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# Thermal analysis of novel third-generation phase-change materials with zinc as a chemical modifier

Vishnu Saraswat,<sup>a</sup> Shiv Kumar Pal,<sup>a</sup> N. Mehta,<sup>b</sup> <sup>\*a</sup> Arun Kumar<sup>b</sup> and M. M. A. Imran<sup>c</sup>

The thermal analysis in the present work is done to analyze the glass/crystal phase transformation in a newly synthesized glassy system (*i.e.*, glassy SeTeSnZn alloys) consisting of chalcogenides Se and Te as major elements, Sn as a third element of the parent alloy and Zn as a chemical modifier. The role of increasing the Zn concentration at the cost of Se has been understood by correlating the kinematics of structural relaxation during the glass transition phenomenon and devitrification during the crystallization phenomenon in the chalcogenide glasses (ChGs) of the quaternary STSZ [*i.e.*,  $\text{Se}_{78-x}\text{Zn}_x\text{Te}_{20}\text{Sn}_2$  ( $0 \leq x \leq 6$ )] system and their different physicochemical properties. A noticeable rise in the crystallization rate is observed after the addition of Zn in the parent SeTeSn glass. With the rise in the zinc content, the values of average heat of atomization and overall mean bond energy are found to be decreased with the decrease in cohesive energy of samples. An inverse correlation is observed between the thermal stability parameter and the enthalpy released during the glass/crystalline phase transformation.

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## 1. Introduction

In recent years, the use of zinc as a dopant or metallic modifier in different advanced functional materials has been enhanced<sup>1–6</sup> because of its efficient role in controlling the properties of the parent materials and achieving fascinating and fruitful results. The zinc chalcogenides have attracted the attention of experimentalists working on metal chalcogenides because of their robust bonding, large energy gaps, high enthalpies of formation, and melting points as compared to most of the other metallic chalcogenides formed by using other elements placed in the II–VI.<sup>1–10</sup> The crystal dynamics of zinc chalcogenides<sup>4–6</sup> suggest their utilization in various applications.<sup>4–10</sup> Particularly, the recent literature survey of the last decade reveals that zinc is one of the building elements of phase change materials (PCMs) for use in phase change optical memory.<sup>11–17</sup> These features compelled various research groups to utilize Zn as a modifier for preparing multicomponent (binary and ternary) ChGs.<sup>18–35</sup>

Mehta *et al.*<sup>18</sup> compared the effective values of thermal diffusivity and corresponding thermal conductivity of binary  $\text{Se}_{90}\text{Zn}_{10}$  and  $\text{Te}_{90}\text{Zn}_{10}$  alloys and found that both parameters are lower for  $\text{Se}_{90}\text{Zn}_{10}$  alloy as compared to  $\text{Te}_{90}\text{Zn}_{10}$  alloy. The outcome of Zn addition on the number of defect states per unit volume in the parent  $\text{Se}_{70}\text{Te}_{30}$  sample was analyzed by Srivastava *et al.*<sup>19</sup> by performing a.c. conductivity measurements.

Singh *et al.*<sup>20,21</sup> observed the Meyer–Neldel relation between the rate constant and activation energy in thermally controlled crystallization during the calorimetric studies of ternary Se–In–Zn glasses. Nasir *et al.*<sup>22</sup> reported the thermal analysis of the kinematics of glass transition as well as the recrystallization of binary  $\text{Se}_{98-x}\text{Zn}_x$  glasses. They noticed the significant impact of zinc on the glass/crystal phase transformation of selenium glass. Nasir and Zulfequar<sup>23,24</sup> performed the electrical measurement too in these glasses to estimate the number of localized states per unit volume and their dielectric behavior. They found the enhancement in the corresponding number density when the concentration of Zn content rises in the selenium glass. They also noticed a sufficient dielectric dispersion as a consequence of Zn incorporation.

The thermophysical and optical characterization of ternary  $\text{Se}_{98-x}\text{In}_x\text{Zn}_2$  and  $\text{Se}_{93-x}\text{Te}_5\text{In}_x\text{Zn}_2$  ( $0 \leq x \leq 10$ ) was done by Singh<sup>25,26</sup> for the comparison between these ternary and quaternary samples. Kishore *et al.*<sup>27</sup> used FTIR spectra of ternary  $\text{Se}_{72-x}\text{S}_{28}\text{Zn}_x$  ( $0 \leq x \leq 4$ ) for analyzing the influence of Zn on the optical parameters of parent  $\text{Se}_{72}\text{S}_{28}$  glass and observed the significant decrement in the optical gap by a small rise in the content of zinc. Ahmad *et al.*<sup>28</sup> used silver ion radiation to expose amorphous thin films of the SeS<sub>2</sub>Zn sample and investigated the irradiation effect on optical, electrical, and structural properties. Abbady *et al.*<sup>29</sup> compared the effect of Zn over two other additives (Ge and Ga) on the optical linearity/non-linearity of selenium by preparing thin-film samples. They found that most of the linear/non-linear optical parameters are optimized for the addition of Zn.

Alwany *et al.*<sup>30</sup> reported the optical properties of ternary  $\text{Ge}_{20}\text{Se}_{70}\text{Zn}_{10}$  alloy by preparing its amorphous thin film. Lu

<sup>a</sup>Physics Department, Banaras Hindu University, Varanasi-221005, India. E-mail: dr\_neeraj\_mehta@yahoo.co.in

<sup>b</sup>Physics Department, IIT Roorkee, Roorkee-247667, India

<sup>c</sup>Physics Department, Al-Balqa Applied University, Al-Salt-19117, Jordan


*et al.*<sup>31</sup> designed a novel  $(\text{As}_2\text{S}_3)_{100-x}(\text{ZnSe})_x$  ( $0 \leq x \leq 30$ ) system for the possible applications in Mid-IR fiber lasers and found that these materials show compatibility with a drawing scheme adopted for preparing fibers showing low-loss of signals. Azhniuk and his research group<sup>32</sup> observed the sign of photoinduced creation of ZnSe nano-crystallites during the investigation of the optical characterization of  $\text{As}_2\text{S}_3$  thin films doped with zinc.

