Synthesis of simple, low cost and benign sol–gel Cu$_{2}$In$_x$Zn$_{1-x}$SnS$_4$ alloy thin films: influence of different rapid thermal annealing conditions and their photovoltaic solar cells†

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Cu$_{2}$In$_x$Zn$_{1-x}$SnS$_4$ (x = 0.4) alloy thin films were synthesized on soda lime glass (SLG) substrate by a simple low-cost sol–gel method followed by a rapid annealing technique. The influence of sulfurization temperature and sulfurization time on the structure, morphology, optical and electrical properties of Cu$_{2}$In$_x$Zn$_{1-x}$SnS$_4$ thin films was investigated in detail. The XRD and Raman results indicated that the crystalline quality of the Cu$_{2}$In$_x$Zn$_{1-x}$SnS$_4$ alloy thin films was improved, accompanied by metal deficiency, particularly tin loss with increasing the sulfurization temperature and sulfurization time. From absorption spectra it is found that the band gaps of all Cu$_{2}$In$_x$Zn$_{1-x}$SnS$_4$ films are smaller than that (1.5 eV) of the pure CZTS film due to In doping, and the band gap of the Cu$_{2}$In$_x$Zn$_{1-x}$SnS$_4$ films can be tuned in the range of 1.38 to 1.19 eV by adjusting the sulfurization temperature and sulfurization time. Hall measurement results showed that all Cu$_{2}$In$_x$Zn$_{1-x}$SnS$_4$ alloy thin films showed p-type conductivity characteristics, the hole concentration decreased and the mobility increased with the increase of sulfurization temperature and sulfurization time, which is attributed to the improvement of the crystalline quality and the reduction of grain boundaries. Finally, the Cu$_{2}$In$_x$Zn$_{1-x}$SnS$_4$ film possessing the best p-type conductivity with a hole concentration of $9.06 \times 10^{16}$ cm$^{-3}$ and a mobility of 3.35 cm$^2$ V$^{-1}$ s$^{-1}$ was obtained at optimized sulfurization condition of 580 °C for 60 min. The solar cell using Cu$_{2}$In$_x$Zn$_{1-x}$SnS$_4$ as the absorber obtained at the optimized sulfurization conditions of 580 °C for 60 min demonstrates a power conversion efficiency of 2.89%. We observed an increment in open circuit voltage by 90 mV. This work shows the promising role of In in overcoming the low Voc issue in Cu-kesterite thin film solar cells.

1. Introduction

Kesterite Cu$_2$ZnSnS$_4$ (CZTS) thin films are being considered as an ideal absorber material for thin film solar cells owing to their wonderful material properties, such as suitable band gap energy of 1.4–1.5 eV, non-toxic constituents and suitable optoelectronic properties. 1,2 CZTS based solar cells have achieved a record device efficiency of 9.2%,3 however this is much lower than that of Cu$_2$ZnSn(S,Se)$_4$ (CZTSSe) (12.6%) and Cu(In,Ga)(S,Se)$_2$ (CIGSSe) (22%) based solar cell devices.4,5 Therefore, to further enhance the conversion efficiency, CZTS thin film solar cells still need a more systematic study. Compared with the device parameters determine the efficiency of CIGSSe and CZTSSe solar cells, it is clear that the lower open-circuit voltage ($V_{oc}$) is currently the major challenge that kesterite devices have to overcome in the near future.6,7 The $V_{oc}$ is relevant to the band gap of CZTS absorber layer.8 Therefore, a lot of endeavors have been done to adjust the band gap of CZTS thin films. For CZTS semiconductor, the band gap can be adjusted by doping Se element in CZTS absorber thin film.9 However, it is very difficult to control precisely the S/Se ratio during the thermal annealing process. Moreover, obtaining a suitable crystal phase and controlling the stoichiometry during preparation still are a great challenge.10 Recent research indicates that partial replacement of the metal ions may be a better way to adapt the band gap of the CZTS.11,12 This idea of partial replacement of the metal ions has been previously explored through substituting Zn with Cd in CZTS.13 Replacing Zn with Cd could adjust the band gap and improve crystallinity of CZTS thin films. Meanwhile, the power conversion efficiency improved significantly from 5.3% to 9.24%, demonstrating partial replacement of the metal ions possesses great potential for enhancing device performance.13 Unfortunately, Cd is recognized as an extremely toxic element,
making it restricted for commercial applications. Hence, it is necessary to find a suitable metal atom for realizing the adjustment of the band gap. The substituting Zn with In atom may be a more advantageous route. Firstly, Zn and In elements have almost similar atomic size. The ionic radius of the Zn$^{2+}$ is 0.74 Å whereas it is about 0.80 Å for In$^{3+}$ (effective ionic radius, IR), so it is more likely to be incorporated into CZTS thin films. Secondly, the substitution of the appropriate number of Zn by In can’t give rise to an obvious change in lattice constant and crystal structure because of the similar radii. Moreover, the 12.7% efficiency of CZTSSe solar cells has got a record open-circuit voltage through applying a double In$_2$S$_3$/CdS emitter, In$_2$S$_3$ buffer was employed as a supplier for indium into the Cs/Ds/CsTeS layers via annealing, which enables the formation of a suitable band alignment between a buffer and an absorber layer, and improves the carrier lifetime as well. However, the real reason of device property enhancement through In$_2$S$_3$ layer has not been demonstrated so far. Therefore, it is necessary to obtain a clear understanding of the real reason of device property enhancement. However, there are few reports about Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ alloy thin films grown by sol–gel technique so far.

