Influence of the Gate/Drain Voltage Configuration on the Current Stress Instability in Amorphous Indium-Zinc-Oxide Thin-Film Transistors With Self-Aligned Top-Gate Structure

Sungju Choi[®], Shinyoung Park, Jae-Young Kim, Jihyun Rhee, Hara Kang, Dong Myong Kim[®], *Member, IEEE*, Sung-Jin Choi[®], and Dae Hwan Kim[®], *Senior Member, IEEE*

Abstract—The influence of V_{GS}/V_{DS} condition on the current stress (CS) instability in amorphous InZnO thin-film transistors (TFTs) with the self-aligned top-gate structure is comprehensively analyzed and quantitatively validated by consolidating: 1) the I-V and C-V characteristics: 2) the extraction of density of states; 3) the decomposition of threshold voltage shift (ΔV_T); and 4) the computer-aided design simulation. It has been found that in a high V_{GS} and low V_{DS} CS condition, electron trapping into the gate insulator globally occurs. However, these effects are combined with a local electron trapping into gate insulator in the source region and the generated peroxide defects in the drain region under a high V_{DS} and low V_{GS} CS condition. The peroxide formation that is followed by the donor generation is clearly distinguished by the activation energy of 0.49 eV from the oxygen vacancies ionization which has been widely modeled for explaining the donor creation in amorphous oxide semiconductor TFTs.

Index Terms—InZnO thin-film transistor, current stress (CS), density-of-state, amorphous oxide semiconductor.

I. INTRODUCTION

MORPHOUS oxide semiconductor (AOS) materials have attracted much attention due to their potential application in high-resolution large-area displays which make use of thin-film transistors (TFTs) [1]. The commercialization of the AOS TFT-driven active-matrix organic light-emitting diode (AMOLED) display has been recently accelerated, especially for devices with the self-aligned top-gate structure. This is due to the high mobility, low source/drain (S/D) resistance, and low parasitic capacitance this technology offers [2]–[5]. However, current stress instability under

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The authors are with the School of Electrical Engineering, Kookmin University, Seoul 02707, South Korea (e-mail: drlife@kookmin.ac.kr).

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practical operating conditions remains a central challenge in the mass production of AOS TFT-driven AMOLED displays. To address this problem, it is necessary to optimize the gate-to-source voltage ($V_{\rm GS}$) and the drain-to-source voltage ($V_{\rm DS}$), during the design process of high performance and high reliability of AMOLED pixels [6]. However, the quantitative analysis of the mechanism and the modeling of the influence of the $V_{\rm GS}/V_{\rm DS}$ configuration on the instabilities of AOS TFTs with self-aligned top-gate structure have not been reported in the semiconductor materials literature.

In this work, the effect of the $V_{\rm GS}/V_{\rm DS}$ configuration on the current stress (CS)-induced instability in the amorphous indium-zinc-oxide (a-IZO) TFTs with the self-aligned top-gate structure is investigated and modeled. The relevant mechanisms are figured out by considering two different $V_{\rm GS}/V_{\rm DS}$ conditions; the saturation operating condition of $V_{\rm GS}/V_{\rm DS} =$ 10 V/30 V, i.e., higher $V_{\rm DS}$ than $V_{\rm GS}$ (*HVDS*), and the linear operating condition of $V_{\rm GS}/V_{\rm DS} =$ 30 V/10 V, i.e., higher $V_{\rm GS}$ than $V_{\rm DS}$ (*HVGS*). Quantitative modeling was carried out using technology computer-aided design simulations(TCAD).

