Role of Boron in Amorphous-InGaZnO Film for Resistance Control Technique

<u>Keisuke Yasuta</u>¹, Toshimasa Ui¹, Tomokazu Nagao¹, Daisuke Matsuo², Toshihiko Sakai², Yoshitaka Setoguchi², Eiji Takahashi², Yasunori Andoh² and Junichi Tatemichi¹

E-mail: ui_toshimasa@nissin.co.jp ¹ NISSIN ION EQUIPMENT CO., LTD. 29, Hinokigaoka, Minakuchi-cho, Koka, Shiga, 528-0068, Japan ² NISSIN ELECTRIC CO., LTD. 47, Umezu-Takase-cho, Ukyo-ku, Kyoto, 615-8686, Japan Keywords: Boron, Implantation, IGZO, Resistance

ABSTRACT

As a next-generation electronics material, amorphous indium-gallium-zinc oxide (a-IGZO) film devices were investigated. In order to further utilize a-IGZO films, we carried out boron ion implantation and analyzed the implanted a-IGZO using Hall measurements with wet etching processes. As a result, we clarified that the implanted boron decreases a-IGZO resistance, which can be applied to drastically sheet resistance reduction methods for a-IGZO device processing.

1 Introduction

Amorphous indium-gallium-zinc oxide (a-IGZO) attracts much attention for next-generation flat-panel displays due to the large band gap, light transparency, printability and low-temperature in film deposition. Using a-IGZO films, thin-film-transistors (TFTs) are realized, which show superior behavior compared with amorphous Si-based TFT [1, 2]. In order to further improve a-IGZO TFT performance via source-drain resistance reduction, plasma treatment [3], excimer laser annealing [4] and ion implantation [5, 6] methods as the sheet resistance reduction techniques were investigated. Among them, the ion implantation methods have depth control ability through insulator layers and microfabrication ability owing to the small channel length reduction. In our previous work we elucidated that a-IGZO sheet resistance reduction can be attributed to oxygen vacancy (Vo) generated by noble gas implantation [7]. However, there is still much room for clarifying and improving the implantation method, since many injectable ion species in a-IGZO are not investigated in detail. In this work, we carried out one of the conventional ion B⁺ implantation and investigation in a-IGZO thin film on glass. Electron transport properties of B⁺ implanted a-IGZO films as a function of the a-IGZO depth were investigated by Hall measurements combined with wet etchings. In addition, Vo concentration profile in the a-IGZO is calculated by a simulation. On the other hand, bonding analysis by X-ray photoelectron spectroscopy (XPS) were carried out. As results, we clarified roles of boron in a-IGZO film for resistance control technique.

2 Experiments and results

We prepared a-IGZO films with a film thickness 50 nm and a sheet resistance ~ 10^{13} Ω /sq. deposited by inductively coupled plasma (ICP) sputtering on a glass (0.5-mm-thick EAGLE XG), in which conditions are gas flow Ar/O₂ = 95/5 sccm, pressure 0.9 Pa, RF power 7 kW and InGaZnO₄ target voltage -400 V [8]. In order to decrease the sheet resistance and fabricate implantation



Fig. 1 (a) B depth profiles in a-IGZO (50 nm)/glass structure of each ion energy E_{ion} measured by secondary mass spectrometry (SIMS) and calculated by a simulator TRIM [10]. (b) electronic and nuclear stopping power as a function of B⁺ energy in the a-IGZO obtained by the SIMS and the TRIM shown in Fig.1 (a).



Fig. 2 Sheet resistances as functions of E_{ion} of 50-nmthick a-IGZO films on glass with B⁺ implantation (B⁺ imp.) and with annealing process in N₂ ambient of 250 °C after B⁺ implantation (B⁺ imp. \rightarrow N₂ 250 °C anl) or and with annealing process in air of 250 °C after B⁺ implantation (B⁺ imp. \rightarrow Air 250 °C anl), respectively, where dose amounts are 1×10¹⁵ cm⁻².

depth control technique, B⁺ ion implantations with the ion energy range of 15-30 keV in the a-IGZO/glass structure were carried out using ion implanter iG series [9] and the implantation depth profiles were measured by secondary mass spectrometry (SIMS) as shown in Fig. 1 (a). From B depth profile in the a-IGZO, we estimated electronic and nuclear stopping power as a function of ion energy Eion of B⁺ in the a-IGZO calculated by a simulator (Transport or lons in Matter TRIM) [10] as shown in Fig.1 (b). Hence, we fabricated B⁺ implantation depth control technique. On the other hand, we carried out B⁺ implantation in the a-IGZO films for the purpose of implantation ion energy optimazation, where the Eion ranges are 5-80 keV with uniform dose 1×10^{15} cm⁻², respectively . In addition, post-annealing (anl) in N2 ambient or the air at 250 °C 1 hour were carried out. From Hall measurement, we obtained the a-IGZO sheet resistance as a function of the Eion as shown in Fig. 2. Although B⁺ implanted a-IGZO sheet resistance decreases with increasing Eion of 10-20 keV, a-IGZO sheet resistance increases with increasing in the range of 20-80 keV, which behaviors almost corresponds to B amounts in a-IGZO films as shown in Fig. 1 (a). In addition, we find that the increase or decrease behaviors are emphasized after the annealing processes. In addition, the annealed a-IGZO films in the air ambient exhibit resistance increase comparison with the N2 annealed sample, since oxidizer H₂O and O₂ in the air reduce Vo in a-IGZO films. In order to analyze the sheet resistance reduction, XPS analyses of B⁺ implanted a-IGZO films were carried out. Figure 3 (a) shows XPS global spectra for a-IGZO without implantation and with B⁺ 15 keV, 1×10^{15} cm⁻² implantation and with the N₂ annealing process after the implantation, where the binding energies were calibrated to common C1s peak at 284.6 eV. Figure 3 (b) shows XPS B1s spectra for a-IGZO without implantation and with the implantation and with the annealing process after the implantation. Considering B1s binding energy 188 eV [11] for B-B bonding and electron negativity relationship of Zn (1.65) < In (1.78) < Ga (1.81) < B (2.04) < O (3.44), we find that B-O bonding effect is dominant for the implanted boron. On the other hand, we previously reported that O1s spectra exhibit Vo decreases after the annealing [12]. Hence, we find a possibility that a-IGZO sheet resistance reduction can be realized by not only Vo but also the boron itself. In order to separate between the boron and the Vo effects in a-IGZO, we carried out the B⁺ 50 keV with 1 × 10¹⁵ cm⁻² implantation in 180-nm-thick a-IGZO film. In addition, we carried out depth profile analyses using Hall measurement combined with wet etching. From Hall measurements, as shown in Fig. 4 (a), (b) and (c), we obtained sheet resistances, Hall mobilities µmeas and sheet concentration n_{meas} as a function of the etching



