

Article

The Impact of Thermal Treatment on the Structural, Optical and Electrochemical Characteristics of Tin Sulfide Films

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Abstract: The development of eco-friendly, cost-effective, and naturally abundant electrode materials for supercapacitors is gaining critical importance in current energy storage research. This study focuses on the synthesis of tin sulfide (SnS_x) films via the eco-friendly successive ionic layer adsorption and reaction (SILAR) method, employing varying quantities of L-ascorbic acid (0.8 and 1.0 g) as a reducing agent. Tin sulfide films were deposited on fluorine-doped tin oxide (FTO) glass substrates and subsequently annealed in an inert atmosphere at temperatures ranging from 200 to 400 °C, resulting in thin films of varying thicknesses (100–420 nm). The structural and compositional characteristics of the films were thoroughly analyzed using Raman spectroscopy to confirm the purity and spectroscopic signatures of the sulfides. Further characterization was performed to assess the films' morphology (scanning electron microscopy, SEM), phase composition (X-ray diffraction, XRD), surface chemical states (X-ray photoelectron spectroscopy, XPS), optical properties (UV–Vis spectroscopy), and electrical properties (Hall measurements). The gathered data were then used to evaluate the potential of tin sulfide films as electrode materials in supercapacitors, highlighting their suitability for sustainable energy storage applications.

Keywords: tin sulfides; supercapacitors; SILAR



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1. Introduction

Energy is a critical factor in global development, with approximately 65% of the world's energy demand currently met by non-renewable fossil fuels [1]. This reliance contributes to greenhouse gas emissions, global warming, and environmental degradation [2]. To mitigate these issues and address the depletion of fossil fuels, a shift toward renewable energy sources is imperative [3]. In addition to the development of renewable energy, advances in energy storage technologies, particularly electrochemical devices, are essential for efficient energy conversion and storage [4].

Supercapacitors (SCs) have emerged as promising energy storage devices due to their high power density, extended life cycle, and superior energy density compared to traditional capacitors [5]. They are suitable for various electrical and electronic applications [6]. SCs are categorized into electric double-layer capacitors (EDLCs) and pseudocapacitors based on their charge storage mechanisms [7]. Pseudocapacitors, which include materials like metal oxides, metal sulfides, and conductive polymers, generally offer higher capacitance than EDLCs but with a shorter life cycle [8].

Among pseudocapacitive materials, metal chalcogenides, particularly tin sulfides, have garnered significant attention due to their advantageous physicochemical properties [9,10]. Tin-based chalcogenides such as SnS and SnS₂ are of particular interest, although research on Sn₂S₃ remains limited [11]. Tin and sulfur are non-toxic and abundant elements, making tin chalcogenides attractive for energy applications due to their robust structure and excellent electrical and optical properties [12].

SnS, with its orthorhombic structure and a bandgap of approximately 1.1 eV, is comparable to silicon [13] and exhibits promising characteristics for near-infrared applications [14]. SnS₂, an n-type semiconductor, adopts a hexagonal structure, while SnS can exhibit both n-type and p-type conductivity, influenced by annealing temperature [15,16]. The properties of these materials make them suitable for a range of applications, including photovoltaic devices [17], supercapacitors [18], optical sensors, and transistors [19].

The synthesis of high-quality thin films using cost-effective and abundant materials remains a challenge. Various deposition methods, such as thermal evaporation [20], chemical bath deposition [21], and spin coating [22], have been explored for tin sulfide films, with photovoltaic applications being the primary motivation [23]. It is well established that post-deposition annealing improves the crystalline quality of tin sulfide films, with optimal results often achieved at temperatures between 300 °C and 600 °C, depending on the desired material properties [14,23–25].

Mixed tin sulfides (SnS_x) have shown versatility in various technological applications. The successive ionic layer adsorption and reaction (SILAR) method is a cost-effective and efficient approach for thin-film deposition, enabling large-area applications with strong substrate adhesion. This study examined the structural, morphological, optical, and electrical properties of tin sulfide films deposited on FTO glass and annealed in a nitrogen atmosphere at 200–400 °C, highlighting the occurrence of “chemical epitaxy” due to the well-defined orientation of tin sulfide on the substrate [26,27].

2. Materials and Methods

2.1. Materials

As substrates for the deposition of thin films, fluorine-doped tin oxide (FTO) glass slides with a parameters of size of 20 mm × 15 mm and a sheet resistance of $9.38 \pm 0.38 \Omega/\text{sq}$ were provided by Ossila.com. Labochemia. It served tin chloride dihydrate with a purity of 98%. 98% sodium sulfide nonahydrate was brought by Honeywell.com (Charlotte, NC, USA). The 99% pure form of L-ascorbic acid was purchased from Sigma-Aldrich (Taufkirchen, Germany).

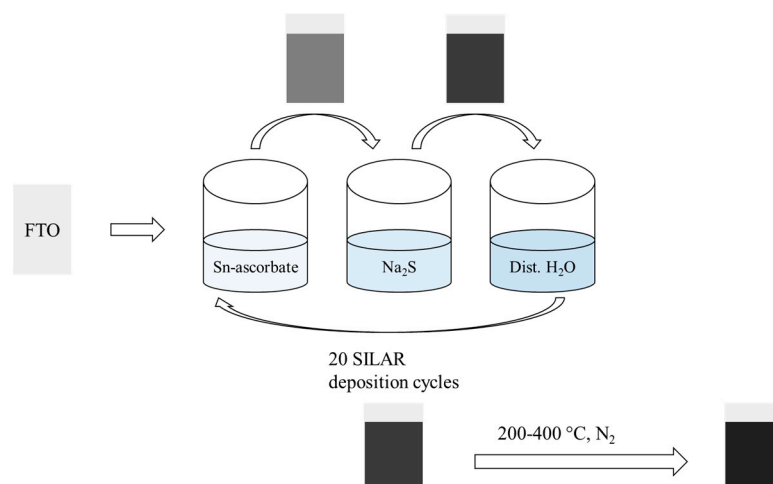
2.2. Film Deposition

For the deposition of a thin tin sulfide film, the SILAR method reported earlier [9] was used. Firstly, the washed FTO glass slides were ultrasonically cleaned in acetone for 10 min. The SILAR method is based on the one-by-one placement of cation and anion precursor solutions and distilled water. The aqueous cationic solution consisted of 0.04 M SnCl₂ with 0.8 g or 1.0 g of L-ascorbic acid, and the aqueous anionic solution was 0.04 M Na₂S. For the synthesis, 20 SILAR cycles at 40 °C were used. The synthesis mechanism is shown in Scheme 1.

