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Thermodynamic Properties of Layered Tetradymite-like Compounds of the Ag–Ge–Sb–Te System

Mykola Moroz,  Fiseha Tesfaye, Pavlo Demchenko, Myroslava Prokhorenko, Daniel Lindberg, Oleksandr Reshetnyak, Leena Hupa

Abstract

The phase equilibria of the Ag–Ge–Sb–Te system in the part Ag_8GeTe_6 –Ge–GeTe– Sb_2Te_3 were investigated by the electromotive force (EMF) method. The determined phase relations were used to express the chemical reactions. The potential-forming reactions were performed by applying electrochemical cells (–) C | Ag | Ag_2GeS_3 -glass | D | C (+), where C is graphite, Ag_2GeS_3 -glass is the fast purely Ag^+ ions conducting electrolyte, and D is an equilibrium mixture of phases. According to the experimental data on the EMF versus temperature of each electrochemical cells, analytical equations for the Gibbs energies of $\text{GeSb}_8\text{Te}_{13}$, $\text{GeSb}_6\text{Te}_{10}$, GeSb_4Te_7 , GeSb_2Te_4 , $\text{Ge}_2\text{Sb}_2\text{Te}_5$, $\text{Ge}_3\text{Sb}_2\text{Te}_6$, and $\text{Ge}_4\text{Sb}_2\text{Te}_7$ compounds were obtained. The thermodynamic properties of silver-saturated tetradymite-like compounds have been calculated for the first time. A good correspondence between experimental values and structure data reported in literature has been established.

Keywords Thermoelectric materials, Layered compounds, Phase equilibria, Thermodynamic properties, EMF method

1. Introduction

In recent decades, the study of thermoelectric materials (TM) has attracted increasing attention both from an energy and environmental point of view. Thermoelectric materials provide a large variety of applications in heating, cooling, and power generation due to direct conversion of heat into electrical energy [1–3]. The efficiency of TM is defined by the figure-of-merit ZT parameter, which is $ZT = S^2\sigma T/(k_{el}+k_{lat})$, where S , σ , k_{el} and k_{lat} represent the Seebeck coefficient, electrical, electronic, and lattice vibrations conductivity, respectively [4]. Many studies in the past decades are focused on increasing of ZT values in several ways by studying the multicomponent materials. The ternary compounds with mixed layered tetradymite-like structure of the homologous series $n\text{GeTe}\cdot m\text{Sb}_2\text{Te}_3$ ($n, m = 1–4$) in the GeTe– Sb_2Te_3 system were identified as good TM [5–10].

The T – x phase diagram of the GeTe– Sb_2Te_3 system has been studied by Abrikosov et al. [11]. The formation of the following compounds was established GeSb_4Te_7 , GeSb_2Te_4 , and $\text{Ge}_2\text{Sb}_2\text{Te}_5$. These ternary compounds melt incongruently at 878, 888, and 903 K, respectively. Skums et al. [12] and Skoroponov et al. [13] reported that the GeSb_4Te_7 and GeSb_2Te_4 compounds melt congruently at 880 and 888 K, respectively. Later on, four additional compounds $\text{Ge}_4\text{Sb}_2\text{Te}_7$, $\text{Ge}_3\text{Sb}_2\text{Te}_6$, $\text{GeSb}_6\text{Te}_{10}$, and $\text{GeSb}_8\text{Te}_{13}$ of the homologous series

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$n\text{GeTe}\cdot m\text{Sb}_2\text{Te}_3$ were found [5–9]. These compounds were identified by high-resolution electron microscopy, electronic diffraction [5, 6, 14, 15] and X-ray diffraction [8, 14] techniques. In the layered ternary compounds Te atoms form a cubic close packing structure with octahedral voids, occupied by Ge and Sb atoms. The hexagonal unit cells are formed by multilayer packets of different types, ordered in alternating directions in the direction perpendicular to hexagonal c axis. The layered compounds can be divided in two groups, based on the compositional variations of “ GeSb_2Te_4 ”. The structures of the GeTe-rich compounds are formed by identical slabs. The structures of Sb_2Te_3 -rich compounds include five- and seven-layer slabs [8].

For the practical application of compounds in electronic cooling devices and transformation of thermal energy into electrical power, absence of structural transformations in the operating temperature range is important. Shelimova et al. [9] measured the Hall coefficient, electrical resistivity, and thermoelectric power of compounds of the homologous series in the range of 100–600 K. According to Skums et al. [12], some of these compounds undergoes a phase transition at $T > 600$ K. Below $T = 600$ K, the energy of thermal vibrations of atoms may not be sufficient to change the lattice crystal structure. Under these conditions, the phase will be in a metastable state. The examples of transition of compounds from the metastable to the thermodynamic equilibrium state. were published in several works [16–21]. Such transition was carried out by including compound as one of the components into a positive electrode of the electrochemical cells (ECCs). A key role in achieving the thermodynamic equilibrium state of the compounds belongs to Ag^+ ions migrated from the Ag-reference electrode to multi-phase mixture of a positive electrode [22].

