Thermal atomic layer deposition of In$_2$O$_3$ thin films using a homoleptic indium triazenide precursor and water

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Abstract

Indium oxide (In$_2$O$_3$) is an important transparent conducting material widely used in optoelectronic applications. Herein, we study the deposition of In$_2$O$_3$ by thermal atomic layer deposition (ALD) using our recently reported indium(III) triazenide precursor and H$_2$O. A temperature interval with self-limiting growth was found between ~270–385°C with a growth per cycle of ~1.0 Å. The deposited films were polycrystalline cubic In$_2$O$_3$ with In:O ratios of 1:1.20, and low levels of C and no detectable N impurities. The transmittance of the films was found to be >70% in visible light and the resistivity was found to be 0.2 mΩcm. The high growth rates, low impurities, high optical transmittance, and low resistivity of these films give promise to this process being used for ALD of In$_2$O$_3$ films that are good candidates for potential display and touch-screen applications.

I. INTRODUCTION

Indium oxide (In$_2$O$_3$) is a material of high interest due to its high electrical conductivity and optical transparency, making it a key material for transparent and optoelectronics, e.g. microelectronic displays on touch screens. The electrical conductivity of In$_2$O$_3$ is controlled by the stoichiometry of the film and is typically enhanced by substituting the In atoms with 5-10 atomic % Sn. All applications of In$_2$O$_3$ as transparent conducting layers require the deposition of high-quality thin films of In$_2$O$_3$. This requirement is the driving force for extensive studies by sputtering, thermal evaporation, chemical vapor deposition, and atomic layer deposition (ALD). ALD of In$_2$O$_3$ is especially interesting as it can deposit high quality films with precise thickness, controlled composition, low impurity contents, and excellent conformality on complex substrates has been demonstrated.
ALD of In$_2$O$_3$ was first reported using InCl$_3$ and either H$_2$O or H$_2$O$_2$ at temperatures between 300-500 °C. The low vapor pressure of InCl$_3$, the generation of HCl as the reaction by-product and the possibility for InCl$_3$ to etch the growing In$_2$O$_3$ layer has motivated the use of other In precursors. Furthermore, there has been limitations on availability of efficient indium precursors that give processes with high growth rates, fully self-limiting reactions, low impurity contents and good film crystallinity. The indium compounds tested as precursors in combination with different oxidants, within different deposition temperature windows are shown in Table 1.

Table 1: Indium precursors and oxidants used for ALD of In$_2$O$_3$ films.

| Indium precursor | Oxidant           | Temperature window (°C) | Ref   |
|------------------|-------------------|-------------------------|-------|
| InCl$_3$         | H$_2$O/H$_2$O$_2$ | 300 – 500               | 11,13,14. |
| InMe$_3$         | H$_2$O            | 200 – 250               | 15,16. |
| InMe$_3$         | O$_3$             | 100 – 200               | 17. |
| InCp             | O$_3$             | 200 – 450               | 12. |
| InCp             | H$_2$O/O$_2$      | 100 – 350               | 18,19,20,21. |
| InCp             | H$_2$O$_2$        | 160 – 200               | 10,22. |
| In(acac)$_3$     | H$_2$O or O$_3$   | 165 – 220(H$_2$O), 160 – 225 (O$_3$) | 23. |
| In(tmhd)$_3$     | O$_2$ plasma      | 100 – 400               | 24. |
| In(guan)$_3$     | H$_2$O$_2$        | 230 – 300               | 25. |
| Et$_2$InN(TMS)$_2$ | H$_2$O        | 175 – 250               | 26. |
| InNCA, DADIn, InEt$_3$ | O$_3$       | 100 – 200               | 27. |
| InNCA            | H$_2$O$_2$        | 125 – 225               | 28. |
| In(dmamp)$_3$    | O$_3$             | 100 – 300               | 29. |
| Me$_2$In(EDPA)   | O$_2$ plasma      | 70 – 250                | 30. |
| In(famd)$_3$ and In(amd)$_3$ | H$_2$O | 150 – 275 and 225 – 300 | 31. |
| In(CH$_3$)$_3$[CH$_3$OCH$_2$CH$_2$NHtBu] | H$_2$O | 150 – 200 | 32. |
| DATIn            | O$_2$ plasma      | 100 – 200               | 33. |
| InCp             | O$_2$ plasma      | 225 – 275               | 34. |

