Rapid Synthesis of SnO$_2$ Thin Films using Monoethanolamine through Wet Chemical Route

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Abstract

Objectives: To develop a unique wet chemical process for the rapid synthesis of SnO$_2$ thin films and hence to study the structural, morphological, optical and electrical properties of the films. Methods/Analysis: Polycrystalline SnO$_2$ thin films having a thickness in the range of 800-1000 nm with crystallite size less than 5 nm were synthesized within a time of 20 minutes and without the need of post-annealing using SILAR technique. Lattice parameters, c/a ratio, cell volume, dislocation density, refractive index, extinction coefficient and porosity of the SnO$_2$ thin films were determined. Findings: Films have a coarse and porous surface morphology with very fine pores distributed nearly uniformly on the film surface. The crystallite size and strain developed in the SnO$_2$ films were computed by the Williamson-Hall technique. Microstrain developed in the films is of the order of 10$^{-3}$. SnO$_2$ films exhibit nearly 70% transmittance in the visible region. The optical band gap of the SnO$_2$ films is 3.65 eV. The refractive index of the films varies from 2.05 to 2.30 in the 450-1900 nm wavelength range. SnO$_2$ films exhibit a resistivity in the order of 10$^{-1}$ Ω cm. Novelty: Reports a unique rapid wet chemical process for the direct preparation of crystalline SnO$_2$ thin films using monoethanolamine at a temperature of 80°C. Literature survey reveals that no other method has yielded such type of good quality SnO$_2$ thin films at this temperature and without post annealing or sintering. First time crystalline SnO$_2$ films were prepared by a wet chemical process within 20 minutes.

Keywords: SnO$_2$ films; SILAR; Monoethanolamine; Strain; Optical band gap

1 Introduction

Wide band gap (3.6 eV) semiconductor tin oxide (SnO$_2$) is a promising material for various technological applications due to its high optical transparency in the visible range of the electromagnetic spectrum with excellent electric conductivity and high chemical stability$^{(1-7)}$. SnO$_2$ exhibits high conductivity due to intrinsic defects like oxygen deficiencies whereas in the stoichiometric form it behaves like a good insulator. The electrical resistivity of undoped SnO$_2$ is limited to 10$^{-2}$-10$^{-3}$ Ω cm and the
conductivity of the films can be enhanced to a great extent by dopants. SnO$_2$ films exhibit good field emission properties. SnO$_2$ can be used as hole blocking layer for perovskite solar cells. SnO$_2$ has been frequently used as resistive-type CO and H$_2$ gas sensors. The oxygen species with trapped negative charges existing in SnO$_2$ usually prompts an upward band bending and are hence promising sensing materials. In TCO applications, both the surface and interface properties of SnO$_2$ plays a vital role. The films have also been employed in a wide range of applications like oxidation catalyst, liquid crystal displays, light emitting diodes, thin film resistors, dye-sensitized solar cells, n-type ohmic electrodes for ink-jet printable electronics, photo-detectors.

SnO$_2$ thin films can be prepared by a variety of depositing processes, such as chemical vapour deposition, sputtering, spin coating, thermal evaporation, Pulsed laser deposition, thermal oxidation, spray pyrolysis, sol-gel method and Dr. Blade method. These methods require sophisticated complex setups and are time consuming. Compared to these deposition methods, the successive ionic layer adsorption and reaction (SILAR) technique is a facile, environment-friendly and low-cost method, as well as appropriate for large area deposition of thin films with closely 100% material exploitation.

Moreover, SILAR opens up the possibility to control the particle size within the nanometer range and to vary the film morphology by varying the different parameters associated with it. On the other hand, there is little given in the literature about the preparation of SnO$_2$ films by a wet chemical process SILAR. Doping is relatively easy in this technique. Fabrication of good quality SnO$_2$ thin films at low processing temperatures is certainly a challenge.

