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Structural, morphological and optical properties of nanocrystalline cadmium selenide thin films

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ABSTRACT

Nanocrystalline cadmium selenide thin films have been deposited on non-conducting glass substrates. The film samples were characterized by XRD, SEM, UV–vis-NIR reflection/absorption spectroscopy and TEP techniques. The annealed film samples showed a crystalline nature with a cubic crystal structure. The optical analysis showed direct band to band type of transition. The band gap of film sample was found to be in the order of 1.7 eV. The electrical conductivity of the film sample was found to be in the order of 10^{-6} (Ω cm)⁻¹. TEP measurements show n-type of conductivity.

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1. Introduction

Cadmium selenide (CdSe) is one of the most important semiconducting materials from the II–VI group. CdSe has potential application in the area of electronic and opto-electronics devices as a transistor, sensors, lasers and photo electrode [\[1–17\].](#page-2-0)

CdSe thin films can be deposited by various techniques such as spray pyrolysis, electro electro-deposition, vacuum evaporation, thermal evaporation, successive ionic layer adsorption and reaction (SILAR) and chemical bath deposition (CBD) [\[18–23,10,24–26\].](#page-2-0)

We report synthesis of nanocrystalline CdSe thin films by CBD techniques because it is simple, cheap and suitable for large area deposition of thin films. The deposited film samples were characterized by various techniques such as XRD, SEM and optical absorption/reflection spectroscopy. The electric as well as the thermo-electric properties of films are also studied.

2. Experimental details

2.1. Deposition of CdSe thin films

Nanocrystalline CdSe thin films have been deposited first time by using malic acid as a complexing agent. In the deposition of CdSe thin films cadmium sulphate octahydrate and sodium selenosulphate were used as source of Cd²⁺ and Se^{2−} ions, respectively. The Cd^{2+} ions are complexed with malic acid. The complex slowly releases Cd^{2+} ions from the complex, which are used for the slow and homogeneous

deposition of CdSe thin films. For the deposition of film samples, the non-conducting glass plates were used as substrates.

In the deposition of CdSe thin films, 10 ml of 0.25 M Cd²⁺ ions were complexed with malic acid. Ammonia and sodium hydroxide were added in the reaction mixture to maintain the pH of the solution. Then 10 ml 0.25 M sodium selenosulphate was added in above reaction mixture. Then resulting solution was diluted up to 50 ml with distilled water. The pH of the reaction was found to be 7.5. The microglass slides were mounted vertically on a specially designed substrate holder and rotated in the reaction mixture with a speed of 55 ± 2 rpm. As the ionic product of Cd²⁺ and Se^{2−} ions exceeds the solubility product of CdSe, then deposition of CdSe thin films is observed.

The microglass substrates were removed from the reaction mixture after 2.0 h. It is then washed with distilled for several times, dried naturally and kept in a glass desiccator over anhydrous CaCl₂. The resultant films were found to be homogenous, well adherent to glass substrate and orange red in color.

2.2. Characterization of film sample

The thickness of the CdSe thin films was measured by weight difference method. Phillips PW-1710 X-ray diffractometer was used for crystallographic analysis of film sample. The film samples were scanned in the range of 20–80 $^{\circ}$ as 2 θ using Cu K α_1 (wavelength = 2.28970 Å). Cambridge Stereo Scan (USA) scanning electron microscope (SEM) was used for topological analysis of film sample. Before the scanning of film sample, samples were coated with gold–palladium layer by using a polaron SEM sputter unit. The optical absorption was recorded in the wavelength range from 400 to 1400 nm using UV–vis-NIR double beam spectrophotometer (Hitachi-330 Japan) at room temperature. The analysis of spectrum was done by computing the values of absorption at every step of 2 nm. The electrical conductivity of the film samples was carried out in the temperature range of 300–550 K on Zintek-502 BC Milliohm meter by using two-probe method. The thermo-electric power measurements were made by maintaining a temperature gradient along the length of the film and measuring, the potential difference across the terminals having a separation of 1 cm with the help of a digital microvolt meter.

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Fig. 1. XRD spectrums of (a) annealed CdSe thin film at 450 ℃ and (b) as deposited CdSe thin film.