Cp = cyclopentadienyl,

acac = acetylacetonate,

tmhd = 2,2,6,6-tetramethyl-3,5-heptanedionate,

guan = N,N-dimethyl-N’N”-diisopropylguanidinato,

famd = N,N’-diisopropylformamidinato,

amd = N,N’-diisopropylamidinato,

DAT = dimethylbutylamino-trimethyl,

TMS = trimethylsilyl,
$NCA = \text{diethyl}[1,1,1\text{-trimethyl-N- (trimethylsilyl)silanaminato},$

$DAD = [3\text{-}(\text{dimethylamino})\text{propyl}]\text{dimethyl},$

$\text{dmamp} = 1\text{-dimethylamino-2-methyl-2-propoxy},$

$EDPA = N\text{-ethoxy-2,2-dimethylpropanamido}.$

Amidinate and guanidinate ligands have been employed to improve the thermal stability of In precursors whilst maintaining sufficient reactivity for surface reactions. This has led to the formation of homoleptic hexacoordinated In–N bonded precursors, In(III) tris-formamidine$^{31}$ (In(famd)$_3$) 1, amidinate$^{35}$ (In(amd)$_3$) 2 and guanidinate$^{36}$ (In(guan)$_3$) 3 (Fig. 1a). As the coordination sphere of the metal centre is all In–N bonds, their use for In$_2$O$_3$ is especially attractive as it makes formation of In–O bonds thermodynamically favourable. This would allow H$_2$O to be used as the co-reactant and remove the need for strong oxidants such as O$_3$ and O$_2$ plasma. Precursors 1-3 have been used with H$_2$O for thermal ALD of In$_2$O$_3$ with all showing self-limiting growth behaviour.$^{25,31}$ Although all precursors gave similar growth rates of In$_2$O$_3$ (0.45-0.55 Å/cycle), 1 was found to have faster kinetics with the surface and H$_2$O than 2 and 3. A surface chemical model was used to explain the faster kinetics of 1, which was thought to be due to its smaller endocyclic carbon substituent. This allowed the exocyclic N-isopropyl groups of the ligand to fold up more, leading to a surface species with less surface repulsion and a more exposed In centre for its subsequent reaction with H$_2$O.$^{31}$ This surface chemical model was further confirmed when we used 1-3 for plasma ALD of InN.$^{37}$ Again, 1 was found to be superior to 2 and 3. To continue this trend, we replaced the substituted endocyclic carbon to a nitrogen atom. This led to our discovery of the indium(III) triazenide (4) (Fig. 1b), which we used with NH$_3$ plasma to render high-quality InN.$^{38}$
Here, we investigate the use of 4 as In precursor in thermal ALD of In$_2$O$_3$ with water as oxidant. The ALD characteristics, optical and electrical properties of the deposited In$_2$O$_3$ films are investigated and a direct comparison between 4 and 1 is done.

II. EXPERIMENTAL

A. Film deposition

Precursors 1 and 4 were synthesised using literature procedures$^{37,38}$. Films were deposited using a homebuilt crossflow ALD reactor at 50 hPa. A flow of N$_2$ (99.999%) was used as the carrier and purging gas. At all times during the deposition, the N$_2$ pressure was kept constant. Precursors 1 or 4 and water were used as the In and O sources, respectively. The precursors were introduced separately into the reactor via stainless steel gas lines, which were extended into the reactor tube to ensure maximum substrate exposure. Precursors 1 and 4 were placed in a stainless-steel bubbler, which was heated to 120 °C, the gas delivery line was heated to 140 °C and flow of precursor into the chamber was aided by N$_2$ carrier gas. The deionized H$_2$O was kept at room temperature and delivered into the chamber without a carrier gas. Film deposition was undertaken on approximately 2×3 cm Si (100) and glass substrates, which were cleaned by dry N$_2$ gas and used without further chemical cleaning. After loading the substrates, the chamber was baked at 155 °C overnight. The deposition zone of the reactor was defined by the

\[ R \quad \text{1 (R = H)} \quad \text{2 (R = Me)} \quad \text{3 (R = NMe}_2\text{)} \]
heated zone of the tube furnace. Unless otherwise noted, a typical ALD cycle consisted of a 4 s pulse of 1 or 4 and 3 s pulse of H$_2$O, with 10s N$_2$ purges after the In precursor and the water pulses. The deposition process was studied between 150 and 520 °C.

