Nuclear Magnetic Resonance Measurement of Ammonia Diffusion in Dense Solid-Liquid Slurries

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Summary

The diffusion of ammonium ions in aqueous solutions was measured by nuclear magnetic resonance (NMR) using the pulsed field gradient (PFG) method. The ammonium ions were obtained from aqueous solutions of ammonium chloride, ammonium sulfate, ammonium bicarbonate, and ammonium hydroxide. The translational diffusion of the ammonium ions was determined by measuring the diffusion of nitrogen nuclei (14N and 15N) in solution. Our results showed that the ammonium diffusion coefficient can be measured in aqueous solutions with concentrations as low as 20 x 10^{-3} M. Typical values measured for the diffusion coefficient of the ammonium ion are 2 x 10^{-5} cm^{2}/s (±10%), similar to the values found for pure water. Due to the effect of the solution pH upon the NMR relaxation parameters for 14N, measurements are constrained to pH values below 8.5. However, 15N labeled ammonia is less sensitive to the solution pH, extending the measurement range to pH of 9.5.

Diffusion measurements were conducted with solutions of varying viscosity and porosity. The results show that the solution viscosity has a measurable impact on the diffusion coefficient. The diffusion coefficient is almost inversely proportional to the relative viscosity of the solution, irrespective of how the viscosity is increased. Further, a randomly-packed porous bed of 200 µm PMMA resulted in a reduction of ~30% in the diffusion coefficient as a result of hindered diffusion.
Acknowledgments

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1.0 Introduction

The flammability and toxicity of ammonia released from the nuclear waste tanks at Hanford have been the subject of several recent studies (e.g., Palmer et al. 1996; Peurrung et al. 1997). These releases may occur episodically, such as the buoyant plume releases occurring in various double-shell tanks (DSTs); gradually through the surface of the waste; or from the partially saturated saltcakes in the single-shell tanks during salt-well pumping.

Scientists at Pacific Northwest National Laboratory have recently set up a one-dimensional convection-diffusion model to capture the gradual transport of ammonia in the DSTs. Using several measured parameters such as the concentration of ammonia in the dome space and within the sludge layer, the model can predict the rate of diffusion of ammonia within the nonconvective solids (sludge) layer. Recent results (Terrones et al. 1997) reveal that the predicted effective diffusion coefficient of ammonia within the sludge layer is comparable to the rate of diffusion of ammonia in pure water. However, this value is nearly 20 times higher than the expected value of the diffusion coefficient of ammonia in a brine solution to be based on the Stokes-Einstein equation.

Inputs to this model are the ammonia concentrations in the dome space and within the waste obtained from the retained gas sampler (RGS) measurements (Shekarriz et al. 1997). Among other problems in measuring the ammonia concentration, the rate at which the ammonia concentration within the RGS extractor volume reaches an equilibrium level has been recognized to be a possible source of difficulty (Shekarriz et al. 1997). Although the waste slurry is agitated in the extractor during the gas extraction process, the flow induced by the mixer could potentially be at a very low Reynolds number, resulting in segregated mixing regimes around the mixing propellers. As a result, the ammonia transport from the slurry would be diffusion-limited and could potentially produce a higher uncertainty in the reported data.

Results of the local RGS measurements for the first five tanks sampled (AW-101, A-101, AN-105, AN-104, and AN-103) show significant variability within the nonconvective layers of each tank, larger than the predicted uncertainty in the measurements (Shekarriz et al. 1997). Whether such variabilities are real or are the artifacts of the measurements depends on the local solubility and diffusion coefficient of ammonia in the nonconvective layers. The solubility of ammonia in high-concentration salt solutions has been measured previously (Norton and Pederson 1995). However, given that diffusion is a crucial parameter quantifying transport, there are currently no experimental data available for ammonia diffusivity in such high-ionic strength, high-pH, and densely loaded sludges.

The nuclear magnetic resonance (NMR) experiments conducted at UC Davis have allowed the diffusion of ammonia in solution to be characterized over a wide range of conditions. In particular, the dependence of the diffusion coefficient upon pH, ammonia concentration, and solution viscosity was investigated. This report summarizes the results, which demonstrate


(b) Hanford waste tanks are designated with the prefix 241-. In this report the prefix is omitted, as it is in common usage.
feasibility for measuring the diffusion of ammonia under conditions similar to those within the tank: high pH, low concentration of ammonia, and solutions within porous media.

