

Structural and Optical Properties of Cd_{0.5}Zn_{0.5}S: Sn Nanocrystalline Thin Films Deposited By Chemical Bath Technique

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Abstract:- Nanocrystalline Cd_{0.5}Zn_{0.5}S:Sn thin films were prepared by chemical bath deposition at 80± 5°C from an aqueous bath containing zinc chloride, cadmium chloride tin chloride and thiourea in the presence of ammonia solution . the optical characteristics of the prepared thin films have been investigated by UV-VIS spectrophotometer in the wave length range (200 – 1100 nm).The energy gap(E_g) decreased from (4.1 to 2.2 eV) with increased doping by Sn. Also the optical constants such as refractive index ,extinction coefficient , real and imaginary dielectrics constants were investigated.The structural properties of obtained films were studied by X-ray diffraction analysis.the diffraction peak existed at 2θ=28.06 of plane (111).no change in the characteristic peak positions was observed in sampls doped with Sn.

I. INTRODUCTION

Cadmium zinc sulphide Cd_{0.5}Zn_{0.5}S thin films have received great attention due to their various optoelectronic applications .They have been used as a wide band-gap window material for hetrojunction solar cells [1,3]. The production of thin films with tailored optical properties is an important challenge for research in optoelectronic application [4,5] . Cd_{0.5}Zn_{0.5}S films are widely used for different applications , especially in photodiodes for solar cell [6,8]. Group 11-V1 compounds can form ternary and quaternary alloys with a direct fundamental band gap assignment over entire alloy composition range and high absorption coefficients , which can be used as materials for fabricating hetero-junction photovoltaic devices [9] .

II. EXPERIMENTAL

The CdCl₂ ,ZnCl₂ and NH₂-CS-NH₂ were used as the source materials of Cd²⁺ , zn²⁺ and s²⁻ ions ,respectively. A catalyst mixture was prepared by dissolving the appropriate amount of CdCl₂ and ZnCl₂ in distilled water .The Cd_{0.5}Zn_{0.5}S ternary thin film was co- precipitated by slowly adding aqueous solution of NH₃ to the mixture of CdCl₂ and ZnCl₂ aqueous solution which was kept stirring with constant 60 rpm through out the reaction . Before making the final solution for the preparation of the films , each of the elemental solutions was stirred for 30 min to obtain homogeneity among the constituents of the solution for uniform coating .for the preparation of good quality films , the concentration of (0.1M) CdCl₂ ,(0.1M) ZnCl₂ and (0.1M) NH₂ – CS –NH₂ were optimized and used as stock solution .The appropriate amount of CdCl₂ , ZnCl₂ and NH₂ – CS –NH₂ solutions were mixed in a 100ml beaker , and made alkaline by the addition of a few drops of NH₃ solution .The mixture we again stirred to form a homogeneous mixture . An appropriate amount of water was added to control the growth kinetics on the formation of Cd_{0.5}Zn_{0.5}S thin films .The glass substrates are cleaned by using hydrofluoric(HF)acid and water having the ration 1:3 volume ratio for 5 min at room temperature and placed vertically inside the chemical bath with the help of Teflon disc .The disc was attached with the constant rotating at 60 rpm and was kept immersed inside the chemical bath. Finally we doping the thin film by 1%,3% and 5% Sn.

Results and discussion

1.X-ray Diffraction Results

The x-ray diffraction patterns are shown in fig (1) .no charge in the character eristic peak positions was observed in samples doped with sn . the diffraction peak existed at 2θ=28.06 of plan (111). Such results suggest the formation of solid solutions between ZnS and CdS in general .However , due to the poor crystallinity the formation of single metal sulfides can not be completely ruled out . these results good agreement with results of Wii et all [11].

2.Optical properties

A representative plot of optical absorbance of film versus wave length of light is shown in the fig(1).the optical studies revealed that the films were highly absorptive with a direct type of transition, which allowed the optical band gap(E_g)to be determined using the following relationship.

$$\alpha = A/h\nu(h\nu - E_g)^{1/2} \dots\dots\dots(1)$$

Where A is a constant.