In our previous work, we have successfully fabricated the Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ alloy thin films with different In content by a simple sol–gel method. And we have investigated the influence of In content on the structure, electrical and optical properties of Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ alloy thin films in detail. The band gap of the Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ alloy thin films can be tuned in the ranges of 1.45–1.29 eV as In content varied from $x = 0$ to 0.5. According to the study results, it is concluded that the Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ alloy thin films should be an ideal light-absorber material for achieving higher efficiency kesterite solar cells. It is well known that the crystalline quality and properties of the semiconductor films are greatly influenced by its annealing condition, for example, the annealing temperature and annealing time greatly affect the properties of the film, such as the crystallinity, electrical and optical properties. That is to say, as-grown films annealed at different sulfurization temperature or sulfurization time often causes the different variation behavior of the properties of the CZTS film.

In the previous literature, many endeavors are done to vary the annealing condition and its effect on the physical properties of CZTS is demonstrated. However, no almost report is available on the study about the influence of the sulfurization temperature and sulfurization time on structural, optical and electrical properties of Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ thin films. Consequently, we concentrated on the influence of sulfurization temperature and sulfurization time on the structural, optical and electrical properties of Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ thin films. In this paper. The Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ (x = 0.4) alloy thin films were synthesized on SLG substrates by sol–gel method followed by rapid annealing technique. The influence of sulfurization temperature and sulfurization time on the structural, optical and electrical properties of the Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ (x = 0.4) alloy thin films was investigated in detail. The Cu$_{1-x}$Zn$_{1-x}$Sn$_2$S$_4$ (x = 0.4) solar cell obtained at an optimized sulfurization condition of 580 °C for 60 min demonstrates a power conversion efficiency of 2.89% and the $J$–$V$ characteristic was studied carefully.

2. Experimental methods

2.1 Preparation of the Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ alloy thin film

The Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ (x = 0.4) alloy thin film was prepared by a two-step process. The first process was to prepare the Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ alloy thin films on soda lime glass (SLG) substrates by the sol–gel technique. The SLG substrate was rinsed in successive ultrasonic baths of trichloroethane, acetone, methanol and de-ionized water for 15 min, respectively. Thereafter, it was washed using de-ionized water, eventually blow-dried it with nitrogen. The formative solution of Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ precursors were made up of zinc acetate, copper(II) acetate monohydrate, tin(II) chloride dihydrate, indium trichloride, thiourea, monoethanolamine (MEA) and 2-methoxyethanol. The MEA and 2-methoxyethanol were employed as the stabilizer and the solvent, respectively. The mole ratios of Cu, In, Zn and Sn in the precursor solution are 2.125 : 0.9 : 0.6 : 1 : 8, respectively. The solution was stirred at room temperature to completely dissolve the metal compounds and the color was colorless and transparent, which could stabilize for a few weeks in air atmosphere. The Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ thin films were synthesized by spin-coating on the SLG at 3000 rpm for 30 s accompanied by preheating at 300 °C for 5 min in air. The spinning and drying process was repeated for a few times to acquire 1μm thick film. The second procedure was to implement the rapid annealing treatment for as-grown Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ thin films under various sulfurization temperature and sulfurization time in the sulfur atmosphere.

2.2 Device fabrication

To fabricate solar cells with a typical structure, CZTS and Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ (x = 0.4) precursors were deposited on the Mo-coated SLG substrates (Mo layer with a thickness about 1 μm was deposited by DC sputtering) and annealed using the aforementioned method. The deposition of 50 nm thick CdS buffer layer on the absorber was completed by the chemical bath deposition (CBD) method. Then 50 nm i-ZnO and 300 nm ITO were sputtered as the window layers and subsequent the Al grid top electrode was evaporated on the ITO layer. Finally, the devices were mechanically divided into 9 small cells, the active area of the cell is 0.19 cm$^2$. The crystal structure was characterized by the Japan Rigaku D/max-ga X-ray diffractometer (XRD) equipped Cu Kz ($\lambda = 0.15406$ nm) source. Raman spectra were measured by a Renishaw system with an excitation wavelength of 514 nm. The composition and the valence state of the elements in the Cu$_{1-x}$In$_x$Zn$_{1-x}$Sn$_2$S$_4$ thin films was detected by X-ray photoelectron spectroscopy (XPS) using an Al Kz monochromatized source (ESCALAB MARK II, VG Inc.). The transmission electron microscope (TEM) image was acquired with JEM-2100 electron microscope. The scanning electron microscope (SEM) image
was performed by a Hitachi S-4800 equipped with an energy dispersive X-ray spectroscopy (EDS) system (EDAX Genesis 2000). The electrical properties were carried out by a Hall-effect measurement system with van der Pauw configuration at room temperature. The optical properties were analyzed using an UV-Vis-near-infrared (NIR) spectrophotometer (UV-3101PC).

