

Cite this: *Dalton Trans.*, 2022, **51**, 4712Received 5th November 2021,
Accepted 22nd February 2022

DOI: 10.1039/d1dt03748j

rsc.li/dalton

Thermal atomic layer deposition of In₂O₃ thin films using a homoleptic indium triazenide precursor and water

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Indium oxide (In₂O₃) is an important transparent conducting material widely used in optoelectronic applications. Herein, we study the deposition of In₂O₃ by thermal atomic layer deposition (ALD) using our recently reported indium(III) triazenide precursor and H₂O. A temperature interval with self-limiting growth was found between ~270 and 385 °C with a growth per cycle of ~1.0 Å. The deposited films were polycrystalline cubic In₂O₃ with In : O ratios of 1 : 1.2, 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 mΩ cm. 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 In₂O₃ films for future microelectronic displays.

1. Introduction

Indium oxide (In₂O₃) is a material of high interest due to its high electrical conductivity and optical transparency, making it a key material for transparent and optoelectronics¹ e.g. microelectronic displays on touch screens. The electrical conductivity of In₂O₃ is controlled by the stoichiometry of the film^{2,3} and is typically enhanced by substituting the In atoms with 5–10 atomic % Sn.^{4,5} All applications of In₂O₃ as transparent conducting layers require the deposition of high-quality thin films of In₂O₃. This requirement is the driving force for extensive studies by sputtering,⁶ thermal evaporation,⁷ chemical vapor deposition,⁸ and atomic layer deposition (ALD).⁹ ALD of In₂O₃ is especially interesting as it can deposit high quality films with precise thickness, controlled composition, low impurity contents, and excellent conformality on complex substrates.¹⁰

ALD of In₂O₃ was first reported using InCl₃ and either H₂O or H₂O₂ at temperatures between 300–500 °C.¹¹ The low vapor pressure of InCl₃, the generation of HCl as the reaction by-product and the possibility for InCl₃ to etch the growing In₂O₃ layer¹² has motivated the use of other In precursors. Furthermore, there has been limitations on availability of efficient In precursors that give processes with high growth rates, fully self-limiting reactions, low impurity contents and good film crystallinity. The In compounds tested as precursors

in combination with different reactants, within different deposition temperature windows are shown in Table 1.

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-formamidinate³¹ (In(famd)₃) **1**, amidinate³⁵ (In(amd)₃) **2** and guanidinate³⁶ (In(guan)₃) **3** (Fig. 1a). As the coordination sphere of the metal centre is all In–N bonds, their use for In₂O₃ is especially attractive as it makes formation of In–O bonds thermodynamically favourable. This would allow H₂O to be used as the co-reactant and remove the need for strong reactants such as O₃ and O₂ plasma. Precursors **1–3** have been used with H₂O for thermal ALD of In₂O₃ with all showing self-limiting growth behaviour.^{25,31} Although all precursors gave similar growth rates of In₂O₃ (0.45–0.55 Å per cycle), **1** was found to have faster kinetics with the surface and H₂O 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 H₂O.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 NH₃ plasma to render high-quality InN.³⁸

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Table 1 Indium precursors and oxidants used for ALD of In₂O₃ films

Indium precursor	Reactant	Temperature window (°C)	Ref.
InCl ₃	H ₂ O/H ₂ O ₂	300–500	11, 13 and 14
InMe ₃	H ₂ O	200–250	15 and 16
InMe ₃	O ₃	100–200	17
InCp	O ₃	200–450	12
InCp	H ₂ O/O ₂	100–350	18–21
InCp	H ₂ O ₂	160–200	10 and 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	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 NH ^t Bu]	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 = diethyl[1,1,1-trimethyl-*N*-(trimethylsilyl)silanaminato], DAD = [3-(dimethylamino)propyl]dimethyl, dmamp = 1-dimethylamino-2-methyl-2-propoxy, EDPA = *N*-ethoxy-2,2-dimethylpropanamido.



Fig. 1 (a) Evolution of In(III) precursors 1–3, used in various indium-based ALD studies, and (b) 4.

