

CONDUCTION STUDIES ON AMORPHOUS InSbX_3 (X = Te OR Se) THIN FILMS

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Amorphous InSbX_3 (X = Te or Se) thin films are obtained by thermal evaporation technique of bulk material on to well cleaned glass substrates. The current –voltage characteristics have been measured in the temperature range (303-393 K) and thickness range (230-490 nm). The obtained I-V curves revealed two types of conduction. The first region is ohmic type in the lower field followed by non-ohmic type of conduction in the high field region. In the high-field region, the field lowering coefficient β is evaluated, and has been analyzed by the anomalous Poole-Frenkel effect. The temperature dependence of ohmic current is that of thermally activated process. The variation of dielectric constant with temperature for the two compounds has been studied.

1. Introduction

In the recent years, investigations of thin film systems of metal-dielectric-metal type led to the establishment of thin film microelectronics as a promising new branch of modern electronics. Thin film microelectronics deals with processes taking place in thin metal, dielectrics and semiconductor films as well as with the properties of the interface between two phases. The knowledge of some physical properties of semiconductor films is necessary. Chalcogenide films are new kind of materials with immense qualities of use in many practical applications. These materials exhibit unique infrared transmission and electrical properties, which makes them potentially useful for applications such as threshold and memory switching [1]. Since a high – speed switching and the memory effect of an amorphous semiconductor were reported [2,3], its properties have been studied very actively. There have been many discussions on the mechanism of high-speed switching and of electric conduction in amorphous semiconductors.

The study of the pre-switching current- voltage characteristics is necessary to identify the electrical conduction mechanism occurring in the off state of chalcogenide glass switches. One possible conduction mechanism is the space charge limited current [4], hopping conduction [5], small polaron conduction [6] and the Poole-Frenkel conduction [7].

The system $\text{In}_2\text{Se}_3 - \text{Sb}_2\text{Se}_3$ is a semiconducting compound with very interesting electrical and optical properties, which has not been sufficiently investigated [8, 9]. Few authors studied the optical [10] and electrical [11,12] properties of InSbSe_3 single crystal, however little attention is devoted to study the physical properties of InSbSe_3 thin films. $\text{In}_2\text{Te}_3\text{-Sb}_2\text{Te}_3$ system ($\text{Sb}_{2-x}\text{In}_x\text{Te}_3$ solid solution) belongs to the family of layered compounds having the structure of space group. Physical properties of $\text{Sb}_{2-x}\text{In}_x\text{Te}_3$ single crystal were described in a number of papers [13-18], however little attention is devoted to study the physical properties of InSbTe_3 thin films.

This paper aimed to investigate the voltage and temperature dependence of current in amorphous InSbSe_3 and InSbTe_3 films produced by vacuum deposition to identify the dominating conduction mechanism of charge transport.

2. Experimental techniques

InSbSe_3 and InSbTe_3 compositions were prepared in bulk form by melting together the stoichiometric amount of the constituent elements of each compositions of purity 99.999% using a specially designed oscillation furnace insure the homogeneity of the samples. The furnaces temperature was raised at a rate of 50 K h^{-1} to 1223 K for InSbSe_3 composition [19] and to 1003 K for InSbTe_3 composition [20]. At this temperature the melt was held for 2 days. Then the ampoules are cooled slowly at a rate of 1 K min^{-1} and 3 K min^{-1} for InSbSe_3 and InSbTe_3 compositions to room

temperature. Thin films of the obtained compositions were then prepared by thermal evaporation technique using a high vacuum coating unit (Edwards type E306A), onto the well cleaned glass substrates of suitable dimensions with the desired mask, and provided with Al electrodes in sandwich configuration for I-V measurements. Thin films with different thicknesses (230 - 490 nm) were deposited at constant rate under vacuum of 1.3 m Pa (10^{-5} torr). The substrate temperature was held at room temperature ~ 300 K. The film thickness was measured by Tolansky's interferometric method.

The X-ray diffractometer of the type Philips (PM 8203) was used to investigate the structure of the obtained samples in bulk and thin film forms. The chemical composition of the obtained samples in bulk and thin films was checked by energy dispersive x-ray analysis (EDX) in the scanning electron microscope (JEOL 5400).