3. Results and discussion

Fig. 1(a) illustrates the XRD diffraction patterns of the four Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) samples (named T500t60, T540t60, T580t60 and T600t60 corresponding to the samples annealed at 500, 540, 580 and 600 °C for 60 min, respectively) obtained on SLG substrates at different sulfurization temperature. For the T500t60, T540t60, T580t60 samples, three sharp distinct diffraction peaks are observed at around 2θ = 28.53°, 47.33° and 56.17°, which are corresponded approximately to the diffraction of the (112), (220) and (312) plane of the kesterite structure CZTS. In addition, some minor peaks become observable at 32.98°, 69.13° and 76.44°, which are corresponded approximately to the diffraction of (200), (008) and (332) plane of CZTS, indicating that the Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) thin films have good crystalline quality. Moreover, except for the CZTS diffraction peak, no other characteristic peak was observed such as indium, zinc or their complex sulfide, suggesting that In doping has not altered the crystal structure of CZTS and will evenly replace the sites of the cation or the interstitial sites in CZTS lattice. In our previous work, we have successfully fabricated the Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ alloy thin films with different In contents by a simple sol–gel method. And we also have investigated the influence of In content on the structure, electrical and optical properties of Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_4$ alloy thin films in detail. According to the study results of XRD, XPS, TEM and EDS, it was proved that the In has been doped into the CZTS and the trivalent In substituted for the divalent Zn site forming the Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ alloy thin films. To describe preferably the influence of annealing condition on the structure properties of Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) thin films, the peak intensity (blue solid circulars) and the full-width half-maximum (FWHM) (red solid stars) are acquired from the (112) peak in the XRD pattern and displayed in Fig. 2. Fig. 2(a) illustrates the structure properties of T500t60, T540t60, T580t60 and T600t60 samples which were annealed at 500, 540, 580, 600 °C for 60 min, respectively. It is seen from Fig. 2(a) that the intensity of (112) peak has been enhanced significantly, manifesting the crystalline quality of Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) thin films are improved as the sulfurization temperature increases up from 500 °C to 580 °C. Interestingly, an opposite variation trend of FWHM was observed simultaneously from Fig. 2(a), and the minimum value of FWHM for Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) thin films was obtained in sample T580t60. When the sulfurization temperature increased to 600 °C, (112) peak intensity of Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) thin film decreases, in addition, a weak peak situated at 31.78° can be observed, as shown in Fig. 1(a), which is ascribed to the (103) diffraction of CuS. It was reported that CZTS can decompose into CuS and ZnS at higher sulfurization temperature. Secondary phase of CuS appeared primarily due to the thermal dissociation of CZTS into binary or ternary phases at higher sulfurization temperature, which reflects the tin loss at higher sulfurization temperature. This also indicates that the higher sulfurization temperature and the presence of secondary phases seems to cause deterioration of the
crystalline quality of the Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) thin films. Consequently, it is reasonable to conclude that the T580t60 sample possesses the best crystalline quality.

Besides the sulfurization temperature, the sulfurization time is also regarded as one of the susceptible elements for effecting the structure and crystallinity of Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) thin films during annealing process. Therefore, we also investigate the influence of the sulfurization time on structure and crystal quality of Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) thin films, and the sulfurization temperature is adopted as 580 °C, which is in the optimized sulfurization temperature. Fig. 1(b) illustrates the XRD patterns of the five Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) samples (named T580t15, T580t30, T580t45, T580t60 and T580t75) which were annealed at 580 °C under various sulfurization time of 15, 30, 45, 60, 75 min, respectively. For the T580t15, T580t30, T580t45 and T580t60 samples, it is found that the four samples have strong diffraction peaks corresponding to (112), (200), (220), (312), (008) and (332) planes of CZTS, particularly, which shows a preferential orientation along the (112) direction implying that the four samples are composed of the CZTS single phase with tetragonal kesterite structure. The intensity and FWHM values of (112) peaks of the CZTS thin films annealed at various sulfurization time were calculated and displayed in Fig. 2(b). It is found from Fig. 2(b) that the intensity of (112) peak has been enhanced significantly, manifesting the crystalline quality of Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) thin films are improved as the sulfurization time increases up from 15 to 60 min. It is observed that (112) peak intensity of Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) thin films reaches a maximum value and the corresponding FWHM displays a minimum value of 0.125 under the sulfurization time of 60 min, indicating that the crystalline quality of Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) film has almost reached the best level under the sulfurization time of 60 min. As the sulfurization time increased to 75 min, a weak peak appeared at 31.8°, which is attributed to the (103) peak of the CuS phase (PDF #06-0464). This phenomenon is associated with the fact that the CZTS decompose into ternary or binary phases at higher temperature, exactly as the earlier explanation for similar phenomenon of XRD patterns mentioned in Fig. 1(a). According to the results of XRD, we hold perspective that the T580t60 sample is the optimal Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ thin film which has the best structure properties.

