

Accepted July 22, 2016

; Published July 30, 2016

Citation: Rani S, Shanthi J, Kashif M, Ayeshamariam A, Jayachandran M (2016) Studies on Different Doped Zn Concentrations of CdSe Thin Films. J Powder Metallurgy 43. © 2016 Rani S, et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Keywords: Electron beam evaporation; Doping effect; XRD of CdSe; PEC solar cells

Introduction

The formation of solid solutions in semiconductors is of great importance in the field of materials science and technology. The formation of solid solutions in semiconductors is of great importance in the field of materials science and technology.

ZnS, CdTe, CuInSe₂ or CuInTe₂, in films of these materials are usually synthesized by chemical methods [1], epitaxy [2], electron beam evaporation [3] and sputtering [4].

Feng Ying et al. reported that few-layer graphene was prepared from highly oriented pyrolytic graphite by micromechanical cleavage

area (0.25 cm²) of CdSe:Zn films on conducting coated glass substrates.

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Zn doped CdSe films were deposited using electron beam evaporation (EBE) technique with optimized substrate temperature 100°C. Different concentrations of zinc (10%, 20% and 30%) were introduced into the CdSe matrix. Figure 1 shows XRD spectra of CdSe:Zn films with Zn concentrations 10%, 20% and 30% deposited at 100°C, respectively. All peaks are well defined and sharp peak

CdSe:Zn films might be due to the presence of high energy gap materials. It is attributed to excess of Cd in the composition over Se, which (ZnSe) than that of undoped CdSe. The variation of n shows a sub-linear behaviour, a lowering phenomenon in the alloy films of CdSe:Zn calculated carrier concentration values are 7.0×10^{18} , 1.4×10^{19} and $3.4 \times 10^{19} \text{ cm}^{-3}$ for CdSe:Zn films with zinc content 10%, 20% and 30% respectively are shown in Figures 6-8. The absorption edge shows a blue-shift with increasing 'Zn' content due to the increase in band gap value. The continuous variation of Zn concentrations confirms the solid solution formation between CdSe and ZnSe so that their optical and electronic properties can be tailored according to the need for device developments.

The electrical resistivity and the interface behaviour of semiconductor thin films with solid solutions are vital in photo-electrochemical solar cell applications. At room temperature the observed resistivity of CdSe:Zn films with 10%, 20% and 30% Zn content was 8.2×10^{-3} , 1.5×10^{-3} and $3.5 \times 10^{-3} \text{ ohm cm}$ respectively. The increase in resistivity is attributed to the fact that band gap increases with introduction of more zinc into the lattice of CdSe. The higher resistivity CdSe:Zn films finds an application as a buffer layer for solid state solar

Figure 9a shows XPS spectra of CdSe:Zn film with zinc content 20% deposited at 100°C. It shows the presence of Cd 3d, Zn 2p and Se 3s peaks confirming the formation of Zn doped CdSe film. Figure 9b-9d shows the narrow spectra of Cd 3d, Zn 2p and Se 3s present in the film. The spectral shift along with the abscissa due to the sample charging is eliminated by moving the measured C 1s peak to the binding energy value of 284.8 eV [17]. In Figure 9b, binding energies of Cd 3d_{5/2} and Cd 3d_{3/2} are observed at 405.14 and 411.74 eV which confirms the presence of Cd in the film. The Se 3s peak is observed at 55.0 eV. The Se 3s peak is assigned to the Se atoms in the lattice.

prepared by the EB evaporation technique which is confirmed by the XRD patterns. The XRD patterns show the formation of linear peaks, which are given under each 2D AFM pictures. Figure 9a-9c shows the elemental analysis of CdSe:Zn films with zinc content 10%, 20% and 30% respectively. It is found that ratios of Cd and Zn elements are in fewer amounts than the percentage present in the initial material, but selenium content is always present in stoichiometric percentage. This is attributed to the fact that the CdSe is first, and then reaches the substrate over which ZnSe gets deposited when they are grown by EB evaporation technique [19,20].

It is well observed that zinc atoms replace cadmium atoms in the crystal lattice and size of the grain is decreased with increasing the Zn contents. It might be caused by the addition of CdSe to ZnSe that the Photoluminescence properties depend on the lateral size of

a pseudo-smooth interface applies when the exciton size is larger. This results in a relatively narrow single PL line, because the exciton properties are averaged over a number of defects. Ternary alloy semiconductor films have provided the possibility of making their band gap (E_g) adjustable in the visible wavelength range and it is therefore useful for optoelectronic applications and devices as laser diodes and light emitting devices operating in the spectral region from red to green [21]. For such applications, a good PL efficiency is required for the CdSe:Zn system to assess them for optoelectronic applications.

Figure 12a-12c shows the PL emission spectra recorded at room temperature. PL peaks are observed at 622, 519 and 482 nm while E_g

