

Increase in current density for metal contacts to n-germanium by inserting TiO₂ interfacial layer to reduce Schottky barrier height

J.-Y. Jason Lin,^{1,a)} Arunanshu M. Roy,¹ Aneesh Nainani,¹ Yun Sun,² and Krishna C. Saraswat¹

¹Department of Electrical Engineering, Stanford University, Stanford, California 94305, USA

²Stanford Synchrotron Radiation Lightsource, SLAC National Accelerator Laboratory, Menlo Park, California 94025, USA

(Received 11 January 2011; accepted 15 February 2011; published online 3 March 2011)

Metal contacts to n-type Ge have poor performance due to the Fermi level pinning near the Ge valence band at metal/Ge interfaces. The electron barrier height can be reduced by inserting ultrathin dielectrics at the metal-semiconductor interface. However, this technique introduces tunneling resistance from the large conduction band offset (CBO) between the insulator and Ge. In this work, the CBO between TiO₂ and Ge is estimated to range from -0.06 to -0.26 eV so tunneling resistance can be reduced. By inserting 7.1 nm TiO₂ between Al and n-Ge, current densities increased by about $900\times$ at 0.1 V and $1200\times$ at -0.1 V compared to contacts without TiO₂. © 2011 American Institute of Physics. [doi:10.1063/1.3562305]

Germanium is a promising material to replace silicon as the channel material in metal oxide semiconductor field effect transistors (MOSFETs) due to its high electron and hole mobility. While p-channel MOSFETs have been demonstrated with superior performance, n-channel MOSFETs have suffered from poor drive current. One reason for this has been the high contact resistance to the n+ source/drain (S/D) regions due to the high electron Schottky barrier height (Φ_{BN}) as a result of Fermi level pinning close to the Ge valence band (VB) at metal/Ge junctions.^{1,2}

Recent experiments have demonstrated Fermi level depinning by inserting thin tunnel barriers such as Si₃N₄,^{3,4} Ge₃N₄,⁵ and Al₂O₃ (Refs. 6 and 7) at the metal-semiconductor interface. Using a low work function metal can, therefore, result in lower Φ_{BN} . However, these insulators introduce a large tunneling resistance due to significant conduction band offset (CBO) to Ge. Simulations performed by Roy *et al.*⁸ have shown that low specific contact resistivity to n-type Ge can only be achieved if the insulating layer is kept sufficiently thin, typically less than 1 nm, so as not to introduce a large tunneling resistance. It was identified that TiO₂ may be a good candidate since it has a nearly zero CBO to Ge for lower tunneling resistance.

In this work, we use TiO₂ as the insulating layer in a metal-insulator-semiconductor (MIS) structure to evaluate the suitability of using TiO₂ for contacts to n+ S/D regions for application in n-channel Ge MOSFETs. The MIS device cross-section is depicted in Fig. 1.

n-type Ge doped with electrically active concentration of $\sim 10^{19}$ cm⁻³ was heteroepitaxially deposited by chemical vapor deposition (CVD) on n-type Si wafers. This relatively high level of doping was used since our aim is to achieve high currents to n+ S/D. 400 nm SiO₂ was then deposited by CVD at 400 °C for device isolation. Standard lithography techniques were used to pattern contact holes. Atomic layer deposition (ALD) of TiO₂ at 150 °C was done using tetrakis(dimethylamido)titanium and water as the precursors. Control samples using ALD of Al₂O₃ at 200 °C were also made

using trimethylaluminum and water as the precursors. Various thicknesses of TiO₂ and Al₂O₃ were deposited by changing the number of ALD cycles. Samples without a dielectric layer were also fabricated to observe the contact characteristics of metal directly on Ge. 100 nm of Al was deposited by e-beam evaporation onto the sample surface. Standard lithography techniques were used to pattern the Al. An argon sputter etch was performed to completely etch exposed TiO₂ using the Al contact as the mask. Due to the low band gap of TiO₂, electrical conduction has been observed in some TiO₂ films;⁹ this etch, therefore, ensures complete isolation between adjacent devices. Finally, 300 nm of Al was deposited by e-beam evaporation onto the backside of the samples to get a good contact.