Moroz and Prokhorenko [16] reported that continuous solid solution of the $\text{PbSe}\text{--}\text{PbTe}$ system at $T < 570$ K is a metastable. Decomposition of the solid solution and formation in the right electrode of ECC the intermediate phase in the $\text{PbSe}_{0.70}\text{Te}_{0.30}\text{--}\text{Pb}_{0.73}\text{Te}_{0.27}$ concentration range was established.

Formation of the Ag_2GeSe_3 compound in the $\text{Ag}_2\text{Se}\text{--}\text{GeSe}_2$ system was found under carefully controlled conditions in ECC [17]. Furthermore, Ag_2GeSe_3 was observed to undergo a phase transition at $T = 535$ K.

The phase diagram of the Bi–Te system in the entire composition range is given in [19]. It is characterized by the presence of Bi_2Te_3 , Bi_4Te_5 , Bi_8Te_9 , BiTe , Bi_6Te_5 , Bi_4Te_3 , Bi_2Te , and $\text{Bi}_{14}\text{Te}_6$ compounds. Data on the temperature and concentration ranges of stability of the Bi_2Te_3 , BiTe , Bi_2Te , and $\text{Bi}_{14}\text{Te}_6$ compounds are given in Abrikosov et al. [20]. It was found that BiTe and Bi_2Te are phases of variable composition. The phase equilibria of the Ag–Bi–Te system and thermodynamic properties of the compounds have been investigated by the electromotive force (EMF) method in the temperature range of 490–550 K [18]. It is characterized by the presence of the $\text{Ag}_2\text{Te}\text{--}\text{Bi}$, $\text{Ag}_2\text{Te}\text{--}\text{Bi}_{14}\text{Te}_6$, $\text{Ag}_2\text{Te}\text{--}\text{Bi}_2\text{Te}$, $\text{Ag}_2\text{Te}\text{--}\text{BiTe}$, and $\text{Ag}_2\text{Te}\text{--}\text{Bi}_2\text{Te}_3$ quasi-binary sections. A good agreement between calculated [18] and literature [23] thermodynamic data for BiTe and Bi_2Te_3 compounds confirm the phase equilibria of the Ag–Bi–Te system presented in [18]. It is mean that in thermodynamic equilibrium BiTe and Bi_2Te are stoichiometric compounds due to the decomposition of their solid solutions under potential-forming process.

According to [21], the solid solubility of PbS in Ag_8GeS_6 does not exceed 13 mol% . Metastable state of this solid solution was established by the EMF method in [24]. It was found that maximum solubility occurs up to 19 mol% PbS . Furthermore, existence of intermediate phase with the approximated composition $\text{Ag}_{6.62}\text{Pb}_{0.16}\text{Ge}_{0.84}\text{S}_{5.20}$ was predicted.

Herein we present the thermodynamic properties of the saturated solid solutions of the $n\text{GeTe}\cdot m\text{Sb}_2\text{Te}_3$ compounds in the phase region $\text{Ag}_8\text{GeTe}_6\text{--}\text{Ge}\text{--}\text{Ge}_4\text{Sb}_2\text{Te}_7\text{--}\text{Sb}_2\text{Te}_3$ of the Ag–Ge–Sb–Te system. These thermochemical data of the compounds can be used to optimize or complete phase diagrams in the investigated system and for selection of thermodynamically stable materials with optimal values of the ZT parameter.

2. Experimental Section

The starting materials for synthesis were high-purity elements: Ag, 99.99 wt% (Alfa-Aesar, Germany); Ge, 99.999 wt% (Alfa-Aesar, Germany); Sb, 99.99 wt% (Alfa-Aesar, Germany); S, 99.999 wt% (Alfa-Aesar, Germany); Te, 99.999 wt% (Alfa-Aesar, Germany).

For the EMF measurements [24–29], the following electrochemical cell was used:



where C is graphite, $\text{Ag}_2\text{GeS}_3\text{-glass}$ is the fast purely Ag^+ ions conducting electrolyte [30, 31], and D is an four-element equilibrium mixture of phases. The vertical lines in ECC indicate phase boundaries or contacts between cell components. The cell polarities and half-cell reactions in the ECCs were established according to rules described in [24, 32].

