



## Fabrication of CZTS thin film on flexible Cu-foil substrate by two-stage process

### İki aşamalı yöntem ile CZTS ince filmlerinin esnek Cu-folyo altlık üzerine büyütülmesi

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#### Abstract

In this research, CZTS thin films were grown on flexible Cu-foil substrates with varying sulfurization times. Distinct characterization methods were employed, including X-ray diffraction (XRD), Raman spectroscopy, Energy-Dispersive X-ray Spectroscopy (EDX), Scanning Electron Microscopy (SEM), optical transmission, and Photoluminescence (PL) measurements. Distinctive diffraction peaks characteristic of the kesterite CZTS phase were observed in the XRD analysis, occurring around at  $2\theta = 28.45^\circ$  (112),  $47^\circ$  (220/204), and  $56^\circ$  (312/116). Additionally, some secondary phases such as  $\text{Cu}_2\text{S}$  and  $\text{SnS}$  were identified. Raman spectroscopy confirmed the presence of the kesterite CZTS phase, with a prominent peak detected at approximately  $\sim 336 \text{ cm}^{-1}$ , attributed to sulfur atom vibrations within the kesterite structure. Apart from CZTS structure, minor peaks suggesting the presence of the  $\text{Cu}_2\text{SnS}_3$  (CTS) phase was detected. EDX analysis revealed compositions with Cu-poor content and Zn-rich content across all samples, with slight variations in sulfurization dwell times affecting the chemical composition. SEM imaging at different magnifications showed alterations in surface morphology and grain structures. Films sulfurized for 30 s and 60 s displayed a granular structure morphology, while extending the dwell time to 120 s resulted in a more compact surface morphology. Optical band gap values ranged between 1.57 and 1.60 eV. PL measurements consistently exhibited strong PL emission around 1.25 eV for all samples, attributed to various transitions within the band structure of CZTS film. The absence of observable band-to-band transitions in the PL measurements indicated the presence of intrinsic defect levels and recombination centers within CZTS. Overall, it was demonstrated in this study that CZTS thin films can be produced on flexible Cu-foils with short sulfurization times, thereby expanding the application areas of CZTS thin-film solar cells.

**Keywords:** CZTS thin film, Cu-foil, Sputtering, RTP, Short sulfurization time

#### 1 Introduction

The CdTe and CIGS based thin film solar cells have already reached conversion efficiency above 20%, which is

#### Öz

Bu araştırmada, farklı sülfürleme süreleri ile esnek Cu-folyo üzerine büyütülmüş CZTS ince filmler incelendi. X-ışını kırınımı (XRD), Raman spektroskopisi, Enerji Dağılımlı X-ışını spektroskopisi (EDX), Taramalı Elektron Mikroskopisi (SEM), optik geçirgenlik ve fotoluminesans (PL) ölçümleri de dahil olmak üzere çeşitli karakterizasyon yöntemleri kullanıldı. XRD analizinde kesterit CZTS fazına özgü belirgin kırınım pikleri  $2\theta = 28.45^\circ$  (112),  $47^\circ$  (220/204) ve  $56^\circ$  (312/116) civarında meydana gelen pikler gözlemlendi. Ayrıca,  $\text{Cu}_2\text{S}$  ve  $\text{SnS}$  gibi bazı ikincil fazlar tespit edildi. Raman spektroskopisi, kesterit CZTS fazının varlığını doğruladı ve kesterit yapısı içindeki kükürt atomu titreşimlerine atfedilen yaklaşık  $\sim 336 \text{ cm}^{-1}$ 'de belirgin bir pik tespit edildi. CZTS yapısının yanı sıra,  $\text{Cu}_2\text{SnS}_3$  (CTS) fazının varlığını işaret eden az sayıda pik de tespit edildi. EDX analizi, tüm örneklerin Cu-fakiri ve Zn-zengini kompozisyona sahip olduğunu göstermiştir. Farklı büyütme oranlarında yapılan SEM görüntülemeleri yüzey morfolojisinde ve tane yapılarında değişiklikler olduğunu gösterdi. 30 ve 60 saniye süreyle sülfürlenmiş filmler taneli bir yapı morfolojisi sergilerken, bekleme süresini 120 saniyeye uzatmak daha kompakt bir yüzey morfolojisine katkı sağladığı tespit edilmiştir. Optik bant aralığı değerleri 1.57 ile 1.60 eV arasında değişkenlik göstermiştir. PL ölçümleri, tüm örneklerde 1.25 eV civarında güçlü bir PL emisyonu sergiledi ve bu, CZTS yapısı içindeki çeşitli geçişlere atfedildi. PL ölçümlerinde gözlemlenen banttan banta geçişlerinin olmaması, CZTS içindeki özden kusur seviyeleri ve rekombinasyon merkezlerinin varlığını işaret ettiği belirlenmiştir. Genel olarak, bu çalışmada CZTS ince filmlerin kısa sülfürleme süreleriyle esnek Cu-folyolar üzerine üretilebileceği ve bu sayede CZTS ince film güneş pillerinin uygulama alanlarının genişletilebileceği gösterilmiştir.