The current- voltage (I-V) characteristics were measured through the temperature range 303-393 K using electrometer (Keithley type 616A) for the potential drop measurements and a microdigit multimeter (TE 924) for the current measurements. The temperature of the sample was monitored using a chromel – alumel thermocouple.

4. Results and discussion

X-ray diffraction pattern obtained for the two systems under investigations the bulk form has a polycrystalline nature while their thin film form has an amorphous nature [21].

EDX analysis indicated that the compositions of the prepared materials under test in powder and thin film forms are closed to the initial compositions with an experimental error $\pm 2\%$ [21].

4.1 I-V characteristics

Fig. 1 (a&b) shows the current –voltage (I-V) characteristics for InSbSe_3 and InSbTe_3 films at different temperatures in the temperature range (303-393K). It is observed that the curves show two different regions; first the linear part for lower voltage, indicating ohmic conduction due to the presence of thermally generated carriers (electrons and/or holes) followed by non-linear part at higher voltage. It was found that at higher temperature the linear regions, extend to higher values of the applied field. The ohmic region is believed to be controlled by electron hopping. Similar results have been obtained previously for other amorphous semiconductor materials [22-25]. The I-V characteristics exhibited an ohmic region at low applied voltage with low field followed by a transition to supra linear at higher field up to the breakdown voltage value.

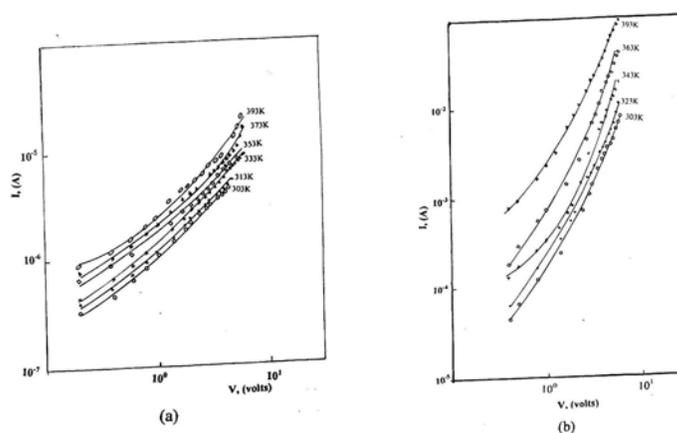


Fig. 1. (a) Current- voltage characteristics for InSbSe_3 film of thickness 250 nm at various temperatures. (b) Current - voltage characteristics for InSbTe_3 film of thickness 230 nm at various temperatures.

In order to analyze the basis of this behavior, the effect of sample thickness and electrode material such as gold and silver on the conductivity of the samples under investigation was

examined. Fig. 2 (a&b) shows the thickness dependence of current for thin film samples of systems under investigations, which reveals that the current is independent of film thickness at constant voltage through the studied range of thickness. Thus we can conclude that the current is independent of these factors, i.e the conductivity is independent of these factors. This suggests that space charge limited conduction and Schottky emission do not specify the conduction mechanism and consequently the field dependence was a property of the bulk material.

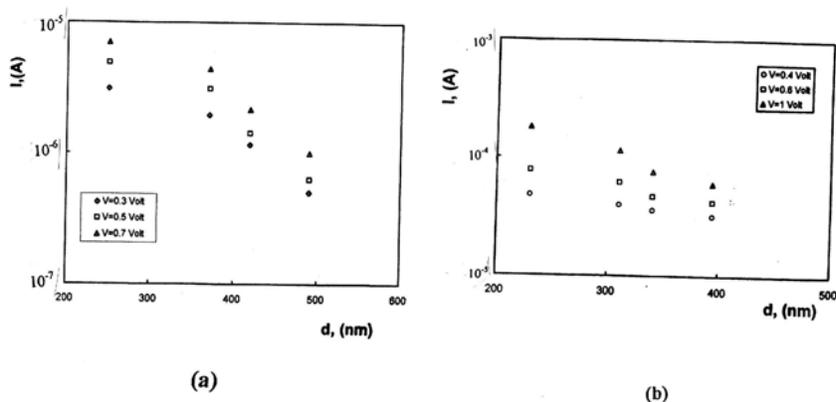


Fig. 2. (a) Thickness dependence of room temperature current for InSbSe_3 film of different voltage. (b) Thickness dependence of room temperature current for InSbTe_3 film of different voltage.

