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 $K_{\rm COMM}$: Electron beam evaporation; Doping e ect; XRD of CdSe; PEC solar cells

e formation of solid solutions in semiconductors is of great

₂S, CdTe, CulnSe or CulnTe, in lms of these materials are usually synthesized blyermidtactritethoda[55].epitaxy [3], electron beam evaporation [4] and

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

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area (0.25 cm of CdSe:Zn Ims on conducting coated glass substrates.

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

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CdSe:Zn Ims might be due to the presence of high energy gap materizells. It is attributed to excess of Cd in the composition over Se, which (ZnSe) than that of undoped CdSe. e variation of shows a sublinear behaviour, a lowering phenomenon in the alloy Ims of CdSe:Zre calculated carrier concentration values are 7.0 x⁸,15.4 × 10⁶ and [15]. e as-prepared Ims exhibited direct band-gap transition and 3.4 × 10⁶ cm³ for CdSe:Zn Ims with zinc content 10%, 20% and 30% the absorption edge shows a blue-shi with increasing 'Zn' content durespectively are shown in Figures 6-8. to the increase in band gap value. e continuous variation of Meth Zn concentrations con rms the solid solution formation between CdSe and ZnSe so that their optical and electronic properties can be tailored posited at 100°C. It shows the presence of Cd '3d, Zn '2p and Se '3 peaks con rming the formation of Zn doped CdSe Im. Figure 9b-9d shows the narrow spectra of Cd 3d, Zn 2p and Se 3d present in the Im. e electrical resistivity and the interface behaviour of e spectral shi along with the abscissa due to the sample charging is semiconductor thin Ims with solid solutions are vital in photo- eliminated by moving the measured C 1s peak to the binding energy electrochemical solar cell applications. At room temperature thealue of 284.8 eV [17]. In Figure 9b, binding energies of Cd and 30% Zmd_{3/2} are observed at 405.14 and 411.74 eV which con rms the presence of beaks on the formation is semiconductor to a stributed to fact that band gap increases wistFi onic 3d

introduction of more zinc into the lattice of CdSe. e higher resistivity

CdSe:Zn Ims nds an application as a bu er layer for solid state solar

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prepared by the EB evaporation technique which is con rmed by the educe the dimension of the lattice leading to the formation of linear line pro le curves given under each 2D AFM pictures. Figure 9a-9grains with reduced grain size. It shows the compactness, pinhole free shows the elemental analysis of CdSe:Zn Ims with zinc content 10% and well adherent nature of these Ims on glass substrates which are 20% and 30% respectively. It is found that ratios of Cd and Zn elementery useful for PEC devices. is is attributed to the fact that the CdSe are in fewer amounts than the percentage present in the initial material consequently sublimation takes place rst, and then reaches the but selenium content is always present in stoichiometric percentage.substrate over which ZnSe gets deposited when they are grown by EE

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

a pseudo-smooth interface applies when the exciton size is larger. is results in a relatively narrow single PL line, because the exciton properties are averaged over a number of defects. Ternary alloy semiconductor Ims has provided 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 e ciency 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 peak are observed at 622, 519 and 482 nm while E

zinc concentration which is attributed to the increase in band-gapcaused by ZnSe composition. Each PL spectrum is characterized by its near band edge (NBE) recombination as suggested for the CdS Ims prepared by pulsed laser deposition technique [22]. PL peak attained at 5K for CdZn_{0.77}Se single quantum well (QW) at about 2.77 eV with 0.4 ML (monolayer) and at 2.73 eV with 1.15 ML [23]. Photoluminescence results con rm solid solution CdSe:Zn Ims deposited by EB evaporation technique can e ciently be used for PEC solar cells and light emitting diode studies.

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Using EB evaporation technique CdSe:Zn thin Ims were prepared on glass substrates at 100°C with Zn concentration of 10%, 20% and 30%, respectively. Nanocrystalline nature of as-prepared lms was characterized using XRD analysis. Optical studies were conducted to record absorbance and transmittance spectra of CdSe:Zn Ims. e absorbance curves, band gap values calculated from² (Vs)(h) curves showed direct band gap nature of all the Ims with band gap values varied from 1.90 eV to 2.42 eV, is con rmed solid solution formation between CdSe and ZnSe compounds and the lattice modi cation is e ected by zinc incorporation. Elemental analysis was carried out by EDAX and XPS. e surface morphology explored by SEM analysis showed highly ne granular grain formation distributed uniformly over the entire surface. e in uence of the variation of zinc content can be explicitly seen by the changing surface grain patterns from AFM studies. Photoluminescence (PL) technique is used to assess for material application to certify that CdSe:Zn Ims are good for the fabrication of PEC cell.

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