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Synthesis and characterization of arrested precipitation assembled Cu₂Se thin films

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Abstract---Nanostructured thin films of copper selenide (Cu₂Se) deposited by arrested precipitation technique (APT) with controlled release of ions at suitable pH=10.5. The deposited Cu₂Se films were characterized for optical, structural, and morphological properties. Optical absorption study indicates that the band gap energy (2.26 eV) of material and shows visible radiation absorption. XRD show that APT is a favorable technique for the synthesis of pure nanocrystalline Cu₂Se thin films with monoclinic crystal structure. An electrochemical impedance spectroscopy analysis confirms the charge transfer resistance. X-ray photoelectron spectroscopy reveals stoichiometry at valance state of Cu₂Se.

Keywords---arrested precipitation technique, thin films, copper selenide, optical properties.

Introduction

Nanostructured thin film solar cells are a favorable and perhaps emergent technology these days [1]. Stoichiometric nanostructured inorganic semiconductor coatings will be superior options for engineering solar cells and limiting the depletion of nonrenewable energy supplies such as fossil fuels, natural gas, and coal oils [2]. Copper-containing chalcogenides compound semiconductors have so concentrated on ultra-thin absorber layers [3], counter electrodes such as Cu_2S [4] and CuSe [5]. Metal chalcogenide thin films are easy to deposit on substrates like glass and plastic, making them ideal for solar building integration as optoelectronic devices. The IB-VIA and II-VIA group chalcogenides are becoming more attractive due to their strong absorption coefficient in visible light and electron-hole separations during solar cell operation [6]. Copper selenide (Cu_2Se , 2.2 eV) [7] is semiconducting and visible radiation active [8-13]. However, they lack generality or need complex equipment, and they are problematic due to the use of hazardous reducing agents, surfactants, solvents, and high temperatures in the synthesis of copper chalcogenide compounds [10]. As a result, solution-processed synthesis utilising the APT approach is easy and cost-effective [11], with structural, compositional, and optoelectrical characteristics equivalent to those of other advanced deposition methods [12].

We used triethanolamine (TEA) as a polydentate ligand in this study, which forms a complex with copper metal ions to prevent bulk precipitation. The goal of this report's innovation is to improve the growth mechanism conditions, particularly the reduction of copper metal ions owing to mild reducing agents (Na_2SO_3), such that Cu_2Se thin films have a homogeneous, highly adherent nanogranular surface shape at room temperature. APT is a chemosynthesis of ternary nanoalloys using a hybrid chemical bath deposition approach in tandem with the controlled chemical growth process (CCGP) [11]. In inorganic metal chalcogenide coating thin films, smaller crystallites serve as nutrition to the directed bigger developing crystallites [14, 15]. Although the possible growth and response mechanisms for Cu_2Se thin film production were briefly explored.

Experimental Details

Materials and Method

Without additional purification, all of the compounds employed were analytical reagent (AR) grade. Precursors included copper sulphate pentahydrate ($\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$), selenium metal powder, sodium sulfite (Na_2SO_3), ammonia solution (NH_3), and triethanolamine ($\text{N}(\text{CH}_2\text{CH}_2\text{OH})_3$). We created a sulfide/polysulfide redox mediator with sodium sulphide (Na_2S), sodium hydroxide pellets (NaOH), and sulphur powder. The substrates were cleaned using a chromic acid solution heated to 500°C for 5 minutes before being ultrasonically cleaned with a combination of isopropanol and deionised water. Cu_2Se thin films were produced on naked glass substrates using a simple APT chemosynthesis method.

