



Preparation and Characterization of CdO Thin Films Prepared by Chemical Method

B. Kavitha^{*}, M. Nirmala, S. Poornachandra, M. Pavithra

Department of Physics, Sri G.V.G Visalakshi College for Women, Udumalpet, TN, India

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*kavitha.sudharsan@gmail.com

ABSTRACT

CdO thin films of different thicknesses were prepared by a simple chemical technique called Successive Ionic Layer Adsorption & Reaction (SILAR) technique onto well cleaned substrates and the thicknesses of the deposited films were determined by gravimetric technique. The films are investigated with X-ray diffraction, scanning electron microscopy and optical spectroscopy. The structural characterization was carried out by X-ray diffraction which confirms the polycrystalline nature of the films with a cubic structure. SEM analysis of the films enabled the conclusion that the prepared films are uniform, smooth and polycrystalline. From the transmittance spectra the type of transition, band gaps of the films, refractive index, dielectric constants of the films were estimated. From the results of the structural and optical analysis, CdO has been identified as an alternate n-type conductor.

Keywords: BANDGAP; CdO; EDAX; SILAR; TCO; XRD.

1. INTRODUCTION

Transparent conductive oxide (TCO) films have been extensively studied because of their use in semiconductor device technology (Yan *et al.* 2001; Tokumoto *et al.* 2002; Chopra and Das, 1983). Particularly, II–VI compound semi-conductor oxides of many metals such as Tin, Indium, Zinc, Cadmium and their alloys, can be used as TCO's possessing transparent conducting property. Most of the studied transparent conducting metal oxides are anion deficient (i.e., Oxygen deficient) and hence are always n- type conductors (Andreas Stadler, 2012). These oxide materials have attracted significant attention due to their potential applications in optoelectronics, ultraviolet light emitting devices, laser diodes, solar cells and optical communications (Ito *et al.* 2006; Dakhel, 2011). However, among these compounds, cadmium oxide (CdO) films received less attention mainly due to their narrow band gap energies compared to other wide band gap oxides (Salunkhe *et al.* 2009; Galicia *et al.* 2000; Zhao *et al.* 2002). The n-type CdO thin films exhibit rock salt structure (FCC) with band gap 2.2eV. It also has good optical conductivity and transmission in the visible range (Ortega *et al.* 1999). CdO films can be synthesized by various methods such as Spray Pyrolysis (Dong Ju Seo, 2004), Sputtering (Subramanyam *et al.* 1998), Sol-gel spin coating (Carballeda-Galicia *et al.* 2000), Activated reactive evaporation (Sravani *et al.* 1991), Metal Organic Chemical Vapour deposition (MOCVD), Pulsed laser deposition. The main goal of the work is to seek a simple, non-vacuum and economic deposition technique for efficient transparent films. In this study CdO thin films were prepared using simple Chemical

Method Successive Ionic Layer Adsorption and Reaction (SILAR) technique, because this method has received immense advantages such as low cost, high growth rate at low temperature and control shape and size of thin films. CdO thin films have prepared at fixed deposition parameters such as concentration, pH value and temperature and annealed at different temperatures. Further, the structural and composition characteristic of annealed CdO thin films are discussed in detail. Optical studies such as transmission, bandgap, refractive index, absorption co-efficient, extinction coefficient, dielectric constant (imaginary and real) were evaluated.

2. EXPERIMENTAL METHODS

Cadmium Oxide (CdO) thin films were deposited onto clean glass substrates using the Successive Ionic Layer Adsorption & Reaction (SILAR) technique. All chemicals used were of Analytical Grade. Before the deposition, the substrates were rinsed with distilled water, washed with detergent and then rinsed with distilled water and then dried in an oven. This process was carried out to ensure clean surface essentially for the formation of nucleation centres that is required for thin film deposition. In SILAR technique, the depositions of CdO thin films were carried out from solutions containing Cadmium nitrate as cationic precursor (3.08 gm) and H₂O₂ (40 ml) as anionic precursor. The deposition was carried out at a temperature of 70 °C. The deposition cycle was varied from 30 to 100 cycles to obtain films of different thicknesses. It has been found that uniform films can be prepared if the deposition cycle range is 30–60 cycles. While varying the pH of the bath, it is found that the

solution having pH value 12 gives uniform films. The deposition process is as follows:

1. Well-cleaned substrates was immersed into the cationic precursor for 20 s and then rinsed with deionized water for 30 s to remove unattached ions.
2. Then the substrates were immersed into anionic precursor solution for 18 s and then rinsed with deionized water for 30 s to remove unattached ions.
3. The above cycle was repeated and the optimized dippings were 40 dipping times to get enough film thickness.

