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# ANALYSIS ON SYNTHESIS, CHARACTERIZATION AND GAS SENSING PROPERTIES OF BINARY OXIDE In2O3:MoO3 THIN FILMS

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Abstract: In 2O3 doped In2O3:MoO3 binary oxide thin films were prepared by using spray pyrolysis technique on glass substrate for various concentrations at 4000C temperature. The effect of In2O3 On structural, morphological and electrical properties has been studied. Thin films were characterized using X-ray diffraction (XRD), SEM and Electrical properties. The obtained film shows good crystalline in nature. The average grain size estimated is 11 nm the other properties were under consideration the sensitivity and selectivity of sensor can be improved by dopants or additives which can change the gas sensing characteristics. Nano composite term contains mixture of two or more nano oxide materials like binary oxide. Nano composite films consists of Nano crystalline or amorphous phase of a least two different materials In2O3 as dopant and MoO3 as a functional material in film. The precursor InCl3 and MoCl5 of concentrations 0.2N:0.1N.

*Keywords:* XRD, SEM EDS, Electrical, Gas sensor, spray pyrolysis technique, binary oxide thin films, In2O3, MoO3, Thin film.

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#### **I INTRODUCTION**

Nanocomposite is mixture of two or more nano oxide materials like binary oxide, ternary oxide . Indium oxide (In2O3) has been identified as the best possible gas sensing material because of its good conductivity with high stability, and is a semiconduction-type n-chemiresistive sensor material with a straight-band gap of "3.55-3.75 eV". Indium oxide was produced as nanostructure, nanosphere, nanoporous and nanoporous, Attractive sensors for different gases, including a variety of nitrogen oxides, methane, ammonia, acetone and ethanol. In2O3 sensing materials can cause dopant element Changes in crystalline structure and size of grain, levels of impurity and surface defects that can significantly enhance In2O3 gas sensing performance. Molybdenum oxide is a wide band gap n-type of semiconductor with a band gap energy of 2.39-2.9 eV. Among the insoluble transition metal oxides, MoO3 has the highest work function value. MoO3 nanoparticles are different from those in the bulk due to their unique physical and chemical properties, especially due to their high surface

area to volume ratio, so they have attracted a lot of attention. Among transition metal oxides, MoO3 was found to be a suitable candidate for many technical applications, such as electrochromic display devices, gas sensors, optical storage, and lithium batteries.

#### **II EXPERIMENTAL METHODS**

Binary oxide In2O3: MoO3 thin films were prepared by spray pyrolysis technique on glass substrate. The spray Pyrolysis process was carried out at substrate temperature 400<sup>o</sup>C.The precursor InCl3 and MoCl5 of concentrations 0.1N, 0.2N and 0.3N were used, designed and assembled in laboratory to overcome limitations of conventionally designed setup; such as number of optimized parameters ,reliability and homogeneity of the deposited films. The thin films of In2O3: MoO3 were prepared for concentration in proportion of 0.2N:0.1N. The study of characteristics such as SEM, EDS,XRD, resistivity, activation energy, TCR and gas sensing property were done to study the changes due to dopant.

# **III RESULTS AND DISCUSSION**

### X-RAY DIFFRACTION ANALYSIS (XRD)

Figure shows XRD pattern of binary oxide In<sub>2</sub>O<sub>3</sub>: MoO<sub>3</sub> thin films on glass substrate fired at  $400^{\circ}$ C of the peak position values of the<sub>2</sub>O-80<sup> $\circ$ </sup> range using X powder 12(CuK $\alpha$ )Radiation.



**Figure 3.1** XRD of Binary oxide In2O3: MoO3 thin film with concentration 0.2N:0.1N

XRD of Binary oxide In2O3: MoO3 thin film with concentration 0.2N:0.1N is shown in table

Plane (hkl)	20	d-spacing	Intensity	I/Io	FWHM
In- 211	21.72	4.09424	801	67.3	1.912
Mo-211	23.01	3.86105	1190	100	0.812
Mo-211	24.74	3.5955	579	48.7	4.285
In- 310	27.21	3.27363	1020	85.7	0.808
In- 321	33.60	2.66577	796	56.2	3.399

The average grain size was determined by using Debye-Scherer formula,

$$D = 0.9\lambda / \beta \cos\theta$$

" $\beta$  is full angular width of diffraction peak at half maximum peak intensity,  $\lambda$  is wavelength of X-radiation."

As per structural analysis the grain size were calculated by using Scherrer formula. The grain size of film at concentrations 0.2N:0.1N, were found 11 nm.

# SCANNING ELECTRON MICROSCOPY (SEM)

Figure shows that the SEM of binary oxide In2O3:MoO3 thin films of 0.2N: 0.1N was deposited on glass substrate using a Spray Pyrolysis Technique and fired at 4000C. The magnifications of all SEM images are taken at 10000X.



**Figure 3.2** SEM of Binary oxide In2O3: MoO3 thin with concentration 0.2N:0.1N

Binary oxide In2O3: MoO3Films prepared by Spray Pyrolysis were observed to be non porous as per SEM analysis. As per SEM analysis, the average particle size of film was calculated by using image j software. The average particle size of film at concentrations 0.2N:0.1N was found as 198 nm.

