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Standard Practice for Determination of Volatiles in Polymers by Static Headspace Gas Chromatography¹

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1. Scope*

1.1 Headspace gas chromatography (GC) involves the determination of volatile components in a polymer solution by gas chromatography of a vapor phase in thermal equilibrium with the sample matrix. Volatiles in finely ground insoluble polymers can also be determined with and without an extracting solvent.

1.2 This practice provides two procedures:

1.2.1 *Procedure A*—Automatic headspace analysis.

1.2.2 *Procedure B*—Manual injection headspace analysis.

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.* Specific precautionary statements are given in Section 6.

NOTE 1—There is no known ISO equivalent to this standard.

2. Referenced Documents

2.1 *ASTM Standards*:²

D4322 *Test Method for Residual Acrylonitrile Monomer Styrene-Acrylonitrile Copolymers and Nitrile Rubber by Headspace Gas Chromatography* (Withdrawn 2010)³

E260 *Practice for Packed Column Gas Chromatography*

E355 *Practice for Gas Chromatography Terms and Relationships*

E594 *Practice for Testing Flame Ionization Detectors Used in Gas or Supercritical Fluid Chromatography*

¹ This practice is under the jurisdiction of ASTM Committee D20 on Plastics and is the direct responsibility of Subcommittee D20.70 on Analytical Methods.

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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 The last approved version of this historical standard is referenced on www.astm.org.

3. Significance and Use

3.1 For various reasons, one may want to measure the level of unreacted or residual monomer, water, or other volatile components in a polymer sample.

3.2 Volatiles of interest are often at trace concentrations. Headspace analysis is suited for determination of these trace components which often cannot be determined by conventional gas chromatography because of sample decomposition or interferences.

3.3 For polymer analysis, sample treatment for headspace analysis is simpler than conventional gas chromatography, where precipitation steps may be required to prevent polymer contamination of the chromatographic column.

3.4 This headspace practice will be able to determine qualitatively any component with sufficient vapor pressure. It is capable of yielding semiquantitative results and can be used for relative comparisons between samples.

4. Apparatus

4.1 *Gas Chromatograph*, equipped with an appropriate detector and backflush valve.

4.1.1 For Procedure A, an *Automated Headspace Sampler*,^{4,5} including backflush capability, thermostated sample tray, and associated accessories fulfill these requirements while providing for automatic sequential sampling of headspace vapors.

4.1.2 Procedure B requires the following additional equipment:

4.1.2.1 *Constant-Temperature Bath*, capable of maintaining $90 \pm 1^\circ\text{C}$.

4.1.2.2 *Gas-Tight Gas Chromatographic Syringes*, which can be heated to 90°C for sampling and injection.

4.1.2.3 *Valve*, 6-port for backflush assembly.

NOTE 2—Appropriate detectors could include the following:

(a) Flame ionization (FID) for general organic volatiles,

⁴ The sole source of supply of the apparatus known to the committee at this time is Perkin-Elmer Corp., Main Ave., Norwalk, CT 06856.

⁵ If you are aware of alternative suppliers, please provide this information to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee,¹ which you may attend.

*A Summary of Changes section appears at the end of this standard

- (b) Electron capture (EC) for halogenated species,
- (c) Nitrogen-phosphorous (NPD) for acrylonitrile,
- (d) Thermal conductivity (TC) for water, and
- (e) Hall electroconductivity or photoionization (PID) for vinyl chloride.

4.2 Chromatographic Columns:

4.2.1 *Packed Column Analysis*—Column packings^{5,6} found to be useful for residual monomers (for example, acrylonitrile, vinyl chloride) include Chromosorb or Porapak porous polymer packings, 0.4 % Carbowax 1500 liquid phase on Carbopak C support, or 0.19 % picric acid on Carbopak C support. Other columns can be used after it is determined that they give the required resolution and accuracy for the component of interest.

4.2.2 *Capillary Column Analysis*—Choose a column capable of achieving the required resolution for the components of interest. For example, for acrylonitrile monomer, the column of choice is a Quadrex 007–2, 25m × 0.32–mm internal diameter fused silica.^{5,7}

4.3 *Integrator, recorder (5–mV full scale), computing integrator or appropriate computer data station and software* capable of measuring and storing peak area data.

