combined with a relatively inexpensive bulk growth method, it pushes the definition of wide bandgap semiconductors to a new level. Taking advantage of controlled doping in epitaxial layers grown on the bulk substrate, the first transistor was reported recently.⁵ Because of its wide bandgap, Bulk β -Ga₂O₃ substrates can also find applications in deep-ultraviolet photodetectors⁶ and light-emitting diodes (LEDs).⁷ Though the bandgap of β -Ga₂O₃ is large, it has a low thermal conductivity of $13 \text{ W m}^{-1} \text{ K}^{-1}$ (Ref. 8) compared to Si ($\sim 150 \text{ W m}^{-1} \text{ K}^{-1}$),⁹ GaN ($\sim 150\text{--}200 \text{ W m}^{-1} \text{ K}^{-1}$),¹⁰ and SiC ($360\text{--}400 \text{ W m}^{-1} \text{ K}^{-1}$).¹¹ For transistors switching high voltages and currents, a high thermal conductivity semiconductor is desirable for efficient dissipation of excess heat in the switching process. It is imperative to explore ways to counter the low thermal conductivity of β -Ga₂O₃ for such applications.

Further, for on-chip power management applications, it would be desirable to investigate the possible integration of wide-bandgap semiconductors on existing platforms. Recent methods of integration of layered semiconductors such as graphene by layer transfer,¹² and of GaN by direct growth on Silicon¹³ points towards similar opportunities with other materials. In this work, we find that rather surprisingly, β -Ga₂O₃ membranes of nanoscale thickness could be easily exfoliated from their bulk crystals, similar to the exfoliation of graphene and other layered crystal materials. We report the demonstration of high-voltage β -Ga₂O₃ nanomembrane transistors on a Silicon platform. We compare these

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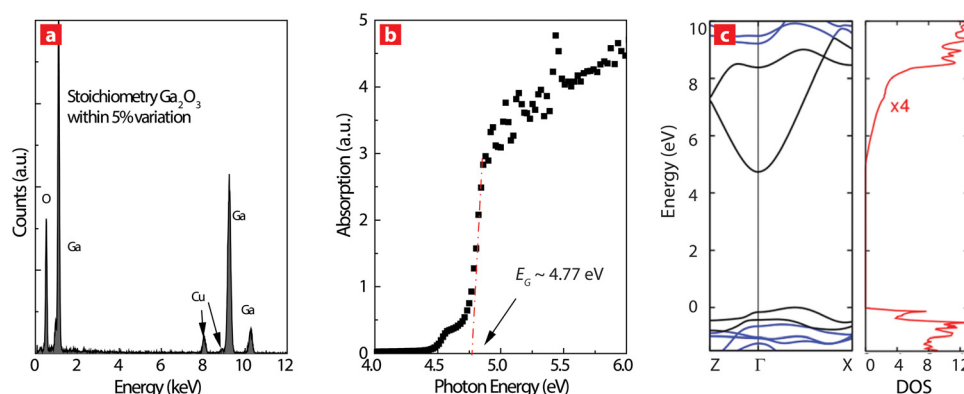


FIG. 1. (a) Normalized intensity of the EDX spectrum for β -Ga₂O₃ nanomembrane. It shows that the β -Ga₂O₃ is formed of stoichiometry structure within 5% variation and the Cu signal is from grid of TEM. The method is too insensitive to measure impurities at the level that impact electrical performance. (b) Optical absorption spectra of β -Ga₂O₃ nanomembrane vs. photon energy, indicating bandgap of ~ 4.77 eV. The absorption is represented by $(\alpha h\nu)^2$. (c) Calculated band structure of Ga₂O₃, plotted along high-symmetry lines in the Brillouin zone. The coordinates of the Z and X points are (0, 0, $1/2$) and (0.266, -0.266 , 0), expressed as fractions of the reciprocal lattice vectors. The highest two valence bands and lowest two conduction bands are highlighted. The right panel shows the density of states (DOS); values for the conduction bands are magnified by a factor of 4 for better visibility.

transistors with similar transistors fabricated from layered MoS₂ semiconductor channels to highlight their high-voltage handling capability. This initial demonstration is intended to motivate research towards the integration of high-voltage transistors with various platforms using nanoscale slivers of wide-bandgap semiconductors. Though nanoscale materials such as nanotubes,¹⁴ graphene,¹² and transition metal dichalcogenides^{15–18} have been actively studied for *low-power* electronic switching applications, the initial findings reported here show that nanoscale materials can also play a role in *high-voltage* electronics.