Here, we investigate the use of **4** as In precursor in thermal ALD of In₂O₃ with H₂O as reactant. The ALD characteristics, optical and electrical properties of the deposited In₂O₃ films are investigated and a direct comparison between **4** and **1** is undertaken.

II. Experimental

A. Film deposition

Precursors **1** and **4** were synthesised using literature procedures.^{37,38} Films were deposited using a homebuilt cross-flow ALD reactor at 50 hPa. A flow of N₂ (99.999%) was used as the carrier and purging gas. At all times during the deposition, the N₂ pressure was kept constant. Precursor **1** or **4** and H₂O 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 N₂ carrier gas. The deionized H₂O 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 N₂ 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 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 H₂O, with 10 s N₂ 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 ($\lambda = 1.54 \text{ \AA}$) in the θ - 2θ mode. A Ni foil was used to filter the K β 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, In₂O₃/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 2 kV using the in-lens detector at a working distance of 3–6 mm. 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 K α radiation ($h\nu = 1486.6$ 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 α is the absorption coefficient (absorbance/film thickness), β is the band tailing parameter, h is the Planck's constant, ν 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 H₂O was studied by varying their pulse lengths at 290 °C. The resulting saturation curves (Fig. 2a) show the deposition process is self-limiting for pulse times ≥ 3 s, resulting in film growth of ~ 1.0 Å per cycle. Varying the deposition temperature whilst using a 4 s pulse of 4 and 3 s pulse of H₂O 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 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 presumably due to steric crowding of the metal centre. Similar behaviour was also reported for ALD of