II. EXPERIMENTAL PROCEDURE

The fabrication of TFT devices with self-aligned topgate structure was conducted by depositing a 50-nm-thick a-IZO active layer on a glass substrate, using the radio frequency magnetron sputtering process. Subsequently, a 100-nm-thick SiO₂ acting as the gate insulator (GI) was deposited using plasma-enhanced chemical vapor deposition (PECVD). A 100-nm-thick layer of molybdenum (Mo) acting as the gate electrode was sequentially sputtered at room temperature. These two layers were continuously dry-etched using a gate pattern. This process was followed by the Ar plasma treatment to form the S/D n^+ regions of the active layer. Next, a 200-nm-thick a-SiO2 interlayer dielectric was deposited by PECVD. Subsequently, a 200-nm-thick layer of Mo acting as the S/D metal was sputtered at room temperature and then patterned by dry etching. Finally, the TFT devices were subjected to thermal annealing at 250 °C for 1 h. The fabricated device is schematically illustrated in the inset of Fig. 1(b). The channel width (W) and length (L) are 40 μ m and 40 μ m.

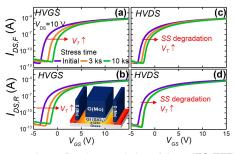


Fig. 1. Measured transfer characteristics of the a-IZO TFTs under (a) forward and (b) reverse V_{DS} modes in *HVGS* and under (c) forward and (d) reverse V_{DS} modes in *HVDS* condition. The inset in (b) is a schematic illustration of the fabricated a-IZO TFTs structure.

III. RESULTS AND DISCUSSION

Figs. 1(a)-(d) show the measured transfer curves of a-IZO TFTs at $V_{DS} = 10$ V during the CS time (t_{str}) up to 10^4 s under the *HVGS* and *HVDS* conditions. In the curves, $I_{DS,F}$ is the drain-to-source current, measured under forward V_{DS} mode (the drain and source positions are identical to the CS positions) in read-out condition, while $I_{DS,R}$ is the drain-to-source current, measured under reverse V_{DS} mode (the drain and source positions are interchanged with respect to the CS positions). The threshold voltage (V_T) was defined as the value of V_{GS} that induces $I_{DS} = 10$ nA at $V_{DS} = 10$ V and the sub-threshold region.

Under CS, the positive V_T shift (ΔV_T) is observed in all cases shown in Fig. 1. It is evident from Figs. 1(a) and (b), that in the case of *HVGS*, the ΔV_T in forward V_{DS} mode $(\Delta V_{T,F})$ is analogous to that in the reverse V_{DS} mode $(\Delta V_{T,R})$ as t_{str} increases, while the *SS* is nearly unchanged. However, in the case of *HVDS*, $\Delta V_{T,F}$ and $\Delta V_{T,R}$ behave differently and a significant degradation of the *SS* is observed [Figs. 1(c)-(d)]. This result implies that the local ΔV_T values near source and drain are evolved differently during CS and the traps are generated near the conduction band minimum level (E_C), especially in the *HVDS* condition.

For more detailed analysis, the small signal C-V(capacitance-voltage) characteristics between the gate and S/D tied to ground, i.e., $C_{G-DS}-V_G$, are measured as shown in Figs. 2(a) and (b). Under the HVDS [Fig. 2(b)], the C-Vslope decreases with a positive $\Delta V_{\rm T}$, while the turn-on voltage of C_{G-DS} decreases during CS compared to the initial state [reference point A in Figs. 2(c) and (d)]. This suggests that a negative $\Delta V_{\rm T}$ mechanism occurs locally in the drain region which is denoted by the blue box in Fig 2(d) [7], [8]. In general, the origin of negative $\Delta V_{\rm T}$ may be due to either hole trapping into GI or donor generation. In our case, a local negative $\Delta V_{\rm T}$ during CS is attributed to donor generation since the holes have short life-time and low mobility in a-IZO TFTs [9]. Thus, as indicated in Fig. 2(d), the CS-induced positive/negative $\Delta V_{\rm T}$ near S/D, individually results from the electron trapping into GI (symbolized by the trapping length L_{trap}) / the donor generation (denoted by the length of donor generation region L_{DG}), respectively. In Figs. 2(c) and (d), point B shows that the source region with L_{trap} is turned on at $V_{\text{G}} = V_{\text{B}}$ while the channel region becomes totally conducting at $V_{\rm G} = V_{\rm C}$.