Nagpal *et al.*<sup>33</sup> investigated the consequences of adding zinc to the various optical parameter of binary  $\text{Se}_{90}\text{Sb}_{10}$  glass ( $0 \leq x \leq 10$ ) and concluded that the optical properties changed noticeably. Chazot and his research team<sup>34</sup> tried to design a fiber laser by using a composite that consists of active crystals of Fe:ZnSe in a glass matrix. For this, they checked the suspension stability of zinc selenide crystals in a specific glass made of  $\text{As}_2\text{S}_3$  and  $\text{As}_2\text{Se}_3$  having compositions 94.6 and 5.4 respectively. Some glass ceramics of the ternary Se-Ge-Zn system were developed by Velea *et al.*<sup>35</sup> for applications in infrared optics that were found capable of the prospective research work in IR optics.

In this way, different groups used zinc as a modifier in various multicomponent ChGs (especially binary and ternary systems) to see its consequence on the physical properties of parent samples. The above literature survey reveals that only a few endeavors were done to study the role of Zn on different physical properties in quaternary systems of ChGs. Keeping in mind this point and the fascinating results reported by the above research groups, we decided to synthesize some ChGs of the quaternary SeTeSnZn system. In this paper, we have reported a systematic analysis of the outcomes derived from the calorimetric experiments done on the present samples of the quaternary STSZ system.

## 2. Experimental

For the present studies, we prepared some glassy samples of SeTeSnZn by quenching their melts in chilled water. The specifics of this synthesis technique are described in ref. 36–38. The non-crystalline character of as-prepared samples was confirmed by the X-ray diffraction (XRD) technique and analysis of scanning electron microscopic (SEM) pictures. Fig. 1 shows such results for as-prepared samples. There were no sharp peaks of sufficient intensity in the XRD patterns of the present samples. Only a few weak peaks of insignificant intensities were observed over the broad humps *i.e.*, the first sharp diffraction peak (FSDP) and second sharp diffraction peak (SSDP) [see Fig. 1(a)] that were the characteristic of intermediate-range ordering in the present samples. However, we have identified some distinguished peaks that can be indexed.

The primary peaks of crystalline t-Se *i.e.*, trigonal Se (COD-9008579) are visible at  $2\theta$  values of  $23.5^\circ$ ,  $29.7^\circ$ , and  $51.7^\circ$ , which correspond to the crystal planes (100), (101), and (201), respectively while the small peaks at  $23.05^\circ$ ,  $27.55^\circ$ ,  $45.88^\circ$ ,  $47.04^\circ$ , and  $49.69^\circ$  are, respectively, associated with the crystal planes (100), (101), (003), (200), and (201) of crystalline t-Te *i.e.*, trigonal Te. The diffraction peaks at  $2\theta$  of  $32^\circ$ , are compared with the JCPDS data and respectively indexed to the

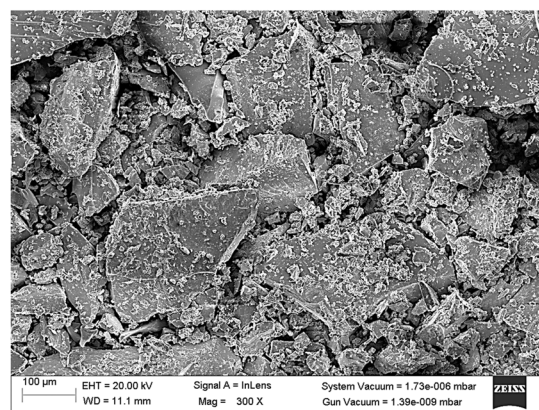
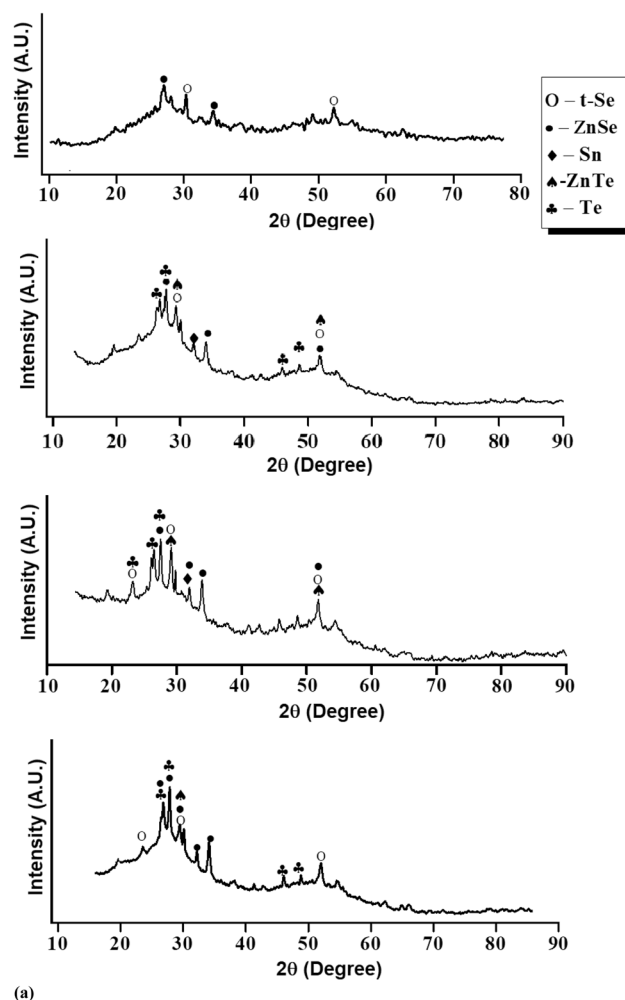


Fig. 1 (a) XRD patterns of the as-prepared samples indicate the glassy nature with evidence of some crystallites embedded in the glass matrix of the samples. (b) SEM picture of a representative sample ( $x = 2$ ) for the confirmation of the glassy nature.

crystallographic planes (101) polycrystalline planes metallic tin (JCPDS card no. 004-0673). Further, the peaks at ( $2\theta$ )  $27.65^\circ$ ,  $45.13^\circ$ , and  $52.6^\circ$  in the ZnSe spectrum correspond to the index planes (111), (220), and (311) for cubic zinc mix structured ZnSe (JCPDS card no. 65-9602). In addition to these peaks, the ZnSe

diffraction pattern also shows peaks at  $(2\theta)$  23.55, 26.4, 28.6, 30.5, 34.9, and 43.8, which correspond to the Miller indices (111), (100), (002), (101), (102) and (220). ZnTe nanocrystals' diffraction peaks at 29.92, 49.52, and 51.88 could be correlated with the (200), (311), and (222) planes of the cubic-phase ZnTe.

