Direct Contact versus Solvent-Shared Ion Pairs in NiCl₂ Electrolytes Monitored by Multiplet Effects at Ni(II) L Edge X-ray Absorption

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We investigate the local electronic structure in aqueous NiCl₂ electrolytes by Ni L edge X-ray absorption spectroscopy. The experimental findings are interpreted in conjunction with multiplet calculations of the electronic structure and the resulting spectral shape. In contrast to the situation in the solid, the electronic structure in the electrolyte reflects the absence of direct contact Ni—Cl ion pairs. We observe a systematic change of the intensity ratio of singlet- and triplet-related spectral features as a function of electrolyte concentration. These changes can be described theoretically by a change in the weight of transition matrix contributions with different symmetries. We interpret these findings as being due to progressive distortions of the local symmetry induced by solvent-shared ion pairs.

Introduction

Most properties of electrolyte solutions depend on the ability of the solvent and solute to interact and hence on the nature of the complex ion formation. One important property is the Gibbs free energy of solvation, which requires an assumption of the effective ionic radius \( +\Delta R \). Here, \( \Delta R \) is a function taking into account the first hydration shell.¹ Due to the importance of ion behavior in electrolyte solutions, ions in electrolytes have been studied using many theoretical and experimental techniques. Molecular dynamics (MD) simulations, for instance, have shown that for a 1 M NaCl solution about 25% of the ions are included in neutral, solvent-shared clusters formed by a minimum of four ions.² MD studies have suggested an association of Na⁺ and Cl⁻ ions in solution via a two-step mechanism, first the establishment of long-distance ion pairs separated by hydration shells, followed by direct contact ion pair formation.³⁻⁴

Several methods have been used to probe the local geometrical structure of ions in solution experimentally. In an X-ray diffraction study of saturated NiCl₂ solutions, Waizumi et al. have reported the existence of mixed-ligand chloroaqua octahedral complexes in addition to 6-fold-water-coordinated Ni ions, i.e., the existence of direct contact ion pairs.⁵ A detailed picture for ion clustering in solution, including the solvent-shared hydration shell, has been given by Fulton et al.⁶ On the basis of their combined X-ray absorption fine structure (EXAFS) and extended X-ray absorption fine structure (EXAFS) studies, these authors have suggested a long-range interaction between Ca²⁺ and Cl⁻ in the solution which leads to a solvent-shared ion pair (Ca²⁺−OH₂−Cl⁻). Direct contact ion pairs between the cations and the anions have been argued to be negligible for CaCl₂ electrolyte solution even at high concentration. This is in agreement with a neutron diffraction study by Badyal et al.,⁷ also providing evidence for the existence of solvent-shared ion pairs.

For Ni²⁺ in aqueous solution, neutron diffraction studies⁸⁻⁹ as well as X-ray diffraction⁵ have shown that Ni²⁺ has a coordination sphere of six water molecules. Inner sphere contact pairs of Ni—Cl have been suggested to exist for 8% of the Ni ions in an X-ray diffraction study of a 3 M NiCl₂ aqueous solution.¹⁰ X-ray diffraction of NiBr₂ aqueous solution has shown strong experimental evidence for ion pair formation at 2 M concentration within octahydration geometry.¹¹ Here we study the range from diluted (50 mM) to concentrated (1.5 M) aqueous NiCl₂ solutions, where we expect a transition in the importance of interionic interactions.

Recently, we have studied the behavior of ions in electrolyte solutions at ambient pressure and temperature, combining XANES studies with density functional theory (DFT) simulations,¹² to investigate the effect of concentration¹³ and solvent¹⁴ on the ion—ion interaction in aqueous NaCl solution. In this paper we present XANES spectra for the Ni²⁺ L edge in aqueous NiCl₂ electrolyte solution as a function of the concentration, starting from solid NiCl₂. Two distinctive spectral features can be assigned as fingerprints of direct contact ion pairs and solvent-shared ion pairs in the solution. The spectra have been analyzed by the means of a charge-transfer multiplet simulation.¹⁵⁻¹⁷ The multiplet approach has been extensively used in the analysis of L edge spectra of transition metals, where it is established as a method for probing the metal—ligand charge transfer.¹⁸⁻²¹

Experimental and Computational Techniques

The experiments were performed at two different light sources. First, measurements were conducted at the Advanced Light Source (ALS), Lawrence Berkeley National Laboratory, using the liquid end station at beamline 7.0.1.²² In this end station, soft X-rays are coupled into a fixed-volume liquid
sample through a Si$_3$N$_4$ membrane of 100 nm thickness and lateral size of 1.0 $\times$ 1.0 mm. The sample holder was mounted in a vacuum chamber with a pressure of approximately 1 $\times$ 10$^{-9}$ mbar. The resolution of the beamline monochromator was set to be 0.3 eV at 850 eV. Further details about the experimental setup have been discussed before.$^{23,24}$

The measurements were repeated using the LIQUIDROM flow jet end station at BESSY, Berlin, at beamline U41-PGM. The liquid was injected into the experimental chamber as a continuous flowing low-pressure liquid jet of about 1 mm diameter. The interaction chamber was filled with 1 atm of helium, with the helium being continuously refreshed. The vacuum parts of the beamline were separated from the atmospheric helium setup via two stages of differential pumping, and a Si$_3$N$_4$ membrane of 100 nm thickness and 0.5 $\times$ 0.5 mm lateral size. The resolution of the monochromator was set to be 0.2 eV at 850 eV.

