



Standard Test Method for Moisture in Wool by Oven-Drying¹

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1. Scope

1.1 This test method covers the determination of the amount of moisture present in ordinary commercial and industrial samples of wool in all forms except grease wool, using the oven-drying technique.

1.2 Formulas for calculating the moisture content (as-received basis) and moisture regain (oven-dried basis) are given. It is always important to use the correct term which corresponds to the basis used in the calculation (see 12.2.1).

NOTE 1—The determination of moisture content for textile materials in general is covered in Test Methods D2654, and an optimal method for determining the moisture in wool by distillation with toluene is covered in Test Method D2462. A method for sampling wool for the determination of moisture in wool is covered in Practice D2525. The oven-drying method has been adapted for cotton in Test Method D2495.

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:*²

D123 Terminology Relating to Textiles

D1060 Practice for Core Sampling of Raw Wool in Packages for Determination of Percentage of Clean Wool Fiber Present

D1776 Practice for Conditioning and Testing Textiles

D2258 Practice for Sampling Yarn for Testing

D2462 Test Method for Moisture in Wool by Distillation With Toluene

D2495 Test Method for Moisture in Cotton by Oven-Drying

D2525 Practice for Sampling Wool for Moisture

¹ This test method is under the jurisdiction of ASTM Committee D13 on Textiles and is the direct responsibility of Subcommittee D13.13 on Wool and Felt.

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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.

D2654 Test Method for Moisture in Textiles (Withdrawn 1998)³

D3333 Practice for Sampling Manufactured Staple Fibers, Sliver, or Tow for Testing

D4845 Terminology Relating to Wool

3. Terminology

3.1 For all terminology related to D13.13, Wool and Felt, see Terminology D4845.

3.1.1 The following terms are relevant to this standard: grease wool, moisture content, moisture-free, moisture regain, oven-dried, pulled wool, raw wool, recycled wool, scoured wool, virgin wool, wool, wool, *as defined in the Wool Products Labeling Act of 1939*.

3.2 For definitions of all other textile terms see Terminology D123.

4. Summary of Test Method

4.1 A specimen of wool material is weighed and then dried to constant mass at $105 \pm 2^\circ\text{C}$ in an oven supplied with ambient air. The loss in mass is considered moisture and reported as either moisture content or moisture regain. Directions are given for the adjustment of the observed results for any change in the moisture content after sampling and before drying.

5. Significance and Use

5.1 Test Method D2462 for the determination of the moisture in wool by distillation with toluene is the preferred method for testing wool for moisture for the acceptance testing of commercial shipments. If, however, the purchaser and the supplier agree, Test Method D1576 for the determination of the moisture in wool by oven drying may be used instead. Comparative tests as directed in 5.1.1, may be advisable.

5.1.1 In case of a dispute arising from differences in reported test results when using Test Method D1576 for acceptance testing of commercial shipments, the purchaser and the supplier should conduct comparative tests to determine if there is a statistical bias between their laboratories. Competent statistical assistance is recommended for the investigation of

³ The last approved version of this historical standard is referenced on www.astm.org.

bias. As a minimum, the two parties should take a group of test specimens which are as homogeneous as possible and which are from a lot of material of the type in question. The test specimens should then be randomly assigned in equal numbers to each laboratory for testing. The average results from the two laboratories should be compared using Student's *t*-test for unpaired data and an acceptable probability level chosen by the two parties before testing is begun. If a bias is found, either its cause must be found and corrected or the purchaser and the supplier must agree to interpret future test results in the light of the known bias.

5.2 This test method is a simple and convenient method for routine process control, in-plant evaluation, estimation of moisture content of a lot of wool, or any other purpose for which a high degree of reproducibility is not necessary (see Section 13).

6. Apparatus

6.1 *Oven*, ventilated and thermostatically controlled in the temperature range of $105 \pm 2^\circ\text{C}$ throughout the enclosure. The oven may be of either the forced draft or the convection type.

6.2 *Weighing Containers*, of perforated metal if weighing is to be performed in the drying enclosure; or containers that can be hermetically sealed (such as glass weighing bottles) if the specimen is to be cooled in a desiccator before weighing in the ambient atmosphere.

6.3 *Sampling Containers*, capable of being sealed. Mason jars have been found to be satisfactory where the sample size is not too great. For larger samples, bags of various plastic materials may be suitable if the wall thickness is sufficient to provide a good moisture vapor barrier (at least 4 mil (approximately 0.1 mm) for polyethylene, for example).