Further, no sign of crystallinity was observed in the SEM picture of the as-prepared samples. Such an SEM picture of a representative sample of  $\text{Se}_{76}\text{Te}_{20}\text{Sn}_2\text{Zn}_2$  is shown here [see Fig. 1(b)]. For the thermal characterization of the samples, we adopted non-isothermal heating and performed the calorimetric measurements in a differential scanning calorimeter (TA Instruments, USA; Model: Q20). DSC runs were performed at four diverse heating rates for all the samples. For doing calorimetric measurement in the DSC unit, the samples were crushed and 5 mg powder of each sample was enclosed in the crimped pans. For the calibration of the DSC cell, indium was used. Nitrogen gas is used as purge gas during the calorimetric measurements. The sensitivity of the employed DSC unit was 1.0  $\mu\text{W}$  and temperature accuracy was  $\pm 0.1^\circ\text{C}$ . The full details of this technique are described in ref. 39.

### 3. Results and discussion

Fig. 2 shows the results of the calorimetric experiment performed on the samples of the STSZ glassy system at a specific heating rate of  $10\text{ K min}^{-1}$ . The appearance of distinct glass transition and crystallization phenomena are seen as the first endothermic peak and the second exothermic peak in each scan of this figure. Further, the homogeneity of each as-prepared sample is reflected in the sharpness of the glass transition and crystallization peaks. When the homogeneity of the samples is poor, there is a strong chance of phase separation,<sup>40–43</sup> which appears as the splitting of the glass

transition peak and/or crystallization peak in double or triple components or the broadening of its width.

Abdel-Wahab<sup>43</sup> prepared a ternary glass ( $\text{Se}_{77.5}\text{Te}_{20}\text{Sn}_{2.5}$ ) whose composition was almost identical to the parent glass ( $\text{Se}_{78}\text{Te}_{20}\text{Sn}_2$ ) of the present SeTeSnZn system. He performed DSC experiments in  $\text{Se}_{77.5}\text{Te}_{20}\text{Sn}_{2.5}$  and found that this sample had two exothermic crystallization peaks as a signature of phase separation. However, such splitting of the glass transition peak and/or crystallization peak is neither observed in the parent  $\text{Se}_{78}\text{Te}_{20}\text{Sn}_2$  ternary glass nor zinc containing SeTeSnZn quaternary glasses. Thus, in our samples, the chance of this unwanted situation is ruled out as a signature of the sharpness of the glass transition peaks. Similar observations were observed in DSC scans obtained by calorimetric measurements at other heating rates. From such DSC scans of all samples at different heating rates, we have determined the glass transition temperature  $T_g$  and the crystallization temperature  $T_c$  by noting down the temperatures corresponding to peak maximum.

To realize the dependency of these characteristic temperatures on heating rate  $\beta$ , we have used the following empirical relations suggested by Lasocka:<sup>44</sup>

$$T_g = A_g + B_g \log \beta \quad (1)$$

Here  $A_g$  and  $B_g$  are Lasocka's parameters for glass transition.

The intercept  $A_g$  in eqn (1) represents the theoretical value of  $T_g$  corresponding to  $1\text{ K min}^{-1}$  heating rate. Lasocka's parameter  $B_g$  represents the slope of the straight line shown by eqn (1). Such linear graphs are shown in Fig. 3. Table 1 tabulates the value of  $A_g$  and  $B_g$  for the present samples.

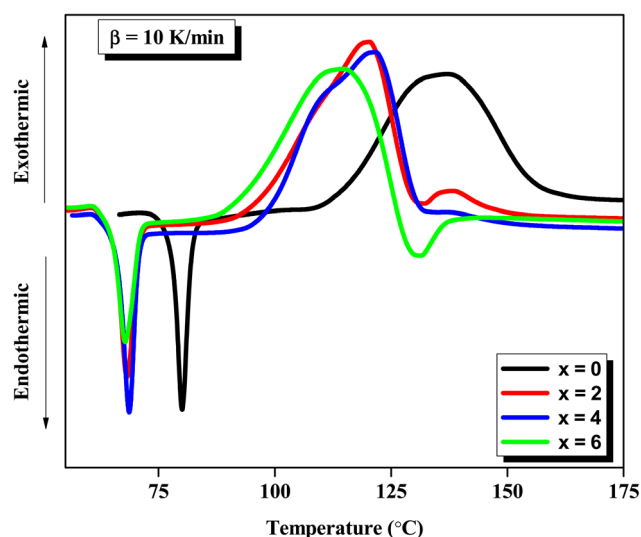


Fig. 2 DSC scans of samples ( $x = 0, 2, 4, 6$ ) of glassy system  $\text{Se}_{78-x}\text{Te}_{20}\text{Sn}_2\text{Zn}_x$  at a particular heating rate ( $10^\circ\text{C min}^{-1}$ ) represent sharp endothermic peaks as a sign of thermal relaxation in the glass transition region.

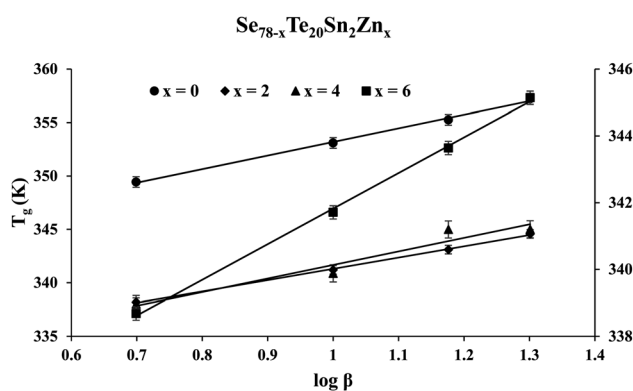


Fig. 3 Plots of  $T_g$  versus  $\log \beta$  for the present samples. The secondary y-axis is plotted for  $x = 6$ .

