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Investigation of Nitrous Oxide Nitridation Temperatures on P-Type Pi-Gate Poly-Si Junctionless Accumulation Mode TFTs

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ABSTRACT In this paper, the influence of nitrous oxide (N_2O) nitridation temperatures on p-type Pi-gate (PG) poly-Si junctionless accumulation mode (JAM) TFTs is experimentally investigated. The tetraethoxysilane (TEOS) gate oxide quality for PG JAM TFTs can be significantly improved by increasing N_2O nitridation temperatures (T_N) from 700 °C to 800 °C in N_2O ambient, resulting in the improvement of average subthreshold swing (A.S.S.), increase of on current (I_{ON}), and enhancement of TEOS gate oxide breakdown E -field (E_{OBD}). PG JAM TFTs by means of a proper channel doping concentration ($N_{ch} = 5 \times 10^{18} \text{ cm}^{-3}$) and a suitable T_N (800 °C) exhibit a steep A.S.S. $\sim 96 \text{ mV/dec.}$ and a large $E_{OBD} \sim 12.1 \text{ MV/cm.}$

INDEX TERMS 3-D integrated circuits (ICs), reverse boron penetration, nitrous oxide (N_2O), Pi-gate (PG), poly-Si, junctionless accumulation mode (JAM).

I. INTRODUCTION

Polycrystalline silicon thin-film transistor (poly-Si TFT) is one kind of field-effect transistors. Poly-Si TFTs have been widely investigated and used in large-area flat panel displays with integrated peripheral circuits as the pixel array, such as active-matrix liquid crystal displays (AMLCDs) and active-matrix organic light-emitting displays (AMOLEDs) [1]–[4]. Moreover, poly-Si TFTs are very suitable for three-dimensional integrated circuit (3-D IC) applications and 3-D NAND flash memory applications owing to those advantages associated with low cost, low thermal budget, simple fabrication, high sequential stacked ability, and CMOS-compatible process [5]–[7].

Before the 0.25 μm technology node, both n-MOSFETs and p-MOSFETs with the same *in situ* n^+ doped poly-Si gates require p-type channel doping regions to acquire a suitable threshold voltage (V_{TH}) for very large scale integration (VLSI) applications [8]. Based on the use of n^+ poly-Si gates, p-MOSFETs are more susceptible to short-channel effects (SCEs) due to the use of p-type channel doping regions for V_{TH} adjustment [8]–[10]. Such

devices are designated as buried-channel MOSFETs (BC-MOSFETs). To enhance SCE immunity as well as attain low and symmetric V_{TH} , n^+ poly-Si gates and p-type channel doping regions for n-MOSFETs as well as p^+ poly-Si gates and n-type channel doping regions for p-MOSFETs in the 0.25 μm technology node are utilized to form surface-channel MOSFETs (SC-MOSFETs) in submicron CMOS devices [11]–[13]. However, boron atoms used to dope the undoped poly-Si gates can rapidly diffuse through the thin gate oxide from the p^+ poly-Si gate into the underlying n-type channel region during a high thermal-budget process, namely, boron penetration. This causes a positive shift in flat-band voltage (V_{FB}), a positive shift in V_{TH} , a degraded subthreshold swing (S.S.), and a degraded Si/SiO₂ interface [11]–[18].

In literature, numerous approaches have been proposed and demonstrated to relieve boron penetration. There are two common methods. The first method is to establish diffusion barrier layers at interfaces of poly-Si gates/gate oxides and/or of gate oxides/n-type channel doping regions. This can be accomplished by nitrogen (N) ion implantations

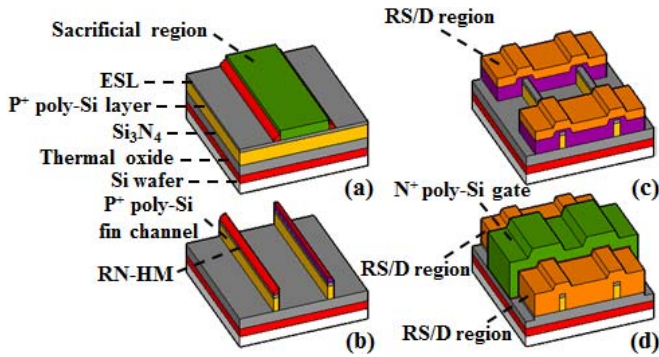


FIGURE 1. (a)–(d) Schematic 3-D configurations of a PG JAM TFT for the key fabrication steps.

