



Designation: D 1603 – 01

Standard Test Method for Carbon Black In Olefin Plastics¹

This standard is issued under the fixed designation D 1603; 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 the determination of the carbon black content in polyethylene, polypropylene, and polybutylene plastics. Its use with acrylic or other polar monomer modifications which might affect the accuracy is not recommended. Determinations of carbon black are made gravimetrically after pyrolysis of the sample under nitrogen. This test method is not applicable to compositions that contain nonvolatile pigments or fillers other than carbon black.

1.1.1 This test method is not applicable to materials containing brominated flame retardant additives at the end.

1.2 The values stated in SI units are to be regarded as the standard. The values in parentheses are given for information only.

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.*

NOTE 1—This test method is similar to ISO 6964-1986(E) in title only. The technical content is significantly different.

2. Referenced Documents

2.1 ASTM Standards:

D 883 Terminology Relating to Plastics²

E 177 Practice for Use of the Terms Precision and Bias in ASTM Test Methods³

2.2 ISO Standard:

ISO 6964-1986(E) Polyolefin Pipes and Fittings—Determination of Carbon Black by Calcination and Pyrolysis—Test Method and Basic Specification⁴

¹ This test method is under the jurisdiction of ASTM Committee D20 on Plastics and is the direct responsibility of Subcommittee D20.70 on Analytical Methods (Section D20.70.01).

In 1993, the scope was revised to clarify limitations related to use of this test method with materials containing brominated additives and to include an ISO equivalency statement. Keywords were also added.

Current edition approved March 10, 2001. Published June 2001. Originally published as D 1603 – 58. Last previous edition D 1603 – 94.

² Annual Book of ASTM Standards, Vol 08.01.

³ Annual Book of ASTM Standards, Vol 14.02.

⁴ ISO/IEC Selected Standards for Testing Plastics, Second Edition, published by ASTM. Also, available from American National Standards Institute, 11 W. 42nd St., 13th Floor, New York, NY 10036.

3. Terminology

3.1 *Definitions*—For definitions of technical terms pertaining to plastics used in this specification, see Terminology D 883.

4. Significance and Use

4.1 The information provided by this test method is useful for control purposes and is required for calculation of optical absorptivity.

5. Apparatus

5.1 *Electric Furnace*, at least 20 cm (7.9 in.) long suitable for use with the tubing described in 5.2.

5.2 *High Temperature Glass Combustion Tube*,⁵ of appropriate diameter and approximately twice as long as the furnace described in 5.1.

5.3 *Stoppers*—Two rubber or neoprene stoppers, to fit the tube described in 5.2, unless the tube is fitted with ground joints and mating connectors.

5.4 *Glass Tubing*, approximately 10 mm (0.39 in.) in diameter, of sufficient amount, and matching rubber or plastic tubing for connections.

5.5 *Combustion Boat*, approximately 8 by 1.9 by 1.3 cm (3.15 by 0.75 by 0.51 in.). Glazed porcelain, quartz high-silica glass, or platinum is suitable.

NOTE 2—A loose-fitting cover for the combustion boat is optional. If used, it shall be considered a part of the boat and handled and weighed with it.

5.6 *Iron-Constantan Thermocouple*, and a potentiometer or millivoltmeter suitable for determining temperatures in the range 300 to 700°C (572 to 1292°F).

5.7 *Flow Meter*,⁶ suitable for measuring gas flow at rates of 1 to 10 L/min.

5.8 *Traps*, three glass traps with removable ground-glass connected heads and 10-mm (0.39-in.) diameter inner and connecting tubes.

NOTE 3—Only one trap is required if the entire apparatus train is placed in a fume hood. None is required if in addition, nitrogen of sufficient

⁵ Pyrex, Vycor, or equivalent tubing has been found satisfactory for this purpose.

⁶ Precision Bore Rotometer Tube No. 2B-25, available from the Fischer & Porter Co., County Line Road, Warminster, PA 18974 has been found satisfactory.

purity is used and produced by the alternative means provided in Section 6.

5.9 *Drying Tube*—A U-shaped drying tube, having an inside diameter of 20 mm (0.79 in.) or larger, fitted with ground glass or neoprene stoppers.

5.10 *Glass Wool*.

5.11 *Desiccator*, with desiccant.

5.12 *Bunsen Burner*.

5.13 *Balance*—An analytical balance having a sensitivity of 0.0001 g.

5.14 *Weights*—A set of Class S weights for use with the balance (if required).

6. Reagents and Materials

6.1 Carbon Dioxide, Solid (Dry Ice).

NOTE 4—The solid carbon dioxide and the trichloroethylene are not required if the entire apparatus train is placed in a fume hood.

6.2 *Desiccant*, such as anhydrous calcium chloride (CaCl₂).