Table 1 Values of Lasocka's parameters of glass transition (i.e.,  $A_g$ ,  $B_g$ ) and crystallization (i.e.,  $A_c$ ,  $B_c$ ) for present samples

Sample	$A_g$ (K)	$B_g$ ( $\text{min}^{-1}$ )	$A_c$ (K)	$B_c$ ( $\text{min}^{-1}$ )
$\text{Se}_{78}\text{Te}_{20}\text{Sn}_2$	340.5	12.7	380.7	29.1
$\text{Se}_{76}\text{Te}_{20}\text{Sn}_2\text{Zn}_2$	330.7	10.6	372.5	20.6
$\text{Se}_{74}\text{Te}_{20}\text{Sn}_2\text{Zn}_4$	329	12.7	364.7	22.2
$\text{Se}_{72}\text{Te}_{20}\text{Sn}_2\text{Zn}_6$	331.1	10.7	370.9	23.3

The similar empirical relation of  $T_c$  for crystallization phenomenon can be expressed in terms of  $\beta$ :<sup>44</sup>

$$T_c = A_c + B_c \log \beta \quad (2)$$

Here  $A_c$ , and  $B_c$  are Lasocka's parameters for crystallization. The linear graphs corresponding to eqn (2) are shown in Fig. 4 for the present glasses. Table 1 tabulates the value of  $A_c$  and  $B_c$  for the present samples.

To find the correlation between  $T_g$  and  $T_c$  in the present study, we have derived the following relation by using eqn (1) and (2):

$$T_g = \left( \frac{B_c A_g - A_c B_g}{B_c} \right) + \left( \frac{B_g}{B_c} \right) T_c \quad (3)$$

Eqn (3) reveals that the kinetic temperatures  $T_c$  and  $T_g$ , corresponding to the crystallization and glass transition phenomena respectively, are mutually dependent on each other. Another interesting fact about such correlation is that we can use this equation for the determination of Kauzmann temperature ( $T_k$ ) which is a significant parameter for the characterization of glassy materials from a thermodynamic lookout.<sup>45</sup> The  $T_k$  represents the value of temperature upon which the entropy of a liquid having glass-forming ability becomes equal to that of its crystalline complement. Thus, keeping in mind the thermodynamic aspects, it signifies a lower limit for  $T_g$  below which the presence of an under-cooled glass-forming liquid does not obey thermodynamic law.<sup>45</sup> We can use eqn (3) for the determination of the Kauzmann temperature in the present case by keeping in mind that the Kauzmann temperature is the bottommost hypothetical borderline for the glass transition corresponding to a permissible assumption  $T_g = T_c = T_k$ <sup>46</sup> at a particular heating rate  $\beta_k$  (known as Kauzmann heating rate).

Solving eqn (3) for  $T_k$  (which is equal to  $T_g$  or  $T_c$ ) and  $\beta_k$ , we obtain the following relations:

$$T_k = \left( \frac{B_c A_g - A_c B_g}{B_c} \right) \quad (4)$$

and

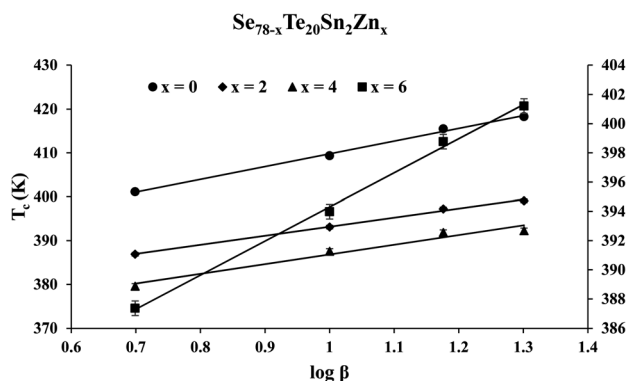


Fig. 4 Plots of  $T_c$  versus  $\log \beta$  for the present samples. The secondary y-axis is plotted for  $x = 6$ .

Table 2 Values of Kauzmann's parameter (*i.e.*,  $T_k$ ,  $\beta_k$ ) for present samples

Sample	$T_k$ (K)	$\beta_k$ (K min <sup>-1</sup> )
Se <sub>78</sub> Te <sub>20</sub> Sn <sub>2</sub>	309.1	$2.12 \times 10^{-6}$
Se <sub>76</sub> Te <sub>20</sub> Sn <sub>2</sub> Zn <sub>2</sub>	286.7	$2.7 \times 10^{-10}$
Se <sub>74</sub> Te <sub>20</sub> Sn <sub>2</sub> Zn <sub>4</sub>	281.2	$2.2 \times 10^{-9}$
Se <sub>72</sub> Te <sub>20</sub> Sn <sub>2</sub> Zn <sub>6</sub>	297.9	$5.8 \times 10^{-8}$

$$\beta_k = 10^{\left[ \frac{A_g - A_c}{B_c - B_g} \right]} \quad (5)$$

Table 2 tabulates the values obtained for  $T_k$  and  $\beta_k$  from eqn (4) and (5) in the present case. From this table, one can realize that it is enormously hard to perform the calorimetric experiments to detect this ideal glass transition because of the very low values of the heating rate.

The composition dependence of different kinetic parameters indicates that there is non-linear variation in different parameters. ChGs are famous for their non-linear physical properties due to their inherent structural flexibility. In ChGs, earlier it was believed that only two critical compositions are having the values 2.4 and 2.67 of mean coordination number ( $\bar{Z}$ ) in the IV–VI chalcogenide glassy system at which such a reversal in the compositional trend of different physical properties is observed. However, we have observed such a reversal in the compositional trend of Lasocka's parameters and Kauzmann's parameters at lower values of  $\bar{Z}$ . In the recent study,<sup>47–49</sup> it has been proposed that the rigidity percolation transition (*i.e.*, a reversal in the compositional trend) can occur over a range of mean coordination numbers leading to three distinct phases namely; floppy, intermediate and rigid. Various researchers<sup>50–52</sup> reported that such transitions occur at different compositions for different physical properties even at lower values of  $\bar{Z}$ . This is probably the reason behind the observation of the reversal in the compositional trend of Lasocka's parameters and Kauzmann's parameters at different compositions.

