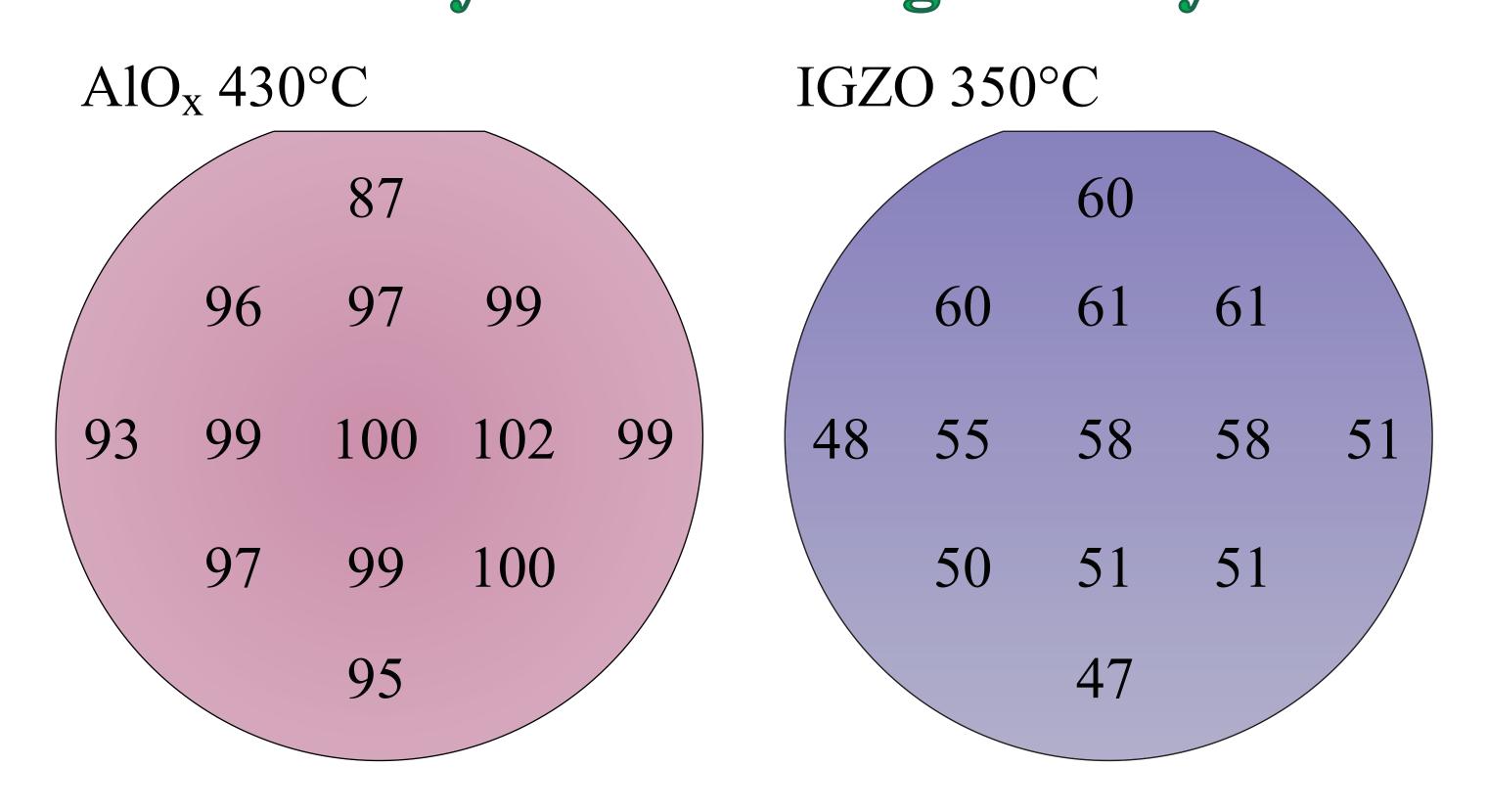
Properties of AlOx thin films grown by Mist CVD $2 \mu m$ Table 1. Deposition conditions of AlO_X thin films 430°C 300°C Solute Aluminum acetylacetonate (AlAcac₃)*2 Solvent (mixing ratio) Distillated water*3, Methanol*4 (10:90) Solution concentration 0.020 mol/LThickness \approx 50 nm, 200 nm $T > 400^{\circ} \text{C}$ 300, 350, 400, 430 °C Substrate temperature field, Substrate $p+-Si^{*5}$ $E_{BD} > 6.0 \text{ MV/cm}$ φ100 mm ver. FC type mist CVD system*6 Growth system k > 6.1Air, $2.5 \text{ L/min} \times 2$ Carrier gas / flow rate Dilution gas / flow rate Air, $10.0 \text{ L/min} \times 2$ RMS = 1.6 nmUltrasonic transducer*1 2.4 MHz, 24 V·0.625 A, 6 *1 HONDA ELECTRONICS, HM-2412 *2 Aluminum acetylacetonate, 99%, Sigma-Aldrich 8.460 nm 11.35 nm *3 Wako Pure Chemical Industries RMS: 1.615 nm RMS: 1.063 nm 400 *4 Methanol, 99.8%, Wako Pure Chemical Industries Temperature (°C) *5 Advantec *6 Refs. 9 and 16. (a) Breakdown field (E_{BD}) , and (b) surface structure of AlO_x thin films grown by mist CVD. The AlO_x thin films grown over 400°C exhibited the breakdown field and the dielectric constant of 6.0 MV/cm and 6.1, respectively. However, the E_{BD} of AlO_x thin film grown at temperatures below 350°C was dramatically declined. Temperature (K) v AlO₆ Compared with the AlO_x thin films deposited at various 700 600 temperature, the difference of the shoulder around 1100 cm⁻¹ 400°C, 206 nm (×12) $v AlO_4$ corresponding to the bending vibration of Al-O-H (v_s (Al)O-H $50 \vdash Ea =$ 350°C, 189 nm (×16) 8.0 (Sed) or v_{as} (Al)O-H) and the difference of peak around 2350 cm⁻¹ 22.5-24.0 kJ/mol corresponding to the stretching vibration of CO₂ adsorbed can (normaliz be seen. The logarithm of deposition ratio and surface roughness are Absorbance (5.0 c) not directly proportion to the reciprocal of the substrate temperature. The line has changed around 350°C. v_{as} (Al)O-H δH_2O v_s (Al)O-H It is suggested that a residual as boehmite $(\gamma-AlO(OH))$ remains in the AlO_x thin film grown at the temperature below 350°C. It is thought that is one of the reasons $1000/T (K^{-1})$ why the E_{BD} dramatically declines. 