6.3 *Nitrogen*, prepurified, having oxygen content below 0.01 %. As a safeguard against accidental leakage, contamination, or inadequate purity, the gas shall be further purified by one of the following procedures:

6.3.1 Passage of the nitrogen through a glass trap inserted ahead of the drying tube (see Fig. 1), filled approximately one third full of potassium hydroxide - pyrogallol solution made to contain 5 g of pyrogallol and 50 g of KOH in 100 mL of water. Technical grade, or better, reagents are satisfactory.

6.3.2 Insertion of a plug, or roll, of clean copper tinsel, foil, or wire 7.5 to 10 cm (3 to 4 in.) long into the combustion tube ahead of the sample (see Fig. 1) so that it is completely within the heated region of the furnace. Take care to prevent channeling of the nitrogen through the plug. The extent of blackening of the copper may be taken as a guide for determining when the plug should be renewed.

6.3.3 Passage of the nitrogen through a combustion tube filled to a length of 15 cm (6 in.) or greater with clean copper

tinsel, foil, or wire, and maintained in a furnace at a temperature around 500°C (932°F).

6.3.4 The need for the procedures described is eliminated if gas having an oxygen content of less than 0.002 % (20 ppm) is used.

6.4 *Trichloroethylene*, technical grade (Note 4).

7. Procedure

NOTE 5—The procedure below assumes that the combustion tube can be easily removed from the furnace. If this is not the case, alternate methods of inserting and removing sample boats are acceptable as long as the temperature, purge time, and flow rate requirements are met.

7.1 Assemble the apparatus as shown in Fig. 1. Both cold traps following the combustion tube shall contain trichloroethylene, but only the first need be cooled with solid carbon dioxide. Alternatively, the entire apparatus may be placed in a fume hood and the two traps following the combustion tube omitted. Fill the drying tube with anhydrous CaCl₂ or other suitable desiccant. Hold between loose plugs of glass wool.

7.2 Heat a clean combustion boat to red heat in a bunsen flame; then transfer the boat to the desiccator and allow it to cool over fresh desiccant for not less than 30 min.

7.3 Remove the boat from the desiccator and weigh it to nearest 0.0001 g (w_1). Immediately place 1.0 ± 0.1 g of the ethylene plastic under test in the boat and quickly weigh to the nearest 0.0001 g (w_2).

7.4 Heat the furnace to a constant temperature of 600°C (1112°F).

7.5 With the combustion tube removed from the furnace, adjust the rate of nitrogen flow through the tube to 1.7 ± 0.3 L/min. Open the inlet end of the combustion tube, quickly place the combustion boat with the sample into the tube positioned so the boat will be at the proper temperature when in the furnace. Close the inlet to the tube and allow the nitrogen to flow for a minimum of 5 min to purge oxygen from the system prior to placing the tube in the furnace. If the furnace is

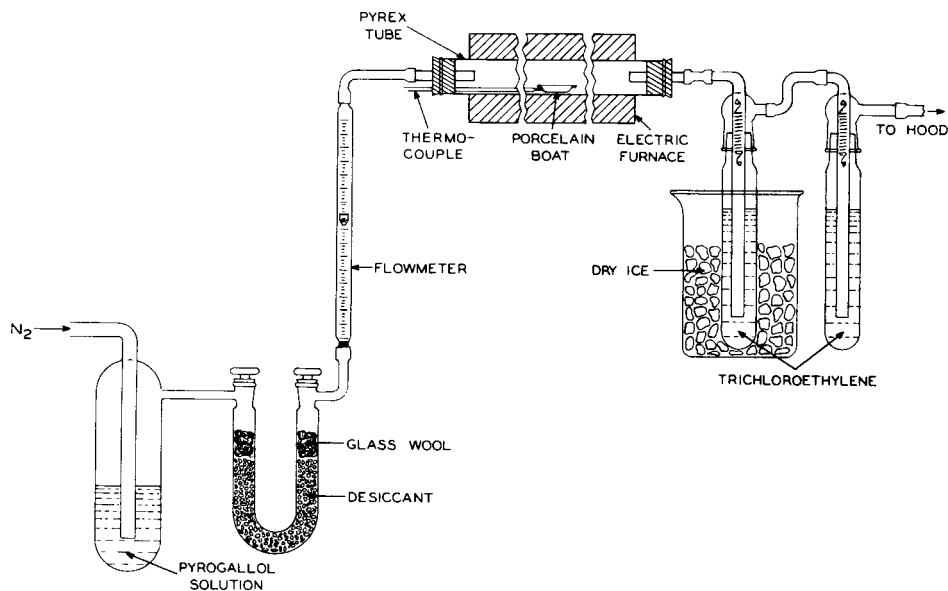


FIG. 1 Assembly of Apparatus