

Designation: D 635 - 98

Standard Test Method for Rate of Burning and/or Extent and Time of Burning of Plastics in a Horizontal Position¹

This standard is issued under the fixed designation D 635; 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 fire-test-response test method covers a small-scale laboratory screening procedure for comparing the relative linear rate of burning or extent and time of burning, or both, of plastics in the form of bars, molded or cut from sheets, plates, or panels, and tested in the horizontal position.

Note 1—This test method, Clause 8 of IEC Draft 695-11-10 and Method A of ISO 1210-1992 are technically equivalent.

Note 2—For additional information on materials which do not burn to the first reference mark by this test, see Test Method D 3801.

- 1.2 This test method was developed for polymeric materials used for parts in devices and appliances. The results are intended to serve as a preliminary indication of their acceptability with respect to flammability for a particular application. The final acceptance of the material is dependent upon its use in complete equipment that conforms with the standard applicable to such equipment.
- 1.3 The classification system described in the appendix is intended for quality assurance and the preselection of component materials for products.
- 1.4 The test method is not intended to cover plastics when used as materials for building construction or furnishing.
- 1.5 This test method should be used to measure and describe the response of materials, products, or assemblies to heat and flame under controlled laboratory conditions and should not be used to describe or appraise the fire hazard for fire risk of materials, products, or assemblies under actual fire conditions. However, results of this test method may be used as elements of a fire risk assessment that takes into account all of the factors that are pertinent to an assessment of the fire hazard of a particular end use.
- 1.6 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. For specific hazards statements, see 9.2.1.

2. Referenced Documents

- 2.1 ASTM Standards:
- D 648 Test Method for Deflection Temperature of Plastics Under Flexural Load²
- D 883 Terminology Relating to Plastics²
- D 3801 Test Method for Measuring the Comparative Burning Characteristics of Solid Plastics in a Vertical Position³
- D 5025 Specification for a Laboratory Burner Used for Small-Scale Burning Tests on Plastic Materials⁴
- D 5207 Practice for Calibration of 20 and 125-mm Test Flames for Small-Scale Burning Tests on Plastic Materials⁴
- E 176 Terminology of Fire Standards⁵
- E 691 Practice for Conducting an Interlaboratory Study to Determine the Precision of a Test Method⁶
- 2.2 IEC Standards:
- IEC 695-11-10 Determination of the Burning Behavior of Horizontal and Vertical Specimens in Contact with a 50 W Nominal Ignition Source⁷
- 2.3 ISO Standards:
- ISO 1210-1992 Plastics—Determination of the Burning Behavior of Horizontal and Vertical Specimens in Contact with a Small-Flame Ignition Source⁷

3. Terminology

- 3.1 Definitions:
- 3.1.1 Definitions used in this test method are in accordance with Terminology D 883, unless otherwise specified. For terms relating to fire, the definitions used in this test method are in accordance with Terminology E 176.

4. Summary of Test Method

4.1 A bar specimen of the material to be tested is supported horizontally at one end. The free end is exposed to a specified gas flame for 30 s. Time and extent of burning are measured

 $^{^{1}}$ This test method is under the jurisdiction of ASTM Committee D-20 on Plastics and is the direct responsibility of Subcommittee D20.30 on Thermal Properties (Section D20.30.03).

Current edition approved July 10, 1998. Published January 1999. Originally published as D 635-41. Last previous edition D 635-97.

² Annual Book of ASTM Standards, Vol 08.01.

³ Annual Book of ASTM Standards, Vol 08.02.

⁴ Annual Book of ASTM Standards, Vol 08.03.

⁵ Annual Book of ASTM Standards, Vol 04.07.

⁶ Annual Book of ASTM Standards, Vol 14.02.

⁷ Publications of the International Electrotechnical Commission (IEC) and International Organization for Standardization (ISO) are available from American National Standards Institute, 11 W. 42nd St., New York, NY 10036.



and reported if the specimen does not burn 100 mm. An average burning rate is reported for a material if it burns to the 100 mm mark from the ignited end.