Because the area of single phase is very limited in CZTS system with Zn-rich and Cu-poor conditions, the secondary phases arises easily in CZTS thin films synthesized by different methods. It is noticed that three XRD peaks located at 2θ = 29.3°, 48.8° and 57.9° of CZTS overlap with those of copper sulfide and zinc sulfide. Thus, the phase purity and crystalline quality are often discerned by XRD and Raman measurements. Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) alloy thin films were further characterized by Raman measurement with a 514 nm excitation wavelength to detect the possible secondary phases, as shown in Fig. 3. Panels (a) and (b) of Fig. 3 describe the Raman patterns of the Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) thin films prepared under various sulfurization temperature and sulfurization time. Fig. 3(a) presents the Raman patterns of Cu$_2$In$_{1-x}$Zn$_x$Sn$_4$ ($x = 0.4$) thin films including T500t60, T540t60, T580t60 and T600t60 samples which were annealed under different sulfurization temperature of 500, 540, 580, 600 °C for 60 min, respectively. It is observed that the T500t60, T540t60 and T580t60 samples exhibit a dominated peak located around at 331 cm$^{-1}$, which corresponded approximately to the Raman peak of 333 cm$^{-1}$ assigned to the A1 vibrational mode from the CZTS phase and other secondary phases were not detected. This dominated peak arises from the A1 vibration mode of the lattice, where only S-anions are involved. It illustrates that the T500t60, T540t60 and T580t60 samples consist of a single phase of kesterite CZTS. As the sulfurization temperature increased up to 600 °C, a weak peak is observed at ~472 cm$^{-1}$, which is attributed to the Cu$_2$S$_4$ phase. This phenomenon also verifies the decomposition of the CZTS arising from the loss of the tin. This result is
consistent with the XRD results. Fig. 3(b) presents the Raman patterns of Cu$_2$In$_x$Zn$_{1-x}$SnS$_4$ ($x = 0.4$) thin films including T580t15, T580t30, T580t45, T580t60 and T580t75 samples which were annealed at 580 °C under various sulfurization time of 15, 30, 45, 60, 75 min, respectively. In the Raman spectra, an evident major peak at 331 cm$^{-1}$ can be clearly observed for all samples expect for the T580t75 sample. The peaks can be attributed to CZTS, and other secondary phases were not detected. It illustrates that all samples expect for T580t75 consist of a single phase of kesterite CZTS. For T580t75 sample, one extra peak at 472 cm$^{-1}$ corresponding to Cu$_2$S can be seen. This can be attributed to the longer sulfurization time, which easily allows the formation of secondary phase of Cu$_2$S. Raman measurements with a 325 nm excitation wavelength was used to further identify the existence of ZnS, as shown in Fig. 3 insert. The peaks at 343, 695 and 1040 cm$^{-1}$ could be assigned to R1LO, R2LO and R3LO vibrational modes of ZnS, respectively. By analyzing Raman spectra, it can be found that the intensity of the ZnS Raman peaks decreases with increasing sulfurization temperature and sulfurization time, as shown in Fig. 3 insert, indicating a decrease of ZnS content, which are in good agreement with the reported literatures. Based on the results of XRD and Raman mentioned above, taking account of crystalline quality of Cu$_2$In$_x$Zn$_{1-x}$SnS$_4$ ($x = 0.4$) thin films, it is deduced that the optimized sulfurization temperature and sulfurization time should be set to 580 °C and 60 min, respectively.

So as to ascertain the phase compositions and valence states of the composing elements, the XPS spectra were measured for the T580t60 sample and shown in Fig. 4. Fig. 4(a) is the full-scan XPS spectrum (0–1200 eV), which showed peaks related to C, Cu, Zn, Sn, S and In. The high-resolution scan XPS spectra of the Cu 2p, Zn 2p, Sn 3d, S 2p and In 3d from the T580t60 sample are shown in Fig. 4(b)-(f), respectively. As shown in Fig. 4(b), the narrow scan XPS spectrum of the Cu 2p displays two peaks at 931.2 eV and 951.7 eV, corresponding to Cu 2p$_{3/2}$ and Cu 2p$_{1/2}$ with a splitting of 20.5 eV, which is consistent with the standard splitting value of Cu$.^3$ So, it can be concluded that Cu exists in the Cu$^+$ state, which is acquired from the reduction of Cu$^{2+}$. The Sn 3d$_{5/2}$ and Sn 3d$_{3/2}$ peaks are revealed at 485.4 eV and 493.5 eV, as shown in Fig. 3(d). The binding energy of the Sn 3d is nearly consistent with that of Sn$^{4+}$ in CZTS alloy thin film. It is observed from Fig. 4(e) that the peaks of S 2p are situated at 160.2 eV and 162.8 eV, which is in good agreement with the binding energy of the S in CZTS.$^{34}$ The binding energies of Zn$^{2+}$, Cu$^+$, Sn$^{4+}$ and sulfides are consistent with the reported values in CZTS alloy thin film. Fig. 4(f) depicts the In 3d XPS spectrum of the T580t60 sample. The two strong characteristic peaks at 443.4 eV and 451.0 eV are ascribed to the core levels of In 3d$_{5/2}$ and In 3d$_{3/2}$, respectively.$^{35}$ The binding energy of the In 3d suggests that the chemical value of In is +3. No else XPS peak of the metallic In or
other valance state was detected, implying that In has been doped into CZTS successfully.