6.4 *Balance*, having a capacity adequate for weighing specimens and containers, and a sensitivity of 0.005 g.

7. Sampling

7.1 *Lot Sample*—As a lot sample for acceptance testing, take at random the number of shipping containers directed in applicable material specification or other agreement between the purchaser and the supplier, such as an agreement to use Practice D2525 for bales of fiber and containers of top or sliver or to use Practice D2258 for beams or cases of yarn. Consider shipping containers to be the primary sampling unit.

NOTE 2—An adequate specification or other agreement between the purchaser and supplier requires taking into account the variability between shipping containers, between laboratory sampling units within a shipping container, and test specimens within a laboratory sampling unit to produce a sample plan with a meaningful producer's risk, consumer's risk, acceptable quality level, and limiting quality level.

7.2 Use extreme care to prevent gain or loss of moisture during the sampling operation and the transfer of material to the sampling container. Weigh each portion of the sample and its container immediately after sampling. Subtract the tare mass of the container to obtain the net mass at time of sampling, *M*.

7.3 *Laboratory Sample*—As a laboratory sample for acceptance testing, proceed as follows:

7.3.1 For wool fiber, take laboratory samples as directed in Practice D1060 for cored samples or Practice D3333 for hand samples.

7.3.2 For wool sliver or top, from each shipping container in the lot sample, take one ball of top. From this ball of top, take approximately 2 m from the inside and 4 m from the outside of the ball.

7.3.3 Take laboratory sampling units which weigh a minimum of 50 g. Follow the instructions in Practice D2525 for reduction of the laboratory samples to specimens.

NOTE 3—Condition the laboratory samples as directed in Section 9 before preparing the specimens from them.

8. Number of Specimens

8.1 Take a number of specimens per laboratory sampling unit that the user can expect at the 95 % probability level that the test result for a laboratory sampling unit will be no more than 0.5 percentage points above or below the true average for the laboratory sampling unit. Determine the number of specimens per laboratory sampling unit as follows:

8.1.1 *Reliable estimate of *s**—when there is a reliable estimate of *s* based upon extensive past records in the user's laboratory as directed in the test method, calculate the required number of specimens per laboratory sampling unit using Eq 1:

$$n = (ts/E)^2 \quad (1)$$

where:

- n* = number of specimens per laboratory sampling unit (rounded upward to a whole number),
- s* = reliable estimate of the standard deviation of individual observations on similar materials in the user's laboratory under conditions of single operator precision,
- t* = the value of Student's *t* for two-sided limits, a 95 % probability level, and the degrees of freedom associated with the estimate of *v*, and
- E* = 0.5 percentage points, the allowable variation.

8.1.2 *No Reliable Estimate of *s**—When there is no reliable estimate of *s* for the user's laboratory, do not use Eq 1 directly. Instead, specify the fixed number of six specimens per laboratory sampling unit. This number of specimens per laboratory sampling unit is calculated using *s* = 0.60 percentage points which is a somewhat larger value of *s* than is usually found in practice. When a reliable estimate of *s* for the user's laboratory becomes available, Eq 1 will usually require fewer than six specimens per laboratory sampling unit.

9. Conditioning

9.1 Condition the lot sample (or laboratory sample(s)) by exposure to moving air in the laboratory atmosphere in which the testing is to be done, until equilibrium for testing is achieved.

NOTE 4—Preconditioning and conditioning as directed in Practice D1776 is acceptable but not necessary, since the object of the conditioning for the purpose of this test is merely to stabilize the sample, that is, to bring all parts of the sample to moisture equilibrium with the prevailing atmosphere in order that changes in moisture level will not occur while the specimens are being prepared and weighed.

9.2 Weigh the conditioned sample(s) to the nearest 0.005 g and record the net mass(es), W .

NOTE 5—The net mass of the conditioned sample, W , and the net mass at the time of sampling, M , will be used to convert the observed moisture content of the conditioned specimen to the moisture content at time of sampling.

9.3 From the weighed conditioned sample(s), take the appropriate size specimen(s) and weigh to the nearest 0.005 g to obtain the specimen mass B .

10. Procedure

10.1 Place the specimen(s) in the oven in a suitable container and dry to constant mass, defined as the absence of any progressive decrease in mass in excess of 0.10 % of the average as determined by three successive weighings using the procedure in either 10.1.1 or 10.1.2 to obtain the oven-dry mass of specimen, D .