**Anahtar kelimeler:** CZTS ince film, Cu-folyo, Saçtırma, RTP, Kısa sülfürleme süresi

competitive with Si-based solar cells. However, toxicity of cadmium (Cd) and the limited availability of indium (In) and gallium (Ga) pose constraints on these thin film solar cells. In this context,  $\text{Cu}_2\text{ZnSnS}_4$  (CZTS) thin film has emerged as

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a highly promising absorber layer for photovoltaic industry (PV). It possesses several advantageous properties including the utilization of earth-abundant, environmental-friendly and cost-effective raw materials. Furthermore, CZTS exhibits a direct optical band gap of approximately 1.5 eV, which can be tuned to 1.0 eV by incorporation of Se into structure [1]. CZTS also demonstrates a high optical absorption coefficient ( $>10^4 \text{ cm}^{-1}$ ) in the visible range and exhibits p-type conductivity [2]. Moreover, CZTS boasts a theoretically high efficiency of approximately 30% according to the theoretical limit [3]. However, the current record conversion efficiency of CZTS stands at around 14% when fabricated on rigid glass substrates [4]. This indicates that there is considerable gap for improvement in enhancing the performance of CZTS as a thin film material for solar cells.

There exist various methods for the growth of CZTS thin films, categorized as physical (vacuum-based) and chemical (non-vacuum-based) growth methods. Physical methods include sputtering techniques such as DC and RF sputtering [5-7], thermal evaporation [8], pulsed laser deposition [9], e-beam evaporation [10], and etc. Chemical methods, on the other hand, encompass spin-coating [11], dip-coating [12], spray pyrolysis [13], electro-deposition [14], and similar approaches. Among these methods, sputtering is widely employed due to its ability to provide homogenous films with controlled thickness and the capability for large-scale production.

There are numerous strategies have been explored to optimize the structural, compositional, optical properties and also electrical characteristics of CZTS-based thin films to enhance conversion efficiency. These approaches involve the manipulation of various factors such as growth parameters (pressure, deposition rate, etc.) [15], chemical compositions (Cu-poor/Zn-rich or Cu-rich/Zn-rich) [16], annealing parameters (temperature, heating rate, dwell time, chalcogen source, etc.) [17], incorporation of different doping atoms (Al, In, Cd, Li, Ge, Ag, Na, K, etc.) [18-20], utilization of diverse back contact materials (Ag, Bi, Ti, Mo, etc.) [21], implementation of different interlayer materials ( $\text{Al}_2\text{O}_3$ , TiN,  $\text{MoO}_x$ , graphene,  $\text{MoS}_2$ , etc.) to achieve clear and well-defined interfaces [22, 23], utilization of a single quaternary target [24, 25] and exploration of different stacking orders [26] to enhance the homogeneity and crystal quality of the films. These strategies offer avenues for fine-tuning the features of the samples and advancing their performance in practical applications.

Thin film solar cells find application in various fields such as building integration, wearable technologies, space applications, and the automotive sector [27]. However, the use of rigid substrates limits their application in these fields, and therefore, flexible substrates are more preferred. The

most common flexible substrates which used to growth CZTS thin film solar cells are stainless steel (SS) [28], aluminum (Al) [29], titanium (Ti) [30], molybdenum (Mo) [31], flexible glass (FG) [32], polyimide (P) [36], etc. Among these, FG and P substrates are organic-based and have a maximum sustainable temperature underlain by 500 °C, which is epically lower as compared with metal foils (Al, Ti, Mo, SS) that can withstand temperatures of 600 °C or higher [33].