To further determine the detailed microscopic structure of the Cu$_2$In$_x$Zn$_{1-x}$SnS$_4$ (x = 0, x = 0.4) thin films, the TEM measurements were performed. Fig. 5 depicts the TEM images of (a) Cu$_2$In$_x$Zn$_{1-x}$SnS$_4$ (x = 0) thin film (CZTS sample) synthesized under the identical experimental conditions as T580t60 sample, (b) Cu$_2$In$_x$Zn$_{1-x}$SnS$_4$ (x = 0.4) thin film (T580t60 sample), respectively. It can be seen from Fig. 5(a) that the measured interplanar spacing of the crystallographic plane is 0.34 nm, which accords well with the interplanar spacing of the (112) plane of CZTS with tetragonal kesterite structure (JCPDS-26-05757). Additionally, the selected-area electron diffraction (SAED) in Fig. 5(c) shows typical polycrystalline diffraction for the sample, which reveals highly crystalline materials. Fig. 5(b) shows that Cu$_2$In$_x$Zn$_{1-x}$SnS$_4$ (x = 0.4) thin films (T580t60 sample) are arrayed in the (112) direction, and a slightly larger lattice spacing of 0.47 nm in comparison with that (0.34 nm) of CZTS film was measured, which is associated approximately with the spacing of (112) crystal plane of CZTS. It is concluded that some Zn sites in CZTS crystal lattice were substituted by In atoms to form the Cu$_2$In$_x$Zn$_{1-x}$SnS$_4$ thin films, since In has a larger ionic radius (0.80 Å) than Zn (0.74 Å), which is consistent with the results of XPS and XRD analysis. The SAED pattern in Fig. 5(d) suggests that the nanocantilever of the sample is single crystalline tetragonal kesterite because of the well-regulated dot pattern. This phenomenon indicates that the T580t60 sample possesses a relatively preferable structure quality, which is consistent with the earlier opinion proposed from the results of XRD analysis.
It is well known that the optical and electrical properties of CZTS thin films display a strong rely on the stoichiometry of Cu, Zn, Sn and S in CZTS thin film. Well-controlled atomic contents of Cu, Zn, Sn and S in CZTS thin film has become a key issue for preparing the kesterite CZTS solar cells with high-performance. In order to check the composition and concentration of elements for Cu₂In₉₋ₓZnₓSn₄ₓ (x = 0.4) thin films. Energy Dispersive Spectrometry (EDS) was performed for the Cu₂In₉₋ₓZnₓSn₄ₓ (x = 0.4) thin films annealed at different sulfurization temperature and sulfurization time. Table 1 summarizes the composition ratios of the Cu, Zn, Sn, In, S, Cu/(Zn + In + Sn) and Sn/(In + Zn) in the Cu₂In₉₋ₓZnₓSn₄ₓ (x = 0.4) thin films obtained at different sulfurization temperature and sulfurization time. The EDS results verified the existence of Cu, Zn, Sn, In and S elements in the prepared films, as shown in Table 1. The atomic percentages of Sn decreased, while the atomic percentages of Cu increased and the atomic percentages of Zn remained relatively constant with the sulfurization temperature and sulfurization time increasing. It manifests that the loss of Sn element during the sulfurization process, which is in good agreement with the opinion proposed from the analysis of XRD results. It also was found that the atomic percentages of Zn and Cu increased and the atomic percentages of Sn decreased in the Cu₂In₉₋ₓZnₓSn₄ₓ (x = 0.4) thin films with increasing the sulfurization temperature and sulfurization time, leading to the ratios of Cu/(Zn + In + Sn) and (Zn + In)/Sn in the film increased, as shown in Table 1.

Well-adopted sulfurization temperature and sulfurization time are crucial to acquire the large grain and a small amount voids in CZTS thin film. Fig. 6 shows the SEM surface images of Cu₂In₉₋ₓZnₓSn₄ₓ thin films annealed at different sulfurization temperature (a) 500 °C, (b) 540 °C, (c) 580 °C and (d) 600 °C for 60 min and annealed at 580 °C with different sulfurization time ((e) 15 min, (f) 30 min, (g) 45 min, (h) 60 min and (i) 75 min). The remarkable influence of the sulfurization temperature and sulfurization time on the microstructure of Cu₂In₉₋ₓZnₓSn₄ₓ thin films is clearly observed. With the sulfurization temperature increasing from 500 to 580 °C, the grain size of Cu₂In₉₋ₓZnₓSn₄ₓ thin films increased and the surface of the thin films became smooth and dense, as shown in Fig. 6(a–c). At higher temperature, the ions have a greater reactivity, which facilitates the growth of the nanocrystals. However, the grain size decreased with sequentially increasing sulfurization temperature up to 600 °C, as shown in Fig. 6(d), which may be ascribed to the loss of Sn element during the sulfurization process. Formation of voids on the surface is evident for the T500t60, T540t60 and T600t60 samples. It is clear that the relatively compact surface with densely packed grains is observed for sample T580t60, as shown in Fig. 6(c). The SEM measurements of the Cu₂In₉₋ₓZnₓSn₄ₓ thin films obtained at different sulfurization time of 15, 30, 45, 60 and 75 min are shown in Fig. 6(e–i). It showed that the Cu₂In₉₋ₓZnₓSn₄ₓ thin film obtained at the sulfurization time of 15 min is composed of nanometer grains, and the grains also stack loosely. Moreover, there are some holes in the Cu₂In₉₋ₓZnₓSn₄ₓ thin film surface, as shown in Fig. 6(e). When increasing the sulfurization time to 30 min, the grain of Cu₂In₉₋ₓZnₓSn₄ₓ thin film increases and gets compact, as shown in Fig. 6(f). However, the surface of Cu₂In₉₋ₓZnₓSn₄ₓ thin film shows some smaller grains, which are not favorable to the formation of the CZTS solar cells with high efficiency. As the sulfurization time increased to 60 min, the grains grew sharply to micrometer scale and stack densely, as shown in Fig. 6(h). For