into poly-Si gates and nitridations for poly-Si gates and/or gate oxides by using nitrous oxide (N₂O), nitric oxide (NO), or ammonia (NH₃). Alternatively, inductive-coupling-nitrogen plasma (ICNP) treatments can be used to incorporate nitrogen-rich layers at interfaces of poly-Si gates/gate oxides and/or of gate oxides/n-type channel doping regions [18]–[23]. The second method is to retard vertical diffusion of boron atoms in gate stack structures. This can be realized by using as-deposited α -Si gates instead of poly-Si gates and stacked (or modified) gate structures with α -Si and poly-Si layers [16], [24]–[26].

Current p-type junctionless (JL) and junctionless accumulation mode (JAM) transistors need ultra-shallow channel doping regions by using boron difluoride (BF₂) ion implantations in order to effectively deplete the conducted carriers in the OFF-state. However, boron diffusion can further aggravate as a result of the presence of fluorine (F) atoms from BF₂ ion implantations with F atoms [11]–[13]. This indicates that reverse boron penetration will become more severe in p-type JL and JAM transistors. As a result, in this study, we will explore the influence of channel doping concentrations (N_{ch}) and N₂O nitridation temperatures (T_N) on p-type PG JAM TFTs to further evaluate the optimum N_{ch} and T_N . Furthermore, we will clarify the major impact factors that affect the threshold voltage, the average subthreshold swing (A.S.S.), the on current (I_{ON}), the S/D series resistance (R_{SD}), and the tetraethoxysilane (TEOS) gate oxide breakdown E -field (E_{OBD}).

II. DEVICE FABRICATION

The schematic 3-D configurations of a PG JAM TFT for the key fabrication steps are shown in Fig. 1(a)–(d). The detailed fabrication steps for the PG JAM TFTs were reported in our previous work [27]. First of all, the 150-nm-thick silicon nitride (Si₃N₄) layer was deposited on the thermal oxide with a film thickness of 500 nm grown on a Si wafer. Subsequently, the 40-nm-thick undoped amorphous-Si (a-Si) layer was deposited and then crystallized during the solid-phase crystallization (SPC) process. Next, the undoped poly-Si layer was implanted with 18-keV boron difluoride ions (BF₂⁺) to a dose of 2×10^{13} cm⁻². After the dopant

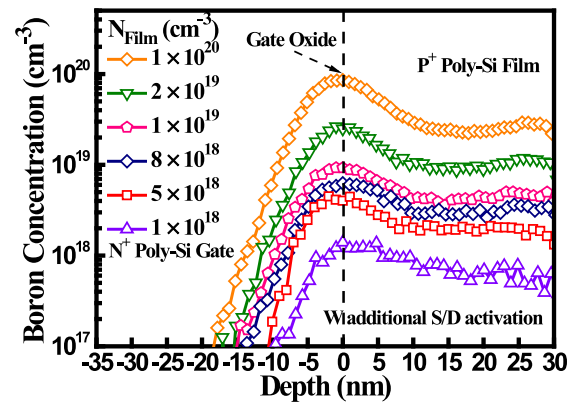


FIGURE 2. SIMS depth profiles of the un-patterned p⁺ poly-Si films with different film doping concentrations after the gate stack formation.

activation, an etching stopping layer (ESL) with a film thickness of 11 nm and a sacrificial layer with a film thickness of 50 nm were sequentially deposited. Following the patterning of the sacrificial regions, the TEOS oxide with a film thickness of 50 nm was deposited and then etched to form the oxide-spacer hard masks (O-SPHMs) [Fig. 1(a)]. After the removal of the sacrificial regions, the O-SPHMs as the first hard masks were utilized to define the rectangular silicon nitride-hard masks (RN-HMs). A 100-nm-thick undoped a-Si layer was deposited and then implanted with the boron ions (B⁺) to a dose of 2×10^{15} cm⁻² after the p⁺ poly-Si fin channel formation [Fig. 1(b)]. Using the RN-HMs as the second hard masks, the twin p⁺ poly-Si fin channels and the raised source/drain (RS/D) regions were simultaneously patterned by means of an anisotropic dry etching process [Fig. 1(c)]. After the RN-HM strip, the gate stack formation consists of the TEOS oxide with a film thickness of 5 nm and the n⁺ poly-Si gate with a film thickness of 200 nm. During the gate stack formations, the TEOS gate oxides were nitridized at different N₂O nitridation temperatures ($T_N = 700$ °C, 750 °C, 800 °C, and 850 °C) in N₂O ambient. The gate formation was accomplished to complete the p-type PG JAM TFTs [Fig. 1(d)]. The channel thickness (T_{ch}) \times the channel width (W_{ch}) for the p⁺ poly-Si fin channel is 36.0 nm \times 26.7 nm for p-type PG JAM TFTs [27].

