



Designation: D4959 – 24

Standard Test Method for Determination of Water Content of Soil By Direct Heating¹

This standard is issued under the fixed designation D4959; 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 reappraisal. A superscript epsilon (ϵ) indicates an editorial change since the last revision or reappraisal.

1. Scope*

1.1 This test method covers procedures for determining the water content of soils by drying with direct heat, such as using a hotplate, stove or a blowtorch, where the heat is applied to the container and not directly to the soils.

1.2 This test method can be used as a substitute for Test Methods **D2216** when more rapid results are desired to expedite other phases of testing and less accurate results are acceptable.

1.3 When questions of accuracy between this test method and Test Methods **D2216** arise, the results of Test Methods **D2216** will be used.

1.4 This test method is applicable for most soil types. For some soils, such as those containing significant amounts of halloysite, mica, montmorillonite, gypsum, or other hydrated materials, highly organic soils or soils that contain dissolved solids, (such as salt in the case of marine deposits), this test method may not yield reliable water content values due to the potential for heating above 110°C or lack of means to account for the presence of precipitated solids that were previously dissolved.

1.5 *Units*—The values stated in SI units are to be regarded as standard. No other units of measure are included in this standard. The sieve designations are identified using the “standard” system in accordance with Specification **E11**, such as 2.0-mm and 19-mm, followed by the “alternative” system of No. 10 and ¾-in., respectively, in parentheses. Reporting of test results in units other than SI shall not be regarded as nonconformance with this standard.

1.6 All observed and calculated values must conform to the guidelines for significant digits and rounding established in Practice **D6026**, unless otherwise superseded by this standard.

1.6.1 The procedures used to specify how data are collected/recorded or calculated, in this standard are regarded as the 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 design.

1.6.2 Significant digits are especially important if the water content will be used to calculate other relationships such as moist mass to dry mass or vice versa, wet unit weight to dry unit weight or vice versa, and total density to dry density or vice versa. For example, if four significant digits are required in any of the above calculations, then the water content must be recorded to the nearest 0.1 %, for water contents below 100 %. This occurs since 1 plus the water content (not in percent) will have four significant digits regardless of what the value of the water content is (below 100 %); that is, 1 plus 0.1/100 = 1.001, a value with four significant digits. While, if three significant digits are acceptable, then the water content can be recorded to the nearest 1 %.

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, health, and environmental practices and determine the applicability of regulatory limitations prior to use.*

1.8 *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:*²

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

¹ This test method is under the jurisdiction of ASTM Committee **D18** on Soil and Rock and is the direct responsibility of Subcommittee **D18.08** on Special and Construction Control Tests.

Current edition approved Feb. 1, 2024. Published February 2024. Originally approved in 1989. Last previous edition approved in 2016 as D4959 – 16. DOI: 10.1520/D4959-24.

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.

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

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

D6026 Practice for Using Significant Digits and Data Records in Geotechnical Data

E11 Specification for Woven Wire Test Sieve Cloth and Test Sieves

3. Terminology

3.1 *Definitions*—For definitions of common technical terms used in this standard, refer to Terminology **D653**.

3.2 *Definitions of Terms Specific to This Standard:*

3.2.1 *direct heating, n*—a process by which the soil is dried by conductive heating from the direct application of heat in excess of 110°C to the specimen container, such as provided by a hot plate, gas stove or burner, heatlamps, or other heat sources. Direct application of heat by flame to the specimen is not appropriate.

4. Summary of Test Method

4.1 A moist soil specimen is placed in a suitable container and its mass is determined. It is then subjected to drying by the application of direct heat until dry by appearance, removed from the heat source, and its new mass is determined. This procedure is repeated until the mass becomes nearly constant within specified limits.

4.2 The difference between the masses of the moist specimen and the dried specimen is used as the mass of water contained in the specimen. The water content (expressed as a percentage) is determined by dividing the mass of water by the dry mass of soil, multiplied by 100. For a given soil type and specimen size, the time to achieve a constant dry mass can be noted and used to estimate drying time for subsequent tests of the same soil type using the same size specimen and drying apparatus.

