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Cite as: Appl. Phys. Lett. **104**, 123505 (2014); <https://doi.org/10.1063/1.4869761>

Submitted: 25 December 2013 . Accepted: 17 March 2014 . Published Online: 26 March 2014

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## Effects of Ta incorporation in $\text{La}_2\text{O}_3$ gate dielectric of InGaZnO thin-film transistor

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(Received 25 December 2013; accepted 17 March 2014; published online 26 March 2014)

The effects of Ta incorporation in  $\text{La}_2\text{O}_3$  gate dielectric of amorphous InGaZnO thin-film transistor are investigated. Since the Ta incorporation is found to effectively enhance the moisture resistance of the  $\text{La}_2\text{O}_3$  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.<sup>1,2</sup> Compared to conventional amorphous silicon or organic TFTs with a field-effect carrier mobility of  $\sim 1 \text{ cm}^2/\text{V}\cdot\text{s}$ ,<sup>3,4</sup> a-IGZO TFTs typically exhibit a mobility higher than  $10 \text{ cm}^2/\text{V}\cdot\text{s}$ ,<sup>5</sup> 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.<sup>6</sup> In order to reduce their operating voltage, high- $k$  materials have been adopted as gate dielectric in a-IGZO TFTs.<sup>7,8</sup> Among them, rare-earth oxide  $\text{La}_2\text{O}_3$  is regarded as one of the most promising candidates due to its relatively high dielectric constant and large band gap ( $\sim 6 \text{ eV}$ ).<sup>9,10</sup> However,  $\text{La}_2\text{O}_3$  is hygroscopic, which can deteriorate both the dielectric constant and surface roughness of  $\text{La}_2\text{O}_3$  film due to the formation of hydroxide  $\text{La}(\text{OH})_3$ ,<sup>11</sup> and thus induce degradations in the electrical characteristics of a-IGZO TFTs.<sup>12</sup> Fortunately, the doping of other elements, for example, Y, was reported to be an effective method to suppress the moisture absorption of  $\text{La}_2\text{O}_3$  film.<sup>13</sup> In this work, the doping of Ta in  $\text{La}_2\text{O}_3$  film is proposed due to the fact that  $\text{Ta}_2\text{O}_5$  can exhibit both very high dielectric constant and excellent step coverage,<sup>14</sup> and accordingly the effects of Ta incorporation in  $\text{La}_2\text{O}_3$  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  $\text{La}_2\text{O}_3$  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  $\text{La}_2\text{O}_3$  and a direct-current (DC) supply for a metal target of Ta in a

mixed ambient of Ar plus  $\text{O}_2$ . 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  $\text{La}_2\text{O}_3$ , sample  $\text{LaTaO}_A$ , sample  $\text{LaTaO}_B$ , and sample  $\text{LaTaO}_C$ , 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  $\text{N}_2$  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 ( $\text{Ga}_2\text{O}_3$ :  $\text{In}_2\text{O}_3$ :  $\text{ZnO} = 1: 1: 1$ ). 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 ( $\text{N}_2$ :  $\text{H}_2 = 95: 5$ ) 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 ( $C_{\text{ox}}$ ). 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  $285.0 \text{ eV}$ . Accordingly to the XPS result, the atomic ratio of Ta/(Ta+La) is 0%, 21.7%, 30.6%, and 69.1% for sample  $\text{La}_2\text{O}_3$ , sample  $\text{LaTaO}_A$ , sample  $\text{LaTaO}_B$ , and sample  $\text{LaTaO}_C$ , respectively. As shown in Fig. 1(a), the  $\text{La}_2\text{O}_3$  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  $\text{La}_2\text{O}_3$ . Furthermore, the La  $3d_{5/2}$  peak of the  $\text{La}_2\text{O}_3$  film (located at  $834.7 \text{ eV}$ ) shifts to a higher binding energy compared to the ideal  $\text{La}_2\text{O}_3$  reference peak (located at  $834.0 \text{ eV}$ ) while being consistent with the reported peak for  $\text{LaOOH}$  film (located at  $834.8 \pm 0.2 \text{ eV}$ ) in peak location,<sup>15</sup> further revealing the