Fig. 3 (a&b) represents the plot of I (in logarithmic scale) against $E^{1/2}$ for the data of Fig. 1 (a&b) for two compositions. It is clear from this figure that there is a dependence of type $I \propto E^{1/2}$ for both InSbSe_3 and InSbTe_3 films over at least two order of magnitude of current. This field dependence of current indicates, that the conduction mechanism may be either the Schottky [26], or Poole-Frenkel [27] type. The following evidence allows us to eliminate Schottky conduction: (a) $\log IT^{-2}$ should vary linearly with T^{-1} at different fixed voltage. This is not found to be the case in any of the two compounds under test, (b) The activation energy for Schottky emission should generally be greater than 0.8 eV. In the present case it is lower than this value. These results together with other features mentioned above, clearly suggest that the conduction is a bulk effect. Moreover the different parameters estimated from graphs found to lie in an acceptable range only if the Poole-Frenkel effect is assumed to be operative.

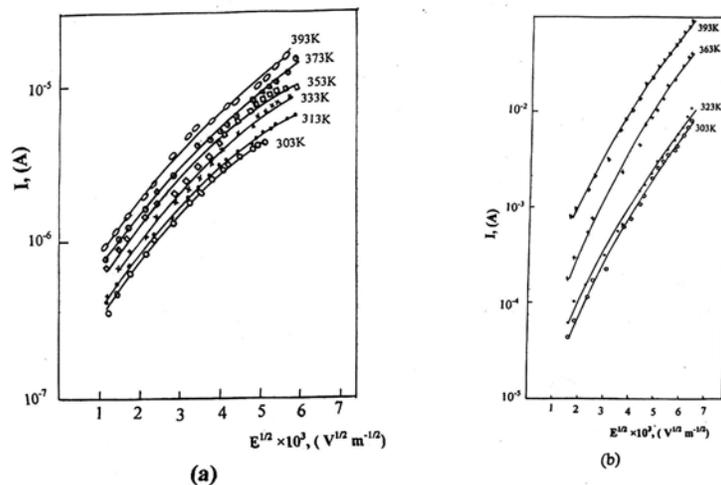


Fig. 3. (a) I (in logarithmic scale) vs $E^{1/2}$ plots for the InSbSe_3 film at different temperatures. (b) I (in logarithmic scale) vs $E^{1/2}$ plots for the InSbTe_3 film at different temperatures.

According to Poole-Frenkel effect, the electric field interacts with the Coulombic potential barrier of a donor center or trap and the height of barrier is lowered. In the presence of an electric field, the electrons are thermally emitted from the randomly distributed traps to the conduction band by the lowering of the Coulombic potential barrier by the external electric field. At higher fields in excess of 10^4 V/m for many dielectric films containing shallow traps exhibits current density (J)-Voltage characteristics is of the form is given by the following equation [28]:

$$J = J_0 \exp(\beta E^{1/2} / kT) \quad (1)$$

Where J_0 is the low field current density, E the applied electric field, k the Boltzmann constant, T the absolute temperature and β the Poole-Frenkel field lowering coefficient which is given by:

$$\beta = (e^3 / a\pi\epsilon\epsilon_0)^{1/2} \quad (2)$$

Where a equal to 1 for Poole-Frenkel emission and equal to 4 for Schottky emission [28], e is the electronic charge, ϵ is the dielectric constant of the material and ϵ_0 the permittivity of the free space.

Now, β_{PF} can be calculated theoretically according to equation (2) (where the dielectric constant ϵ has been determined) and calculated experimentally from the slope of the linear part in the non-ohmic region of Fig.3(a&b).