The VB offsets (VBO) of TiO₂/Ge and Al₂O₃/Ge were measured using synchrotron radiation photoelectron spectroscopy (SRPES). Low energy ($h\nu=120-160$ eV) photons from the Stanford Synchrotron Radiation Lightsource beam line 8-1 allows precise measurement of the VBO by taking the difference between the VB spectrum of TiO₂ or Al₂O₃ and the VB spectrum of bulk Ge, using the Ge 3d peak for alignment. Figure 2(a) shows the VB spectra of TiO₂ on Ge and bulk Ge, indicating a VBO of 2.9 eV. The band gap of amorphous TiO₂ has been reported to be 3.3–3.5 eV,¹⁰ while the crystalline phases of TiO₂ have smaller band gaps.¹¹ We

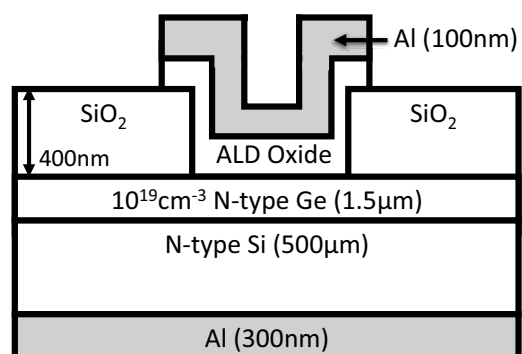


FIG. 1. Schematic cross-section of the MIS device. ALD of both TiO₂ and Al₂O₃ were used.

^{a)}Electronic mail: jasonlin@stanford.edu.

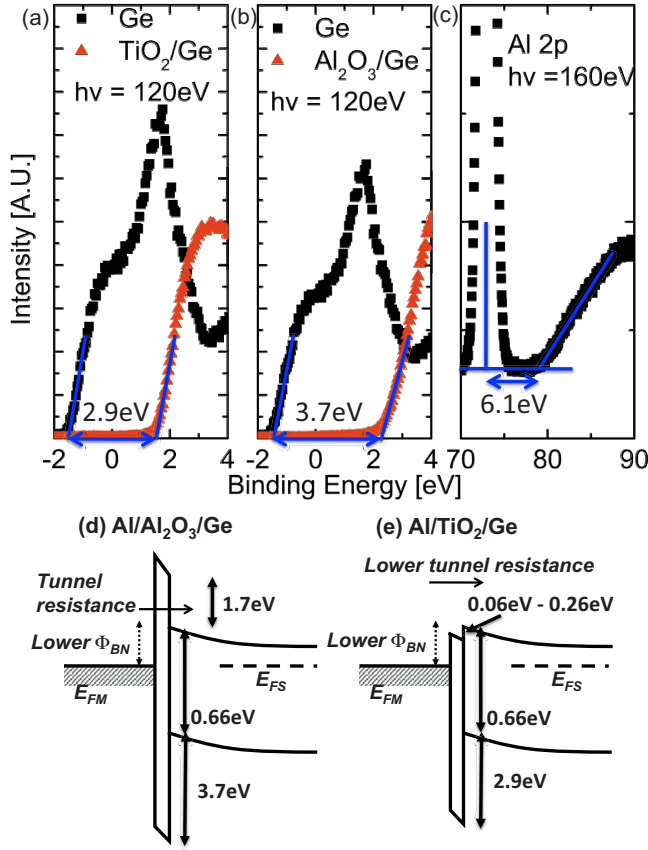


FIG. 2. (Color online) SRPES spectra for (a) TiO₂ and Ge VBs, (b) Al₂O₃ and Ge VBs, and (c) Al 2p peak. The photon energy used is indicated for each plot. The TiO₂/Ge and Al₂O₃/Ge VBO was determined to be 2.9 eV and 3.7 eV, respectively. The band gap of Al₂O₃ was confirmed to be 6.1 eV. Parts (d) and (e) are band diagrams summarizing the VBO and CBO data from SRPES. The metal Fermi level is drawn close to the conduction band, which assumes Φ_{BN} can be reduced.

