Development of a Flow Injection System with Two Analytical Paths for Ammonium Determination in Soil Extracts by Conductometry

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A flow injection system with two analytical paths for ammonium determination in soil extracts by conductometry is described. The ammonia generated on line in an alkaline medium is permeated through a hydrophobic membrane (TPFE), collected by a deionized water stream and directed toward the conductivity flow cell. The flow network was designed with two analytical paths in order to increase the analytical throughput. The procedure was used to determine ammonium in 2 mol L$^{-1}$ KCl soil extracts and allowed 100 determinations per hour with a relative standard deviation (r.s.d) of about 2% and an average recovery of 99.9% for a typical sample with concentration of 2.1 mg NH$_4^+$ L$^{-1}$.

Keywords: ammonium determination, flow analysis, conductometric detection, analysis of soil extracts

Introduction

Available ammonium ions in soils is an important parameter for consideration in plant nutrition and soil fertility studies. Usually, a 2 mol L$^{-1}$ KCl solution is used to extract ammonium ions in soil samples$^1$. This high salt concentration may cause difficulties in employing spectrophotometric determinations, mainly those based on the Nessler$^2$ or the Berthelot methods$^3$. In addition, the reagents used in both methods contain mercury and phenol, respectively, and should be avoided. Ammonium determination can also be performed using titrimetry but this procedure presents low throughput since a distillation step is necessary.

Studies of soil fertility usually require the analysis of the ammonium ion in a large number of samples. Therefore, the availability of automated analytical methods presenting high throughput is very important. Among the automated analytical procedures, flow injection could be selected, considering its versatility, its inherent high throughput, and its precision$^4,5,6$.

Soil extracts can be considered a complex matrix, presenting some analytical difficulties when analyzed by spectrophotometric methods, *i.e.* for ammonium determination by the chemical reaction occurring in alkaline medium where a separation step should be incorporated$^2,3$. The indirect determination by atomic absorption spectrometry presented high sensitivity, however the high concentration...
of KCl (2 mol L\(^{-1}\)) could cause interference as indicated elsewhere\(^6\).

Flow injection modules with the on-line ammonia generation to separate it from the sample bulk has been proposed earlier. This was done by diffusing the gas through an air gap that separates the sample carrier and the acceptor solution streams\(^7\), or by permeating the gas through a hydrophobic membrane\(^8,9\). Using these procedures ammonia is collected by an aqueous stream which flows parallel with the sample carrier stream inside the separation chamber. In the acceptor stream, ammonia is again converted to ammonium ions that are detected by conductometry\(^8\) or by spectrophotometry\(^7,9\). Ammonia separation with a hydrophobic membrane is more commonly used than that with an air gap and has been employed for conductometric analysis of plant digests with high ammonium concentrations\(^8\).

The advantages of ammonium determination by conductometry, including the instrumentation simplicity and the requirement of a hydroxide solution as the only reagent, make it an attractive alternative to replace methods using mercury or phenol\(^2,3\). In this work, a flow injection procedure employing conductometry was developed to determine ammonium in soil extracts. The content of ammonium in some types of soils is generally lower than 1 mg L\(^{-1}\). Therefore, to attain the appropriate limit of detection, it is necessary to use a large volume of sample for analysis. Ammonium permeation through a hydrophobic membrane is a slow process, resulting in an analytical method with a low sampling rate in comparison with other flow procedures. This disadvantage becomes more drastic for samples with low ammonium content and when a large number of samples must be analyzed. Considering this fact, the flow network was designed with two analytical paths in order to increase the analytical throughput by sequential sampling and detection.

**Experimental**

**Reagents**

Chemicals of analytical-reagent grade and freshly distilled and deionized water were used throughout. All reagents, standard solutions, and samples were stored in clean polyethylene bottles.

The stock ammonium solution 1000 mg NH\(_4^+\) L\(^{-1}\) was prepared by dissolving 3.666 g of ammonium sulfate in water and diluting to 1000 mL.

The soil extracting solution, 2 mol L\(^{-1}\) KCl, was prepared by dissolving 149.1 g of the salt in water and diluting to 1000 mL.

The alkaline solution, 1.5 mol L\(^{-1}\) NaOH was prepared by dissolving 54.9 g NaOH in water and diluting to 1000 mL.

Working standard solutions of 0.00, 0.25, 0.50, 1.00, 2.00, 3.00 and 4.00 mg NH\(_4^+\) L\(^{-1}\) were prepared by dilution of the stock solution with the 2 mol L\(^{-1}\) KCl solution.

Soil extracts in 2 mol L\(^{-1}\) KCl were prepared as described elsewhere\(^1\) and were refrigerated until analysis.