Section 2 contains a description of the principles of operation of the NMR diffusion measurement technique. The particular technique used for ammonia diffusivity measurements are presented in Section 3 and the results in Section 4. Section 5 contains the conclusions drawn from this research and Section 6 the cited references.
2.0 Nuclear Magnetic Resonance: Principle of Operation

The most common way to measure diffusion by NMR is to use magnetic field gradient pulses to encode and decode the NMR signal, or the so-called PFG (pulsed field gradient) method. In its simplest version, a PFG experiment consists of applying a pair of field gradient pulses of duration $\delta$ separated by a time $\Delta$, as depicted in Figure 2.1.

The time $\Delta$ is the time in which the spins are allowed to diffuse and is often referred to as the “diffusion time.” The NMR signal is irreversibly attenuated due to the random translational motion of the liquid molecules (self-diffusion) during the period $\Delta$. This attenuation can be written as (Callaghan 1991):

$$S \propto e^{-\gamma^2 g^2 \delta^2 D (\Delta - \delta/3)}$$

$$\propto e^{-q \cdot D}$$

(2.1)

where $\gamma$ is the gyromagnetic ratio of the nucleus, $g$ is the amplitude of the gradient, $\delta$ is the duration of the gradient $g$, $D$ is the diffusion coefficient, and $\Delta$ is the gradient pulse spacing. Thus the signal attenuation can be obtained by varying $g$, $\delta$, or $\Delta$.

To verify the accuracy of the system in measuring the diffusion coefficient, two separate and independent measurements were performed. Figure 2.2 shows the NMR signal attenuation in a PFG experiment for pure water. These experiments were performed on the same sample within a 10-day interval and showed excellent reproducibility. The horizontal axis is the parameter $q$ depicted in Equation (2.1). Note that the signal maintains its linearity over almost three decades on a log-log plot, which indicates that it obeys the exponential law presented in Equation (2.1), and the diffusion coefficient is constant throughout the whole range.

![Figure 2.1. Pulsed Field Gradient (PFG) Experiment](image-url)
Figure 2.2. Signal Attenuation for Water (PFG experiment)
3.0 NMR in Aqueous Ammonia Solutions

3.1 $^1$H Nucleus

The first attempt at measuring ammonia diffusion made use of the $^1$H nucleus, based on the fact that NMR spectroscopy allows one to discern nuclei from different molecules. This effect is called "chemical shift" and is due to the local magnetic fields created by the local molecular surroundings. Moreover, $^1$H has a high gyromagnetic ratio (hence a high NMR sensitivity; see Table 3.1), and it is abundant in the system under study. Unfortunately, as shown in Figure 3.1, the spectrum acquired for $^1$H in ammonia aqueous solutions did not give a unique or distinguishable signal. This is explained by the fact that the studied nucleus ($^1$H) is in fast chemical exchange. In other words, $^1$H is "jumping" from the ammonium ions to the water molecules constantly, faster than the time scale of the NMR measurement. The effect of rate in chemical exchange is depicted in Figure 3.2.

Table 3.1. NMR Properties of $^1$H and $^{14-15}$N

<table>
<thead>
<tr>
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<th>$^1$H</th>
<th>$^{14}$N</th>
<th>$^{15}$N</th>
</tr>
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<tbody>
<tr>
<td>Spin</td>
<td>1/2</td>
<td>1</td>
<td>1/2</td>
</tr>
<tr>
<td>Gyromagnetic Ratio (10⁴ s⁻¹ T⁻¹)</td>
<td>2.675</td>
<td>0.193</td>
<td>0.272</td>
</tr>
<tr>
<td>Natural Abundance (%)</td>
<td>99.99</td>
<td>99.63</td>
<td>0.365</td>
</tr>
<tr>
<td>Relative Sensitivity for Equal Number of Spins</td>
<td>1.00</td>
<td>1.01x10⁻³</td>
<td>1.04x10⁻³</td>
</tr>
</tbody>
</table>

Figure 3.1. $^1$H NMR Spectrum of Ammonium Chloride Aqueous Solution
Several relatively recent techniques have been developed to investigate and characterize systems in fast exchange by NMR (Knüttel and Balaban 1991) but were not investigated for this report. Because of the problem of the chemical exchange rate, $^1$H has been discarded as an NMR probe for the diffusion of ammonia in this study.