B. Film characterization

The crystallographic phases of the films were characterized by a X-ray diffractometer (XRD) PANalytical X’pert Pro equipped with a Cu Kα X-ray source ($\lambda = 1.54$Å) in the $\theta$-2$\theta$ mode. A Ni foil was used to filter the Kβ radiation. Film thickness was measured using the same PANalytical X’Pert Pro X-ray diffractometer in X-ray reflectivity (XRR) mode. The film thickness was obtained from the XRR data using X’Pert Reflectivity software and a two-layer model, In$_2$O$_3$/Si, to fit the data. The film morphology was analyzed by scanning electron microscopy (SEM) using a LEO 1550 scanning electron microscope with an acceleration energy of 2kV using the in-lens detector at a working distance of 3-6mm. SEM was also used to obtain film thickness on samples that were too thick or rough for XRR measurements. The composition and impurity levels of the deposited films were probed with high resolution X-ray photoelectron spectroscopy (XPS), Kratos AXIS Ultra DLD, equipped with an Ar (0.30 eV) sputtering source for 600 s, employing monochromatic Al Kα radiation (hν =1486.6 eV). Sputter-cleaning (sputter-etching) was undertaken to eliminate surface contamination, which resulted after exposure of the substrate to air. Analysis of the photoelectron spectra was completed using the CasaXPS software package. By quantitative analysis, the signals originating from the substrate and the thin film could be de-convoluted and the chemical composition was obtained. Gaussian-Laurentius functions and Shirley background were used to fit the experimental XPS data.

The optical, transmittance, absorbance, and bandgap properties of the films were obtained using a Shimadzu UV-2450 UV-VIS spectrophotometer in a wavelength range of 250–800 nm. The UV-VIS spectrophotometer was operated in the reflectance mode in analyzing films deposited on glass substrates. Determination of band gap was determined using the Tauc plot formalism from data obtained by UV-Vis spectrophotometry based on the Tauc relation: $(\alpha h\nu)^{1/y} = \beta (h\nu – E_g)$ where $\alpha$ is the absorption coefficient (absorbance/film thickness), $\beta$ is the band tailing parameter, $h$ is the Planck’s constant, $\nu$ is the frequency of incident light, $E_g$ is the energy of the optical band gap and $y$ is the
power factor, which depends upon the nature of the transition (semi-conducting materials, \( y = 0.5 \) for direct allowed transitions). Therefore: \((\alpha h\nu)^2 = \beta(h\nu - E_g)\).

By drawing a tangential line from the plotted graph of \( h\nu \) versus \((\alpha h\nu)^2\), the value of the band gap was determined as the value of \( h\nu \) at the point where this line crosses the axis i.e. the intercept of the extrapolation to near zero absorption with photon energy axis i.e. \((\alpha h\nu)^2 \rightarrow 0\). Electrical characterization by resistivity measurements were performed using the 4-point probe technique. Measurements were done on a Jandel, Model RM3000 test unit or probing system, which was first calibrated using a standard glass sample before sample resistivities were determined.