Moynihan and coworkers<sup>53</sup> developed a method to evaluate the glass transition activation energy. This relation is generally known as the Moynihan relation and can be expressed as:

$$\ln(\beta) = \left( -\frac{E_g}{RT_g} \right) + \text{constant} \quad (6)$$

Although the Kissinger equation<sup>54</sup> is primarily derived for the evaluation of kinetic activation energy  $E_c$  involved in thermally governed crystallization, one can adopt an analogous relation for the glass transition to estimate the corresponding activation energy  $E_g$ . The premise of employing this relation for the assessment of  $E_g$  originates from the shifting of maxima at  $T_g$  towards the higher temperature side with the rise in  $\beta$  values. Consequently, the analogous Kissinger equation takes the form:<sup>54</sup>

$$\ln \left( \frac{\beta}{T_g^2} \right) = -\frac{E_g}{RT_g^2} + \text{constant} \quad (7)$$





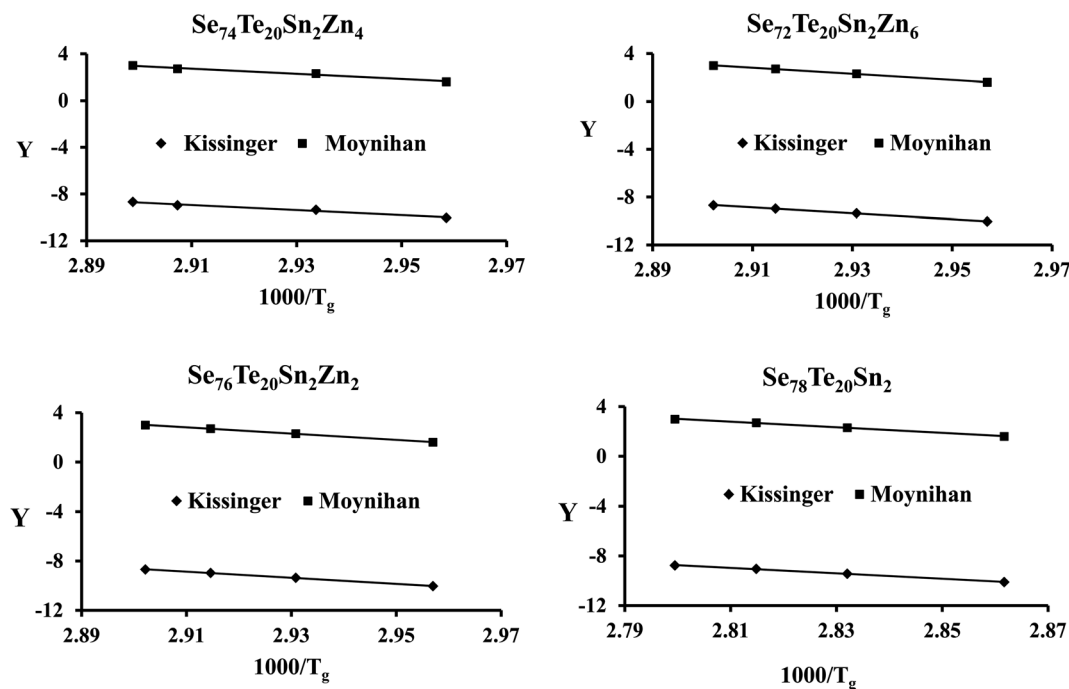


Fig. 5 Plots of  $\ln(\beta)$  vs.  $10^3/T_g$  and  $\ln(\beta/T_g^2)$  vs.  $10^3/T_g$  for the present samples. On the y-axis,  $Y = \ln(\beta)$  and  $\ln(\beta/T_g^2)$  for the Moynihan method and Kissinger method, respectively.

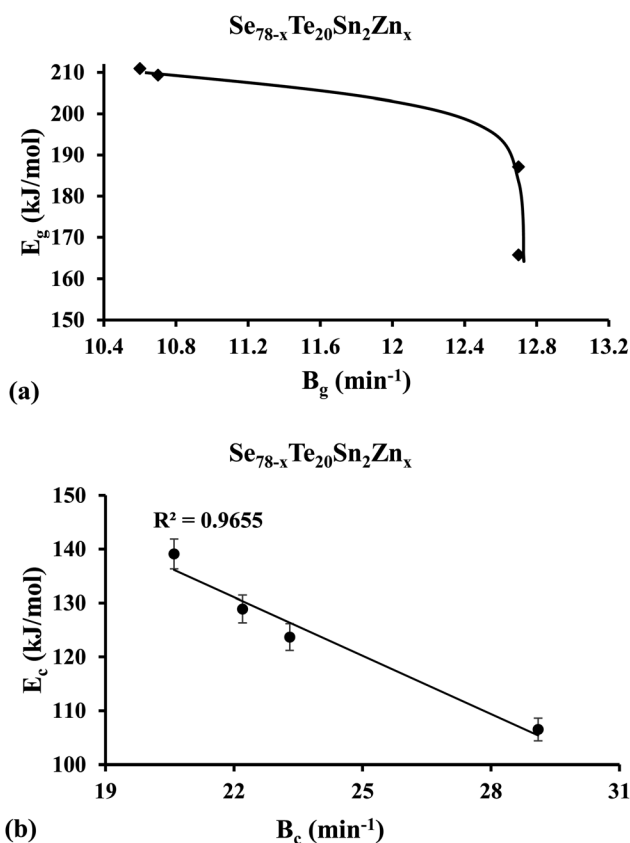


Fig. 6 Plots of (a) glass transition activation energy  $E_g$  against Lasocka parameter  $B_g$ , and (b) crystallization activation energy  $E_c$  against Lasocka parameter  $B_g$ .

The plots of  $\ln(\beta)$  and  $\ln(\beta/T_g^2)$  against  $1000/T_g$  are shown in Fig. 5. The linear nature of the graphs confirms the applicability of eqn (6) and (7). From the gradients of such linear graphs, we have determined  $E_g$  for all the samples.

Lasocka<sup>44</sup> observed that  $B_g$  is subject to the cooling rate of the melt unswervingly. He suggested that parameter  $B_g$  signifies the retroactivity of the configurational alterations arising in the vicinity of the temperature range corresponding to the glass transformation. Keeping in mind the fact that glass transition activation energy  $E_g$  is related to the movements of structural species throughout the thermal relaxation in the neighborhood of the glass transition, we have plotted the graph of  $E_g$  against parameter  $B_g$  and found that  $E_g$  of the samples is decreased non-linearly with increasing parameter  $B_g$  [see Fig. 6(a)] in the present case. This indicates that the higher the value of  $B_g$ , the more transient responsiveness of the local configurational deviations and so the glass network requires less activation energy. A similar correlation between glass transition activation energy  $E_g$  and Lasocka parameter  $B_g$  was observed earlier in the glassy  $\text{Se}_{80-x}\text{Te}_{20}\text{Sn}_x$  system by Kumar *et al.*<sup>55</sup> Thus, our present results of the glassy  $\text{Se}_{80-x}\text{Te}_{20}\text{Sn}_2\text{Zn}_x$  system are consistent with our previous results of the glassy  $\text{Se}_{80-x}\text{Te}_{20}\text{Sn}_x$  system.<sup>55</sup>