In this work, we report a unique process for the rapid synthesis of adherent, dense and transparent SnO$_2$ thin films by SILAR on glass substrates using the cost-effective precursor tin (II) chloride (SnCl$_2$) and the complexing agent monoethanolamine (MEA). MEA (H$_2$NCH$_2$CH$_2$OH) used in this work is a clear, water-white, hygroscopic liquid with a mild ammoniacal odour and having a pH of 11.8. MEA undergoes fair biodegradation and is not expected to be persistent in the environment. Hence, MEA can be treated as an environment-friendly complexing agent.

Fabrication of polycrystalline SnO$_2$ thin films within a time of 20 minutes at a low processing temperature of 353 K was achieved without the need for post-annealing or sintering. Films having a thickness in the range of 800-1000 nm with crystallite size less than 5 nm were deposited. Structural, morphological, optical and electrical properties were investigated to illustrate the quality of the films.

2 Materials and Methods

Analytical grade chemicals (Merck) were used as received. The details of successive ionic layer adsorption and reaction (SILAR) have been described elsewhere. In this work, immersion time in cationic and anionic solution was optimized to 4 s. The optimized immersion time for rinsing the substrates was 8 s.

SnO$_2$ thin films were synthesized from the cationic precursor prepared by dissolving 1 mg of SnCl$_2$ in distilled water, followed by the addition of 3 ml hydrochloric acid (35%) and 2 ml MEA. The cationic precursor solution was kept at room temperature. 2 ml hydrogen peroxide (30%) mixed distilled water kept at 353 K served as an anionic precursor. Magnetic stirring was performed for thorough mixing while each of the chemical reagents was added to prepare the precursor solutions.

Films were synthesized by choosing 80 deposition cycles to get highly adherent films. Accordingly, the deposition time was almost around 20 minute. After deposition, the samples were rinsed in distilled water and dried under hot airflow before characterization.

The crystalline properties of the as-prepared and annealed SnO$_2$ films were examined by X-ray diffractometer Bruker AXS-8 advance and the surface morphology by JEOL Model JSM-6490LV microscope. The energy-dispersive X-ray (EDAX) spectroscopic measurements were done by the JEOL Model JED-2300 spectrometer, which was coupled with the scanning electron microscope. Optical spectra were recorded using a Hitachi-U-3410 UV-Vis-NIR spectrophotometer. The thickness of the films was determined by a gravimetric method using an analytical balance (OHAUS Model PA64C) of readability 0.1 mg. Electrical studies were carried out by the two probe method with electrodes made from silver paste using the Keithley source measure unit (Model SMU Keithley 2400).

3 Results and Discussion

3.1 Effect of MEA and film formation mechanism

SnCl$_2$·2H$_2$O used was dissolved in distilled water and followed by the addition of HCl acid solution and thereby adequate amounts of Sn$^{2+}$ ions were made available in the solution. Like triethanolamine, when MEA was added, it complexes with Sn$^{2+}$ to form a chelate compound [Sn(MEA)$_2$]$_2^{2+}$. MEA wraps around the Sn$^{2+}$ ions and averts the ions from further chemical changes or early release into the solution.
During the first SILAR process, on immersing the substrate in the cationic solution the \([\text{Sn(MEA)}_k]^2^+\) complex was adsorbed on it. On the second process, i.e. when rinsing the substrate in distilled water kept at room temperature, \([\text{Sn(MEA)}_k]^2^+\) on the substrate reacts with \(\text{H}_2\text{O}\) and forms tin hydroxide \((\text{Sn(OH)}_2)\). Excess and loosely held species on the substrate were removed during this process. In the third process, the strong oxidizer \(\text{H}_2\text{O}_2\) present in the distilled water kept at 353 K was oxidizing \((\text{Sn(OH)}_2)\) into \(\text{SnO}_2\) and the excess \(\text{H}_2\text{O}_2\) was simultaneously removed at this temperature on exposure to air. In the last immersion process, loosely held chemical species were removed from the film surface. Rapid deposition of \(\text{SnO}_2\) films was achieved within 20 minute, which is a noticeable property of the unique MEA bath used here. This was well supported by a report of the rapid deposition of \(\text{ZnO}\) films from a chemical bath containing MEA\(^{(10)}\). \(\text{SnO}_2\) film formation reaction may be as follows:

