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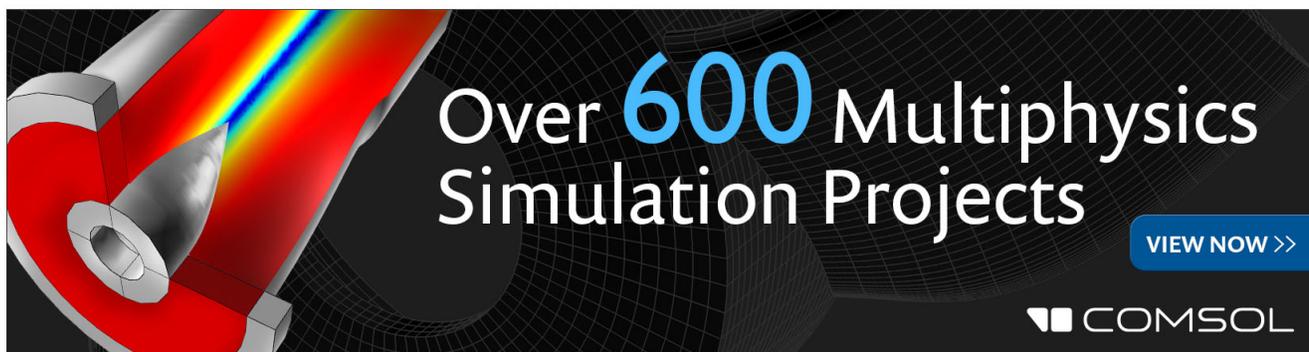
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Effects of Ta incorporation in La_2O_3 gate dielectric of InGaZnO thin-film transistor

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The effects of Ta incorporation in La_2O_3 gate dielectric of amorphous InGaZnO thin-film transistor are investigated. Since the Ta incorporation is found to effectively enhance the moisture resistance of the La_2O_3 film and thus suppress the formation of $\text{La}(\text{OH})_3$, both the dielectric roughness and trap density at/near the InGaZnO/dielectric interface can be reduced, resulting in a significant improvement in the electrical characteristics of transistor. Among the samples with different Ta contents, the one with a Ta/(Ta+La) atomic ratio of 21.7% exhibits the best performance, including high saturation carrier mobility of $23.4 \text{ cm}^2/\text{V}\cdot\text{s}$, small subthreshold swing of $0.177 \text{ V}/\text{dec}$, and negligible hysteresis. Nevertheless, excessive incorporation of Ta can degrade the device characteristics due to newly generated Ta-related traps. © 2014 AIP Publishing LLC. [<http://dx.doi.org/10.1063/1.4869761>]

Recently, amorphous InGaZnO (a-IGZO) thin-film transistors (TFTs) have been extensively explored for the application in various flat-panel displays.^{1,2} Compared to conventional amorphous silicon or organic TFTs with a field-effect carrier mobility of $\sim 1 \text{ cm}^2/\text{V}\cdot\text{s}$,^{3,4} a-IGZO TFTs typically exhibit a mobility higher than $10 \text{ cm}^2/\text{V}\cdot\text{s}$,⁵ which can translate to higher switching speed for electronic devices. In addition, a-IGZO TFTs offer better uniformity in device characteristics compared with polycrystalline silicon TFTs and have more excellent transparency to visible light than all the silicon-based devices.⁶ In order to reduce their operating voltage, high-k materials have been adopted as gate dielectric in a-IGZO TFTs.^{7,8} Among them, rare-earth oxide La_2O_3 is regarded as one of the most promising candidates due to its relatively high dielectric constant and large band gap ($\sim 6 \text{ eV}$).^{9,10} However, La_2O_3 is hygroscopic, which can deteriorate both the dielectric constant and surface roughness of La_2O_3 film due to the formation of hydroxide $\text{La}(\text{OH})_3$,¹¹ and thus induce degradations in the electrical characteristics of a-IGZO TFTs.¹² Fortunately, the doping of other elements, for example, Y, was reported to be an effective method to suppress the moisture absorption of La_2O_3 film.¹³ In this work, the doping of Ta in La_2O_3 film is proposed due to the fact that Ta_2O_5 can exhibit both very high dielectric constant and excellent step coverage,¹⁴ and accordingly the effects of Ta incorporation in La_2O_3 gate dielectric of a-IGZO TFTs are investigated. Three samples of a-IGZO TFTs with different Ta/(Ta+La) atomic ratios are prepared while one sample with pure La_2O_3 gate dielectric is also fabricated as the control sample.