To modify the deposited films, annealing in an inert atmosphere at different temperatures (200 °C, 250 °C, 300 °C, 350 °C, and 400 °C) was applied. The deposition parameters and sample numbers are listed in Table 1.

Table 1. Parameters and sample numbers.

Quantity of L-ascorbic Acid, g	Annealing Temperature, °C				
	200	250	300	350	400
0.8	1	2	3	4	5
1.0	6	7	8	9	10



Scheme 1. Graphic representation of the preparation of thin tin sulfide films using the SILAR method.

2.3. Characterization of Films

Thin tin sulfide films were characterized using XRD, Raman, SEM, and XPS measurements. UV–Vis spectroscopy and electrochemical calculations were performed on the samples. A description of the methods used is provided in the Supplementary Data.

3. Results

Tin sulfide nanoparticles were synthesized by changing the mass of the capping agent (L-ascorbic acid) via the SILAR route, as shown in Scheme 1. To improve crystallinity, the samples were annealed in an inert atmosphere in the temperature range of 200–400 °C. As-deposited and annealed tin sulfide thin films adhere strongly to the substrate. Furthermore, it was noted that with an increase in the annealing temperature, the obtained films became smoother and thinner.

This section may be divided into subheadings. A concise and precise description of the experimental results, their interpretation, and the conclusions that can be drawn should be provided.

X-ray diffraction. X-ray diffraction (XRD) analysis was utilized to examine the crystalline structure and phase purity of the thin films. The XRD data of the as-prepared tin sulfide films deposited previously under the same conditions [1] agree with the typically reported structure of herzenbergite (ICDD PDF No. 39-0354). Meanwhile, the XRD curves (Figure 1) of Sn-S annealed at different temperatures identify some phase transformations and increased crystallinity. In general, all the annealed films are characterized by the main and most intense peaks at $2\theta = 26.64^\circ$, 37.88° , and 51.65° , which are assigned to orthorhombic SnS (ICDD PDF No. 83-47). However, SnO₂ peaks (ICDD PDF No. 46-1088) can also be identified by these peaks due to the very close values of the interplanar spacing (Table 2). Therefore, other instrumental analysis methods have been used to clarify the prevailing phase of tin sulfide.

Starting at 200 °C, a certain hill at $2\theta = 18\text{--}35^\circ$ is barely visible compared to the as-prepared films, which were revealed to be more amorphous [1]. At 250 °C and below, a small peak appears at $2\theta = 31.68^\circ$ (hkl index 310), which is assigned to Sn₂S₃ (ICDD PDF No. 75-2183). This phase decomposes to SnS₂ and SnS at temperatures above 300 °C [2,3] via the following reaction route: Sn₂S₃ → SnS₂ + SnS. A further increase in the annealing temperature highlights the presence of hexagonal SnS₂ (mineral berndite, ICDD PDF No. 83-1705) with a small peak at $2\theta = 16.00^\circ$ (hkl index 001). Although the XRD spectra of films annealed at 300 and 350 °C do not disclose any SnS₂ peaks, this particular phase shows poor crystallinity and can be intangible by the XRD technique [4–6]. Furthermore, due to the desulfurization of SnS₂ to SnS and S above 350 °C [4,7], this phase no longer exists in the 400 °C spectrum.

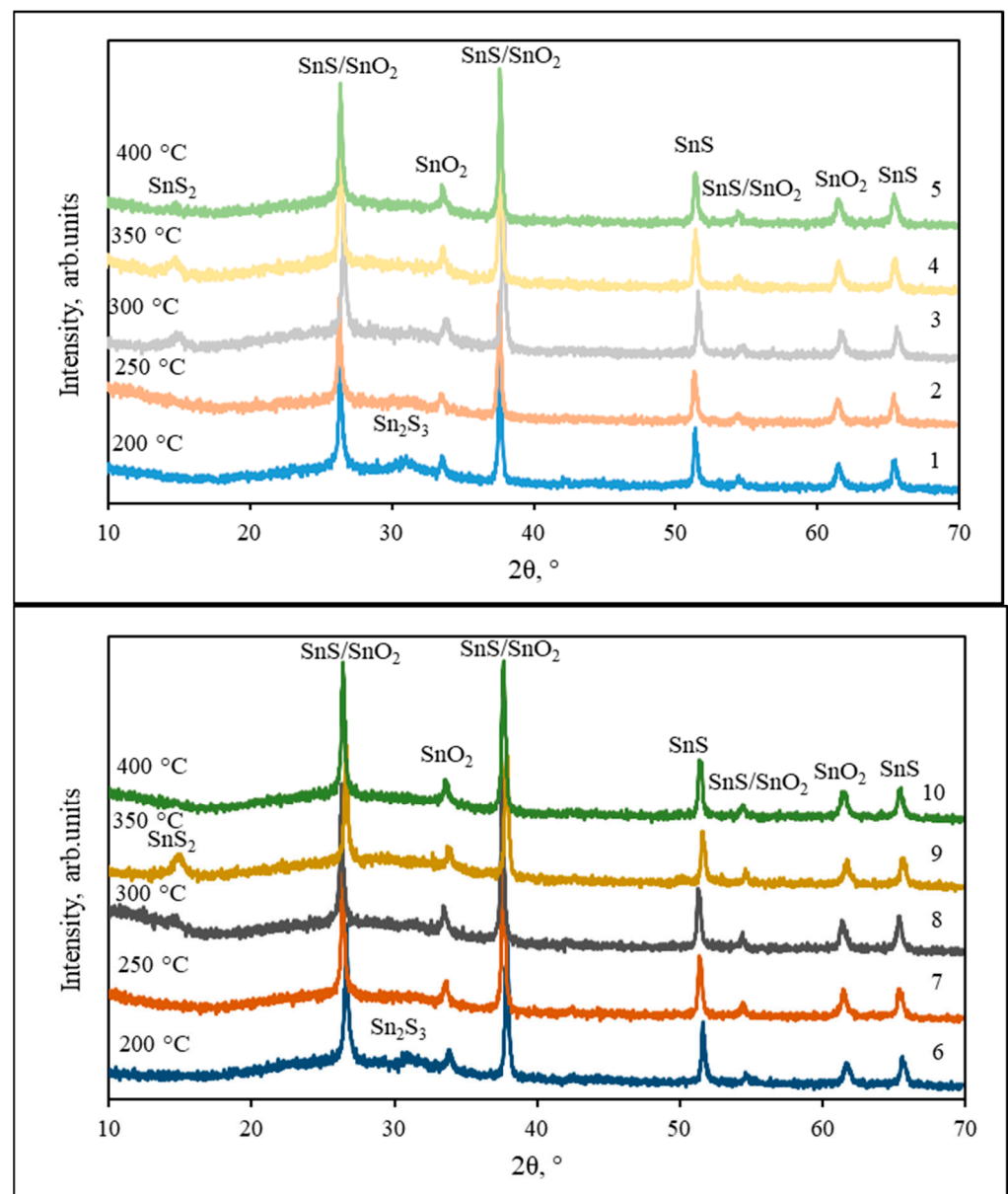


Figure 1. X-ray diffraction patterns of samples prepared with 0.8 g (top) and 1.0 g (bottom) of L-ascorbic acid, annealed at temperatures ranging from 200 to 400 °C.