expect our ALD-deposited TiO₂ films to be amorphous, although this was not confirmed. Since we use the band gap data only to calculate CBO, the larger band gap of amorphous TiO₂ represents a worst-case since it leads to a larger CBO. Using the range of band gaps for amorphous TiO₂, we obtain a CBO between -0.26 and -0.06 eV at the TiO₂/Ge interface. This low CBO value confirms the possibility that inserting TiO₂ at the metal/Ge interface may be able to achieve high current as long as Φ_{BN} can be reduced. Figure 2(b) shows the VB spectra of Al₂O₃ on Ge and bulk Ge, indicating a VBO of 3.7 eV. This is slightly larger than the typical value of ~ 3.3 eV obtained for Al₂O₃/Ge VBO.¹² Since our goal is to study TiO₂, we did not examine this further, but it is likely that different deposition conditions may give slightly different VBO. The ALD Al₂O₃ band gap was found to be 6.1 eV as determined from the energy loss spectrum of the Al 2p peak shown in Fig. 2(c), which is consistent with published results for ALD Al₂O₃.¹² We could not use this technique to determine the band gap of TiO₂ since the Ti 3p peak was too wide and therefore masked the energy loss onset. The VBO and CBO results are summarized in Figs. 2(d) and 2(e). Based on these band diagrams, it appears that TiO₂ MIS devices should be able to achieve higher currents than Al₂O₃ devices on the basis of CBO values, as long as the insertion of TiO₂ can also reduce Φ_{BN} . We, therefore, examine Φ_{BN} next.

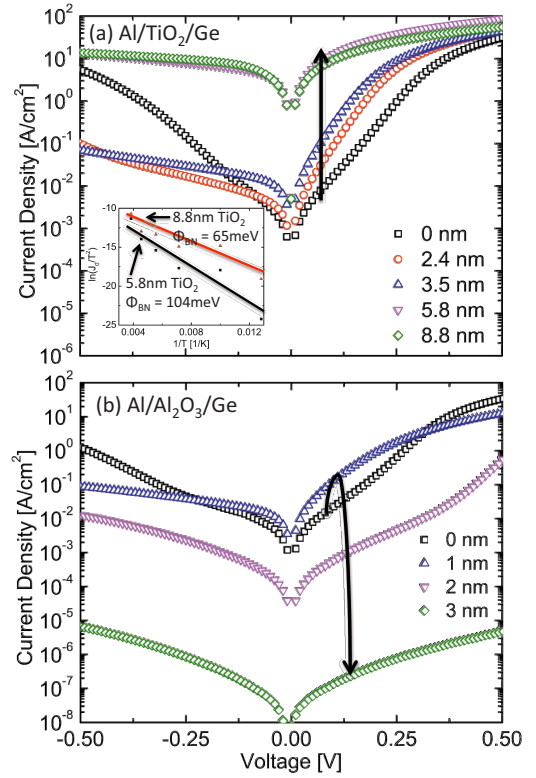


FIG. 3. (Color online) I-V characteristics of (a) Al/TiO₂/Ge and (b) Al/Al₂O₃/Ge MIS devices. With TiO₂, the forward current increases up to 5.8 nm, with no significant current degradation even up to 8.8 nm. With Al₂O₃, the forward current increases only up to 1 nm then starts decreasing rapidly. The inset of (a) shows temperature-dependent I-V measurements of the 5.8 nm and 8.8 nm TiO₂ devices, which show electron barrier heights of 104 meV and 65 meV, respectively.