Table 1 provides information on the price, electrical resistivity, and conductivity properties of various materials. Among the available metal foils, Mo foil is commonly used due to its linear thermal expansion coefficient of  $5.2 \times 10^{-6} \text{ K}^{-1}$ . However, Mo foil is associated with drawbacks such as high price (76.67 USD/kg) [34], high resistivity ( $4.99 \times 10^{-6} \Omega \cdot \text{cm}$ ) [35], and low conductivity ( $17.9 \times 10^6 \text{ S/m}$ ) [36] when compared to other materials. Therefore, there is a demand for a foil material that exhibits low cost, low resistivity, and high conductivity, enabling the fabrication of high-performance CZTS thin film solar cells. Among the listed materials, copper (Cu) foil possesses the most desirable properties, including lower cost (9.17 USD/kg) [34], lower resistivity ( $1.68 \times 10^{-8} \Omega \cdot \text{cm}$ ) [36], and higher electrical conductivity ( $5.96 \times 10^7 \text{ S/m}$ ) [37]. Due to these advantageous properties, Cu foil is widely used as an electrical conductor in circuit boards, batteries, and solar energy appliances.

Work function of back contact, absorber layer, etc. is crucial for enhancing cell parameters and this can be important property to select the flexible foils for CZTS based thin film solar cells. If the work function of metal foil is smaller than semiconductor absorber layer, the Schottky contact is formed between the metal and semiconductor according to the Schottky contact model [38]. The work function of CZTS absorber layer is  $\sim 5.7 \text{ eV}$  [39] and work function of Cu, Mo, Ti and Al foil which presented in Table 1 is determined as 5.10 eV, 5.0 eV, 4.33 eV, and 4.2 eV, respectively. The enhanced work function is belong to the Cu foil and the higher work function of back contact (foil) is enhance to the built-in potential of device [40]. The built-in potential is a consequence of the potential decrease across the width of the absorber layer. This gradual decrease in potential is responsible for generating an electric field directed towards the p-n junction. Consequently, this electric field drives the charge carriers towards the junction. Moreover, the electric field induces an increase in the diffusion length of carriers, enhancing their collection on the n-side of the junction. In the literature, the open-circuit voltage ( $V_{OC}$ ) exhibits high sensitivity to changes in  $\Phi_M$  values, while the short-circuit current ( $J_{SC}$ ) and fill factor (FF) demonstrate relatively minor dependence on  $\Phi_M$  [38].

**Table 1.** Properties of some metals

Subs.	Price (USD/kg) [34, 41]	Resistivity ( $\Omega \cdot \text{cm}$ ) [35, 42]	Conductivity (S/m) [36, 43]	Work Function (eV) [44, 45]	Therm. Exp. ( $\text{K}^{-1}$ ) at $>500 \text{ K}^\circ$ [40, 46, 47]
Cu	9.17	$1.68 \times 10^{-8}$	$5.96 \times 10^7$	5.10	$16 \times 10^{-6}$
Mo	76.67	$4.99 \times 10^{-6}$	$17.9 \times 10^6$	5.0	$4.8 \times 10^{-6}$
Ti	8.26	$4.20 \times 10^{-7}$	$2.38 \times 10^6$	4.33	$8.4-8.6 \times 10^{-6}$
Al	2.56	$2.82 \times 10^{-8}$	$3.5 \times 10^7$	4.2	$21 \times 10^{-6}$

In light of all this information, in this study, CZTS thin films will be produced for the first time on Cu foil using Cu-foil/ZnS/Cu/Sn/Cu stacking order (and RTP method. To investigate the impact of different sulfurization times on the properties of the CZTS thin films, we tested short dwell times (30, 60 and 120 sec.) at reaction temperature of 425 °C. By varying the dwell times, we aimed to understand how it affected the structural, optical, and electrical properties of the CZTS thin films. It should be noted that higher reaction temperatures (> 425°C) have been tried for fabrication of CZTS thin films. However, the most suitable temperature has been determined to be 425°C due to occurrences of fragmentation and detachment on the substrate surface.

## 2 Material and methods

### 2.1 Cleaning of substrates

The study involved a specific pre-treatment process for the glass and flexible Cu-foil substrates before the CZTS thin film deposition. The pre-treatment steps are as follows:

- The glass substrate was cleaned for 5 minutes in an ultrasonic bath with acetone.
- Then, it was cleaned for another 5 minutes in an ultrasonic bath with isopropanol alcohol (IPA).
- Finally, it was cleaned for 10 minutes in an ultrasonic bath with distilled water (DW).
- After the cleaning process, all substrates (both glass and flexible Cu-foils) were dried using N<sub>2</sub> gas.
- All substrates, after the drying process, were exposed to an ultrasonic plasma cleaner system for 30 minutes. The ultrasonic plasma cleaning is a method used to remove any remaining impurities and contaminants from the substrate surfaces, ensuring a clean surface for thin film deposition.
- Once the plasma cleaning was completed, the substrates were placed in a PVD system. Before starting the deposition process, a bias voltage was applied to the substrates for 10 min. This step was carried out under a vacuum level of < 2x10<sup>-6</sup> Torr, ensuring a low-pressure environment to prevent contamination during the deposition process.