Table 2. Comparison of observed d-values with standard d-values.

Observed d-Values, Å	SnS (83-47) d-Values, Å	Sn ₂ S ₃ (75-2183) d-Values, Å	SnO ₂ (46-1088) d-Values, Å	SnS ₂ (83-1705) d-Values, Å
5.9416				5.8800
3.3757	3.3774		3.3510	
2.6680		2.6521	2.6518	
2.3727	2.3751	2.3707	2.3799	
1.7762	1.7792	1.7752	1.7649	
1.6841	1.6810			
1.5064		1.5001	1.5012	
1.4270	1.4342	1.4251		

The quantity of L-ascorbic acid was found to not affect the phase composition as the spectra are similar when 0.8 g and 1.0 g are used. However, it appears that a higher amount of ascorbic acid improves the crystallinity of the films because the spectra obtained with 1.0 g are smoother.

In summarizing the results of the XRD analysis, it can be emphasized that the diffraction patterns indicate a mixed-phase composition of the samples, whereas the main peaks are assigned to SnS. The annealing temperature promotes the desulfurization process of the secondary phases, so it is likely that the purest films with a predominant SnS are obtained at a temperature of 400 °C. However, XRD is a one-point analysis technique [8]; therefore, the possibility of the existence of secondary phases (SnS₂ and Sn₂S₃) should not be ignored. To check the reliability of the results, other methods of instrumental analysis, Raman mapping and XPS, have been conducted in this work. Since more pronounced changes in the structure of the films occur approximately every 100 °C, the samples heated at temperatures of 200, 300, and 400 °C were selected for further analysis, eliminating intermediate temperatures.

Crystallite size was determined using the most prominent XRD peaks. Compared to the findings in a previous report [9], annealing resulted in an increase in the crystallite size. When utilizing 0.8 g of L-ascorbic acid, there was a slight variation in particle size, ranging between 33 and 35 nm. Elevated temperatures influenced the crystallite size, leading to a reduction. Conversely, a higher amount (1.0 g) of L-ascorbic acid led to the formation of larger particles. As the annealing temperature increased, the particle size exhibited significant variation within a broader range of 30–35 nm. To generate smaller crystallites of tin sulfide, higher annealing temperatures and reduced quantities of L-ascorbic acid are required.

Raman studies. The Raman study, along with the X-ray diffraction analysis, confirms the growth of mixed-phase tin sulfide films. It was expected from previous results [9] that samples are not entirely homogeneous; in that case, to obtain more accurate data, Raman mapping was a better method than acquiring spectra from single points of the sample. Figure 2 represents the Raman results of the samples. According to the literature [10–12], the peaks at 70, 90, 191, and 224 cm⁻¹ (attributed to the different Raman modes of A_{1g} and B_{3g}) are indicators of the presence of SnS, while the peak at 312 cm⁻¹ indicates SnS₂ [13,14] and is present at the point where the spectrum was obtained. However, the peak that appeared at 308 cm⁻¹ was independent of the Raman spectra of the SnS single crystal and represented Sn₂S₃ [15,16]. Thermal treatment at 200 °C shows the formation of mixed phases where SnS peaks and a low-intensity peak of Sn₂S₃ were observed. The peak shift from Sn₂S₃ to SnS₂ was visible after annealing of samples at the temperature of 300 °C. Here, a new peak with no clear form peak at 191 cm⁻¹, assigned to SnS, was obtained. A high-intensity peak of SnS₂ appeared for the sample annealed at 400 °C. No additional peaks assigned to other phases were observed, and the spectra of the entire surface show the formation of SnS₂.

One broad peak around 310 cm⁻¹ was detected in the samples; it was challenging to analyze SnS₂ and Sn₂S₃ separately. The local maxima on the spectral average curve were 306 cm⁻¹ for sample 1, 311 cm⁻¹ for sample 3, 312 cm⁻¹ for sample 8, and 314 cm⁻¹ for sample 5. In the case of sample 5, the peak was significantly thinner, and SnS₂ mainly caused this peak.

To demonstrate the homogeneity of the samples, colored maps were graphed based on a single variable analysis method. The proportion of the intensities of the most intense peaks at 90 cm⁻¹ and 312 cm⁻¹ are plotted on two complementary maps. These two peaks were chosen for analysis because they represent the two endpoints of the oxidation states of the Sn atoms. The map on the left of the table was calculated by the following equation for each point of the map:

$$\frac{\text{Intensity at } 90 \text{ cm}^{-1}}{\text{Intensity at } 90 \text{ cm}^{-1} + \text{Intensity at } 312 \text{ cm}^{-1}} * 100 \quad (1)$$

while map on the right:

$$\frac{\text{Intensity at } 312 \text{ cm}^{-1}}{\text{Intensity at } 90 \text{ cm}^{-1} + \text{Intensity at } 312 \text{ cm}^{-1}} * 100 \quad (2)$$