Thus, deposition parameters such as temperature, concentration of ions in the bath and dipping cycle have been optimized and then the films were taken out and dried naturally. The thicknesses of the prepared films determined by the gravimetric technique and CdO thin films were annealed at 100 °C, 200 °C, 300 °C for one hour and then used for the analysis. Structural characterization of these films was carried out by using Shimadzu (Lab X-6000) x-ray diffractometer with Cu K α ($\lambda = 1.5406 \text{ \AA}$) line in 2θ range from 20 to 80 degrees. A JASCO (V570: UV-vis-NIR) double beam spectrophotometer was used for optical studies in the wavelength range 400–2500 nm. Energy dispersive x-ray analysis (Thermo SuperDry II) is used to carry out semi-quantitative elemental analysis of the annealed thin film samples.

3. RESULTS & DISCUSSION

3.1 Compositional Analysis

Fig. 1 shows the EDAX spectra of representative CdO thin film prepared by SILAR technique and the spectrum shows the presence of the chemical constituents (Cd and O) of CdO thin films. EDAX quantitative results confirm the atomic percentage of constituents in the prepared films as in table 1.

Table 1. Elemental composition of CdO thin films

Annealing temperature: 100 °C and thin thickness: 530 nm	
Element	Atomic %
O k	36.45
CdL	63.55

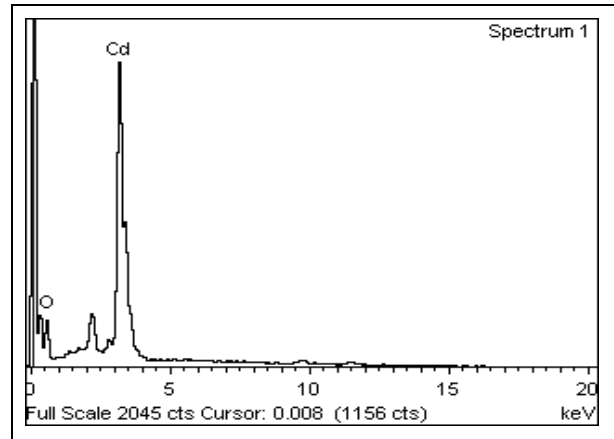


Fig. 1: Representative EDAX spectra of CdO thin film

3.2 XRD Analysis of CdO Thin Films

Fig. 2 shows the x ray diffractograms of CdO thin films prepared at 70 °C containing cadmium nitrate, H₂O₂ and deionized water in same molar concentrations. Uniform and adherent CdO thin films are prepared and annealed at different temperatures of 100 °C, 200 °C and 300 °C respectively and then used for structural analysis. From the XRD profiles, the interplanar spacing d_{hkl} was calculated using the Bragg's relation,

$$d_{hkl} = n\lambda / 2\sin \theta \quad \dots\dots\dots(1)$$

The crystallite size (D) was calculated using the formula from the full width at half maximum (FWHM).

$$D = k\lambda / \beta \cos \theta \quad \dots\dots\dots(2)$$

where the constant K is the shape factor ≈ 0.98 , ' λ ' is the wavelength of the x-rays (1.5406 \AA for Cu K α), ' θ ' is the Bragg's angle and ' β ' is the FWHM. The dislocation density (δ) can be evaluated from the crystallite size (D) by the following relation.

$$\delta = 1 / D \quad \dots\dots\dots (3)$$

where the constant K is the shape factor ≈ 0.98 , ' λ ' is the wavelength of the x-rays (1.5406 \AA for Cu K α), ' θ ' is the Bragg's angle and ' β ' is the FWHM. The dislocation density (δ) can be evaluated from the crystallite size (D) by the following relation.