Each sample was fabricated on a p-type (100) silicon wafer with a resistivity of $0.01\text{--}0.02 \text{ }\Omega\cdot\text{cm}$ which acted as both the substrate and gate electrode. First, a 40-nm dielectric film was deposited by a sputtering system under a radio-frequency (RF) power supply for a ceramic target of La_2O_3 and a direct-current (DC) supply for a metal target of Ta in a

mixed ambient of Ar plus O_2 . The RF power was fixed at 40 W while the DC supply was set to be 0 A, 0.03 A, 0.04 A, and 0.05 A for sample La_2O_3 , sample LaTaO_A , sample LaTaO_B , and sample LaTaO_C , respectively, so as to realize different atomic ratios of Ta/(Ta+La) in dielectric films. Second, an annealing treatment at $400 \text{ }^\circ\text{C}$ in an N_2 ambient for 30 min followed. Subsequently, the four samples received the deposition of a 60-nm a-IGZO active layer through RF sputtering from a ceramic target (Ga_2O_3 : In_2O_3 : $\text{ZnO} = 1: 1: 1$). After that, a lift-off process was utilized to form the source/drain electrodes, which were composed of 20-nm Ti and 80-nm Au. The channel width (W) and channel length (L) were $100 \text{ }\mu\text{m}$ and $10 \text{ }\mu\text{m}$, respectively. Finally, all the samples were annealed in a forming-gas (N_2 : $\text{H}_2 = 95: 5$) ambient at $350 \text{ }^\circ\text{C}$ for 20 min so that the contact resistance of the source/drain electrodes was reduced. In addition, metal-insulator-semiconductor capacitor was also prepared beside each sample to monitor the gate-oxide capacitance per unit area (C_{ox}). The current-voltage (I-V) curves of the TFTs were measured by a HP 4145B semiconductor parameter analyzer. Furthermore, the structural properties of the dielectric films were studied based on X-ray photoelectron spectroscopy (XPS) and atomic force microscopy (AFM).

Fig. 1 shows the XPS spectra of (a) La $3d_{5/2}$ and (b) O $1s$ core levels for the dielectric films. The binding energies have been corrected for sample charging effect with reference to the C $1s$ line at 285.0 eV . Accordingly to the XPS result, the atomic ratio of Ta/(Ta+La) is 0%, 21.7%, 30.6%, and 69.1% for sample La_2O_3 , sample LaTaO_A , sample LaTaO_B , and sample LaTaO_C , respectively. As shown in Fig. 1(a), the La_2O_3 film exhibits an obvious shoulder at the high binding energy side of the La $3d_{5/2}$ main peak, suggesting the presence of La-OH bond due to the moisture absorption of La_2O_3 . Furthermore, the La $3d_{5/2}$ peak of the La_2O_3 film (located at 834.7 eV) shifts to a higher binding energy compared to the ideal La_2O_3 reference peak (located at 834.0 eV) while being consistent with the reported peak for LaOOH film (located at $834.8 \pm 0.2 \text{ eV}$) in peak location,¹⁵ further revealing the

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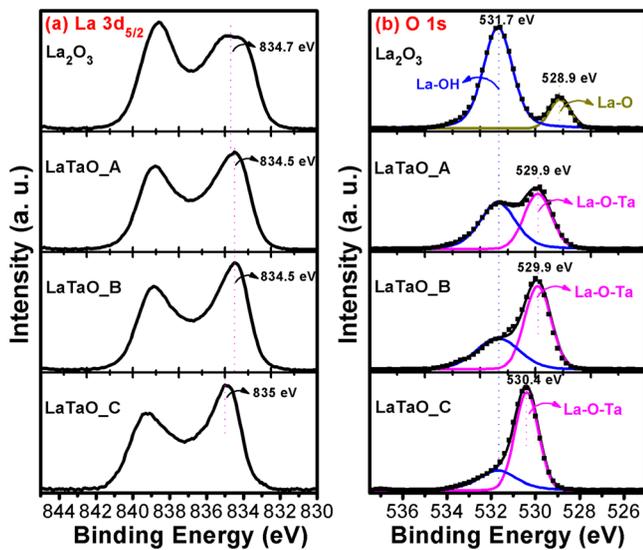


FIG. 1. XPS spectrum of (a) La $3d_{5/2}$ and (b) O $1s$ for the dielectric films in sample_La₂O₃, sample_LaTaO_A, sample_LaTaO_B, and sample_LaTaO_C.