The current-voltage (I-V) characteristics of Al/TiO₂/Ge and Al/Al₂O₃/Ge MIS devices were measured, using the backside as the second contact. Figure 3(a) shows the I-V characteristics of the TiO₂ devices. Without TiO₂ (0 nm curve) we observe low current as expected due to the large Φ_{BN} . The I-V characteristics are Ohmic-like due to the relatively high semiconductor doping used. As the TiO₂ thickness was increased to 2.4 and 3.5 nm, a noticeable increase in current was seen at forward and low reverse biases. This is evidence of a slight decrease in Φ_{BN} , which increases thermionic emission. At larger reverse bias, however, electron tunneling from the metal to the semiconductor through the semiconductor barrier becomes dominant. Although Φ_{BN} has been decreased, it is likely still a significant fraction of the band gap; the added TiO₂ reduces tunneling and results in current lower than the 0 nm case. With 5.8 nm TiO₂, the current saturates at values significantly higher compared to the 0 nm sample. This high current is due to a significant reduction in Φ_{BN} , which we verified later using temperature-dependent measurements. The fact that the I-V characteristics of the 5.8 and 8.8 nm samples look similar is an indication that thicker TiO₂ does not increase tunneling resistance significantly, as was predicted by our CBO estimation using SRPES.

Temperature-dependent I-V measurements were made between 78 and 260 K for the 5.8 and 8.8 nm TiO₂ devices. The inset of Fig. 3(a) is a Richardson plot of $\ln(J_0/T^2)$ versus $1/T$, where J_0 is the reverse saturation current and T is the temperature. The slope of such a plot is $-q\Phi_{BN}/k$, where q is

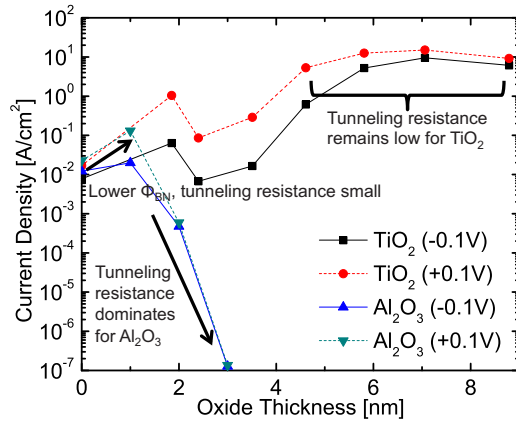


FIG. 4. (Color online) Current density of TiO_2 and Al_2O_3 devices as a function of oxide thickness at ± 0.1 V. TiO_2 devices show a substantial increase in current from reduction in Φ_{BN} . Al_2O_3 devices show an initial increase in current for the same reason but tunneling resistance with thicker Al_2O_3 quickly begins to dominate and reduces current density.

the electronic charge and k is the Boltzmann constant. The extracted Φ_{BN} was 0.104 eV (5.8 nm) and 0.065 eV (8.8 nm). This is a substantial reduction in Φ_{BN} from the 0 nm case, where $\Phi_{\text{BN}} \approx 0.58$ eV.² The reduction in Φ_{BN} combined with the low CBO explains the high current of the 5.8 and 8.8 nm TiO_2 devices.

In Fig. 3(b), the I-V characteristics of Al/ Al_2O_3 /Ge devices are shown. There is a slight current increase with 1 nm of Al_2O_3 at low biases, which can be attributed to reduction in Φ_{BN} . However, with 2 nm Al_2O_3 or more, the resistance due to tunneling through Al_2O_3 starts to become significant and the current drops markedly. In a very large scale integration (VLSI) application, it would be difficult to achieve the optimum Al_2O_3 thickness uniformly across a 300 mm wafer since the current depends exponentially on thickness. This illustrates an advantage of using TiO_2 to relax the uniformity requirements, since a wide range of thicknesses still yielded high currents.

