



Total Nitrogen and Crude Protein in Feed by Segmented Flow Analysis (SFA)

(Cartridge Part #A002726)

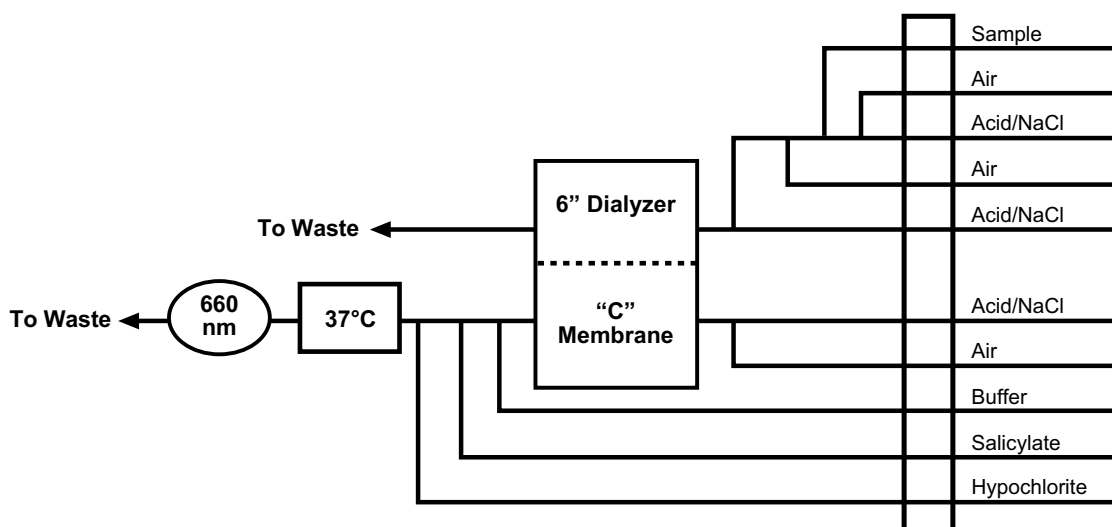
1.0 Scope and Application

- 1.1 This method is used for the determination of total nitrogen and crude protein in animal feed.
- 1.2 The Method Detection Limit (MDL) of this method is 1.0 mg/L nitrogen (N). The applicable range of the method is 3.0–150 mg/L nitrogen or approximately 2–70% protein in a 0.5–1.0 g sample. The range may be extended to analyze higher concentrations by sample dilution.

2.0 Summary of Method

- 2.1 Samples are digested prior to analysis by heating in the presence of the appropriate digestion solution (Reference 15.2). Free ammonia and organic nitrogen compounds are converted to ammonium ion. The samples are then automatically diluted, dialyzed, and finally reacted with salicylate and hypochlorite in a buffered alkaline solution in the presence of sodium nitroferricyanide. The salicylic acid analog of indophenol blue is formed, and the absorbance is measured at 660 nm (Reference 15.2).
- 2.2 The quality of the analysis is assured through reproducible calibration and testing of the Segmented Flow Analysis (SFA).

- 2.3 A general flow diagram of the SFA system is shown below (see Section 17.0 for a detailed flow diagram).



3.0 Definitions

Definitions for terms used in this method are provided in Section 16.0, "Glossary of Definitions and Purposes."

4.0 Interferences

- 4.1 On-line dialysis eliminates turbidity and background color interferences.

5.0 Safety

- 5.1 The toxicity or carcinogenicity of each compound or reagent used in this method has not been fully established. Each chemical should be treated as a potential health hazard. Exposure to these chemicals should be reduced to the lowest possible level.
- 5.2 For reference purposes, a file of Material Safety Data Sheets (MSDS) for each chemical used in this method should be available to all personnel involved in this chemical analysis. The preparation of a formal safety plan is also advisable.
- 5.3 The following chemicals used in this method may be highly toxic or hazardous and should be handled with extreme caution at all times. Consult the appropriate MSDS before handling.
- 5.3.1 Ammonium Phosphate Dibasic, $(\text{NH}_4)_2\text{HPO}_4$ (FW 132.06)
- 5.3.2 Chloroform, CHCl_3 (FW 119.38)