existence of La(OH)₃ in the La₂O₃ film. With Ta incorporation, the La $3d_{5/2}$ main peak becomes sharper, which is more obvious for higher Ta/(Ta + La) atomic ratio in the LaTaO film. This result is presumably due to the enhancement in the moisture resistance of the La₂O₃ film and accordingly the suppression in the formation of La(OH)₃ after Ta incorporation. Nevertheless, the La $3d_{5/2}$ peak related to sample_LaTaO_C (located at 835.0 eV) shifts to an even higher binding energy in comparison to sample_La₂O₃, suggesting the formation of La-O-Ta bond in the LaTaO film. The O $1s$ spectra of the La₂O₃ and LaTaO films are shown in Fig. 1(b), and each

fitting peak follows the general shape of the Lorentzian–Gaussian function. As for the La₂O₃ film, the two O $1s$ peaks correspond to La-O bond (located at 528.9 eV)¹⁵ and La-OH bond (located at 531.7 eV), respectively. Moreover, the O $1s$ peak corresponding to La-O bond has a much higher intensity compared with that corresponding to La-OH bond, indicating that most of La atoms in the La₂O₃ film have been transformed into La(OH)₃ due to moisture absorption. With Ta incorporation, the intensity of the O $1s$ peak corresponding to La-OH bond decreases. Moreover, this effect becomes more obvious for higher Ta/(Ta + La) atomic ratio in the LaTaO film. This result further demonstrates the suppressed formation of La(OH)₃ due to the enhanced moisture resistance of the La₂O₃ film after Ta incorporation. Meanwhile, the O $1s$ peak corresponding to La-O bond has been completely replaced by that corresponding to La-O-Ta bond (located at a higher binding energy) for each Ta-incorporated sample. Moreover, the O $1s$ peak corresponding to La-O-Ta bond shifts to a higher binding energy with increased Ta/(Ta + La) atomic ratio as reflected by the comparison between sample_LaTaO_C and the other two Ta-incorporated samples, and similar effect also occurs to the La $3d_{5/2}$ spectrum in Fig. 1(a).

Fig. 2 shows the AFM images of the La₂O₃ and LaTaO films with a measurement area of $1 \mu\text{m} \times 1 \mu\text{m}$. The La₂O₃ film, with a RMS of 1.28 nm, exhibits the roughest surface among the dielectric films, which should result from non-uniform volume expansion of the La₂O₃ film after moisture absorption.¹¹ With Ta incorporation, the dielectric roughness is significantly reduced, which is more obvious for higher Ta/(Ta + La) atomic ratio. Accordingly, the RMS value of LaTaO film in sample_LaTaO_A, sample_LaTaO_B, and