Figure 4 shows the current densities at ± 0.1 V for the TiO_2 and Al_2O_3 devices as a function of oxide thickness. Both positive and negative voltages were plotted since in a MOSFET, one contact will be forward biased while the other will be reverse biased. The trend is similar for both voltages. We focus on low biases since the high current levels at high biases causes large potential drop across the lightly doped Si substrate. The TiO_2 devices show an increase in current density due to reduction in Φ_{BN} , until about 5.8 nm TiO_2 . With thicker TiO_2 , the current remains roughly constant. The highest current achieved was with 7.1 nm of TiO_2 , where the current increased by about $900\times$ at 0.1 V (1.7×10^{-2} to

15.1 A/cm^2) and $1200\times$ at -0.1 V (7.8×10^{-3} to 9.6 A/cm^2) compared to the 0 nm sample. In contrast, the Al_2O_3 devices show only a small initial increase in current up to 1 nm, again due to Φ_{BN} reduction. Tunneling resistance dominates with thicker Al_2O_3 , and the current drops significantly. These results illustrate the advantage of using a low CBO material such as TiO_2 in MIS contacts.

In summary, the TiO_2 /Ge interface was estimated to have a nearly zero CBO using SRPES, which makes it an excellent candidate material in an MIS contact due to its lower tunneling resistance. An ALD TiO_2 interfacial layer was used to decrease Φ_{BN} , as verified through temperature-dependent I-V measurements. We experimentally demonstrate that a TiO_2 interfacial layer is effective at significantly increasing current density for contacts to n-type Ge. This is, therefore, a promising method to increase the performance of n-channel Ge MOSFETs.

This research was supported by Intel Corporation and the MSD Focus Center (Contract No. 2009-MT-2051, Subcontract No. 5710002717-02). J.-Y. J. Lin acknowledges financial support from a Stanford Graduate Fellowship. Device fabrication was performed at the Stanford Nanofabrication Facility. Portions of this research were carried out at the Stanford Synchrotron Radiation Lightsource, a Directorate of SLAC National Accelerator Laboratory and an Office of Science User Facility operated for the U.S. Department of Energy Office of Science by Stanford University.

¹A. Dimoulas, P. Tsipas, A. Sotiropoulos, and E. K. Evangelou, *Appl. Phys. Lett.* **89**, 252110 (2006).

²T. Nishimura, K. Kita, and A. Toriumi, *Appl. Phys. Lett.* **91**, 123123 (2007).

³D. Connelly, C. Faulkner, P. A. Clifton, and D. E. Grupp, *Appl. Phys. Lett.* **88**, 012105 (2006).

⁴M. Kobayashi, A. Kinoshita, K. Saraswat, H.-S. P. Wong, and Y. Nishi, *J. Appl. Phys.* **105**, 023702 (2009).

⁵R. R. Lieten, S. Degroote, M. Kuijk, and G. Borghs, *Appl. Phys. Lett.* **92**, 022106 (2008).

⁶T. Nishimura, K. Kita, and A. Toriumi, *Appl. Phys. Express* **1**, 051406 (2008).

⁷Y. Zhou, M. Ogawa, X. Han, and K. L. Wang, *Appl. Phys. Lett.* **93**, 202105 (2008).

⁸A. M. Roy, J.-Y. J. Lin, and K. C. Saraswat, *IEEE Electron Device Lett.* **31**, 1077 (2010).

⁹B.-S. Jeong, D. P. Norton, and J. D. Budai, *Solid-State Electron.* **47**, 2275 (2003).

¹⁰K. Eufinger, D. Poelman, H. Poelman, R. De Gryse, and G. B. Marin, *Appl. Surf. Sci.* **254**, 148 (2007).

¹¹J. Aarik, A. Aidla, A.-A. Kiisler, T. Uustare, and V. Smmelselg, *Thin Solid Films* **305**, 270 (1997).

¹²V. V. Afanas'ev, A. Stesmans, A. Delabie, F. Bellenger, M. Houssa, and M. Meuris, *Appl. Phys. Lett.* **92**, 022109 (2008).