

X-RAY ABSORPTION SPECTROSCOPY STUDY OF AQUEOUS  
ELECTROLYTE SOLUTION CRITICAL PROPERTIES.

M. IRAR<sup>1</sup>, E. BAZARKINA<sup>1,2</sup>, D. TESTEMALE<sup>1</sup>, O. PROUX<sup>3</sup>,  
A. AGUILAR-TAPIA<sup>1</sup>, I. KIEFFER<sup>3</sup>, W. DEL NET<sup>3</sup>,  
E. LAHERA<sup>3</sup>, M. ROVEZZI<sup>1</sup>, AND J.L. HAZEMANN<sup>1\*</sup>

<sup>1</sup>Inst. Néel, UPR 2940 CNRS - UGA, F-38000 Grenoble,  
France

<sup>2</sup>IGEM RAS, 119017 Moscow, Russia

<sup>3</sup>OSUG, UMS 832 CNRS - UGA, F-38041 Grenoble, France

The goal of this study is quantify the properties of electrolytes at near-critical conditions using X-ray Absorption Spectroscopy (XAS) techniques on BM30b FAME beamline at ESRF (Grenoble, France) using hydrothermal spectroscopy cell and high-pressure autoclave (Testemale et al., 2005). Two types of measurements were performed: transmission XAS density measurements and High Energy Resolution Fluorescence Detection (HERFD) XAS measurements via crystal analyzers (Proux et al., 2017).

With heating from 25 to 500°C at constant pressure (280, 300, 345 and 400bar), the absorption coefficients of chloride and bromide solutions decreases slowly until ~373°C (similarly with pure water), but then increases up to ~380°C, and finally decreases to gas-like values at higher temperatures. These absorption measurements reflect the anomalous density behavior at near-critical T-P-x region. At the same electrolyte concentration 0.3 mol/kg of H<sub>2</sub>O, the relative density increase in this critical zone is more pronounced in order Li < Na < K < Rb < Cs for both bromides and chlorides. Complementary HERFD XAS measurements at Br K-edge in bromide solutions at similar T-P-x indicate that this density phenomenon is probably accompanied by structural changes (ion-pairing). Our new data complement previous synchrotron small angle X-ray scattering measurements (Da Silva-Cadoux et al., 2012) and open new perspectives for studies on electrolyte aqueous fluid properties in near-critical state.

Testemale D. et al. (2005) *Rev. Sci. Instrum.* **76**, 43905 ;

Proux O. et al. (2017) *J. Environ. Quality* (in press);

Da Silva Cadoux et al. (2012) *J. Chem. Phys.* **136**, 044515.