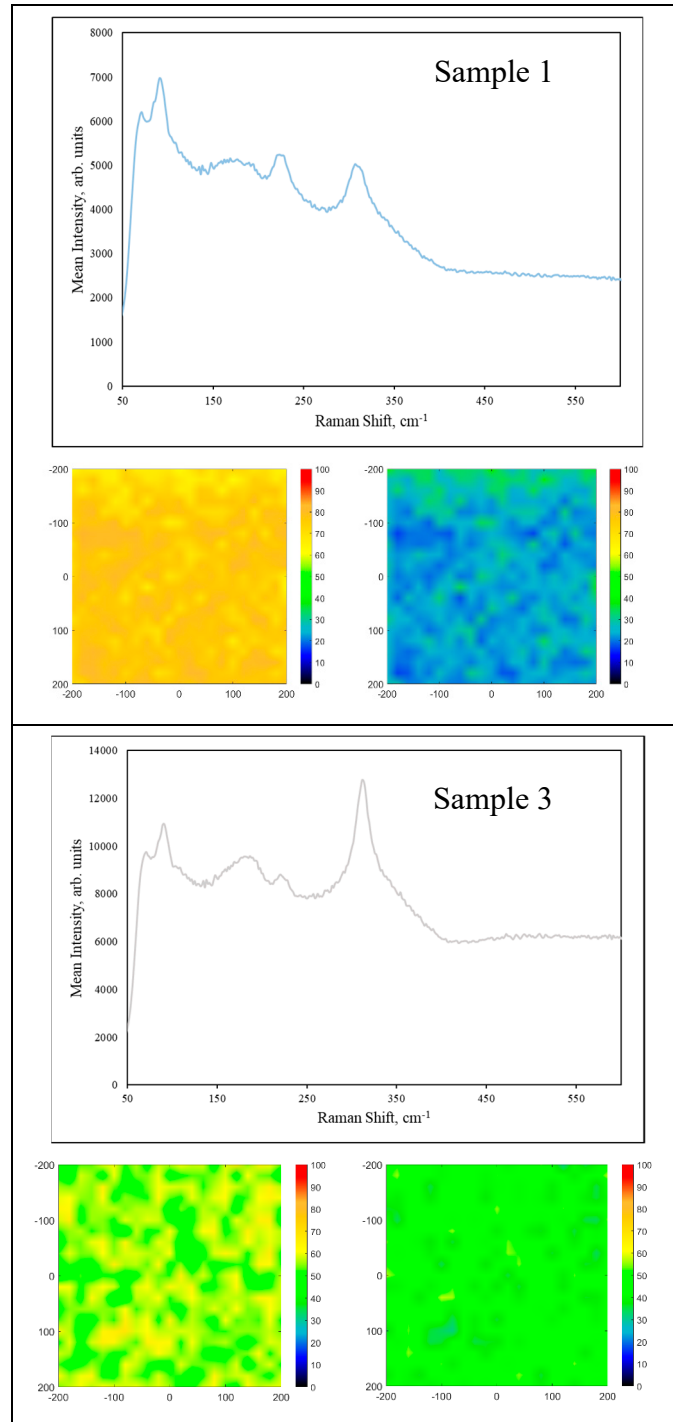


Figure 2. Cont.

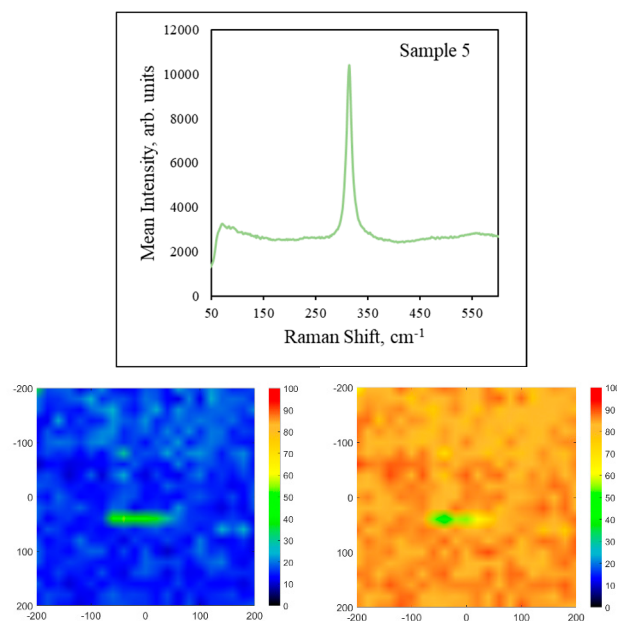


Figure 2. Results of Raman spectroscopy of samples prepared with 0.8 g L-ascorbic acid and annealed.

A color scale is included in these matrices and plotted as a graphical map. These two maps were arranged together with the mean spectra and an array of curves with all spectra obtained from each sample. These diagrams can provide detailed data on the oxidation state of tin atoms, the components present, and how different materials are distributed on the surface of the sample.

Annealing affects the oxidation state of the tin atoms, and the Raman measurements clearly show this. Sample 1, annealed at 200 °C, has very similar chemical properties to the non-annealed sample in our previous reports [1,9]. In addition, the effects of annealing are significant only at higher temperatures. However, it is very hard to synthesize pure SnS; impurities like SnS₂ and Sn₂S₃ are present in annealed samples (and non-annealed ones), but the proportion of Sn(II) in sample 1 is much greater compared to that in other samples. Sample 3, annealed at 300 °C, is a great example of mixed tin sulfides; the proportion of the examined peaks is close to equal, and all possible materials are present in great amounts. The difference in the amount of ascorbic acid used has little effect compared to the annealing temperature. Sample 5, annealed at 400 °C, has a higher amount of SnS₂ than SnS. Probably, annealing at higher temperatures causes the SnS phase transformation to SnS₂ [17].

X-ray photoelectron spectroscopy.

To investigate the oxidation state, chemical composition, and constituent elements of the formed films at various post-annealing temperatures, XPS measurements were conducted. The films were found to consist of a few phases: SnS, SnS₂, and Sn₂S₃, as determined by XRD and supported by Raman analyses. Figure 3 shows the high-resolution XPS spectra of the Sn 3d, S 2p, and O 1s peaks of thermally treated films at 200–400 °C. The high-resolution spectra of Sn 3d for the two phases obtained are shown in Figure 3. The formation of tin sulfide SnS is confirmed by the component with the binding energy of 485.6–486.1 eV, while the reference value for tin disulfide SnS₂ is higher than 486.8 eV [6,18]. According to the XPS database [19], the indication of Sn²⁺ ions marks the peak at a binding energy of 486.8 eV. Additionally, the peak observed at 486.4 eV corresponds to Sn²⁺, aligning with the formation of SnS. An E_b of approximately 486.1 eV is characteristic of tin bonded with sulfur in SnS [20–22]. Here, the deconvolution of the Sn 3d peak showed binding energies of 485.8 and 486.4 eV, corresponding to SnS and SnS₂, respectively [23]. The location of the tin peak depends on the annealing temperature; a lower temperature indicates the formation of Sn²⁺. A light shift to higher energies is due to the increase in the oxidation state of tin to Sn⁴⁺ with increasing annealing temperature. For the comparison

of the binding energies of the films, it should be mentioned that the binding energy corresponding to the Sn^{4+} state appeared after thermal treatment at $300\text{ }^\circ\text{C}$, while the Sn^{2+} state E_b appeared with annealing at $200\text{ }^\circ\text{C}$ [24]. Properly, the phase conversion from SnS to SnS_2 occurs with an increase in the annealing temperature, and the mixture of SnS_2 and SnS appears with annealing at $300\text{ }^\circ\text{C}$, forming the binding energies between Sn^{4+} and Sn^{2+} states.