The purpose of these pre-treatment steps is to create a clean and well-prepared surface for the subsequent CZTS thin film deposition. Proper cleaning and surface preparation are crucial in ensuring good adhesion and quality of the deposited CZTS thin films, which, in turn, will impact the performance of the fabricated CZTS thin film solar cells.

### 2.2 Deposition of CZTS precursor and thin films

The CZTS samples were grown on glass and flexible Cu-foils using sputtering and RTP methods, respectively. Firstly, high purity Cu (5N), Sn (5N) and ZnS (4N) sputter targets were used to form stacks on flexible Cu-foils via magnetron sputtering (DC and RF). Cu layers were deposited using a DC power supply, while the Sn and ZnS layers were deposited using an RF power supply. Deposition parameters such as 2x10<sup>-6</sup> mTorr base pressure, 10<sup>-3</sup> mTorr operating pressure, 0.9-1.0 Å/s deposition rate was used for the deposition of Cu, Sn and ZnS precursor films. The thickness

of ZnS, Cu, Sn, and Cu layers was given in our previously study [48]. The second stage of fabrication is annealing of precursor layers to form CZTS thin films by RTP system. Annealing parameters of Cu foil/CZTS were illustrated in Table 2. Cu-foil/CZTS samples were labeled as CZTS-30, CZTS-60, and CZTS-120.

**Table 2.** The sample ID, stacking order and annealing parameters of CZTS precursor films

Sample ID	Stacking Order	Annealing Parameters
CZTS-30	Cu-foil/ZnS/Cu/Sn/Cu	425 °C, 1 °C/s, 30 s, 5 mg S
CZTS-60		425 °C, 1 °C/s, 60 s, 5 mg S
CZTS-120		425 °C, 1 °C/s, 120 s, 5 mg S

### 2.3 Characterization of CZTS thin films

The chemical composition and structural properties of all samples were determined using various techniques. Energy-dispersive X-ray (EDX) spectroscopy, X-ray diffractometry (XRD) (PANalytical-EMPYREAN instrument), and Raman spectroscopy (Renishaw inVia Spectrometer) were employed for structural characterization. EDX measurements were performed on a 115×135 μm<sup>2</sup> area, and XRD patterns were obtained using CuKα (λ = 1,5406 Å) radiation. Raman spectroscopy measurements performed by 633 nm of wavelength. Additionally, scanning electron microscopy (SEM) using a ZEISS-GeminiSEM instrument was utilized to examine the surface properties of the samples.

The optical properties of the samples were determined using two techniques: spectroscopic ellipsometry (J.A. Woollam Co., Inc.) and photoluminescence (PL). Transmittance measurements were conducted on samples deposited on glass substrates ranging from 620 nm to 1240 nm. Room temperature PL measurements were obtained using a hybrid Raman spectroscopy system employing laser excitation wavelength of 633 nm.

## 3 Result and discussion

The EDX analysis of precursor films annealed using 30 s, 60 s, and 120 s dwell times deposited on flexible Cu-foil at 425 °C reaction temperature to form CZTS films are given in Table 3. In Table 3, regardless of dwell times, all samples exhibited Cu-poor (Cu/(Zn+Sn)<1) chemical compositions. However, it was observed that, only sample which annealed using 30 dwell time had Zn-rich (Zn/Sn>1) composition, while samples which annealed by using 60 and 120 s dwell times had stoichiometric Zn (Zn/Sn=1) composition. The Cu-poor and Zn-rich chemical composition is a desired property for CZTS-based thin film solar cells, as this composition gives rise to the formation of Cu vacancies (V<sub>Cu</sub>) and Cu<sub>Zn</sub> antisite defects. These defects contribute to achieving strong p-type conductivity and, consequently, higher solar cell performance [48].

