

Effect of Top-Gate Dielectric Deposition on the Performance of Indium Tin Oxide Transistors

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Abstract—We report ultrathin (~4 nm) channel indium tin oxide (ITO) transistors, comparing different precursors for atomic layer deposition (ALD) of the Al₂O₃ top-gate dielectric, and analyze the role of dielectric deposition on transistor performance and gate bias stress stability. Waterbased ALD leads to very negative threshold voltage (V_T), with devices remaining in the on-state. In contrast, both ozone and O₂-plasma precursors yield devices that can turn off, but ozone-based ALD devices have less negative $V_{\rm T}$ shift at short channel lengths, and relatively more positive V_{T} at all channel lengths. We achieve maximum drive current, $I_{max} \approx 260 \ \mu A/\mu m$ at $V_{DS} = 1$ V, on/off current ratio of >10¹⁰ (limited by the instrument's noise floor) for $L \approx 700$ nm ozone-Al₂O₃ top-gated transistors. Across multiple devices, the effective mobility is \sim 42 cm²V⁻¹s⁻¹ and contact resistance is ~376 $\Omega \cdot \mu$ m. The transistors also show good gate bias stability with normalized V_T shift of +0.12 V(MV/cm)⁻¹ at gate stress field >3 MV/cm, a \sim 3x improvement vs. our previous reports of uncapped ITO transistors.

Index Terms-ITO, transistors, atomic layer deposition, effective mobility, contact resistance, bias stress stability.

I. INTRODUCTION

XIDE semiconductors are compatible with back-end-of-Unite (BEOL) processing (≤ 500 °C) [1] and monolithic three-dimensional (3D) integration due to their large-scale, low-temperature deposition methods [2], [3], [4], [5]. They are promising candidates as channel materials for field-effect

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transistors (FETs), due to high drive currents and low off-state currents [2], [3], [6], [7]. Recently, ultra-short channel (~8 nm) 2.5 nm thin In₂O₃ FETs have shown \sim 3 mA/ μ m drive currents, but they suffer from poor on/off current ratio $<10^2$ [4]. On the other hand, indium gallium zinc oxide (IGZO) devices have demonstrated high on/off current ratio of $\sim 10^{12}$ but lower on-state current $\sim 6 \,\mu A/\mu m$ [5]. Among this family of materials, indium tin oxide (ITO) transistors have shown both good on- and off-state performance, with effective mobility $\mu_{\rm eff} \sim 55 \ {\rm cm^2 V^{-1} s^{-1}}$ [2] and high on/off current ratio $\sim 10^{11}$ [3], [8]. However, almost all ITO transistors have backgated (BG) structures with no channel encapsulation. Thus, the effect of top dielectric deposition on the performance and stability of ITO FETs remains unexplored.

In this work, we fabricate top-gated (TG) ITO transistors with three different precursors: H₂O, O₂-plasma, and ozone (O_3) , for atomic layer deposition (ALD) of the Al₂O₃ TG dielectric. We compare the role of the TG dielectric deposition on device behavior, under DC measurements and positive gate bias stress. In doing so, we determine the optimum precursor for Al₂O₃ top dielectric deposition on ITO channels, based on parameters like threshold voltage (V_T), on/off current ratio, mobility, drive current, minimum subthreshold swing and $V_{\rm T}$ stability.