5. Significance and Use

- 5.1 Tests made on a material under conditions herein prescribed are of value in comparing the rate of burning or extent and time of burning characteristics, or both, of different materials, in controlling manufacturing processes, or as a measure of deterioration or change in these burning characteristics prior to or during use. Correlation with flammability under actual use conditions is not implied.
- 5.2 The rate of burning and other burning phenomena will be affected by such factors as density, pigments, any anisotropy of the material and the thickness of the specimen. Test data shall be compared only for specimens of similar thickness, whether comparisons are being made with the same or different materials. The rate of burning and other burning phenomena will vary with thickness.
- 5.3 It is feasible that sheet materials that have been stretched during processing will relax during burning and give erratic results unless they are first heated above their deflection temperature, in accordance with Test Method D 648, for a time sufficient to permit complete relaxation.
- 5.4 Burning tests require that certain variables be arbitrarily fixed, for example, specimen size, energy source and application time, and end points. Materials will be found that are unusually sensitive to one or more of the conditions chosen for this method leading to highly variable results. Additional burning characterization by other methods is highly desirable in such cases (see Note 2).
- 5.5 In this procedure, the specimens are subjected to specific laboratory test conditions. If different test conditions are substituted or the end-use conditions are changed, it will not always be possible by or from this test to predict changes in the fire-test-response characteristics measured. Therefore, the results are valid only for the fire-test-exposure conditions described in this procedure.

6. Apparatus

6.1 Test Chamber, enclosed laboratory hood, or chamber free of induced or forced draft during test, having an inside

volume of at least 0.5 m³. An enclosed laboratory hood with a heat-resistant glass window for observing the test and an exhaust fan for removing the products of combustion after the tests is recommended. The atmosphere in and around the test chamber shall be maintained between 15 to 35°C and 45 to 75 % relative humidity.

Note 3—The amount of oxygen available to support combustion is naturally important for the conduct of these fire-test-response tests. For tests conducted by this test method when burning times are protracted, chamber sizes less than $1\ m^3$ may not provide accurate results.

Note 4—Some laboratory hoods have induced drafts even with the exhaust fan off. A positive-closing damper is recommended.

Note 5—A mirror in the chamber, to provide a rear view of the specimen, has been found useful in some enclosures.

6.2 Test Fixture, A laboratory ring stand or test fixture equipped with a means of holding a 125 mm² wire gauze horizontal and a small clamp permitting the specimen to be held with its longitudinal axis horizontal and its transverse axis inclined at $45 \pm 2^{\circ}$ angle as illustrated in Fig. 1.

Note 6—A pan of water may be placed on the floor of the hood in position to catch any burning particles that may drop during the test.

- 6.3 *Laboratory Burner*, constructed in accordance with Specification D 5025.
- 6.4 Gas Supply, a supply of technical-grade methane gas with suitable regulator and meter for uniform gas flow. Natural gas mixtures having an energy density of approximately 37 MJ/m³ have been found to provide similar results. However, technical-grade methane gas shall be used as the referee in cases of dispute.
- 6.5 Wire Gauze, 20-mesh (approximately 20 openings per 25 mm), made with 0.43 ± 0.03 mm diameter iron wire cut to approximately 125 mm², to sustain burning or glowing particles falling from the specimens.
 - 6.6 Timing Device, accurate to 0.5 s. 64/astm-d635-98
 - 6.7 Scale, graduated in millimeters.
 - 6.8 Micrometer, accurate to 0.05 mm.
- 6.9 Conditioning Room or Chamber, capable of being maintained at 23 \pm 2°C and 50 \pm 5 % relative humidity.
- 6.10 Flexible Specimen Support Fixture, used to facilitate the testing of specimens that sag and touch the wire gauze. (See 9.4 and Fig. 2.)

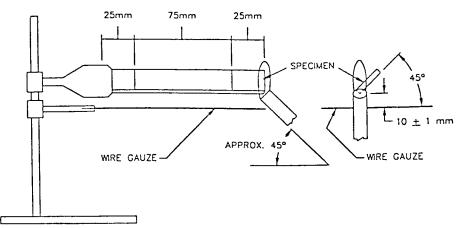


FIG. 1 Test Fixture



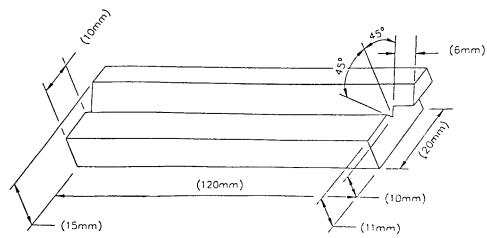


FIG. 2 Flexible Specimen Support Fixture

7. Test Specimens

7.1 All test specimens shall be cut from a representative sample of the material (sheet or end products), or shall be cast or injection-, compression-, transfer- or pultrusion-molded to the necessary form. After any cutting operation, care shall be taken to remove all dust and any particles from the surface; cut edges shall be fine sanded to have a smooth finish. Unless otherwise agreed, fabrication of test specimens shall be in accordance with the specifications of the material being tested.