The experimental values of β_{PF} calculated from the slope of the graphs in Fig. 3 (a&b) for the two systems under investigation are listed in Tables (1&2). Values of the dielectric constant ϵ can be calculated according to equation (2) and are also given in Tables (1&2). The obtained value of β_{PF} at room temperature (303K) is about $2.77 \times 10^{-5} \text{ eVm}^{1/2}\text{V}^{-1/2}$ for InSbSe₃ films, but if using the estimated value of $\epsilon = 8.8$ [29], the theoretical values of β_{PF} calculated from equation (2) is $2.6 \times 10^{-5} \text{ eVm}^{1/2}\text{V}^{-1/2}$ and the Schottky coefficient $\beta_s = \frac{1}{2} \beta_{PF} = 1.38 \times 10^{-5} \text{ eVm}^{1/2}\text{V}^{-1/2}$. Similarly, the experimental value of β_{PF} at room temperature (303K) is about $1.86 \times 10^{-5} \text{ eVm}^{1/2}\text{V}^{-1/2}$ for InSbTe₃ films, but if using the estimated value of $\epsilon = 35$ obtained from ac measurements (using a PM 6404 programmable automatic RCL meter bridge) the theoretical value of β_{PF} is $1.65 \times 10^{-5} \text{ eVm}^{1/2}\text{V}^{-1/2}$ and the Schottky coefficient $\beta_s = 8.25 \times 10^{-6} \text{ eVm}^{1/2}\text{V}^{-1/2}$. These calculations suggested that the experimental values of β_{PF} is in agreement with the Poole-Frenkel mechanism rather than the Schottky mechanism. However this agreement between the experimental and theoretical values of β_{PF} can not be taken for suggesting the conduction mechanism which will be operated.

Table 1. Variation of β and ϵ with temperature for InSbTe₃ film of thickness 230 nm.

	Poole-Frenkel Model		Jonscher Model	
	Log I vs $E^{1/2}$	Log I vs $E^{-1/2}$	Log I vs $E^{1/2}$	Log I vs $E^{-1/2}$
Temperature K	$\beta/10^{-5} \text{ eVm}^{1/2}\text{V}^{-1/2}$	Dielectric constant ϵ	$\beta/10^{-5} \text{ eVm}^{1/2}\text{V}^{-1/2}$	Dielectric constant ϵ
303	1.86	16.64	1.93	15.46
323	2.15	12.46	2.13	12.69
343	2.6	8.52	2.39	10.08
363	3.24	5.48	3.22	5.55
393	3.97	3.65	3.81	3.97

Table 2. Variation of β and ϵ with temperature for InSbSe₃ film of thickness 250 nm.

Temperature K	Poole-	Frenkel	Model		Jonscher	Model		
	Log I vs	$E^{1/2}$	Log I E_1^{-1}	vs $E^{1/2}$	Log I $E_2^{1/2}$	vs $E^{1/2}$	Log I $E_2^{-1/2}$	vs $E^{1/2}$
	$\beta/10^{-5}$ $eVm^{1/2}V^{-1/2}$	Dielectric constant ϵ	$\beta/10^{-5}$ $eVm^{1/2}V^{-1/2}$	Dielectric constant ϵ	$\beta/10^{-5}$ $eVm^{1/2}V^{-1/2}$	Dielectric constant ϵ	$\beta/10^{-5}$ $eVm^{1/2}V^{-1/2}$	Dielectric constant ϵ
303	2.77	7.5	2.05	13.7	2.77	7.5	2.99	6.44
313	3.3	5.28	2.11	13.09	3.55	4.57	3.38	5.04
333	4.3	3.11	2.64	10.33	3.8	3.99	3.97	3.65
353	4.8	2.5	3.33	5.19	4.3	3.11	4.36	3.03
373	5.8	1.71	4.37	3.02	5.17	2.15	5.38	1.99
393	7.1	1.14	5.09	2.22	6.53	1.35	7.22	1.11

It is clear also from Table (1&2) that the experimental values of β_{PF} for both InSbSe₃ and InSbTe₃ films show an apparent dependence on temperature which increase with increasing temperature through the investigated range.

Various models are put forward to explain the anomalous values of β . The first one [30] is based on the presence of shallow neutral traps at energy E_t and a deep donor level at energy E_d , with the Fermi level lying centrally between them. E_t and E_d were measured from the bottom of the conduction band. The second model [31] is based on donor composition and the presence of acceptors such that the relative densities of donor and acceptor sites control the slope of log I versus $E^{1/2}$ graph.