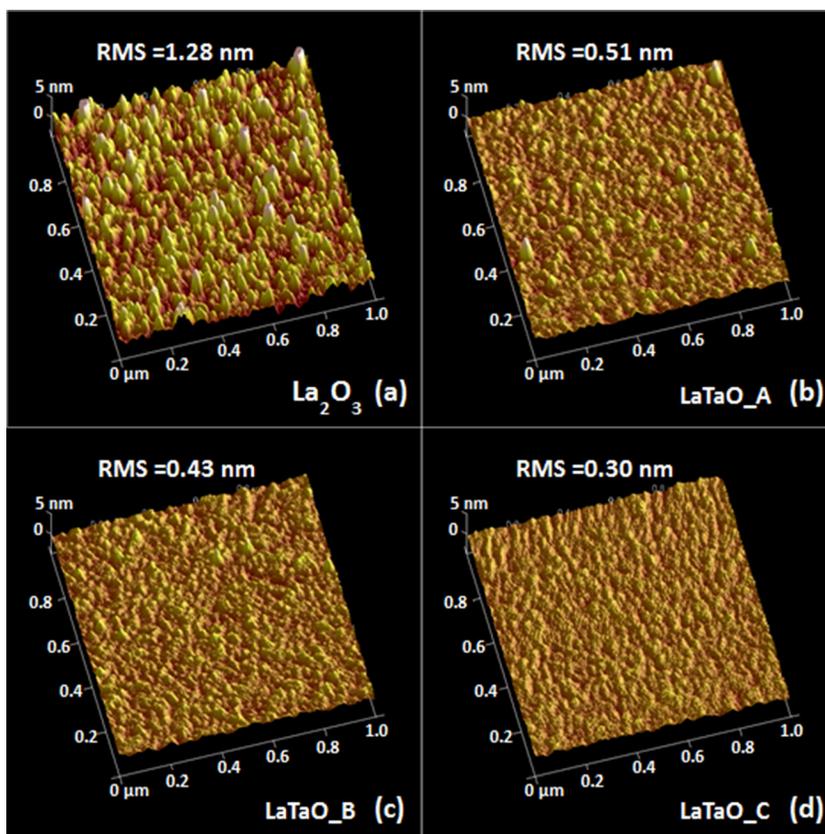


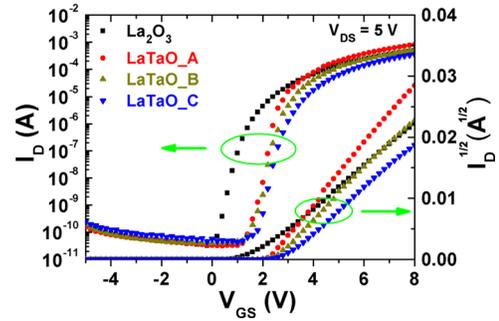
FIG. 2. AFM image of dielectric films in (a) sample_La₂O₃, (b) sample_LaTaO_A, (c) sample_LaTaO_B, and (d) sample_LaTaO_C.

TABLE I. Extracted electrical parameters of the a-IGZO TFT's.

Sample	La ₂ O ₃	LaTaO_A	LaTaO_B	LaTaO_C
La deposition (RF/W)	40	40	40	40
Ta deposition (DC/A)	0	0.03	0.04	0.05
Atomic ratio of Ta/(Ta + La)	0%	21.7%	30.6%	69.1%
μ_{sat} (cm ² /V·s)	12.1	23.4	16.3	11.0
V_{TH} (V)	1.85	2.40	2.66	2.97
SS (V/dec)	0.234	0.177	0.201	0.217
ΔV_{H} (V)	-0.76	0.10	1.29	2.34
I_{on} (μA)	494	810	520	349
$I_{\text{on}}/I_{\text{off}}$	1.5×10^7	2.6×10^7	1.3×10^7	8.6×10^6
C_{ox} ($\mu\text{F}/\text{cm}^2$)	0.231	0.262	0.269	0.279
Dielectric constant	10.4	11.8	12.2	12.6

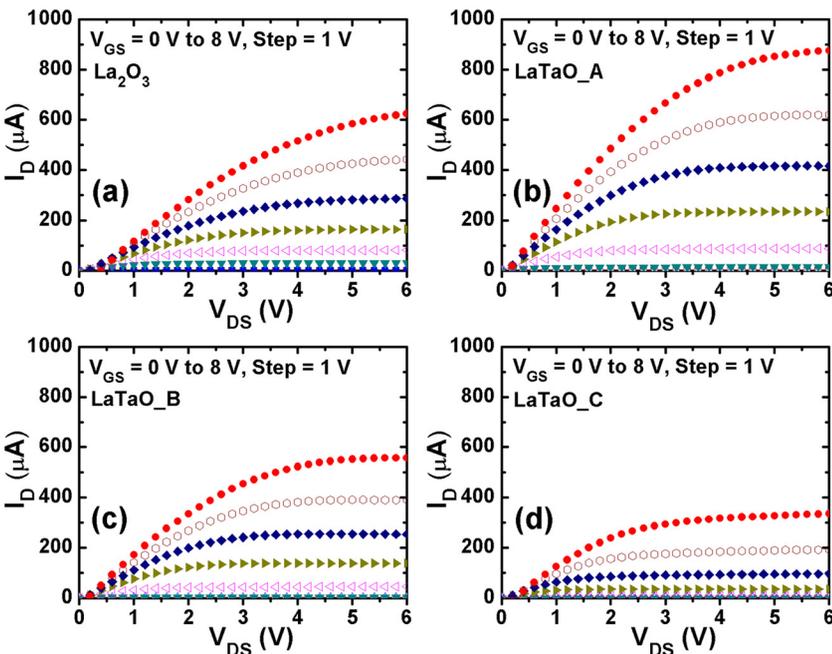
sample_LaTaO_C is 0.51 nm, 0.43 nm, and 0.30 nm, respectively, further demonstrating that Ta incorporation is an effective way to enhance the moisture resistance of La₂O₃ film and accordingly reduce its surface roughness. In addition, the enhanced moisture resistance of the La₂O₃ film also effectively suppresses the deterioration of its dielectric constant, and thus results in a continuous increase of dielectric constant associated with increasing Ta incorporation as listed in Table I. As compared to 3.9 of conventional SiO₂ dielectric, a larger dielectric constant of the LaTaO film (~ 12) is conducive to achieving higher-performance TFT with smaller operating voltage and larger output current.