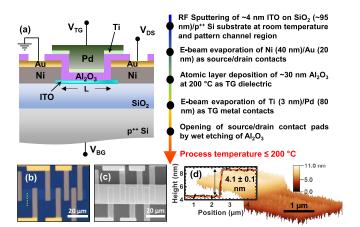
II. FABRICATION AND MEASUREMENT

The cross-sectional schematic of a device along with the fabrication steps are shown in Fig 1(a). First, we deposit \sim 4 nm thick ITO channels by radio frequency (RF) magnetron sputtering on SiO₂ (95 nm)/p⁺⁺ Si substrates, which are used as global back-gates for initial BG measurements. The ITO deposition is done at room temperature with 100 W RF power, 5 mTorr deposition pressure, and 9:1 Ar:O₂ ratio. The channel region is patterned by lift-off or wet etching in diluted HCl (1.7%). Next, we e-beam evaporate and lift-off Ni (40 nm)/Au (20 nm) as source/drain contacts. For the TG dielectric, ALD of ~30 nm Al₂O₃ film is done at 200 °C, with three different precursors: H₂O, O₂-plasma, and ozone (O₃). Then, we e-beam evaporate and lift-off Ti (3 nm)/Pd (80 nm) as TG metal contact, before opening the source/drain contact pads by wet etching of Al₂O₃. The temperature remains \leq 200 °C throughout all fabrication steps.

We fabricate transfer length method (TLM) structures with transistors of different channel lengths ($L \sim 0.7, 1.6, 2.8, 4.7,$ 7.3, and 9.8 μ m). Figs. 1(b,c) show an optical and a scanning electron microscope image of a TLM before and after TG dielectric and metal deposition, respectively. Fig. 1(d) shows

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(a) Schematic cross-section of the device structure and fabrication process flow of ITO transistors. (b) Optical image (before TG dielectric and metal deposition) and (c) scanning electron microscopy image (after TG dielectric and metal deposition) of transfer length method (TLM) structure. (d) Atomic force microscopy image across the green dotted line shown in (b); orange dashed line fitting the height profile (inset) shows the ITO channel thickness.

the atomic force microscopy profile across the green dotted line in Fig. 1(b), confirming the ITO thickness ~ 4 nm. All electrical measurements were done at room temperature, in air, using a probe station with a Keithley 4200-SCS.

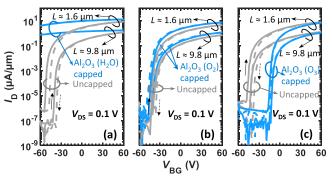
III. RESULTS AND DISCUSSION

A. Back-Gated (BG) Measurements

For initial analysis, we performed BG measurements using the p⁺⁺ Si substrate as a back-gate. The gray and blue curves in Fig. 2 show the BG transfer characteristics $(I_D - V_{BG})$ of the uncapped and Al₂O₃-capped devices, respectively, for two channel lengths ($L \sim 1.6 \ \mu m$ and 9.8 μm). The effect of ALD capping is shown with the three different precursors, H₂O, O₂, and O₃, one in each panel of Fig. 2. The transfer curves for all uncapped, as-deposited samples are similar, with similar $V_{\rm T}$. After ALD of H₂O-Al₂O₃ the $V_{\rm T}$ strongly shifts negatively, and the devices lose BG modulation [Fig. 2(a)]. This conductive channel could be caused by H-doping with residual H from the H₂O precursor, or by the highly reactive trimethylaluminum precursor stripping off chemisorbed oxygen or -OH groups from the oxide channel, generating oxygen vacancies which donate free electrons [9], [10]. However, after O₂-plasma and O₃-based ALD, the transistors retain good BG modulation, suggesting passivation of oxygen vacancies or reintroduction of -OH groups into the channel, unlike H₂Obased ALD [9]. For O₂-plasma ALD, $V_{\rm T}$ remains close to the as-deposited case [Fig. 2(b)], while O_3 -ALD shifts the V_T positively, closer to 0 V [Fig. 2(c)]. A control experiment with the same number of *only* oxidant (O_2 and O_3) pulses revealed that both types of reactants shift the $V_{\rm T}$ positively, and thus the duration, order, and number of O2 or O3 oxidant cycles may be tuned to achieve a desired $V_{\rm T}$.