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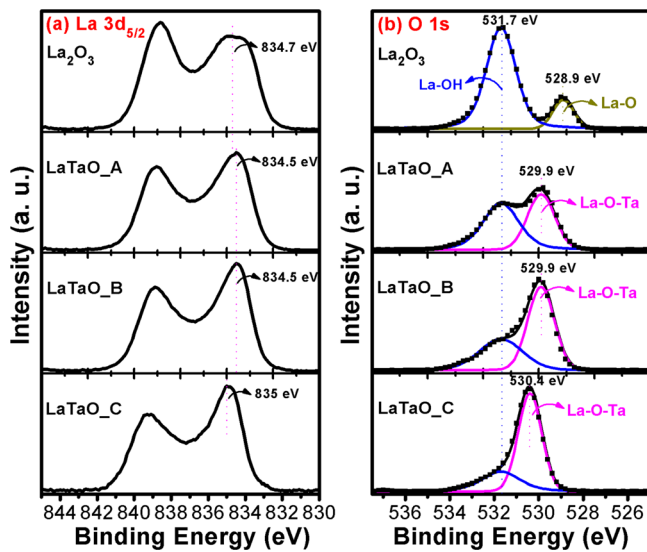


FIG. 1. XPS spectrum of (a) La 3d<sub>5/2</sub> and (b) O 1s for the dielectric films in sample\_La<sub>2</sub>O<sub>3</sub>, sample\_LaTaO\_A, sample\_LaTaO\_B, and sample\_LaTaO\_C.

existence of La(OH)<sub>3</sub> in the La<sub>2</sub>O<sub>3</sub> film. With Ta incorporation, the La 3d<sub>5/2</sub> 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 La<sub>2</sub>O<sub>3</sub> film and accordingly the suppression in the formation of La(OH)<sub>3</sub> after Ta incorporation. Nevertheless, the La 3d<sub>5/2</sub> peak related to sample\_LaTaO\_C (located at 835.0 eV) shifts to an even higher binding energy in comparison to sample\_La<sub>2</sub>O<sub>3</sub>, suggesting the formation of La-O-Ta bond in the LaTaO film. The O 1s spectra of the La<sub>2</sub>O<sub>3</sub> 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 La<sub>2</sub>O<sub>3</sub> film, the two O 1s peaks correspond to La-O bond (located at 528.9 eV)<sup>15</sup> and La-OH bond (located at 531.7 eV), respectively. Moreover, the O 1s peak corresponding to La-OH bond has a much higher intensity compared with that corresponding to La-O bond, indicating that most of La atoms in the La<sub>2</sub>O<sub>3</sub> film have been transformed into La(OH)<sub>3</sub> 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)<sub>3</sub> due to the enhanced moisture resistance of the La<sub>2</sub>O<sub>3</sub> 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<sub>5/2</sub> spectrum in Fig. 1(a).

Fig. 2 shows the AFM images of the La<sub>2</sub>O<sub>3</sub> and LaTaO films with a measurement area of 1 μm × 1 μm. The La<sub>2</sub>O<sub>3</sub> 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 La<sub>2</sub>O<sub>3</sub> film after moisture absorption.<sup>11</sup> 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