Fig. 3 exhibits the transfer characteristics of the a-IGZO TFTs: drain current (I_{D}) vs. gate-to-source voltage (V_{GS}) and $I_{\text{D}}^{1/2}$ vs. V_{GS} at a drain-to-source voltage (V_{DS}) of 5 V. The saturation carrier mobility (μ_{sat}), threshold voltage (V_{TH}), subthreshold swing (SS), on current (I_{on}), and on-off current ratio ($I_{\text{on}}/I_{\text{off}}$) of the devices are extracted from Fig. 3 and listed in Table I. Among them, μ_{sat} and V_{TH} are calculated from a linear fitting to the plot of $I_{\text{D}}^{1/2}$ versus V_{GS} , which is based on the I-V equation of field-effect transistor operating in the saturation region

FIG. 3. Transfer characteristics of the a-IGZO TFTs in sample_La₂O₃, sample_LaTaO_A, sample_LaTaO_B, and sample_LaTaO_C.

$$I_{\text{D}} = (\mu_{\text{sat}} C_{\text{ox}} W / 2L) (V_{\text{GS}} - V_{\text{TH}})^2. \quad (1)$$

By comparing sample_La₂O₃ and sample_LaTaO_A, μ_{sat} is nearly doubled from 12.1 cm²/V·s to 23.4 cm²/V·s with a reduction of SS from 0.234 V/dec to 0.177 V/dec due to the Ta incorporation in the La₂O₃ gate dielectric. It is believed that the reduction in dielectric roughness can induce a smoother a-IGZO/dielectric interface, thus resulting in an increase in carrier mobility due to reduced surface-roughness scattering on the carriers.¹⁶ In addition, carrier mobility can also be improved by reducing the trap density at/near the a-IGZO/dielectric interface because of the restraint of electron trapping. Hence, the increase in μ_{sat} mentioned above can be attributed to smoother dielectric surface as well as lower trap density at/near the a-IGZO/dielectric interface, which are supported by the smaller values of RMS and SS, respectively.^{1,17} Furthermore, it was reported that a large number of oxygen vacancies are easily generated in rare-earth oxide film due to the formation of hydroxide after reacting with moisture.¹⁸ Hence, it is believed that the high trap density at/near the a-IGZO/dielectric interface in sample_La₂O₃ is related to the oxygen vacancies originated from the hygroscopicity of La₂O₃. In addition, a smaller V_{TH} (1.85 V) of sample_La₂O₃ in comparison to V_{TH} (2.40 V) of

FIG. 4. Output characteristics of the a-IGZO TFTs: (a) sample_La₂O₃; (b) sample_LaTaO_A; (c) sample_LaTaO_B; and (d) sample_LaTaO_C.

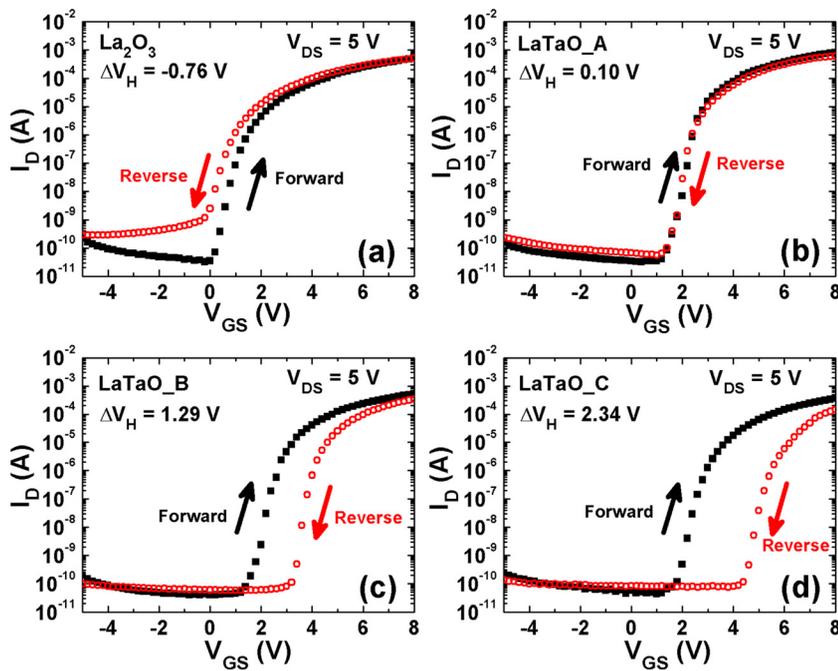


FIG. 5. Transfer characteristics of the a-IGZO TFTs measured under the forward and reverse V_{GS} sweepings: (a) sample_La₂O₃; (b) sample_LaTaO_A; (c) sample_LaTaO_B; and (d) sample_LaTaO_C.