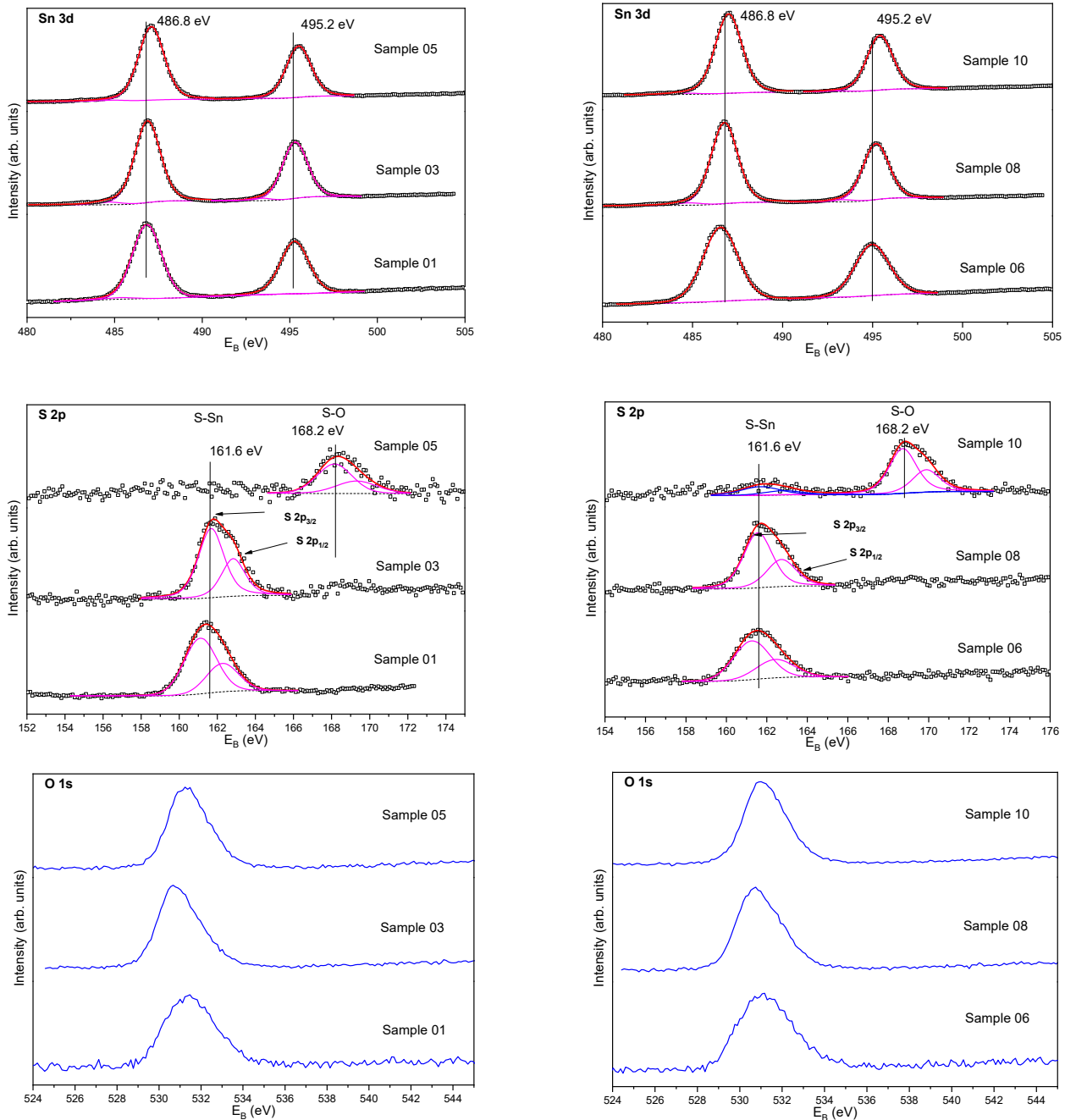


Figure 3. High-resolution XPS spectra of tin, sulfur, and oxygen of samples annealed at $200\text{ }^\circ\text{C}$ (samples 01 and 06), $300\text{ }^\circ\text{C}$ (samples 03 and 08), and $400\text{ }^\circ\text{C}$ (samples 05 and 10).

Taking into account the XPS spectra of sulfur, the thin film thermally treated at $200\text{ }^\circ\text{C}$ has two peaks at 162.4 and 161.6 eV, showing the formation of both tin sulfides (SnS and SnS_2). Thermally treated film at $300\text{ }^\circ\text{C}$ demonstrates a light chemical shift when SnS_2 was formed. Annealing at $400\text{ }^\circ\text{C}$ gave significant results due to the appearance of the S-O

junction at 168.2 eV. Ascorbic acid will probably dissociate, and its fragments will react with sulfur. In addition, two peaks at 161.6 and 162.8 eV were observed, indicating the formation of SnS₂.

The oxygen spectra consist of one very weak peak, corresponding to the Sn-O bond or the C-O bond, which could be due to the small amount of oxygen and carbon impurities in the deposited thin films [17,25].

Furthermore, the X-ray diffraction analysis did not reveal any signal of sulfate formation, but only low-intensity peaks of secondary phases, such as SnS₂ and Sn₂S₃. In addition, for SnS in the films annealed at different temperatures, XPS analysis established the presence of sulfate and secondary phases. An explanation for the absence of these phases might be due to the growth of the amorphous phase [26–29]. Although a possible reason is for the diffraction cards, the peak intensity of sulfate and SnS₂ could be shorter than that of SnS, and their position is closer to the SnS peaks. Therefore, they could overlap each other, and no peaks could be detected against the SnS peaks. Furthermore, there is a possibility that such compounds, as well as sulfate and SnS₂, could appear only in small amounts and could be detected only by XPS analysis due to their higher sensitivity compared to XRD [23].

Scanning electron microscopy

The morphology of the annealed SnS thin film has been studied using SEM. Tin sulfide crystals grow, form thin films, and adhere to the substrate by van der Waals forces. Tin sulfide particles are formed and adhere to the substrate as agglomerates [23]. SEM images of thin films with a magnification of 100k in Figure 4 show densely packed grains. These images show spherical SnS nanoparticles, some of which are connected to each other to form short chains. SEM analysis clearly demonstrates that annealing has a significant effect on grain size and surface roughness. SEM images in our previous report [1] show the presence of very irregular shapes and forms of tin sulfide that have fewer deep valleys. It is clear that tin sulfide is formed in irregular shapes and forms. At lower annealing temperatures (samples 1 and 6), the particles combined into larger agglomerates than those annealed at high temperatures (samples 5 and 10). Moreover, with an increase in the annealing temperature, the fragmentation of agglomerates also increases. Furthermore, particles are formed more densely, and the surface does not look so deep. Annealing of the films improved their morphology and compactness [30].

The film thickness depends on the quantity of ascorbic acid and the annealing temperature. It should be noted that the annealing temperature clearly affects the thickness of the films. The film thickness values are provided in Table 3.