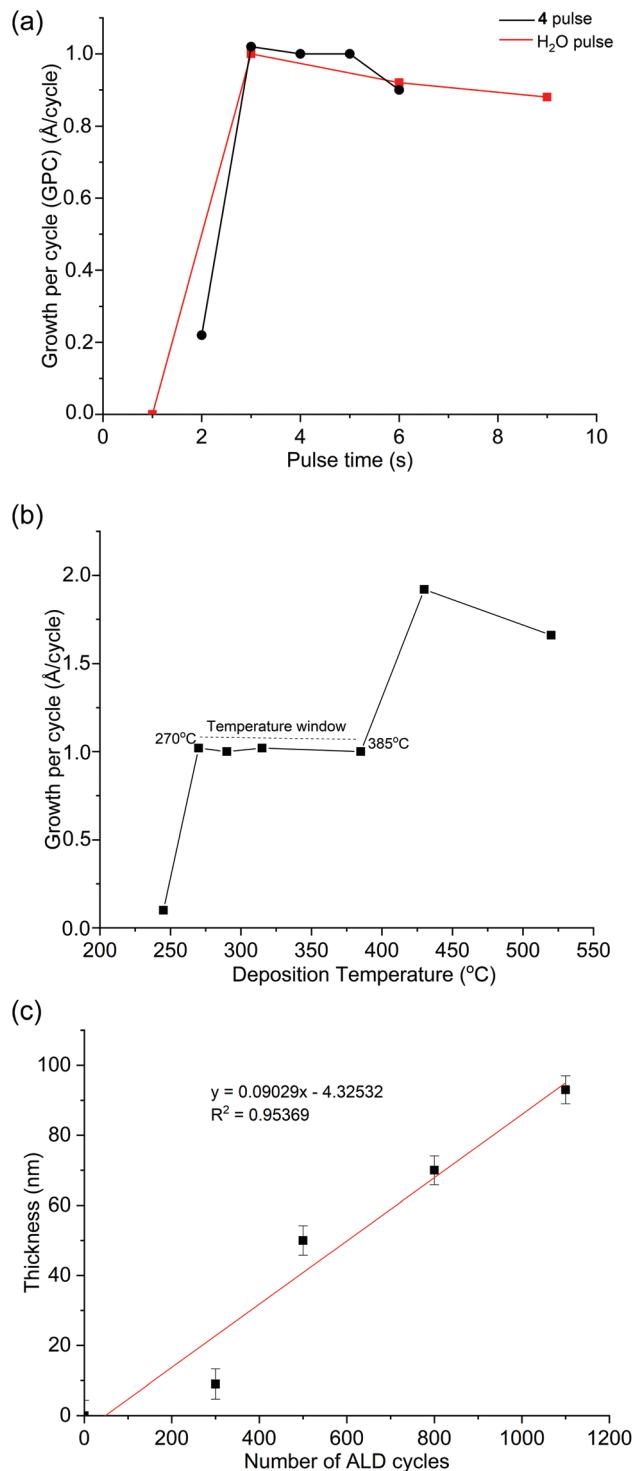


Fig. 2 (a) Growth per cycle behaviour of 4 and H₂O pulses at 290 °C on Si (100). (b) Growth dependence on process temperature using a 4 s pulse of 4 and 3 s pulse of H₂O on Si(100). (c) Growth behaviour of 4 at 290 °C dependent on the number of cycles using a 4 s pulse of 4 and 3 s pulse of H₂O on Si(100).

InN from 1, whereby multiple pulses of 1 was used to circumvent this problem.³⁷ Attempts to fit a linear trend line results in an average growth per cycle of 0.9 Å.



B. Film properties

The θ - 2θ XRD analysis of films deposited at 315 °C and 430 °C (Fig. 3) showed cubic In_2O_3 as the only crystalline phase except the Si substrate. In addition, the films are polycrystalline with a preferred (222) orientation.

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.¹⁰ The elemental composition of the films displayed in Table 2 indicates that higher deposition temperatures and H_2O pulse times lower the impurity concentrations. A longer H_2O 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_2O_3 films. Oxygen vacancies are point defects³⁹ critical for explaining the optoelectrical properties of metal oxide thin films.⁴⁰

Top view SEM images of 50 nm and 96 nm thick films deposited at 290 °C and 430 °C respectively (Fig. 4) showed smooth films at lower temperatures (Fig. 4a), and larger, flattened, and faceted grains at higher temperatures (Fig. 4b). Cross-section micrographs further indicated smooth and uniform films grown at 315 °C (Fig. 4c).

The results above show In_2O_3 can be deposited from **4** and H_2O 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_2O_3 from **1** and water.³¹ Presumably, **4** loses one or two triazenide ligands upon chemisorption by reacting with surface hydroxyl groups, forming In–O bonds and protonated triazenide ligands. A similar ligand exchange

Table 2 Elemental percentages of 50 nm thick In_2O_3 films deposited at different temperatures (H_2O pulse of 3 s) and different H_2O 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)				
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

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_2O_3 films deposited on glass at 290 °C with different H_2O 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_2O_3 is reported to be mainly controlled by the oxygen stoichiometry of the film.⁴¹ A perfect In_2O_3 stoichiometry is not ideal since it has no oxygen vacancies, which results in increased resistivity.⁴² In_2O_3 typically exhibits n-type semiconducting properties from the oxygen vacancies,¹⁹ which increase carrier density and hence act as donors.¹ The deposited films in this study show a general decrease in resistivity, *i.e.*, an increase in conductivity, with longer H_2O pulse time. The improved conductivity for films deposited with increased H_2O dosage may be attributed



Fig. 3 The θ - 2θ XRD analysis of deposited films at 315 °C (50 nm thick) and 430 °C (96 nm thick), showing polycrystalline cubic In_2O_3 films.



