

Studies on Chemically Synthesis of Polycrystalline CdTeO₃ Thin Films

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Abstract: - Polycrystalline cadmium telluride oxide (CdTeO₃) thin films have been achieved by Chemical synthesis process and annealed at 300⁰C in air. XRD of films shows the peaks corresponding to single phase face centered cubic polycrystalline CdTeO₃. The compressive strain energy of the film is 0.02569J calculated by angle of contact. Optical absorption study revealed that CdTeO₃ is a direct band gap of 2.3eV for the film annealed at 300⁰C and it is effect of oxygen concentration of the deposited material. The resistivity of the films was decreases with increase the temperature.

Keywords: - Chemical synthesis, cadmium telluride oxide, strain energy.

I. INTRODUCTION

Semiconductor Oxides thin films have number of practical applications in the different devices, for the fabrication of field effect transistors [1] and surface passivation[2]. Thin films of group of II- VI are generally used in many semiconducting compounds which are used in many semiconductor devices because of their optoelectronic properties and the development of flexible and light weight solar cells which are highly stable[3-4]. Due to wide energy band gap of semiconductor oxides, they can be considered as absorbent layers even in the ultraviolet region and used for optoelectronic applications[5]. A polycrystalline cadmium telluride thin film is one of the most promising low cost candidates for the terrestrial applications. The studies on cadmium telluride and its oxides are very important compound because of its technological importance because they may the part of semiconductor device structures with specific functionality. It is direct band gap material with high absorption coefficient [6-7]. Cadmium telluride oxides can be obtained with controlled energy band gap available values higher than 1.5eV for CdTe[8] and upto ~ 3.8eV accoding to oxygen concentration[1]. The addition of oxygen to CdTe at dopent level has been aimed to improve the stability of CdTe-based photovoltaic devices [9-12]. Not much work has been done on CdTeO material till few researchers pay their attention on CdTeO. The synthesis of Oxygenated CdTe films has been reported by different techniques as pulse laser deposition [10], R.F. sputtering [5, 13] etc. Some researcher have conclude that tellurium oxygen (Te-O) bonds were associated with CdTeO[5-14] and other researchers conclude that it is co-existence of CdTe₂O₅ [15] but more groups also available with different composition [1,16]. In this communication we report and analyzed For the present work, Chemical bath deposition (CBD) method was used for deposition of the films, as it is a simple, convenient for the large area deposition, capable of yielding good quality films, it forms ion-by-ion formation, controlled thickness by varying time and the most important is, it does not requires sophisticated instrumentations.

In the present paper, polycrystalline oxygenated thin films of CdTe were deposited at room temperature on glass substrate by CBD and annealed at 300⁰C. The morphological, structural and optical properties of annealed films were studied.

II. EXPERIMENTAL

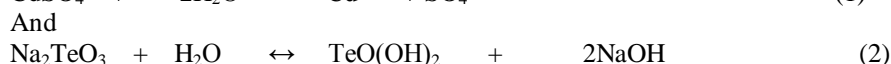
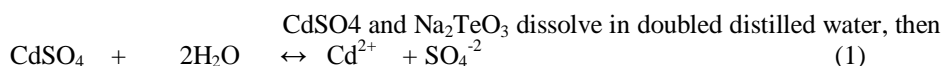
Polycrystalline CdTeO₃ thin films have been deposited on glass substrates by CBD method from aqueous solution of 0.1 M CdSO₄, 0.1 M Na₂TeO₃ and 20% of ammonia used as a complexion agent prepared in doubled distilled water. There were no further treatments on the chemicals. Both solutions of CdSO₄ and Na₂TeO₃ were taken independently in different containers, they were clear solutions. When both solutions were mixed together, white precipitate formed in container. Before dipping the glass substrates into mixtent solution, ammonium hydroxide added drop-by-drop into precipitate and the precipitate completely dissolves and the solution becomes clear. The glass substrates were cleaned with lebolene solution and by ultrasonic cleaner with doubled distilled water for removal of contaminants from the surface of the films. These cleaned glass substrates were mounted on substrate holder and immersed into the solution for deposition at room temperature for different time intervals. The deposited films were homogeneous without cracks and off white in colored. The deposited film dried later in air and annealed at 300⁰C for 2 hours. The annealed films were characterized by