\[
\text{SnCl}_2 + k(\text{MEA}) \leftrightarrow [\text{Sn(MEA)}_k]^2^+ + 2\text{Cl}^{-}
\]

\[
[\text{Sn(MEA)}_k]^2^+ + 2\text{OH}^- \rightarrow \text{Sn(OH)}_2 + k(\text{MEA})
\]

\[
\text{Sn(OH)}_2 + \text{H}_2\text{O}_2 \rightarrow \text{SnO}_2 + 2\text{H}_2\text{O}
\]

### 3.2 Structural and morphological analysis

The X-ray diffraction (XRD) profile of the \(\text{SnO}_2\) thin films is depicted in [Figure 1](#). These films were not treated after deposition. High-intensity XRD peaks at 26.58°, 33.87°, 37.95°, 51.77° and 65.96° corresponding to the planes \((110), (101), (200), (211)\) and \((301)\) respectively show that the films are polycrystalline. All the peaks can be indexed to tetragonal structured \(\text{SnO}_2\), which are in good agreement with the JCPDS data file (File No. 71-0652). The XRD analysis detected no other impurity peaks. Literature study reveals that no other wet chemical method had yielded such type of crystalline films at a low processing temperature of 353 K without extensive post-annealing or sintering\(^{(1,7)}\). Moreover, crystalline quality of the films are in par with the films prepared by other sophisticated methods at high processing temperature\(^{(2,3,6)}\).

For estimation of the crystallite size ‘\(D\)’ of the films, the Scherrer formula was used:

\[
D = \frac{k\lambda}{\beta\cos\theta}
\]

Where ‘\(k\)’ is the shape factor which is 0.94, ‘\(\lambda\)’ is the wavelength of X-rays which is 1.5406 nm for CuK\(_\alpha\), ‘\(\beta\)’ is the full width at half maximum of diffraction peak measured in radians and ‘\(\theta\)’ is the Bragg’s angle. The grain size estimated from the main peak \((110)\) is 4.7 nm and the value is in tune with the reported values obtained from other wet chemical methods\(^{(4,11)}\). The reported crystallite size varies between 4.796 nm to 11.563 nm in the case of films prepared by Spray Pyrolysis\(^{(2)}\). When the grain size of the films drops under 10 nm, accordingly, the size effect in electronic, optical and sensing properties can be anticipated\(^{(17)}\).

![Fig 1. XRD patterns of the SnO2 films](https://www.indjst.org/1567)
The lattice parameters of the samples were calculated using the observed values of 2θ and d-values for the tetragonal structure using the equation,

\[
\frac{1}{d^2} = \left( \frac{h^2 + k^2}{a^2} \right) + \frac{l^2}{c^2}
\]  

(5)

The primitive cell volume (\(V_c\)) was calculated using the equation,

\[
V_c = a^2 c
\]  

(6)

The obtained value of lattice parameters \(a = 0.4683\) nm and \(c = 0.3124\) nm matches with the ideal values \(a = 0.4738\) nm and \(c = 0.3186\) nm. The primitive cell volume computed is \(0.0685\) nm\(^3\) and is slightly less than that of the ideal value \(0.0715\) nm\(^3\) \((17)\). The crystallite size and strain developed in the SnO\(_2\) films were further determined by the technique developed by the Williamson and Hall (W-H plot) \((18)\). The W-H equation is given as Eq. (7).

\[
\beta \cos \theta = \frac{k\lambda}{D} + 4\varepsilon \sin \theta
\]  

(7)