5. Significance and Use

5.1 The water content of a soil is used throughout professional practice both in the laboratory and in the field. The use of Test Methods **D2216** for water content determination can be time consuming and there are occasions when a more expedient method is desirable. Drying by direct heating is one such method. Results of this test method have been demonstrated to be of satisfactory accuracy for use in field control testing, such as in the determination of water content, and in the determination of in-place dry unit weight of soils.

5.2 The principal objection to the use of the direct heating for water content determination is the possibility of overheating the soil, thereby yielding a water content higher than would be determined by Test Methods **D2216**. While not eliminating this possibility, the incremental drying procedure in this test method will reduce its effects. Some heat sources have settings or controls that can also be used to reduce overheating. Loose fitting covers or enclosures can also be used to reduce overheating while assisting in uniform heat distribution.

5.3 The behavior of a soil when subjected to direct heating is dependent on its mineralogical composition, and as a result, no one procedure is applicable for all types of soils or heat sources. The general procedure of this test method applies to all soils, but test details may need to be tailored to the soil being tested.

5.4 When this test method is to be used repeatedly on the same or similar soil from a given site, a correction factor can usually be determined by making several comparisons between the results of this test method and Test Methods **D2216**. A correction factor is valid when the difference is consistent for several comparisons, and is reconfirmed on a regular specified basis.

5.5 This test method is not appropriate when precise results are required, or when minor variations in water content will affect the results of other test methods, such as borderline situations where small variations in the measured water content could affect acceptance or rejection.

5.6 This test method is not appropriate for specimens known to contain flammable organics or contaminants, and other test methods should be utilized in these situations.

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 facility used. Agencies that meet the criteria of Practice **D3740** are generally considered capable of competent and objective testing/sampling/inspection/etc. Users of this standard are cautioned that 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 those factors.

6. Interferences

6.1 When testing sand and gravel size particles, additional care should be taken to avoid the possibility of particle shattering.

6.2 Due to the localized high temperatures in the soil during testing, the physical characteristics of the soil may be altered. Degradation of individual particles may occur, along with vaporization, chemical transition, or loss of organics. Therefore, specimens used in this test method must not be used for other tests after drying.

7. Apparatus

7.1 *Direct Heat Source*—Any source of heat that can be directed to the soil specimen to raise the specimen temperature to or above 110°C. Commonly used sources include electric, gas, butane or oil-fired stoves, and hotplates, blowtorches, heat lamps, hair driers, and space heaters. Heat sources that directly apply open flame to the specimen may cause extreme degradation of the specimen along with oxidation of and depositing of soot in the specimen and must not be used.

7.2 *Balances*—All balances must meet the requirements of Guide **D4753** and this section. A Class GP2 balance of 0.1 g readability is generally advised. However, the balance used may be controlled by the number of significant digits needed (see 1.6).

7.3 *Specimen Containers*—Suitable containers made of material resistant to corrosion and a change in mass upon repeated

heating, cooling, and cleaning. One container is needed for each water content determination.

7.4 *Container Handling Apparatus*—Gloves or suitable holder for moving hot containers after drying.

7.5 *Desiccator (optional)*—A desiccator cabinet or jar of suitable size containing silica gel, anhydrous calcium phosphate, or equivalent. It is preferable to use a desiccant that changes color to indicate when it needs to be replaced.

7.6 *Miscellaneous (as needed)*—Mixing tools such as spatulas or spoons; eye protection, such as safety glasses or goggles; dry light-weight paper or tissue, and knives.

8. Hazards

8.1 Container holders or gloves are recommended for handling hot containers. Some soil types can retain considerable heat, and serious burns could result from improper handling.

8.2 Suitable eye protection such as safety glasses or goggles is recommended due to the possibility of particle shattering during heating, mixing, or mass determinations.