different techniques as X-ray diffraction (XRD) micrographs were obtained by using X-ray diffractometer (XRD Rigaku D/max-2400 with Cu-K α = 0.154 nm) at 40kV with 35mA and the scanning rate 0.02^o with counting time of 8 s. UV-VIS-NIS spectrophotometer (Hitachi 330, Japan) was used to record optical absorption spectra at room temperature angle of contact on substrate were used for calculating the strain energy on the films.

III. RESULTS AND DISCUSSION

CdTeO₃ films have been deposited by using cadmium sulphate (CdSO₄) as a source of Cd ions and sodium telluride as a source of Te. Ammonium hydroxide is a complexing agent for adherence and to adjust the pH of reaction bath.

The kinematics of the film formula can be understood from following reaction.



From eqⁿ (1) and (2),

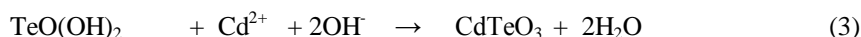


Figure 1 represents the structural analysis of annealed CdTeO₃ films on glass substrates, the crystalline structure of the sample in the lower part of figure and it shows low intensity peaks on diffractogram. The peaks appearing at 2 θ angles (23.3) and (30.3) with (100), (102) planes respectively were single phase face-centered cubic in nature and they were good agreement with JCPDS card No.80-0090 of CdTe. The diffraction features were observed which having greater intensities with sharp picks at (33.2), (38.3) and (55.3) and these peaks are of oxygenated CdTe and they were good agreement with JCPDS card No.77-1906 of oxygenated CdTe. However R.C. Rodriguez et al [2] reported the films of CdTeO₃ are single phased face-centered

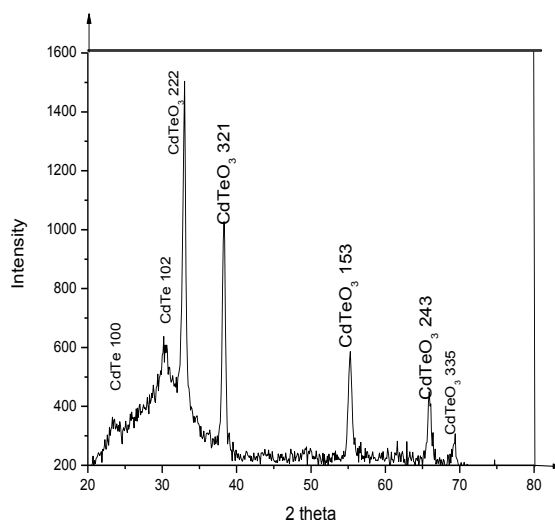


Figure 1- XRD of CdTeO₃ on glass substrate of annealed film at 300^o C.

cubic. The diffraction peaks of CdTeO₃ can be clearly identified and these results well agree with F.J. Beltrant et al [13] and Rhiger et al [17].

The particle size was approximately 3.25nm of the deposited film and it is determined by Debye-Scherrer's formula,

$$t = \frac{K\lambda}{\beta \cos\theta} \quad \text{----- (I)}$$

where, t be the individual crystalline size, K be Scherrer's constant which is from 0.89 to 0.9, λ be the wavelength, β be the FWHM of the peak and θ be the Bragg angle.