III. RESULTS AND DISCUSSION

Fig. 2 illustrates the secondary ion mass spectroscopy (SIMS) depth profiles of the un-patterned p⁺ poly-Si films with different film doping concentrations after the gate stack formation. The film doping concentration (N_{film}) values for the un-patterned p⁺ poly-Si films are, respectively, 1×10^{20} cm⁻³, 2×10^{19} cm⁻³, 1×10^{19} cm⁻³, 8×10^{18} cm⁻³, 5×10^{18} cm⁻³, and 1×10^{18} cm⁻³. It is noteworthy that more boron atoms can penetrate through the TEOS gate oxide from the p⁺ poly-Si film toward the *in situ* n⁺ doped poly-Si gate with an increase in N_{film} , namely, channel doping concentration-induced reverse boron penetration. This implies that more

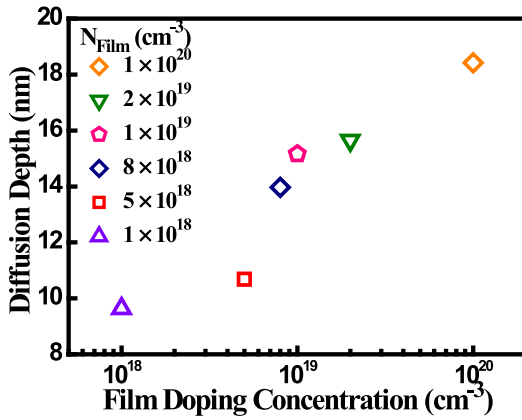


FIGURE 3. Diffusion depth of boron atoms versus N_{film} in the *in situ* n⁺ doped poly-Si gate.

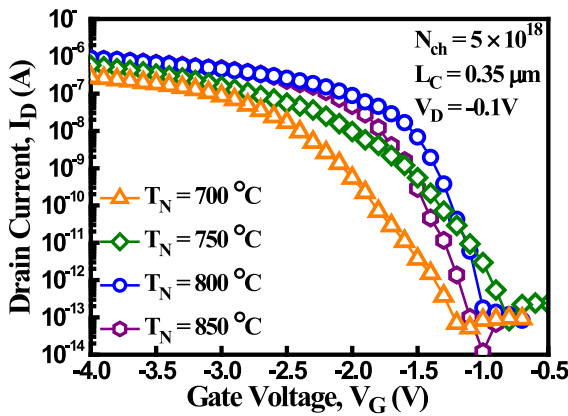


FIGURE 4. Comparison of the transfer characteristics of PG JAM TFTs with different T_N values.

boron atoms can stay at the TEOS gate oxide with the increase of N_{film} . In addition, the diffusion depth of boron atoms in the *in situ* n⁺ doped poly-Si gate becomes long with an increase in N_{film} , as shown in Fig. 3. According to the above-mentioned SIMS depth profiles, the PG JAM TFTs with a higher N_{ch} will undergo more severe channel doping concentration-induced reverse boron penetration. Based on an evaluation between device performance and channel doping concentration-induced reverse boron penetration, an N_{ch} of $5 \times 10^{18} \text{ cm}^{-3}$ is suitable for PG JAM TFTs [27].

Fig. 4 displays the transfer characteristics of PG JAM TFTs with different T_N values. V_{TH} values are defined by a constant current of $5 \times 10^{-9} \text{ A}$. With increasing T_N from 700 °C to 800 °C in N₂O ambient, the positive shift in V_{TH} for PG JAM TFTs might be largely attributed to two causes. One is the enhancement of gate oxide interface quality since the deposition temperature for the TEOS gate oxides was 700 °C. The other is the effective suppression of reverse boron penetration by N₂O treatments [20], [22]. In PG JAM TFTs, the negative shift in V_{TH} could result from the further generation of positive fixed oxide charge with an increase in T_N from 800 °C to 850 °C in N₂O ambient [28]–[33].