2000 3000 1000 4000 Deposition rate and surface roughness Wavenumber (cm⁻¹) of AlO_x thin films grown by mist CVD. FT-IR spectra of 200 nm thick AlO_x thin films grown by mist CVD. - Properties of IGZO thin films grown by Mist CVD $2 \mu m$ Table 2. Deposition conditions of IGZO thin films Temperature (K) (a) SPT 200°C (b) Mist 300°C (c) Mist 350°C 500 700 600 Solute1 Indium acetylacetonate (InAcac₃)*2 Gallium acetylacetonate (GaAcac₃)*³ Solute2 Solute3 Zinc acetylacetonate (ZnAcac₂)*4 Solvent Distillated water*5, Methanol*6 (mixing ratio: 10:90) 20 $0.030 (1:1:1)^{*7} \text{ mol/L}$ Solution concentration Deposition $\approx 200 \text{ nm}$ Thickness 150, 200, 250, 300, 350, 400, 430 °C Substrate temperature raito Substrate Quartz*8, Eagle XG*9 φ100 mm ver. FC type mist CVD system*10 Growth system Carrier gas / flow rate Air, $2.5 \text{ L/min} \times 2$ 121.5 nm 22.03 nm 9.360 nm Dilution gas / flow rate Air, $10.0 \text{ L/min} \times 2$ RMS: 19.34 nm Surface RMS: 1.060 nm RMS: 3.637 nm Ultrasonic transducer*1 2.4 MHz, 24 V·0.625 A, 6 roughness *1 HONDA ELECTRONICS, HM-2412 Surface structure of the IGZO thin films. *2 Indium(III) acetylacetonate, 99.99%, Sigma-Aldrich *3 Gallium(III) acetylacetonate, 99.99%, Sigma-Aldrich *4 Zinc acetylacetonate, >95%, nacalai tesque Surface of M-IGZO is rough compared with that of *5 Wako Pure Chemical Industries $1000/T (K^{-1})$ SPT-IGZO. Especially, in this case, Surface of *6 Methanol, 99.8%, Wako Pure Chemical Industries M-IGZO is roughest at 300°C. *7 In:Ga:Zn, atomic ratio Deposition ratio and surface roughness of *8 Mitorika Glass the IGZO thin films grown by the mist CVD *9 Corning Deposition ratio of M-IGZO does not depend on *10 Refs. 9 and 16. temperature. This is because the dominant reaction shifts at each temperature domain due to each precursor with variant chemical reaction rate. $(b)_{60}$ (a) Both M-IGZO and SPT IGZO were amorphous $T = 350^{\circ} \text{C}$ since no diffraction peak was detected in XRD Mist Spt amorphous measurement. deposition rate > 25 nm /min — Ga From RBS measurement, the composition ratio of SPT-IGZO — Zn In:Ga:Zn = 16:37:47M-IGZO and SPT-IGZO were so different from ideal Component elemental ratio. In a SPT-IGZO, each metal composition ratio was 36:27:37 (In:Ga:Zn) owing to different sputtering ratio of each metal oxide contained M-IGZO, 350°C Change of dominant reaction in the target. In the mist CVD, composition ratio of depending on precursor is observed. films depends on growth temperature and mixture ratio 300 of source materials at preparation. Hence, in a Temperature (°C) 2θ Cu Kα (degree)