The positive (right) electrodes D of the EECs were synthesized by melting of chemical elements in thin-walled evacuated quartz glass ampoules at $T = 1070$ K for 5 h. Slowly cooled to room temperature samples were ground into a fine powder with the particle size of ≤ 5 μm , evacuated, and then annealing at $T = 550$ K for 250 h. The composition of positive electrodes of ECCs were calculated based on equations of electrochemical reactions for each of 7 four-phase regions of the $\text{Ag}_8\text{GeTe}_6\text{--}\text{Ge}\text{--}\text{Ge}_4\text{Sb}_2\text{Te}_7\text{--}\text{Sb}_2\text{Te}_3$ system. The $\text{Ag}_2\text{GeS}_3\text{-glass}$ [30, 33] was obtained by melt quenching of the corresponding elements from $T = 1200$ K in ice water. The phase compositions of the right electrode compounds were characterized by differential thermal analysis (Paulik-

Paulik-Erdey derivatograph fitted with chromel-alumel thermocouples and an H307-1 XY recorder) and by X-ray powder diffraction (STOE STADI P diffractometer, transmission mode, $\text{CuK}\alpha_1$ radiation, a bent Ge (111) monochromator on primary beam, $2\theta/\omega$ scan mode) techniques. X-ray phase analysis was performed using STOE WinXPOW [34] and PowderCell [35] program packages using crystallographic data on the structures of known phases taken from databases [36, 37].

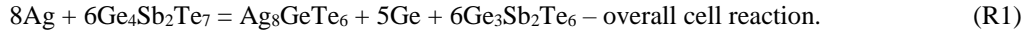
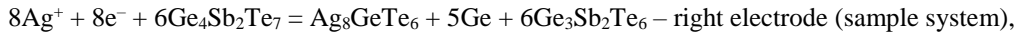
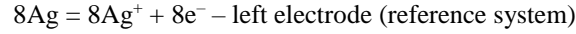
Components of the ECCs in powder form were pressed at 10^8 Pa through a 2 mm diameter hole arranged in the fluoroplast matrix up to density $\rho = (0.93 \pm 0.02) \cdot \rho_0$, where ρ_0 is the experimentally determined density of cast samples. Five-fold thermal cycling of ECCs in the temperature range of 450–510 K was performed to eliminate possible defects due to plastic deformation during sample pressing. The heating and cooling rates were of 2 K min^{-1} . Experiments were performed in a horizontal resistance furnace, similar to that described in [38]. As protective atmosphere, we used a continuously flowing highly purified (0.9999 volume fraction) Ar(g) at $P = 1.2 \cdot 10^5$ Pa, with a flow rate of $2 \cdot 10^{-3} \text{ m}^3 \text{ h}^{-1}$ from the right to left electrode of the ECCs. The temperature was maintained with an accuracy of ± 0.5 K. The EMF of the cells were measured by high-resistance (input impedance of $>10^{12} \Omega$) universal U7-9 digital voltmeter. The equilibrium in ECCs at each temperature was achieved within 2 h. After equilibrium has been attained, the EMF values were constant or their variation did not exceed ± 0.2 mV.

In our previous works [39, 40] we have described in details the scheme of ECCs and procedure of the EMF measurements.

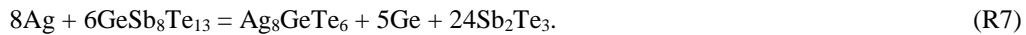
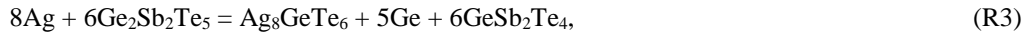
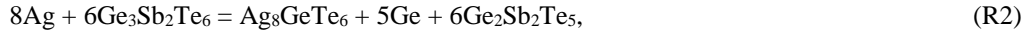
3. Results and Discussion

As can be seen in Table 1 and Fig. 1, concentration space of the Ag–Ge–Sb–Te system in the Ag_8GeTe_6 –Ge– $\text{Ge}_4\text{Sb}_2\text{Te}_7$ – Sb_2Te_3 part consists of 7 four-phase regions. In Fig. 1, the phase regions borders are marked by two-phase equilibrium lines. The lines of two-phase equilibria were determined in [9, 41, 42] as well as by our investigation by the EMF method. The spatial location of the established phase regions relative to composition of Ag were used to determine the thermodynamic properties of ternary compounds.

In the phase region (1), listed in Table 1, the electrochemical process of the formation of Ag_8GeTe_6 , Ge, and $\text{Ge}_3\text{Sb}_2\text{Te}_6$ from the pure Ag and $\text{Ge}_4\text{Sb}_2\text{Te}_7$ compound follows:



For the phase regions (2)–(7), listed in Table 1, the overall cell reactions can be expressed as:



Reactions (R1)–(R7) were realized to take place in the ECCs at different temperatures. Based on equations (R1)–(R7), the positive electrodes D of ECCs were prepared from pure elements Ag, Ge, Sb, and Te taken in molar ratios: 1) 8 : 3 : 10 : 22, 2) 8 : 4 : 14 : 29, 3) 8 : 4 : 10 : 23, 4) 8 : 4 : 6 : 17, 5) 8 : 5 : 4 : 15, 6) 8 : 7 : 4 : 17, and 7) 8 : 9 : 4 : 19, respectively.