### Table 1

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<td>0.87</td>
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<td>46.37</td>
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<tr>
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<td>10.29</td>
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<tr>
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<tr>
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<td>9.82</td>
<td>4.24</td>
<td>44.29</td>
<td>0.92</td>
<td>1.94</td>
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Fig. 6 SEM images of the Cu₂In₉₋ₓZnₓSn₄ₓ (x = 0.4) alloy thin films annealed at 500 °C (a), 540 °C (b), 580 °C (c), 600 °C (d), 15 min (e), 30 min (f), 45 min (g), 60 min (h), and 75 min (i).
solar cell application, it is known that the CZTS absorber thin film with large grain size, smooth surface and less grain boundaries are desirable to the preparation of high PCE solar cell, because the larger grains will promote the transport of photo-generated carriers, which could lead to minimize grain boundary recombination. So, the T580t60 sample is well suited for application as the absorption layer in photovoltaic devices. Further increasing the sulfurization time to 75 min, the grain size decreases and the grains stack loosely, as shown in Fig. 6(i). The noticeable change for Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ ($x = 0.4$) morphology corresponds well to the XRD results discussed above. The decrease of the grain size may be because of the loss of Sn during the sulfurization process, and there were some secondary phases (like CuS) for T580t75 sample.

Table 2 depicts the electrical properties of as-grown CZTS, as-grown Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ ($x = 0.4$) and Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ ($x = 0.4$) thin films which were annealed at various sulfurization temperature and sulfurization time, respectively. In order to better compare and clarify the influence of annealing condition and In doping on electrical properties of Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ ($x = 0.4$) thin films, pure CZTS thin films were deposited on SLG substrates under the same experimental conditions as Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ ($x = 0.4$) thin film, which were showed in Table 2, too. As summarized in Table 2, the as-grown CZTS thin film exhibits a bad p-type conductivity due to its numerous intrinsic defects. With the incorporation of In, the as-grown Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ ($x = 0.4$) thin film remained p-type behavior with a carrier concentration of 9.46 $\times$ 10$^{17}$ cm$^{-3}$, a mobility of 0.91 cm$^2$ V$^{-1}$ s$^{-1}$ and a resistivity of 8.99 $\times$ 10$^{2}$ $\Omega$ cm, which are considerably weaker than that of pure CZTS. According to the results of XRD, XPS and TEM mentioned above, In has been doped into the CZTS and the trivalent In substituted for the divalent Zn site to form the In$_{2n}$ donor, then introducing some electrons into this system. For this reason, the hole concentration of Cu$_2$Zn$_{1-x}$Sn$_x$S$_4$ would be decreased by the electrons that In$_{2n}$ donor provided, consequently leading to the worse p-type conduction property.

Table 2 also displays the electrical properties of the Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ ($x = 0.4$) thin films annealed at different sulfurization temperature. It was found that all Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ ($x = 0.4$) thin films showed the p-type (hole mediated) conductivity behavior. It has been reported that Cu substituting Zn site (Cu$_{2n}$) are the main of acceptor in stoichiometric CZTS due to their low formation energies and shallow level, which is responsible for the p-type conduction of CZTS. It was observed that the carrier concentration decreased from 5.47 $\times$ 10$^{17}$ to 9.06 $\times$ 10$^{16}$ cm$^{-3}$ as the sulfurization temperature increased from 500 to 580 $^\circ$C. However, the change tendency of the resistivity and the mobility is contrary with that of the carrier concentration. This may be due to that with the sulfurization temperature increasing, the crystalline quality is improved and the grain boundaries become less, leading to the reduction of the carrier concentration and increment of the mobility.

Xiao et al. have pointed out that the conduction characteristics of the CZTS thin films are also affected greatly by sulfurization time. Hence, the effect of sulfurization time on the electrical properties of Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ ($x = 0.4$) thin films was investigated under the fixed sulfurization temperature of 580 $^\circ$C, as displayed in Table 2. It is observed that all the Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ thin films synthesized at different sulfurization time behave p-type conductivity. Also, it was found that the carrier concentration of the film decreased obviously from 1.32 $\times$ 10$^{17}$ to 9.06 $\times$ 10$^{16}$ cm$^{-3}$ with the sulfurization time increasing from 15 to 60 min, attributed to the increase of the atomic ratio of Cu/[Zn + Sn] in the film, which has been proved in the foregoing EDS result. The increase of the atomic ratio of Cu/[Zn + Sn] could produce the decrease of the Cu$_{2n}$ antisite acceptor defect and the increase of Cu$_{2n}$ + Sn$_{2n}$ deep donor defect, leading to the reduction of the carrier concentration.