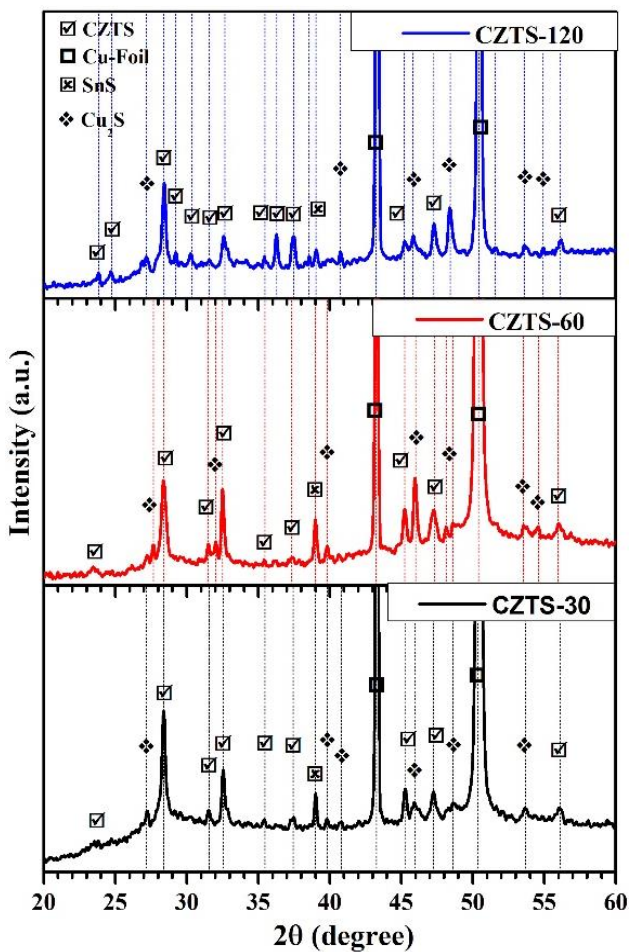
The diffraction pattern of annealed samples for 30 s, 60 s, and 120 s dwell times on flexible Cu-foil was presented in Figure 1. The main diffraction peaks of CZTS samples observed at around 2θ = ~28.45° (112), ~47° (220/204) and ~56° (312/116) that are attributed to kesterite CZTS structure (98-018-4358). The other observed peaks are also marked on the Figure 1. Also, the diffraction peaks at 2θ = ~43.25°, and



~50.40° positions stemming from the Cu-foil (98-062-7117) were observed. Moreover, the peak we determined at  $2\theta = \sim 38.9^\circ$  position was attributed to the SnS secondary phase (98-006-7442) and peaks at around  $2\theta = 27.20^\circ, 39.8^\circ, 40.75^\circ, 45.75^\circ, 48.41^\circ, 53.71^\circ$  and  $54.96^\circ$  are attributed to the  $\text{Cu}_2\text{S}$  secondary phase (98-002-3596). It is predicted that the formation of the  $\text{Cu}_2\text{S}$  phase which detected in the diffraction patterns is caused by the interaction of the sulfur vapor with the Cu foil. In addition, the formation of SnS phase which observed in XRD patterns is due to the inability to provide sufficient sulfur vapor pressure in the structure due to low reaction temperature. It was determined that the diffraction peaks of the mentioned secondary phases decreased significantly by increasing the dwell time from 30 to 120 s [49].

**Table 3.** The EDX measurements of the sulfurized samples on flexible Cu-foil using different dwell times

Sample	Atomic Weight (%)				Atomic Ratio		
	Cu	Zn	Sn	S	Cu/ (Zn+Sn)	Zn/ Sn	S/ Metal
CZTS-30	22.4	12.4	12.0	53.2	0.92	1.03	1.13
CZTS-60	22.1	12.3	12.2	53.4	0.90	1.00	1.15
CZTS-120	21.9	12.1	12.1	53.8	0.91	1.00	1.17



**Figure 1.** The XRD pattern of annealed samples for 30 s, 60 s and 120 s sulfurization times

In order to investigate and compare crystal quality of the samples, some structural parameters of CZTS thin films were determined by using main diffraction peak belongs to (112) diffraction plane of kesterite CZTS structure. First, the Debye-Scherrer formula (Equation (1)) were used to calculate the crystallite size ( $D$ ) of films by using the Full Width Half Maximum (FWHM) values ( $\beta$ ) of the (112) diffraction peak. In addition, the micro-strain values ( $\epsilon$ ) inside the lattice were calculated by Equation (2). The dislocation density ( $\delta$ ) and the number of crystals per unit surface area ( $N$ ) were determined with Equation (3) and Equation (4).  $T$  represents thickness of the film. Moreover, the inter-planar spacing and lattice parameters such as  $a$  and  $c$  were calculated by using Equation (5) and Equation (6).