$$\varepsilon = \beta \cos \theta / 4 \quad \dots\dots\dots(4)$$

From (hkl) planes, the lattice constant can be evaluated using the relation

$$1 / d = 4/3 \{h^2+hk+k^2/a^2\} + 1^2/c \dots (5)$$

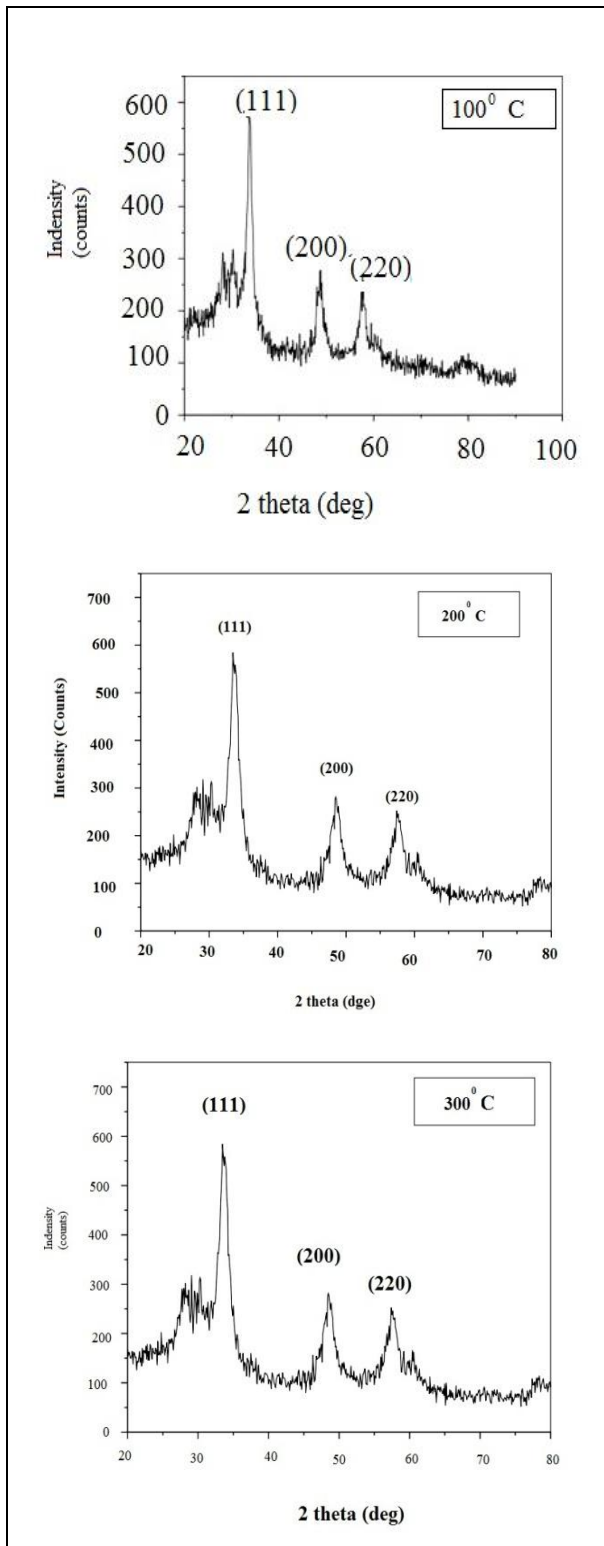


Fig. 2: X-ray diffractograms of CdO thin films

All the films shows polycrystalline nature containing cubic structure of pure CdO phase. The lattice spacing 'd' has been determined and it has been found that it is in well agreement with ASTM DATA (75-0594), earlier researchers and are presented in table 2. The predominant peak (111) and weak intense peaks (200)

and (220) plane were observed from the XRD profile. When annealing temperature increases from 100 °C to 300 °C, the crystallite size also increases from 5.5 nm to 5.8 nm. The intensity of the diffraction peak were also found to increase with increase in annealing temperature and get sharper with decreasing full width half maximum(FWHM). This can be attributed to the improvement in crystallinity of CdO thin films. The structural parameters of CdO thin films are calculated and given in table 2.