Table 3. Thickness of annealed tin sulfide films.

Sample Number	Sample Thickness	
	The Area of Sample Thickness, nm	The Average Size of Sample Thickness, nm
1	35–590	240
2	30–670	235
3	20–460	230
4	30–470	160
5	90	90
6	50–1900	430
7	40–910	340
8	30–590	250
9	20–450	210
10	100	100

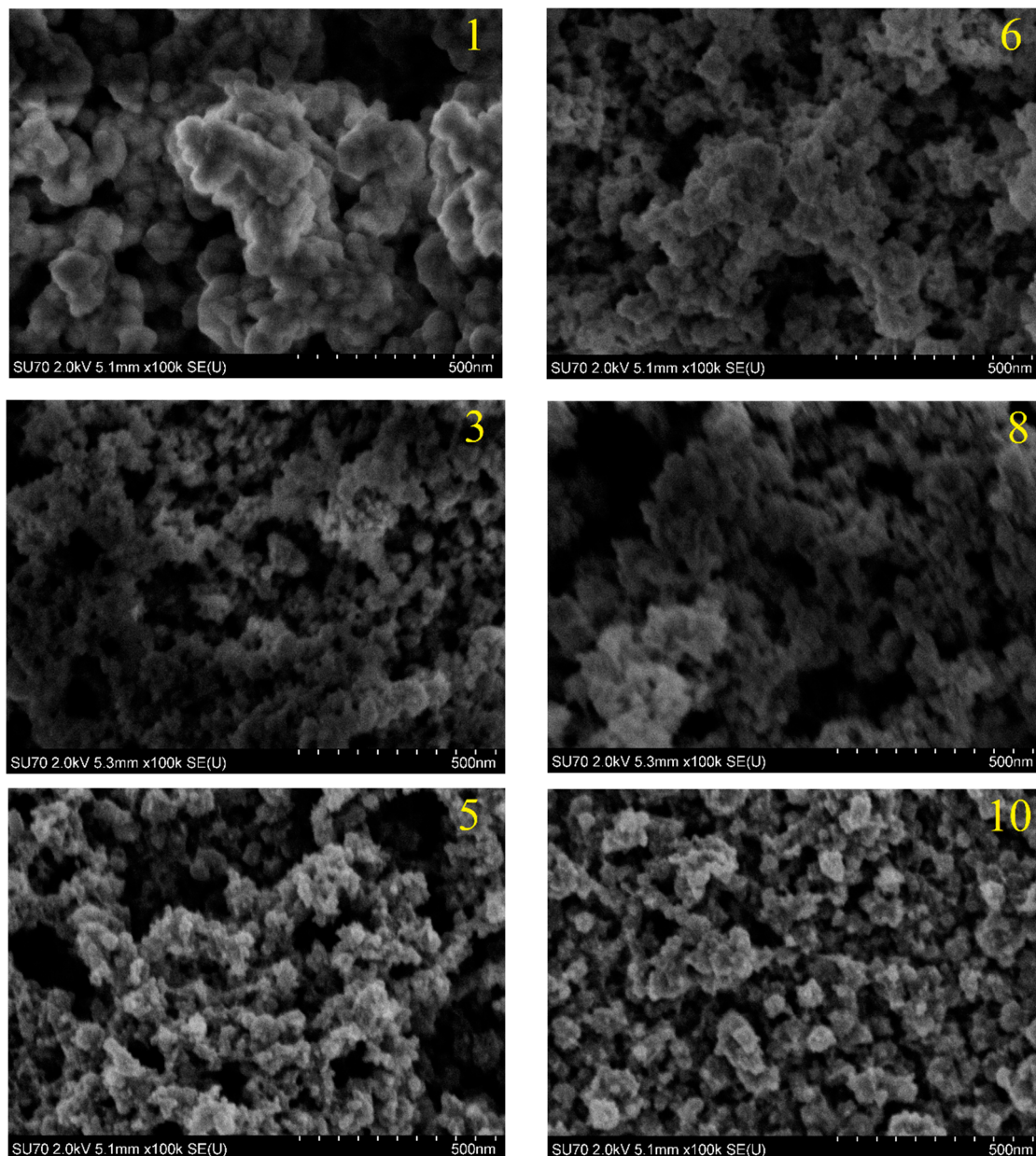


Figure 4. SEM images of samples annealed at 200 °C (1 and 6), 300 °C (3 and 8), and 400 °C (5 and 10).

From the data presented in Table 3, it is evident that the films exhibit roughness, displaying a significant variability in thickness within the same sample. Visual examination with an interferometer confirms that the obtained tin sulfides are uneven, featuring higher elevations and irregularities. It is noteworthy that as the annealing temperature increases, the films become smoother and thinner. The film thickness decreases with higher annealing temperatures, ranging from 240 to 90 nm for samples prepared with 0.8 g of L-ascorbic acid, and 430 to 100 nm for samples prepared with 1.0 g of L-ascorbic acid. Notably, an annealing temperature of 400 °C yields consistent thickness values, indicating smooth films. When comparing films annealed at the same temperature, but with different quantities of L-ascorbic acid (0.8 g vs. 1.0 g), an increase in film thickness is observed, highlighting the impact of the precursor quantity. Temperatures exceeding 120 °C result in L-ascorbic acid degradation [30], indicating the initiation of its destruction within the tin complex [31]. This significant reduction in thickness with increasing annealing temperature is likely associated with the degradation of L-ascorbic acid. The structure becomes more compact, possibly due to intercalated ions or the removal of ascorbic acid fragments.

Bandgap. The optical properties of the thermally treated thin films were analyzed using UV–Vis analysis. The bandgap is a crucial parameter for semiconducting materials, defining their fundamental properties. Typically, semiconductors possess either a direct or indirect bandgap, each with distinct applications. Materials with a direct bandgap find utility in optoelectronic devices such as light-emitting diodes (LEDs) and laser diodes due to their efficient light emission characteristics. Conversely, materials with an indirect bandgap are commonly employed in solar cells and batteries, leveraging their properties for energy conversion and storage applications. [31]. The measurement and analysis of bandgaps are crucial for both fundamental research and practical applications.

The value of the bandgap could affect the annealing temperature. With an increase in the annealing temperature, the bandgap value decreases [32]. Moreover, increasing the quantity of L-ascorbic acid decreases the value of the bandgap.