$$D = \frac{K\lambda}{\beta \cdot \cos\theta} \quad (1)$$

$$\epsilon = \frac{\beta \cdot \cos\theta}{4} \quad (2)$$

$$\delta = \frac{1}{D^2} \quad (3)$$

$$N = \frac{T}{D^3} \quad (4)$$

$$d = \frac{\lambda}{2 \cdot \sin(\theta)} \quad (5)$$

$$\frac{1}{d^2} = \frac{h^2 + k^2}{a^2} + \frac{l^2}{c^2} \quad (6)$$

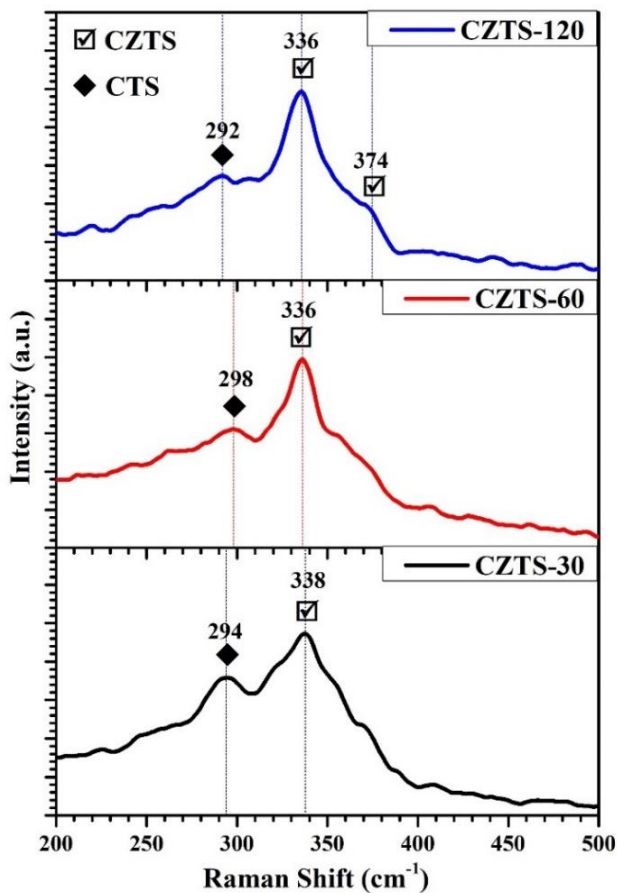
The crystallite sizes of kesterite CZTS films sulfurized for 30, 60, and 120 s dwell times were found to be 34.51 nm, 22.88 nm, and 32.18 nm, respectively. In addition, for 30 s, 60 s and 120 s sulfurization dwell times,  $1.05 \times 10^{-3}$ ,  $1.58 \times 10^{-3}$ ,  $1.13 \times 10^{-3}$  micro-strain,  $8.4 \times 10^{-4}$ ,  $1.9 \times 10^{-3}$ ,  $9.6 \times 10^{-4}$  dislocation density,  $3.4 \times 10^{-2}$ ,  $11.6 \times 10^{-2}$ ,  $4.2 \times 10^{-2}$  number of crystal per unit surface area values were calculated and presented in Table 4. As a result of the calculations, it was seen that CZTS-30 thin film exhibited higher crystallite size and better structural properties. The smaller crystallite size means that formation of more grain boundaries and accordingly more recombination centers for charge carriers [50]. Furthermore, in the lattice, increasing or decreasing of the strain can change all the physical properties of the films (for example, the position of the atoms). Additionally, secondary phase formations ( $\text{Cu}_2\text{S}$ , SnS, etc.) causes to decrease the active region in the films, the shunt resistance and accordingly cell performance [51].

The diffraction patterns of kesterite CZTS phase is similar with some other secondary phases (CTS, ZnS, etc.) [52]. Thus, Raman spectra was used as a complementary technique. The Raman spectra of annealed films for different sulfurization times (30, 60, and 120 s) are given in Figure 2. Considering the Figure 2, irrespective of the sulfurization times, the main peak position was observed at around  $\sim 336$ - $338 \text{ cm}^{-1}$  which is attributed to the kesterite phase in all samples. This peak is ascribed to vibration of S atoms in kesterite structure [53, 54]. The other peaks are marked on the graphs [55, 56]. Apart from Raman modes of kesterite

phase, the minor peaks determined in all samples at around 292-298  $\text{cm}^{-1}$  are associated with the  $\text{Cu}_2\text{SnS}_3$  (CTS) phase [57, 58].