(a) XRD $2\theta/\omega$ spectra and (b) composition ratio of the IGZO thin films

- Uniformity of thin film grown by Mist CVD

grown by the mist CVD



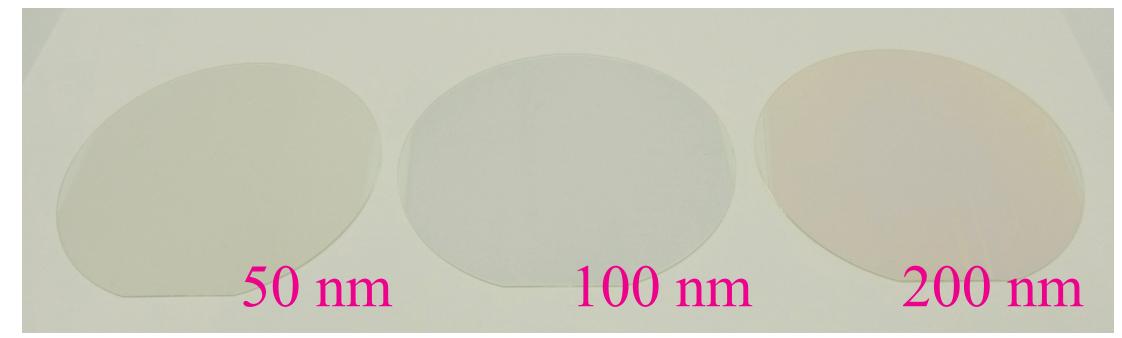
The photograph of IGZO thin films grown on the substrate of ϕ 100 mm.

16:37:47 (In:Ga:Zn).

M-IGZO grown at 350°C, which is a same as the

condition of IGZO thin film used for the channel layer

of the oxide TFT, each metal composition ratio was



AlO_x: Approximately-uniformity IGZO: $\pm 10\%$

In mist CVD, uniformity thin films can be get.