The specific surface area of Binary oxide In2O3: MoO3thin film was calculated using BET method for spherical particles using the equation,

$$S_{w} = \frac{6}{\rho d}$$

"Where, *d* is the diameter of the particles,  $\rho$  is the density of the particles."

The specific Surface area with different concentrations of binary oxide In2O3: MoO3 was found as  $26.49002 \text{ m}^2/\text{g}$ .

# IV ENERGY DISPERSIVE X-RAY ANALYSIS (EDS)

Figure shows that the count (along Y- axis) Verses KeV (along X-axis) EDS of 0.2N:0.1N concentration of binary oxide In2O3: MoO3 thin films.



Figure 3.3 EDS of Binary oxide In2O3: MoO3 thin film with concentration 0.2N:0.1N

From the EDAX spectra, it is found that mass% and at. wt.% of In, Mo and O is nearly matched.

EDS of Binary oxide In<sub>2</sub>O<sub>3</sub>: MoO<sub>3</sub> thin film with Concentration 0.2N:0.1N is shown in table

Element	Atomic %
0	85.48
Мо	11.44
In	3.08

#### ELECTRICAL CHARACTERIZATION

The electrical characterization was done to measure the variation in electrical resistance at operating temperatures in air atmosphere, the resistivity, TCR and activation energy.

#### RESISTIVITY

The DC resistance of In2O3:MoO3 thin films with normality 0.2N:0.1N on glass substrate and fired at 400<sup>o</sup>C was measured by using half bridge method as a function of temperature. Figure shows resistance variation of In2O3:MoO3thin films with normality 0.2N:0.1N temperature variation in an atmosphere. There is decrease in resistance with increase in temperature indicating semiconductor behavior, obeying R= R0e<sup>- $\Delta E/KT$ </sup> in the temperature range of 40-350<sup>o</sup>C.

The resistance In2O3:MoO3thin films with normality 0.2N:0.1N falls rapidly, decreases linearly up to certain transition temperature and after resistance decreases exponentially with increase in temperature and lastly saturates to steady level.

The resistivity of In2O3:MoO3 thin films at constant temperature is calculated using the relation,

$$\rho = (R \ge A) / l$$
$$\rho = (R \ge b \ge t) / l$$

Where,  $R = \text{Resistance of In}_{2O3}$  :MoO3 thin film with constant temperature

ohm-m

"*t* = thickness of the film sample"

"l =length of the thin film"

"b = breadth of the thin film"





The resistivity of binary oxide In<sub>2</sub>O<sub>3</sub>:MoO<sub>3</sub> sample with concentrations 0.2N:0.1N, of MoO<sub>3</sub> as additives in TiO<sub>2</sub> film was calculated 14.980  $\times 10^{3}\Omega$ -m.

## **ACTIVATION ENERGY**

Figure shows plot of log( R) versus reciprocal of temperature, (1/T) for In2O3:MoO3 thin films with normality 0.2N:0.1N.



Figure 3.5 Activation energy of Binary oxide In2O3:MoO3 thin film with concentration 0.2N:0.1N

This plot is reversible in both heating and cooling cycles obeying the Arrhenius equation

Where,

Ro = the constant = Resistance at room temperature

 $\Delta E$  = The activation energy of the electron transport in the conduction band,

"K = Boltzman constant

"T = Absolute temperature"

The Activation energy at high temperature and at low temperature were found 0.1707eV and 0.7976 eV respectively at 0.2N:0.1N.

# TEMPERATURE COEFFICIENT OF RESISTANCE (TCR)

Temperature coefficient of resistance (TCR) of In<sub>2</sub>O<sub>3</sub>: MoO<sub>3</sub> thin films prepared at  $400^{\circ}$ C is calculated by using the following relation,

TCR:-





**Figure 3.6** TCR graph of binary oxide In2O3: MoO3 thin film with concentration 0.2N:0.1N

The temperature coefficient of resistance (TCR) was found as 0.0179 /  $^{\rm 0}$  C.

#### GAS SENSING PROPERTIES

The main feature is the optimization of film sample operating temperature for test gases. The sensitivity and selectivity of the thin film sensor are estimated on the basis of measured data in the air condition for a fixed gases concentration of 1000 ppm.

The variation in sensitivity of binary oxide In<sub>2</sub>O<sub>3</sub>:MoO<sub>3</sub>thin films as a function of temperature and for LPG, Ethanol, NH<sub>3</sub>, CO and NO<sub>2</sub> gases [1000 ppm concentration]. The operating temperature was varied at the interval of  $50^{\circ}$ C. The sensitivity of gas was measured at specific operating temperatures using the following equation based both on measured resistance in the air and in the gas atmosphere.

$$Sensitivity(S) = \frac{\left|\frac{R_a - R_g}{R_a}\right| \times 100$$

Where,

 $R_a$  – resistance of thin film in air atmosphere,

 $R_g$  – resistance of thin film in gaseous atmosphere.