### III. RESULTS AND DISCUSSION

#### A. Film deposition

The ALD process with 4 and water was studied by varying their pulse lengths 290 °C. The resulting saturation curves (Fig. 2a) show the deposition process is self-limiting for pulse times \( \geq 3 \) s, resulting in film growth of \(~1.0\ \text{Å}/cycle\). Varying the deposition temperature whilst using a 4 s pulse of 4 and 3 s pulse of water revealed a temperature interval where the growth per cycle (GPC) is constant with the temperature between 270 and 385 °C (Fig. 2b). The film thickness for films deposited at 290 °C using 300, 500, 800 and 1100 ALD cycles (Fig. 2c) follows a linear trend line, however, it shows a significantly slower initial deposition interpreted as a nucleation delay of approximately 49 cycles. This result suggests slow surface kinetics of 4 at low temperatures due to steric crowding of the metal centre. Similar behaviour was also reported for 1, whereby a ‘fill-empty’ approach was used to circumvent this problem.\(^{37}\) Attempts to fit a linear trend line results in a line suggesting an average growth per cycle of 0.9 Å.
a) Graph showing growth per cycle (GPC) as a function of pulse time.

b) Graph showing growth per cycle (GPC) as a function of deposition temperature.

- 4 pulse
- H$_2$O pulse

Temperature window: 270°C to 385°C
Figure 2. a) Growth per cycle behaviour of 4 and H₂O pulses at 290 °C on Si (100). b) Growth dependence on process temperature using a 4 s pulse of 4 and 3 s pulse of H₂O on Si(100). c) Growth behaviour of 4 at 290 °C dependent on the number of cycles using a 4 s pulse of 4 and 3 s pulse of H₂O on Si(100).

B. Film properties

The XRD analysis, acquired in the θ-2θ mode, of films deposited at 315 °C and 430 °C (Fig. 3) showed cubic In₂O₃ as the only crystalline phase except the Si substrate. In addition, the films are polycrystalline with a preferred (222) orientation.
Figure 3. θ-2θ XRD analysis of deposited films at 315 °C (50nm thick) and 430 °C (96nm thick), showing polycrystalline cubic In$_2$O$_3$ films.

XPS analysis of the films showed the C impurities range from 0.8 to 3.5 at. % and N impurities are below the detection limit. As summarized in Table 2, XPS analysis also shows the films are oxygen deficient, which is comparable to literature. The elemental composition of the films displayed in Table 2 indicates that higher deposition temperatures and water pulse times lower the impurity concentrations. A longer water pulse also increases the O content and reduces the impurity content of the films. The oxygen deficient nature of the deposited films opens the possibility of O vacancies in the In$_2$O$_3$ films. Oxygen vacancies are point defects critical for explaining the optoelectrical properties of metal oxide thin films.
**Table 2: Elemental percentages of 50nm thick In\textsubscript{2}O\textsubscript{3} films deposited at different temperatures (H\textsubscript{2}O pulse of 3 s) and different H\textsubscript{2}O pulse times (at temperature 290 °C).**

| Deposition temperature (°C) | C (at. %) | In (at. %) | O (at. %) | Atomic ratio In:O (Expected 1:1.5) |
|-----------------------------|-----------|------------|-----------|-----------------------------------|
| 290                         | 3.5       | 43.5       | 53.0      | 1:1.2                             |
| 315                         | 1.4       | 45.0       | 53.6      | 1:1.2                             |
| 385                         | 0         | 51.5       | 48.5      | 1:0.9                             |

| Water pulse length (s) | C (at. %) | In (at. %) | O (at. %) | Atomic ratio In:O (Expected 1:1.5) |
|------------------------|-----------|------------|-----------|-----------------------------------|
| 3                      | 4.3       | 43.2       | 52.5      | 1:1.2                             |
| 6                      | 1.3       | 45.3       | 53.4      | 1:1.2                             |
| 9                      | 0.8       | 45.6       | 53.6      | 1:1.2                             |
| 12                     | 1.3       | 44.7       | 54.0      | 1:1.2                             |

Top view SEM images of 50nm and 96nm thick films deposited at 290 °C and 430 °C respectively (Fig. 4) show smooth films at lower temperatures (Fig. 4a), and larger, flattened, and faceted grains at higher temperatures (Fig. 4b). Cross-section micrographs further indicate smooth and uniform films grown at 315 °C (Fig. 4c).
The results above shows that In$_2$O$_3$ can be deposited from 4 and water in an ALD process. We speculate that the primary reaction mechanism for the film deposition is a surface chemical reaction of ligand exchange type, similar to what has been suggested for ALD of In$_2$O$_3$ from 1 and water.\textsuperscript{31} Presumably, 4 loses one or two triazenide ligands upon chemisorption to the surface by reacting with surface hydroxyl groups, forming In–O bonds and protonated triazenide ligands. A similar ligand exchange takes place during the water pulse to remove the remaining surface triazenide ligands and replacing them with new hydroxyl groups.