Where \(\varepsilon\) is the strain developed in the films. W-H plot for the films was constructed by plotting \(\beta \cos \theta\) vs \(4 \sin \theta\) and is shown in Figure 2. The crystallite size and strain were determined from the intercept at the \(\beta \cos \theta\)-axis and the slope of the W-H plot respectively according to Eq. (7). The microstrain extracted from the W-H analysis is the non-uniform strain resulting from the systematic shifts of atoms from their ideal positions. This shift may be originating from defects like point defects, site disorder, vacancies and also due to plastic deformation \((19)\). Microstrain associated with the films was further determined using Eq. (8) directly.

\[
\varepsilon = \frac{\beta}{4 \tan \theta}
\]  

(8)

![Fig 2. W-H plot and the microstrain (inset) of the SnO\(_2\) films](https://www.indjst.org/)

The microstrain obtained from the W-H plot and Eq. (8) is plotted for comparison and is illustrated in the inset of Figure 2. The reported value of microstrain is between \(6 \times 10^{-3}\) and \(12 \times 10^{-3}\) \((20)\). The value of strain obtained is small and compressive (negative slope of W-H plot) in nature. The grain size obtained from the W-H plot is slightly less than that determined using Scherrer’s formula, but establishes lower crystallite size and hence the possibilities of the size effect \((17)\). Figure 3 shows the grain size obtained from both techniques.
The defects in the SnO$_2$ films were quantified by computing the dislocation density ($\delta$) using Williamson Smallman relation \(^{(21)}\),

$$\delta = \frac{1}{D^2} \quad (9)$$

The dislocation density of the films was determined using the grain size obtained from both Scherrer’s formula and the W-H plot. The values obtained from both techniques show no much difference. The values obtained are depicted in Table 1. The computed values of dislocation density from the values of crystallite size obtained from Scherrer formula and W-H plot are $4.9 \times 10^{15}$ lines/m$^2$ and $6.5 \times 10^{15}$ lines/m$^2$ respectively. The reported values of dislocation density varies between $7.484 \times 10^{15}$ lines/m$^2$ to $43.592 \times 10^{15}$ lines/m$^2$ \(^{(2,20)}\). The values obtained in the rapid synthesis are also in par with this. The smaller value of dislocation densities may be an indication of the crystalline nature of the SnO$_2$ films.

**Figure 4** illustrates the scanning electron microscopic (SEM) images of the SnO$_2$ films. Some aggregated structures with undefined morphologies were observed. However, the films have a coarse and porous surface morphology with very fine pores distributed nearly uniformly on the film surface. This kind of dense morphology is capable of affording a large inherent adsorptive surface area. SEM images divulge agglomerated grains, which contributes to an augmented surface-to-volume ratio, making the films potential ones in the field of gas sensing \(^{(17)}\).

The compositional purity of the SnO$_2$ films has been studied by EDAX analyses (not shown here), which indicate that the films contain mainly Sn and O, with no significant signal from other elements being detected.
3.3 Thickness of the films

Film thickness was determined by the gravimetric method. The samples were weighed before and after the film deposition to determine the mass of the films. Knowing the dimensions of the substrates used, the film thickness (t) was computed using Eq. (10).

\[ t = \frac{m}{pA} \]  

Where 'm' is the mass of the film deposited on area 'A' and 'p' is the bulk density of the material. The bulk density of the material is assumed as 6.99x10^3 kg/m^3. The thickness obtained for the films was 900 nm.

3.4 Optical properties

The optical transmittance and reflectance spectra of the samples are shown in Figure 5(a). The SnO_2 films exhibit nearly 70% transmittance in the visible region. The transmittance of the films is decreasing with an increase in wavelength in the near-infrared (NIR) region of the spectrum. Optical transmission through a rough surface is much affected by the scattering of light having wavelengths near the magnitude of the film surface features [22-24]. The reflectance spectra of the SnO_2 films in Figure 5(a) shows reflectance below 20% and exhibits a slight increase in reflectance with an increase in wavelength in the NIR region.