3. Results and discussion

3.1. Growth kinetics

The deposition of CdSe film takes place when the ionic product of Cd2+ and Se2[−] ions exceeds to solubility product of CdSe (Ks_p = 10⁻³³). The growth kinetics of CdSe thin films is given below.

$$
Cd^{2+} + \rightarrow nMA \rightarrow Cd(MA)_n + 2H^+ \tag{1}
$$

where MA = malic acid.

$$
Na_2SeSO_3 + 2^-OH \rightarrow Se^{2-} + Na_2SO_4 + H_2O
$$
 (2)

$$
Cd(MA)n + Se2- + 2H+ \rightarrow CdSe + nMA
$$
 (3)

The color of the deposited CdSe film was found to be orange red. The thickness of the resultant film sample was found to be $0.5 \mu m$.

3.2. Structural analysis

XRD was used for crystallographic analysis of CdSe thin films. Xray spectrums of annealed CdSe film at 450 ◦C and as deposited film sample are shown in Fig. 1. A broad hump peak is observed in the 1 1 1 reflection of as deposited film sample due to the presence of amorphous material. The observed d-values and respective prominent peaks correspond to the reflection from the (1 1 1), (2 2 0) and (3 1 1) planes, which coincide well with the JCPDS data [\[27\]. T](#page-3-0)he results of X-ray analysis well agree with earlier investigators report [\[1–2,16,21–23,28–34\]. X](#page-2-0)-ray diffraction analysis showed that CdSe film sample exhibit in cubic crystal structure.

The lattice parameter 'a' has been calculated by using following equation:

$$
a = d\sqrt{h^2 + k^2 + l^2} \tag{4}
$$

where 'd' is the interplanar distance and 'h, k , l ' are Miller indices of the lattice planes. Crystallographic parameters of CdSe thin films are given in Table 1.

Table 1

Crystallographic parameters of CdSe thin films.

Fig. 2. SEM photograph of annealed CdSe thin film at 450◦C.

The crystallite size (D) in the films has been evaluated by using Scherrer's formula:

$$
D = \frac{K\lambda}{\beta \cos \theta} \tag{5}
$$

where K is constant (0.94), $($ is the wavelength of the X-ray used, $($ is broadening of diffraction line measured at half of its maximum intensity (in rad) and (is Bragg's diffraction angle. The crystallite size of CdSe thin film was found to be 27 nm.

3.3. SEM analysis

SEM was used for surface morphological analysis of film sample. The SEM micrograph of annealed CdSe thin films at 450° C is shown in Fig. 2. The film samples were uniform, pinhole free and well covered to the glass substrate surface. From the micrograph, it is observed that the film composed of minute grains and uniformly distributed over a smooth homogenous background. The grain size of CdSe film by SEM analysis was found to be 28 nm.

3.4. Optical studies

Optical absorbances/reflections of CdSe films were used for the calculation of band gap energy. It is recorded on UV–vis-NIR double beam spectrophotometer at room temperature. All the absorbances of the film samples were measured in the wavelength range of 400–1400 nm. A careful observation of the spectrum shows the presence of a broad absorption edge in the 600–800 nm range. The absorption data were analyzed using the classical relation for near edge optical absorption of semi-conductors. The relation between the absorption coefficient ' α ' and the incident photon energy 'hv' can be given as [\[35\]](#page-3-0)

$$
\alpha = \frac{A(h\nu - E_g)^n}{h\nu} \tag{6}
$$

where 'A' is a constant, ' E_g ' is optical band gap of the material and the exponent 'n' depends upon the type of transition. The values of 'n' for direct allowed, indirect allowed and direct forbidden transitions are $n = 1/2$, 2, and $3/2$, respectively. Optical analysis of film samples

Fig. 3. Plot of $(\alpha h v)^2$ versus $h v$ of CdSe thin film.

Fig. 4. Plot of $log(\sigma)$ versus (1000/T) of CdSe thin film.

showed direct transitions. The band gap (E_g) was determined by plotting a graph of $(\alpha h v)^2$ versus hv. The band gap plot is shown in Fig. 3. The band gap energy ' E_g ' was determined by extra plotting the straight line to the energy axis whose intercept to the x-axis gives the optical band gap. The band gap energy of CdSe film was found to be 1.7 eV which agrees well with earlier investigators results [1–4,36–38].

