

# Standard Test Method for Cement Content of Hardened Soil-Cement Mixtures <sup>1</sup>

This standard is issued under the fixed designation D806; the number immediately following the designation indicates the year of original adoption or, in the case of revision, the year of last revision. A number in parentheses indicates the year of last reapproval. A superscript epsilon ( $\varepsilon$ ) indicates an editorial change since the last revision or reapproval.

## 1. Scope\*

1.1 This test method covers the determination by chemical analysis of cement content of hardened soil-cement mixtures.

1.2 *Units*—The values stated in SI units are to be regarded as standard. No other units of measurement are included in this standard.

1.3 All observed and calculated values shall conform to the guidelines for significant digits and rounding established in Practice D6026.

1.3.1 For purposes of comparing a measured or calculated value(s) with the specific limits, the measured or calculated value(s) shall be rounded to the nearest decimal or significant digits in the specific limits.

1.3.2 The procedures used to specify how data are collected/ recorded or calculated in the standard are regarded as industry standard. In addition they are representative of the significant digits that generally should be retained. The procedures used do not consider material variation, purpose for obtaining the data, special purpose studies, or any considerations for the user's objectives; and it is common practice to increase or reduce significant digits of reported data to be commensurate with these considerations. It is beyond the scope of this standard to consider significant digits used in analysis methods for the engineering design.

1.4 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, health, and environmental practices and determine the applicability of regulatory limitations prior to use. For specific hazard precautions, see Section 7.

1.5 This international standard was developed in accordance with internationally recognized principles on standardization established in the Decision on Principles for the Development of International Standards, Guides and Recommendations issued by the World Trade Organization Technical Barriers to Trade (TBT) Committee.

## 2. Referenced Documents

- 2.1 ASTM Standards:<sup>2</sup>
- C114 Test Methods for Chemical Analysis of Hydraulic Cement
- C125 Terminology Relating to Concrete and Concrete Aggregates
- C219 Terminology Relating to Hydraulic and Other Inorganic Cements
- D653 Terminology Relating to Soil, Rock, and Contained Fluids
- D2216 Test Methods for Laboratory Determination of Water (Moisture) Content of Soil and Rock by Mass
- D2901 Test Method for Cement Content of Freshly Mixed Soil-Cement
- D3740 Practice for Minimum Requirements for Agencies Engaged in Testing and/or Inspection of Soil and Rock as Used in Engineering Design and Construction
- D4753 Guide for Evaluating, Selecting, and Specifying Balances and Standard Masses for Use in Soil, Rock, and Construction Materials Testing
- D5982 Test Method for Determining Cement Content of Fresh Soil-Cement (Heat of Neutralization Method)

D6026 Practice for Using Significant Digits in Geotechnical

### 3. Terminology

#### 3.1 Definitions:

3.1.1 For definitions of common technical terms used in this standard relating to soil, refer to Terminology D653

3.1.2 Refer to Terminology C125 for definitions of common technical terms relating to concrete.

3.1.3 Refer to Terminology C219 for definitions of common technical terms relating to cement.

#### 4. Summary of Test Method

4.1 Samples of the raw soil, cement and soil-cement mixtures are obtained and analyzed. Hydrochloric acid is added to each sample and the resulting precipitate discarded. The

<sup>&</sup>lt;sup>1</sup> This test method is under the jurisdiction of ASTM Committee D18 on Soil and Rock and is the direct responsibility of Subcommittee D18.15 on Stabilization With Admixtures.

Current edition approved Dec. 1, 2019. Published January 2020. Originally approved in 1944. Last previous edition approved in 2011 as D806–11. DOI: 10.1520/D0806-19.

<sup>&</sup>lt;sup>2</sup> 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.

remaining filtrate is made slightly ammonaical and the hydroxides are allowed to settle out. The precipitate is washed with ammonium nitrate and the hydroxides are dissolved with hydrochloric acid. The solution is again made slightly ammonaical, the precipitate washed with ammonium nitrate and hydroxide precipitate discarded. Ammonium hydroxide is added to the filtrate, the solution is heated and ammonium oxalate is added. The calcium oxalate is filtered off, washed into a beaker, diluted and sulfuric acid then added. This is then titrated with a standard potassium permanganate solution. The cement content is then computed by means of equations given in Section 10.

#### 5. Significance and Use

5.1 This test method determines cement content in mixtures of cement with soil or aggregate by chemical analysis. It was developed primarily for testing samples for which a significant degree of cement hydration or hardening has taken place. Test Methods D2901 or D5982 may be used for determining cement content of freshly mixed soil-cement mixtures.

