

Standard Test Methods for Determination of the Water Absorption Coefficient by Partial Immersion¹

This standard is issued under the fixed designation C1794; 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 (ε) indicates an editorial change since the last revision or reapproval.

1. Scope

1.1 This test method defines a procedure to determine the water absorption coefficient of a material by partial submersion. The scope is to evaluate the rate of absorption of water due to capillary forces for building materials in contact with normal or driving rain above grade. The procedure is typically suitable mainly for masonry material, plaster, or a coating in combination with a substrate; but it can also be used for insulation materials. This test method is designed to be used only on homogeneous materials and does not apply to materials that are composites or non-homogeneous (for example, Faced Rigid Closed-cell Insulation). It is not within the scope of this standard to determine liquid uptake phenomena in below-grade applications. The water absorption coefficient is mainly used as an input datum for numerical simulation of the combined heat and moisture transport in building envelopes for design and forensic investigation purposes.

1.2 The values stated in either SI units or inch-pound units are to be regarded separately as standard. The values stated in each system may not be exact equivalents; therefore, each system shall be used independently of the other. Combining values from the two systems may result in non-conformance with the standard. However, derived results can be converted from one system to the other using appropriate conversion factors (see Table 1).

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

C168 Terminology Relating to Thermal Insulation

2.2 Other Standards:³

- ISO 9346 Hygrothermal performance of buildings and building materials—Physical quantities for mass transfer— Vocabulary
- ISO 15148 Determination of water absorption coefficient by partial immersion

3. Terminology

3.1 *Definitions of Terms*—For definitions associated with thermal insulation issues refer to Terminology C168. For definitions associated with water absorption refer to the terms and definitions given in ISO 9346.

3.2 Definitions of Terms Specific to This Standard:

3.2.1 *homogeneous material*, *n*—materials, which properties are uniform on a macroscopic scale.

3.2.2 *water absorption coefficient, n*—mass of water absorbed by a test specimen per face area and per square root of time (see Eq 2).

3.3 *Symbols and Units*—the Symbols and Units shown in Table 2 are used.

4. Summary of Test Method and Use

4.1 Water absorption by partial submersion will be calculated by measuring the change in weight of the specimen in a situation when the bottom surface is in contact with liquid water within a time range of at least 24 h. Liquid water that is only adsorbed on the surface shall not be taken into account and shall be removed before weighing, for example., by blotting with a sponge.

4.2 The purpose is of this test is to derive reliable data on the capillary water uptake of building materials in appropriate units using a simple apparatus. These data can also be used as material properties for hygrothermal simulation of the building envelope for design and forensic purposes.

5. Significance and Use

5.1 The purpose of these tests is to obtain, by means of simple apparatus, reliable and easy to determine values of

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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.

³ Available from American National Standards Institute (ANSI), 25 W. 43rd St., 4th Floor, New York, NY 10036, http://www.ansi.org.

TABLE 1 Metric Units and Conversion Factors

Multiply	by	To Obtain (for the same test condition)		
Area				
m ²	10.8	ft ²		
ft ²	9.29 × 10 ⁻² m ²			
Mass				
kg	2.20	lb		
lb	0.454	kg		
Mass per Area				
kg/m ²	0.205	lb/ft ²		
lb/ft ²	4.88	kg/m ²		
Water Absorption Coefficient				
kg/(m²⋅√s)	0.205	lb/(ft²⋅√s)		
lb/(ft²·√s)	4.88	kg/(m²⋅√s)		
$kg/(m^2 \cdot \sqrt{h})$	0.205	lb/(ft ² ·√h)		
$lb/(ft^2 \cdot \sqrt{h})$	4.88	$ka/(m^2 \cdot \sqrt{h})$		

^A The IP unit system includes several meanings of the pound unit lb. In this standard the unit lb refers to pound mass only.