Table 1 Four-phase regions of the Ag–Ge–Sb–Te system in the Ag_8GeTe_6 –Ge– $\text{Ge}_4\text{Sb}_2\text{Te}_7$ – Sb_2Te_3 part and the EMF values of ECCs at $T=480$ K in corresponding phase areas

No	Phase region	E / mV
1	$\text{Ge}_4\text{Sb}_2\text{Te}_7$ –Ge– Ag_8GeTe_6 – $\text{Ge}_3\text{Sb}_2\text{Te}_6$	275.09
2	$\text{Ge}_3\text{Sb}_2\text{Te}_6$ –Ge– Ag_8GeTe_6 – $\text{Ge}_2\text{Sb}_2\text{Te}_5$	272.44
3	$\text{Ge}_2\text{Sb}_2\text{Te}_5$ –Ge– Ag_8GeTe_6 – GeSb_2Te_4	270.17
4	GeSb_2Te_4 –Ge– Ag_8GeTe_6 – GeSb_4Te_7	265.98
5	GeSb_4Te_7 –Ge– Ag_8GeTe_6 – $\text{GeSb}_6\text{Te}_{10}$	258.38
6	$\text{GeSb}_6\text{Te}_{10}$ –Ge– Ag_8GeTe_6 – $\text{GeSb}_8\text{Te}_{13}$	254.52
7	$\text{GeSb}_8\text{Te}_{13}$ –Ge– Ag_8GeTe_6 – Sb_2Te_3	247.50

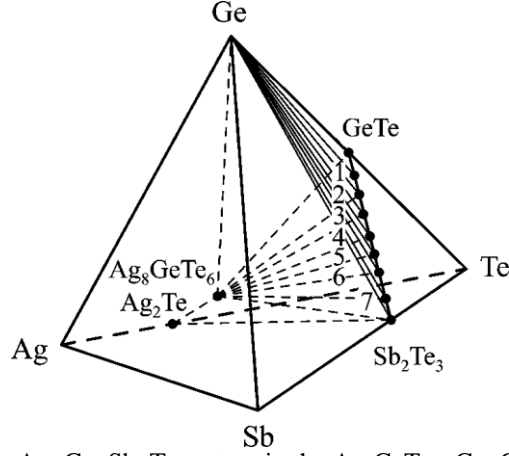


Fig. 1 The phase equilibria of the Ag–Ge–Sb–Te system in the Ag_8GeTe_6 –Ge–GeTe– Sb_2Te_3 part, below $T = 520$ K. 1 is $\text{Ge}_4\text{Sb}_2\text{Te}_7$, 2 is $\text{Ge}_3\text{Sb}_2\text{Te}_6$, 3 is $\text{Ge}_2\text{Sb}_2\text{Te}_5$, 4 is GeSb_2Te_4 , 5 is GeSb_4Te_7 , 6 is $\text{GeSb}_6\text{Te}_{10}$, 7 is $\text{GeSb}_8\text{Te}_{13}$

The relationship of EMF versus temperature of reactions (R1)–(R7) in ECCs were approximated by Eqs. (1)–(7) and are shown in Fig. 2. The crystal lattices of tetrademite-like compounds are highly disordered, due to the presence of mixed occupancy of cations in octahedral voids [43]. The bends of the curves at different temperatures in Fig. 2 arises from individual activation changes of occupancy sites by Ge, Sb, and Ag cations that are characteristic of each of the studied phases.

$$E_{(1)}/\text{mV} = (162.98 \pm 0.21) + (233.56 \pm 0.43) \cdot 10^{-3} T/\text{K} \quad 462 \leq T/\text{K} \leq 503, \quad (1)$$

$$E_{(2)}/\text{mV} = (171.25 \pm 0.21) + (210.82 \pm 0.44) \cdot 10^{-3} T/\text{K} \quad 458 \leq T/\text{K} \leq 495, \quad (2)$$

$$E_{(3)}/\text{mV} = (185.46 \pm 0.16) + (176.48 \pm 0.34) \cdot 10^{-3} T/\text{K} \quad 457 \leq T/\text{K} \leq 499, \quad (3)$$

$$E_{(4)}/\text{mV} = (139.20 \pm 0.17) + (264.13 \pm 0.35) \cdot 10^{-3} T/\text{K} \quad 460 \leq T/\text{K} \leq 499, \quad (4)$$

$$E_{(5)}/\text{mV} = (139.39 \pm 0.18) + (247.89 \pm 0.38) \cdot 10^{-3} T/\text{K} \quad 457 \leq T/\text{K} \leq 496, \quad (5)$$

$$E_{(6)}/\text{mV} = (147.71 \pm 0.22) + (222.53 \pm 0.46) \cdot 10^{-3} T/\text{K} \quad 465 \leq T/\text{K} \leq 501, \quad (6)$$

$$E_{(7)}/\text{mV} = (145.75 \pm 0.19) + (211.97 \pm 0.39) \cdot 10^{-3} T/\text{K} \quad 465 \leq T/\text{K} \leq 497. \quad (7)$$

