
Impact of structural water on sub-T_g relaxations in glass

Joachim Deubener^{*1}, Stefan Reinsch², Harald Behrens³, and Ralf Müller²

¹Institute of Non-Metallic Materials, Clausthal University of Technology (TU Clausthal) –
Zehntner-Straße 2A, D-38678-Clausthal-Zellerfeld, Germany

²Federal Institute for Materials Research and Testing (BAM) – Richard-Willstätter-Straße 11, D-12489
Berlin, Germany

³Institute of Mineralogy, Leibniz University Hannover (LU Hannover) – Callinstr. 3, D-30167
Hannover, Germany

Abstract

Structural relaxation at temperatures below glass transition provides the basis for aging and fatigue phenomena of glasses. Different water species are discussed to contribute to sub-T_g relaxation. Their dynamics are found to be decoupled from the cooperative rearrangements of the glassy network (alpha-relaxation) and the local motions of diffusional transport of ionic species (gamma-relaxation). Using mechanical spectroscopy, two types of new relaxations peaks (beta-relaxations) are evident in hydrous soda-lime silicate and hydrous sodium borosilicate glasses, which are assigned to the contributions of OH-groups and H₂O molecules. Projected characteristic times at ambient temperatures of water-induced internal friction are ca. 10¹ seconds for beta(H₂O) and ca. 10³ seconds for beta(OH). In general, beta-relaxations are assumed to involve stress accommodating rearrangements that are located at non-bridging oxygen (NBO) containing silicon tetrahedra, but in case of the faster beta(H₂O)-relaxation, the discussion also addresses possible jumps of H₂O molecules between network cavities besides a hopping mechanism of protons between H₂O and NBO.

Keywords: Relaxation, Water species, Mechanical spectroscopy

^{*}Speaker