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Vibrational dynamics and structural relaxation in aqueous electrolyte solutions in the liquid, undercooled liquid and glassy states

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Summary

In very concentrated aqueous electrolyte solutions, the competition between dipolar and ionic forces and most of the system properties, are finely monitored by the concentration. For LiCl, RH₂O (R being the number of water moles per mole of salt) two domains of concentration exist around even hydration numbers R = 6 and R = 4, which can be reversibly undercooled from the liquid to the glassy state. At other concentrations, for R<12, metastability is achieved only in a restricted temperature range.

The collective behaviour as well as individual atomic motions, have been studied by neutron inelastic scattering in the metastable state of LiCl, 6H₂O and LiCl, 4H₂O down to the glass transition.

The study of the structural relaxation by neutron spin echo (NSE) allows us to show that the temperature dependence of the response of the system follows a scaling relation with a cross-over at a phase transition present in the liquid phase.

Increasing hydrodynamic interactions in a liquid is certainly the best manner to analyse the intrinsic and microscopic correlations into the system. So the various motions, collective, relaxational, individual and their coupling can be revealed by approaching glass formation conditions.

LiCl, RH₂O (or LiCl, RD₂O) is one of the most suitable system because by changing R (the number of water molecules per molecule of salt) we change strongly the nature of the medium and by using isotopic substitution on Li⁺, Cl⁻, D we can differentiate the dynamics of the various atoms or ions.

Changing R gives the possibility to analyse two types of liquids: the first one, for R>6 is intrinsically inhomogeneous and described by a regular mixture of pure water and the hydrate LiCl, 6H₂O (1). The second one, for R<6 is a mixture in which the competitions between dipolar forces and coulombic forces are finely monitored.

In this paper, we summarize some recent analysis of different kinds of dynamical responses which lead to some questions on their non linear behaviour.
The **total dynamical responses** of the electrolyte LiCl, 6H2O and LiCl, 4H2O have been analyzed between 0.3 meV and 120 meV, i.e. in a domain of frequencies covering collective vibrational response and individual motion of ions.

1. **Dynamical structure factor.**

From the **dynamical structure factor** $S(q,\omega)$ measured with the IN 6 instrument (ILL, Grenoble) at a resolution of 80 μeV, for LiCl, R H2O (with R=6), two types of informations can be derived:

(i) - the **frequency distribution** $G(E)$ ($E=\hbar\omega$) (extracted as in ref (3), neglecting the incoherent contribution to the scattering), which permits an analysis of the characteristic frequencies (2) which define this low energy "vibrational network" (3).

(ii) - The **central mode**, i.e. diffusive processes from the quasi elastic behaviour around $E=0$.

- In the frequency distribution, three fundamental modes of the H-bonded network dominate: bending at 7 meV, restricted translational motion or stretching at 23 meV, and restricted rotational motion or libration at 85 meV (see figure 1). The identification of these modes has been made by comparison with experiment on pure water (2) (10).

Fig. 1. Generalized density of states obtained for LiCl, 6H2O in the glassy and liquid states and assignments of the principal bands for the liquid state (see 2).

The **influence of temperature** on the different modes has been followed in this disordered system down to the non ergodic situation ($T < 142$ K for LiCl, 6H2O). (Fig. 2)

Fig. 2. Evolution of the three fundamental bands characterizing the bending, stretching, librational motions assigned to H bonded H2O in LiCl, 6H2O versus temperature.
The broadening of the bending motion which is large in the liquid state is correlated to the existence of different microscopic dynamical configurations. This mode has a significant asymmetry towards high energies which increases with temperature, giving some support to a delocalization of the excitations or a fluctuating coupling between different microscopic configurations (12).

Among these modes, stretching is the most influenced by the glass transition or equivalently by T₀ i.e. the lowest temperature limit of liquid metastability (2). However the limited range of energy that we were able to reach, does not permit us to fully analyze the low temperature behaviour for librational modes. There is another temperature of interest which is characteristic of a third order phase transition in the liquid (quoted by Tₚ = peritexy at 207 K). It appears that the stretching mode is nearly temperature independent in the low temperature supercooled liquid for T₀ < T < Tₚ. This temperature range is characterized by additional microscopic configurations (hydration 5) compared to those of the room temperature liquid.

B - Quasi elastic data, i.e. the central mode, permits to obtain informations on the individual motion of H or D in H₂O and D₂O. At the composition R=6, the system is molecular and although there is no correct diffusion's analysis of the different elements Li⁺, Cl⁻ and H₂O, the trends give D_Li⁺ < D_Cl⁻ < D_H₂O at 25°C (4).