The direct evidence of the CS instability was experimentally verified, by extracting the subgap density-of-state (DOS) (shown in Fig. 3) using the multi-frequency method [10].

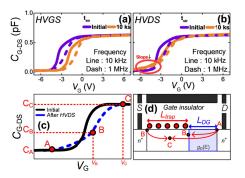


Fig. 2. (a) t_{str} -dependent C_{GDS} - V_G during (a) the *HVGS* and (b) the *HVDS*. Scheme of (c) the *C*-*V* curve under *HVDS* and (d) the spatial evolution of the TFT conducting channel during the *HVDS* depending on the V_G -dependent C_{G-DS} characteristic.

The energy distribution of the AOS *DOS* is described by the following equation models [6], [10]:

$$g_A(E) = g_{TA}(E) + g_{DA}(E) = N_{TA} \times \exp\left(\frac{E - E_C}{kT_{TA}}\right) + N_{DA} \times \exp\left(\frac{E - E_C}{kT_{DA}}\right)$$
(1)
$$g_D(E) = g_{TD}(E) + g_{SD}(E) = N_{TD} \times \exp\left(\frac{E_V - E}{kT_{DA}}\right)$$

$$b(E) = g_{TD}(E) + g_{SD}(E) = N_{TD} \times \exp\left(\frac{kT_{TD}}{kT_{TD}}\right) + N_{SD} \times \exp\left(-\left(\frac{(E_V - E) + E_{SD}}{kT_{SD}}\right)^2\right)$$
(2)

where the donor-like tail states (g_{TD}) [not shown in Fig. 3], shallow donor states (g_{SD}) , acceptor-like tail (g_{TA}) and deep states (g_{DA}) are individually modeled with the parameters of densities/characteristic energies and the Gaussian peak center energy of N_{TD}/kT_{TD} , N_{SD}/kT_{SD} , N_{TA}/kT_{TA} , N_{DA}/kT_{DA} , and E_{SD} . In the HVGS condition [Fig. 3(a)], the DOS remains unchanged during CS compared to that in the initial state, i.e., $N_{TD} = 1 \times 10^{20} \text{ cm}^{-3} \text{eV}^{-1}$, $kT_{TD} = 0.15 \text{ eV}$, $N_{TA} = 1 \times 10^{18} \text{ cm}^{-3} \text{eV}^{-1}$, $kT_{TA} = 0.04 \text{ eV}$, $N_{DA} = 2.5 \times 10^{17} \text{ cm}^{-3} \text{eV}^{-1}$, $kT_{DA} = 0.25 \text{ eV}$. The change in g_{SD} is negligible (the extracted values of parameters are within the range of those reported in the literature [9], [11], [12]), while $g_{SD}(E)$ noticeably increases to be $N_{SD} = 1.3 \times$ $10^{17} \text{ cm}^{-3} \text{eV}^{-1}$, $E_{SD} = E_{C} - 0.3 \text{ eV}$, and $kT_{SD} = 0.5 \text{ eV}$ after $t_{\rm str} = 10^4$ s only in the *HVDS* condition [Fig. 3(b)]. This result is consistent with the HVDS CS-induced donor generation mentioned above and presented in Figs. 2(c) and (d). The increase of $g_{SD}(E)$ may be associated with excessive oxygen [13], [14], oxygen interstitials [15], ionized oxygen vacancy defects [16], [17], and intrinsic (In*-M) electrontrapped centers [18]. Among them, the increased Gaussian distribution of DOS near E_C is usually related to the ionization of oxygen vacancies $(V_o + 2h^+ \rightarrow V_o^{2+})$ or the formation of peroxide defects $(O^{2-} + O^{2-} \rightarrow O_2^{2-} + 2e^-)$ [13], [19], [20]. These two phenomena can occur locally near the drain region by self-heating [17], [21], the high electric field [22], and the negative bias stress caused by V_{GD} near the drain side [23].