β -Ga₂O₃ single crystals were obtained by the Czochralski method with use of an iridium crucible. The melting point of β -Ga₂O₃ is around 1820 °C. A dynamic, self-adjusting growth atmosphere was used to minimize the decomposition of β -Ga₂O₃ and oxidation of iridium at the same time, as described in detail in Ref. 4. The crystal structure of β -Ga₂O₃ belongs to the monoclinic system with the space group C2/m and lattice constants $a = 12.214$ Å, $b = 3.0371$ Å, $c = 5.7981$ Å, and $\beta = 103.83^\circ$.¹⁹ It contains two crystallographically inequivalent Ga positions, one with tetrahedral geometry (Ga1) and one with octahedral geometry (Ga2). The octahedra share edges to form double chains parallel to the b-axis, which are connected by corner-sharing tetrahedral.¹⁹ Such structure forms two cleavage planes parallel to (100) and (001) planes. Single crystals of 20 mm diameter were grown along the b-axis, i.e., parallel to both

(100) and (001) cleavage planes. From the single crystal, an oriented cube of $1 \times 1 \times 1$ cm³ size was prepared with an exposed (100) plane for easy-cleaving or exfoliation purpose. Hall effect measurements of bulk β -Ga₂O₃ in the van-der-Pauw configuration with use of In-Ga ohmic contacts showed a free electron concentration = 5.5×10^{17} cm⁻³, resistivity = 0.1 Ω cm, and electron mobility = 112 cm² V⁻¹ s⁻¹, all at room temperature.

Nanomembranes of ~ 20 to 100 nm thickness could be mechanically exfoliated from the bulk crystal, much like the exfoliation of graphene and other transition-metal dichalcogenide layered crystals.¹² This is surprising, because β -Ga₂O₃ is a 3D crystal. Mechanical exfoliation is possible in layered materials due to the highly *anisotropic* chemical bonding characterized by strong in-plane covalent bonds and much weaker interlayer van der Waals bonds. Though β -Ga₂O₃ is not a layered material in the conventional sense, it has a monoclinic structure, with a much larger lattice constant in the (100) direction. The ease of exfoliation could be attributed to this property, though further careful study is necessary to confirm this hypothesis. The Energy-dispersive X-ray (EDX) spectrum of a nanomembrane transferred onto Cu transmission electron microscopy (TEM) grids is shown in Fig. 1(a). It reveals the presence of Ga and O and no other impurities. The absorption spectrum of the β -Ga₂O₃ (100) nanomembrane is shown in Fig. 1(b). The extracted optical band gap matches the expected value obtained from

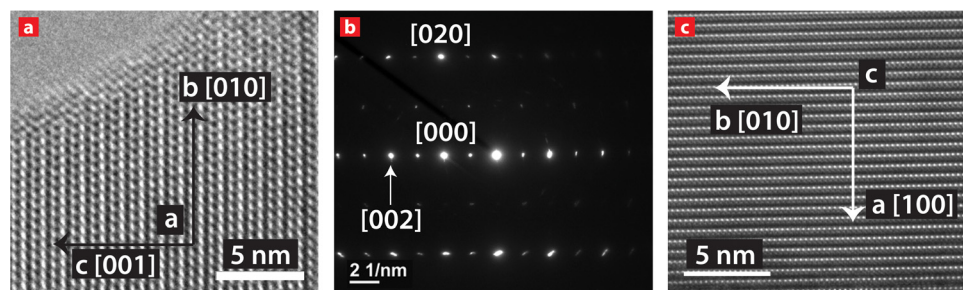


FIG. 2. TEM images of β -Ga₂O₃ nanomembrane (a) HRTEM image of bc-plan-view. (b) Corresponding electron diffraction pattern of (a), the units of 2 1/nm means 2 reversed nanometers. It allows to estimate the g vectors and corresponding lattice spacing as $1/g$ in nm. (c) Cross-sectional HRTEM view of β -Ga₂O₃ nanomembrane FET.

TABLE I. Comparison of bulk lattice parameter¹⁹ and nanomembrane lattice parameter. The value of the nanomembrane lattice parameter is obtained from Fig. 2 and the standard deviation of that is less than 0.005 nm, respectively.