Synthesis of Cu₂Se thin films

Simple chemical processes were the focus of a recent synthetic route. To create the clear translucent solution, 0.05 M copper sulphate solutions were employed to form a complex with triethanolamine (TEA) as a polydentate ligand. At alkaline pH=10.5, the Cu²⁺-(TEA)₂ complex interacts with the Se²⁻ chalcogen ions, releasing slowly by dissociation of Sodium selenosulfite (Na₂SeSO₃). Refluxing selenium metal powder with anhydrous sodium sulfite (Na₂SO₃) at 70 °C for 10 hours yielded Na₂SeSO₃. The nucleation and Ostwald's ripening mechanisms were used to deposit Cu₂Se thin films. Different preparative factors such as ligand concentration, Cu²⁺ precursor concentration, pH, deposition period, single-solution phase, and bath temperature are mostly responsible for these mechanisms. As a result, all of these parameters were optimised at the thin film deposition initiative stage. The precursor concentration of 0.05 M Cu²⁺ was kept constant, as was the deposition duration at ambient temperature (RPM). A surface profiler was used to measure the thickness of the Cu₂Se films after they were formed. After the depositions are completed, the substrate peels away from the overgrowth surface, with no increase in film thickness. All of the deposited thin films were homogenous, highly adhering translucent blackish tinted at the terminal development phase.

Characterizations

A UV-Vis-NIR spectrophotometer was used to capture optical absorption spectra (Shimadzu, UV-1800). An X-ray diffractometer (Bruker AXS, D8) was used to validate structural characteristics using Cu K α ($\lambda = 1.5418 \text{ \AA}$) radiation for 2θ from 20° to 80°. Field-emission scanning electron microscopy (FE-SEM) and X-ray photoelectron spectroscopy (XPS, Thermo Scientific, Multilab-2000) with a multi-channel detector that can withstand high photon energies from 0.1 to 3.0 keV were used to demonstrate the surface morphology and elemental composition of the deposited thin films. The auto lab PGSTAT 100 FRA 32 was used to conduct the EIS.

Results and Discussion

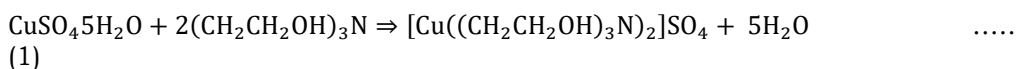
Thin films formation and its growth mechanism

It may be feasible to analyze attributes of deposited material with chosen morphology and crystallinity in order to determine how shape evolution processes are predictable. Cu²⁺ metal ions are complexed utilizing complexation (TEA) and released slowly at the optimal precursor concentration, temperature, and pH in the current process. By raising the concentration of OH⁻ ions and the proportionate rise in chalcogen ions Se²⁻ releasing ability in the reaction solution, aqueous NH₃ is employed to adjust the pH [11]. However, when the temperature rises, the rate of metal ions released rises, and the reaction accelerates, resulting in bulk precipitation rather than the desired quality thin film production. As a result, we must first improve depositions at room temperature (300 K), Cu²⁺ precursor concentrations of 0.05 M, and pH of 10.5. Cu₂Se thin films of exceptional quality and uniformity were synthesized. APT approaches are based on nucleation and the Ostwald ripening mechanism. When the ionic product (K_p)

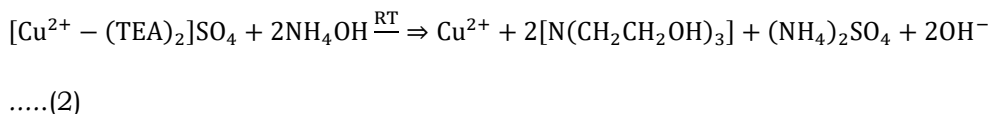
of the metal ions exceeds the solubility product (K_{sp}), thin films are formed. Following the ion-by-ion condensation of metal ions onto the substrate surface, the nucleation process begins [11]. A delayed reaction rate produces good stochastic compositions with high-quality and adhesive thin films in general [16]. As a result, as compared to traditional chemical bath methods, the hybrid created arrested precipitation methodology is more effective.