**Table 4.** The structural parameters of CZTS samples sulfurized for 30 s, 60 s and 120 s dwell times

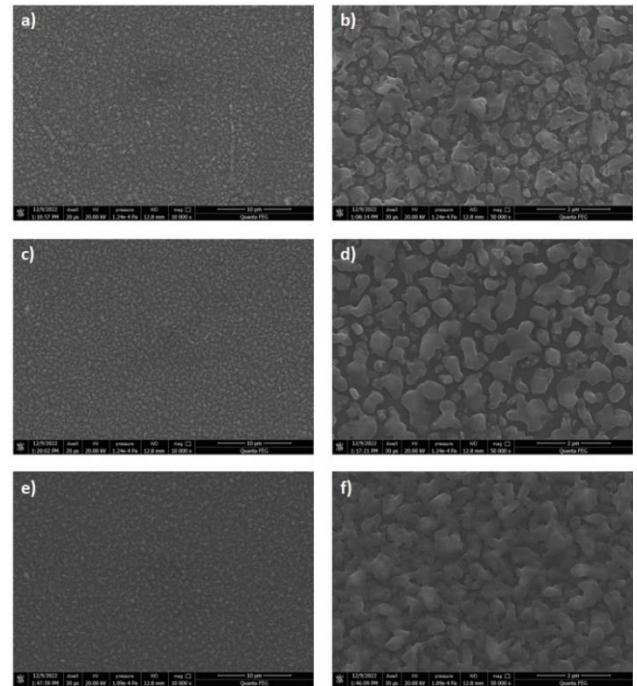
Parameters	CZTS-30	CZTS-60	CZTS-120
$2\theta$ ( $^\circ$ )	28.39	28.40	28.41
FWHM ( $^\circ$ )	0.248	0.374	0.266
D (nm)	34.51	22.88	32.18
$\varepsilon$ (a.u.)	$1.05 \times 10^{-3}$	$1.58 \times 10^{-3}$	$1.13 \times 10^{-3}$
$\delta$ ( $\text{nm}^{-2}$ )	$8.4 \times 10^{-4}$	$1.9 \times 10^{-3}$	$9.6 \times 10^{-4}$
N ( $\text{nm}^{-2}$ )	$3.4 \times 10^{-2}$	$11.6 \times 10^{-2}$	$4.2 \times 10^{-2}$
d ( $\text{Å}$ )	3.1412	3.1401	3.1339
a ( $\text{Å}$ )	5.4356	5.4269	5.4356
c ( $\text{Å}$ )	10.9023	10.9261	10.8797
c/2a	1.0028	1.0067	1.0008



**Figure 2.** The Raman spectra of annealed samples with different sulfurization dwell times

The effect of sulfurization time on the surface morphologies and microstructure features of all samples were determined by SEM measurements employing two different magnifications (10 kX and 50 kX). The SEM images of CZTS thin films grown on the flexible Cu-foil using different sulfurization times are given in Figure 3. It was observed that all samples had dense and homogenous polycrystalline surface structure. CZTS samples which are sulfurized for 30 and 60 s dwell times have a granular structure morphology (Figure 3a-d). After sulfurization time

increased from 60 to 120 s (Figure 3e, f), the grain boundaries were disappeared, and the more compact surface morphology was formed.



**Figure 3.** The SEM images of kesterite samples sulfurized for different dwell times; (a-b) 30 s, (c-d) 60 s, (e-f) 120 s

Optical properties of CZTS thin films grown on Cu foil using various sulfurization times (30 s, 60 s, and 120 s) characterized by two distinct approaches. In the first method, CZTS thin films were deposited on glass substrates to ascertain the optical band gaps by employing optimized sulfurization parameters for Cu-foil based samples. The optical band gap of the CZTS thin films was calculated based on the absorption coefficient ( $\alpha$ ), as determined by applying the Lambert-Beer law outlined in Equation (7) [59]:

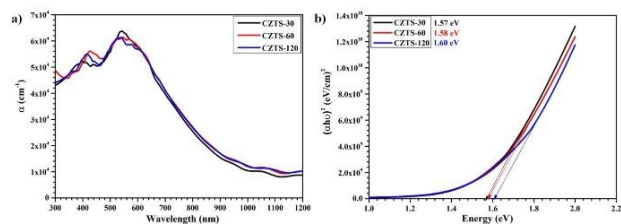
$$\alpha = \frac{1}{d} \ln\left(\frac{1}{T}\right) \quad (7)$$

where, T and d are transmission and thickness parameters of the film. After calculating  $\alpha$ , absorption coefficient spectra were given in Figure 4a. It was observed that all samples have absorption coefficient above  $10^4 \text{ cm}^{-1}$ . However, CZTS-30 thin film has slightly higher than the others. In addition, the Tauc plot [60] was used to estimate the optical band gap of the films using Equation (8);