	Lattice parameter (nm)		
	a(100)	b(010)	c(001)
Bulk	1.22	0.30	0.58
Nanomembrane	1.22	0.30	0.57

first-principles calculations. The band structure in Fig. 1(c) is calculated using the same hybrid density functional theory approach as in Refs. 20 and 21 but using a corrected assignment of the high-symmetry lines in the Brillouin zone. Figure 1(c) shows that the conduction-band minimum is at Γ , with a small and almost isotropic electron effective mass of $0.28m_e$. The band gap is indirect, with a value of 4.85 eV and the valence-band maximum located along the I-L line. The direct band gap at Γ is only slightly larger, at 4.88 eV; the absorption spectra shows both the direct and indirect signatures. Calculations of matrix elements indicate that direct optical transitions at Γ are allowed, consistent with the experimentally observed sharp absorption onset.²² In addition to the small effective mass, another feature that renders Ga_2O_3 attractive as an electronic material is the fact that secondary conduction-band minima are at least 2.6 eV higher in energy. The EDX spectrum and the absorption spectrum of the (100) $\beta\text{-Ga}_2\text{O}_3$ nanomembrane thus reveals that the bulk properties are preserved in the nanomembrane.

In order to further investigate the structural properties of the $\beta\text{-Ga}_2\text{O}_3$ nanomembrane, TEM was performed. Figure 2(a) shows the HRTEM image of the bc-plane view, the major cleavage plane of $\beta\text{-Ga}_2\text{O}_3$. The electron diffraction pattern of the bc-plane is shown in Fig. 1(b), revealing the lattice symmetry and the lattice parameters of the bc-plane. The lattice parameters of the [100] a-direction can be

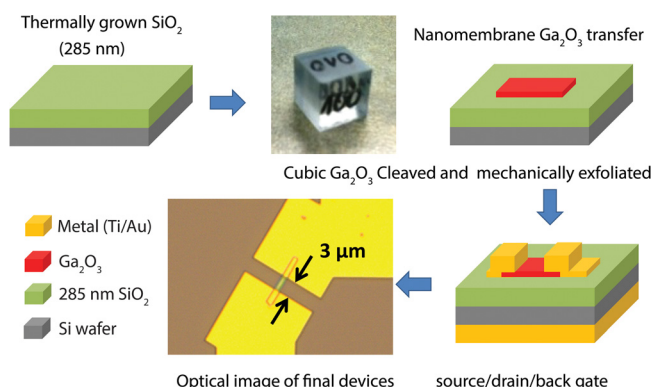


FIG. 3. Schematic process flow for nanomembrane $\beta\text{-Ga}_2\text{O}_3$ field-effect transistors. The nanomembrane thicknesses are in the range of 20 to 100 nm

observed from Fig. 2(c), which is the cross-section HRTEM image of Fig. 4.

The atomic properties of $\beta\text{-Ga}_2\text{O}_3$ nanomembrane were compared with those of bulk in Table I. It shows that the atomic properties of $\beta\text{-Ga}_2\text{O}_3$ nanomembrane are identical to those of bulk, indicating that no obvious deformation or stress is introduced in the nanomembrane during the processing.

Based on the information above, $\beta\text{-Ga}_2\text{O}_3$ nanomembrane high-voltage FETs were fabricated as shown in Fig. 3. Nanomembranes were mechanically exfoliated and transferred onto a back-gated Silicon substrate. The source and drain contacts were defined by electron beam lithography (EBL) using Ti/Au (5/150 nm) metal stacks. The final device went through an annealing process in Ar/H_2 at 300 °C for 3 h to reduce the contact resistance.

Figure 4 shows the cross-section sketch of the back-gated transistor, and the corresponding TEM images of various regions of the device. The highly crystalline nature of the $\beta\text{-Ga}_2\text{O}_3$ channel region, and atomically smooth interfaces at the bottom with SiO_2 and on the top with the Ti/Au ohmic metal is evident in this picture. The lattice parameters of the $\beta\text{-Ga}_2\text{O}_3$ nanomembrane is identical in the channel,

