

Abstract Submitted
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Molecular origin of the giant conductivity enhancement in $(\text{Ag}_2\text{S})_x(\text{As}_2\text{S}_3)_{1-x}$ glasses¹ CHAD HOLBROOK, P. BOOLCHAND, University of Cincinnati — The solid electrolyte additive Ag_2S is found to homogeneously alloy with base As_2S_3 glass at low concentrations ($x < 6\%$, single: $T_g = T_g^{high} \sim 210\text{C}$), but it rapidly segregates as a Ag-rich glass phase at medium concentrations ($6\% < x < 20\%$, bimodal : T_g^{high} and $T_g^{low} \sim 170\text{C}$), and becomes the principal glass phase populated at higher $x > 35\%$ (single: T_g^{low}) as revealed by modulated calorimetric measurements. The stoichiometry of the Ag-rich (T_g^{low} phase) is suggested to be near AgAs_3S_7 at $x \sim 25\%$ but becomes closer to that of Smithite (AgAsS_2) at $x > 40\%$, as revealed by Raman scattering. In the $9\% < x < 14\%$ composition range, one observes, in calorimetric experiments, the opening of a reversibility window, and a pronounced increase in the fractional population, $R(x)$ of the Ag-rich glass phase, both of which correlate well with the 5-orders of magnitude increase in electrical conductivity^{1,2} across this compositional interval. In the same interval molar volumes on our samples show a local plateau. These observations suggest a new interpretation of the giant electrical conductivity enhancement observed at $x > 15\%$ in the present electrolyte glass system. ¹ E.A. Kazakova and Z.U.Borisova, Fiz. Khim.Stekla **6**, 424(1980).

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