sample_LaTaO_A can further reveal the existence of oxygen vacancies in the La₂O₃ gate dielectric as well as their reduction after Ta incorporation because oxygen vacancies can act as donor-like traps, inducing a negative shift of transfer curve and a degradation of SS in a-IGZO TFTs.¹⁹ Besides, I_{on} and I_{on}/I_{off} are increased from 494 μ A and 1.5×10^7 to 810 μ A and 2.6×10^7 , respectively, by Ta incorporation due to the improvement in carrier mobility. However, for further increase of Ta incorporation in the gate dielectric, the electrical characteristics of the TFT start to degrade even with a smoother dielectric surface, as reflected by the results of sample_LaTaO_B and sample_LaTaO_C. This should be ascribed to the creation of new Ta-related traps because a high density of defect states generally exists in Ta₂O₅ film,²⁰ which can be supported by the continual degradation of SS with increasing Ta/(Ta + La) atomic ratio in the LaTaO gate dielectric. Fig. 4 displays the output characteristics of the samples, which clearly exhibit an n-type enhancement mode. According to the comparison between sample_La₂O₃ and sample_LaTaO_A, the output current of the TFT is significantly increased by the Ta incorporation in the La₂O₃ gate dielectric due to the increase in carrier mobility. A continual reduction in output current with increasing Ta/(Ta + La) atomic ratio in the LaTaO gate dielectric is observed, which is consistent with the degradation in SS due to the generation of new Ta-related traps.

As shown in Fig. 5, the hysteresis properties of the samples are investigated according to the transfer characteristics measured under forward and reverse V_{GS} sweepings successively. ΔV_H is defined as the V_{TH} shift in the hysteresis loop. It is found that sample_La₂O₃ exhibits an obvious anticlockwise hysteresis with a negative ΔV_H (-0.76 V), further revealing the existence of donor-like traps at/near the a-IGZO/dielectric interface,²¹ which is due to the introduction of oxygen vacancies in the La₂O₃ film after moisture absorption. These donor-like traps can induce electron-detrapping or hole-trapping and become positively charged¹⁹ during the forward V_{GS} sweep of the hysteresis measurement. As a result, a

decrease of V_{TH} is observed during the subsequent backward V_{GS} sweep. With Ta incorporation in the La₂O₃ gate dielectric, negligible hysteresis is exhibited by sample_LaTaO_A ($\Delta V_H = 0.10$ V), which further demonstrates the reduction in the trap density at/near the a-IGZO/dielectric interface due to the enhanced moisture resistance of the dielectric film and thus suppressed generation of oxygen vacancies. In addition, the generation of new Ta-related traps, which are acceptor-like and prefer to capture electrons, has also been revealed by the continual enhancement of clockwise hysteresis with increasing Ta/(Ta + La) atomic ratio in the LaTaO gate dielectric. As a result, larger ΔV_H is exhibited by sample_LaTaO_B ($\Delta V_H = 1.29$ V) and sample_LaTaO_C ($\Delta V_H = 2.34$ V). A similar phenomenon of different signs of ΔV_H for hysteresis related to donor-like and acceptor-like traps has also been observed in other work.²²

In this work, the impact of Ta incorporation in La₂O₃ gate dielectric on the electrical characteristics of a-IGZO TFT has been studied. It is found that Ta incorporation can effectively enhance the moisture resistance of the La₂O₃ film and suppress the formation of La(OH)₃, thus reducing the dielectric roughness as well as the trap density at/near the a-IGZO/dielectric interface. Accordingly, the electrical characteristics of the TFT are significantly improved as reflected by nearly doubled μ_{sat} , reduced SS, suppressed hysteresis, and increased output current. However, excessive incorporation of Ta in the gate dielectric can degrade the device characteristics due to the creation of new Ta-related traps. In summary, these results demonstrate the potential use of LaTaO gate dielectric for making high-performance a-IGZO TFTs.

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