10.1.1 If the weighings of the dried specimen(s) are to be obtained with the specimen(s) inside the oven, perform the weighings with any forced-air circulation turned off. Space the weighings so that the drying intervals between readings will be equal to 20 % of the normal cycle with a minimum interval of 5 min. Determine the normal cycle by running rate-of-drying curves for similar specimens using the equipment under the same conditions that will be used for ordinary testing. Continue readings of mass until the conditions specified in 10.1 are achieved.

10.1.2 If the weighings of the dried specimen(s) are to be obtained outside the oven, dry the specimen(s) in a container provided with a tight-fitting cover with this cover removed while in the oven. At the end of the drying period, cover the container and remove it from the oven. Place the covered container in a desiccator, loosen the cover, and cool the specimen and container to approximately room temperature. When cooling is completed, set the cover firmly on the container and weigh the container, cover and specimen. Replace the container and specimen in the oven, remove the cover, and dry for an additional 30 % of the normal cycle. Repeat the cooling and weighing procedures. Continue this procedure until the conditions specified in 10.1 are achieved.

11. Calculation

11.1 Calculate to the nearest 0.01 percentage point the percent moisture present in the sample as taken, using Eq 2 for moisture content or Eq 3 for moisture regain.

$$\text{Moisture content, percentage points} \quad (2)$$

$$= [1 - ((W \times D)/(M \times B))] \times 100$$

$$\text{Moisture regain, percentage points} \quad (3)$$

$$= [((M \times B)/(W \times D)) - 1] \times 100$$

where:

M = net mass of subsample at time of sampling,
 W = net mass of subsample at time of measurement,
 B = net mass of specimen before drying, and
 D = oven-dry mass of specimen.

11.2 Calculate the average moisture content (or moisture regain) of all specimens tested for one lot to the nearest 0.1 percentage point.

11.3 The following equations may be used to convert moisture regain in percentage points to moisture content in percentage points and vice versa:

$$R = [C/(100 - C)] \times 100 \quad (4)$$

$$C = [R/(100 + R)] \times 100 \quad (5)$$

where:

R = moisture regain, percentage points, and
 C = moisture content, percentage points.

12. Report

12.1 State that the specimens were tested as directed in Test Method D1576. Describe the material or product sampled and state the method of sampling used.

12.2 Report the following information:

12.2.1 The average value of the results calculated for a particular lot to the nearest 0.1 percentage point, stating whether the reported value is the moisture content or the moisture regain.

12.2.2 The number of specimens tested.

12.2.3 The range of the moisture contents or moisture regains (difference between the largest and smallest observed results).

13. Precision and Bias

13.1 *Interlaboratory Test Data*⁴—An interlaboratory test was carried out in 1963 in which 4 laboratories tested 12 specimens each of a nominally uniform wool material for moisture content. The components of variance expressed as standard deviations were calculated to be:

Within-laboratory component	0.236 percentage point
Between-laboratory component	0.469 percentage point

13.1.1 The within-laboratory component includes the single-operator component which was not determined separately. The components listed above do not include any sampling error. This error must be added in any application of the method.

⁴ Supporting data have been filed at ASTM International Headquarters and may be obtained by requesting Research Report RR:D13-1016.

13.2 *Precision*—For the components of variance reported in 13.1, two averages of observed values should be considered significantly different at the 95 % probability level if the difference equals or exceeds the critical differences listed below:

Number of Observations in Each Average	Critical Differences, Percentage Points, for the Conditions Noted ^{A,B}	
	Within-Laboratory Precision	Between-Laboratory Precision
2	0.46	1.38
3	0.38	1.35
5	0.29	1.33
10	0.21	1.32

^A The values for the critical differences were calculated using $t = 1.960$ which is based on infinite degrees of freedom.

^B The values of the critical differences listed constitutes a general statement particularly with respect to between-laboratory precision. Before a meaningful statement can be made about two specific laboratories, the amount of statistical bias, if any, between them must be established, with each comparison being based on recent data obtained on randomized specimens from one sample of the type of material to be tested.

13.3 *Bias*—The procedure in Test Method D1576 for determination of the amount of moisture present in wool by oven-drying has no bias because the value of that property can be defined only in terms of a test method.

14. Keywords

14.1 moisture content and wool

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