Fig 5. (a) Transmittance and reflectance spectra and (b) plot of (αhν)^2 versus hν of the SnO_2 films
According to inter-band absorption theory, the relation between the absorption coefficient near the threshold and the incident photon energy for allowed direct transitions is given by the following relation (25):

$$\alpha = \frac{A(h\nu - E_g)}{h\nu} \frac{1}{2}$$

(11)

Where 'A' is the probability parameter for the transition and 'E_g' is the optical band-gap energy. In Figure 5(b), ($\alpha h\nu$)$^2$ versus $h\nu$ plot of the films is shown. The value of 'E_g' was determined by extrapolating the linear region of ($\alpha h\nu$)$^2$ vs $h\nu$ plots. The 'E_g' value of the SnO$_2$ films obtained is 3.65 eV. The 'E_g' values obtained were in good agreement with the reports that SnO$_2$ films usually exhibit band gap energy in the range of 3.4 - 4.6 eV (26-28). Optical band gap of the films can be tuned by suitable doping or annealing (2,11,29).

Refractive index of the material influences the film optical characteristics such as transmission spectrum, reflection spectrum etc. The Refractive index, n, of the films was determined from the extinction coefficient 'k' and the reflectance 'R' using Eqns. (12) and (13) and is found in agreement with values reported in the literature (29-32).

$$k = \frac{\alpha \lambda}{4\pi}$$

(12)

$$n = \frac{(1 + R) + \sqrt{4R - (1 - R)^2k^2}}{(1 - R)}$$

(13)

Figure 6 shows the variation of the refractive index of the SnO$_2$ films as a function of wavelength. The values of 'n' obtained for the films vary from 2.05 to 2.30 in the 450-1900 nm wavelength range. A small gradual increase in 'n' can be noted with an increase in wavelength in the visible-NIR region and attains a steady value in the NIR region of the spectrum. The higher transmission in the visible region shown by the films may be due to the low refractive index possessed by it. The values of the refractive index obtained are in agreement with the reported values (26,31,32). The extinction coefficient of the films shows a gradual linear increase with the increase in wavelength in the visible-NIR region of the spectrum.

The porosity of the films was determined from the refractive index of the film and the refractive index ($n_s$) of SnO$_2$ ($n_s \sim 2.006$) using Eq. (14) (33).

$$P(\%) = 1 - \frac{n^2 - 1}{n_s^2 - 1}$$

(14)
As depicted in the inset of Figure 6, SnO$_2$ films possess a porosity of more than 25%. Hence, these films with excellent porous morphology ensure a large solid-to-gas interface due to the high surface-to-volume ratio making it a potential candidate for gas sensing \((2,3,34,35)\).

### 3.5 Electrical properties

The electrical behaviour of the SnO$_2$ films was examined by determining the electrical resistivity of the films. Experimental measurements were carried out by dc two-point probe method using silver electrode contacts. SnO$_2$ films exhibit a high resistivity in the order of $10^3$ Ωcm. Usually, metal oxide thin films prepared by wet chemical method SILAR exhibit high resistivity due to the rough structure and high porosity \((14,15)\). Due to the small grain size, the density of the grain boundaries in the SnO$_2$ films may be large. Grain boundaries act as traps for the free charge carriers. These grain boundaries block the transport of charge carriers. Moreover, the SEM images discussed in section 3.2 exposed that the films have a surface morphology with pores. All these factors contributed to the higher electrical resistivity in the films than the reported values of resistivity of $10^2$ Ωcm \((4)\) and $10^3-10^4$ Ωcm \((11,31)\). The prepared SnO$_2$ films are potential candidates for various thin film resistor applications, which requires high resistivity \((36)\). Literature study reveals that the film resistivity was reduced by Fluorine doping from $10^{-2}$ to $10^{-3}$ Ωcm \((4)\). The electrical conductivity of the SnO$_2$ films can be improved by doping with suitable elements like F, Nb, In, Al and Co \((7,12,24,28,31)\) or by annealing \((30,37)\).