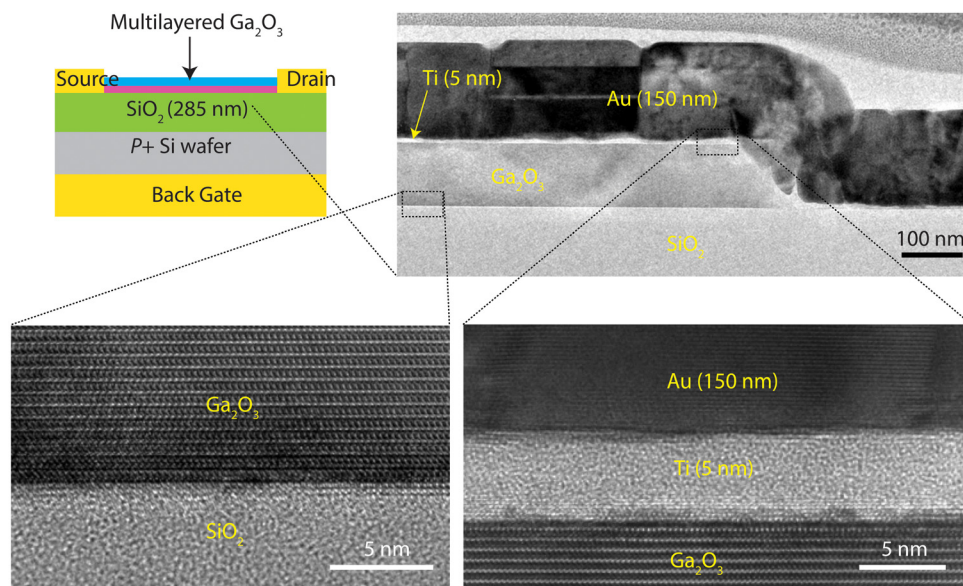


FIG. 4. Cross-sectional TEM image of $\beta\text{-Ga}_2\text{O}_3$ FETs, showing a flat interface between $\beta\text{-Ga}_2\text{O}_3$ and the SiO_2 dielectrics as well as between the $\beta\text{-Ga}_2\text{O}_3$ and the Ti/Au electrode.

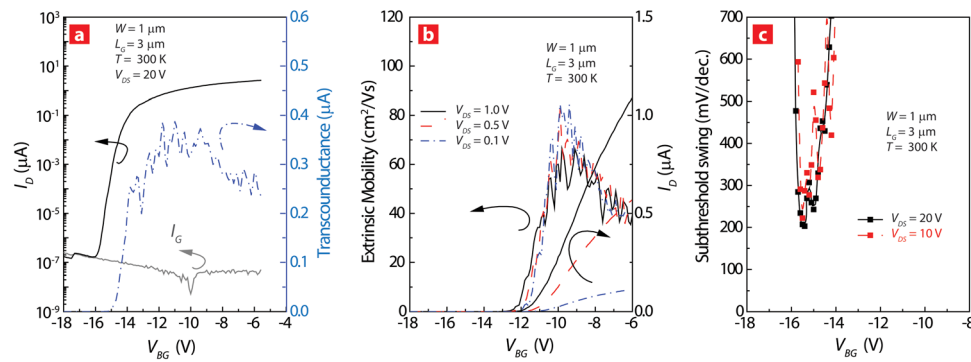


FIG. 5. Transport properties of β -Ga₂O₃ field-effect transistor with $W/L = 1/3 \mu\text{m}$. (a) Drain current, I_D vs. back gate-to-source voltage, V_{BG} , showing $\sim 10^7$ on/off current ratio and n-type semiconductor behavior. (b) Field-effect mobility and (c) subthreshold swing vs. V_{BG} .

under the metal contacts and near SiO₂, and all of them are identical to the bulk value, indicating minimal strain and damage in the transfer and device fabrication process.

Figure 5(a) shows the drain current versus gate-source bias, I_D vs. V_{BS} , at room temperature at a high drain bias $V_{DS} = 20 \text{ V}$. Nanomembrane (100 nm) β -Ga₂O₃ FETs show a high gate modulation of $\sim 10^7 \times$ even under a high drain voltage of 20 V; this modulation is limited not by the channel material, but the gate leakage current. The high current modulation is attractive for high-power and high-voltage device applications. From the transfer characteristics, we extract an “extrinsic” field-effect mobility of $\sim 70 \text{ cm}^2/\text{Vs}$ as shown in Fig. 5(b). This value represents a lower limit because it is not corrected for the contact resistance. The real electron mobility is expected to be around $\sim 130 \text{ cm}^2/\text{Vs}$ closer to the bulk value. A low contact resistance can thus boost the current drive significantly.⁵ The subthreshold swing (SS) of the device shown in Fig. 5(c) approaches $\sim 200 \text{ mV/dec}$. The low SS, though far from the ideal 60 mV/decade is encouraging, considering the unoptimized interfaces and the thick 285 nm SiO₂ back-gate dielectric. This device shows an unintentional n-type behavior which can be caused by atomic defects and/or impurities, etc., in the β -Ga₂O₃.^{23,24}