### 3.2 Nitrogen NMR

The only remaining possibility is to use nitrogen ($^{14}$N or $^{15}$N) as the NMR probe for the study of ammonia solutions. One drawback of using these nuclei is that they have low gyromagnetic ratios and thus generate a poor NMR signal-to-noise ratio. Different NMR characteristics of the relevant nuclei in this study are listed in Table 3.1. Nitrogen is rather an uncommon nucleus in NMR, and consequently a new radiofrequency coil needed to be designed and constructed for these experiments.

### 3.3 The Radiofrequency Coil

Different considerations must be given to constructing the specific transmitter-receiver radiofrequency coil for detecting nitrogen nuclei. It must meet all the hardware specifications to work at the 7 Tesla spectrometer of the NMR facility at UC Davis. These specifications are

- All the parts must be made of non-magnetic material (e.g., plastic, copper, aluminum).
The components should fit into a 5 in. (12.7 cm) bore.
The circuit must be impedance-matched at 50 Ω.
The working resonances must be 21.7 MHz for $^{14}$N and 30.4 MHz for $^{15}$N.
The coil must be optimized for the nature and geometry of the samples.

In this study, where sensitivity is our major concern, a solenoid shaped coil is the best choice (Hoult and Richards 1976). This geometry has been optimized to improve sensitivity, as suggested by Idziac and Haeberlen (1982). The coil constructed in the laboratory, as shown in Figure 3.3 (a), meets all the specifications cited above. Typical samples are shown in the picture and are 7–10 mL in volume (c). A second coil (b), tuned to work at the proton frequency of 300 MHz, was also constructed to help maintain the settings of the experiment (i.e., homogeneity of the main magnetic field, position of the sample).

Figure 3.3. Image of the Radiofrequency Coil Constructed for Diffusion Experiments at UC Davis: a) $^{14}$-15N Coil, b) 1H Coil, and c) Representative Samples
4.0 Results and Discussion

4.1 Signal-to-Noise Ratio (SNR) Versus Ammonia Concentration

The upper limit of the ammonia concentration is governed by the solubility of the salt. For example, the maximum amount of ammonium sulfate that can be dissolved in water at 20°C is around 40 wt%, which corresponds to a solution of 7.4 M of ammonium ion (see, for example, CRC 1976). This value is about two orders of magnitude higher than the ammonia concentration found in the Hanford waste tanks (Shekarriz et al. 1997).

The lowest concentration limit is constrained by the signal-to-noise ratio (SNR) of the NMR experimental system. The SNR depends upon several factors such as the working field, nucleus, geometry of the coil, and NMR relaxation times. In practice, once the principal parameters are fixed by the experiment, the limits of the ammonia concentration can be determined easily from a single NMR scan. The SNR can be roughly formulated as shown:

$$\text{SNR} \propto \sqrt{N_{\text{acq}} \cdot C} \quad (4.1)$$

where $N_{\text{acq}}$ is the number of acquisitions and $C$ is the concentration of ammonium ion in the solution. On the other hand, we know that quantitative NMR parameters can be extracted if the SNR is greater than or equal to 80. This is about half of what we have for a single scan on a 17 wt% ammonium chloride solution. Each acquisition requires between 20 and 300 seconds, and to extract the diffusion parameter at least four experiments are required. These considerations show that it would be difficult to work with concentrations lower than 50 mM (0.1 wt% of salt). However, this value is about 20 to 50 times the value measured in the Hanford tanks, so our method does allow measurements at realistic concentrations. Because the sensitivity of $^{14}$N and $^{15}$N are close, this discussion holds for both nuclei.

4.2 NMR Relaxation Times

The “life-time” of the NMR signal is characterized by two relaxation times: the spin-lattice relaxation time ($T_1$) and the spin-spin relaxation time ($T_2$). They always satisfy $T_1 \geq T_2$ inequality and are generally about the same order of magnitude in bulk liquids. It is well known that the relaxation process can be greatly enhanced in presence of paramagnetic substances or if the fluid is confined in a porous medium (Callaghan 1991).