By using the JMA model, various methods are developed by different authors to calculate  $E_c$  for different kinds of glasses. As derived by Kissinger,<sup>54</sup> we can express  $E_c$  in terms of  $\beta$  as follows:

$$\ln\left(\frac{\beta}{T_c^2}\right) = -\frac{E_c}{RT_c^2} + \text{constant} \quad (8)$$

Matusita and Sakka<sup>56,57</sup> set up the following equation for the determination of  $E_c$  from the classical JMA relation:



$$\ln(\beta) = -\frac{E_c}{RT_c} + \text{constant} \quad (9)$$

Augis and Bennett<sup>58</sup> derived a slightly different non-isothermal relation to calculate the crystallization activation energy:

$$\ln\left(\frac{\beta}{T_c}\right) = -\frac{E_c}{RT_c} + \ln K_0 \quad (10)$$

By using eqn (8)–(10) we have plotted the graphs of  $\ln(\beta/T_c^2)$  [*i.e.*, Kissinger method],  $\ln(\beta)$  [*i.e.*, Matusita–Sakka method] and  $\ln(\beta/T_c)$  [*i.e.*, Augis–Bennett method] against  $1000/T_c$ . Such plots are shown in Fig. 7. By using the slopes of these graphs, we have evaluated the  $E_c$  values in the present study for the various samples. Keeping in mind the observed correlation between Lasocka parameter  $B_g$  and glass transition activation energy  $E_g$ , we have plotted a similar plot for the Lasocka parameter  $B_c$  and crystallization activation energy  $E_c$  and observed that  $E_c$  is also decreased with a rise in Lasocka parameter  $B_c$  [see Fig. 6(b)]. However, the decrement in  $E_c$  with  $B_c$  is almost linear.

The crystallization rate of ChGs can be determined by studying its temperature dependence which is governed by the Arrhenius equation:<sup>59,60</sup>

$$K = K_0 \exp\left(-\frac{E_c}{RT}\right) \quad (11)$$

Since the rate  $K$  represents the growth rate of the crystalline species during the devitrification of the samples, we can understand the influence of zinc incorporation on the rate constant  $K$  of crystallization for the parent glass by substituting

the value of the crystallization activation energy  $E_c$  in eqn (11) at the temperatures lying in the crystallization region of the SeTeSnZn samples.

Knowing the value of  $E_c$  and pre-factor  $K_0$ , we have determined the rate of crystallization for the present samples of the SeTeSnZn system using eqn (11). The composition dependence of  $\ln K$  is shown in Fig. 8 for all the samples at different heating rates. From this figure, we see that the crystallization rate of the quaternary alloys is larger in comparison to parent SeTeSn glass and it attains an almost constant higher value after zinc incorporation at all the heating rates.

We have used the following relation for the determination of the average heat of atomization  $\bar{H}_A$  of the samples in the present study:<sup>61</sup>

$$\bar{H}_A = \frac{(jH_{\text{Se}} + kH_{\text{Te}} + lH_{\text{Sn}} + mH_{\text{m}})}{(j + k + l + m)} \quad (12)$$

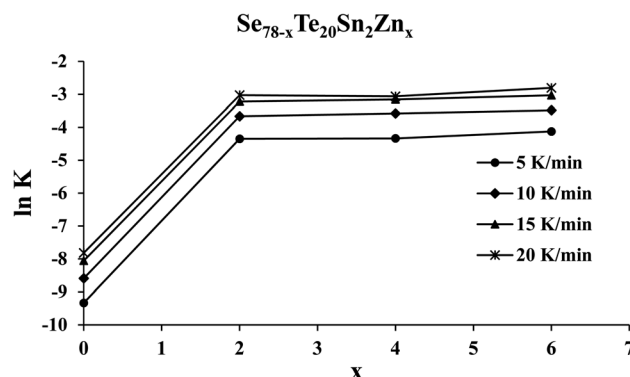


Fig. 8 Composition dependence of the rate of crystallization  $K$ .

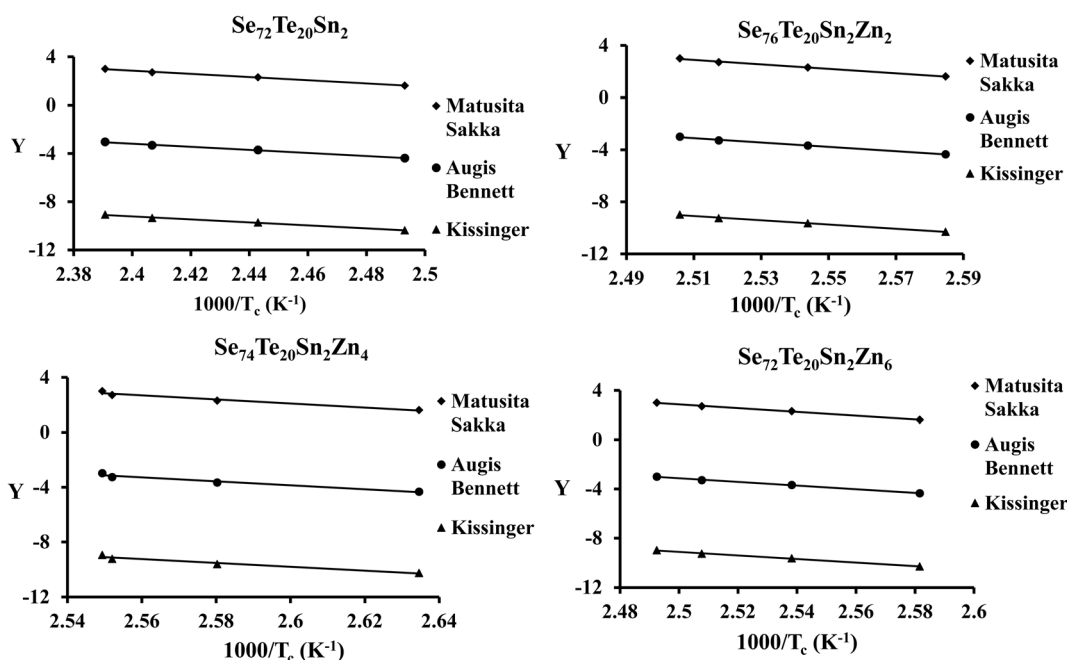


Fig. 7 Plots of  $\ln(\beta/T_c^2)$ ,  $\ln(\beta)$ , and  $\ln(\beta/T_c)$  versus  $1000/T_c$  for the present samples. On the y-axis,  $Y = \ln(\beta/T_c^2)$ ,  $\ln(\beta)$ , and  $\ln(\beta/T_c)$  for the Kissinger method, Matusita–Sakka method, and Augis–Bennett method respectively.