B. Top-Gated (TG) Measurements

Fig. 3(a) shows the TG transfer characteristics $(I_{\rm D}-V_{\rm TG})$ of 1.6 μ m devices with the three types of Al₂O₃ as TG dielectric. As for the BG measurement, the H₂O-precursor transistors show no TG modulation. On the other hand, O₂ and O_3 -Al₂O₃ devices are modulated by the TG (like the BG), but the O_2 -plasma Al_2O_3 device displays more negative $V_{\rm T}$ (-1.1 V vs. 0.03 V) and degraded minimum subthreshold



Back-gated (BG) transfer characteristics for ITO transistors Fig. 2. (with L \sim 1.6 and 9.8 μ m) before (gray) and after (blue) capping with Al₂O₃ by ALD using (a) H₂O, (b) O₂-plasma, and (c) O₃ precursor. Solid and dashed small arrows indicate the voltage sweep directions and correspond to the solid and dashed curves, respectively. For all measurements, BG leakage currents were $< 10^{-5} \mu A/\mu m$.

swing (~430 mV/dec vs. ~230 mV/dec). Fig. 3(b) shows $I_{\rm D}$ - V_{TG} curves for O₂- and O₃-Al₂O₃ devices with $L \approx 700$ nm. Here, the O₃-Al₂O₃ device shows $I_{\rm max} \approx 260 \ \mu {\rm A}/\mu {\rm m}$ at $V_{\text{TG}} = 20 \text{ V}$ and $V_{\text{DS}} = 1 \text{ V}$, on/off current ratio >10¹⁰ (limited by the noise floor of the instrument), extrinsic field-effect mobility $\mu_{\rm FE,ext} \approx 38 \text{ cm}^2 \text{V}^{-1} \text{s}^{-1}$ and $V_{\rm T} = -1.6$ V. The $\mu_{\rm FE,ext}$ and $V_{\rm T}$ are extracted at peak transconductance. Note, at this shorter channel length, the $V_{\rm T}$ shifts more negatively for O_2 -plasma Al₂O₃ device (-7.9 V vs. -1.6 V for O_3), together with increased hysteresis and drain-induced barrier lowering (|DIBL| $\approx 261 \pm 182$ mV/V for O₂-plasma vs. $128 \pm 66 \text{ mV/V}$ for O₃). Such degradation may be explained by the oxygen vacancies in the channel and under the contacts, acting as dopants and defect centers, which can cause charge trapping and affect the semiconductor capacitance. However, this needs to be investigated in more detail in future studies.

Figs. 3(c,d) show output characteristics $(I_D - V_{DS})$ for \sim 700 nm long devices with O₂-plasma and O₃-based Al₂O₃ TG dielectric, respectively. The O₃-Al₂O₃ device has a less negative $V_{\rm T} \approx -3$ V vs. -10 V for O₂-plasma. In addition, for the same TG overdrive voltage $(V_{TG} - V_T = 7 \text{ V})$ at $V_{\rm DS} = 4$ V, the drive current of the O₃-Al₂O₃ sample is almost double that of the O₂-Al₂O₃ device (498 μ A/ μ m vs. 266 μ A/ μ m).

Fig. 4(a) displays $V_{\rm T}$ roll-off *i.e.* decreasing $V_{\rm T}$ at shorter channel lengths, ostensibly due to oxygen scavenging at the contacts [5], [11] for both O₂ and O₃-Al₂O₃ samples. The O₃ precursor sample shows less $V_{\rm T}$ roll-off (-4.4 V) compared to the O_2 -plasma (-15.9 V), which could be due to the passivation of a dissimilar number of oxygen vacancies in the channel during ALD. Here, the $V_{\rm T}$ roll-off is the $V_{\rm T}$ difference between the longest (~9.8 μ m) and shortest (~700 nm) channels. Notably, $V_{\rm T} > 0$ V for the O₃-Al₂O₃ sample down to ~1.6 μ m channel length, only becoming negative ($V_{\rm T,avg} \approx -2.2$ V) for the shortest channel (\sim 700 nm). On the other hand, the average $V_{\rm T}$'s for the O₂-Al₂O₃ devices remain negative for all channel lengths. Further reduction of the dielectric and channel thickness could achieve a positive $V_{\rm T}$ with turn-on voltage close to 0 V, at the expense of lower on-state current due to the thinner channel [12].