4.3 Limitations on Determination of Diffusion Coefficient

Several issues regarding hardware limitations in the PFG experiment merit some discussion. As shown in Equation (2.1), the attenuation of the signal depends upon the amplitude of the gradient, $g$, the duration of these gradients, $\delta$, and the diffusion time, $D$. The strength of the gradients is limited by the power supplied to the coils. In standard imaging systems the practical
gradient ranges from 2 to 40 Gauss/cm. Our system is capable of going up to a maximum gradient of 30 Gauss/cm. The other major limitation is the duration of the PFG experiment or \( \Delta + 2\delta \) (see Figure 2.1). This duration cannot be much longer than the relaxation times. A reasonable timing would be \( \Delta = T_2/2 \) and \( \delta = T_2/4 \) (or \( \Delta + 2\delta = T_2 \)). We can then substitute \( T_2 \) for \( \Delta \) and \( \delta \) in Equation (2.1). Moreover, to obtain an accurate and reproducible value of \( D \), we know that the attenuation of the signal has to be at least 1/2 (Bain 1990). Thus the experimental limitation to determine the diffusion coefficient can be expressed by the following inequality:

\[
2 \geq e^{-\gamma^2 g_{\text{max}}^2 T_2^2 D/16}
\]

For diffusion coefficients of \( 2 \times 10^{-5} \text{ cm}^2/\text{s} \), the relaxation times \( T_1 \) and \( T_2 \) need to be 50 ms or larger for both nuclei \(^{14}\text{N}\) and \(^{15}\text{N}\). This constraint determines the conditions under which we can and cannot measure diffusion, as discussed in the next section.

### 4.4 Self-Diffusion Coefficient of \(^{14}\text{N}\) Ammonia in Aqueous Solutions

The diffusion coefficient has been measured for different samples of ammonia solution. Representative data are shown in Figure 4.1 for a solution of \( 3.3 \text{ M} \) of ammonium chloride with a pH of 5.16. The acquisition time was approximately eight minutes, and the attenuation of the signal is well described by Equation (2.1). The diffusion coefficient is found to be \( 2.15 (\pm 0.01) \times 10^{-5} \text{ cm}^2/\text{s} \).

![Figure 4.1. Diffusion of \(^{14}\text{N}\) in Aqueous Solution of Ammonium Chloride](image)

\[
D = 2.15 (\pm 0.01) \times 10^{-5} \text{ cm}^2/\text{s}
\]
4.4.1 Effect of pH

A series of experiments was performed to study the dependence of relaxation times of $^{14}$N as a function of solution pH. In an aqueous solution of ammonium chloride, the ammonium concentration was maintained at approximately 3.3 M, while the pH was varied by adding small amounts of acid to lower it and ammonium hydroxide to raise it. In Figure 4.2 note that the relaxation times strongly depend on the pH, especially for basic solutions (pH >7). Also note that on this figure the relaxation time plunges below 50 ms for pH >8.5. Thus it would not be possible to measure diffusion of ammonia solutions having a pH higher than 8.5, based on the detectibility arguments presented in the previous sections. This strong pH dependence of the relaxation times is probably due to the nature of the relaxation process, which is quadrupolar and very sensitive to the electronic environment of the molecules (Mason 1987).

Figure 4.3 shows the effect of pH on the diffusion coefficient measured for ammonium chloride solutions. Although some variation is observed, the pH does not dramatically affect the diffusion in such solutions. The small differences (=4%), slightly larger than the experimental error, can be attributed to the slight differences in viscosity of the solutions to which we added acid or ammonium hydroxide.

![Figure 4.2. $^{14}$N Relaxation Times Versus pH in Aqueous Solution of Ammonium Chloride](image-url)
4.4.2 Effect of Ammonia Concentration

Measurements were performed for solutions with different concentrations of ammonia. In this case ammonium sulfate was chosen because of its high solubility in water (maximum of 7.4 M/L). The concentration was varied from 5.3 to 0.18 M, and the results are shown in Figure 4.4. The variation of the diffusion coefficient observed is almost linear with concentration and is directly correlated with the viscosity of the solution (CRC 1976).

4.4.3 Effect of Viscosity

To outline this correlation we performed a measurement on a 3.3 M ammonium chloride solution to which we added some corn syrup to increase the viscosity to match that of the 2.6 M/L ammonium sulfate solution. The two diffusion coefficients are in quantitative agreement: $D_{\text{NH}_4 \text{Cl}} = 1.65 \pm 0.03 \times 10^{-5}$ cm$^2$/s and $D_{\text{SO}_4(\text{NH}_4)_2} = 1.69 \pm 0.04 \times 10^{-5}$ cm$^2$/s. This suggests that viscosity is the principal parameter affecting the diffusion of ammonia in a solution. It is possible that the increase in the viscosity of the sludge due to the presence of a large number of interacting colloidal particulates will result in a significant drop in the diffusion coefficient. However, the viscosity of the interstitial liquid still remains the same as or close to that of water. Currently, there is no consensus on how we should expect the diffusion coefficient to vary as a function of the colloidal particle characteristics and loading. Thus this issue merits further investigation and verification.