5.2 This test method is based on determination by chemical analysis of the calcium oxide (CaO) content of the sample. The method may not be applicable to soil-cement materials containing soils or aggregates which yield significant amounts of dissolved calcium oxide (CaO) under the conditions of the test.

Note 1—The quality of the result produced by this standard is dependent on the competence of the personnel performing it, and the suitability of the equipment and facilities used. Agencies that meet the criteria of Practice D3740 are generally considered capable of competent and objective testing and sampling. Users of this standard are cautioned the compliance with Practice D3740 does not in itself assure reliable results. Reliable results depend on many factors: Practice D3740 provides a means of evaluating some of these factors.

NOTE 2—Analysis using X-ray fluorescence in accordance with C114 may also be used for determination of calcium oxide (CaO) content of hardened soil-cement.

#### 6. Apparatus

6.1 *Analytical Balance*—An analytical balance conforming to the requirements of Class GP2 in Specification D4753 and with Class S weights.

6.2 *Filter Paper*—Qualitative filter paper including Whatman No. 1, 11 and 15 cm in diameter; Whatman No. 41, 15 cm in diameter; and Whatman No. 2, 11 or 15 cm in diameter.

6.3 Fifty-Millilitre-Pipet.

6.4 Drying Oven—Thermostatically controlled, preferably of the forced-air type, meeting the requirements of Test Method D2216 and capable of maintaining a uniform temperature of  $110 \pm 5^{\circ}$ C throughout the drying chamber.

6.5 *Miscellaneous Apparatus*—Supplementary equipment, such as electric ovens, hot plates, a small riffle, a 425  $\mu$ m (No. 40) sieve with bottom pan and cover, a cast iron mortar and pestle, and a ball mill if possible.

#### 7. Reagents

7.1 *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 Commit-

tee on Analytical Reagents of the American Chemical Society, where such specifications are available.<sup>3</sup> 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.

7.2 Potassium Permanganate, Standard Solution (0.1 N)— Prepare a 0.1 N KMnO<sub>4</sub> solution and standardize against primary standard grade oxalic acid, sodium oxalate or iron (II) ammonium sulfate hexahydrate.

Note 3—The use of a standard  $0.1 N \text{ KMnO}_4$  solution is not necessary when the samples are titrated in accordance with 9.9 and the results are calculated in accordance with 10.2. However, the actual reagent concentration must be determined by titration against primary standard grade oxalic acid, sodium oxalate or iron (II) ammonium sulfate hexahydrate.

7.3 Ammonium Nitrate Solution—Dissolve 20 g of  $NH_4NO_3$  in 1L of distilled water.

7.4 *Hydrochloric Acid* (1 + 3)—Add 200 mL of HCl (sp gr 1.19) to 600 mL of distilled water.

7.5 *Hydrochloric Acid* (1 + 1)—Add 25 mL of HCl (sp gr 1.19) to 25 mL of distilled water.

7.6 Nitric Acid—See Note 4.

7.7 Ammonium Oxalate Solution (5 %) —50 g of ammonium oxalate. (Warning —In addition to other precautions, this is done by adding the acid, slowly while stirring, to the water to avoid a sudden temperature rise that could cause boiling and spattering of the acid solution.)

7.8 Ammonium Hydroxide, NH<sub>4</sub>OH (sp gr 0.90).

7.9 Sulfuric Acid (1 + 1)—Add 500 mL H<sub>2</sub>SO<sub>4</sub> (sp gr 1.84) to 500 mL of distilled water.

### 8. Samples

8.1 Samples of the following shall be selected for the test: 8.1.1 *Raw Soil*, representative of the soil phase of the soil-cement mixture.

8.1.2 *Cement*, representative of the cement phase of the soil-cement mixture, and

8.1.3 Soil-cement Mixture to be analyzed.

8.2 The gross laboratory sample of each component shall be approximately 200 g. This may be obtained by reducing the sample in bulk and, if necessary, in particle size through the use of drying, riffling and grinding processes.

#### 9. Procedure

9.1 Dry 25 g of each of the samples in an oven to constant mass at  $110 \pm 5^{\circ}$ C to remove free water. Reduce the samples to pass a 425 µm (No. 40) sieve.

9.2 Using an analytical balance, prepare the following amounts for each of the samples: raw soil, 5 g; soil-cement mixture, 5 g; and cement, 1 g. Place each of the weighed

<sup>&</sup>lt;sup>3</sup> "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 "Reagent Chemicals and Standards," by Joseph Rosin, D. Van Nostrand Co., Inc., New York, NY, and the "United States Pharmacopeia."