Figure 5 shows the curves of $(\alpha h\nu)^{1/2}$ as a function of photon energy for samples treated at different post-annealing temperatures. The bandgap values are provided in the legend. Theoretically, tin(II) sulfide has band gap values of 1.3 eV [24], 1.3–1.6 eV [33], 1.35 eV [30], 1.2 eV [34], and 1.1 eV [35]. As mentioned in a previous report [1], the bandgap could affect the substrate mirroring and the secondary phase, which is detected in some samples. SnO₂ typically exhibits relatively high bandgap values ranging from 3.2 to 3.72 eV [36,37], 3.6 eV [38], and 3.7 eV [39], but the bandgap values observed for these samples are notably lower. This suggests that tin sulfide is densely packed on the substrate without substrate mirroring. Additionally, the formation of Sn₂S₃ could influence the bandgap value, as it has been reported to have a bandgap ranging from 1.09 eV [3], 1 eV [40], 1.46–1.64 eV [41], 1.25 eV (indirect) and 1.75 eV (direct, forbidden) [42]. These values closely resemble the bandgap values of SnS.

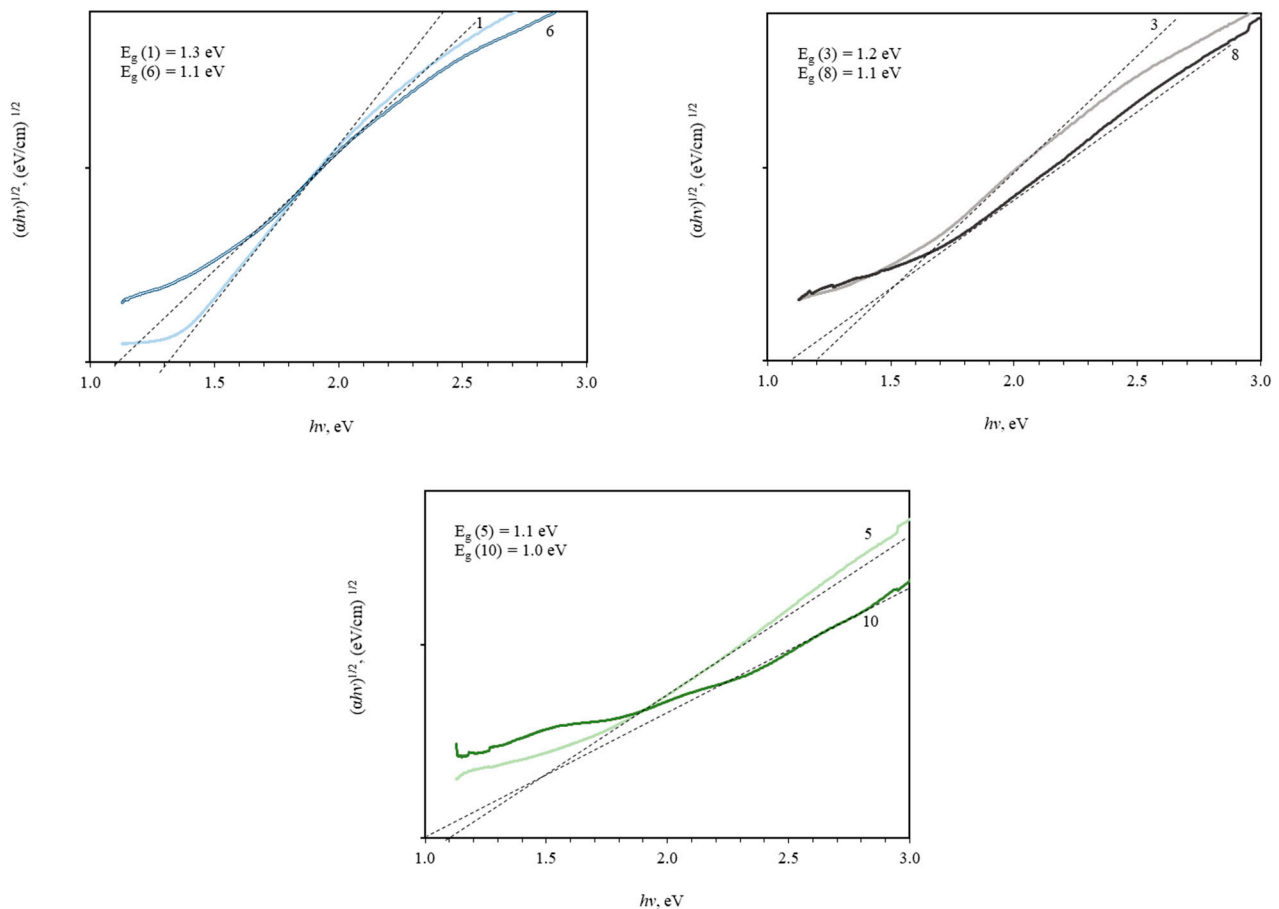


Figure 5. The bandgap values of annealed tin sulfide films.

Furthermore, SnS_2 , which is in Sn_2S_3 , has the possibility to change the value of bandgap. Contrary to other tin sulfides, this one has only indirect transition and the highest value of 2.24 eV [3], 2.4 eV [43], 2.18–2.44 eV [44], and 2.55 eV [45]. As it could be seen, SnS_2 could increase the bandgap value of the samples. Our results showed that the obtained tin sulfide bandgap values are very close to the tin(II) sulfide values. It can be seen in Figure 5 that in this study, there are two factors that affect the bandgap value. Firstly, the quantity of L-ascorbic acid reduces the bandgap, and secondly, this value decreases with an increase in the annealing temperature.