**Apparatus**

The flow set up consisted of a B330 Micronal conductometer, a REC 61 Radiometer strip chart recorder, a MP13 Ismatec peristaltic pump furnished with pumping Tygon tubes, a proportional injector made in Perspex with two switching sections\(^9\), two electric switches with double contacts being one normally open and the other closed, helicoidal reaction coils and transmission lines (0.8 mm i.d.) of polyethylene tubing, two gas permeation units, and two stainless steel conductivity cells.

The gas permeation units were constructed from two Perspex blocks (20 x 5 x 2 cm). A silicone rubber strip (20 x 2.5 x 0.005 cm) was glued to the face of each Perspex block. A channel (8 x 0.02 cm) was made in the rubber strip as indicated in Fig. 1. A PTFE tape was placed on one of the rubber strips covering the channel. The unit was assembled by attaching the two Perspex blocks together with screws. The PTFE tape was sandwiched between them, so the channel on both sides forms the flow pathway in the permeation units where the donor and acceptor streams flow. The rubber strips were used to avoid fluid leakage and to provide a soft contact with PTFE tape to increase life time of the unit.

The conductivity flow cells were constructed from two Perspex cylinders of 40 mm diameter and 30 mm depth. In each cylinder, an axial hole of 15 mm diameter was drilled, and a stainless steel cylinder was inserted in this hole as indicated in Fig. 2. The metal cylinder was sealed with epoxy glue to avoid fluid leakage. A disk of silicone rubber (25 mm diameter) with a slit (10 x 2 mm) was placed on the metallic surface (detail in Fig. 2). The conductivity flow cell was assembled by attaching the two Perspex cylinders together with screws, sandwiching the rubber disk between them. The slit on the rubber disk provides the electric

**Figure 1.** Diagram of the permeation chamber. The detail shows the inside of the permeation chamber e = silicone rubber strip, 0.5 mm thickness and a PTFE membrane. D indicates the donor stream and A the acceptor.
contact between both electrodes during flow of solution, allowing the detection of the solution conductivity.

Flow set up and experimental variables

The flow set up is depicted in Fig. 3. In the position shown, the solution inside loop L₁ is displaced by the carrier stream (Cs) toward the gas permeation unit (D₁). At the confluence point (X), sodium hydroxide solution (R) is added, and the ammonium ions are converted to ammonia (NH₃) while passing through the reaction coil (B₁). While the solutions flow through the permeation unit toward waste (W), part of the generated ammonia permeates through the PTFE membrane and is collected into the acceptor stream (Rc). Ammonia is converted to ammonium ions in the acceptor stream solution while carried to the conductivity flow cell (C₁). This device is electrically connected to the conductometer (DET) by means of the electric switches ch₁ and ch₂ for detection. Variation in conductivity is converted to a variation in electric potential, which is recorded by the strip chart recorder (Reg) as a function of time. While the described steps are occurring, the sample S₂ flows through the sampling loop L₂ toward waste.

Changing the injector to the other resting position by displacing the sliding bar, the sampling loop L₁ is loaded with another sample (S₁) and the sampling loop L₂ is inserted into the analytical path composed of reaction coil (B₂), gas permeation unit (D₂) and conductivity flow cell (C₂). A mechanism attached to the sliding bar of the injector (dashed line) performs the switching of the electric switches ch₁ and ch₂ to connect the conductometer to the conductivity flow cell C₂ instead of C₁. Sample processing and signal measurement occur as previously described.

To establish the optimal operational conditions, experiments were performed by varying the sampling loop length and the carrier and acceptor streams flow rates.
Results and Discussion

The analytical signal obtained by flow systems with phase separator devices like gas permeation membranes must be optimized by adjusting the carrier and acceptor stream flow rates. Experimental results used to determine the optimal conditions with respect to residence time and dispersion are shown in Figs. 4, 5 and 6.

An increase in the analytical signal greater than 100% was observed when a sampling loop of 50 cm was used and the sample carrier flow rate changed from 6.0 to 1.6 mL min⁻¹ as shown in Fig. 4. When a 100 cm sampling loop was employed using the same flow rates, the observed increase in signal was about 40% as shown in Fig. 6. In both cases the acceptor flow rate was maintained at 1.6 mL min⁻¹.

By increasing the sample carrier flow rate, less ammonia permeates the membrane due to the decrease in average residence time. On the other hand, when the sampling loop length was doubled, two effects must be considered: the sample zone broadening and dispersion in the carrier solution. Broadening the sample zone requires a long time-interval for transport through the permeation unit. In addition, decreasing the sample dispersion increases the ammonia partial pressure of the carrier solution due to the enhancement of the permeation rate. These two parameters favorably affect the ammonium permeation through the membrane. The permeation conditions were not a problem for determination of ammonium permeation through the membrane. The permeation conditions were not a problem for determination of ammonium in plant digests due to the high analyte concentrations of the samples, however, in soil extracts this parameter must be considered.