Figure 4.3. Ammonium Diffusion Coefficient Versus pH in Ammonium Chloride Aqueous Solution
4.4.4 Apparent Diffusion Coefficient of Ammonia in Porous Media

One set of experiments has been conducted in a porous medium to show feasibility of diffusion coefficient measurements using the current PFG NMR technique. The porous medium was composed of randomly packed PMMA beads (200 μm) saturated with a 3.3 M ammonium chloride solution (pH 4.67). As is seen in Figure 4.5, the attenuation of the NMR signal of the bulk solution is greater than that for the solution in the porous media. This will lead to an approximately 30% smaller diffusion coefficient for the fluid in the porous medium. In fact, the diffusion time (D) is long enough to let a large portion of the molecules encounter the surface of the particles within the pores of porous media, causing an “observed” restricted or hindered diffusion. The corresponding diffusion coefficient is called the NMR apparent diffusion coefficient and is a function of D in the NMR experiment. This particular property of the PFG measurement has been used to characterize the structural properties of porous systems (Callaghan 1991; Chui et al. 1995).

4.4.5 15N Labeled Ammonia Solutions

The 15N isotope has become an increasingly accessible nucleus for NMR related work. Enriched salts such as ammonium chloride or ammonium sulfate and even ammonium hydroxide solutions have become commercially available. Although 15N labeled products are relatively expensive (around $100 per gram or mL), the 15N nucleus has a spin 1/2 and presents several advantages from the spectroscopy point of view (Mason 1987). This nucleus has a higher
gyromagnetic ratio, leading to a higher sensitivity to diffusion measurements (see Table 3.1 and Equation 2.1). Also, relaxation is caused by dipolar interactions instead of quadrupolar for $^{14}$N, leading to longer relaxation times. Table 4.1 shows the relaxation times of $^{15}$N labeled ammonium chloride solutions with different pH.

Because the relaxation times are less sensitive to the pH of the solution, we should be able to measure the diffusion coefficient for solutions up to pH 9.5 (relaxation times over 50 ms). The diffusion coefficient measured for the solution of 3.3 M/L ammonium concentration, pH 5.3, is of $2.20 \pm 0.01 \times 10^{-5}$ cm$^2$/s, which is in excellent agreement with the $2.15 \pm 0.01 \times 10^{-5}$ cm$^2$/s value measured for standard ammonium chloride (see Figure 4.1).

**Table 4.1. PH and Relaxation Times for $^{15}$N Ammonium Chloride Solutions**

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<th>pH</th>
<th>$T_1$ (ms)</th>
<th>$T_2$ (ms)</th>
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<tr>
<td>5.3</td>
<td>39</td>
<td>500</td>
</tr>
<tr>
<td>9.1</td>
<td>37</td>
<td>54</td>
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5.0 Conclusions and Recommendations

Diffusion coefficients of ammonium aqueous solutions have been successfully determined by PFG NMR methods in bulk solutions and for confined solutions within porous media. Feasibility for relatively low concentrations (50 mM/L) and high pH (9.5) have been demonstrated. The diffusion coefficients measured are close to those found in pure water (≈ 2 x 10^{-5} cm^2/s). There is no appreciable dependence of the diffusion coefficient upon concentration or pH within the range of conditions studied. However, as expected, it has been found that the diffusion coefficient is correlated to the viscosity of the solution and the porosity of the porous medium.

These attractive results will lead to further characterization of the rate of diffusion in model samples that closely resemble the conditions in the underground waste storage tanks at Hanford. Future experiments are recommended in which two different levels of complexity are added: 1) varying the viscosity of the medium by adding colloidal particulates and 2) imposing a temperature gradient in the medium to examine the effect of thermal gradient on the ammonia diffusivity. Both these variables will create a more realistic model of the Hanford tank waste conditions, especially in the nonconvective layer.

Other things to look for are 1) diffusion of NH_4^+ Cl^- in high concentration of NaCl solution and 2) diffusion of NH_4^+ Cl^- in high concentration of NaOH solution, if possible. It is expected that these two salts will have different effects on the solution viscosity and, as a result, might give rise to different values for diffusion coefficients.
6.0 References


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