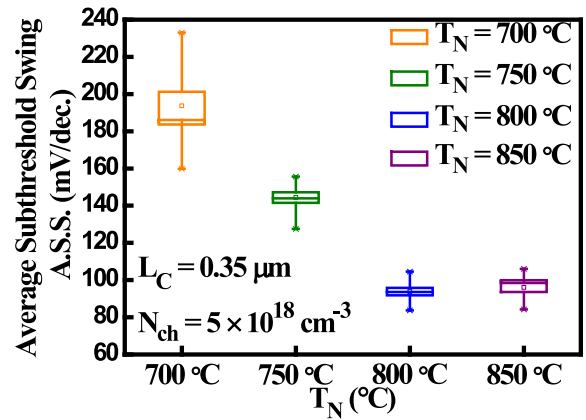


FIGURE 5. Statistical distributions of A.S.S. for the PG JAM TFTs with different T_N values.

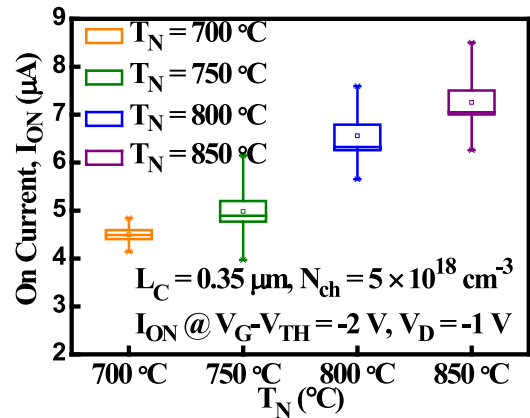


FIGURE 6. Statistical distributions of I_{ON} for the PG JAM TFTs with different T_N values.

Fig. 5 shows the statistical distributions of A.S.S. for PG JAM TFTs with different T_N values. Degradation of A.S.S. for PG JAM TFTs can be attributed to the reduction of gate oxide interface quality and the aggravation of reverse boron penetration [11]–[13], [27]. With increasing T_N from 700 °C to 800 °C in N₂O ambient, the obvious improvement in A.S.S. suggests that the enhancement of gate oxide interface quality is the major impact factor, which is consistent with the above-mentioned analyses and explanations for V_{TH} . Furthermore, PG JAM TFTs have a similar A.S.S. as the TEOS gate oxides are nitridized at a T_N of 800 °C and a T_N of 850 °C in N₂O ambient. This indicates that both PG JAM TFTs have a similar interface state density.

Fig. 6 exhibits the statistical distributions of I_{ON} for PG JAM TFTs with different T_N values. PG JAM TFTs utilizing a T_N of 850 °C in N₂O ambient show the highest I_{ON} . There are two causes that are associated with the increase of I_{ON} for PG JAM TFTs. One is the decrease in coulomb scattering due to the enhancement of gate oxide interface quality. This can be proved through the obvious improvement for A.S.S., as presented in Fig. 5. The other is the decrease in R_{SD} with an increase in T_N . So as to further understand and analyze

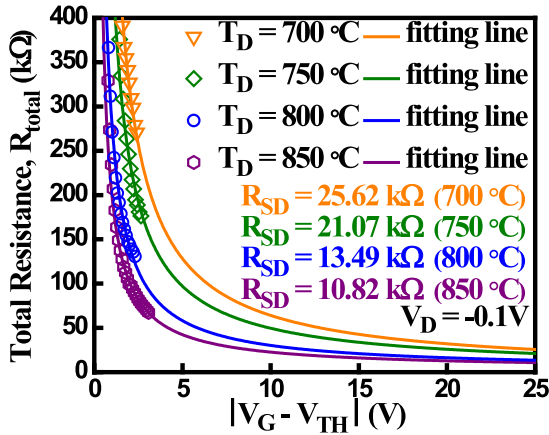


FIGURE 7. Extracted R_{total} as a function of V_G for the PG JAM TFTs with different T_N values.