3.5. Electrical properties

3.5.1. Electric conductivity

The two-point probe method was used to measure electrical conductivity of film sample. All electrical measurements were carried out in the temperature range of 300–550 K. The electrical conductivity of film sample is mainly depends upon structure and composition of material [\[39,40\]. T](#page-3-0)he conductivity of film sample increases with increase in temperature indicates semi-conducting nature of sample. The variation of $\log \sigma$ against (1000/T) for the film sample is shown in Fig. 4. The non-linear nature of the plot is also indicates two types conduction mechanism. The electrical conductivity of the film sample was found to be in the order of 10⁻⁶ (Ω cm)⁻¹ [1,2,21]. The activation energy (E_a) is determined

by using Arrhenius equation which can be given as

$$
\sigma = \sigma_0 \exp^{-E_a/kT} \tag{7}
$$

where σ is measured electrical conductivity, σ_0 is the conductivity pre exponential factor, k is Boltzmann's constant and T is absolute temperature. The ' E_a ' was found to be 0.650 eV at the high temperature and 0.120 eV at the lower temperature region [2,37,38].

3.5.2. Transport number property

Thermo-electric power measurements (TEP) were carried out for the investigation of transport property of film sample. In the TEP measurements the open circuit thermo voltage was generated by the film sample when a temperature gradient was applied across a length of the sample and was measured by using a potentiometer. From the sign of the potentiometer terminal connected at the cold end, one can deduce the sign of the predominant charge carriers. In our investigation of film sample, the negative terminal was connected to the cold end; hence film sample shows n-type of conductivity.

4. Conclusions

- (i) Binary CdSe thin films were grown onto non-conducting glass substrates by using malic acid as a complexing agent.
- (ii) XRD analysis revealed that CdSe film sample exhibit in the cubic crystal structure.
- (iii) The average grain size of the CdSe films was found to be 27–28 nm.
- (iv) The optical absorption study shows direct type of transition and band gap energy was found to be in the order of 1.7 eV.
- (v) The electric conductivity of CdSe film sample was found to be in the order of 10^{-6} ((cm)⁻¹.
- (vi) TEP measurement of the CdSe film indicates n-type of conduction mechanism.