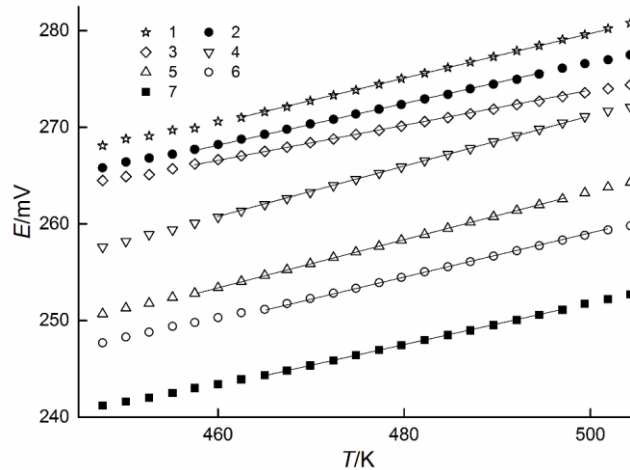


Fig. 2 The temperature dependences of EMF of the cells with the positive electrodes D of the phase regions: 1 is $\text{Ge}_4\text{Sb}_2\text{Te}_7$ –Ge– Ag_8GeTe_6 – $\text{Ge}_3\text{Sb}_2\text{Te}_6$, 2 is $\text{Ge}_3\text{Sb}_2\text{Te}_6$ –Ge– Ag_8GeTe_6 – $\text{Ge}_2\text{Sb}_2\text{Te}_5$, 3 is $\text{Ge}_2\text{Sb}_2\text{Te}_5$ –Ge– Ag_8GeTe_6 – GeSb_2Te_4 , 4 is GeSb_2Te_4 –Ge– Ag_8GeTe_6 – GeSb_4Te_7 , 5 is GeSb_4Te_7 –Ge– Ag_8GeTe_6 – $\text{GeSb}_6\text{Te}_{10}$, 6 is $\text{GeSb}_6\text{Te}_{10}$ –Ge– Ag_8GeTe_6 – $\text{GeSb}_8\text{Te}_{13}$, 7 is $\text{GeSb}_8\text{Te}_{13}$ –Ge– Ag_8GeTe_6 – Sb_2Te_3

In addition, Eqs. (1)–(7) confirm the correctness of the spatial location of the established phase regions. In particular, EMF values of ECCs remain constant independent of the general composition in the phase region, but change drastically on their boundaries [24]. Furthermore, as can be seen in Table 1, the phase region further away from the composition of Ag is characterized by higher EMF value at $T = \text{const}$.

The Gibbs energies, entropies, and enthalpies of the reactions (R1)–(R7) can be calculated by combining the measured EMF values of each ECC and the thermodynamic Eqs. (8)–(10) [49,59,60]:

$$\Delta_r G = -n \cdot F \cdot E, \quad (8)$$

$$\Delta_r H = -n \cdot F \cdot [E - (dE/dT)T], \quad (9)$$

$$\Delta_r S = n \cdot F \cdot (dE/dT), \quad (10)$$

where $n = 8$ is the number of electrons involved in the reactions (R1)–(R7), $F = 96485.33289 \text{ C} \cdot \text{mol}^{-1}$ is Faraday constant, and E is the EMF of the electrochemical cells.

By assuming $\left(\frac{\partial \Delta_r H}{\partial T}\right)_p = 0$ and $\left(\frac{\partial \Delta_r S}{\partial T}\right)_p = 0$, which implies that $\Delta_r C_p = 0$ [32], the thermodynamic function of reactions (R1)–(R7) were calculated through extrapolation of the linear temperature dependences of the EMF of ECCs to $T = 298 \text{ K}$ and using Eqs. (8)–(10). The results of the calculations are listed in Table 2.

