



Designation: **D 4972 – 95a01**

Standard Test Method for pH of Soils¹

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1. Scope*

1.1 This test method covers the measurement of the pH of soils for uses other than for corrosion testing. Such measurements are used in the agricultural, environmental, and natural resources fields. This measurement determines the degree of acidity or alkalinity in soil materials suspended in water and a 0.01 *M* calcium chloride solution. Measurements in both liquids are necessary to fully define the soil's pH. This variable is useful in determining the solubility of soil minerals and the mobility of ions in the soil and assessing the viability of the soil-plant environment. A more detailed discussion of the usefulness of this parameter is not

¹ This test method is under the jurisdiction of ASTM Committee D-18 on Soil and Rock and is the direct responsibility of Subcommittee D18.22 on Soil as a Medium for Plant Growth.

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

warranted here; however, it can be found in many discussions of the subject. A few such discussions are given as Refs (1-6)² at the end of the text.

1.2 The values given in SI units are to be regarded as the standard.

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:

C 670 Practice for Preparing Precision and Bias Statements for Test Methods for Construction Materials³

D 1193 Specification for Reagent Water 653 Terminology Relating to Soil, Rock, and Contained Fluids⁴

D 1193 Specification for Reagent Water⁵

D 3740 Practice for Minimum Requirements for Agencies Engaged in the Testing and/or Inspection of Soil and Rock as Used in Engineering Design and Construction⁶

G 51 Test Method for pH of Soil for Use in Corrosion Testing⁷

3. Summary Terminology

3.1 Definitions:

3.1.1 For common definitions of Test Method

3.1 Measurement of the pH of soils in both suspensions of water and a calcium chloride solution are made with either a potentiometer using a pH sensitive electrode system (Method A), or pH sensitive paper (Method B). The potentiometer is calibrated with buffer solutions of known pH. The pH sensitive paper is a less accurate measurement and should only be terms used for a rough estimate of the soil pH. The electrode must be used for in this measurement unless the pH sensitive paper is specified. standard, refer to Terminology D 653.

4. Significance and Use

4.1 The pH Summary of the soil is a useful variable in determining the solubility Test Method

4.1 Measurement of soil minerals and the mobility pH of ions soils in the soil and assessing the viability of the soil-plant environment.

4.2 pH measurements are made in both suspensions of water and a calcium chloride solution because the calcium displaces some of the exchangeable aluminum. are made with either a potentiometer using a pH sensitive electrode system (Method A), or pH sensitive paper (Method B). The low ionic strength counters the dilution effect on the exchange equilibrium by setting the salt concentration potentiometer is calibrated with buffer solutions of the solution closer to that expected in the soil solution. known pH. The pH values obtained in the solution sensitive paper is a less accurate measurement and should only be used for a rough estimate of calcium chloride are slightly lower than those measured in water due to the release of more aluminum ions which then hydrolyses. Therefore, both measurements are required to fully define the character of the soil's pH.

4.3 For the purpose of this test method the test soil pH. The electrode must be sieved through a No. 10 sieve (2 mm sieve mesh openings). Measurements on soils or soil fractions having particle sizes larger than 2 mm by this test method may be invalid. If soil or soil fractions with particles larger than 2 mm are used, it must be stated in the report since the results may be significantly different.

4.4 AH water used for this test method must be ASTM Type III or better. Type III water measurement unless the pH sensitive paper is defined by Specification D 1193. It is prepared by distillation, ion exchange, reverse osmosis, or a combination thereof. specified.

5. Significance and Use

5.1 The pH of the soil is a useful variable in determining the solubility of soil minerals and the mobility of ions in the soil and assessing the viability of the soil-plant environment.

5.2 pH measurements are made in both water and a calcium chloride solution because the calcium displaces some of the exchangeable aluminum. The low ionic strength counters the dilution effect on the exchange equilibrium by setting the salt concentration of the solution closer to that expected in the soil solution. The pH values obtained in the solution of calcium chloride

² The boldface numbers in parentheses refer to a list of references at the end of the test method.

³ *Annual Book of ASTM Standards*, Vol 04.02.

⁴ *Annual Book of ASTM Standards*, Vol 11.01: 04.09.

