

Structural and microscopic relaxations in a charged colloidal glass under aging

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The Q-dependence of the fast τ_1 and slow τ_2 relaxation times under aging of an anisotropic charged colloidal system, Laponite, has been studied. This complex system, characterized by a microscopic competition between attractive and repulsive interactions, ages spontaneously from an initial liquid state towards multiple arrested states such as gel and glass depending on clay and salt concentrations [1,2]. Here samples in salt free water conditions at weight concentration $C_w = 3.0$ wt% where the system forms a Wigner glass in few hours have been investigated. Through X-ray Photon Correlation Spectroscopy (XPCS) at the Troika beamline ID10 of ESRF, Neutron Spin Echo (NSE) at the spectrometer IN15 of ILL and a five angle Dynamic Light Scattering (DLS) a wide range of exchanged wavevectors Q and times has been explored. The microscopic relaxation time τ_1 does not change significantly with waiting time and it scales always as Q^{-2} , depicting the Brownian diffusive nature of particles motion. On the other hand, the structural relaxation time τ_2 shows two distinct behaviors: an initial exponential growth with waiting time associated to a Q^{-2} dependence again related to the Brownian diffusive dynamic, and a subsequent power law increase with waiting time characterized by a Q^{-1} dependence, depicting the discontinuous hopping diffusion of caged particles [3].

References

- [1] - B. Ruzicka and E. Zaccarelli, *Soft Matter* **7**, 1268 (2011).
 - [2] - B. Ruzicka, E. Zaccarelli, L. Zulian, R. Angelini, M. Sztucki, A. Moussaid, T. Narayanan and F. Sciortino, *Nature Materials* **10**, 56 (2011)
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