An experimental decomposition technique [24], can be used for a more quantitative analysis on the influence of the $V_{\rm GS}/V_{\rm DS}$ configuration, the decomposition of a total $\Delta V_{\rm T}$ ($\Delta V_{\rm T,tot}$) into the $\Delta V_{\rm T}$ due to the electron trapping into GI ($\Delta V_{\rm T,GI}$) [illustrated as $L_{\rm trap}$ in Fig. 2(d)] and the $\Delta V_{\rm T}$ resulting from the g_{SD} generation ($\Delta V_{\rm T,SD}$) [denoted by $L_{\rm DG}$ in Fig. 2(d)]. The symbols in Fig. 4, indicate that the only

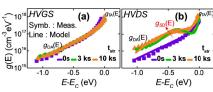


Fig. 3. t_{str}-evolution of DOS under (a) the HVGS and (b) the HVDS.

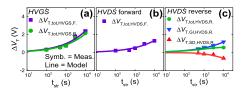


Fig. 4. Decomposed $\Delta V_{T}(t_{str})$ under (a) the *HVGS*, (b) forward and (c) reverse mode of the *HVDS* (Symbol: measurement and line: MSEF model).

Тне М	ISEF Parai	TABL METERS F		CS CONDITION
Parameters	HVGS,F/R	HVDS,F	HVDS,R	HVDS,R
$\Delta V_{\rm T}[{ m V}]$		Positive		Negative
$\Delta V_{\rm T0}$ [V]	30	10	10	-0.7

ß		
2	0.4 - 0.5	0.95
τ [s]	$3 \times 10^{6} - 6 \times 10^{6}$	5×10^3
$E_{\rm a}[{\rm eV}]$	0.83 - 0.85	0.49
Mechanism	Charge-trapping into GI	Peroxide defect

and $\Delta V_{T,SD,HVDS,R}$ should be decomposed into $\Delta V_{T,GI,HVDS,R}$ and $\Delta V_{T,SD,HVDS,R}$, whereas the only $\Delta V_{T,GI}$ contributes to $\Delta V_{T,tot}$ in all cases of $\Delta V_{T,tot,HVGS,F}$, $\Delta V_{T,tot,HVGS,R}$, and $\Delta V_{T,tot,HVDS,F}$. Here, the complicated subscripts identify the $\Delta V_{T,tot}/\Delta V_{T,tot}/\Delta V_{T,tot}$, the CS type of HVGS/HVDS, and $\Delta V_{T,F}/\Delta V_{T,R}$, respectively. Consistently with the donor generation locally near the drain, it is noteworthy that the $\Delta V_{T,SD}$ is observed only in $\Delta V_{T,tot,HVDS,R}$.

The mechanism on the donor generation under the *HVDS* CS can be elucidated, by fitting the experimentally decomposed $\Delta V_{T,tot}(t_{str})$ with the multiple stretched-exponential functions (MSEFs) [lines in Fig. 4(c)], as follows [26]:

$$\Delta V_{\mathrm{T,tot}}(t) = \Delta V_{\mathrm{T0,SD}} \left[1 - \exp\left(-\left(\frac{t}{\tau_{\mathrm{SD}}}\right)^{\beta_{\mathrm{SD}}}\right) \right] + \Delta V_{\mathrm{T0,GI}} \left[1 - \exp\left(-\left(\frac{t}{\tau_{\mathrm{GI}}}\right)^{\beta_{\mathrm{GI}}}\right) \right]$$
(3)

where ΔV_{T0} is ΔV_T at infinite time, τ is the characteristic time constant, and β is the stretching exponent. Then, $\Delta V_{T,SD}$ is the negative component of ΔV_T caused by the creation of shallow donor states, while $\Delta V_{T,GI}$ is the positive component of ΔV_T caused by the charge-trapping in the GI (because the $\Delta V_{T0,GI}$ has a positive value). The lines in Fig. 4 and Table I show that all of $\Delta V_{T,tot}$, $\Delta V_{T,GI}$, and $\Delta V_{T,SD}$ are well fitted with the MSEFs using individually selected parameters.