4.4 *Headspace Vials, septa, ring closures, and vial sealer.*

4.5 *Analytical Balance, capable of weighing to ±0.0001 g.*

4.6 *Soap Film Flowmeter and Stopwatch, or other means of measuring gas flow rates.*

4.7 *Pressure Regulators* for all required gas cylinders.

4.8 *Filter-Dryer Assemblies* for each required gas cylinder.

5. Reagents and Materials

5.1 *Solvent, Reagent Grade, or Best Available*—The solvent should be free of interferences at the retention time of volatile compounds determined by this technique. Typical polymer solvents include water, dimethylformamide (DMF), dimethylacetamide (DMAC), propylene carbonate, and *o*-dichlorobenzene. These solvents are readily backflushed from most chromatographic columns.

NOTE 3—Volatile contaminants can often be removed by sparging with an inert gas for 24 to 48 h.

5.2 *Nitrogen, oxygen-free.*

NOTE 4—Helium may also be used as the carrier gas and argon/methane is the preferred carrier gas for use with electron capture detectors.

5.3 *Hydrogen, prepurified, for use with flame ionization detectors.*

5.4 *Air, breathing or water pumped, for use with flame ionization or nitrogen-phosphorus detectors.*

5.5 *Standards, best available, for volatile compounds to be analyzed using this technique.*

⁶ The sole source of supply of the apparatus known to the committee at this time is Supelco, Inc., P.O. Box 628, 146 S. Water St., Bellefonte, PA 16823.

⁷ The sole source of supply of the apparatus known to the committee at this time is Quadrex Corp., P.O. Box 3881, Woodbridge, CT 06525.

6. Safety Precautions

6.1 Volatiles such as vinyl chloride and acrylonitrile should not be released to the laboratory atmosphere. Prepare standards and handle samples in a well-ventilated fume hood.

6.2 Some of the recommended solvents are flammable. Dimethylacetamide is also absorbed through the skin, so avoid contact.

6.3 Be careful not to come into contact with heated chromatograph parts, such as the detector, column, rotating sample tray, hot sample bottles, etc.

6.4 Once heated, sample vials containing polymer volatiles are under pressure. After analysis, vent the pressure with a hypodermic syringe needle connected to a charcoal slug or vented hood *before* removing vials from the constant temperature bath.

7. Storage and Sampling

7.1 To prevent loss of volatiles from polymer pellets, or resin, store in tightly sealed jars with minimal headspace or heat-sealed pouches constructed of an outer polyester layer, an aluminum foil middle layer, and a polyolefin inner sealing layer.

7.2 For rolled polymer film samples, discard 3 to 5 m of film before taking the sample for headspace analysis because volatiles can be lost by diffusion through the outer film layers.

7.3 Powdered resin samples (for example, poly(vinyl chloride)) can be weighed directly into a tared headspace vial. A 4-g sample (weighed to the nearest 0.0001 g) should be adequate for determination of many volatiles at the parts per million (ppm) levels using sensitive ionization detectors.

7.4 Polymer pellets or granules and film samples can be dissolved, depending on chemistry of polymer, to facilitate attaining thermal equilibrium in a reasonable period of time. These samples are weighed (0.5 ± 0.001 g) into headspace vials, then dissolved in 10 mL of an appropriate solvent. Using ionization detection, these 5 % solutions provide sensitivity for many volatiles at ppm levels. Sample size can be increased to attain sensitivity at lower levels.

NOTE 5—Freeze grinding polymer pellets or granules with liquid nitrogen and screening through a thirty-mesh screen often works as well as testing powdered resin. However, some volatiles may be lost even at these low temperatures.

NOTE 6—This technique of sample preparation requires analysis of the solvent blank to ensure freedom from interferences at volatile component retention times.

NOTE 7—Increased sample sizes can result in viscous solutions. Warming these solutions is often necessary to ensure complete sample dissolution.

8. Preparation of the Gas Chromatograph

8.1 Refer to Recommended Practice E260 and the operator and service manuals of the instrument being used for details regarding column installation, flow rate adjustment, and optimization of detector sensitivity.

8.2 Refer to Practices E355 and E594 for information on terms, relationships, and procedures for testing flame ionization detectors.