Fig. 3 (a) X-ray photoelectron spectroscopy (XPS) global spectra for 50-nm-thick a-IGZO films without implantation and with B⁺ 15 keV, 1×10^{15} cm⁻² implantation and with the annealing process in N₂ ambient after the implantation. **(b)** XPS B1s spectra for 50-nm-thick a-IGZO films without implantation and with the B⁺ implantation and with the annealing process after the implantation.



Fig. 4 (a) Sheet resistance, **(b)** Hall mobilities and **(c)** sheet concentrations as a function of B⁺ implanted 180-nm-thick a-IGZO etching depth *x* from the a-IGZO surface before the etching processes, where the implantation condition is ion energy 50 keV and dose 1×10^{15} cm⁻², respectively.



Fig. 5 Local electron concentration n_j and local electron mobility μ_j model of the a-IGZO.

depth *x* from the a-IGZO surface. The sheet resistance slowly increases with increasing in *x* < 150 nm. In addition, both μ_{meas} and n_{meas} slowly decrease with increasing in *x* < 150 nm also, which indicates B⁺ penetration ability and resistance reduction ability are higher than that of Ar⁺ reported previously [12]. In order to estimate the local electron concentration n_j and local Hall mobility μ_j profiles as functions of a-IGZO depth, we considered a parallel conductance model in the a-IGZO film shown in Fig. 5. In case of the electrical transport properties development in each etching layer and a weak-magnetic-field ($\mu_{meas}B << 1$, where $\mu_{meas} = 1-20$ cm²/Vs as shown in the Fig. 4 and the magnetic flux density $B \sim 0.5$ Tesla) approximation applicable, two independent equations consist as follows:

$$\sigma_{\text{meas}}(x) = n_{\text{meas}}(x) \cdot \mu_{\text{meas}}(x), \ \sigma_j = n_j \cdot \mu_j,$$

$$\sigma_{\text{meas}}(x) \simeq \sum \sigma_j, \qquad (1)$$

$$\sigma_{\text{meas}}(x) \cdot \mu_{\text{meas}}(x) \simeq \sum \sigma_j \mu_j^2, \qquad (2)$$

where the σ_{meas} , n_{meas} , and μ_{meas} are conductance, electron concentration and Hall mobility obtained by the Hall measurements as shown in Fig. 4, respectively. Hence, n_i and μ_i as functions of a-IGZO depth can be estimated by the simultaneous equation (1) and (2). Figure 6 (a) and (b) shows concentrations and local Hall mobilities as functions of a-IGZO depth, where the depth zero corresponds to the a-IGZO surface. In addition, we estimated boron and Vo concentration profiles as shown in Fig. 6 (a), where the boron and Vo profiles are calculated by TRIM using the electronic and nuclear stopping powers for B in a-IGZO as shown in Fig.1 (b) and the bonding energies of In, Ga, Zn and O are ~ 3 eV [13], respectively. From comparison between the n_i and Vo and B concentrations, the Vo concentration profile is not agreement with $n_{\rm l}$ profile only, but also agreement the B profiles at the deep position around 150 nm of a-IGZO. On the other hand, μ_j profile also almost uniform value in the a-IGZO deep position, which indicates also that boron itself contribute the a-IGZO resistance reduction. We find that the Hall measurement with the wet etching results are good agreement with the Hall measurement of each ion energy combined the SIMS results and the XPS results.



Fig. 6 (a) Concentrations and **(b)** local electron mobilities as functions of a-IGZO depth, where the depth zero corresponds to the a-IGZO surface.

3 Summary

We investigated B⁺ implantation 50-nm-thick a-IGZO films on the glass, in which a-IGZO films were prepared by ICP sputtering system. On the other hand, the B⁺ implantations were carried out by using iG series. From Hall measurement, B⁺ implanted a-IGZO sheet resistances have bottom value in the range of 10-80 keV, which behaviors corresponds to B amounts in a-IGZO films, from which we obtained one of the minimum sheet resistance realized by optimized Elon. From the SIMS results and the XPS results and Hall measurements combined wet etching processes, we find a role of implanted boron in a-IGZO, which exhibits a-IGZO sheet resistance reduction can be realized by implanted boron itself. We expect that the B⁺ implantation technology for a-IGZO sheet resistance control technique is much useful for a-IGZO device processing.

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