Furthermore, the effective energy barrier, i.e., the activation energy (E_a) corresponding to the respective physical mechanism, can be extracted by using the Laplace transform method [27] (data not shown). In Table I, the values of E_a are found to be 0.83, 0.83, 0.85, 0.85, and 0.49 eV corresponding to $\Delta V_{T,tot,HVGS,F}$, $\Delta V_{T,tot,HVGS,F}$, $\Delta V_{T,tot,HVGS,F}$, $\Delta V_{T,GI,HVGS,R}$, and $\Delta V_{T,SD,HVGS,R}$, respectively. The values of $E_{a,GI}$'s = 0.83 – 0.85 eV are in good agreement with

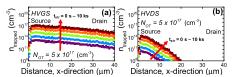


Fig. 5. Simulated n_{trapped} during (a) the HVGS and (b) the HVDS.

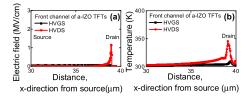


Fig. 6. Simulated distribution of (a) electric field and (b) temperature under CS.

the E_a values reported in studies which have considered the charge-trapping in the GI as the dominant mechanism on ΔV_T . These observed E_a values ranged from 0.60 to 1.33 eV [25], [27], [28]. On the other hand, it is noteworthy that the value of $E_{a,SD} = 0.49$ eV extracted at negative ΔV_T under the *HVDS* condition coincides with the E_a value associated with the peroxide defect formation through electron de-trapping $(O^{2-}+O^{2-} \rightarrow O_2^{2-}+2e^-)$. This E_a value has been calculated to be 0.46 eV [29]. The extracted $E_{a,SD} = 0.49$ eV does not match the E_a value for the reaction of ionized oxygen vacancies $(V_o + 2h^+ \rightarrow V_o^{2+}, E_a = 0.88 - 0.97 \text{ eV})$ [30]. Therefore, the donor generation under the *HVDS* CS condition is attributed to the formation of peroxide defects rather than the ionization of oxygen vacancies.

The Atlas-2D TCAD software [31] was used to simulate the local charge-trapping model and the change in the *DOS*, in order to thoroughly verify this physical behavior. The simulations were incorporated into by physical models, a transient charge-trapping model, the GI trap model, F-N tunneling, band-to-band tunneling, trap-assisted tunneling, and hot-carrier injection. The spatial distribution of the density of trapped electrons (n_{trapped}) in the GI over CS time at a fixed density of oxide bulk trap ($N_{\text{OT}} = 5 \times 10^{17} \text{ cm}^{-3}$) is shown in Figs. 5(a) and (b). In the *HVGS* case, global charge-trapping occurs throughout the GI as t_{str} increases. However, in the *HVDS* case, n_{trapped} increases locally only in the drain region as t_{str} increases.

Finally, the higher electric field and the temperature of a-IZO TFTs were observed in the drain side, under the *HVDS* condition as shown in Figs. 6(a) and (b). These simulation results verify that the formation of peroxide defects becomes more activated due to a locally higher electric field and temperature in the drain side as $V_{\rm DS}$ becomes higher than $V_{\rm GS}$, i.e., in the *HVDS* condition.

IV. CONCLUSION

The effect of V_{GS}/V_{DS} configuration on the CS instability in a-IZO TFTs with self-aligned top-gate structure is controlled by the combination of the GI charge trapping and the local formation of peroxide defects. The latter becomes more boosted as the V_{DS} becomes higher than V_{GS} , i.e., *HVDS* rather than *HVGS* due to locally higher electric field and self-heating near the drain edge. Our results provide a guide-line to optimize the bias point of current-driving TFTs with the self-aligned top-gate structure in the AOS-driven AMOLED backplanes.

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