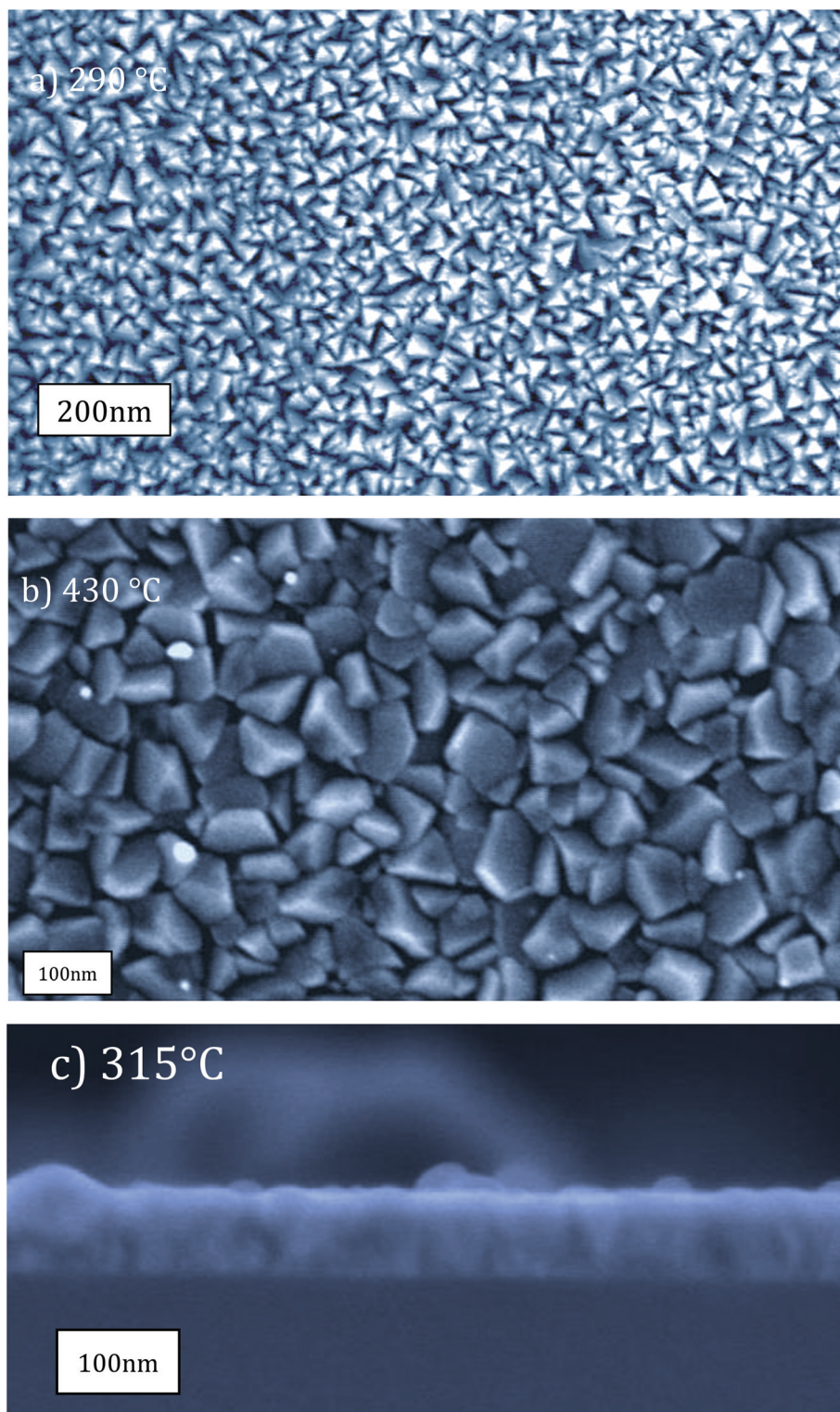


Fig. 4 SEM micrographs for films deposited at (a) 290 °C, (b) 430 °C and at (c) 315 °C.

to incorporation of hydroxyl groups which cause self-doping.⁴³ The resistivities of different film samples, all with approximately 50 nm thicknesses, deposited with 6 s, 9 s and 12 s of H₂O pulse were 1.2; 2.6 and 0.16 mΩ cm, 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 led 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 pre-



Table 3 Resistivities of In₂O₃ films grown at different oxidant doses, all 50 nm 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

vious reports on the importance of the oxygen vacancies for the electrical conductivity of the In₂O₃ films.