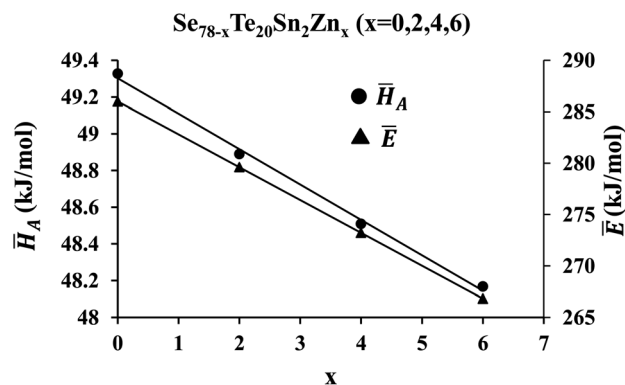


Fig. 9 Composition dependence of average heat of atomization  $\bar{H}_A$  and overall mean bond energy  $\bar{E}$ .

Here  $H_{Se}$ ,  $H_{Te}$ ,  $H_{Sn}$ , and  $H_{Zn}$  are the respective atomization heat values of Se, Te, Sn, and Zn in the present samples ( $0 \leq x \leq 6$ ) of the  $Se_{78-x}Zn_xTe_{20}Sn_2$  system, while  $j$ ,  $k$ ,  $l$ , and  $m$  are the corresponding concentrations. The dependency of  $\bar{H}_A$  on the zinc content is revealed in Fig. 9.

To determine the overall mean bond energy of the samples, we have used a model recommended by Tichy and Ticha based on the approach chemical bond theory.<sup>62</sup> According to the Tichy–Ticha model, the ChGs can be categorized into three categories based on a parameter  $R$  according to the following conditions:

$$R = 1; R > 1 \text{ or } R < 1 \quad (13)$$

For  $R$  is equal to one (*i.e.*,  $R = 1$ ), Tichy and Ticha suggested that the glass network attains the stoichiometric composition because of the presence of heteropolar bonds solitary and this composition is corresponding to the chemical threshold where we observe frequently the utmost values of glass transition temperature  $T_g$ . If  $R$  is greater than one (*i.e.*,  $R > 1$ ), then the corresponding glassy system will be a member of the chalcogen-rich family. In such a case, the glassy system consists of both chalcogen–chalcogen bonds and heteropolar bonds. If  $R$  is smaller than one (*i.e.*,  $R < 1$ ), then the corresponding glassy system will be a member of the chalcogen-poor family. In such cases, the chalcogen–chalcogen bonds are almost absent and the glassy system consists of only metal–metal bonds and heteropolar bonds.<sup>62</sup> Since in the present studies, the concentration of chalcogen elements Se and Te is very high as compared to the other two elements (Sn and Zn), our glassy system belongs to the chalcogen-rich family.

For the chalcogen-rich region, the Tichy–Ticha model further suggests that the degree of cross-linking can be determined by using the expression:

$$P_r = \frac{kN_{Te} + lN_{Sn} + mN_{Zn}}{(k + l + m)} \quad (14)$$

In eqn (14),  $N_{Zn}$ ,  $N_{Sn}$ , and  $N_{Te}$  are the respective coordination numbers of Zn, Sn, and Te. The average bond energy ( $E_c$ ) corresponding to the mean cross-linking per atom is given by:<sup>62</sup>

$$E_c = P_r \times E_{hb} \quad (15)$$

Here,  $E_{hb}$  is the mean hetero-polar bond energy that can be expressed as:<sup>62</sup>

$$E_{hb} = \frac{kN_{Te}(E_{Se-Te}) + lN_{Sn}(E_{Se-Sn}) + mN_{Zn}(E_{Se-Zn})}{(kN_{Te} + lN_{Sn} + mN_{Zn})} \quad (16)$$

In eqn (16),  $E_{Se-Te}$ ,  $E_{Se-Sn}$ , and  $E_{Se-Zn}$  represent the hetero-polar bond energies of Se atoms with Te, Sn, and Zn atoms respectively. For the ‘remaining glass matrix’, we now express the mean bond energy per atom  $E_{rm}$  as follows:<sup>62</sup>

$$E_{rm} = \frac{(E_{Se-Se}) \times (\bar{r} - 2P_r)}{\bar{r}} \quad (17)$$

In eqn (17)  $E_{Se-Se}$  represents the strength of the homopolar Se–Se bond in the chalcogen-rich region.

Knowing the values of  $E_c$  and  $E_{rm}$ , we can determine the overall value  $\bar{E}$  of mean bond energy by adding them, *i.e.*,<sup>62</sup>

$$\bar{E} = E_c + E_{rm} \quad (18)$$

Fig. 9 shows the dependency of  $\bar{E}$  on the concentration of zinc also. This figure reveals that both average heat of atomization  $\bar{H}_A$  and overall mean bond energy  $\bar{E}$  are decreased with the intensification of the zinc atoms in the parent SeTeSn glass. This is probably understood through the chemically ordered network model and chemical bond approach<sup>63</sup> as described in the next paragraph.

The bonds of Se–Se ( $330.5 \text{ kJ mol}^{-1}$ ) are sturdier as compared to the bonds of Te–Te ( $293.3 \text{ kJ mol}^{-1}$ ).<sup>64</sup> Similarly, the Se atom has higher electro-negativity as compared to the Te atom. Thus, the bond strength of Se–Zn bonds is greater than Te–Zn bonds. Consequently, we can expect that zinc as a foreign element gives preference to Se after entering the glass network of parent glass and so there is a drop in the number of Se–Se bonds ( $330.5 \text{ kJ mol}^{-1}$ ) as an outcome of the dissolution of Zn atoms in polymeric Se-chains.<sup>64</sup> Thus, the creation of Se–Zn bond ( $170.7 \text{ kJ mol}^{-1}$ ) and Zn–Zn bonds ( $22 \text{ kJ mol}^{-1}$ ) occur.<sup>64</sup> Accordingly, the cohesive energy of the system decreases with a rise in the Zn concentration in the parent glass. This is probably the reason for the reduction in the overall value of mean bond energy and average heat of atomization of the preset samples with the rise in Zn concentration.