Hill [7] proposed that donors and traps co-exist and the density of ionized donor is only a small fraction of the total density of ionized donor. If this is the case, then ionization of donor by Poole-Frenkel emission can stimulate further ionization by other processes. The initial ionized donor acts as a capture center for the electron released from the second donor, giving rise to an effective mobile donor and anomalous values of β .

The variation of I (in logarithmic scale) vs $E^{1/2}$ at room temperature (303K) for both InSbSe₃ and InSbTe₃ films are shown in Fig. (4) It is clear from this figure that for a constant value of the applied voltage, a decreases in electrical conductivity for InSbSe₃ than for InSbTe₃ films is observed. The presence of a high concentration of localized states in the band structure is responsible for the larger value electrical conductivity the case of InSbTe₃ films [36].

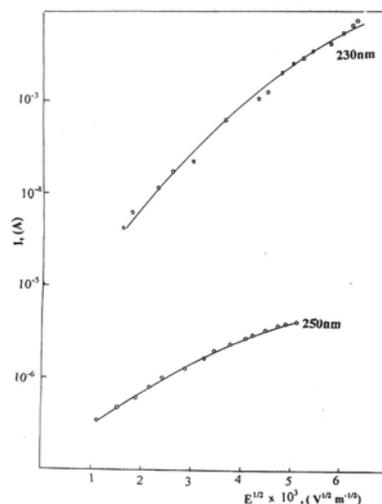


Fig. (4) I (in logarithmic scale) vs $E^{1/2}$ plots for the InSbSe₃ and InSbTe₃ film of thickness 250 and 230 nm respectively at room temperature.

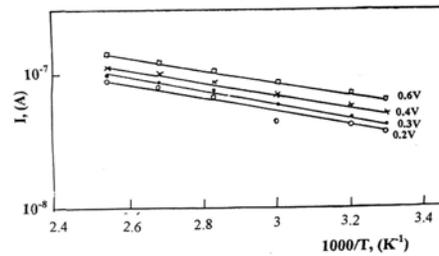
Using the values of β_{PF} (listed in Tables (1&2)), the values of dielectric constant at different temperatures were estimated according to equation (2) and are listed in the same Table. It is observed that the values of ϵ at any given temperature for InSbTe_3 films are higher than those there values for InSbSe_3 films. This can be explained on the basis of chemical bonds present in the two system [32,33] as well as the electro negativity of Se atoms is stronger than Te atom [34,35] (see Table (3)). Accordingly the bonding energy In—Se and Sb—Se are stronger than In—Te and Sb—Te bonds respectively. So InSbTe_3 can be response to the electric field much easier than InSbSe_3 .

Table 3.

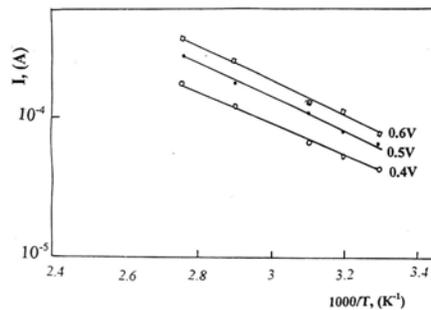
Elements	In—Te	Sb—Te	In—Se	Sb—Se
Electronegativity Difference	0.4	0.2	0.7	0.5

4.2 I-T characteristics

The temperature dependence of current was measured for several constantly applied voltages, having in mind the importance of these characteristics for the proper choice of charge transfer model. Fig. 5 (a&b) show the temperature dependence of the current was studied for the InSbSe_3 and InSbTe_3 films respectively, at different constant applied voltages in the temperature range (303-393K). It is clear from this figure that all obtained relations are straight lines indicating that the conduction in these films takes place through an activated process having a single activation energy in the considered temperature range.



(a)



(b)

Fig. 5. (a) I (in logarithmic scale) vs $1000/T$ for InSbSe_3 film at different applied voltage.
(b) I (in logarithmic scale) vs $1000/T$ for InSbTe_3 film at different applied voltage.