### 4 Conclusion

A novel rapid synthesis of polycrystalline SnO$_2$ thin films was carried at a relatively low temperature 353 K within 20 minutes by SILAR technique using the complexing agent MEA. Films with crystallite size less than 5 nm exhibit a coarse and porous surface morphology. This inventive approach discards the traditional way of post-annealing or sintering to prepare crystalline SnO$_2$ films. Lattice parameters, cell volume, strain and dislocation density of the SnO$_2$ films illustrates the quality of the films. The Williamson-Hall technique was utilized for establishing crystallite size and strain of the films.

Microstrain developed in the films is of the order of $10^{-3}$. Transmittance is relatively high in the visible region and the films exhibit nearly 70% transmittance. The optical band gap of the SnO$_2$ films is 3.65 eV. The refractive index of the films varies from 2.05 to 2.30 in the 450-1900 nm wavelength range. The extinction coefficient of the films shows a gradual linear increase in the entire visible-NIR region of the spectrum. The SnO$_2$ films exhibit a resistivity in the order of $10^3$ Ωcm. The results obtained from this low-cost method are in par with the results obtained from other sophisticated methods except the electrical conductivity.

The nanostructured SnO$_2$ films are suitable for various applications such as gas sensors, thin film resistors and dye-sensitized solar cells. Modulation of the electrical and optical behaviour of the films can be explored by doping and annealing. Sensing performance of these films can be further investigated.

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### References

1. Jadhav H, Suryawanshi S, More MA, Sinha S. Pulsed laser deposition of tin oxide thin films for field emission studies. *Applied Surface Science*. 2017;419:764–769. Available from: https://dx.doi.org/10.1016/j.apsusc.2017.05.020.
2. Kamble DL, Harale NS, Patil VL, Patil PS, Kadam LD. Characterization and NO2 gas sensing properties of spray pyrolyzed SnO2 thin films. *Journal of Analytical and Applied Pyrolysis*. 2017;127:38–46. Available from: https://dx.doi.org/10.1016/j.jaap.2017.09.004.
3. Sharma B, Sharma A, Joshi M, ha Myung J. Sputtered SnO2/ZnO Heterostructures for Improved NO2 Gas Sensing Properties. *Chemosensors*. 2020;8(3):67–67. Available from: https://dx.doi.org/10.3390/chemosensors8030067.
4. P K, Vijayanarayanan D, M S, T AKR, C S, K J. Optimization of substrate temperature and characterization of tin oxide based transparent conducting thin films for application in dye-sensitized solar cells. *Thin Solid Films*. 2017;631:1–11. Available from: https://dx.doi.org/10.1016/j.tsf.2017.04.003.
5. Ponja SD, Williamson BAJD, Sathanavay S, Scanlon DO, Parke IP, Carmalt Cfl. Enhanced electrical properties of antimony doped tin oxide thin films deposited via aerosol assisted chemical vapour deposition. *Journal of Materials Chemistry C*. 2018;6(27):7257–7266. Available from: https://dx.doi.org/10.1039/c8tc01929k.
6. Tao H, Ma Z, Yang G, Wang H, Long H, Zhao H, et al. Pingli Qin,Guojia Fang. Room-temperature processed tin oxide thin film as effective hole blocking layer for planar perovskite solar cells. *Applied Surface Science*. 2018;434:1336–1343. Available from: https://dx.doi.org/10.1016/j.apsusc.2017.11.161.
7. Basyouni MA, Eker YR, Yilmaz M. Structural, optical, electrical and room temperature gas sensing characterizations of spin coated multilayer cobalt-doped tin oxide thin films. *Superlattices and Microstructures*. 2020;140:106465. Available from: https://dx.doi.org/10.1016/j.spmi.2020.106465.
8. Ingole SM, Navale ST, Navale YH, Bandgar DK, Stadler FJ, Mane RS, et al. Nanostructured tin oxide films: Physical synthesis, characterization, and gas sensing properties. *Journal of Colloid and Interface Science*. 2017;493:162–170. Available from: https://dx.doi.org/10.1016/j.jcis.2017.01.025.
9) Hunashimarad BG, Bhat JS, Raghavendra PV, Bhanjani RF. Photoluminescence in Strontium doped tin oxide thin films. *Optical Materials*. 2021;114(110962). Available from: https://dx.doi.org/10.1016/j.optmat.2021.110962.