- 5.3.3 Potassium Sodium Tartrate Tetrahydrate, $C_4H_4O_6NaK \cdot 4H_2O$ (FW 282.23)
- 5.3.4 Salicylic Acid Sodium Salt, $C_7H_5O_3Na$ (FW 160.11)
- 5.3.5 Sodium Hydroxide, NaOH (FW 40.00)
- 5.3.6 Sodium Hypochlorite, 5.25% available chlorine (household bleach), NaOCl (FW 74.44)
- 5.3.7 Sodium Nitroferrocyanide Dihydrate, $Na_2Fe(CN)_5NO \cdot 2H_2O$ (FW 297.95)
- 5.3.8 Sodium Phosphate Dibasic Heptahydrate, $Na_2HPO_4 \cdot 7H_2O$ (FW 268.07)
- 5.3.9 Sulfuric Acid, concentrated, H_2SO_4 (FW 98.08)
- 5.4 Unknown samples may be potentially hazardous and should be handled with extreme caution at all times.
- 5.5 Proper personal protective equipment (PPE) should be used when handling or working in the presence of chemicals.
- 5.6 This method does not address all safety issues associated with its use. The laboratory is responsible for maintaining a safe work environment and a current awareness file of OSHA regulations regarding the safe handling of the chemicals specified in this method.

6.0 Apparatus, Equipment, and Supplies

- 6.1 Segmented Flow Analysis (SFA) System (OI Analytical Flow Solution® IV) consisting of the following:
 - 6.1.1 Model 502 Multichannel Peristaltic Pump
 - 6.1.2 Random Access (RA) Autosampler
 - 6.1.3 Expanded Range (ER) Photometric Detector with 5-mm path length flowcell and 660-nm optical filter
 - 6.1.4 Data Acquisition System (PC or Notebook PC) with WinFLOW™ software
 - 6.1.5 Total Nitrogen and Crude Protein in Feed Cartridge (Part #A002726)
- 6.2 Sampling equipment—Sample bottle, amber glass, with polytetrafluoroethylene (PTFE)-lined cap. Clean by washing with detergent and water, rinsing with two aliquots of reagent water, and drying by baking at 110°–150°C for a minimum of one hour.
- 6.3 Standard laboratory equipment including volumetric flasks, pipettes, syringes, etc. should all be cleaned, rinsed, and dried per bottle cleaning procedure in Section 6.2.

7.0 Reagents and Calibrants

7.1 Raw Materials

- 7.1.1 Ammonium Phosphate Dibasic, $(\text{NH}_4)_2\text{HPO}_4$ (FW 132.06)
- 7.1.2 Brij[®]-35, 30% w/v (Part #A21-0110-33)
- 7.1.3 Chloroform, CHCl_3 (FW 119.38)
- 7.1.4 Deionized Water (ASTM Type I or II)
- 7.1.5 Potassium Sodium Tartrate Tetrahydrate, $\text{C}_4\text{H}_4\text{O}_6\text{NaK} \cdot 4\text{H}_2\text{O}$ (FW 282.23)
- 7.1.6 Salicylic Acid Sodium Salt, $\text{C}_7\text{H}_5\text{O}_3\text{Na}$ (FW 160.11)
- 7.1.7 Sodium Chloride, NaCl (FW 58.44)
- 7.1.8 Sodium Hydroxide, NaOH (FW 40.00)
- 7.1.9 Sodium Hypochlorite, 5.25% available chlorine (household bleach), NaOCl (FW 74.44)
- 7.1.10 Sodium Nitroferricyanide Dihydrate, $\text{Na}_2\text{Fe}(\text{CN})_5\text{NO} \cdot 2\text{H}_2\text{O}$ (FW 297.95)
- 7.1.11 Sodium Phosphate Dibasic Heptahydrate, $\text{Na}_2\text{HPO}_4 \cdot 7\text{H}_2\text{O}$ (FW 268.07)
- 7.1.12 Sulfuric Acid, concentrated, H_2SO_4 (FW 98.08)