Lattice constant (a) is found to be 0.461nm, 0.462 nm and 0.464 nm for the films annealed at 100 °C, 200 °C and 300 °C, respectively. However a shift in lattice constant towards its standard value (0.4695 nm) is observed when the films annealed from 100 °C to 300 °C. From table 2, it is evident that the post annealing process improves the quality of the crystalline thin films.

The individual crystalline size (D_c) in the films of different thicknesses have been estimated equation (2) are in the range of 520 – 650 nm and in very good agreement with the reported values. Using the size of the crystallites, the dislocation density, the number of crystallites per units surface area, volume and strain have been determined equation (2-5) and presented in table 2. From the table 3 it is observed that both dislocation density and strain decrease with increasing annealing temperature. This can be attributed to the improvement in crystallinity due to the regular arrangements of atoms in the crystal lattice.

Linear increase in texture coefficient is observed for all thin films with increase in annealing temperature. For a preferential orientation, the texture coefficient should be greater than unity. It observed that the texture coefficient for (111) plane is greater than unity and increases with increase in annealing temperature.

Table 2. XRD data of CdO thin films

Annealing Temp (°C)	Film thickness	hkl plane	2θ (degree)	d spacing (Å)
			observed	observed
100	530	111	33.63	2.66
		200	48.43	2.40
		220	57.77	1.70
200	620	111	33.48	2.66
		200	48.33	2.49
		220	57.55	1.65
300	650	111	33.64	2.65
		200	48.52	2.41
		220	57.55	1.59

Table 3. Structural parameters of CdO thin films

Annealing Temp (°C)	Film Thickness (nm)	Lattice constant		Volume a^3 (10^{-29})	Crystalline Size D_c (nm)	No of Crystalline per unit Area ($10^{15}m^{-2}$)	Dislocation Density(δ) ($10^{15}/lines/m^2$)	Strain(ϵ) (10^{-3})	Textured coefficient
		a(nm)							
		observed	ASTM						
100	530	0.461		9.79	5.49	67.25	3.46	6.72	2.25
200	610	0.462	0.465	9.86	5.60	31.81	3.33	6.38	2.60
300	650	0.464		9.98	5.79	29.75	3.19	5.38	2.77

3.3 Optical Analysis of CdO Thin Films

Absorption coefficient was calculated using the transmittance (T) value measured for a particular wavelength and the film thickness (t) using the relation,

$$\alpha = -\ln(T)/t$$

The absorption index or the extinction coefficient (k), which is the attenuation per unit radian, can be written as

$$k = \alpha\lambda/4\pi$$

where λ is the wavelength of the monochromatic light.

The optical transmittance spectra of CdO thin films of different annealing temperatures are shown in fig 3. All the films show transmittance above 80% in the IR region. The film annealed at 300 °C exhibited maximum transmittance and minimum transmittance for 100 °C. The absorption edge is found to shift towards longer wavelength as a function of annealing temperature.

Important optical parameters such as type of transition, band gap etc. can be satisfactorily analyzed on the basis of formulae derived for 3D and 2D models. By 3D crystal model, nature of transition in film composition can be obtained by plotting $(\alpha hv)^{1/2}$ versus (hv) for various values of r [α is the absorption coefficient, hv is the photon energy and exponent r determines the type of transition and dimensionality of the bands]. r has values $1/2$ (direct allowed), $3/2$ (indirect allowed), 2 (direct forbidden). Extrapolation of straight-line portion of $(\alpha hv)^{1/2}$ versus (hv) plot at $(hv > E_g; E_g = \text{direct band gap})$ to zero absorption (hv -axis) gave the value of energy gap.

Plot of $(\alpha hv)^2$ versus (hv) (fig. 4) for all CdO thin films were plotted and the straight line portion is extrapolated to cut the x axis which gives the band gap. The estimated band gaps are found to be 2.2eV, 2.06eV and 1.86eV for the films annealed at 100 °C, 200 °C and 300 °C respectively (table 4). It was observed that the band gap decreases with increase in the annealing temperatures and in agreement with the earlier

researchers. The decrease in optical band gap energy with increasing annealing temperature may be due to the increase in the carrier concentration and also may be due to its quantum confinement effect since the crystallite size is found to be very small.