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References

- [1] P.P. Hankare, P.A. Chate, D.J. Sathe, B.V. Jadhav, J. Alloys Compd 503 (2010) 220–223.
- [2] A.S. Khomane, P.P. Hankare, J. Alloys Compd. 489 (2010) 605–608.
- [3] H.E. Esparza-Ponce, J. Hernandez-Borja, A. REyes-Rojas, M. Cervantes-Sanchez, Y.V. Vorbier, R. Ramirez-Bon, J.F. Perez-Robles, J. Gonzalez-Hernandez, Mater. Chem. Phys. 113 (2009) 824–828.
- [4] M. Dhanam, R.R. Prabhu, P.K. Manoj, Mater. Chem. Phys. 107 (2008) 289–296.
- [5] M.A. Hernan dez-Perez, J. Aguilar Hernandez, G. Centreras-Puente, J.R. Vargas-Garcia, E. Rangel-Salinas, Phys. E: Low-dimens. Syst. Nanostruct. 40 (2008) 2535–2539.
- [6] S.M. Pawar, A.G. Moholkar, P.S. Shinde, K.Y. Rajpure, C.H. Bhosale, J. Alloys Compd. 459 (2008) 515–520.
- [7] R. Wang, H. Peter Jakobson, R. Kou, J. Tang, R.Z. Fineman, D. Yu, Y. Lu, Chem. Mater. 18 (18) (2006) 4231.
- [8] G. Zou., H. Ju, Anal. Chem. 76 (23) (2004) 6871.
- [9] D.P. Padiyam, A. Marikani, K.R. Murali, Mater. Chem. Phys. 78 (2002) 51.
- [10] K. Subba Ramaiah, Y.K. Su, S.J. Chang, F.S. Juang, K. Ohdaira, Y. Shiraki, H.P. Liu, I.J. Chen, A.K. Bhatnagar, J. Cryst. Growth 224 (2001) 74.
	- [11] M. Bouroshian, J. Charoud Got, Z. Loizos, N. Spyrellis, G. Maurin, Thin Solid Films 381 (2001) 39.
	- R.M. Abdel-Latif, Phys. B 270 (1999) 366.
	- [13] D. Samantha, B. Samantha, A.K. Chudhuri, S. Ghorai, U. Pal, Semi-Cond. Sci. Technol. 11 (1996) 548.
	- [14] J.W. Edmund, E.M. Dale, R.N. Donald, B.E. Arthur, F.G. John., T.F. Kucch, J. Cryst. Growth 148 (1995) 63.
- [15] T. Gruszecki, B. Holmstrom, Sol. Energy Mater. Sol. Cells 31 (1993) 227.
- [16] K.C. Sathyalatha, S. Uthanna, P. Jayarama Reddy, Thin Solid Films 174 (1989) 233.
- [17] A. Van Calster, A. Vervat, I. De Rycke, J. De Baets, J. Cryst. Growth 86 (1989) 924.
- [18] Yu.V. Melelera, N.A. Redy Chev, G.F. Nevikor, Inorg. Mater. 43 (5) (2007) 455.
- [19] K.N. Shreekanthan, B.V. Rajendra, V.B. Kasturi, G.K. Shivakumar, Cryst. Res. Technol. 38 (1) (2003) 31.
- [20] O. Yomamoto, T. Sasamoto, M. Inagaki, J. Mater. Res. 13 (1998) 3394.
- [21] P.P. Hankare, S.D. Delekar, M.R. Asabe, P.A. Chate, V.M. Bhuse, A.S. Khomane, K.M. Garadkar, B.D. Sarwade, J. Phys. Chem. Solids 67 (2006) 2506–2511.
- [22] P.D. More, G.S. Shahane, L.P. Deshmukh, P.N. Bhosale, Mater. Chem. Phys. 80 (2003) 48.
- [23] R.C. Kainthila, D.K. Pandya, K.L. Chopra, J. Electrochem. Soc. (1980) 277.
- [24] C.D. Lokhande, E.H. Lee, K.D. Jung, O.S. Joo, Mater. Chem. Phys. 91 (2005) 399.
- [25] S.M. Rashwan, S.M. Abdul-Wahab, M.M. Mohammed, J. Mater. Sci. Mater. Electron. 1818 (2007) 575.
- [26] T. Elango, V. Subramanium, K.R. Murali, Surf. Coat. Technol. 123 (2000) 8.
- [27] JCPDS Data File No. 00-019-0191.
- [28] S. Evat, H. Metin, M. Ari, Mater. Chem. Phys. III (2008) 114–120.
- [29] R.B. Kale, C.D. Lokhande, Semi-Cond. Sci. Technol. 20 (2005) 1–9.
- [30] M.E. Hernandez Torres, R.S. Gonzates, H.N. Contreras, M.A. Vidal, J.M. Gracia, Mod. Phys. Lett. B 15–19 (2001) 741.
- [31] P. Nemec, D. Mikes, J. Rohorec, E. Uhllirova, F. Trojanek, P. Maly, Mater. Sci. Eng. B 69–70 (2000) 500.
- [32] H. Padmanabha Sarma, N. Rangarajan, K.R. Murali, Phys. Status Solidi (A) 148 (1995) K.77.
- [33] S. Gorer, G. Hodes, J. Phys. Chem. 98 (1994) 5338.
- [34] G. Hodes, A.A. Yaran, F. Decker, P. Motisuke, Phys. Rev. B 36 (1987) 4215.
- [35] D.S. Sutrave, G.S. Shahane, V.B. Patil, L.P. Deshmukh, Mater. Chem. Phys. 65
- (2000) 298. [36] C.S. Shahane, D.S. Sutrve, L.P. Deshmukh, Ind. J. Pure Appl. Phys. 34 (1996) 153–157.
- [37] P.P. Hankare, A.D. Jadhav, V.M. Bhuse, A.S. Khomane, K.M. Gradkar, Mater. Chem. Phys. 80 (2003) 102.
- [38] P.P. Hankare, V.M. Bhuse, K.M. Garadkar, S.D. Delekar, I.S. Mulla, Semi-Cond. Sci. Technol. 19 (2004) 70–75.
- [39] K.L. Chopra, Thin Film Phenomenon, McGraw-Hill, New York, 1969.
- [40] C.N. Rao, Modern Aspects of Solid State Chemistry, Plenum Press, New York, 1970, p. 531.