Reaction mechanism of Cu₂Se thin film formation

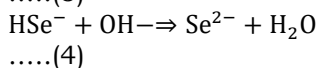
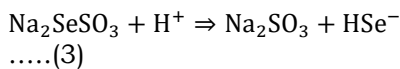
Solution of Cu²⁺ ions are complexes with Triethanolamine (TEA) as complexion (1)



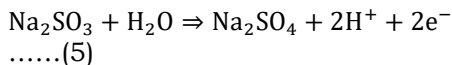
With addition of basic solution of 30 % liquor NH₃, the Cu²⁺ ions released due to the dissociation of metal complex bonds at alkaline pH = 10.5 as explained in reaction (2)



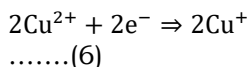
Sodium selenosulfite dissociates into sodium sulfite and highly active HSe⁻ species at the same pH and ambient temperature. By absorbing protons in alkaline media reactions (3), this highly active species swiftly transforms into reactive Se²⁻ ions (4) [17]



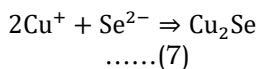
The sodium sulfites released are actions as mild reducing agents and forms the sodium sulfate is shown in eqn. (5) [18].



The ionic product of the Cu²⁺ and Se²⁻ ions surpasses the solubility product in the reaction bath during the creation of Cu₂Se thin films, resulting in a gradual ion-by-ion condensation of the metal and chalcogen ions according to the Ostwald ripening law. The moderate reducing agents Na₂SO₃ make the condensed Cu²⁺ metal ions more reactive and easy to reduce, as illustrated in reactions (6) [18].



Finally the free anions combined with cations to form Cu₂Se are summarized in below as (7)



Optical studies

The absorption spectra of UV-Vis-NIR (ultraviolet-visible-near infrared) in the wavelength range 300–900 nm were shown in Fig. 1. The Cu_2Se thin films' optical absorption spectra were obtained and clearly demonstrated that the highest visible light absorption occurs between 400 and 550 nm. All of the samples had an absorption coefficient on the order of 10^5 cm^{-1} . The optical band gap energy may be calculated using electron excitation from the valance band to the conduction band. The term "optical data" refers to the use of the equation (8)

$$\alpha = \frac{A(h\nu - E_g)^n}{h\nu} \quad \dots\dots(8)$$

When A denoted as a transition probability parameter, h denoted as the Planck constant, E_g denoted as the material's optical band gap energy, and the exponent (n) is dependent on the kind of transition. 1/2, 2, 3/2, and 3 are the values of (n) for direct allowed, indirect allowed, direct prohibited, and indirect forbidden transitions, respectively. The optical band gap energy values of Cu_2Se thin films were calculated using the graph of $(ah\nu)^2$ versus $(h\nu)$ ((eV/cm)² vs (eV)), as shown in the inset of Fig. 1.

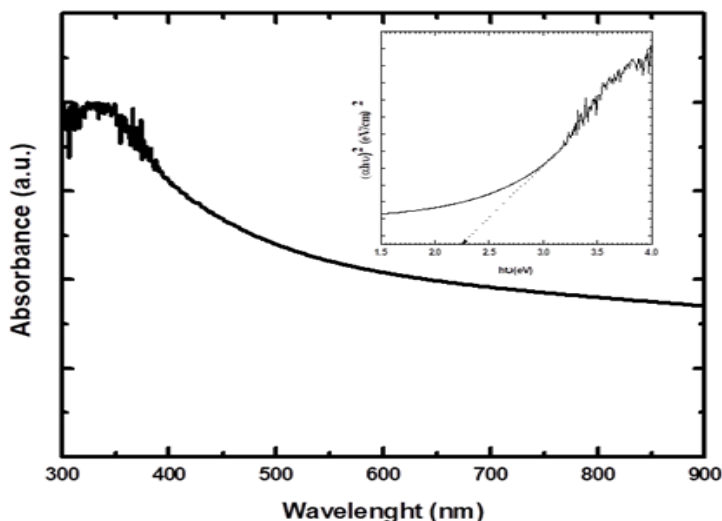


Fig. 1. Optical absorption spectra of the Cu_2Se thin films, inset is plot of $(ah\nu)^2$ vs $(h\nu)$.