The Hall mobility of the Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ thin films were increased from 1.22 to 3.35 cm$^2$ V$^{-1}$ s$^{-1}$, which is possibly ascribed to the fact that the crystalline quality is enhanced and the grain boundaries become less as the sulfurization time increases.

For the sake of studying the effect of the sulfurization temperature and sulfurization time on the band gap ($E_g$) of Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ ($x = 0.4$) thin films, the absorption characteristic of the Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ ($x = 0.4$) prepared at different sulfurization temperature and sulfurization time were researched by an UV-Vis-NIR spectrophotometer. According the optical absorption spectra, the band gap value of the Cu$_2$In$_{1-x}$Zn$_{1-x}$Sn$_x$S$_4$ ($x = 0.4$) is obtained by using the relation:

$$ (a h \nu)^{1/n} = B(h \nu - E_g)^{1/2}/h \nu $$

where $a$ is the absorption coefficient, $h$ is Planck’s constant, $\nu$ is the frequency of light, $E_g$ is the band gap energy, and $n$ is the Tauc’s index (1 for direct gap and 2/3 for indirect gap).
where \( B \) is a constant, \( \alpha \) is the absorption coefficient and the \( hv \) is photon energy. The plot of \((\alpha hv)^2 \) against \( hv \) for \( \text{Cu}_2\text{In}_{x-1}\text{Zn}_1\text{Sn}_x \) (\( x = 0.4 \)) thin films annealed at various sulfurization temperature and sulfurization time has been illustrated in Fig. 7. By utilizing the eqn (1) and the data in Fig. 7, the band gap of the \( \text{Cu}_2\text{In}_{x-1}\text{Zn}_1\text{Sn}_x \) (\( x = 0.4 \)) prepared at different sulfurization temperature are estimated to be 1.38, 1.26 and 1.19 eV for \( T500t60, T540t60 \) and \( T580t60 \) samples, respectively, which are smaller than that (1.5 eV) of pure CZTS thin film due to the In doping. Base on the well-known equation in polynomial form, the \( E_g \) variations versus the sulfurization temperature (denoted as \( x \) in the inset here) was fitted into a quadratic function of \( E_g(x) = 1.52 - 0.15x + 0.17x^2 \) (eV), as shown in the inset of Fig. 7(a). It can be found that the \( E_g \) is decreased from 1.38 to 1.19 eV as the sulfurization temperature increases from 500 to 580 \( ^\circ \)C, by utilizing the data in Fig. 7(b), the absorption spectra of \( \text{Cu}_2\text{In}_{x-1}\text{Zn}_1\text{Sn}_x \) (\( x = 0.4 \)) films which were annealed at 580 \( ^\circ \)C for 15, 30, 45 and 60 min, respectively. The band gaps of the \( T580t15, T580t30 \), \( T580t45 \) and \( T580t60 \) samples were determined as 1.35, 1.24, 1.23 and 1.19 eV, respectively. Base on the well-known equation in polynomial form, the \( E_g \) variations versus sulfurization time (denoted as \( x \) in the inset here) was fitted into a quadratic function of \( E_g(x) = 1.46 - 0.13x + 0.01x^2 \) (eV), as shown in the inset of Fig. 7(b). It can be found that the \( E_g \) also is decreased from 1.35 to 1.19 eV as the sulfurization time increases from 15 to 60 min. The band gap of the \( \text{Cu}_2\text{In}_{x-1}\text{Zn}_1\text{Sn}_x \) (\( x = 0.4 \)) alloy thin films was decreased with increasing the sulfurization temperature and sulfurization time, the reasons may be as follows: one is that, residual organic precursor compounds in the films were decreased with the sulfurization temperature and sulfurization time increasing, leading to the reduction of the band gap.\(^{39}\) The other one may be due to the increase of \( \text{Cu}/(\text{Zn} + \text{Sn}) \) atomic ratio with increasing the sulfurization temperature and sulfurization time, which has been verified in the results of the EDS measurement. It is reported that Cu-rich CZTS films have smaller optical band gap than Cu-poor CZTS films. It is also reported that the band gap of CZTSe decrease with the \( \text{Cu}/(\text{Zn} + \text{Sn}) \) atomic ratio increasing.\(^{41}\) They ascribed this to the variations in the p–d hybridization degree between the Cu d-level and Se p-level.
Table 3 Parameters of the device performance