10) Abdulla N, Ismail NA, Nuruzzaman DM. Preparation of tin oxide (SnO2) thin films using thermal oxidation. *IOP Conference Series: Materials Science and Engineering*. 2018;319(012022). Available from: https://dx.doi.org/10.1088/1757-899x/319/1/012022.

11) ELCANGOVAN E, SINGH M, RAMAMURTHI R. Studies on structural and optical properties of spray deposited SnO2:F thin films as a function of film thickness. *Materials Science and Engineering B*. 2004;112(1):143–148. Available from: https://dx.doi.org/10.1016/j.mseb.2004.03.085-x.

12) Misra M, Hwang DK, Kim YC, Myoung JM, Lee TI. Eco-friendly method of fabricating indium-tin-oxide thin films using pure aqueous sol-gel. *Ceramics International*. 2018;44(3):2927–2933. Available from: https://dx.doi.org/10.1016/j.ceramint.2017.11.041.

13) Asadzadeh M, Tajabadi F, Dastan D, Sangpour P, Shi Z, Taghavinia N. Facile deposition of porous fluoride doped tin oxide by Dr. Blade method for capacitive applications. *Ceramics International*. 2021;47(4):5487–5494. Available from: https://dx.doi.org/10.1016/j.ceramint.2020.10.131.

14) Tolstoy VP. The porosity route of the successive ioni2er layer deposition procedure for synthesizing nanolayers of metal oxides, hydroxides and peroxides. *Thin Solid Films*. 1997;307(1-2):10–13. Available from: https://dx.doi.org/10.1016/s0040-6090(97)00256-3.

15) Pathan HM, Lokhande CD. Deposition of metal chalcogenide thin films by successive ioni2er layer adsorption and reaction (SILAR) method. *Bulletin of Materials Science*. 2004;27(2):85–111. Available from: https://dx.doi.org/10.1007/bf82708491.

16) Shariffuddin SS, Herman SH, Rusop M. Layer-by-Layer Nanoparticles ZnO Thin Films Prepared by Sol-Gel Method. *Advanced Materials Research*. 2011;403(408):1178–1182. Available from: https://dx.doi.org/10.4028/www.scientific.net/amr.403-408.1178.

17) Nowak P, Maziarz W, Rydosz A, Kowalski K, Ziajka M, Zakrzewska K. SnO2/TiO2 Thin Film n-n Heterostructures of Improved Sensitivity to NO2. *Sensors*. 2020;20(23). Available from: https://dx.doi.org/10.3390/s20236830.

18) Williamson GK, Hall WH. X-ray line broadening from filed aluminium and wolfram. *Acta Metallurgica*. 1953;11(1):22–31. Available from: https://dx.doi.org/10.1016/0001-6160(53)90006-6.

19) Fell CR, Chi M, Meng YS, Jones JL. Ion induced effects. *Ceramic Society of Japan*. 2009;25(1):221–226. Available from: https://dx.doi.org/10.2525/csj.2008.2708044.2018.1477559.

20) Deyu GK, Muñoz-Rojas D, Rapenne L, Deschanvres JL, Klein A, Jiménez C, et al. SnO2 Films Deposited by Ultrasonic Spray Pyrolysis: Influence of Al Incorporation on the Properties. *Molecules*. 2019;24(15). Available from: https://dx.doi.org/10.3390/molecules24152797.

21) Khadraoui M, Benramdane N, Mathieu C, Bouzidi A, Milora R, Kebbab Z. Optical and electrical properties of thin films grown by spray pyrolysis. *Solid State Communications*. 2010;150(5-6):297–300. Available from: https://dx.doi.org/10.1016/j.ssc.2009.09.032.