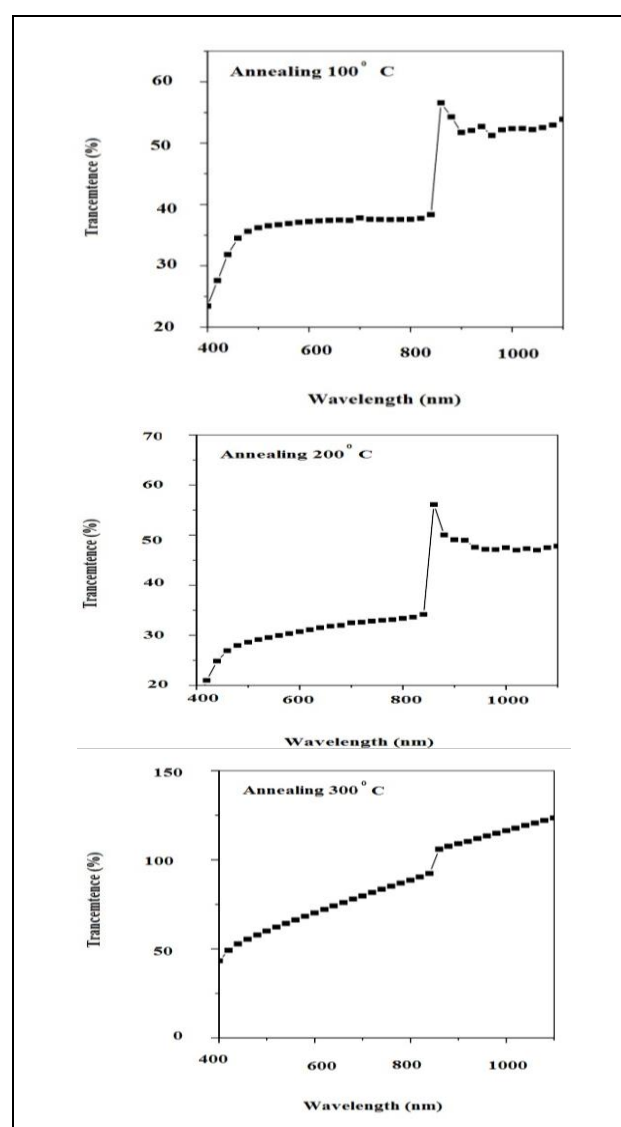


Fig. 3: Transmittance spectra of CdO thin films prepared and annealed at different temperatures

Plot of (αhv) versus $(\alpha hv)^{1/2}$, $(\alpha hv)^{1/3}$ and $(\alpha hv)^{2/3}$ reveals that CdO films did not have line above $hv > E_g$. Since extrapolation of it did not touch the zero absorption axis which confirms the fact that CdO phase

do not have indirect allowed, direct forbidden and indirect forbidden transitions. The optical parameters such as absorption coefficient, extinction coefficient, reflectance, refractive index, band gap and dielectric constant are estimated and are presented in table 4.