In both experiments, the X-ray absorption spectra from liquid samples were recorded in fluorescence yield (FY) mode using a gallium arsenide photodiode 5 $\times$ 5 mm$^2$ in size. The beamline energy was calibrated with respect to the 852 eV peak from a NiO single-crystalline sample.

The samples were prepared from commercially available NiCl$_2$·6H$_2$O salts, purchased from Sigma-Aldrich. The powder has 99.8% purity, and was used without further purification. For the measurements, a layer of powder was held on a flat substrate using electrically conductive carbon adhesive tape. All Ni(II) solutions were freshly prepared before the experiment.

We use charge-transfer multiplet (CTM) calculations based on a combination of the Cowan code$^{25}$ for atomic multiplets, the crystal field program of Butler,$^{26}$ and a charge-transfer model Hamiltonian$^{15}$ to calculate the electronic structure at the Ni site and the resulting XANES spectra. The calculations include the electronic Coulomb interactions and the spin–orbit coupling on every open shell. The XANES spectra were simulated on the basis of the crystal field strengths and the charge-transfer parameters, as explained in detail before.$^{15}$ All spectra shown in this work have been broadened by Lorenzians of 0.2 and 0.3 eV for L$_3$ and L$_2$ edges, respectively, and a Gaussian of 0.25 eV to account for lifetime broadening and the experimental resolution, respectively.

There are several codes calculating the single-particle X-ray absorption which can explain K edge absorption well. Nevertheless, these codes typically agree poorly with the experiment in the case of the L$_2,3$ edge. The reason for this is that in general the density of states calculated theoretically is not observed in such an X-ray process. The density of states is affected by the strong overlap between the core wave function and the valence wave function. When the excitation takes place, the core orbitals are partially filled as in Ni(II) 2p$^6$3d$^9$, and different pd multiplets can be excited. This multiplet effect was shown to be of the same order of magnitude as the spin–orbit coupling.

As we will see below, our experimental observations can be understood in a framework of distorted symmetry. Distortion of the local symmetry around the Ni ions influences which multiplet states can contribute to the spectra (depending on their symmetry). It is thus important to briefly discuss the symmetry of the involved states and operators on a group-theory basis.

In X-ray absorption, the 2p core electron is transferred to 3d orbitals, and the transition can be described as 2p$^6$3d$^9$ $\rightarrow$ 2p$^5$3d$^9$. The ground state has $^3$F$_4$ symmetry, and there are 12 term symbols for the final state ($^1$P$_1$, $^1$D$_2$, $^3$F$_5$, $^3$P$_{0,1,2}$, $^3$D$_{1,2,3}$, and $^3$F$_{2,3,4}$). The energies of the final states are described by the 2p3d Slater–Condon parameters, the 2p spin–orbit coupling, and the 3d spin–orbit coupling. According to the selection rules, the 12 final states are reduced to 4 accessible final states, $^3$F$_5$, $^3$D$_{3,5}$, and $^3$F$_{3,4}$. By applying a cubic crystal field, the symmetry changes from spherical (SO$^3$) to octahedral (O$_h$), causing the p orbital to be branched to a T$_{1u}$ state. A d orbital splits into E$_g$ and T$_{2g}$ states. The dipole transition operator has p symmetry on the atomic level, and by applying the crystal field, it branches to T$_{1u}$ symmetry. The 3d wave function ground state splits into $^3$A$_2$, $^3$T$_1$, and $^3$T$_2$ wave functions in O$_h$ symmetry, where $^3$A$_2$ is the lowest energy state. In other words the eight 3d electrons fill all T$_{2g}$ states plus the two spin-up E$_g$ states, leaving two spin-down E$_g$ holes. The symmetry of these two E$_g$ holes is written as $^3$A$_2$. The CTM program works in double group symmetry, after the inclusion of spin–orbit coupling. This implies that the spin multiplicity must be branched as well. The $^3$A$_2$ state has S = 1, which is branched to T$_1$ in O$_h$ symmetry. Applying spin–orbit coupling, i.e., multiplying T$_1$ and A$_2$ symmetry, yields a T$_2$ overall symmetry ground state. Four dipole matrix elements as classified by their symmetry can contribute to the 2p$^5$3d$^9$ $\rightarrow$ 2p$^5$3d$^9$ transition, namely, (T$_2$|T$_1$|T$_1$), (T$_2$|T$_1$|E), (T$_2$|T$_1$|T$_2$), and (T$_2$|T$_1$|A$_2$). The final spectra are then obtained as a linear superposition of these contributions. Charge transfer from Cl$^-$ to Ni$^{2+}$ is calculated by taking into account the 2p$^5$3d$^9$L and 2p$^5$3d$^9$L configurations (L referring to a ligand hole) based on the Anderson impurity model$^{28–33}$ and related short-range model Hamiltonians that are applied to core level spectroscopies.