8.3 Highly organic soils, and soils containing oil or other contaminants may ignite during drying with direct heat sources. Means for smothering flames to prevent operator injury or equipment damage should be available during testing. Fumes given off from contaminated soils or wastes may be toxic, and should be vented accordingly.

8.4 Due to the possibility of steam explosions, or thermal stress shattering of porous or brittle aggregates, a vented covering over the sample container may be appropriate to prevent operator injury or equipment damage. This also prevents loss of the test specimen during the drying cycle while aiding in uniform heating of the specimen.

9. Samples

9.1 Perform the water content determination as soon as practical after sampling to prevent water loss and damage to potentially corrodible containers.

9.2 Prior to testing, store samples in non-corrodible airtight containers at a temperature between approximately 3 and 30°C and in an area that prevents direct exposure to sunlight.

10. Test Specimens

10.1 Select a representative portion of the total sample. If a layered soil or more than one soil type is encountered, select an average portion or individual portions of each, and note which portion(s) were tested in the report of the results.

10.1.1 For cohesionless soils, mix the material thoroughly, and select a test specimen having a mass of moist material in accordance with [Table 1](#).

10.1.2 For cohesive soils, remove about 3 mm of material from the exposed periphery of the sample and slice the remaining specimen in half (to check if the material is layered), prior to selecting a test specimen having a moist mass in accordance with [Table 1](#). If the soil is layered, see [10.1](#). Breaking or cutting of cohesive samples to approximately 6 mm particles speeds drying and prevents crusting or overheating the surface while drying the interior.

TABLE 1 Test Specimen Masses

Sieve Size Retaining More Than 10 % of Sample, mm	Minimum Mass of Moist Specimen, g ^A
2.0 (No. 10)	200 to 300
4.75 (No. 4)	300 to 500
19.0 (No. ¾)	500 to 1000

^A Larger specimens may be used and are encouraged. Generally, inherent test inaccuracies are reduced by using specimens with as large a mass as practical.

10.2 Using a test specimen smaller than the minimum mass indicated in [Table 1](#) requires discretion, though it may be adequate for the purpose of the test. Note a specimen having a mass less than the previously indicated value in the report of results.

NOTE 2—When working with a small sample containing a relatively large coarse-grained particle, it may be appropriate not to include this particle in the test specimen, depending on the use of test results. If this is done, such exclusion should be noted in the report of the results.

10.3 When the result of a water content determination by the use of this test method is to be compared to the results of another method, such as Test Methods [D2216](#), obtain a second specimen during selection of the specimen for this comparison. Take precautions to obtain a specimen that represents the same water content as closely as practicable. Protect the comparison specimens from water loss by transporting and storing the specimens in sealed containers. A correction factor can be determined for use on subsequent water content determinations on the same soil types from the same site when the difference is relatively constant using several comparisons. Check the correction factor on a regular, specified basis. Recognize that different technicians, heat sources, and such may result in different correction factors.

11. Conditioning

11.1 Prepare, process, and test all specimens quickly to reduce unrecorded moisture loss.

11.2 Cut or break up the soil into small size aggregations to aid in obtaining more uniform drying of the specimen, taking care to avoid any loss of soil.

11.3 If the specimens are not being tested immediately, place the specimens in containers that can be closed and stored in an area not exposed to direct sunlight, to prevent loss of moisture prior to initial mass determinations.

12. Procedure

12.1 Determine the mass of a clean, dry specimen container, and record.

12.2 Place the soil specimen in the container, and immediately determine and record the mass of the soil and container.

12.3 Apply heat to the soil specimen and container, taking care to avoid localized overheating. Continue heating while stirring the specimen to obtain even heat distribution. Continue application of heat until the specimen first appears dry. A comparatively uniform color should result. Avoid localized burnt or darkened appearance of any part of the soil by intermittent mixing and stirring.