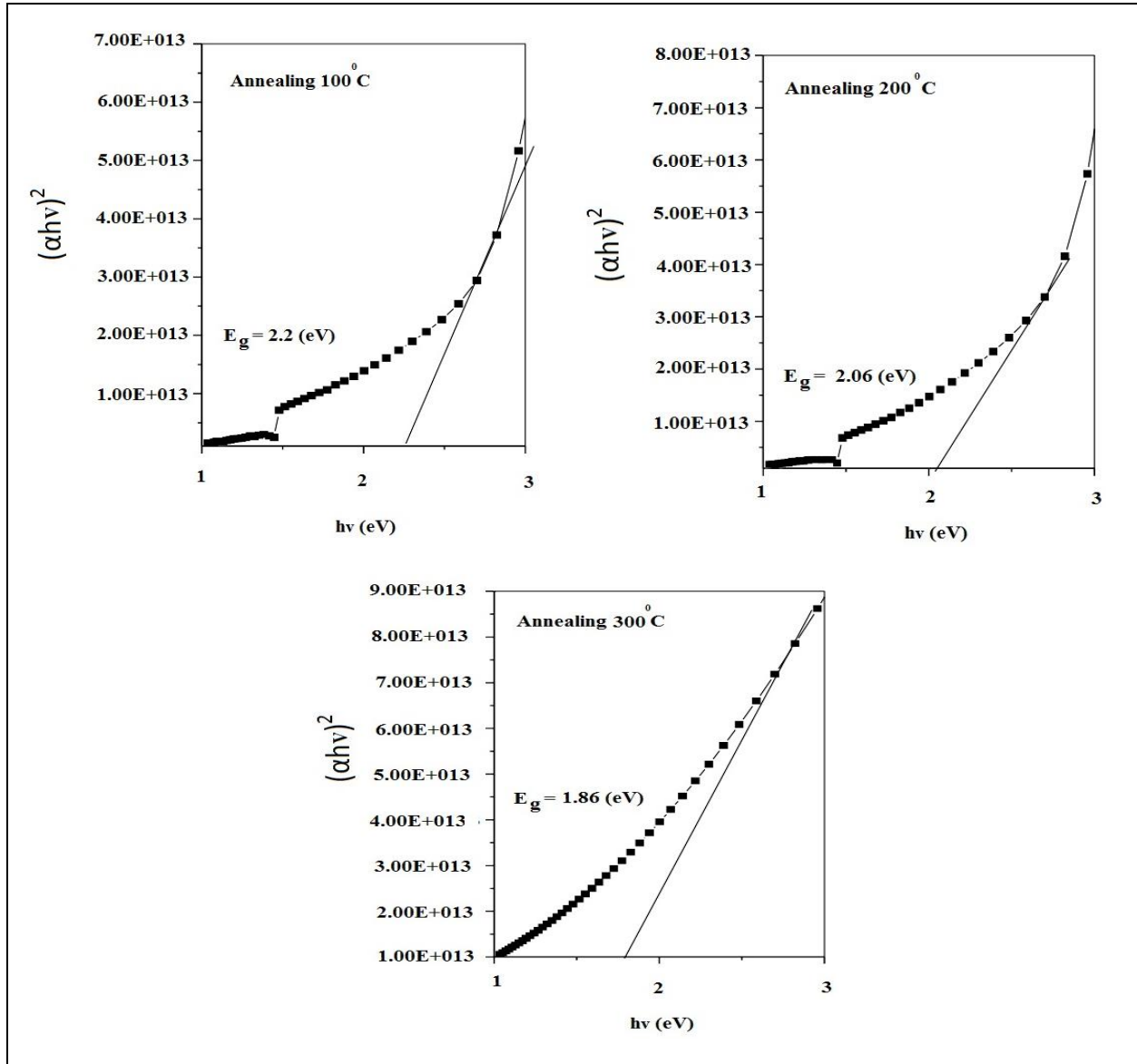


Fig. 4: Plot of $(\alpha hv)^2$ vs hv of CdO thin films

Table 4. Optical parameters of CdO thin films prepared and annealed at different temperatures

Annealing Temp °C	Film thickness	α (10^6 m^{-1}) [$\lambda=1000\text{nm}$]	% Reflectance R	Refractive Index n	Extinction coefficient k [$\lambda=1000\text{nm}$]	Dielectric Constant		E_g (eV)
						ϵ_r	ϵ_i	
100	530	3.77	180.46	1.186	0.3	2.20	0.2887	2.20
200	610	1.22	179.83	1.485	0.0972	2.256	0.2925	2.06
300	650	1.22	1156.80	1.505	0.0972	1.134	0.66	1.86

Refractive index of CdO thin films has been calculated using the relation,

$$n = 1 + R/1 - R \dots\dots\dots (8)$$

Fig. 5 shows the wavelength dependence on refractive index at different annealing temperatures. Refractive index of the CdO films increased from 1 to 2.2 with increase in temperature from 100 to 300°C in the wave length region of 300 to 600 nm. Thereafter, refractive index of the CdO films annealed at 100 °C and 200 °C has a constant refractive value in the wavelength region of 600 to 1200 nm, but the refractive index of the film annealed at 300 °C decreases in the same wavelength region. This unusual increase and decrease in the refractive index as a function of wavelength may be due to the damping of CdO thin films (table 4). Reflectance of CdO thin films has been calculated using the relation, $R=1-(A+T)$ (9)