Optical transparency of the films was obtained from the transmission spectrum of 50 nm thick In₂O₃ films deposited at 290 °C on glass referenced to the spectrum of the uncoated glass substrate. The resultant average transmission of the In₂O₃ film in the wavelength range 250–800 nm was calculated using the relationship $A = 2 - \log_{10}(\%T)$, where A is absorbance and T is transmittance of light. The spectra in Fig. 5 shows the optical transmittance for the In₂O₃ 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 In₂O₃ films is comparable to values already reported in literature.^{23,44} With uncoated glass being the control, it would be $71/74 \times 100\%$ 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 In₂O₃ 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.^{41,45}

Tauc plots of the In₂O₃ films deposited with different H₂O pulse lengths are shown in Fig. 6, indicating that the direct optical band gap varies between 4.0 and 4.2 eV with higher band gap for shorter H₂O 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 deposition temperatures used in our study, which are higher than what is typically employed in ALD of In₂O₃. 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 vacancies⁴¹ that can supply free electrons⁴² 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.⁴⁶ This increase of

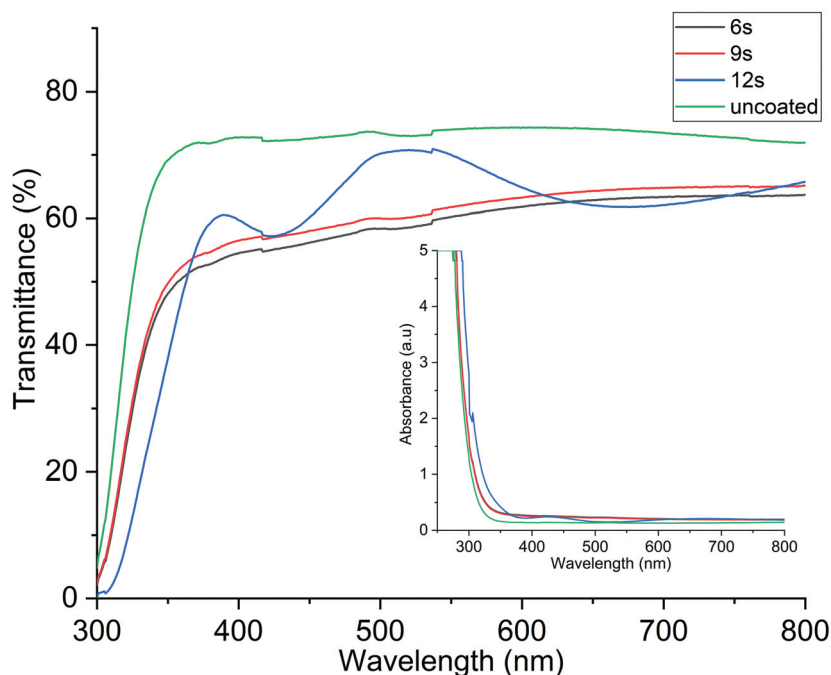


Fig. 5 Optical transmittance of the In₂O₃ films deposited with different lengths of the H₂O pulse. The transmittance increases with increasing H₂O pulse time.



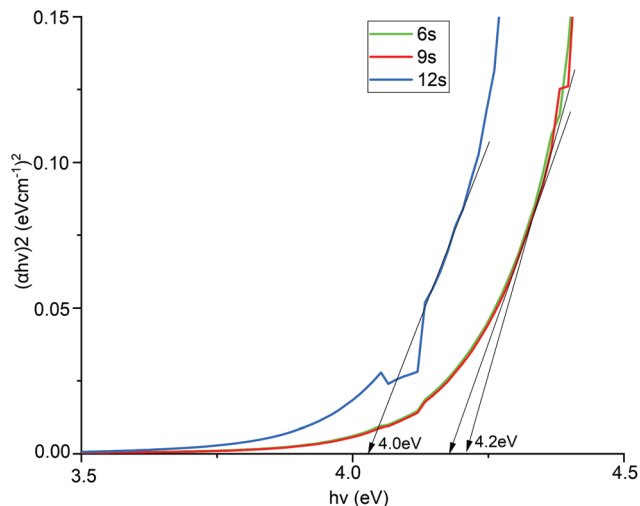


Fig. 6 Tauc plots for estimating the optical band gaps of In_2O_3 films deposited with different H_2O pulse times.