TABLE 2 Symbols and Units^A

Symbol	Quantity	SI unit	IP unit
Α	Face Area	m ²	ft ²
A_{w}	Water Absorption Coefficient	kg/(m²⋅√s)	lb/(ft²⋅√s)
A_{wt}	Water Absorption Coefficient, related	kg/(m²⋅√s)	lb/(ft²⋅√s)
	to a specific time, t, in seconds		
Δm_t	mass gain per face area after time t	kg/m ²	lb/ft ²
M_i	Initial mass of specimen	kg	lb
m_t	Mass of specimen after time t	kg	lb
m _i	Starting mass	kg	lb
t	Time	s or h	h
W_{w}	Water Absorption Coefficient	kg/(m²⋅√h)	lb/(ft²⋅√h)
W_{wt}	Water Absorption Coefficient, related	kg/(m²⋅√h)	lb/(ft²⋅√h)
	to a specific time, t, in hours		

^A The IP unit system includes several meanings of the pound unit lb. In this standard the unit lb refers to pound mass only.

liquid water transport for capillary active materials expressed in suitable units. These values are for use as part of the material properties in hygrothermal analysis tools for building envelope design and forensic studies. As the topic of liquid transport phenomena in porous materials is very complex, Appendix X1 in ISO 15148 shows some more detailed background information.

6. Apparatus

6.1 The apparatus must contain the following:

6.1.1 A scale to measure the weight of the specimen with an accuracy within ± 0.1 % of the specimen's mass.

6.1.2 A water tank with a regulation system to keep the water level constant within $\pm 3 \text{ mm} (\pm \frac{1}{8} \text{ in.})$ and equipment to keep the position of the specimen at least 5 mm ($\frac{1}{4} \text{ in.}$) above the bottom of the tank without harming the specimen.

6.1.3 Equipment to measure the time with accuracy of at least 1 second.

7. Test Specimen

7.1 *Shape*—The shape shall represent typical material dimensions and must have a constant area to ensure onedimensional moisture flow. The surfaces shall be as flat as possible.

7.2 *Area*—The area which is in contact with the water must be at least 50 cm² (8 in.²). However, in the case of materials

including macroscopic particles such as aggregates, the side of a square specimen or the smallest diameter of the face shall be at least ten times the largest particle size. Influences of irregular surface structures shall be neglected if those irregular surface structures are part of the materials' design.

Note 1—Larger specimens, preferably with a face area of at least 100 $\rm cm^2$ (16 in.²), are advised as they will lead to greater accuracy.

7.3 *Thickness*—Where possible, use a specimen thickness which represents the full product thickness. When specimens are cut from products they shall be representative of the material to be assessed and thick enough to enable handling without damage. In the case of materials including macroscopic particles such as aggregates, it is preferable that the thickness be at least ten times the largest particle size, but shall be no less than five times, the largest particle size.

7.4 *Number*—At least three specimens shall be tested. If the water contact area of the individual specimens is less than 100 cm² (16 in.²), at least six specimens shall be tested representing a total area of at least 300 cm² (47 in²).

7.5 Preparation—Test specimens shall be representative of the whole material and shall be cut so that they do not include product edges. In the case of materials known to be nonisotropic, sets of test specimens shall be prepared in all orientations of the potential use of the material. The test specimens shall be prepared by methods that do not change the original structure of the product; any skins, facings or coatings shall be retained. In the case of products such as coatings, thin rendering or plasterwork that are normally adhered to a substrate in use, specimens shall be made up from the product and a normal substrate combined. The total thickness then is the sum of the coating and the substrate. The sides of a solid specimen shall be sealed with a water and vapor tight sealant that does not react chemically with it or significantly penetrate the pores of the product as shown in Fig. 1. It is especially important that the sides of specimens with surface coatings are sealed to prevent bypassing of the coating. If sealing is not possible in the case of very low density fibrous or loose fill materials place them in a tightly fitting tube supported on a wire mesh placed over the mouth of the tube. The open area of the mesh shall be as large as possible while completely supporting the sample during the whole course of the test. In this case, to minimize the edge effects, the face area of the specimen shall be at least 100 cm^2 (16 in²). The surface in contact with the water shall be plane, allowing for the normal surface roughness of the material.