7.2 Specimens shall be 125 ± 5 mm long by 13.0 ± 0.5 mm wide, and provided in the minimum thickness and in the 3.0 (-0.0 + 0.2) mm thickness. The 3.0 mm thick specimens are not necessary if the minimum thickness is greater than 3.0 mm, or the maximum thickness is less than 3.0 mm. The maximum thickness shall not exceed 13 mm. The maximum width shall not exceed 13.5 mm. The edges shall be smooth, and the radius on the corners shall not exceed 1.3 mm.

8. Conditioning

8.1 Condition ten bar specimens for each material and thickness to be tested in accordance with Test Method D 618 at $23\pm2^{\circ}\text{C}$ and 50 ± 5 % relative humidity for a minimum of 48 h. Once removed from the conditioning atmosphere test the specimens within 1 h.

8.2 Conduct testing in a laboratory atmosphere of 15 to 35°C and 45 to 75 % relative humidity.

9. Procedure

- 9.1 Prepare at least ten bar specimens. After measuring and recording the specimen thickness, mark each specimen with two lines perpendicular to the longitudinal axis of the bar, 25 ± 1 and 100 ± 1 mm from the end that is to be ignited.
- 9.2 Conduct the burning test in a chamber, enclosure, or laboratory hood free of induced or forced draft.

Note 7—Warning: Products of combustion may be toxic. An enclosed laboratory hood and an exhaust fan for removing the products of combustion after the tests are recommended. The exhaust fan is turned off during the test and turned on immediately following the test in order to remove products of combustion.

9.3 Clamp the specimen at the end farthest from the 25 mm reference mark, in a support with its longitudinal axis horizon-

tal and its transverse axis inclined at $45 \pm 2^{\circ}$ as illustrated in Fig. 1. Clamp the wire gauze horizontally beneath the specimen, with a distance of 10 ± 1 mm between the lowest edge of the specimen and the wire gauze, and with the free end of the specimen even with the edge of the gauze. Any material remaining on the wire gauze from the previous test must be burned off or a new section of wire gauze used for each test.

9.4 If the specimen sags at its free end during the initial set up and is not able to maintain the distance of 10 ± 1 mm as specified in 9.2, the flexible specimen support fixture illustrated in Fig. 2 shall be used. Position the support fixture under the specimen with the small extending portion of the support fixture at least 20 mm from the free end of the specimen. Provide enough clearance at the clamped end of the specimen so that the support fixture can be moved freely sidewards. As the flame front progresses along the specimen, withdraw the support fixture at the same approximate rate, preventing the flame front from contacting the flexible specimen support fixture, so that there is no effect on the test flame or on the burning of the specimen.

9.5 With the central axis of the burner tuber in the vertical position, place the burner remote from the specimen, ignite, and adjust it to produce a blue flame 20 mm high. Adjust the gas supply and the air ports of the burner until a 20-mm yellow-tipped blue flame is produced, and then increase the air supply until the yellow tip just disappears. Measure the height of the flame. If the flame height is not 20 ± 2 mm, adjust the burner gas supply to give the proper flame height. Once the flame has been properly set to a height of 20 ± 2 mm wait for at least 5 min to allow the burner conditions to reach equilibrium.

Note 8—See Practice D 5207 for recommended back pressure and flow rate for the gas supply and calibration procedure for the 20 mm flame.

9.6 Place the burner so that the test flame impinges on the free end of the test specimen to a depth of approximately 6 mm starting the timing device simultaneously. The central axis of the burner tube is to be in the same vertical plane as the longitudinal bottom edge of the specimen and inclined toward the end of the specimen at an angle of approximately 45 ± 2 degrees to the horizontal. See Fig. 1. Apply the flame for 30 ± 1 s without changing its position. If the test specimen