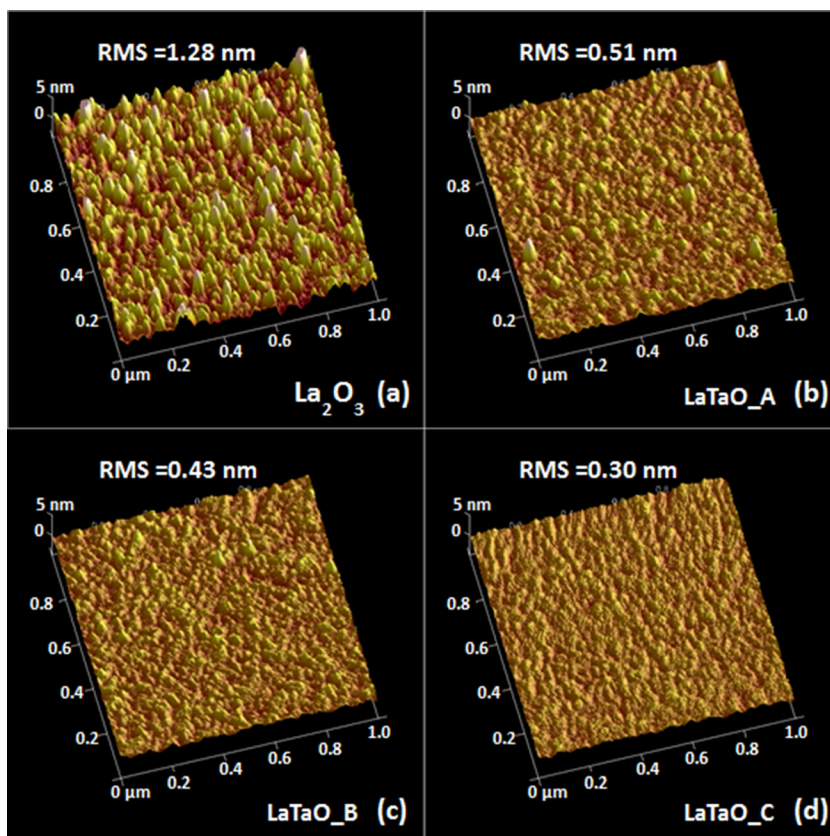


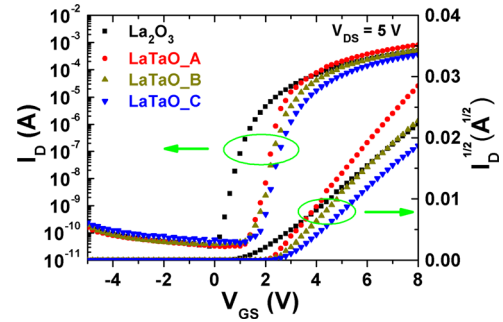
FIG. 2. AFM image of dielectric films in (a) sample\_La<sub>2</sub>O<sub>3</sub>, (b) sample\_LaTaO\_A, (c) sample\_LaTaO\_B, and (d) sample\_LaTaO\_C.

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

Sample	La <sub>2</sub> O <sub>3</sub>	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%
$\mu_{\text{sat}}$ (cm <sup>2</sup> /V·s)	12.1	23.4	16.3	11.0
$V_{\text{TH}}$ (V)	1.85	2.40	2.66	2.97
SS (V/dec)	0.234	0.177	0.201	0.217
$\Delta V_{\text{H}}$ (V)	-0.76	0.10	1.29	2.34
$I_{\text{on}}$ ( $\mu\text{A}$ )	494	810	520	349
$I_{\text{on}}/I_{\text{off}}$	$1.5 \times 10^7$	$2.6 \times 10^7$	$1.3 \times 10^7$	$8.6 \times 10^6$
$C_{\text{ox}}$ ( $\mu\text{F}/\text{cm}^2$ )	0.231	0.262	0.269	0.279
Dielectric constant	10.4	11.8	12.2	12.6