band gap can be further explained by the gap widening phenomenon called the Burstein–Moss shift.⁴⁷

C. Comparison to the $\text{In}(\text{famd})_3$ precursor (1)

ALD using **4** and H_2O renders In_2O_3 films with properties comparable to previous studies using **1**, albeit **4** seems to require higher deposition temperatures. A temperature window of 150–275 °C for In_2O_3 films using **1** and H_2O was reported, but slow surface kinetics of **1** were observed.³¹ This was then circumvented using a stop-flow process to give **1** more time to react with the surface. We could not utilize a stop-flow process in our ALD reactor, therefore undertook a direct comparison between **1** and **4** by studying **1** and H_2O in our reactor. We

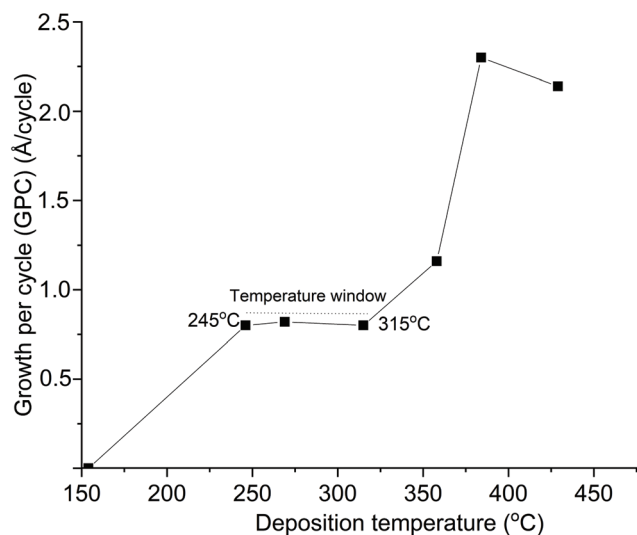


Fig. 7 Growth per cycle vs. deposition temperature for ALD of In_2O_3 using **1** and H_2O in our ALD reactor, which does not have stop flow mode.

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_2O (Fig. 7). This is higher than the 150–275 °C reported for **1**, again showing its slow surface kinetics.

A stop flow mode was used to allow for a lower deposition temperature of In_2O_3 using **1**, whilst we employed a multiple-pulse approach using **1** for depositing InN by ALD.³⁷ The higher temperature interval found for ALD of In_2O_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. 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 $\text{In}(\text{III})$ triazenide precursor in combination with H_2O to deposit In_2O_3 films in a thermal ALD process. Polycrystalline In_2O_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 mΩ cm. A temperature interval with self-limiting growth was found between 270–385 °C. This is higher than that for ALD of In_2O_3 using the previously reported **1**. 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.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was financially supported by the Swedish Foundation for Strategic Research through the project “Time-resolved low temperature CVD for III-nitrides” (no. SSF-RMA 15-0018). PM acknowledges the Swedish Institute for funding his master program studies at Linköping University.

Notes and references

- O. Bierwagen, *Semicond. Sci. Technol.*, 2015, **30**, 24001.
- M. F. Bhopal, D. won Lee, A. Rehman and S. H. Lee, *Vacuum*, 2016, **133**, 108–113.
- N. Balasubramanian and A. Subrahmanyam, *J. Phys. D: Appl. Phys.*, 1989, **22**, 206–209.
- H. Kim, C. M. Gilmore, A. Piqué, J. S. Horwitz, H. Mattoussi, H. Murata, Z. H. Kafafi and D. B. Chrisey, *J. Appl. Phys.*, 1999, **86**, 6451–6461.