The slopes of curve I in Fig. 6 and curve II in Fig. 5, are approximately equal (ΔV/ΔC ≅ 12). These two curves were obtained using the same parameters. Similar behavior is demonstrated by curves II in Figs. 4 and 6. Considering these results, flow rates of 3 mL min⁻¹ and 2.5 mL min⁻¹ for the sample carrier and ammonia acceptor, respectively, were established for further experiments.

Figure 4. Effect of the sample carrier flow rate. Curves I, II and III obtained with sample flow rates of 1.6, 3.0 and 6.0 mL min⁻¹, respectively and sampling loop of 50 cm long. Other parameters as in Fig. 3.

Figure 5. Effect of the sample carrier flow rate. Curves I, II and III obtained with sample flow rates of 1.6, 3.0 and 6.0 mL/min, respectively and sampling loop of 100 cm long. Other parameters as in Fig. 3.

Figure 6. Effect of the sampling loop length. Curves I and II obtained with sampling loop of 100 and 50 cm long, respectively. Other parameters as in Fig. 3.
Peaks recorded in Fig. 7 demonstrate how the analytical throughput was improved using the proposed flow network. The recorder tracing a was obtained by operating the flow system without switching ch1 and ch2 from C1 to C2, thus a FIA peak with a long tail is obtained due to the sample dispersion into the carrier stream. The peak profile reflects the intrinsic characteristic of the flow system dimensions and the chemical solutions employed. Peak b was obtained by switching ch1 and ch2 from C1 to C2 immediately after attaining the maximum peak height of the analytical signal. As a result, the signal returned abruptly to the baseline. The time interval required to obtain peak profiles a and b, were about 60 and 20 s, respectively. This time difference could be decreased by increasing the overall flow rate, but in this work the flow rates were optimized for maximum sensitivity as shown in Figs. 4 and 5. The simultaneous movement of the injector and switches between the conductivity flow cells defined the analytical throughput. In routine analysis an average of 120 determinations per hour could be achieved.

The set of recorder tracings at c in Fig. 7 shows the sequence of signals obtained by using both analytical channels. The proposed flow network comprises two analytical paths, and as their dimensions are not identical, this set of peaks reveals differences in the baseline and in sensitivity. The gas permeation rate through PTFE membrane is expected to be proportional to the membrane surface interfacing the donor and acceptor fluids. In this instance, some difference in the channel dimensions of the permeation unit (Fig. 1) could generate the difference in sensitivity observed. Also, some difference in the conductivity area could contribute to the effect on the baseline. The difficulty of constructing two identical analytical channels to achieve the same baseline and signal sensitivity made it imperative to calibrate both channels independently. The routine graph

![Figure 7](image-url) Effect of switching the conductivity flow cell. From right to left, a peak profile obtained from one conductivity flow cell, b peak profile obtained by switching from C1 to C2 (Fig. 3) at peak maximum followed by a set of recorder tracing obtained running the flow set up with a 3.0 mg NH4+ L⁻¹ standard solution. Other parameters as in Fig. 3.

![Figure 8](image-url) Recording related to a set of standards and samples. From right to left peaks in duplicate obtained with standard solutions of 0.5, 1.0, 2.0 and 3.0 mg NH4+ L⁻¹ for each analytical path, followed by those of 12 samples in duplicate and the standards again. Samples signed by odd numbers were analyzed by the analytical path comprised by reaction coil B1, permeation unit D1 and conductivity flow cell C1.

<table>
<thead>
<tr>
<th>Soilextract</th>
<th>Concentration (mg/L)*</th>
<th>Recovery (%)</th>
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<tbody>
<tr>
<td>1</td>
<td>0.94</td>
<td>96</td>
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<tr>
<td>2</td>
<td>1.43</td>
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<td>98</td>
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*Average concentration (n = 3). Recovery 98.9 ± 2.9.

Table 1. Results of the recovery test after addition of 0.5 mg NH4+ L⁻¹ to different soil extracts.
showing a sequence of standard and sample solutions for the two independent analytical paths is shown in Fig. 8. Soil extract results are shown in Fig. 8 where it is possible to identify, without difficulty, the peaks of each sample. By observing the peaks corresponding to the two sets of standard solutions (right and left in Fig. 8), one can conclude that the system is very stable. The reliability was tested by standard addition, and results are characterized by recoveries of 96-105% (Table 1). Another feature was a relative standard deviation better than 2% calculated from 8 sequential measurements obtained for a sample of 1.04 mg NH₄⁺ L⁻¹.

Conclusions

The flow system with two analytical channels for ammonium determination using sequential detection from two conductivity flow cells, allows a higher sample throughput than the single channel configuration. The conductometric method is robust for ammonium determination in high saline samples (2 mol L⁻¹ KCl soil extracts) even at low concentrations. This is a clean method and could be employed instead of those based on mercuric or phenolic reagents.

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