The experimentally observed value of $(\alpha h\nu)^2$ plotted against $h\nu$ is shown in fig(2) for different doping. The linear nature of the plots at the absorption edge confirmed that Cd_{0.5}Zn_{0.5}S is a semiconductor with a direct band gap. The optical band gap is decreased from (4.1 to 2.2) eV but at 5% doping by Sn the energy gap increased to 3.9 eV that may be due to the polarization or new orientation of the atoms. The extinction coefficient has been calculated by using the following relation [10].

$$K = \alpha\lambda/4\pi \dots\dots\dots(2)$$

Fig (5) shows the spectral dependence of extinction coefficient for Cd_{0.5}Zn_{0.5}S:Sn thin films.

The real part of refractive index (n) of thin films can be calculated for the relation.

$$n = [(1+R)/(1-R) + [4R/(1-R)^2 - K^2]^{1/2}]^{1/2} \dots\dots\dots(3)$$

Fig (4) shows the variation of the (n) with wavelength. The real and imaginary parts of dielectric constants were determined by the following relations in the absorption region.

$$\epsilon_1 = n^2 k^2 \dots\dots\dots(4)$$

where ϵ_1 is the real part of the dielectric constant represents the normal dielectric constant

$$\epsilon_2 = 2\text{ink} \dots\dots\dots(5)$$

ϵ_2 is the imaginary part of dielectric constant represents the absorption associated with radiation by free carrier. Figures (6 and 7) show the variation of ϵ_1 and ϵ_2 respectively.

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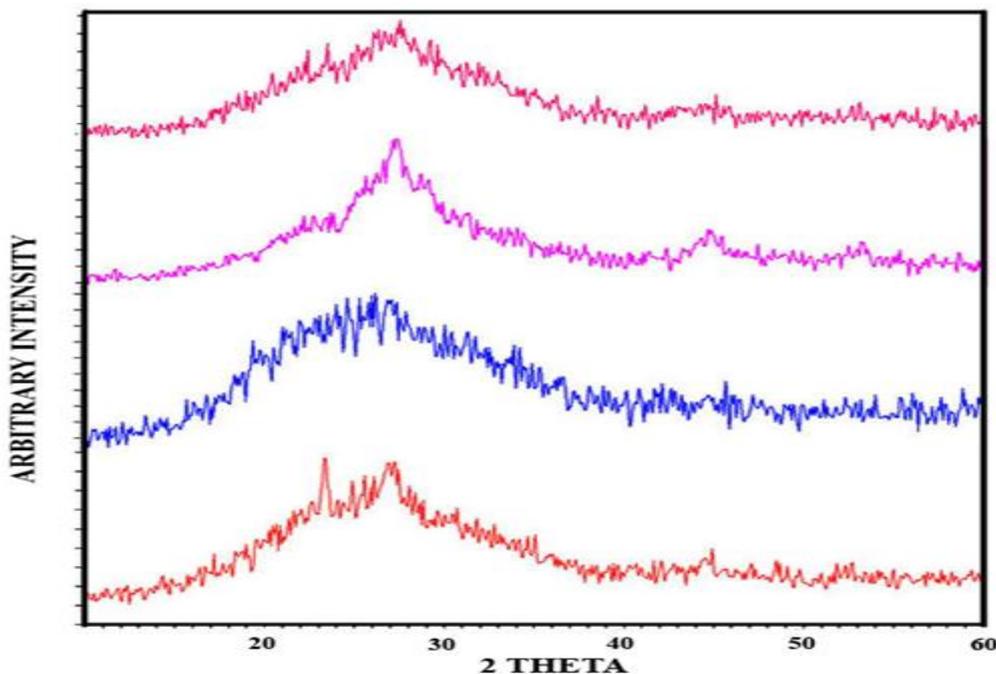
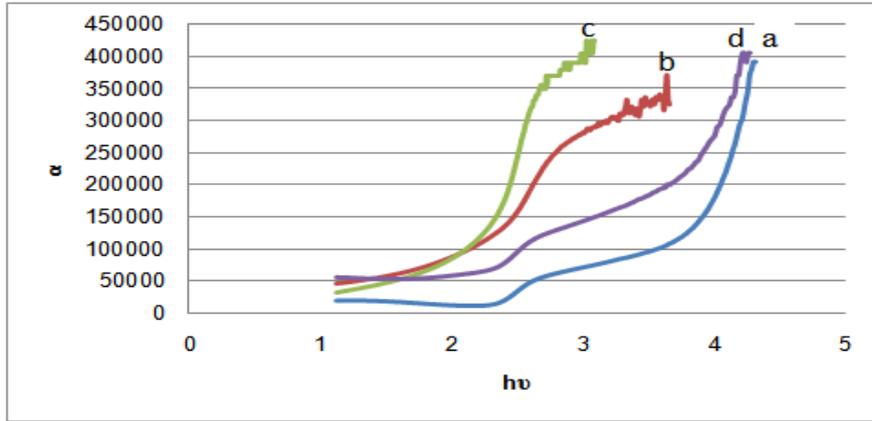
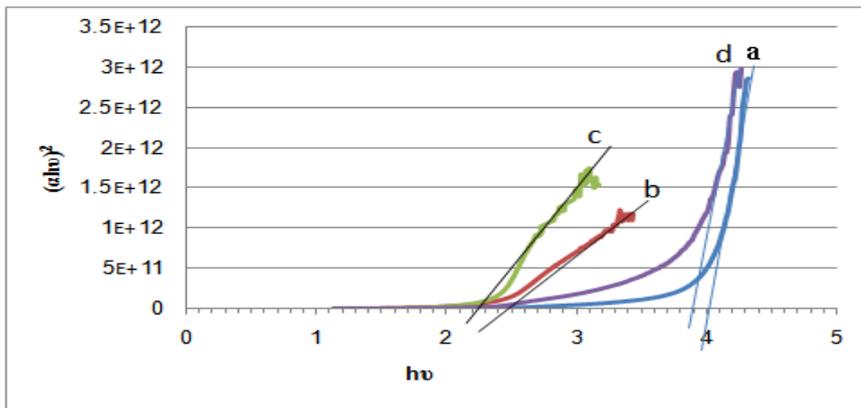