- 5 M. Mazur, D. Kaczmarek, J. Domaradzki, D. Wojcieszak, S. Song and F. Placido, *Conf. Proc. - 8th Int. Conf. Adv. Semicond. Devices Microsystems, ASDAM 2010*, 2010, pp. 65–68.
- 6 Y. Liu, W. Ren, P. Shi, D. Liu, M. Liu, W. Jing, B. Tian, Z. Ye and Z. Jiang, *AIP Adv.*, 2017, **7**, 1–8.
- 7 P. Nath and R. F. Bunshah, *Thin Solid Films*, 1980, **69**, 63–68.
- 8 M. R. Karim, Z. Feng, J. M. Johnson, M. Zhu, J. Hwang and H. Zhao, *Cryst. Growth Des.*, 2019, **19**, 1965–1972.
- 9 J. H. Han, B. K. Park and T. M. Chung, *Ceram. Int.*, 2020, **46**, 3139–3143.
- 10 Q. Ma, H. M. Zheng, Y. Shao, B. Zhu, W. J. Liu, S. J. Ding and D. W. Zhang, *Nanoscale Res. Lett.*, 2018, **13**, 4, 0–8.
- 11 M. Ritala, *Electrochem. Solid-State Lett.*, 1999, **1**, 156.
- 12 J. W. Elam, A. B. F. Martinson, M. J. Pellin and J. T. Hupp, *Chem. Mater.*, 2006, **18**, 3571–3578.
- 13 M. Ritala, T. Asikainen, M. Leskelä and J. Skarp, *MRS Online Proc. Libr.*, 1996, **426**, 513–518.
- 14 T. Asikainen, M. Ritala and M. Leskelä, *J. Electrochem. Soc.*, 1995, **142**, 3538–3541.
- 15 A. W. Ott, J. M. Johnson, J. W. Klaus and S. M. George, *Appl. Surf. Sci.*, 1997, **112**, 205–215.
- 16 D. J. Lee, J. Y. Kwon, J. Il Lee and K. B. Kim, *J. Phys. Chem. C*, 2011, **115**, 15384–15389.
- 17 A. U. Mane, A. J. Allen, R. K. Kanjolia and J. W. Elam, *J. Phys. Chem. C*, 2016, **120**, 9874–9883.
- 18 J. W. Elam, J. A. Libera and J. N. Hryn, *ECS Trans.*, 2019, **41**, 147–155.
- 19 J. A. Libera, J. N. Hryn and J. W. Elam, *Chem. Mater.*, 2011, **23**, 2150–2158.
- 20 B. Macco, H. C. M. Knoop and W. M. M. Kessels, *ACS Appl. Mater. Interfaces*, 2015, **7**, 16723–16729.
- 21 Y. Wu, B. Macco, D. Vanhemel, S. Kölling, M. A. Verheijen, P. M. Koenraad, W. M. M. Kessels and F. Roozeboom, *ACS Appl. Mater. Interfaces*, 2017, **9**, 592–601.
- 22 Q. Ma, Y. Shao, Y. P. Wang, H. M. Zheng, B. Zhu, W. J. Liu, S. J. Ding and D. W. Zhang, *IEEE Electron Device Lett.*, 2018, **39**, 1672–1675.
- 23 O. Nilsen, R. Balasundaraprabhu, E. V. Monakhov, N. Muthukumarasamy, H. Fjellvåg and B. G. Svensson, *Thin Solid Films*, 2009, **517**, 6320–6322.
- 24 R. K. Ramachandran, J. Dendooven, H. Poelman and C. Detavernier, *J. Phys. Chem. C*, 2015, **119**, 11786–11791.
- 25 M. Gebhard, M. Hellwig, H. Parala, K. Xu, M. Winter and A. Devi, *Dalton Trans.*, 2014, **43**, 937–940.
- 26 W. J. Maeng, D. W. Choi, K. B. Chung, W. Koh, G. Y. Kim, S. Y. Choi and J. S. Park, *ACS Appl. Mater. Interfaces*, 2014, **6**, 17481–17488.
- 27 W. J. Maeng, D. W. Choi, J. Park and J. S. Park, *J. Alloys Compd.*, 2015, **649**, 216–221.