The family of I_D - V_{DS} curves at various V_{BG} in Fig. 6(a) shows typical transistor performance characterized by linear behavior at low V_{DS} and current saturation at high V_{DS} . The contact resistance was extracted to be $\sim 55 \Omega \text{ mm}$ at low V_{DS} . The value is comparable to MoS₂ FETs with an

identical process flow,²⁵ but is much higher than that of conventional Si and III-V FETs of $\sim 0.1 \Omega \text{ mm}$.²⁶ The high contact resistance is responsible for the lower-than-expected on-current for a long-channel FET of similar geometry and transport properties. The resistance can be further lowered by using low work function metals and by increasing the electron density by local ion-implantation doping under the contacts. In Fig. 6(a), the transistor characteristics of the β -Ga₂O₃ FET is compared to a multilayer MoS₂ channel FET of identical geometry and dimensions. Owing to the much larger bandgap, the $1 \mu\text{m}$ gate length β -Ga₂O₃ channel FET shows a much higher voltage handling capability than the MoS₂ channel FETs. In this comparison, both MoS₂ and β -Ga₂O₃ channel FETs went through the same process flow with identical gate stack structure. Fig. 6(b) shows the bandgaps, and band alignments of the two channel materials. The output characteristics of the nanomembrane β -Ga₂O₃ maintain a robust current saturation up to 70 V with no signs of output conductance. In comparison, the sharp increase in the output current of the MoS₂ channel FET around 10–15 V drain bias is a characteristic signature of avalanche or impact-ionization breakdown, as is expected of semiconductors of comparable bandgaps. This result shows that nanomembranes β -Ga₂O₃ channel transistors can sustain and switch high voltages even when integrated in thin layer forms on foreign substrates. High thermal conductivity but electrically insulating layers such as AlN or BN can be used to help circumvent the low thermal conductivity of the β -Ga₂O₃ channel. The high thermal conductivity insulating layers can also serve as the gate insulator for the transistor.

In summary, nanomembrane high-voltage FETs with β -Ga₂O₃ channels were fabricated and characterized. The large bandgap of $\sim 4.9 \text{ eV}$ leads to high on/off ratio and high voltage transistors with low subthreshold slopes. The high breakdown field of β -Ga₂O₃ allows significantly higher voltages to be switched by orders of magnitude compared to lower-bandgap layered MoS₂ transistors. Since β -Ga₂O₃ has been found to have a rather poor thermal conductivity, the nanomembrane topology offers opportunities for efficient heat removal. A primary challenge is the formation of ohmic contacts. Mechanical exfoliation is indeed not a scalable method, but methods similar to smart-cut technology²⁷ used in Silicon-on-insulator (SOI) wafer manufacture can potentially enable controlled release of large nanomembranes of the wide bandgap material. Such a method can potentially

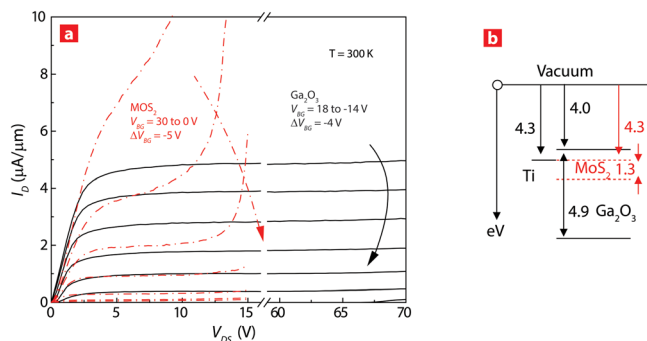


FIG. 6. (a) Common-source transistor characteristics, I_D vs. V_{DS} linear region and current saturation under high V_{DS} , and comparison of breakdown voltage of Ga₂O₃ with $W/L = 1/3 \mu\text{m}$ and MoS₂ with $W/L = 1/3 \mu\text{m}$. (b) Band diagram of β -Ga₂O₃ comparing with MoS₂, indicating the formation of Schottky barrier contact between metal and β -Ga₂O₃.

enable the integration of nanomembrane high-voltage transistors on multiple platforms for high-voltage switching and power management.

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