The activation energy ΔE_σ was calculated using the following relation:

$$I = I_0 \exp - \Delta E_\sigma / kT \quad (3)$$

Where I_0 is the current extrapolated to $1/T = 0$. The slopes of the lines do not vary appreciably with the applied voltage (see ΔE_σ values in Table (4)). It was found that the electrical conductivity of InSbTe_3 is higher than that of InSbSe_3 . This result is attributed the nature of chemical bonding in the system being tested. Since the bonding energies of In—Se and Sb—Se are stronger than In—Te and Sb—Te bonds [32,33]. This is confirmed by the values of electro negativity difference for elements given in Table (3). Larger the difference, the more likely bond will form [35].

Table 4. Activation energy of InSbTe_3 film of thickness 230 nm and InSbSe_3 film of thickness 250 nm at various applied bias.

InSbTe_3		InSbSe_3	
Applied bias (V)	Activation energy (ΔE_σ), eV	Applied bias (V)	Activation energy (ΔE_σ), eV
0.4	0.176	0.2	0.108
0.5	0.190	0.3	0.099
0.6	0.194	0.4	0.095
		0.6	0.108

4.3 Poole-Frenkel mechanism and its modification

To determined whether the Poole-Frenkel effect is the dominant mechanism in the high field region, the data of Fig. 3 (a&b) are plotted in accordance with normal high electric field theory to establish the Poole-Frenkel effect if it dominant. Fig 6(a&b) shows a plots of IE^{-1} (in logarithmic scale) vs $E^{1/2}$ as required by the Poole-Frenkel equation [27]. The derived value of β_{PF} and ϵ are listed in Table (1&2) for both systems. It is observed that the field lowering coefficient β_{PF} increases with increasing temperature through the investigated range.

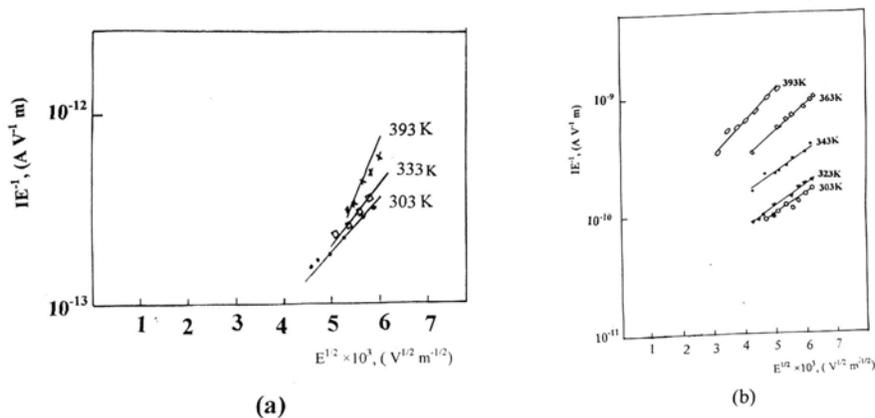


Fig. 6. (a) IE^{-1} (in logarithmic scale) vs $E^{1/2}$ plots for the InSbSe_3 film at different temperatures. (b) IE^{-1} (in logarithmic scale) vs $E^{1/2}$ plots for the InSbTe_3 film at different temperatures.

Based on the Jonscher's model [37], which takes account of the emission of electrons from sites in one particular direction in space with respect to the applied field, the data given in Fig.2(a&b) for both InSbSe_3 and InSbTe_3 films can be analyzed. Accordingly, plots of $IE^{1/2}$ (in logarithmic scale) vs $E^{1/2}$ are shown Fig.7 (a&b) for the two systems. The obtained values of β and ϵ are listed in Tables (1&2). Fitting the room temperature data to the Jonscher's model is an indication that a modified Poole-Frenkel process is operating.