Structural studies

X-ray diffraction measurements in the range 2θ of 20° to 70° were used to record the crystal structure analysis of the sample. The Cu_2Se thin film's XRD pattern is shown in Figure 2. The Cu_2Se thin films are nanocrystalline by nature, as evidenced by the XRD pattern, which exhibits both wide and strong peaks [19].

The primary diffraction peaks were concentrated at $2\theta = 27.44^\circ$, 43.82° , 51.34° , 51.81° , and 54.44° , which corresponded to the monoclinic crystal system's (111), (220), (311), (311), and (220) planes.

Peaks emerge at same sites in a typical XRD examination of Cu_2Se thin films, with changes to the intensity and expansion of the diffraction peaks, indicating the development of pure phase materials with increased crystallinity. Some of the minor intensity peaks demonstrate the amorphous character of the films due to the presence of glass beads in the materials. Furthermore, no further peaks are seen, indicating that the substance is in its pure phase. Strain or micro-strain might explain the observed widening of the diffraction peaks. The crystallite size was also determined using the Debye Scherer formula equation (9) [19].

$$D = \frac{0.9\lambda}{\beta \cos\theta} \quad \dots\dots(9)$$

where D denotes the crystallite size, $\lambda=1.5406$ is the X-ray wavelength, FWHM denotes the radians is the full-width-at-half-maximum (FWHM), and θ denotes the Bragg's angle and the crystallite size calculated is 24 nm.

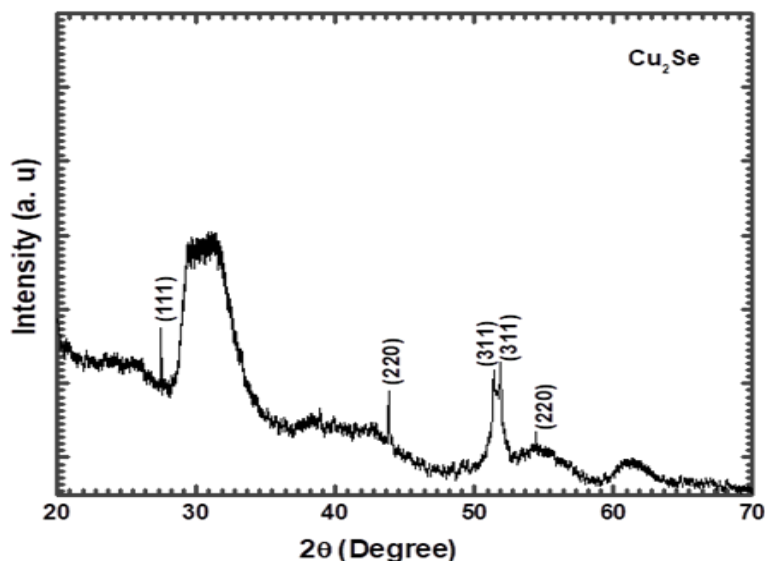


Figure 2: XRD pattern of the Cu_2Se thin film.

Electrochemical impedance spectroscopy (EIS)

The Cu_2Se thin film's PEC kinetics were investigated using EIS. EIS measurements were carried out in the dark using a 0.2 M polysulfide mediator and a 0.5 V forward bias voltage. The ac frequency range that was measured was 0.1 Hz to 1000 Hz. The Nyquist plot of Cu_2Se thin film is shown in Figure 3. The solution resistance ($R_s = 25 \Omega \text{ cm}^{-2}$) R_1 is the charge transfer resistance between the Pt-FTO counter/electrolyte interface, whereas R_2 is the charge transfer resistance due to photo-excited electron hole recombination at Cu_2Se /mediator.

