

Abstract Submitted
for the MAR14 Meeting of
The American Physical Society

Consequences of the superstrong nature of chalcogenide glass-forming liquids at select compositions¹ KAPILA GUNASEKERA, SIDDHESH BHOSLE, PUNIT BOOLCHAND, University of Cincinnati, MATTHIEU MICOULAUT, University of Paris VI — Growth of homogeneous melts of stoichiometric compositions of chalcogenides is facilitated by underlying crystalline phases. Such is not the case for non-stoichiometric melt compositions in which, for example, variation of fragility (m) from complex specific heat measurements show global minimum [1] at an extremely low value ($m=14.8(0.5)$) in the 21.5% $<x <23\%$ range of Ge in homogenized $\text{Ge}_x\text{Se}_{100-x}$ melts. This has unwittingly led to variability of results in physical properties of *non-stoichiometric* melts/glasses due to their heterogeneity. By directly mapping melt stoichiometry variation along a quartz tube as a function of reaction time of starting materials at a fixed temperature $T > T_g$ over days, we have observed a slowdown [1] of melt-homogenization by the super-strong melt compositions, 21.5% $<x <23\%$. This range, furthermore, appears to be correlated to the one observed between the β -exible and stressed rigid phase in network glasses.

[1] K.Gunasekera et al. , J.Chem Phys. 139, 164511 (2013).

¹Supported by NSF grant DMR 08-53957.

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Date submitted: 15 Nov 2013

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