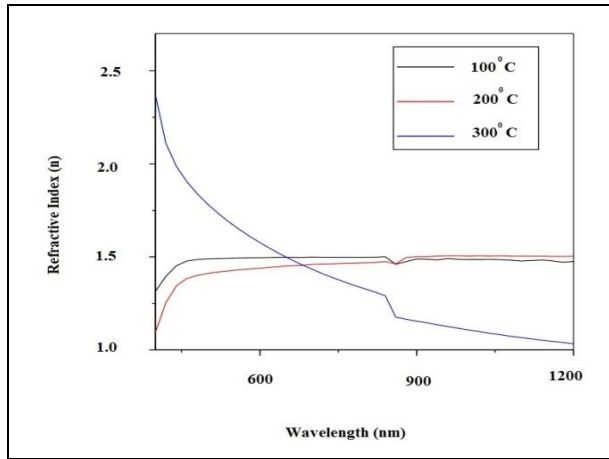


Fig. 5: Plot of Wavelength vs Refractive index of CdO thin films

Fig. 6 shows the wavelength dependence on percentage of reflectance at different annealing temperatures. Film annealed at 300 °C shows the reflectance of the film increases as wavelength increases with highest reflectance upto 900 nm. The higher reflectance exhibited by this material makes it useful in the manufacture of highly reflectance mirrors commonly found in desktop scanners, photocopy machines, car head lamps and halogen lamps. But the films annealed at 100 and 200 °C shows the minimum reflectance in the same wavelength region shown in table 4.

Real part of the dielectric constant is related to the n value and imaginary part of the dielectric constant to the k value. The values of ϵ_r and ϵ_i were calculated using the relations,

$$\begin{aligned} \epsilon_r &= n^2 - k^2 \\ \epsilon_i &= 2nk \end{aligned} \dots\dots\dots (11)$$

The imaginary ϵ_r and real ϵ_i parts of the dielectric constants are plotted as a function of photon energy and are presented in fig. 7 & table 4. The imaginary part shows the absorption of energy of a dielectric material from an electric field due to dipole motion. The real dielectric constant (fig 7) increases from 1 to 2.5 as annealing temperature increases. The imaginary part of dielectric constant decreases with increase in photon energy (fig 8).