22) Xiao J, Nie L, Qin J, Kang T, Li C, Xie J. Effect of oxygen stoichiometry on the structure, optical and epitaxy-near-zero properties of indium tin oxide films. *Optics Express*. 2019;27(20). Available from: https://dx.doi.org/10.1364/oe.27.028618.

23) Czapa A, Kusior E, Bucko M. Optical properties of non-stoichiometric tin oxide films obtained by reactive sputtering. *Thin Solid Films*. 1989;182(1-2):15–22. Available from: https://dx.doi.org/10.1016/0040-6090(89)90239-3.

24) Wang L, Yu J, Niu X, Wang L, Fu C, Qiu R, et al. Effect of F and Nb co-doping on structural, electrical and optical properties of spray deposited tin oxide thin films. *Thin Solid Films*. 2018;649:147–153. Available from: https://dx.doi.org/10.1016/j.tsf.2018.01.035.

25) Vinodkumar R, Lethy KJ, Arunkumar PR, Krishnan RR, Pillai NV, Pillai VPM, et al. Effect of cadmium oxide incorporation on the microstructural and optical properties of pulsed laser deposited nanostructured zinc oxide thin films. *Materials Chemistry and Physics*. 2010;121(3):406–413. Available from: https://dx.doi.org/10.1016/j.matchemphys.2010.01.004.

26) Woo SW, Seo HB, Choi J, Bae BS, Yun EJ. Effects of the Process Parameters on the Properties of Sputter-Deposited Tin Oxide Thin Films. *Journal of Nanoscience and Nanotechnology*. 2019;19(1):1301–1307. Available from: https://dx.doi.org/10.1166/jnn.2019.16241.

27) Ching-Prado E, Watson A, Miranda H. Optical and electrical properties of fluoride doped tin oxide thin film. *Journal of Materials Science: Materials in Electronics*. 2018;29(18):15299–15306. Available from: https://dx.doi.org/10.1007/s10854-018-8979-8.

28) Maheswari S, Karunakaran M, Chandrasekar LB, Kasirajan K. Ammonia sensors on the base of gadolinium doped tin oxide thin films. *Thin Solid Films*. 2019;602. Available from: https://dx.doi.org/10.1016/j.matchemphys.2020.110962.

29) Yoldas BE, Partlow DP. Formation of broad band antireflective coatings on fused silica for high power laser applications. *Thin Solid Films*. 1985;129(2-1):14. Available from: https://dx.doi.org/10.1016/0040-6090(85)90008-6.

30) Kaur M, Dadhich BK, Singh R, Ganapathi SK, Bagwaiya T, Bhattacharya S, et al. RF sputtered SnO2:NiO thin films as sub-ppm H2S sensor operable at room temperature. *Sensors and Actuators B: Chemical*. 2021;292:389–403. Available from: https://dx.doi.org/10.1016/j.snb.2021.116544.

31) Bhownick T, Ambarekar V, Ghosh A, Dewan M, Bandyopadhyay PP, Nag S. Multilayered and Chemiresistive Thin and Thick Film Gas Sensors for Air Quality Monitoring. In: Multilayer Thin Films - Versatile Applications for Materials Engineering. Partha Pratim Bandyopadhyay. 2020. doi:10.5772/intechopen.89710.

32) Park KW, Hur SG, Ahn JK, Seong NJ, Yoon SG. High-Resistance Thin-Film Resistors Grown Using CrB2-Si-SiC Materials by Radio-Frequency Magnetron Sputtering. *IEEE Transactions on Electron Devices*. 2010;57(6):1475–1480. Available from: https://dx.doi.org/10.1109/TED.2010.2045673.

33) Djamil R, Aicha K, Souiff A, Fayçal D. Effect of annealing time on the performance of tin oxide thin films ultraviolet photodetectors. *Thin Solid Films*. 2017;623:1–7. Available from: https://dx.doi.org/10.1016/j.tsf.2016.12.035.