$$(\alpha h\nu)^2 = A(h\nu - E_g) \quad (8)$$

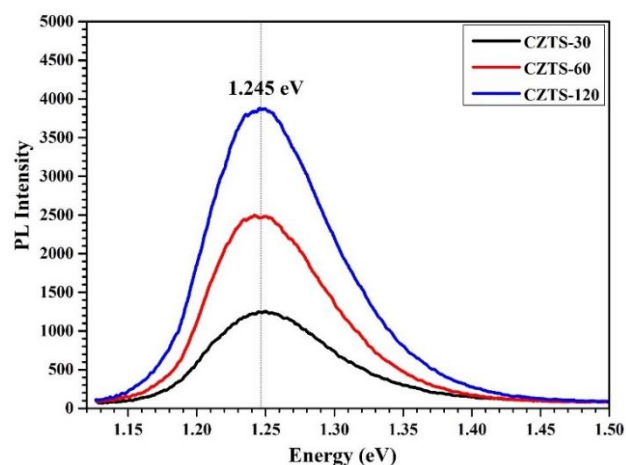
where, A represents a constant,  $h\nu$  denotes photon energy, and  $E_g$  signifies the optical band gap energy of the films. The optical band gap was ascertained through the plot of  $(\alpha h\nu)^2$  against photon energy (eV), as depicted in Figure 4b. Figure 4b displays the determination of the optical band gap for CZTS thin films, yielding values of 1.57 eV, 1.58 eV, and 1.60 eV for sulfurization times of 30 s, 60 s, and 120 s,

respectively. Notably, these obtained optical band gap values align with previously reported findings in the literature [61]. The minor variations observed in the optical band gaps (ranging from 1.57 eV to 1.60 eV) have been attributed to the presence of secondary phases within the kesterite structure. Consequently, these slight differences in the chemical composition of the films and the stresses induced within the structure by these secondary phases are believed to be contributing factors [62].



**Figure 4.** The plots for absorption coefficient and optical band gap of CZTS samples

The optical properties of CZTS thin films subjected to various sulfurization dwell times were additionally assessed through room temperature PL measurements, as illustrated in Figure 5. In all cases, the films displayed a prominent PL emission at approximately 1.25 eV. When comparing the band gap values and PL emissions of these samples (as depicted in Figure 4), it is reasonable to suggest that the PL emissions can be ascribed to transitions occurring from the conduction band to acceptor energy levels, band tailing, or other defect levels (such as  $V_{Cu}$ ,  $Cu_{Zn}$ , etc.) [63]. In summary, the lack of observable band-to-band transitions in the PL measurements can be linked to the intrinsic defect levels and recombination centers present within the CZTS structure [64].



**Figure 5.** Room temperature PL measurements of CZTS thin films sulfurized with different dwell times

#### 4 Conclusion

This study presents a detailed investigation of kesterite CZTS thin films deposited on flexible Cu-foil with varying sulfurization dwell times. The XRD analysis identified distinctive kesterite CZTS phase peaks at  $2\theta = 28.45^\circ$  (112),  $47^\circ$  (220/204), and  $56^\circ$  (312/116). Additional peaks

corresponding to Cu-foil,  $Cu_2S$  secondary phase, and  $SnS$  secondary phase were also observed. Raman spectroscopy confirmed the kesterite phase, notably with a dominant peak at  $\sim 336\text{ cm}^{-1}$  attributed to S atom vibrations within the kesterite structure. Minor peaks indicated the presence of  $Cu_2SnS_3$  (CTS) phase not discernible by XRD. EDX analysis revealed consistent Cu-poor and Zn-rich compositions across for some samples, with slight variations due to sulfurization dwell times. SEM imaging at different magnifications (10 kX and 50 kX) displayed changes in surface morphology and grain structures. Films sulfurized for 30 s and 60 s exhibited a granular structure, whereas extending to 120 s resulted in a more compact surface. Optical band gap values ranged from 1.57 to 1.60 eV. PL measurements consistently showed strong emission at around 1.25 eV for all samples, indicating transitions related to defect levels within the CZTS structure. The absence of observable band-to-band transitions in PL suggested the presence of intrinsic defect levels and recombination centers within CZTS. These findings provide significant insights into developing sustainable and efficient solar energy utilizing flexible Cu-foil as a promising substrate for CZTS-based thin film solar cells.

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#### Conflict of interest

The authors declare that there is no conflict of interest.

**Similarity rate (iThenticate):** 17 %

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