

Excess light scattering in glass formers with a liquid-liquid phase transition

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Several glass-forming liquids are known to exhibit excess light scattering due to the long-range density fluctuations which can exceed 100 nm [1]. These fluctuations are known as “Fischer clusters” which can not be understood by considering density fluctuations only.

Triphenyl phosphite (TPP) is known to show a liquid-liquid phase transition and it can be explained by the existence of a second order parameter coupled to the density [2, 3]. To investigate the relationship between Fischer clusters and the liquid-liquid phase transition, we have carried out dynamic and static light scattering measurements on TPP.

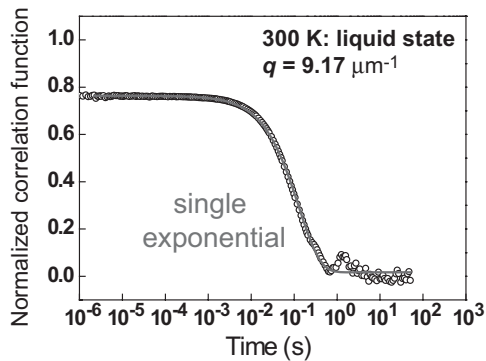


Figure 1: Ultra-slow mode of TPP obtained by dynamic light scattering at wavenumber $q=9.17\mu\text{m}^{-1}$.

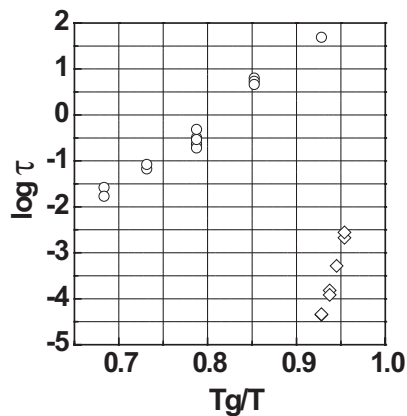


Figure 2: Relaxation time of ultra-slow mode (\circ) at $q=25.1\mu\text{m}^{-1}$ and structural relaxation (\diamond) for TPP.

We found that TPP shows properties common to Fischer clusters. Figure 1 shows that TPP has an ultra-slow mode in the liquid state. The time scale of the ultra-slow mode is much slower than that of the structural relaxation (Fig. 2). The relaxation time depends on q^2 , which suggests the mode is diffusive.

Static light scattering intensity increases with a decrease in temperature in Fig. 3. The intensity at each temperature shows the Ornstein-Zernike-type q -dependence.

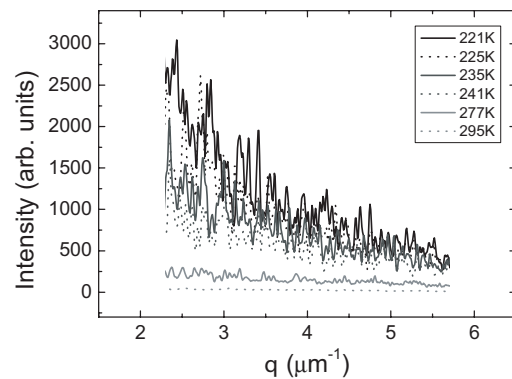


Figure 3: Static light scattering intensity for TPP at several temperatures. (melting point $T_m=297$ K, glass transition temperature $T_g=205$ K.)

Our preliminary results show that the correlation length obtained from the static light scattering increases near the spinodal temperature of the liquid-liquid phase transition. This suggests that Fischer clusters might reflect critical phenomena associated with the liquid-liquid phase transition.

References

- [1] E.W. Fischer, *Physica A* **201**, 183 (1993).
- [2] H. Tanaka, R. Kurita, and H. Mataka, *Phys. Rev. Lett.* **92**, 025701 (2004).
- [3] H. Tanaka, *Phys. Rev. E* **62**, 6968 (2000).