Electrochemical measurements. The shapes of both the CV (Figure 6) and GCD (Figure 7) curves exhibit a dual charge storing mechanism. A triangle-shaped GCD with rectangular CV is usually attributed to electric double-layer capacitors (EDLCs). Meanwhile, Figure 6 shows a tendency to store charge due to Faradaic (redox) reactions, which are more characteristic of pseudocapacitors (PC) [46]. However, no significant anodic or cathodic peak is detected that is so common to PC, and only an even increase in current densities is noticed. According to the results of XRD, Raman mapping, and XPS, the thin films present a multiphase structure with Sn(II) dominating at lower temperatures and Sn(IV) at 300 °C and higher. Tin sulfide thermally treated at 300 °C and 350 °C should be the most active with respect to its current response to the applied potential (Figure S1). This result might be associated with the variety of oxidation states that originate from tin, which has +2 and +4 in these films. The variety of oxidation states is usually a significant factor that affects the performance of pseudocapacitors. However, the phase structure is not necessarily the determining factor for differences in the activities of the films. SnS has a layered structure that is favorable for ion intercalation and therefore demonstrates enhanced capacitive properties [47]. With increasing temperature, the thickness of the films decrease. This is probably because intercalated ions or fragments of ascorbic acid are removed, and the structure becomes more compact. Furthermore, the crystallites can be located more closely with a surface, turning to a smoother one. Optimized porosity is known to facilitate the accessibility of electrolyte ions and increase the active surface area. However, compact films may help overcome unwanted side processes that occur in pores, such as slow diffusion [48]. The results of the CV and GCD techniques suggest that optimal morphology is achieved when the films are annealed at around 300 °C. A summary comparison of the basic energetic parameters of all coatings is given in Table 4. The variation in the quantitative values of the specific capacitance (SC) generally agrees with the regularities of the CV and GCD curves. An increase in temperature results in improved capacitive behavior, but only until 300 °C is reached. Later, with further annealing, the parameters start to worsen. This may indicate a lower capacity for SnS_2 charge accumulation since tin has the highest oxidation state and does not tend to participate in redox reactions. When considering the impact of ascorbic acid, a higher quantity appears to have a positive effect. Presumably, this can be associated with a higher probability of intercalation of ions and organic fragments between the layers during SILAR synthesis. This assumption is also supported by the increase in film thickness when 1.0 g of ascorbic acid is used. When the capacitive behavior of the annealed and prepared films is compared, which has been previously discussed [1], a clear advantage of the annealed films can be seen. The values of the untreated tin sulfide synthesized under the same conditions were 0.36 F g^{-1} (with 0.8 g of ascorbic acid) and 0.15 F g^{-1} (with 1.0 g of ascorbic acid). Such low values are mainly attributed to the more amorphous structure and the presence of Sn_2S_3 , which transforms into SnS and SnS_2 during the heat treatment. Another benefit of thermal treatment worth mentioning is the widening of the potential window. When it changes from 2 V to 3 V, it marks an improvement in the important characteristic of a capacitor in an aqueous electrolyte.

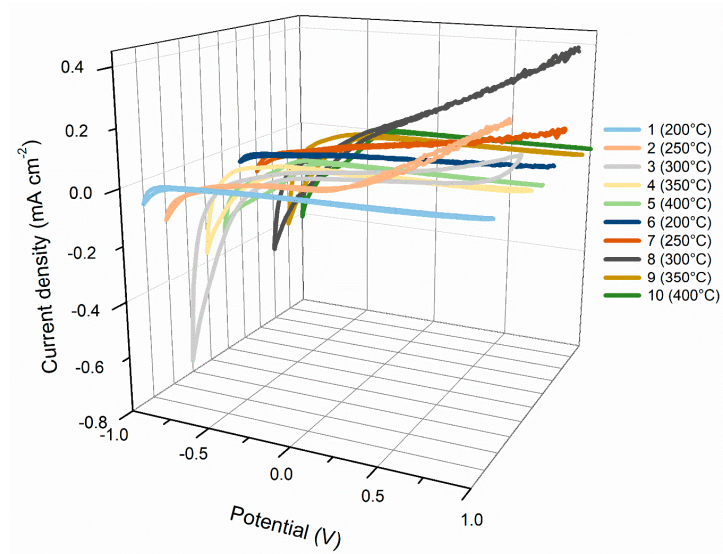


Figure 6. CV curves of the tin sulfide coatings synthesized with 0.8 g of ascorbic acid (1–5) or 1.0 g (6–10) and annealed at different temperatures. Supporting electrolyte—0.1 M NaCl.

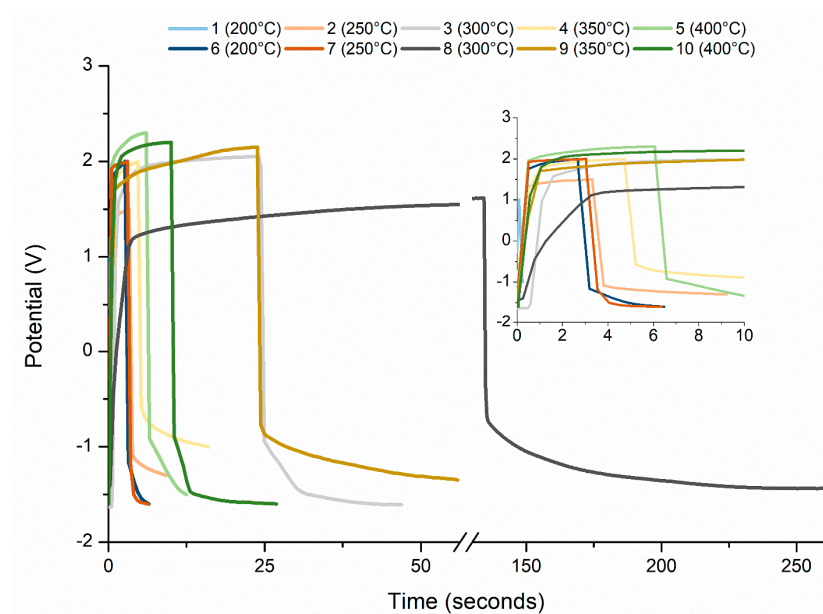


Figure 7. GCD curves of the tin sulfide coatings synthesized with 0.8 g of ascorbic acid (1–5) or 1.0 g (6–10) and annealed at different temperatures. Supporting electrolyte—0.1 M NaCl, applied current density—1 A g⁻¹.

Table 4. The results of energetic parameters for the samples.

Sample Number	Temperature (°C)	SC (F g ⁻¹)	SE (Wh kg ⁻¹)	SP (W kg ⁻¹)
1	200	0.07	0.04	1000
2	250	1.05	1.14	697
3	300	6.90	13.11	1847
4	350	3.50	4.37	1498
5	400	1.71	3.43	1898

Table 4. Cont.

Sample Number	Temperature (°C)	SC (F g ⁻¹)	SE (Wh kg ⁻¹)	SP (W kg ⁻¹)
6	200	1.02	1.84	1800
7	250	1.11	2.00	1798
8	300	39.00	50.06	1521
9	350	12.67	22.17	1773
10	400	4.44	8.90	1899

4. Conclusions

This study investigated the impact of annealing temperature on the structural, optical, and electrochemical properties of tin sulfide films. XRD analysis revealed the formation of mixed sulfide phases, indicating the presence of tin in both the +2 and +4 oxidation states. Raman and XPS analyses confirmed that SnS was predominantly formed at annealing temperatures of 250 °C and below. At 300 °C, the proportions of SnS and SnS₂ were approximately equal, whereas higher annealing temperatures favored the formation of pure SnS₂.

The highest energetic parameters were observed at an annealing temperature of 300 °C. However, as the annealing temperature increased beyond this point, the energetic values declined.

Supplementary Materials: The following supporting information can be downloaded at <https://www.mdpi.com/article/10.3390/coatings14101284/s1>, Figure S1: CV curves of the tin sulfide coatings synthesized with 0.8 g of ascorbic acid (a) or 1.0 g (b) and annealed at different temperatures. Supporting electrolyte—0.1 M NaCl.

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