The film of binary oxide In2O3:MoO3was exposed to various gases. The film of binary oxide In2O3: MoO3was exposed to various gases. The film of In2O3: MoO3 at 0.2N:0.1N showed 64.50% sensitivity for CO gas at operating temperature  $150^{\circ}$ C and CO gas concentration was at 300 ppm.

#### V CONCLUSION

The Binary oxide In2O3: MoO3 Nano crystalline thin films have been grown successfully on to glass substrate . The average particle size of film at concentrations0.2N:0.1N was found 198 nm. Specific Surface area with different concentrations of binary oxide In2O3:MoO3 was found as 26.49002 m<sup>2</sup>/g. The atomic % of O, Mo, In were found as 85.48% ,11.44% and 3.08 % respectively. XRD gives the grain size of film 11 nm. The resistivity of sample was calculated 14.980x10<sup>3</sup>  $\Omega$ -m. The Activation energy at high temperature and at low temperature were found as 0.1707eV and 0.7976 eV respectively .The temperature coefficient of resistance were found was 0.0179/<sup>0</sup> C. The film of In2O3: MoO3 at 0.2N:0.1N showed 64.50 % sensitivity for CO gas at operating temperature 150<sup>o</sup>C and CO gas concentration was at 300 ppm.

#### REFERENCES

- 1. S. Basu, A. Dutta, Materials Chemistry and Physics 47(1997) 93-96
- G. UozumiM.Miyayama, H.Yanagida, Journal of Materials Science 32 (11) (1997) 2991-2996.
- 3. Sian.T.S and Reddy.G.B, Appl.Surf.Sci., 2004,236, 1-5.
- 4. W. W. Chen, Y. K. Liu, Z. J. Qin, Y.
- 5. X. Chen, S.S. Mao, Chem. Rev. 107 (2007) 2891–2959.
- Y.F. Sun, S.B. Liu, F.L. Meng, J.Y. Liu, L.T. Kong, J.H. Liu, Sensors 12 (2012) 2610–2631.
- S. Matsushima, Y. Teraoka, N. Miura, N. Yamazoe, Jpn.J.Appl.Phys.27 (1988) 1798-1802
- J. G. Duh, J. W. Jou, B. S. Chiou, Electrochem.Soc.136 (1989) 2740-2747

- 9. P. Prathap, N. Revathi, K. T. R. Reddy, and R. W. Miles, Thin SolidFilms 518, 1271 (2009).
- T. Brezesnski, J. Wang, S. H. Tolbert and B. Dunn, Nat. Mater., 9 (2010) 146.
- 11. D. V. Ahire, S. D. Shinde, G. E. Patil, K. K. Thakur, V. B. Gaikwad, V. G. Wagh1 and G. H. Jain, International Journal On Smart Sensing And Intelligent Systems, Vol. 5, No. 3, September 2012, Issn 1178-5608.
- E. B. Santos, F. A. Sigoli, I. O. Mazali, J. Solid State Chem. 190 (2012) 80-84.
- 13. Granqvist.C.G, Handbook of Inorganic Electrochromic Materials, Elsevier, 115.
- 14. Sian.T.S and Reddy.G.B, Appl.Surf.Sci., 2004,236, 1-5.
- 15. X. Chen, S.S. Mao, Chem. Rev. 107 (2007) 2891-2959.
- 16. Y.F. Sun, S.B. Liu, F.L. Meng, J.Y. Liu, L.T. Kong, J.H. Liu, Sensors 12 (2012) 2610–2631.
- 17. L.Gao, Li, Q., Song, Z., Wang Sens. Actuators B71 (2000) 179-183
- 18. S. Cho, J. Korean Phys. Soc. 60, 2058 (2012).
- 19. D.R. Patil, L.A.Patil, Sensors and Actuators B 123 (2007) 546–553.
- 20. L. Satyanarayana, K. Madhusudan Reddy, S.V. Manorama, Sensors and Actuators B 89(2003) 62-67.
- 21. D.M. Smyth, Solid State 129 (2000) 5-12.
- Titkov.IE, Delimova.LA, Zubrilov.AS, Seredova.NV, Liniichuk.IA and GrekhovIV: J Mod Opt., 2009, 56, 653– 660.
- 23. G. Korotcenkov, Sens. Actuators B: chem., 107, 2005, 209–232.
- 24. S. Matsushima, Y. Teraoka, N. Miura, N. Yamazoe, Jpn.J.Appl.Phys.27 (1988) 1798-1802
- 25. J. G. Duh, J. W. Jou, B. S. Chiou, Electrochem.Soc.136 (1989) 2740-2747
- S. Basu, A. Dutta, Materials Chemistry and Physics 47(1997) 93-96.
- 27. X. P. Shen, L. J. Guo, G. X. Zhu, C. Y. Xi, Z. Y. Ji and H. Zhou, RSC Adv., 2015, 5, 64228–64234.
- 28. X. Y. Lai, P. Li, T. L. Yang, J. C. Tu and P. Xue, Scr. Mater., 2012, 67, 293–296.