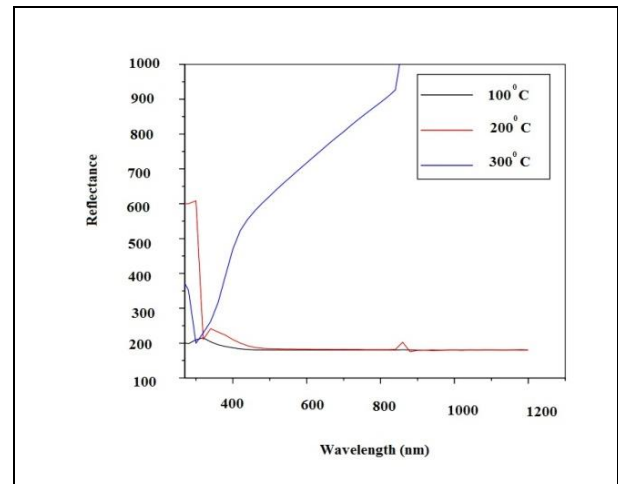


Fig. 6: Plot of Wavelength vs Reflectance of CdO thin films

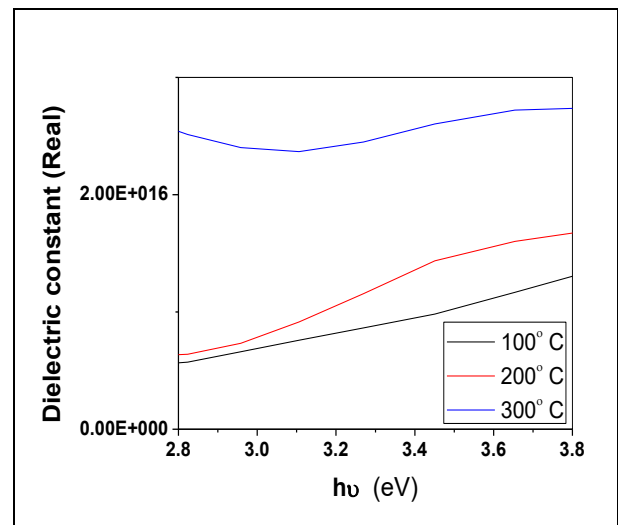


Fig. 7: Plot of (hv) vs Dielectric constant (Real) of CdO thin films

The extinction coefficient (k) plot against the photon energy in fig.9 shows that the extinction coefficient (also called the attenuation coefficient) is high for low photon energy and it decreases with increase of photon energy and as well as for increase in annealing temperature. Stronger absorbing medium shows more extinction coefficient. The energy loss of electromagnetic radiation through that medium is measured by the extinction coefficient of a particular substance (table 4).

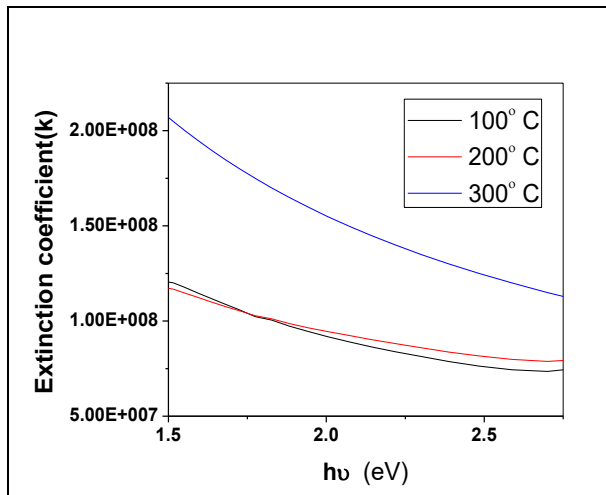


Fig. 8: Plot of $(h\nu)$ vs Dielectric constant (Imaginary) of CdO thin films

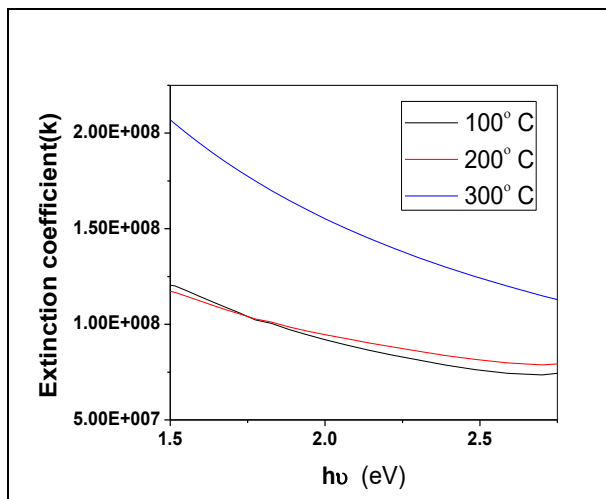


Fig. 9: Plot of $(h\nu)$ vs Extinction coefficient (k) of CdO thin films

4. CONCLUSION

CdO Thin Films are prepared by Successive Ionic Layer Adsorption and Reaction method. Thickness of the prepared films are calculated by Gravimetric Method. Structure of the prepared films has been analyzed by XRD. It reveals that the prepared films are polycrystalline in nature with cubic structure. The characteristics peaks are identified and the structure parameters are calculated and presented. EDAX analysis confirms the presence of the constituents Cd and O and the wavelength percentage is presented. The type of transition and band gap has been estimated from optical analysis. The band gap is found to be in the range 2.2 to 1.8 eV respectively. Then optical parameters such as absorption coefficient, extinction coefficient, reflectance, refractive index and dielectric constant are calculated and presented.

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CONFLICTS OF INTEREST

The authors declare that there is no conflict of interest.

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