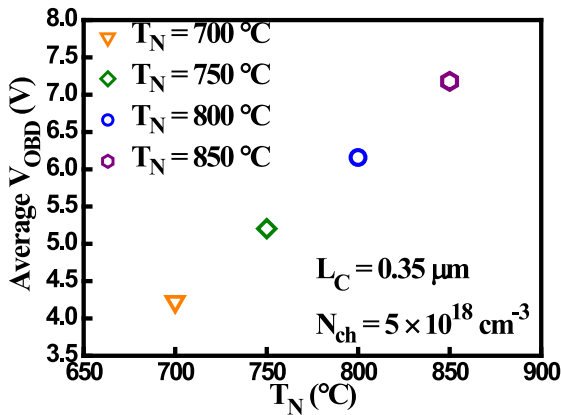


FIGURE 8. Extracted average V_{OBD} versus T_N for the PG JAM TFTs with different T_N values.

the impact of R_{SD} on I_{ON} for PG JAM TFTs with different T_N values, R_{SD} is derived from $R_{\text{total}} = R_{\text{SD}} + (L_G)/[W_{\text{eff}} \times \mu_{\text{eff}} \times C_{\text{ox}} \times (V_G - V_{\text{TH}})]$ at $V_{\text{DS}} = -0.1$ V [34], where R_{total} is the total resistance, W_{eff} is the effective channel width, μ_{eff} is the effective mobility, and C_{ox} is the gate oxide capacitance. The second term on the right-hand side of the equation can be ignored as $V_G - V_{\text{TH}}$ is close to infinity, i.e., R_{total} is equal to R_{SD} . Fig. 7 compares the extracted R_{total} as a function of V_G for PG JAM TFTs with different T_N values. The extrapolated R_{SD} for PG JAM TFTs are, respectively, 25.62 k Ω , 21.07 k Ω , 13.49 k Ω , and 10.82 k Ω at $V_G - V_{\text{TH}} = -25$ V with the increase of T_N from 700 °C to 850 °C in N₂O ambient. Using a T_N of 850 °C in N₂O ambient, PG JAM TFTs exhibit the lowest extrapolated R_{SD} , which is consistent with the previous result shown in Fig. 6.

Fig. 8 displays the extracted average TEOS gate oxide breakdown voltage (V_{OBD}) versus T_N for PG JAM TFTs with different T_N values. It is worth noting that the extracted average V_{OBD} increase with an increase in T_N . To further check the TEOS gate oxide quality for the TEOS gate oxides with different T_N values, E_{OBD} is extracted by $(V_{\text{OBD}} - V_{\text{TH}})/T_{\text{ox}}$. Here, T_{ox} is the TEOS gate oxide thickness. The

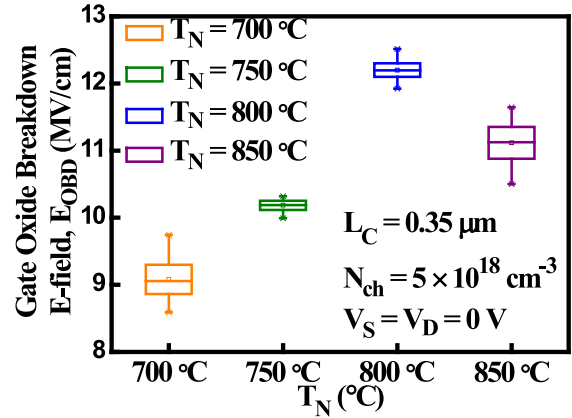


FIGURE 9. Statistical distributions of E_{OBD} for the PG JAM TFTs with different T_N values.

statistical distributions of E_{OBD} for PG JAM TFTs with different T_N values are presented in Fig. 9. Utilizing a T_N of 800 °C in N₂O ambient, PG JAM TFTs can achieve the largest E_{OBD} owing to these devices with the best TEOS gate oxide quality [17], [18]. This is consistent with the previous result shown in Fig. 8.

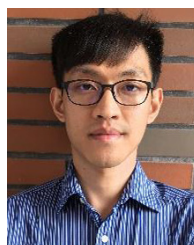
IV. CONCLUSION

In this paper, the influence of N₂O nitridation temperatures on the electrical characteristics of p-type PG JAM TFTs was experimentally investigated by N₂O treatments. For the first time, it was found that more boron atoms can penetrate through the gate oxide from the p⁺ fin channel toward the n⁺ poly-Si gate with an increase in N_{ch} . By increasing T_N from 700 °C to 800 °C in N₂O ambient, the TEOS gate oxide quality can be obviously improved, leading to an improvement of A.S.S., an increase of I_{ON} , and an enhancement of E_{OBD} . By means of a proper N_{ch} (5×10^{18} cm⁻³) and a suitable T_N (800 °C), PG JAM TFTs can exhibit the steepest A.S.S. and the largest E_{OBD} .

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KUN-CHENG LIN, photograph and biography not available at the time of publication.



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