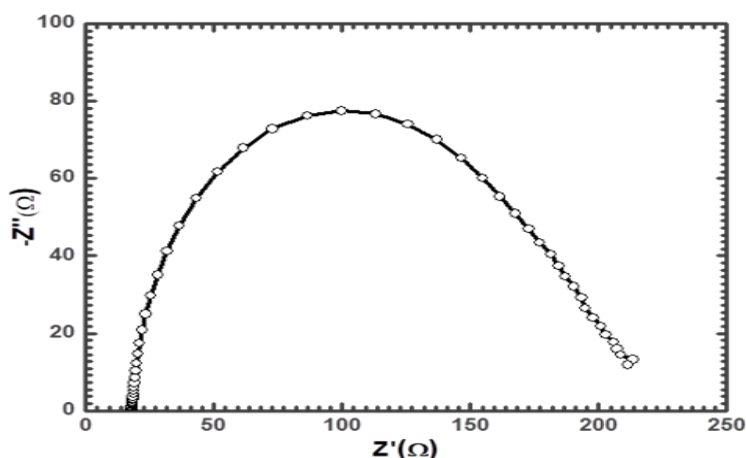


Figure 3: Nyquist plot of Cu_2Se thin film.

Morphological Studies

The surface morphology of Cu_2Se thin films may be seen using field-emission scanning electron microscopy (FE-SEM). FE-SEM micrographs of Cu_2Se thin films at low and high magnification are shown in Fig. 4. FE-SEM micrographs revealed that nanoflakes were evenly distributed over the whole surface. At lower precursor concentrations, a beaded nanoflakes-like morphology was observed. Cu_2Se materials with tiny grain sizes develop at a lower concentration and are distributed throughout the whole substrate surface. The grain size of these nanoflakes-like materials is less than ~ 40 nm.

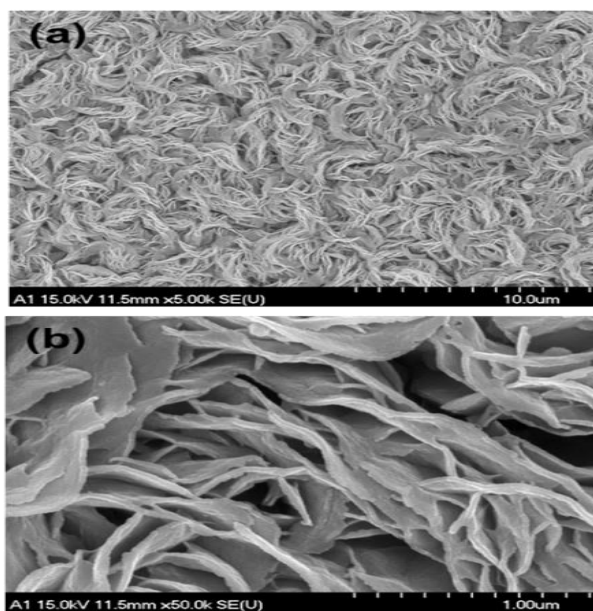


Figure 4: Field-emission scanning electron microscopy (FE-SEM) (a) low and (b) high magnification images of Cu_2Se .

Compositional Analysis

In order to check the surface elemental composition of the Cu_2Se thin film. The XPS survey and high resolution core level spectrum demonstrated in Fig. 5. The two distinct peaks confirm the presence of Copper and Selenium elements respectively. Additionally, elements valance state and the existing form of the Cu_2Se thin film was investigated by XPS study as shown in Fig. 5 (a), peaks along with Carbon and Oxygen in the sample.