The resistivity of approximately 50 nm In$_2$O$_3$ films deposited on glass at 290 °C with different water pulse lengths was determined by four-point probe measurements. The film resistivity reflects the electrical properties of a film since it is determined by the carrier concentration and carrier mobility. Electrical conductivity of In$_2$O$_3$ is reported to be mainly controlled by the oxygen stoichiometry in the film.\textsuperscript{42} A perfect In$_2$O$_3$ stoichiometry is not ideal since it has no oxygen vacancies, which results in increased resistivity.\textsuperscript{43} In$_2$O$_3$ typically exhibits n-type semiconducting properties from the oxygen vacancies\textsuperscript{19}, which increase carrier density and hence act as donors.\textsuperscript{1} The deposited films in this study show a general decrease in resistivity, i.e., an increase in conductivity, with longer water pulse time. The improved conductivity for films deposited with increased H$_2$O dosage may be attributed to incorporation of hydroxyl groups which cause self-doping.\textsuperscript{44} The resistivities of different film samples, all with approximately 50 nm
thicknesses, deposited with 6 s, 9 s and 12 s of water pulse were 1.2; 2.6 and 0.16 mΩcm, respectively (Table 3). Upon postdeposition annealing at 520 °C for 2 hours in air, the In:O ratio in the film changed from 1:1.2 to 1:1.3 but this lead to an increase in the resistivity from e.g., 1.2 to 2.0 mΩ.cm (shown in Table 3). This result is in line with previous reports on the importance of the oxygen vacancies for the electrical conductivity of the In$_2$O$_3$ films.

Table 3: Resistivities of In$_2$O$_3$ films grown at different oxidant doses, all 50nm thick.

| H$_2$O pulse (s) | Average sheet resistance as deposited (Ohms per square) | Resistivity (mΩ.cm) | H$_2$O pulse (s) | Average sheet resistance after post-annealing at 520°C (Ohms per square) | Resistivity (mΩ.cm) |
|----------------|----------------------------------------------------------|---------------------|----------------|---------------------------------------------------------------------|--------------------|
| 6              | 238.8                                                    | 1.2                 | 6              | 400.6                                                               | 2.0                |
| 9              | 506.8                                                    | 2.6                 | 9              | 1500.0                                                              | 7.5                |
| 12             | 32.2                                                     | 0.16                | 12             | 40.4                                                                | 0.2                |

Optical transparency of the films was obtained from the transmission spectrum of 50 nm thick In$_2$O$_3$ films deposited at 290 °C on glass referenced to the spectrum of the uncoated glass substrate. The resultant average transmission of the In$_2$O$_3$ film in the wavelength range 250–800 nm was calculated using the relationship $A = 2 \cdot \log_{10} \left(\frac{T}{100}\right)$, where $A$ is absorbance and $T$ is transmittance of light. The spectra in Fig. 5 shows the optical transmittance for the In$_2$O$_3$ films with uncoated glass included for reference. The highest obtained optical transmittance was 71 %, which is very similar to an uncoated glass substrate, which showed 74 % transmittance. The optical transmittance of the In$_2$O$_3$ films is comparable to values already reported in literature.$^{45,46}$ With uncoated glass being the control, it would be 71/74 x 100% which equals 96% transmittance in relation to glass. Fig. 5 shows that the transmittance increases with longer water pulse, which from Table 2, renders higher oxygen content in the film.

The results from the resistivity and transmittance measurements are in line with the literature on In$_2$O$_3$ where the oxygen stoichiometry in the films which has been shown to impact both optical transmission and electrical conductivity of the films.$^{19}$ The high concentration of oxygen vacancies causes our films to be more conducting than those reported in previous studies.$^{42,47}$
Figure 5. Optical transmittance of the In$_2$O$_3$ films deposited with different lengths of the water pulse. The transmittance increases with increasing water pulse time.