**Results and Discussion**

In Figure 1, the Ni(II) L edge electron yield XANES spectrum for NiCl$_2$ solid is presented as a reference for the liquid measurements. The 2p (L edge) XANES spectrum of Ni(II) has...
a number of peaks that split into two main regions, one at 853 eV and one at 870 eV. The structures around 853 eV are related to the L3 edge (P1 and P2 peaks). The features around 870 eV are related to the L2 edge (P4 and P5 peaks); the 2p spin–orbit coupling leads to a splitting of approximately 18 eV. Peak P1 and peak P4 relate to a 2p5 3d9 final state of triplet character, i.e., where the spins of the 2p shell and 3d electron are parallel. In contrast, peaks P2 and P5 relate to a singlet final state. A satellite peak above the L3 edge at around 858 eV appears for the solid Ni(II) (P3 peak). The spectrum agrees well with published data by G. van der Laan et al.32 Within the L3 edge, two peaks, P1 and P2, split by 1.6 eV are observed. The P3 satellite peak is known to be due to ligand–metal charge transfer (LMCT) from Cl− to Ni2+,15 which is possible due to the close proximity of Ni and Cl in the solid. An overview spectrum for NiCl2 aqueous solution is shown for comparison. As for all liquid spectra, the XANES spectrum was measured by the FY. The changes in the peak intensities and splittings will be discussed as a function of concentration in the context of Figure 2.

As a function of electrolyte concentration from 50 to 1500 mM, a systematic change in the XANES spectral features is observed as seen in Figure 2. Even at the highest concentration investigated, peak P3 is not detectable against the background noise level in the spectra of the electrolytes. The absence of this peak compared to its presence in the solid reflects the nonexisting or at least strongly reduced amount of direct interaction between Ni2+ and Cl− in the electrolyte solution as compared to the solid. However, given an uncertainty of about 2% in our baselines, our results are still consistent with 8% of the Ni atoms in direct contact pairs with Cl−, as reported for a 3 M NiCl2 solution.10 As a second important difference between bulk and electrolyte spectra, the energy splitting within the L3 edge in the electrolyte solutions is 2.4 eV compared to 1.6 eV for the solid. This increase of the L3 splitting energy can be assigned to the absence of direct contact ion pairs as well, as we will discuss later.

The most prominent intensity change is the increasing P2/P1 intensity ratio for increasing NiCl2 concentration in the solvent. For geometrical reasons, FY-XANES spectra can exhibit distorted intensities if the edge absorption of interest is larger than the background absorption.34,35 This effect becomes important in concentrated samples and will be strong in solid NiCl2. For the solid, we have thus used XANES spectra in electron yield (EY) mode to measure the absorption cross-section (Figure 1), which due to its surface sensitivity does not suffer from these distortions. This approach is not feasible for the liquid samples. Here, we have quantified the concentration-dependent saturation effects. On the basis of the atomic absorption cross-sections, we found the distortion to be negligible for 50 mM NiCl2 solution. We calculate how the absorption cross-section of the 50 mM solution would appear at elevated concentrations, which can be done analytically in conjunction with the known experimental geometry and atomic absorption cross-sections.34,35 These simulated spectra are presented in Figure 2 along with the measured spectra for different electrolyte concentrations.

Clearly, this geometrical saturation effect alone cannot account for the observed intensity changes as a function of electrolyte concentration. It is the remaining effect of concentration on the XANES spectra which will be discussed in detail in this paper. This remaining effect after correction for geometrical
saturation corresponds to the gray shaded areas in Figure 2. It is evident from the XANES spectra that the electronic structure locally at the Ni(II) atoms in the solution changes as a function of electrolyte concentration. The change is such that peak P2 increases relative to P1 for increasing electrolyte concentration. Furthermore, a change in multiplet splitting energy was observed between solid and liquid spectra. In the following, we will rationalize these changes in the electronic structure on the basis of electronic structure calculations, where symmetry parameters are being varied.