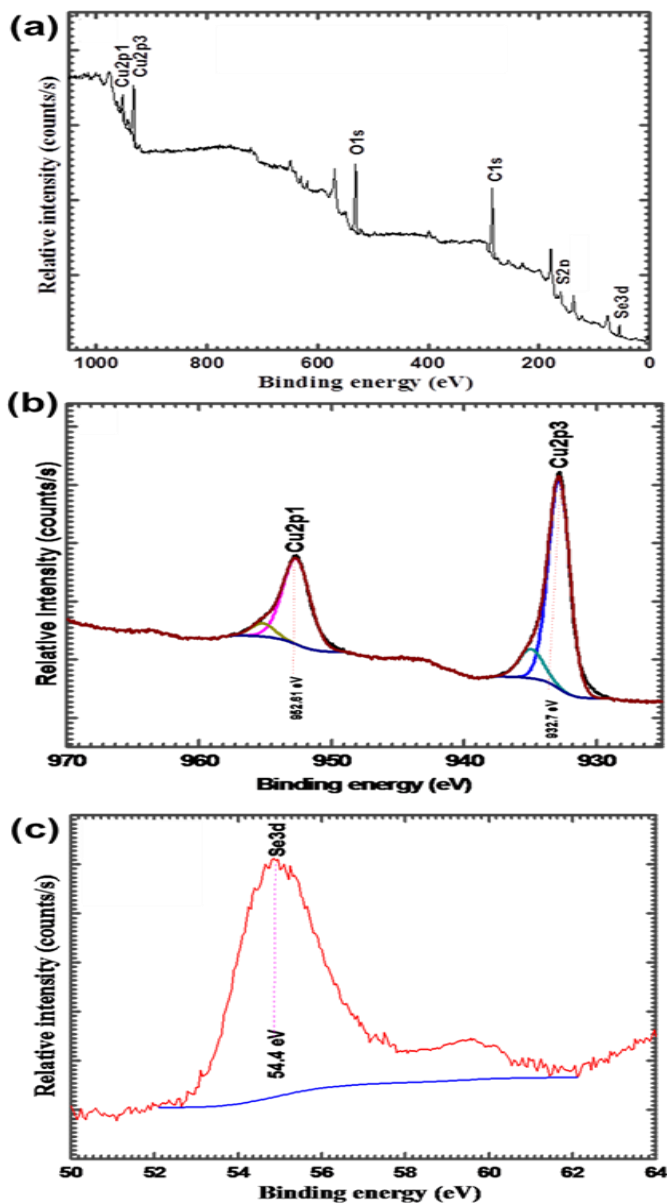


Figure 5: (a) XPS survey spectrum of Cu_2Se thin film, (b) high-resolution core level XPS spectrum of Cu, (c) high- resolution core level XPS spectrum of Se.

Copper's spectrum at its most fundamental level Cu2p peaks can be readily recognised in Fig.5 (b), with two peaks ascribed to Cu2p_{3/2} and Cu2p_{1/2} with binding energies of 932.7 eV and 952.61 eV, respectively, indicating the existence of Cu⁺ in the sample. Cu2p_{3/2} and Cu2p_{1/2} were also assigned two peaks with binding energies of 932.7 eV and 952.61 eV, respectively, indicating the existence of Cu⁺ with a splitting energy of 19.91 eV [7] [20]. The presence of Se²⁻ is verified by a significant peak at 54.4 eV ascribed to the Se3d transition in Fig. 5 (c). For the Cu⁺ and Se²⁻ states, the binding energies for Cu and Se are at their corresponding locations. It denotes the presence of Cu⁺ and Se²⁻. It means that Cu⁺, and Se²⁻ exists in the stochastic formula of deposited Cu₂Se thin film.

Conclusion

All of the above discussion and analysis indicate that nanoflakes Cu₂Se thin films were effectively created for the first time employing the single-solution phase arrested precipitation approach. Cu₂Se thin film has control over directed crystal formation due to vital function including TEA, the band gap energy 2.26 eV, according to opto-structural and morphological research. The crystalline nature of the sample is shown by the XRD pattern, which reveals a pure monoclinic crystal structure that governs the ion insertion kinetics leading to a charge transfer process. XPS confirms the stochastic Cu₂Se thin film at chemical valance state.

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