⁵ *Annual Book of ASTM Standards*, Vol 03.02: 11.01.

⁶ *Reagent Chemicals, American Chemical Society Specifications*, American Chemical Society, Washington, DC. For suggestions on the testing *Annual Book of reagents not listed by the American Chemical Society, see Analytical ASTM 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: Vol 04.08.

⁷ These data satisfy the 1S and D2S requirements outlined in Practice C 670.

⁷ *Annual Book of ASTM Standards*, Vol 03.02.

are slightly lower than those measured in water due to the release of more aluminum ions which then hydrolyses. Therefore, both measurements are required to fully define the character of the soil's pH.

5.3 For the purpose of this test method the test soil must be sieved through a No. 10 sieve (2 mm sieve mesh openings). Measurements on soils or soil fractions having particle sizes larger than 2 mm by this test method may be invalid. If soil or soil fractions with particles larger than 2 mm are used, it must be stated in the report since the results may be significantly different.

5.4 All water used for this test method must be ASTM Type III or better. Type III water is defined by Specification D 1193. It is prepared by distillation, ion exchange, reverse osmosis, or a combination thereof.

6. Interferences

56.1 This test method as measured by a pH probe has possible interferences due to a suspension effect or sedimentation potential. Users interested in a detailed discussion of the mechanism of this effect can find it in Refs (5) and (6).

56.2 This effect is the main reason Test Method G 51 can not be used for general measurement of pH outside of that for corrosion analysis. Test Method G 51 measures pH (an aqueous parameter) without adding any aqueous phase to the soil. This results in excessive soil particle-pH probe contact that overestimates the activity of the hydrogen ions in solution and is therefore unacceptable for general soil analysis.

56.3 The suspension effect can be mitigated by careful attention to 9.10.1.

67. Apparatus

67.1 *Method A, pH Meter*—Potentiometer equipped with glass-calomel electrode system. Follow the manufacturer's instructions for the pH meter used. A silver/silver chloride electrode system or similar is also acceptable.

67.2 *Method B, pH Paper*—pH paper sensitive to a pH range from 1 to 12, with resolution to the nearest 0.2 pH unit.

78. Reagents

78.1 *Purity of Reagents*—Reagent grade chemicals should be used in all tests. Unless otherwise indicated, it is intended that all reagents should conform to the specifications of the Committee on Analytical Reagents of the American Chemical Society, where such specifications are available.⁸ Other grades may be used, provided it is first ascertained that the reagent is of sufficient purity to permit its use without lessening the accuracy of the determination.

78.2 *Purity of Water*—All water used for this test method must be ASTM Type III or better. Type III water is defined by Specification D 1193. It is prepared by distillation, ion exchange, reverse osmosis, or a combination thereof.

78.3 *Acid Potassium Phthalate Buffer Solution (0.05 M)*—Dissolve 10.21 g (dried 1 h at 105°C) of potassium phthalate in water and dilute to 1 L. The pH of this solution should be 4.0 at 20°C. Protect the solution against evaporation and against contamination with molds. Replace the solution when mold is noticed. The effect of temperature is as follows:

°C	pH
5 to 37	4.0

This illustrates that the pH of the solution does not change over the range in temperature from 5 to 37°C.

78.4 *Calcium Chloride Stock Solution (1.0 M)*—Dissolve 147 g of CaCl₂·2H₂O in water in a 1-L volumetric flask, cool, dilute to volume with water, and mix.

78.5 *Calcium Chloride Solution (0.01 M)*—Dilute 20.0 mL of stock 1.0 M CaCl₂ solution to 2 L with water. The pH of this solution should be between 5 and 7.

78.6 *Phosphate Buffer Solution (0.025 M)*—Dissolve 3.40 g of KH₂PO₄ and 3.55 g of KH₂HPO₄ in water and dilute to 1 L. Dry salts 2 h at 130°C before use. The pH of this solution should be 6.9 at 20°C. The effect of temperature is as follows:

°C	pH
0	7.0
10	6.9
20	6.9
30	6.8
40	6.8

8. Calibration of pH Meter

8.1 Calibrate the pH meter using the acid potassium phthalate and phosphate buffer solutions. Adjustment of the pH meter should follow the manufacturer's direction.

