



Designation: D 5012 – 01

Standard Guide for Preparation of Materials Used for the Collection and Preservation of Atmospheric Wet Deposition¹

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1. Scope

1.1 This guide presents recommendations for the cleaning of plastic or glass materials used for collection of atmospheric wet deposition (AWD). This guide also presents recommendations for the preservation of samples collected for chemical analysis.

1.2 The materials used to collect AWD for the analysis of its inorganic constituents and trace elements should be plastic. High density polyethylene (HDPE) is most widely used and is acceptable for most samples including samples for the determination of the anions of acetic, citric, and formic acids. Borosilicate glass is a collection alternative for the determination of the anions from acetic, citric, and formic acid; it is recommended for samples for the determination of other organic compounds.

1.3 *This standard does not purport to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this standard to establish appropriate safety and health practices and determine the applicability of regulatory limitations prior to use.*

2. Referenced Documents

- 2.1 *ASTM Standards:*²
- D 883 Terminology Relating to Plastics
 - D 1125 Test Methods for Electrical Conductivity and Resistivity of Water
 - D 1129 Terminology Relating to Water
 - D 1193 Specification for Reagent Water
 - D 1356 Terminology Relating to Sampling and Analysis of Atmospheres
 - D 1695 Terminology of Cellulose and Cellulose Derivatives
 - D 2914 Test Methods for Sulfur Dioxide Content of the Atmosphere (West-Gaeke Method)
 - D 4453 Practice for Handling of Ultra-Pure Water Samples

¹ This guide is under the jurisdiction of ASTM Committee D22 on Air Quality and is the direct responsibility of Subcommittee D22.03 on Ambient Atmospheres and Source Emissions.

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² For referenced ASTM standards, visit the ASTM website, www.astm.org, or contact ASTM Customer Service at service@astm.org. For *Annual Book of ASTM Standards* volume information, refer to the standard's Document Summary page on the ASTM website.

3. Terminology

3.1 Definitions:

3.1.1 For definitions of terms used in this guide, refer to Terminology D 1129.

3.1.2 For definition of **plastic** refer to Terminology D 1695 and Terminology D 883.

3.1.3 For definition of AWD (**precipitation, meteorological**) refer to Terminology D 1356.

4. Significance and Use

4.1 Some chemical constituents of AWD are not stable and must be preserved before chemical analysis. Without sample preservation, it is possible that analytes can be lost through decomposition or sorption to the storage bottles.

4.2 Contamination of AWD samples can occur during both sample preservation and sample storage. Proper selection and cleaning of sampling containers are required to reduce the possibility of contamination of AWD samples.

4.3 The natural sponge and talc-free plastic gloves used in the following procedures should be recognized as potential sources of contamination. Individual experience should be used to select products that minimize contamination.

5. Apparatus

5.1 Instruments shall be selected in accordance with an applicable test method given in Test Methods D 1125.

5.2 The conductivity cell shall be pipet or dip type with a cell constant (K) of 0.1 .

6. Reagents and Materials

6.1 *Purity of Reagents*—Reagent grade acids and other chemicals shall be used to reduce the risk of contaminating the AWD samples. Unless otherwise indicated, it is intended that all reagents conform to the specifications of the Committee on Analytical Reagents of the American Chemical Society where such specifications are available.³

³ *Reagent Chemicals, American Chemical Society Specifications*, American Chemical Society, Washington, DC. For suggestions on the testing of reagents not listed by the American Chemical Society, see *Analar Standards for Laboratory Chemicals*, BDH Ltd., Poole, Dorset, U.K., and the *United States Pharmacopeia and National Formulary*, U.S. Pharmaceutical Convention, Inc. (USPC), Rockville, MD.