Fig. 4(b) shows the plot for total resistance (R_{Total} , averaged over 8 to 10 devices at each channel length) vs. channel length L for both O_2 and O_3 -Al₂O₃ samples. This allows the extraction of contact resistance ($R_{\rm C}$) and effective mobility ($\mu_{\rm eff}$) at a given electron density, $n \approx 2.6 \times 10^{13}$ cm⁻², estimated as Authorized licensed use limited to: Stanford University. Downloaded on May 24,2023 at 18:38:47 UTC from IEEE Xplore. Restrictions apply.

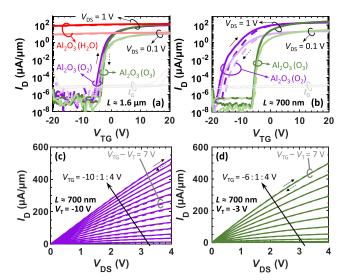


Fig. 3. Top-gated (TG) transfer characteristics for (a) $L \approx 1.6 \ \mu m$ and (b) $L \approx 700 \ nm$. For all measurements, TG leakage currents ($I_{\rm G}$) were $< 10^{-5} \ \mu A/\mu m$. (c) Output characteristics ($I_{\rm D}$ - $V_{\rm DS}$) for \sim 700 nm devices with O₂ and (d) O₃-Al₂O₃ TG dielectric. Solid and dashed small arrows mark voltage sweep directions and correspond to the solid and dashed curves, respectively. Purple and green are used to denote O₂-plasma and ozone (O₃) Al₂O₃ samples, respectively, in all panels.

 $n = C_{\rm TG}(V_{\rm TG} - V_{\rm T})/q$, where $C_{\rm TG} \approx 0.28 \ \mu {\rm F/cm^2}$ for O_2 -Al₂ O_3 and 0.29 μ F/cm² for O_3 -Al₂ O_3 is the top-gate capacitance and q is the elementary charge. The O₃-Al₂O₃ sample shows lower $R_{\rm C} \sim 376~\Omega \cdot \mu {
m m}$ and higher $\mu_{\text{eff}} \sim 41.7 \text{ cm}^2 \text{V}^{-1} \text{s}^{-1}$ compared to the O₂-Al₂O₃ sample. The difference in the $R_{\rm C}$ and mobility values for the two samples is also present in the BG measurement before ALD capping, as evident from the lower drive current of the O2-plasma sample before ALD [Fig. 2(b)]. Before ALD capping, the samples have $R_{\rm C}$ and $\mu_{\rm eff}$ of ~495 $\Omega \cdot \mu m$ and ~28.6 cm²V⁻¹s⁻¹ [Fig. 2(b)] vs. $\sim 313 \Omega \cdot \mu m$ and $\sim 45.2 \text{ cm}^2 \text{V}^{-1} \text{s}^{-1}$ [Fig. 2(c)] respectively, at $n = 1.1 \times 10^{13}$ cm⁻². This difference could be due to variation in as-deposited ITO films and/or Ni contacts, possibly caused by different chamber conditions during different runs. Notably, the Al_2O_3 capping introduces further changes in these parameters, possibly due to passivation of oxygen vacancies in the ITO film during ALD.