Tauc plots of the In$_2$O$_3$ films deposited with different water pulse lengths are shown in Fig. 6, indicating that the direct optical band gap varies between 4.04 and 4.22 eV with higher band gap for shorter water pulses. The bandgap is lower for higher oxygen content in the films. The obtained band gap (4.0 - 4.2 eV) is comparable to previously reported values but is slightly higher than the expected range (3.5 - 3.9 eV). The higher band gap is ascribed to the higher deposition temperatures used in our study, which are higher than what is typically employed in ALD of In$_2$O$_3$. From our XPS results (Table 2), higher deposition temperatures render lower oxygen content and carbon incorporation in the films. We speculate that this leads to an increase of oxygen vacancies$^{42}$ that can supply free electrons$^{43}$ in the material leading to an increase in the optical band gap. A higher temperature can also increase the optical band gap by decreasing the density of grain boundaries, leading to fewer electrons being trapped in grain boundaries, increasing the
number of free electrons.\textsuperscript{48} This increase of band gap can be further explained by the gap widening phenomenon called the Burstein–Moss shift\textsuperscript{49}.

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{figure6.png}
\caption{Tauc plots for estimating the optical band gaps of In\textsubscript{2}O\textsubscript{3} films deposited with different water pulse times.}
\end{figure}

C. Comparison to the In(famd)\textsubscript{3} precursor (1).

From the results above, ALD using 4 and water renders In\textsubscript{2}O\textsubscript{3} films with properties comparable to the previously reported In\textsubscript{2}O\textsubscript{3} films deposited using 1, albeit 4 seems to require higher deposition temperatures. A temperature window of 150 – 275 °C for In\textsubscript{2}O\textsubscript{3} films using 1 and H\textsubscript{2}O was reported, but slow surface kinetics of 1 were observed.\textsuperscript{31} This was then circumvented using a stop-flow process to give 1 more time to react with the surface. We cannot do stop-flow ALD in our ALD reactor, and therefore undertook a direct comparison between 1 and 4 by studying ALD of In\textsubscript{2}O\textsubscript{3} using 1 and water in our reactor. We found a temperature interval of 245 – 315 °C where the GPC is constant with the temperature when using a 5 s pulse of 1 and 4 s pulse of H\textsubscript{2}O (Fig. 7). This is higher than the 150 – 275 °C reported for 1, again suggesting a slow surface kinetics for 1.
Figure 7. Growth per cycle vs. deposition temperature for ALD of In$_2$O$_3$ using 1 and water in our ALD reactor, which does not have stop flow mode.

A stop flow mode was used to allow for a lower deposition temperature of In$_2$O$_3$ using 1, whilst we employed a multiple-pulse approach using 1 for depositing InN by ALD.\textsuperscript{37} The higher temperature interval found for ALD of In$_2$O$_3$ using 1 in our ALD reactor suggest that the temperature window obtained for 4 could possibly also be lowered by using stop flow mode in the ALD process. What perhaps speaks against this is that ALD of InN using 4 did not require the multiple pulse approach used for ALD of InN using 1.
IV. SUMMARY AND CONCLUSIONS

In summary, we used our recently reported In(III) triazenide precursor in combination with H\textsubscript{2}O to deposit In\textsubscript{2}O\textsubscript{3} films in a thermal ALD process. Polycrystalline In\textsubscript{2}O\textsubscript{3} films with a cubic structure were successfully deposited. The GPC, impurities, and In/O ratios are in line with the literature for other In precursors. The transmittance was found to be >70% in visible light and the resistivity was found to be 0.2 mΩcm. A temperature interval with self-limiting growth was found between 270 – 385°C. This is higher than that for ALD of In\textsubscript{2}O\textsubscript{3} using the previously reported In(famd\textsubscript{3}) precursor. After a direct comparison between the two In precursors used in this study, we suggest the temperature interval for 4 could be lowered by using a stop flow mode ALD process.

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