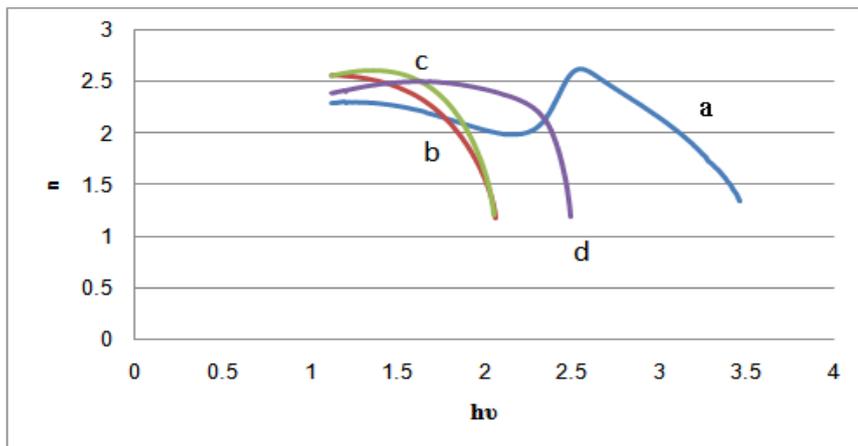
fig (1) x-ray diffractograms of Cd_xZn_{1-x}S:Sn films at X=0.5 a- without doping b- doping 1% by Sn c- doping 3% by Sn d- doping 5% by Sn.



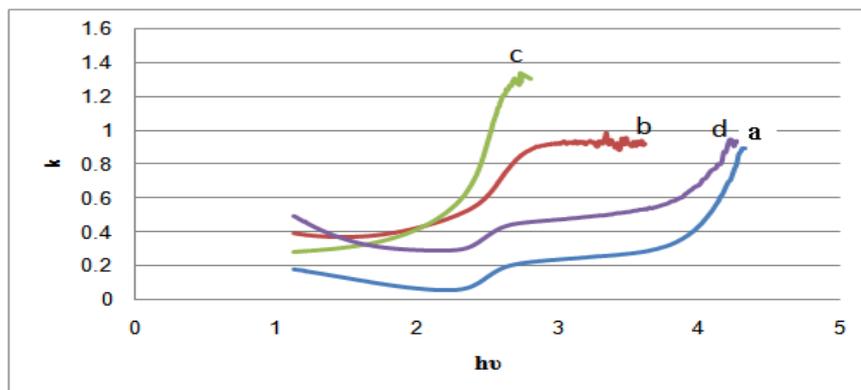
Fig(2) the variation of the absorption coefficient (α) with ($h\nu$) for thin films at $X=0.5$ a-without doping b- doping 1% by Sn c- doping 3% by Sn d- doping 5% by Sn.