7.6 Conditioning—The test specimens shall be stored under the test conditions (see 8.1) until the mass of each specimen has stabilized to within 0.1 % of its total mass, when measured over 24 h. In case the stability of 0.1 % of mass cannot be reached, additionally the relative humidity shall be controlled, for example, by using a climatic-controlled room or chamber, to be within 50 % \pm 5 % and furthermore the specimen must be exposed to these conditions until the end of the measurement except of the short time of the weighing process.

8. Procedure

8.1 The procedure must be conducted within the conditions stated in Table 3.



FIG. 1 Example of a Suitable Testing Apparatus

TABLE 3 Test Conditions

	Temperature
Allowed range of test conditions	18 to 23 °C (64 to 73 °F)
Allowed variation during test	± 2 °C (± 4 °F)

8.2 Weigh each specimen at steady state with an accuracy of ± 0.1 % of mass to determine the initial mass m_i after conditioning.

8.3 Fill the tank with water to the specified level stated in step 8.4. Wait until the temperature of the water has reached the specimen temperature.

8.4 Put the specimen into the tank and ensure a distance between the specimen and the tank bottom of at least 5 mm (1/4 in.) by using distance pins as shown in Fig. 1. The pins shall not affect the liquid uptake at the specimen surface. It must be ensured that no air pockets are trapped between the surface and the water. If necessary, place a weight on the top of the specimen. The water level shall be kept at a constant 5 ± 2.5 mm (0.2 \pm 0.1 in.) above the highest point of the absorbing surface of the specimen.

8.5 If a weight is used to fix the specimen's position, it must be ensured that the specimen is not mechanically influenced by the weight.

8.6 To avoid anisotropic effects half the specimens of a homogeneous material cut from the same sample must be placed with one major face downwards. The other half with this must face upwards. Specimens of non-homogeneous materials shall be placed with the face which is normally exposed to driving rain or other expected water sources downwards. This is especially important in the case of renders and coatings which are tested on their substrate.

8.7 Start the time measurement precisely when the specimen first contacts the liquid water.

8.8 After approximately 5 min, remove the specimen from the water and remove surface-adhered water drops using a

moist sponge that has been squeezed out beforehand. Weigh the specimen with an accuracy of ± 0.1 % of the mass, m_i , at time *t*. Put the specimen back into the water. Repeat the measurement at least two more times within 24 h to derive a 1 h, 2 h, 4 h, 8 h series of measurements for m_i at time *t*. Be sure to take a measurement at 4 h.

Note 2—The measurement shall be accomplished as quickly as possible and the specimen returned to the water immediately afterwards. A time of less than a minute expected to be appropriate.

8.9 The method of evaluation depends on the shape of the curve stated in Section 9, and the shape of the curve depends almost entirely on how drying is handled. Conduct the calculations and plots of the curve immediately after the weighing to determine the end of the weighing process.

Note 3—Some materials can cause some nonlinear effects especially at the beginning.

8.10 If liquid water is visible on the top surface of the specimen, the measurement shall be terminated.

8.11 If the liquid uptake is less than 0.001 kg/m² (0.0002 lb/ft²) of the absorbing surface, the measurement can be terminated and the material can be specified as resistant against liquid water uptake.

8.12 After termination of the measurement the specimen shall be investigated for cracking of the sealant which was applied to the specimen in 7.5.

9. Calculation and Expression of Results

9.1 Calculate the area A in m^2 (ft²) perpendicular to the direction of flow of the water. In the case of a low-density fiber or loose fill material, the area of the tube shall be used.

9.2 Calculate the difference between the area-related mass at each weighing according to the equation:

$$\Delta m_t = \frac{m_t - m_i}{A} \tag{1}$$

and plot these against the square root of the time \sqrt{t} .

9.3 The graph resulting from step 8.2 will be one of two different types (type A or B) shown in Figs. 2 and 3. It must be distinguished whether liquid water was seen on the top surface of the specimen after the test was terminated.