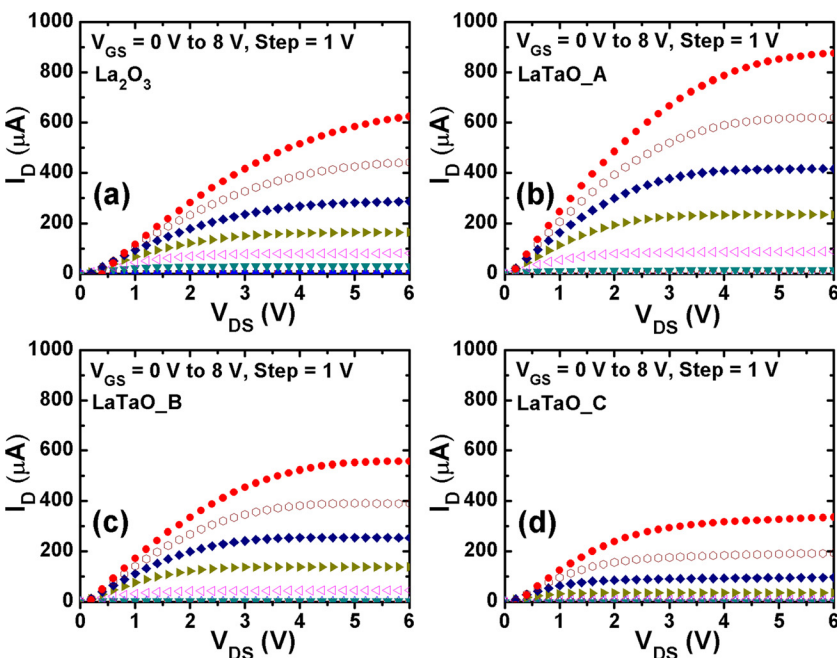
sample\_LaTaO\_C is 0.51 nm, 0.43 nm, and 0.30 nm, respectively, further demonstrating that Ta incorporation is an effective way to enhance the moisture resistance of La<sub>2</sub>O<sub>3</sub> film and accordingly reduce its surface roughness. In addition, the enhanced moisture resistance of the La<sub>2</sub>O<sub>3</sub> 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 SiO<sub>2</sub> dielectric, a larger dielectric constant of the LaTaO film ( $\sim 12$ ) 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 ( $I_{\text{D}}$ ) vs. gate-to-source voltage ( $V_{\text{GS}}$ ) and  $I_{\text{D}}^{1/2}$  vs.  $V_{\text{GS}}$  at a drain-to-source voltage ( $V_{\text{DS}}$ ) of 5 V. The saturation carrier mobility ( $\mu_{\text{sat}}$ ), threshold voltage ( $V_{\text{TH}}$ ), subthreshold swing (SS), on current ( $I_{\text{on}}$ ), 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,  $\mu_{\text{sat}}$  and  $V_{\text{TH}}$  are calculated from a linear fitting to the plot of  $I_{\text{D}}^{1/2}$  versus  $V_{\text{GS}}$ , 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\_La<sub>2</sub>O<sub>3</sub>, 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\_La<sub>2</sub>O<sub>3</sub> and sample\_LaTaO\_A,  $\mu_{\text{sat}}$  is nearly doubled from 12.1 cm<sup>2</sup>/V·s to 23.4 cm<sup>2</sup>/V·s with a reduction of SS from 0.234 V/dec to 0.177 V/dec due to the Ta incorporation in the La<sub>2</sub>O<sub>3</sub> 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.<sup>16</sup> 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  $\mu_{\text{sat}}$  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.<sup>1,17</sup> 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.<sup>18</sup> Hence, it is believed that the high trap density at/near the a-IGZO/dielectric interface in sample\_La<sub>2</sub>O<sub>3</sub> is related to the oxygen vacancies originated from the hygroscopicity of La<sub>2</sub>O<sub>3</sub>. In addition, a smaller  $V_{\text{TH}}$  (1.85 V) of sample\_La<sub>2</sub>O<sub>3</sub> in comparison to  $V_{\text{TH}}$  (2.40 V) of

FIG. 4. Output characteristics of the a-IGZO TFTs: (a) sample\_La<sub>2</sub>O<sub>3</sub>; (b) sample\_LaTaO\_A; (c) sample\_LaTaO\_B; and (d) sample\_LaTaO\_C.