9. Calibration of pH Meter

9.1 Calibrate the pH meter using the acid potassium phthalate and phosphate buffer solutions. Adjustment of the pH meter should follow the manufacturer's direction.

⁸ *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.

10. Procedure

910.1 When making measurements with the pH electrode, place the electrode into the partially settled suspension to mitigate the suspension effect.

910.2 For both methods, begin with an air dried soil that has been sieved through a No. 10 sieve (2 mm holes) to remove the coarser soil fraction. Air drying the soil is necessary to accomplish sieving and to control the amount of water present at the time of measurement.

910.3 *pH in Distilled Water*—For both methods, weigh out approximately 10 g of air dried soil. Place the soil into a glass container and add approximately 10 mL of water. Mix thoroughly and let stand for 1 h.

910.4 *Method A*—Read pH on pH meter.

910.5 *Method B*—Read pH on pH paper.

910.6 *pH in 0.01 M Calcium Chloride Solution*—For both methods weigh out approximately 10 g of air dried soil. Place the soil into a glass container and add approximately 10 mL of the 0.01 M CaCl₂ solution. Mix thoroughly and let stand for 1 h.

910.7 *Method A*—Read pH on pH meter.

910.8 *Method B*—Read pH on pH paper.

910.9 The mixture should be at approximately room temperature (15 to 25°C) at the time of pH measurement.

10. Report

10.1 Report the pH of the soil to the first decimal place. Specify which of the pH measurements is in water and which is in the calcium chloride solution. Also specify whether the determinations were made with Method A or Method B. If size fractions other than sieved through the No. 10 sieve are used, it must be stated in the report since the results may be significantly different.

11. Report

11.1 Report the pH of the soil to the first decimal place. Specify which of the pH measurements is in water and which is in the calcium chloride solution. Also specify whether the determinations were made with Method A or Method B. If size fractions other than sieved through the No. 10 sieve are used, it must be stated in the report since the results may be significantly different.

12. Precision and Bias

12.1 Precision:

12.1.1 *Within-Laboratory Precision*—The within laboratory standard deviations for Method A are 0.031 (pH units) for the water mixture and 0.139⁹ (pH units) for the calcium chloride mixture. Therefore, results of two properly conducted tests in the same or different laboratories should not differ by more than 0.065⁹ (pH units) for the water mixture and 0.389 pH units for the calcium chloride mixture. The within-laboratory standard deviations for Method B are 0.189 (pH units) for the water mixture and 0.212⁹ (pH units) for the calcium chloride mixture. Therefore, results of two properly conducted tests in the same or different laboratories should not differ by more than 0.53⁹ (pH units) for the water mixture and 0.60 pH units for the calcium chloride mixture.

12.1.1.1 The precision of Method A presented was determined by the National Technical Center of the United States Department of Agriculture. In their evaluation they used 174 replicates for the water mixture and 32 replicates in testing the calcium chloride mixture.

12.1.1.2 The precision of Method B presented was determined by the United States Army Environmental Hygiene Agency. In their evaluation they used 25 replicates in testing each mixture.

12.1.2 *Between-Laboratory Precision*—The between-laboratory standard deviation has not been determined for either method. Subcommittee D18.22 is actively seeking data to evaluate the between laboratory precision of this test method.

12.2 *Bias*—There is no accepted reference value for this test method; therefore, bias cannot be determined.

123. Keywords

123.1 acidity; alkalinity; pH; reaction; soil

⁹These data satisfy the 1S and D2S requirements outlined in Practice C 670.

REFERENCES

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- (3) Sposito, G., *The Thermodynamics of Soil Solutions*, Clarendon, Oxford, United Kingdom, 1981.
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- (5) Hunter, R. J., *Zeta Potential in Colloid Science*, Academic, New York, NY, 1981.
- (6) Perrin, D. D., and Dempsey, B., *Buffers for pH and Metal Ion Control*, Chapman and Hall, London, United Kingdom, 1974.

SUMMARY OF CHANGES

This section identifies location of changes to this practice since the last edition.

- (1) Added Practice D 3740 and Terminology D 653 to Referenced Documents.
- (2) Added Section 3 on Terminology.

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