9.4 *Type A Graph*—As shown in Fig. 2 left, type A graphs show a linear relationship until the end of the test; a regression curve can be used through the points of Δm_t against \sqrt{t} . Extend the regression curve to the vertical axes to determine $\Delta m'_0$. The water absorption coefficient A_w and W_w will then be calculated from:

$$A_{w} = \frac{\Delta m'_{tf} - \Delta m'_{0}}{\sqrt{t_{f}}}$$
(2)

where:

 $\Delta m'_{tf}$ = value of Δm on the regression curve at time t_{f} , in kg/m² (lb/ft²), and

 t_f = duration of the test (typically 1 day) in seconds.

or

$$W_{w} = \frac{\Delta m'_{tf} - \Delta m'_{0}}{\sqrt{t_{f}}}$$
(3)

where:

 t_f = the duration of the test (typically 1 day) in hours.

9.4.1 If the graph shows a slope with a sudden decrease of the slope, as can be seen at right in Fig. 2, the water front inside the specimen has most likely reached the top surface. In this case, the points after the sudden decrease of the slope shall be ignored.

9.5 Type B Graph—As shown in Fig. 3, type B graphs show a nonlinear relationship between Δm_t and t. In this case, the measured value of Δm_t at 4 h ($2\sqrt{h}$) shall be used for Δm_{tf} and A_w or W_w shall be calculated according to:

$$A_{w,4} = \frac{\Delta m'_{tf} - \Delta m'_{0}}{\sqrt{14400s}}$$
(4)

$$W_{w,4} = \frac{\Delta m'_{tf} - \Delta m'_0}{2\sqrt{h}} \tag{5}$$

9.5.1 If liquid water has reached the top surface of the specimen before the 4 h measurement, the time shall be noted and no value for A_w or W_w shall be reported.

10. Report

10.1 The report must contain:

10.1.1 Reference to this standard.

10.1.2 Product identification:

10.1.2.1 Name of manufacturer or distributor,

10.1.2.2 Type of product according to manufacturer specification,

10.1.2.3 Product code number, if applicable,

10.1.2.4 Packing,

10.1.2.5 Condition in which the specimen arrived in the laboratory, and

10.1.2.6 Nominal physical properties, such as bulk density, thickness, etc.

10.1.3 Test procedure, including the following:

10.1.3.1 Date and duration of test,

10.1.3.2 Method of sampling and preparation,

10.1.3.3 Temperture during the test and whether a climatecontrolled chamber was used,

10.1.3.4 Immersion depth if greater than 7 mm ($\frac{1}{4}$ in.),

10.1.3.5 Type of result graph (type A or B), and

10.1.3.6 Statement whether cracks in the coating were detected.

10.1.4 Test results:

10.1.4.1 All values for A_w and W_w in a type A graph, or A_{w4} and W_{w4} in a type B graph, and

10.1.4.2 Graphs of Δm_t versus \sqrt{t} , including regression curves for all specimens.

11. Precision and Bias

11.1 At this time, no measurement data are available for precision and bias. The plan is to complete a round robin within 2 years after this standard is accepted.

11.2 Three parameters measured during this test are:



FIG. 2 Type A Graphs without Liquid Water on the Top Surface (left) and with Liquid Water on the Top Surface (right)

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11.2.1 Record the time at which the specimen is removed for weighing to the nearest minute, which over the time scale of about 24 h gives an accuracy of better than 0.1 %.

11.2.2 The mass of the specimen. The mass, as specified in step 7.2, shall be recorded with an accuracy of ± 0.1 % of the mass of the specimen. However, it is possible that problems in handling and drying the specimen before weighing will introduce substantially larger inaccuracies. It is recommended that the values of mt be calculated and plotted, as specified in Section 8, after each weighing so that the quality of the data can be assessed, and that additional weighings be carried out if better accuracy is required.

11.2.3 The area of the specimen. Given a regular specimen, it will be possible to measure the area to within 1 %. It will be more difficult to measure the area of irregular specimens as precisely. If better accuracy is needed, additional specimens can be tested.

12. Keywords

12.1 hygrothermal simulation; hygrothermics; liquid transport coefficient; liquid water uptake; material properties; partial immersion; water absorption

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