A crucial parameter to determine the permanency of glassy substances in the vitreous state is the released amount of the enthalpy ( $\Delta H_{gc}$ ) through the phase transformation from the non-crystalline state (*i.e.*, glassy phase) to the crystalline state (*i.e.*, crystal phase). It can be determined by subtracting the glass transition enthalpy ( $\Delta H_g$ ) from crystallization enthalpy ( $\Delta H_c$ ), so that:

$$\Delta H_{gc} = \Delta H_c - \Delta H_g \quad (19)$$

The glass having lower values of  $\Delta H_{gc}$  possesses higher thermal stability. To confirm this, we have determined thermal stability by using the Saad–Poulin relation:<sup>65</sup>



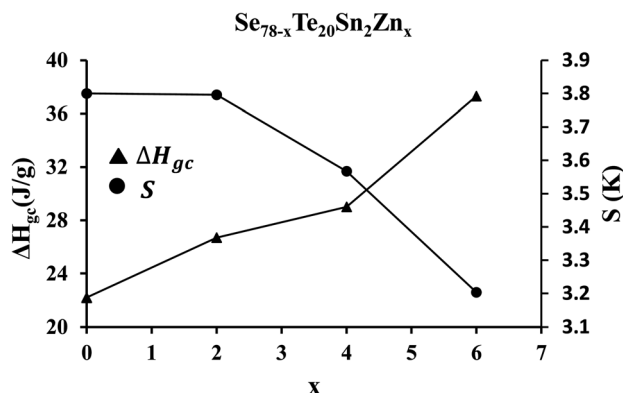


Fig. 10 Composition dependence of enthalpy ( $\Delta H_{gc}$ ) released during phase transformation from the glassy state to the crystalline state and thermal stability parameter  $S$ .

$$S = \frac{(T_c - T_o)(T_c - T_g)}{T_g} \quad (20)$$

Here  $T_o$  represent the on-set temperature of the crystallization peaks for the present samples. Knowing the values of stability parameter  $S$ , we have plotted the graphs of  $S$  and  $\Delta H_{gc}$  against the composition of the zinc (see Fig. 10). The figure reveals that the thermal stability parameter  $S$  increases with decreasing  $\Delta H_{gc}$ . A similar correlation was observed by Naqvi and Saxena in some samples of the glassy  $\text{Se}_{80-x}\text{Te}_{20}\text{Zn}_x$  system.<sup>66</sup>

In the last two decades, various research groups have studied the thermal stability of zinc containing multicomponent (*i.e.*, binary, ternary, and quaternary) ChGs of Se–Te to optimize the durability and reduce aging effects.<sup>22,66–68</sup> Hence, we have compared the results of the present study with that are reported

in ref. 22 and 66–68. The comparison of such data is tabulated in Table 3. This table reveals the better durability of the present samples as compared to others zinc containing multicomponent ChGs due to their thermal stability values.

## 4. Conclusions

The incorporation of zinc in SeTeSn helps in the synthesis of new third-generation samples of quaternary SeTeSnZn system that are free from the problem of phase separation. This is confirmed by the appearance of the sharp solo peaks at glass transition temperature in zinc-containing derived samples. Further analysis reveals that the activation energies corresponding to the crystallization and glass transition phenomena are decreased with a rise in the corresponding Lasocka parameters.

The crystallization rate is increased appreciably in the quaternary alloys after zinc incorporation in the parent SeTeSn ternary sample. The average heat of atomization and overall mean bond energy of the present samples is decreased with the reduction in cohesive energy when we increase the concentration of zinc. The thermal stability parameter is found to be increased with the drop in enthalpy released during the phase transformation from glass to the crystalline state.

## Conflicts of interest

There are no conflicts to declare.

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Table 3 Comparisons of thermal stability for zinc containing ChGs in the present study and with the available data in literature

S. no.	Glass composition	$S$ (K)	Reference
1	$\text{Se}_{98}\text{Zn}_2$	1.6	22
	$\text{Se}_{95}\text{Zn}_5$	2.7	
	$\text{Se}_{90}\text{Zn}_{10}$	2.5	
	$\text{Se}_{80}\text{Zn}_{20}$	2.0	
2	$\text{Se}_{70}\text{Te}_{28}\text{Zn}_2$	3.1	67
	$\text{Se}_{70}\text{Te}_{26}\text{Zn}_4$	3.6	
	$\text{Se}_{70}\text{Te}_{24}\text{Zn}_6$	2.9	
	$\text{Se}_{70}\text{Te}_{22}\text{Zn}_8$	3.3	
3	$\text{Se}_{78}\text{Te}_{20}\text{Zn}_2$	3.1	66
	$\text{Se}_{76}\text{Te}_{20}\text{Zn}_4$	3.2	
	$\text{Se}_{74}\text{Te}_{20}\text{Zn}_6$	2.9	
	$\text{Se}_{72}\text{Te}_{20}\text{Zn}_8$	2.4	
	$\text{Se}_{70}\text{Te}_{20}\text{Zn}_{10}$	1.9	
4	$\text{Se}_{93}\text{Zn}_2\text{Te}_5$	1.8	68
	$\text{Se}_{91}\text{Zn}_2\text{Te}_5\text{In}_2$	2.8	
	$\text{Se}_{89}\text{Zn}_2\text{Te}_5\text{In}_4$	3.9	
	$\text{Se}_{87}\text{Zn}_2\text{Te}_5\text{In}_6$	4.2	
	$\text{Se}_{83}\text{Zn}_2\text{Te}_5\text{In}_{10}$	3.0	
5	$\text{Se}_{78}\text{Te}_{20}\text{Sn}_2$	3.8	Present work
	$\text{Se}_{78}\text{Te}_{20}\text{Sn}_2\text{Zn}_2$	3.8	
	$\text{Se}_{76}\text{Te}_{20}\text{Sn}_2\text{Zn}_4$	3.7	
	$\text{Se}_{74}\text{Te}_{20}\text{Sn}_2\text{Zn}_6$	3.2	





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