Table 2 Standard thermodynamic values of reactions (R1)–(R7) in ECCs at $T = 298 \text{ K}$

Reaction	$-\Delta_r G^\circ$	$-\Delta_r H^\circ$	$\Delta_r S^\circ$
	$\text{kJ} \cdot \text{mol}^{-1}$		$\text{J} \cdot \text{mol}^{-1} \cdot \text{K}^{-1}$
(R1)	179.53 ± 0.62	125.80 ± 0.16	180.28 ± 0.33
(R2)	180.68 ± 0.66	132.18 ± 0.16	162.73 ± 0.34
(R3)	183.75 ± 0.57	143.15 ± 0.12	136.22 ± 0.26
(R4)	168.20 ± 0.48	107.45 ± 0.13	203.88 ± 0.27
(R5)	164.61 ± 0.52	107.59 ± 0.14	191.34 ± 0.29
(R6)	165.20 ± 0.64	114.01 ± 0.17	171.77 ± 0.36
(R7)	161.26 ± 0.56	112.50 ± 0.15	163.62 ± 0.30

Standard Gibbs energy and entropy of reactions (R1)–(R7) are related to the Gibbs energy of formation and entropy of compounds and pure elements the following equations:

$$\Delta_{r(1)} G^\circ = \Delta_f G_{\text{Ag}_8\text{GeTe}_6}^\circ + 6\Delta_f G_{\text{Ge}_3\text{Sb}_2\text{Te}_6}^\circ - 6\Delta_f G_{\text{Ge}_4\text{Sb}_2\text{Te}_7}^\circ, \quad (11)$$

$$\Delta_{r(1)} S^\circ = S_{\text{Ag}_8\text{GeTe}_6}^\circ + 5S_{\text{Ge}}^\circ + 6S_{\text{Ge}_3\text{Sb}_2\text{Te}_6}^\circ - 8S_{\text{Ag}}^\circ - 6S_{\text{Ge}_4\text{Sb}_2\text{Te}_7}^\circ, \quad (12)$$

$$\Delta_{r(2)} G^\circ = \Delta_f G_{\text{Ag}_8\text{GeTe}_6}^\circ + 6\Delta_f G_{\text{Ge}_2\text{Sb}_2\text{Te}_5}^\circ - 6\Delta_f G_{\text{Ge}_3\text{Sb}_2\text{Te}_6}^\circ, \quad (13)$$

$$\Delta_{r(2)} S^\circ = S_{\text{Ag}_8\text{GeTe}_6}^\circ + 5S_{\text{Ge}}^\circ + 6S_{\text{Ge}_2\text{Sb}_2\text{Te}_5}^\circ - 8S_{\text{Ag}}^\circ - 6S_{\text{Ge}_3\text{Sb}_2\text{Te}_6}^\circ, \quad (14)$$

$$\Delta_{r(3)} G^\circ = \Delta_f G_{\text{Ag}_8\text{GeTe}_6}^\circ + 6\Delta_f G_{\text{GeSb}_2\text{Te}_4}^\circ - 6\Delta_f G_{\text{Ge}_2\text{Sb}_2\text{Te}_5}^\circ, \quad (15)$$

$$\Delta_{r(3)} S^\circ = S_{\text{Ag}_8\text{GeTe}_6}^\circ + 5S_{\text{Ge}}^\circ + 6S_{\text{GeSb}_2\text{Te}_4}^\circ - 8S_{\text{Ag}}^\circ - 6S_{\text{Ge}_2\text{Sb}_2\text{Te}_5}^\circ, \quad (16)$$

$$\Delta_{r(4)} G^\circ = \Delta_f G_{\text{Ag}_8\text{GeTe}_6}^\circ + 6\Delta_f G_{\text{GeSb}_4\text{Te}_7}^\circ - 12\Delta_f G_{\text{GeSb}_2\text{Te}_4}^\circ, \quad (17)$$

$$\Delta_{r(4)} S^\circ = S_{\text{Ag}_8\text{GeTe}_6}^\circ + 5S_{\text{Ge}}^\circ + 6S_{\text{GeSb}_4\text{Te}_7}^\circ - 8S_{\text{Ag}}^\circ - 12S_{\text{GeSb}_2\text{Te}_4}^\circ, \quad (18)$$

$$\Delta_{r(5)} G^\circ = \Delta_f G_{\text{Ag}_8\text{GeTe}_6}^\circ + 12\Delta_f G_{\text{GeSb}_6\text{Te}_{10}}^\circ - 18\Delta_f G_{\text{GeSb}_4\text{Te}_7}^\circ, \quad (19)$$

$$\Delta_{r(5)} S^\circ = S_{\text{Ag}_8\text{GeTe}_6}^\circ + 5S_{\text{Ge}}^\circ + 12S_{\text{GeSb}_6\text{Te}_{10}}^\circ - 8S_{\text{Ag}}^\circ - 18S_{\text{GeSb}_4\text{Te}_7}^\circ, \quad (20)$$

$$\Delta_{r(6)} G^\circ = \Delta_f G_{\text{Ag}_8\text{GeTe}_6}^\circ + 18\Delta_f G_{\text{GeSb}_8\text{Te}_{13}}^\circ - 24\Delta_f G_{\text{GeSb}_6\text{Te}_{10}}^\circ, \quad (21)$$