<table>
<thead>
<tr>
<th>Device</th>
<th>Active area</th>
<th>$V_{oc}$ (V)</th>
<th>$J_{sc}$ (mA cm$^{-2}$)</th>
<th>FF</th>
<th>PCE</th>
</tr>
</thead>
<tbody>
<tr>
<td>CZTS</td>
<td>0.19 cm$^2$</td>
<td>0.25</td>
<td>18.69</td>
<td>30.59</td>
<td>1.44</td>
</tr>
<tr>
<td>Cu$<em>2$In$</em>{2-x}$Zn$_{1+x}$Sn$_4$ (x = 0.4)</td>
<td>0.19 cm$^2$</td>
<td>0.34</td>
<td>25.73</td>
<td>32.62</td>
<td>2.89</td>
</tr>
</tbody>
</table>

Fig. 8 shows the current–voltage ($J$–$V$) characteristics of Cu$_2$In$_{2-x}$Zn$_{1+x}$Sn$_4$ (x = 0) device (CZTS) (red solid circulars) and Cu$_2$In$_{2-x}$Zn$_{1+x}$Sn$_4$ (x = 0.4) (blue solid stars) device which were synthesized under the identical experimental conditions, respectively. All device parameters like $J_{sc}$, $V_{oc}$ and fill-factor corresponding to the Cu$_2$In$_{2-x}$Zn$_{1+x}$Sn$_4$ (x = 0.4) absorber are much higher compared to the pure CZTS absorber and yields a power conversion efficiency of 2.89% under 1 sun condition, as shown in Table 3. From $J$–$V$ curves, the series resistance ($R_s$) is calculated to be ~42 $\Omega$ cm$^2$ and shunt resistance ($R_{sh}$) is ~137 $\Omega$ cm$^2$ in the Cu$_2$In$_{2-x}$Zn$_{1+x}$Sn$_4$ (x = 0.4) device whereas for CZTS device $R_s$ ~ 65 $\Omega$ cm$^2$ and $R_{sh}$ ~ 88 $\Omega$ cm$^2$ for the CZTS device. The decrease in series resistances and increase of shunt resistance could be a result of increased grain size and reduction of defect density in the Cu$_2$In$_{2-x}$Zn$_{1+x}$Sn$_4$ (x = 0.4) device. These improvements in series and shunt resistance results in the improved current density and fill factor. We also observed a significant improvement in $V_{oc}$ of about 90 mV. This is mainly due to the reduction of Cu$_{2-z}$ antisite defects which could reduce band-tailing and yield high band bending at the interface of the absorber and buffer layer. This yielded an enhancement in $V_{oc}$, $J_{sc}$ and fill factor of the device. Thus, In substitution improves the overall performance of CZTS solar cell and should be pursued as a solution to the $V_{oc}$ deficit issue in the Cu-kesterite thin film solar cells.

4. Conclusion

In conclusion, the Cu$_2$In$_{2-x}$Zn$_{1+x}$Sn$_4$ (x = 0.4) alloy thin films have been successfully synthesized on SLG substrates at different sulfurization temperature and sulfurization time by a simple low-cost sol–gel method followed by rapid annealing technique. The structure, optical and electrical properties of Cu$_2$In$_{2-x}$Zn$_{1+x}$Sn$_4$ (x = 0.4) alloy thin films are greatly dependent on sulfurization temperature and sulfurization time. XRD results indicated the Cu$_2$In$_{2-x}$Zn$_{1+x}$Sn$_4$ (x = 0.4) alloy thin film has the best crystalline quality at the optimized sulfurization condition of 580 °C for 60 min, respectively. Hall measurements results showed that all Cu$_2$In$_{2-x}$Zn$_{1+x}$Sn$_4$ alloy thin films showed p-type conduction characteristics, the hole concentration decreased and the mobility increased with the increase of sulfurization temperature and sulfurization time, ascribing to the improvement of the crystalline quality and the reduction of the grain boundaries. The optimal p-type Cu$_2$In$_{2-x}$Zn$_{1+x}$Sn$_4$ (x = 0.4) film with a mobility of 3.35 cm$^2$ V$^{-1}$ s$^{-1}$ and carrier concentration of 9.66 $\times$ 10$^{16}$ cm$^{-3}$ also was obtained at the optimized sulfurization condition of 580 °C for 60 min, respectively. The $E_g$ of all the Cu$_2$In$_{2-x}$Zn$_{1+x}$Sn$_4$ alloy thin films is smaller than that of the undoped CZTS on account of the In doping. The $E_g$ of Cu$_2$In$_{2-x}$Zn$_{1+x}$Sn$_4$ (x = 0.4) alloy thin films is decreased from 1.38 to 1.19 eV with the increase of the sulfurization temperature and sulfurization time, which is attributed to the decrease of the residual organic precursor compounds and the increase of Cu/(Zn + Sn) atomic ratio in the films. The Cu$_2$In$_{2-x}$Zn$_{1+x}$Sn$_4$ (x = 0.4) solar cell obtained at an optimized sulfurization condition of 580 °C for 60 min demonstrates a power conversion efficiency of 2.89%.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors would like to thank financial support of the National Natural Science Foundation of China under Grant No. 61505067, 11254001, 61475063, 61605059, 21576111, 61378085. The National Key Research and Development Program of China under Grant No. 2017YFF0108607, the Program for the development of Science and Technology of Jilin province Grant No. 20150520086JH.

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