- The proton delocalization due to vibrations is deduced from the Debye Waller component. It is obtained from the total quasielastic intensity, I and is shown in fig. 3a and fig. 3b for LiCl, 6H₂O and LiCl, 4H₂O.

Insert: Mean square displacement of H₂O in LiCl, 6H₂O and in LiCl, 4H₂O in liquid and supercooled liquid versus (T - T₀), T₀ being the lowest temperature limit of the liquid metastability.

* Proton delocalization in pure water (11)
The mean square displacement obtained for $H_2O$ in LiCl, $6H_2O$ and LiCl, $4H_2O$ and referred to $T_0$, lowest temperature limit of the metastability (10) is temperature dependent in the liquid state and nearly temperature independent in the supercooled liquid. This effect is induced by a restricted translational motion due to interactions of $H_2O$ with Li$^+$ and Cl$^-$. 

- The diffusion coefficient of $H_2O$ can be obtained from $S(q, \omega)$ and compared to the ordinary diffusion law (5). As a matter of fact, the diffusion can be coupled to the hydrodynamics of the medium, the coupling being function of the kinematic viscosity $D_T$, according to $\delta = D / (D + D_T)$ (5). The $q$ dependence of the half width of the central mode given in fig.4 for LiCl, $6H_2O$ permits to obtain diffusion coefficient in agreement with literature (4).

\[ \Delta E / q^2 \text{ (meV A}^2) \]

**LiCl, 6H$_2$O**

<table>
<thead>
<tr>
<th>Temperature</th>
<th>223K</th>
<th>192K</th>
</tr>
</thead>
<tbody>
<tr>
<td>$D$ (10$^{-9}$ m$^2$/s)</td>
<td>0.13</td>
<td>0.05</td>
</tr>
</tbody>
</table>

The narrow of the central mode for all $q$ values inferior to $q_{max} (= 1.9$ Å$^{-1}$) confirms the coupling of this mode with hydrodynamics (6).

![Fig. 4. FWHM of $S(q, \omega)$ divided by $q^2$ for LiCl, 6H$_2$O at different temperatures above and below $T_p$.](image)

**Fig. 5.** $H_2O$ diffusion coefficient evolution with temperature in LiCl, RH$_2$O for various $R$. 

- The $H_2O$ diffusion coefficient obtained in this molecular system, for $R = 6$ and $R = 4$ permits to confirm the strong influence of the solvation on the dynamics of $H_2O$ when $R \leq 6$ (fig 5).
2. Relaxational behaviour.

The relaxational behaviour has been obtained on an "adequate" range of time (10^-12 sec. to 10^-9 sec) by NSE experiment performed on IN11 at ILL(9) on LiCl, 6D_2O at q = q_{max} = 1.9 Å⁻¹, q_{max} being the momentum transfer of the main peak of the structure factor at room temperature.

We deduce from this data (7), the evolution of the characteristic relaxational time τ_{NSE} which can be compared to the one obtained from viscosity (8) (fig. 6). Two scaling laws (10) give the evolution of τ, with two characteristic temperatures of the system: lowest limit of the metastability T_0 and crystallisation temperature for the first hydrate formed: R = 5

![Figure 6a](image)

Fig. 6a. Neutron spin echo response at q = q_{max} = 1.9 Å for LiCl, 6D_2O measured and analyzed according to ref. (7).

![Figure 6b](image)

Fig. 6b. Relaxational time deduced from NSE (•) for LiCl, 6D_2O compared to τ_S obtained from kinematic viscosity (s) according to ref. (8) for LiCl, 5.75 H_2O.

This result confirms the connexion between the relaxational process and the third order phase transition, at T_p. Slowing down is observed in the supercooled liquid scaled to the lowest limit T_0 of the metastability and the non-ergodic character appears down to T_0 (7).

In conclusion, in these molecular aqueous solutions, collective vibrational motion of H_2O such as stretching, mean square displacement and relaxational response in the hydrodynamic regime and at finite momentum transfer are sensible to the approach of some characteristic temperatures of the system such as T_0 and T_p an apparent third order transition in the vicinity of the equilibrium melting point. The existence of diffusive motion confirms the coupling with the hydrodynamics. The extension of these studies to the more concentrated domain, offers the opportunity to study the coupling between ionic conduction and relaxational slowing down.
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" H-Bond characteristic in supercooled and vitreous aqueous solutions"
" Ice nucleation and growth in Electrolytes : role of ionic forces"