- 28 J. Sheng, D. W. Choi, S. H. Lee, J. Park and J. S. Park, *J. Mater. Chem. C*, 2016, **4**, 7571–7576.
- 29 J. H. Han, E. A. Jung, H. Y. Kim, D. H. Kim, B. K. Park, J. S. Park, S. U. Son and T. M. Chung, *Appl. Surf. Sci.*, 2016, **383**, 1–8.
- 30 H. Y. Kim, E. A. Jung, G. Mun, R. E. Agbenyeke, B. K. Park, J.-S. Park, S. U. Son, D. J. Jeon, S.-H. K. Park, T.-M. Chung and J. H. Han, *ACS Appl. Mater. Interfaces*, 2016, **8**, 26924–26931.
- 31 S. B. Kim, A. Jayaraman, D. Chua, L. M. Davis, S. L. Zheng, X. Zhao, S. Lee and R. G. Gordon, *Chem. – Eur. J.*, 2018, **24**, 9525–9529.
- 32 J. H. Lee, J. Sheng, H. An, T. H. Hong, H. Y. Kim, H. K. Lee, J. H. Seok, J. W. Park, J. H. Lim and J. S. Park, *Chem. Mater.*, 2020, **32**, 7397–7403.
- 33 S.-H. Choi, H.-J. Jeong, T. Hong, Y. H. Na, C. K. Park, M. Y. Lim, S. H. Jeong, J. H. Lim and J.-S. Park, *J. Vac. Sci. Technol., A*, 2021, **39**, 032406.
- 34 M. J. Zhao, Z. X. Zhang, C. H. Hsu, X. Y. Zhang, W. Y. Wu, S. Y. Lien and W. Z. Zhu, *Nanomaterials*, 2021, **11**, 978.
- 35 M. Gebhard, M. Hellwig, A. Kroll, D. Rogalla, M. Winter, B. Mallick, A. Ludwig, M. Wiesing, A. D. Wieck, G. Grundmeier and A. Devi, *Dalton Trans.*, 2017, **46**, 10220–10231.
- 36 S. T. Barry, P. G. Gordon, M. J. Ward, M. J. Heikkilä, W. H. Monillas, G. P. A. Yap, M. Ritala and M. Leskelä, *Dalton Trans.*, 2011, **40**, 9425–9430.
- 37 P. Rouf, N. J. O'Brien, K. Rönnyby, R. Samii, I. G. Ivanov, L. Ojamaä and H. Pedersen, *J. Phys. Chem. C*, 2019, **123**, 25691–25700.
- 38 N. J. O'Brien, P. Rouf, R. Samii, K. Rönnyby, S. C. Buttera, C. W. Hsu, I. G. Ivanov, V. Kessler, L. Ojamaä and H. Pedersen, *Chem. Mater.*, 2020, **32**, 4481–4489.
- 39 G. Pacchioni, *Solid State Sci.*, 2000, **2**, 161–179.
- 40 F. Gunkel, D. V. Christensen, Y. Z. Chen and N. Pryds, *Appl. Phys. Lett.*, 2020, **116**, 120505.
- 41 R. E. Agbenyeke, E. A. Jung, B. K. Park, T. M. Chung, C. G. Kim and J. H. Han, *Appl. Surf. Sci.*, 2017, **419**, 758–763.
- 42 Q. Ma, H.-M. Zheng, Y. Shao, B. Zhu, W.-J. Liu, S.-J. Ding and D. W. Zhang, *Nanoscale Res. Lett.*, 2018, **13**, 4.
- 43 P. Karthik, V. Vinesh, A. R. Mahammed Shaheer and B. Neppolian, *Appl. Catal., A*, 2019, **585**, 117208.
- 44 M. Thirumoorthi and J. Thomas Joseph Prakash, *J. Asian Ceram. Soc.*, 2016, **4**, 124–132.
- 45 T. Asikainen, M. Ritala and M. Leskelä, *Appl. Surf. Sci.*, 1994, **82–83**, 122–125.
- 46 H. R. Fallah, M. Ghasemi and A. Hassanzadeh, *Phys. E*, 2007, **39**, 69–74.
- 47 K. G. Saw, N. M. Aznan, F. K. Yam, S. S. Ng and S. Y. Pung, *PLoS One*, 2015, **10**, e0141180.