Fig(3)the optical energy gap(E_g) value of films at $X=0.5$ a-without doping b- doping 1% by Sn c- doping 3% by Sn d- doping 5% by Sn .



fig(4) the refractive index (n) with ($h\nu$) at at $X=0.5$ a-without doping b- doping 1% by Sn c- doping 3% by Sn d- doping 5% by Sn .



fig(5) the extinction coefficient (K) with (hν) at X=0.5 a-without doping b- doping 1% by Sn c- doping 3% by Sn d- doping 5% by Sn.

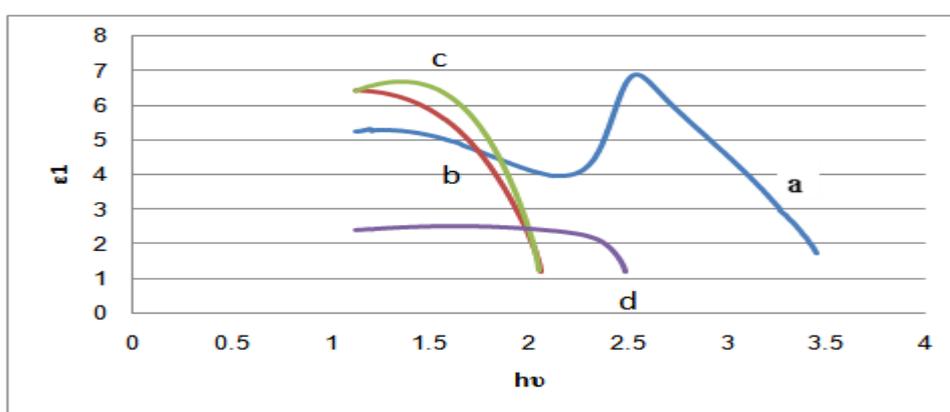


Fig (6) the real part of dielectric constant ϵ_1 with (hν) at X=0.5 a-without doping b- doping 1% by Sn c- doping 3% by Sn d- doping 5% by Sn.

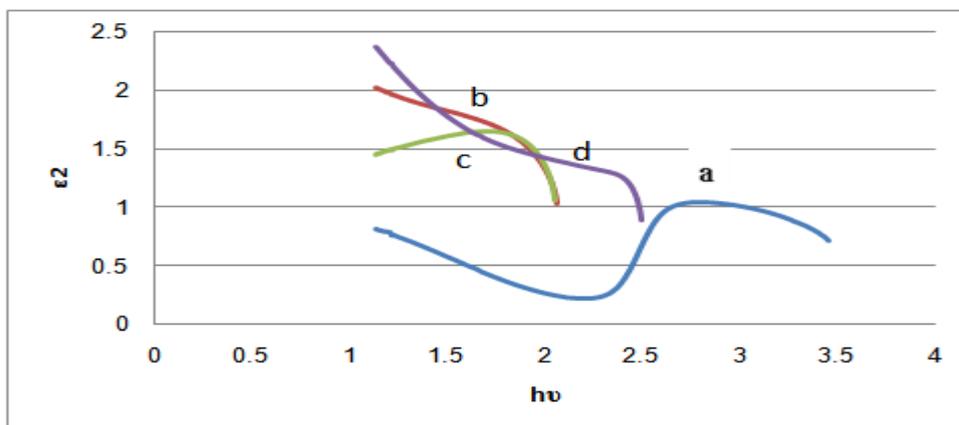


Fig (7) the imaginary part of dielectric constant ϵ_2 with (hν) at X=0.5 a-without doping b- doping 1% by Sn c- doping 3% by Sn d- doping 5% by Sn