Figs. 4(c,d) show the positive bias stress stability of $\sim 1.6 \ \mu m \log O_2$ and O_3 -Al₂O₃ samples at a TG bias stress field of ~3.33 MV/cm. Interestingly, both samples show similar $V_{\rm T}$ shift ($\Delta V_{\rm T} \approx +0.4$ V) due to positive bias stress. This corresponds to a positive normalized $V_{\rm T}$ shift of 0.12 V(MV/cm)⁻¹, which is $\sim 3 \times$ more stable compared to uncapped ITO devices under similar back-gate stress field reported in our prior work [13]. Figs. 4(e,f) show the evolution of $V_{\rm T}$ and $\Delta V_{\rm T}$ with stress time ($t_{\rm stress}$) averaged across 5 devices each from O₂ and O₃-Al₂O₃ samples. The dashed lines in Fig. 4(f) are the stretched-exponential model of $\Delta V_{\rm T}$ time dependence [14], based on charge trapping and redistribution from the channel interface into the dielectric. The model is $\Delta V_{\rm T} = \Delta V_{\rm T0} \{1 - \exp[-(t/\tau)^{\beta}]\}$, where $\Delta V_{\rm T0}$, τ , and β represent $\Delta V_{\rm T}$ at infinite time, the average trapping time of electrons, and a fitting exponent, respectively. In our work, these parameters are ~ 0.6 V, 314 s and 0.21 for O₂-Al₂O₃ and ~ 0.42 V, 4.4 s and 0.39 for O₃-Al₂O₃, respectively. During bias stress testing, we took DC measurements of the transfer curves, which took ~ 14 s with no additional hold or delay

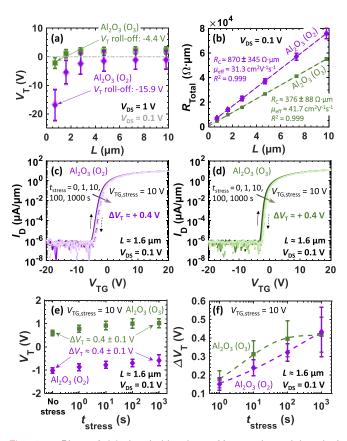


Fig. 4. Plots of (a) threshold voltage $V_{\rm T}$ vs. channel length *L*, (b) total resistance $R_{\rm Total}$ vs. *L* at electron density, $n = 2.6 \times 10^{13}$ cm⁻². Error bars represent the standard deviation among 8 to 10 devices at each channel length. Insets in (b) list the estimated contact resistance $R_{\rm C}$ (from the vertical axis intercept of the linear fit) and the effective mobility, $\mu_{\rm eff}$, whereas R^2 denotes the goodness of fit. (c) Positive bias stress analysis at $V_{\rm TG, stress} = 10$ V for ~1.6 μ m devices with O₂ and (d) O₃-Al₂O₃ TG dielectric. Solid and dashed small arrows indicate the voltage sweep directions and correspond to the solid and dashed curves, respectively. Evolution of (e) $V_{\rm T}$ and (f) $V_{\rm T}$ shift, $\Delta V_{\rm T}$, vs. stress time, $t_{\rm stress}$ for $V_{\rm TG, stress} = 10$ V. Error bars in (e, f) mark the variability among 5 devices. Dashed lines in (f) are fit to the stretched-exponential model in [14]. Purple and green are used to denote O₂-plasma and ozone (O₃) Al₂O₃ samples, respectively, in all panels.

time in between the stress and measurement. The stress time was accumulated over all cycles.

IV. CONCLUSION

We report top-gated ITO transistors with good $R_{\rm C} \sim 376 \ \Omega \cdot \mu \text{m}$ and $\mu_{\rm eff} \sim 42 \ \text{cm}^2 \text{V}^{-1} \text{s}^{-1}$. We achieve $I_{\rm max} \approx 260 \ \mu \text{A}/\mu \text{m}$ at $V_{\rm DS} = 1 \ \text{V}$ in a $\sim 700 \ \text{nm}$ long channel with O₃-based Al₂O₃ as the top dielectric. We show that, both O₃ and O₂-plasma precursors during the ALD of the top-gate dielectric produce bias stress stable devices. Using O₃ precursor over O₂-plasma demonstrates the advantage for minimizing the $V_{\rm T}$ roll-off after ALD as well as reducing short channel effects. These results are important for ITO channel transistor applications in BEOL and 3D integration. They also open a pathway towards enhancing oxide transistor performance by channel length scaling and improved gate control, *i.e.* dual-gating or gate-all-around structures.

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