



Standard Test Method for Formic Acid in Glacial Acetic Acid¹

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1. Scope

1.1 This test method covers the determination of the formic acid (and other reducing substances) content of glacial acetic acid by oxidation with lead tetraacetate.²

1.2 *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.*

1.3 For hazard information and guidance, see the supplier's Material Safety Data Sheets.

2. Referenced Documents

2.1 ASTM Standards:

D 1193 Specification for Reagent Water³

E 200 Practice for Preparation, Standardization, and Storage of Standard Solutions for Chemical Analysis⁴

3. Summary of Test Method

3.1 In an aqueous acetic acid solution each molecule of formic acid is oxidized to one molecule of carbon dioxide by reduction of one molecule of lead tetraacetate. The amount of lead tetraacetate remaining is then determined by adding potassium iodide and titrating the liberated iodine with standard sodium thiosulfate solution to the starch end point.

4. Significance and Use

4.1 This test method is useful for determining the formic acid content of glacial acetic acid by chemical means.

4.2 Low molecular weight organic acids (such as acetic and propionic), aldehydes (including formaldehyde and acetaldehyde), ketones, and alcohols (including methyl alcohol) do not interfere with the test.

4.3 Formic acid (and other reducing substances) may be present as a result of contamination during storage, distribution, and manufacture. This test method may be used in assessing compliance with a specification.

5. Reagents

5.1 *Purity of Reagents*—Reagent grade chemicals shall be used in all tests. Unless otherwise indicated it is intended that all reagents shall conform to the specifications of the Committee on Analytical Reagents of the American Chemical Society, where such specifications are available.⁵ 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.

5.2 *Purity of Water*—Unless otherwise indicated reference to water shall be understood to mean reagent water conforming to Type IV of Specification D 1193.

5.3 *Lead Tetraacetate Solution*—Dissolve completely 40.0 g of sodium acetate in 1 L of glacial acetic acid and then dissolve 40.0 g of lead tetraacetate.

5.4 *Sodium Thiosulfate Standard Solution (0.1 N)*—Prepare and standardize 0.1 N sodium thiosulfate ($\text{Na}_2\text{S}_2\text{O}_3$) as described in Practice E 200.

5.5 *Starch Indicator Solution*—Prepare the starch indicator solution as described in Practice E 200.

5.6 *Stopping Solution*—Dissolve 100 g of potassium iodide and 500 g of sodium acetate in 1 L of water.

6. Procedure

6.1 Pipet 25 mL of the lead tetraacetate solution into each of two 500-mL Erlenmeyer flasks. Pipet the specimen (Note 1) into one of the flasks, swirl, and allow to stand for 20 to 30 min. The second flask is the blank.

NOTE 1—The following are suggested specimen sizes for varying concentrations of formic acid:

Formic Acid, %	Specimen Size, mL
0.0 to 0.10	25
0.10 to 0.30	10
0.30 to 0.60	5
0.60 to 1.00	3
1.00 to 2.00	2
2.00 to 3.00	1

6.2 Add to each flask a slight excess of the stopping solution as indicated by the formation of the yellow lead iodide precipitate. Add enough water just to dissolve the yellow

¹ This test method is under the jurisdiction of ASTM Committee D-1 on Paint and Related Coatings, Materials, and Applications and is the direct responsibility of Subcommittee D01.35 on Solvents, Plasticizers, and Chemical Intermediates.

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² Perlin, A. S., "Determination of Formic Acid by Oxidation with Lead Tetraacetate," *Analytical Chemistry*, Vol 26, 1954, pp. 1053–1054.

³ *Annual Book of ASTM Standards*, Vol 11.01.

⁴ *Annual Book of ASTM Standards*, Vol 15.05.

⁵ *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.