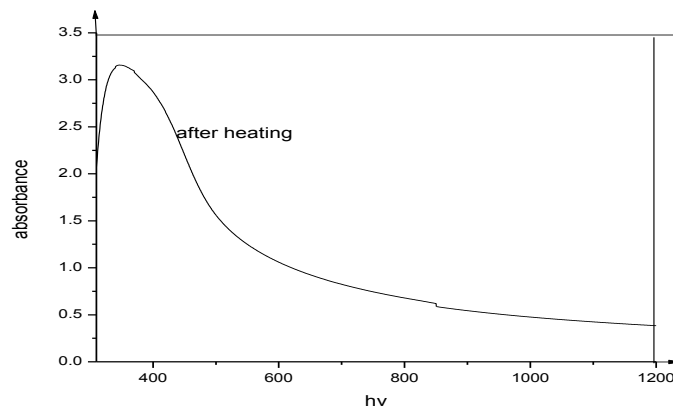


Figure: 2 Absorption spectra of CdTeO₃ glass substrate annealed at 300⁰C

Optical absorption study (Figure 2) of annealed films on glass substrates were in the wavelength range of 300 to 1200 nm. As the CdTe is a direct band gap material, CdTe oxide can be obtained with band gap values (Figure 3) higher than 1.5 eV for CdTeO₃ and it is 2.3eV, according to the oxygen concentration [18-19].

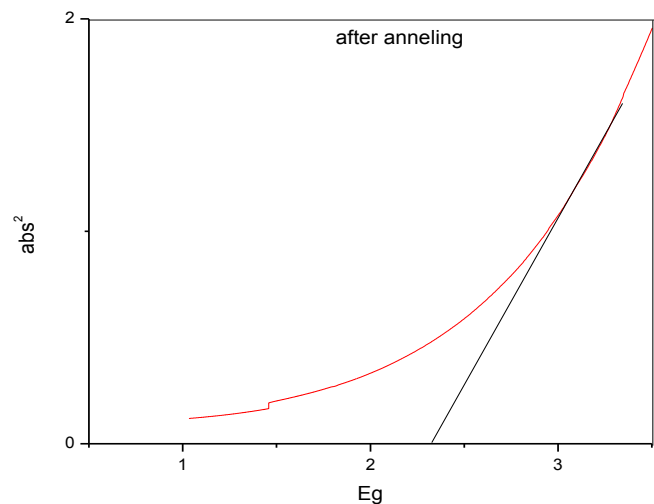


Figure: 3 Plot of $(\alpha hv)^2$ verses $h\nu$ for film on glass substrate



Figure: 4 Contact angle between CdTeO₃ films on the glass with water

The band gap shift may be accounted for one or more of the hexagonal to the cubic phase transformation taking place at temperature can reduce the energy band gap of films. The optical absorption behavior of polycrystalline CdTeO₃ thin films deposited on glass substrates depends upon the size and the concentration of the CdTe polycrystalline. The angle of contact of the CdTeO₃ film (Figure 4) was measured and the surface energy of the film was measured and found 0.02569 J.

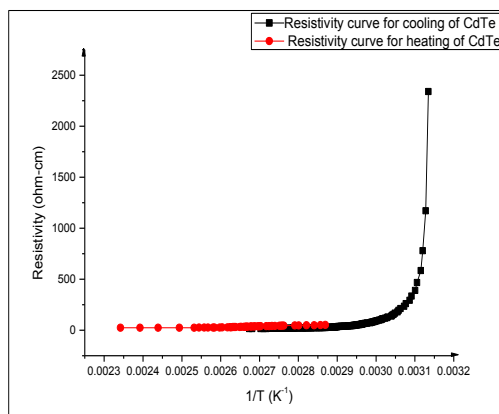


Figure 5 Resistivity of CdTeO₃ films during heating and cooling.

IV. CONCLUSIONS

Chemical bath deposition of CdTeO₃ thin films on glass substrates is simplest method for polycrystalline films. X-ray diffraction study revealed that, CdTeO₃ films are single face centered cubic structures and polycrystalline in nature. The band gap energy were slightly higher than band gap of CdTe ($E_g = 1.5$ eV), due to oxygen concentration. SEM micrograph shows the particles was not single crystal grains but it composed of number of crystallites. Surface energy is calculated by angle of contact and it is 0.02569J. The resistivity of the films (Figure 5) was decreases with increase the temperature.

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