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High frequency acoustic modes in vitreous beryllium fluoride probed by inelastic x-ray scattering

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Inelastic x-ray scattering measurements of the dynamics structure factor have been performed on vitreous beryllium fluoride $(v\text{-BeF}_2)$ at T=297 K in the momentum transfer, Q, range Q=1.5-10 nm⁻¹. We find evidence of well defined high frequency acoustic modes. The energy position and linewidth of the excitations disperse with Q as $\propto Q$ and $\propto Q^2$, respectively, up to about one half of the first maximum of the static structure factor. Their magnitude compares favorably with low-frequency sound velocity and absorption data. The results indicate worth mentioning similarities of the high frequency collective dynamics of different network forming glasses such as $v\text{-B}_2\text{O}_3$ and $v\text{-SiO}_2$. © 2003 American Institute of Physics. [DOI: 10.1063/1.1526097]

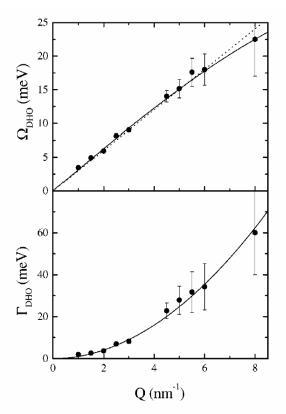


FIG. 2. Q-dependence of the sound frequency and attenuation. Upper panel: experimental sound dispersion (full dots) and best fit through a sine function (full line). The adiabatic sound speed, measured by ultrasonics, is also reported (dotted line). Lower panel: Acoustic attenuation as determined by the Brillouin linewidth (full dots). The solid line is a power law fit to the experimental data and gives $\Gamma_{\rm DHO} = 1.05 Q^{1.96}$.

In conclusion, a room temperature IXS study of glassy BeF₂ has been undertaken in the present work. In accordance with all previous studies in strong and fragile glasses, evi-

dence has been presented for well-defined propagating (highfrequency) acoustic modes, whose frequency position and linewidth scale as $\propto Q$ and $\propto Q^2$, respectively. The longitudinal speed of sound for glassy BeF2 has been estimated to exceed its low-frequency (ultrasonic) limit by almost 5%; a case analogous to that found in studies of vitreous silica. The extrapolation of the high-frequency linewidth conforms nicely with the value obtained from ultrasonic studies, and exhibits a scenario similar to that of vitreous silica. Another similarity which deserves further study is that the temperature dependence (in a T_g -scaled plot) of the nonergodicity factor, as determined from the ratio of the elastic to the total scattered intensity, follows the behavior exhibited by SiO₂ (Fig. 5) while for less strong glasses the drop of f_O is much faster with increasing temperature. Unfortunately, for BeF2 it is up to now available as only one point in such a T_g -scaled plot, which however coincides with the f_O data for silica. Further temperature-dependence studies on BeF₂ are expected to shed more light on this issue.