In an XANES experiment, a spatial ensemble average over temporally frozen configurations is observed. Rigorously, such a system should be described by an ensemble average over a multitude of different local Ni environments in the liquid, obtained, e.g., by molecular dynamics simulations. Here, we will follow a much simpler route and try to qualitatively understand the observed spectral changes by analyzing one model. To simulate the observed spectral changes, multiplet calculations were performed, where Ni(II) in an $O_h$ symmetric crystal field is considered. This approach is justified by the fact that the highly directed Ni d orbitals favor a quite rigid local symmetry, as known from complex chemistry. For highly concentrated NiCl$_2$ solutions, local structural data were well described in $O_h$ symmetry, as expected for Ni$^{2+}$ with a d$^8$ electron configuration.

In Figure 3 we display the simulated spectra obtained using a crystal field of 1.1 eV for the d$^8$ configuration, and a d$^0$L charge transfer state was included to describe the electron transfer from Cl$^-$ to the valence d orbitals of Ni(II). This configuration interaction with a contribution of the d$^0$L configuration of about 14% can explain the appearance of the P3 peak in the experimental spectra. This d$^0$L charge transfer can only be significant with Ni(II) and Cl$^-$ being in direct contact, as in the solid NiCl$_2$. The significant reduction of the P3 peak intensity upon dissolving the solid NiCl$_2$ in water is consequently explained by the absence of d$^0$L charge transfer; that is, the Cl$^-$ ions have moved out of Ni(II) in the first hydration shell. Furthermore, this treatment reproduces the experimentally observed P1–P2 peak energy splitting, which is 1.6 eV for solid NiCl$_2$ and 2.4 eV for the solution. The simulated spectra have peak separations of 1.65 eV for mixed d$^8$ with d$^0$L configurations and 2.3 eV for a pure d$^8$ configuration. This implies that the ions in solution have rather weak charge transfer, since charge transfer compresses the multiplet configuration. The simulated spectra in Figure 3 reproduce the experimental XANES spectra for the solid NiCl$_2$ well, including the spectral finger print (P3) for the existence of ion pairs. Within pure $O_h$ calculations, the change of the P2/P1 peak ratio in the salt solutions as a function of concentration (Figure 2) cannot be reproduced. Only the most dilute 50 mM Ni(II) solution can be described in perfect $O_h$ symmetry with a pure d$^8$ configuration.

To explain the change of the fluorescence yield as a function of Ni(II) concentration, we mention that the calculated absorption for the d$^8$ configuration in an $O_h$ crystal field consists of four dipole matrix element components added equally, e.g., $(|p_i|) = \alpha|T_2|T_1|E\rangle + \beta|T_2|T_1|E\rangle + \sigma|T_2|T_1|E\rangle + \varphi|T_2|T_1|A_2\rangle$, where $\alpha$, $\beta$, $\sigma$, and $\varphi$ are equal to 1. The origin of these transition dipole matrices has been discussed in detail in the Experimental and Computational Techniques. We observe that, to reproduce the experimental XANES spectra for Ni$^{2+}$ solution as a function of concentration, these four matrix elements are added not equally. Furthermore, analysis shows that the main channels for the singlet states (855 and 872 eV) have $T_1$ character, while the triplet 2p$^3$3d$^9$ states (853 and 870 eV) have mixed $T_2$, E, and $A_2$ character. This implies that one can simulate the variation in triplet (P1 peak) versus singlet (P2 peak) states by varying the ratio of the different coefficients.
α through ϕ and in particular by varying α with respect to the remaining coefficients.

In Figure 4, we plot the resulting variation of the spectral shape of the coefficients α through ϕ, within limits such that the overall spectral shape starts to become unphysical for the extreme values of α and ϕ. This defines a variation interval from 0.5 to 3.0; outside this interval the overall intensity ratio of L1-related vs L2-related peaks clearly disagrees with the experimental spectra in the literature and in this work. Over this parameter interval, both the (T2|T1|T0) and (T2|T1|T2) matrix elements show a strong variation of the P2/P1 intensity ratio, with an inverted sign of the intensity ratio dependence on α and ϕ, respectively. A variation of α from 3.0 to 0.5 while the other coefficients are kept constant at 1.0 does reproduce the variation in spectral shape that is observed experimentally; i.e., it increases the P2/P1 peak intensity ratio. Consequently, linear combinations with variation of all four coefficients can generate theoretical spectra which describe the experimental findings as a function of electrolyte concentration well. The change of the relative contributions of the (T2|T1|T0) matrix elements to the total transition probability reflects a changing weight of the electronic configuration at the Ni2+ ion pair (or ion cluster) formation. In a simple picture one could describe our findings as evidence of local distortions due to the increase of the ion concentration in the electrolyte.

This interpretation is based on experimental electronic structure XANES data in comparison to the multiplet calculations of the spectral shape in O h symmetry and with distortions which one would encounter, e.g., if the crystal field strength and the total symmetry around the Ni ions.

References and Notes