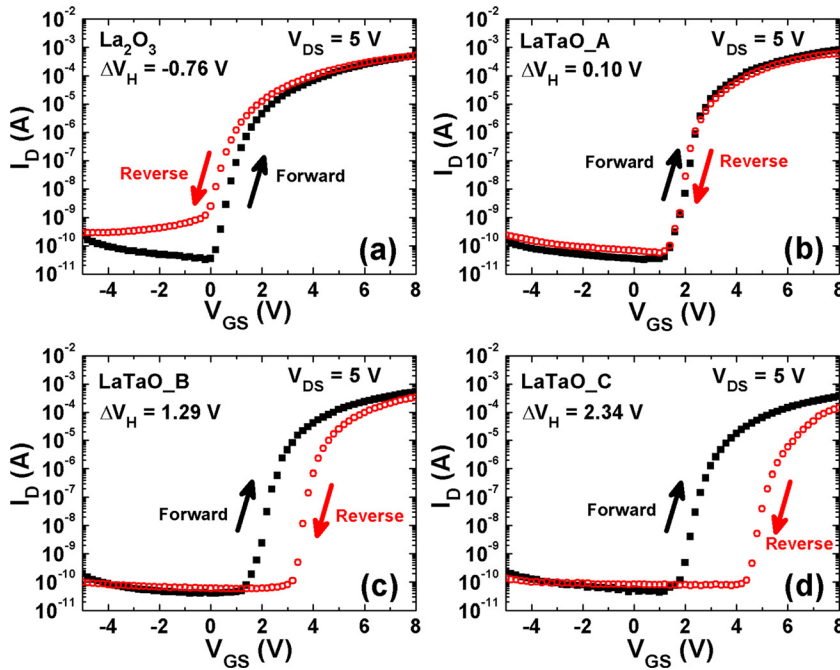


FIG. 5. Transfer characteristics of the a-IGZO TFTs measured under the forward and reverse  $V_{GS}$  sweepings: (a) sample\_La<sub>2</sub>O<sub>3</sub>; (b) sample\_LaTaO\_A; (c) sample\_LaTaO\_B; and (d) sample\_LaTaO\_C.

sample\_LaTaO\_A can further reveal the existence of oxygen vacancies in the La<sub>2</sub>O<sub>3</sub> 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.<sup>19</sup> Besides,  $I_{on}$  and  $I_{on}/I_{off}$  are increased from 494  $\mu$ A and  $1.5 \times 10^7$  to 810  $\mu$ A and  $2.6 \times 10^7$ , 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 Ta<sub>2</sub>O<sub>5</sub> film,<sup>20</sup> 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\_La<sub>2</sub>O<sub>3</sub> and sample\_LaTaO\_A, the output current of the TFT is significantly increased by the Ta incorporation in the La<sub>2</sub>O<sub>3</sub> 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.  $\Delta V_H$  is defined as the  $V_{TH}$  shift in the hysteresis loop. It is found that sample\_La<sub>2</sub>O<sub>3</sub> exhibits an obvious anticlockwise hysteresis with a negative  $\Delta V_H$  ( $-0.76$  V), further revealing the existence of donor-like traps at/near the a-IGZO/dielectric interface,<sup>21</sup> which is due to the introduction of oxygen vacancies in the La<sub>2</sub>O<sub>3</sub> film after moisture absorption. These donor-like traps can induce electron-detrapping or hole-trapping and become positively charged<sup>19</sup> 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 La<sub>2</sub>O<sub>3</sub> 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  $\Delta V_H$  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  $\Delta V_H$  for hysteresis related to donor-like and acceptor-like traps has also been observed in other work.<sup>22</sup>

In this work, the impact of Ta incorporation in La<sub>2</sub>O<sub>3</sub> 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 La<sub>2</sub>O<sub>3</sub> film and suppress the formation of La(OH)<sub>3</sub>, 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  $\mu_{sat}$ , 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.

This work was supported by the University Development Fund (Nanotechnology Research Institute, 00600009) of the University of Hong Kong.

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