

Designation: D 5194 – 04

Standard Test Method for Trace Chloride in Liquid Aromatic Hydrocarbons¹

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1. Scope*

1.1 This test method covers the determination of total chloride (organic and inorganic) in liquid aromatic hydrocarbons and cyclohexane.

1.2 The test method is applicable to samples with chloride concentrations of 1 to 25 mg/kg.

1.3 Bromides and iodides, if present, will be calculated as chlorides.

1.4 Materials, such as styrene, that are polymerized by sodium biphenyl reagent cannot be analyzed by this test method.

1.5 The following applies to all specified limits in this test method: for purposes of determining conformance with this test method, an observed value or a calculated value shall be rounded off "to the nearest unit" in the last right-hand digit used in expressing the specification limit, in accordance with the rounding-off method of Practice E 29.

1.6 The values stated in SI units are to be regarded as standard. No other units of measurement are included in this standard.

1.7 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. For a specific hazard statement, see Section 7.

2. Referenced Documents

2.1 ASTM Standards:

D 891 Test Methods for Specific Gravity, Apparent, of Liquid Industrial Chemicals²

D 1193 Specification for Reagent Water

- D 3437 Practice for Sampling and Handling Liquid Cyclic Products
- D 3505 Test Method for Density or Relative Density of Pure Liquid Chemicals
- D 4052 Test Method for Density and Relative Density of Liquids by Digital Density Meter
- E 29 Practice for Using Significant Digits in Test Data to Determine Conformance with Specifications
- 2.2 Other Documents:
- OSHA Regulations, 29 CFR, paragraphs 1910.1000 and 1910.1200^3

3. Summary of Test Method

3.1 A known amount of hydrocarbon sample is transferred into a separatory funnel containing toluene. Sodium biphenyl reagent is added to convert organic halogens into inorganic halides. The excess reagent is decomposed with water and the phases are separated. The aqueous phase is acidified, washed, and concentrated. Acetone is added and the solution is titrated with silver nitrate solution.

4. Significance and Use

4.1 Organic and inorganic chlorine compounds can have a deleterious effect on equipment and reactions in processes involving aromatic hydrocarbons.

4.2 Maximum chloride levels are often specified for process streams and for aromatic hydrocarbon products.

5. Apparatus

5.1 *Titrator*, potentiometric, recording, + 2000 mV range, 1 mV resolution with dispenser having a volume readout of 0.00 to 9.99 mL or 0.00 to 99.99 mL and 0.01 % resolution.

- 5.2 *Electrode*, glass, reference.
- 5.3 *Electrode*, silver, billet type.

6. Reagents and Materials

6.1 *Purity of Reagents*—Reagent grade chemicals shall be used in all tests. Unless otherwise indicated, it is intended that

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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 Available from U.S. Government Printing Office Superintendent of Documents, 732 N. Capitol St., NW, Mail Stop: SDE, Washington, DC 20401.

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all reagents conform to the specifications of the Committee on Analytical Reagents of the American Chemical Society, where such specifications are available.⁴ Other grades may be used, provided it is first ascertained that the reagent is of sufficiently high purity to permit its use without lessening the accuracy of the determination.

6.2 *Purity of Water*—Unless otherwise indicated, references to water shall be understood to mean reagent water as defined by Types II or III of Specification D 1193.

6.3 Acetone, 99.9 % purity.

6.4 Congo Red Paper.

6.5 Detergent, residue free.

6.6 Isobutanol, 99.9 % minimum purity.

6.7 Isooctane.

6.8 Nitric Acid, concentrated.

6.9 *Nitric Acid*, 5-M. Dilute 160 mL concentrated nitric acid to 500 mL with water.

6.10 Potassium Chloride, primary standard.

6.11 Potassium Chloride Solution, saturated.

6.12 Scouring Powder, cleanser.

6.13 Silver Nitrate, 99.99 % minimum purity.

6.14 Silver Nitrate Solution, 0.01 N, standardized to 0.1 %.

Note 1—This solution may be obtained as follows:

(1) Purchase from a laboratory supply company, (2) Weigh to fourplaces, 1.680 to 1.720 g silver nitrate, transfer quantitatively into a 1000-mL volumetric flask, make to mark with water, and mix well.