TABLE 1 Preservation of AWD Samples Collected for Inorganic Cation and Anion Determinations

| Preservation Technique | Species Determined | Remarks | Reference |
|-------------------------|--|--|-----------|
| No preservation | All inorganic cations and anions | Rapid analysis is required after collection because ion concentrations may change in samples. Ammonium, nitrate, and ortho-phosphate concentrations may be reduced in samples that are biologically active. Cation and trace metal concentrations may be reduced by sorption onto container surfaces. | (10-14) |
| Refrigerate 4°C | All inorganic cations and anions | Chilling may reduce the loss of ammonium, nitrate, and ortho-phosphate in samples that are biologically active. Samples must be allowed to come to ambient temperature (23–27°C) before performing pH and specific conductance determinations. Specific conductance and pH determinations should be performed on-site as soon as possible after sample collection. | (15-17) |
| HNO ₃ pH ≤ 2 | Ca ²⁺ , Mg ²⁺ , Na ⁺ , K ⁺ | Samples must first be filtered, or acid addition may dissolve particles in the AWD samples (see 7.4.1). Acid addition will interfere with anion determination, so a separate aliquot will be needed for other ion determinations. | (11,12) |
| HCl, 5 mL/L | All dissolved Hg and total Hg | Samples for dissolved Hg are filtered through 0.45 µm capsule filter. Samples collected directly into specially cleaned, pretested, fluoropolymer bottle. | (19,20) |
| Filtration | All inorganic cations and anions | pH and specific conductance determinations may be affected by filtration. Care must be taken to minimize the possibility of sample contamination during filtration. | (18) |

6.2 Nitric acid, ultra pure.

6.3 Nitric acid (1+9)—Dilute 1 volume concentrated nitric acid with 9 volumes of water.

6.4 Chloroform, high performance liquid chromatography reagent.

6.5 Hydrochloric acid, pretested for total and methyl mercury.

6.6 *Purity of Water*—Unless otherwise indicated, references to water shall be understood to mean reagent water as defined by Type I of Specification D 1193. AWD samples collected for organic analysis may require Type II (distilled) reagent water.

7. Sample Preservation

7.1 Interaction between the sample and the atmosphere must be minimized. The sample container should be sealed as soon as possible after collection or sub-sampling. AWD samples can be easily contaminated because of the low concentration levels of their analytes. Trace metals, and possibly other ions in AWD samples, can be lost through sorption with the bottle or container in which they are stored. AWD samples may contain biologically active microorganisms which could affect the concentration of many analytes. Immediate analysis of AWD samples is best and chemical preservation should be used only when the preservation is shown not to interfere with the analysis being performed. Samples collected for total and

methyl mercury determinations are to be collected directly into specially cleaned, pretested, fluoropolymer bottle(s) (19 and 20).⁴

7.2 *Samples Collected for Inorganic Cation and Anion Determinations:*

7.2.1 Samples collected for pH, specific conductance, calcium, magnesium, potassium, sodium, chloride, fluoride, and sulfate analysis are often only placed in pre-cleaned plastic containers (see Section 8) before analysis. If there is a delay between time of collection and time of analysis, a preservation technique may eliminate or moderate chemical and biological changes in the AWD samples. Table 1 summarizes the recommended sample preservation techniques for AWD samples.

7.3 *Samples Collected for Organic Acids:*

7.3.1 Samples collected for the analysis of acetate, citrate, formate and other low molecular weight organic acid anions (C₁-C₁₂) should be preserved within minutes after collection. Organic acids have been determined in AWD samples collected from locations around the world (1-7). These compounds (in particular formic and acetic acids) can constitute from a small fraction to mostly all of the free acidity in AWD samples.

⁴ The boldface numbers in parentheses refer to the list of references at the end of this standard.

TABLE 2 Preservation of AWD Samples Collected for Organic Acid Determinations

| Preservation Technique | Species Determined | Remarks | Reference |
|------------------------|--|--|-----------|
| Chloroform | Acetate, citrate, formate, C ₁ -C ₅ | Analysis is usually by ion-exclusion chromatography because chloroform may interfere with capillary GC-MS analysis. This preservative is often coupled with sample storage at 4°C. When chloroform treatment is combined with storage in the dark at 4°C, samples are reportedly stable for 60 days. Final chloroform concentration in the AWD sample must be 0.2 % by weight. | (3,4) |
| Freeze at – 20°C | Short-chain dicarboxylic acids C ₂ -C ₁₂ | Analysis is by capillary gas chromatography or GC-MS. Sample must be completely in the liquid state before sample is prepared for analysis. Maximum storage time has not been reported. | (6) |
| Chill at 4°C | Low molecular weight acids | This is a minimum procedure for organic acid preservation. This preservation technique will not preserve acetate and formate longer than 3 days. Degradation may occur in less than 3 days. | (7) |