$$\Delta_{r(6)} S^\circ = S_{\text{Ag}_8\text{GeTe}_6}^\circ + 5S_{\text{Ge}}^\circ + 18S_{\text{GeSb}_8\text{Te}_{13}}^\circ - 8S_{\text{Ag}}^\circ - 24S_{\text{GeSb}_6\text{Te}_{10}}^\circ, \quad (22)$$

$$\Delta_{r(7)} G^\circ = \Delta_f G_{\text{Ag}_8\text{GeTe}_6}^\circ + 24\Delta_f G_{\text{Sb}_2\text{Te}_3}^\circ - 6\Delta_f G_{\text{GeSb}_8\text{Te}_{13}}^\circ, \quad (23)$$

$$\Delta_{r(7)} S^\circ = S_{\text{Ag}_8\text{GeTe}_6}^\circ + 5S_{\text{Ge}}^\circ + 24S_{\text{Sb}_2\text{Te}_3}^\circ - 8S_{\text{Ag}}^\circ - 6S_{\text{GeSb}_8\text{Te}_{13}}^\circ. \quad (24)$$

The entropy of formations of the $\text{GeSb}_8\text{Te}_{13}$ compound can be calculated as:

$$\Delta_f S_{\text{GeSb}_8\text{Te}_{13}}^\circ = S_{\text{GeSb}_8\text{Te}_{13}}^\circ - S_{\text{Ge}}^\circ - 8S_{\text{Sb}}^\circ - 13S_{\text{Te}}^\circ. \quad (25)$$

For $\text{Ge}_3\text{Sb}_2\text{Te}_6$, $\text{Ge}_4\text{Sb}_2\text{Te}_7$, $\text{Ge}_2\text{Sb}_2\text{Te}_5$, GeSb_2Te_4 , GeSb_4Te_7 , and $\text{GeSb}_6\text{Te}_{10}$ compounds the corresponding reactions to determine $\Delta_f S^\circ$ can be written similar to Eq. (25) with their respective moles.

By combining Eqs. (8)–(25) and using data of the pure elements [23], Ag_8GeTe_6 , and Sb_2Te_3 [23, 24], the standard Gibbs energy of formations of the layered tetradyomite-like compounds of the homologous series $n\text{GeTe} \cdot m\text{Sb}_2\text{Te}_3$ were calculated to be:

$$\Delta_f G_{\text{GeSb}_8\text{Te}_{13}}^\circ / (\text{kJ} \cdot \text{mol}^{-1}) = -(248.06 \pm 2.17) - (3.90 \pm 0.03) \cdot 10^{-3} T / \text{K}, \quad (26)$$

$$\Delta_f G_{\text{GeSb}_6\text{Te}_{10}}^\circ / (\text{kJ}\cdot\text{mol}^{-1}) = -(191.51 \pm 1.56) - (1.74 \pm 0.02) \cdot 10^{-3} T / \text{K}, \quad (27)$$

$$\Delta_f G_{\text{GeSb}_4\text{Te}_7}^\circ / (\text{kJ}\cdot\text{mol}^{-1}) = -(135.32 \pm 1.01) + (0.75 \pm 0.01) \cdot 10^{-3} T / \text{K}, \quad (28)$$

$$\Delta_f G_{\text{GeSb}_2\text{Te}_4}^\circ / (\text{kJ}\cdot\text{mol}^{-1}) = -(79.14 \pm 0.53) + (3.54 \pm 0.02) \cdot 10^{-3} T / \text{K}, \quad (29)$$

$$\Delta_f G_{\text{Ge}_2\text{Sb}_2\text{Te}_5}^\circ / (\text{kJ}\cdot\text{mol}^{-1}) = -(96.16 \pm 1.21) + (6.50 \pm 0.06) \cdot 10^{-3} T / \text{K}, \quad (30)$$

$$\Delta_f G_{\text{Ge}_3\text{Sb}_2\text{Te}_6}^\circ / (\text{kJ}\cdot\text{mol}^{-1}) = -(114.99 \pm 2.13) + (10.79 \pm 0.16) \cdot 10^{-3} T / \text{K}, \quad (31)$$

$$\Delta_f G_{\text{Ge}_4\text{Sb}_2\text{Te}_7}^\circ / (\text{kJ}\cdot\text{mol}^{-1}) = -(134.90 \pm 3.29) + (15.96 \pm 0.31) \cdot 10^{-3} T / \text{K}. \quad (32)$$

The thermodynamic values of the saturated solid solutions of compounds in the $\text{Ag}_8\text{GeTe}_6\text{-Ge-Ge}_4\text{Sb}_2\text{Te}_7\text{-Sb}_2\text{Te}_3$ part are given in Table 3 and Fig. 3.