Normality of solution =
$$\frac{\text{Weight AgNO}_3}{169.9}$$

or (3) Dissolve 8.5 g silver nitrate in 500 mL water to give a 0.1 N solution. Weigh 0.09 to 0.10 g of dried (105°C) potassium chloride to the nearest 0.1 mg into a 250-mL electrolytic beaker, add 100 mL of water and a stirring bar. While stirring, titrate with the silver nitrate solution.

Normality of AgNO₃ solution = $\frac{\text{Weight KCl}}{0.0746 \times \text{mL AgNO}_3}$

Pipet 50.00 mL of the solution into a 500-mL volumetric flask, dilute to mark with water, and mix well. Divide the calculated normality of the 0.1 N solution by 10 to give the normality of final AgNO $_3$ solution.

6.15 *Sodium Biphenyl Reagent*—The reagent is packed in 15-mL vials that contain 0.2 to 0.4 g of active sodium each. 6.16 *Toluene*, 99.9 % minimum purity.

7. Hazards

7.1 A material, such as styrene, which is polymerized by sodium biphenyl can cause a violent reaction and should never be used as the sample.

7.2 Consult current OSHA regulations and suppliers' Material Safety Data Sheets, and local regulations for all materials used in this test method.

8. Sampling

8.1 Refer to Practice D 3437 for proper sampling and handling of liquid hydrocarbons analyzed by this test method.

9. Electrode Preparation

9.1 Clean the surface of the silver electrode with mild detergent and scouring powder, and rinse with water.

9.2 Immerse the electrode in the saturated potassium chloride solution until the electrode tip turns light gray.

9.3 Rinse well with water and attach to the titrimeter.

9.4 Repeat the electrode preparation when the silver chloride film begins to peel from the surface, or if the film becomes discolored.

10. Procedure for Total Chloride

10.1 Extreme care must be used to prevent contamination and all glassware should be exclusively reserved for this analysis. Just prior to use, the glassware should be rinsed with water followed by acetone and then air dried.

10.2 Place 50 mL of toluene into a 250-mL separatory funnel and pipet in the amount of the liquid sample that corresponds to the estimated chloride content as prescribed in Table 1.

NOTE 2—It is generally more convenient to measure the liquid samples by volume and then convert to mass using density or relative density. Table 2 lists the relative densities of several pure hydrocarbons. Densities of unknowns may be determined by using Test Methods D 891, D 3505 or D 4052.

NOTE 3—Alternately, place the sample into a 125-mL bottle and weigh. From the contents of this bottle add the appropriate amount of the sample to the toluene in the separatory funnel. Reweigh the bottle, and determine the weight of the analytical specimen.

10.3 Add the contents of one vial of sodium biphenyl reagent, stopper the separatory funnel, and gently swirl to mix thoroughly, venting the funnel from time to time. If the resulting solution or suspension is not blue-green, add more sodium biphenyl reagent (one vial at a time) until the blue green color persists.

NOTE 4—The sodium biphenyl reagent has a limited shelf life, given as six months by the manufacturer. This can be extended, in most cases, to approximately one year by keeping the reagent under refrigeration. If this is done, the reagent should be kept at room temperature for several days just prior to use to dissolve any sodium biphenyl that may have precipitated upon cooling.

10.4 Allow the mixture to stand for approximately 10 min. Slowly add 20 mL water and swirl gently with the funnel unstoppered until the blue-green color changes to white. Stopper the funnel again and rock it gently for 1 min, venting the pressure frequently through the stopcock.

10.5 Add 10 mL 5 N nitric acid, and then 5 mL isobutanol. Shake gently, releasing the pressure frequently through the stopcock.

TABLE 1 Specimen Size

Estimated chloride, mg/kg	Specimen volume, mL
0 to 5	100
5 to 25	50

⁴ 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. Pharmacopeial Convention, Inc. (USPC), Rockville, MD.