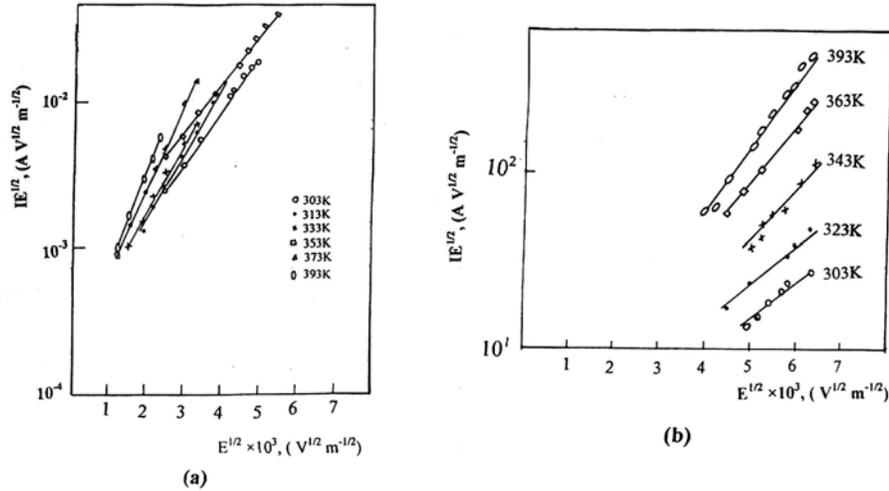


Fig. 7.(a) $IE^{1/2}$ (in logarithmic scale) vs $E^{1/2}$ plots for the $InSbSe_3$ film at different temperatures. (b) $IE^{1/2}$ (in logarithmic scale) vs $E^{1/2}$ plots for the $InSbTe_3$ film at different temperatures.

Fig. 8(a&b) shows the $IE^{-1/2}$ (in logarithmic scale) vs $E^{1/2}$ characteristics for conduction model in which carrier motion are assumed to travel for a constant period of time before being trapped. The deduced values of β and ϵ are also given in Tables (1&2). From these Tables it is notice that a large change in β and ϵ occurred with increasing temperature for the two systems.

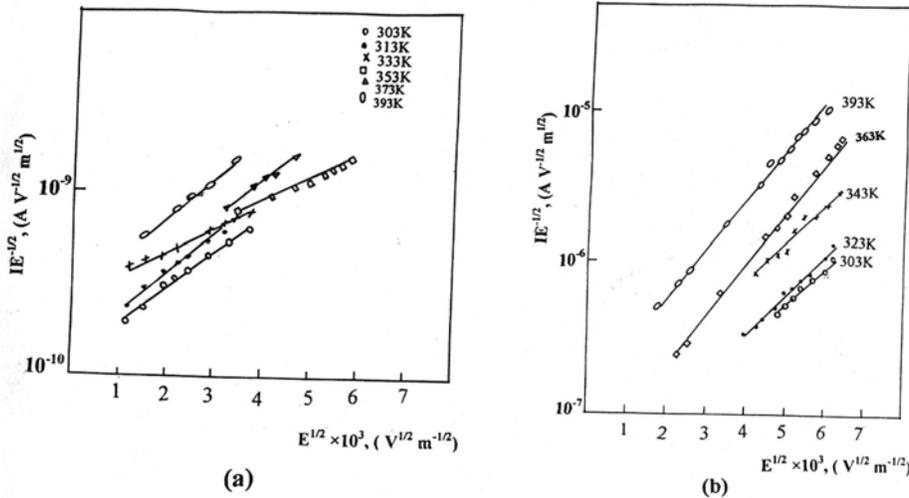


Fig. 8. (a) $IE^{-1/2}$ (in logarithmic scale) vs $E^{1/2}$ plots for the $InSbSe_3$ film at different temperatures. (b) $IE^{-1/2}$ (in logarithmic scale) vs $E^{1/2}$ plots for the $InSbTe_3$ film at different temperatures.

Both normal Poole-Frenkel and Jonscher models are capable to explain a change in β and ϵ for the tested systems with temperature in the investigated range as shown in Tables (1&2).

5. Conclusion

In the above sections we analyzed the electrical conduction in $InSbX_3$ ($X = Te$ or Se) films. The I-V characteristics of thermally evaporation $InSbX_3$ ($X = Te$ or Se) amorphous thin films were obtained in the temperature range (303-393K) and thickness range (230-490nm). The characteristics

exhibited a transition from an ohmic region at low applied voltage to a non-linear region at higher voltage arises from lowering of the potential barriers by a high electric field. The behavior of the non-ohmic region of the I-V characteristics can be understood in terms of both simple and modified Poole-Frenkel mechanisms.

The dependence of ohmic current on the temperature corresponds to a thermally activated process with single activation energy. The values of activation energy and the dielectric constant for both InSbSe₃ than for InSbTe₃ films were investigated.

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