Table 3 Summary of the standard thermodynamic properties of layered tetradymite-like compounds of the $\text{GeTe-Sb}_2\text{Te}_3$ system at $T=298$ K determined in this work

Phase	$-\Delta_f G^\circ$	$-\Delta_f H^\circ$	$T\Delta_f S^\circ$	S°	$\Delta_f S^\circ$
	$\text{kJ}\cdot\text{mol}^{-1}$			$\text{J}\cdot\text{mol}^{-1}\cdot\text{K}^{-1}$	
$\text{GeSb}_8\text{Te}_{13}$	251.96 ± 1.38	248.06 ± 2.17	3.90 ± 0.03	1051.80 ± 14.73	13.09 ± 0.09
$\text{GeSb}_6\text{Te}_{10}$	193.25 ± 1.28	191.51 ± 1.56	1.74 ± 0.02	805.02 ± 13.61	5.84 ± 0.04
GeSb_4Te_7	134.57 ± 1.18	135.32 ± 1.01	-0.75 ± 0.01	557.16 ± 6.48	-2.50 ± 0.02
GeSb_2Te_4	75.60 ± 1.09	79.14 ± 0.53	-3.54 ± 0.02	308.25 ± 3.35	-11.88 ± 0.06
$\text{Ge}_2\text{Sb}_2\text{Te}_5$	89.66 ± 1.21	96.16 ± 1.21	-6.50 ± 0.06	378.86 ± 3.34	-21.84 ± 0.21
$\text{Ge}_3\text{Sb}_2\text{Te}_6$	104.20 ± 1.38	114.99 ± 2.13	-10.79 ± 0.16	445.06 ± 5.91	-36.23 ± 0.53
$\text{Ge}_4\text{Sb}_2\text{Te}_7$	118.94 ± 1.57	134.90 ± 3.29	-15.96 ± 0.31	508.32 ± 8.99	-53.54 ± 1.03

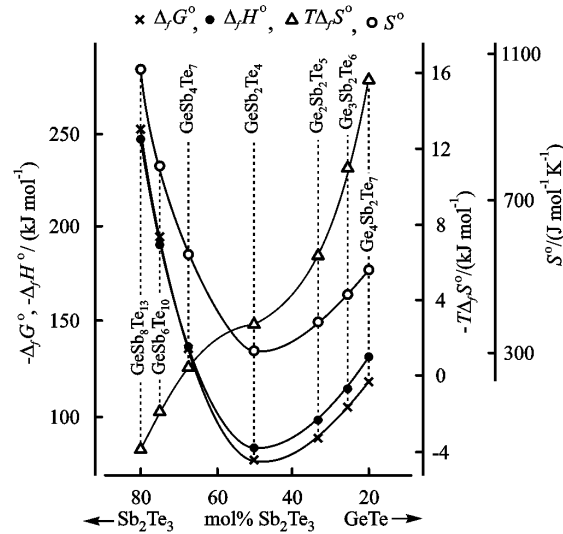


Fig. 3 Concentration changes of thermodynamic functions of layered tetradymite-like compounds of the $\text{GeTe-Sb}_2\text{Te}_3$ system

As can be seen in Fig. 3, at 50 mol% Sb_2Te_3 , a composition that corresponds to GeSb_2Te_4 compound, the $\Delta_f G^\circ$, $\Delta_f H^\circ$, S° values are minimum and the sign of the second derivative of $T\Delta_f S^\circ$ function changes. Gibbs energy, enthalpy, and entropy of the compounds is observed to increase with the addition of GeTe or Sb_2Te_3 to GeSb_2Te_4 composition. Results shown in Table 3 and Fig. 3 are in a good agreement with those data reported in [7–9] regarding the difference in the crystal structures of the compounds of the $\text{GeTe-Sb}_2\text{Te}_3$ system on either side of the GeSb_2Te_4 . The unit cell of the GeSb_2Te_4 compound contains 3 seven-layer slabs across the c axis. According to [7], in GeTe -rich compounds thermodynamic values increasing is accounted with more quantities of atomic layers in the packet and number of the same type packets that determine the unit cell parameter c . For the Sb_2Te_3 -rich compounds, increases are due to the number and alternation of 5- and 7-layer packets along the c axis.

4. Conclusions

1. The reproducibility of the values $E_{(1)-(7)}(T)$ of the ECCs during the heating and cooling cycles characterizes saturated solid solutions of the compounds of the GeTe–Sb₂Te₃ system as thermodynamically stable phases in the temperature range of 455–505 K.
2. The measured EMF values were applied to calculate the Gibbs energies, enthalpies, and entropies of silver-saturated solid solutions of the $n\text{GeTe}\cdot m\text{Sb}_2\text{Te}_3$ ($n, m = 1-4$) compounds.
3. Concentration dependences of the thermodynamic values of compounds confirm the previously established difference between the crystalline structures of GeTe-rich and Sb₂Te₃-rich phases of the GeTe–Sb₂Te₃ system.

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Conflict of Interest

The authors declare that they have no conflict of interest.

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