



# Standard Test Method for Thermal-Oxidative Stability of Polypropylene Using a Specimen Rotator Within an Oven<sup>1</sup>

This standard is issued under the fixed designation D 3012; 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 provides a means for estimating the resistance of polypropylene, in molded form, to accelerated aging by heat in the presence of air using a forced draft oven.

1.2 The stability determined by this test method is not directly related to the suitability of the material for use when different environmental conditions prevail and shall not be used to predict performance.

NOTE 1—The specified thermal levels in this test method are considered sufficiently severe to cause failure of commercial grades of heat-stable polypropylene within a reasonable period of time. If desired, lower temperatures can be applied to estimate the performance of polypropylene with lower heat stabilities.

1.3 The values stated in SI units are to be regarded as the standard. The values in brackets are for information only.

1.4 *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 2—This test method and ISO 4577–1983 are technically similar but different in preparation of test specimens, thickness of test specimen, measurement of the number of air flow changes in the ovens, and the number of air changes per hour required.

## 2. Referenced Documents

### 2.1 ASTM Standards:<sup>2</sup>

- D 618 Practice for Conditioning Plastics and Electrical Insulating Materials for Testing
- D 883 Terminology Relating to Plastics
- D 3641 Practice for Injection Molding Test Specimens of Thermoplastic Molding and Extrusion Materials

<sup>1</sup> This test method is under the jurisdiction of ASTM Committee D20 on Plastics and is the direct responsibility of Subcommittee D20.15 on Thermoplastic Materials.

Current edition approved March 1, 2004. Published April 2004. Originally approved in 1972. Last previous edition approved in 2000 as D 3012 - 00.

<sup>2</sup> For referenced ASTM standards, visit the ASTM website, [www.astm.org](http://www.astm.org), or contact ASTM Customer Service at [service@astm.org](mailto:service@astm.org). For *Annual Book of ASTM Standards* volume information, refer to the standard's Document Summary page on the ASTM website.

- D 4101 Specification for Polypropylene Injection and Extrusion Materials
- D 5374 Test Methods for Forced-Convection Laboratory Ovens for Evaluation of Electrical Insulation
- E 1 Specification for ASTM Thermometers
- E 77 Test Method for Inspection and Verification of Thermometers
- E 220 Test Method for Calibration of Thermocouples by Comparison Techniques
- E 608 Specification for Metal-Sheathed Base-Metal Thermocouples
- E 644 Test Methods for Testing Industrial Resistance Thermometers
- E 691 Practice for Conducting an Interlaboratory Study to Determine the Precision of a Test Method
- E 1137 Specification for Industrial Platinum Resistance Thermometers
- 2.2 *ISO Standard:*<sup>3</sup>
  - ISO 4577–1983 Plastics—Polypropylene and Propylene Copolymers—Determination of Thermal Oxidative Stability in Air-Oven Method
  - ISO 1873 Plastics—Polypropylene and Propylene-Copolymer Thermoplastics:
    - Part 1: Designation
    - Part 2: Determination of Properties

## 3. Terminology

3.1 *Definitions*—The definitions of plastics used in this test method are in accordance with Terminology D 883 unless otherwise indicated.

## 4. Summary of Test Method

4.1 Aging is accelerated by exposing the specimens to an elevated temperature in a forced draft oven equipped with a biaxial or uniaxial rotating specimen holder.

4.2 Visual examination is used to determine the time to failure. The time to failure of the material is taken as the

<sup>3</sup> Available from American National Standards Institute (ANSI), 25 W. 43rd St., 4th Floor, New York, NY 10036.

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

number of days after which the specimen shows localized crazing, crumbling, or discoloration, or a combination thereof.

## 5. Significance and Use

5.1 Under the severe conditions of this test method, the specimens undergo degradation at a rate that is dependent upon the thermal endurance of the polypropylene material under examination.

5.2 The thermal level of this test method is considered sufficiently severe to cause failure of commercial grades of heat-stable polypropylene within a reasonable period of time. If desired, lower temperatures can be applied to estimate the performance of polypropylene materials with lower heat stability.

5.3 The technique of specimen rotation described in this test method provides an estimate of the life-temperature relationship of polypropylene. If this test method is conducted at different temperatures on the same material, a more reliable estimate of the life-temperature relationship of polypropylene is determined. This test method can be conducted at several temperatures and the data interpreted through use of the Arrhenius relation, by plotting the logarithms of times to failure against the reciprocals of the temperatures in kelvins (K). Temperatures in the range from 100 to 150°C, with intervals of 10°C, are suggested for this purpose.

5.4 The stability as determined under the prescribed test method is not directly related to the suitability of the compound for a use where different conditions prevail.

5.5 The specimen rotation technique of thermal aging increases the probability that all specimens will be exposed similarly and that the effect of temperature gradients in an oven will be minimized.

## 6. Apparatus

6.1 *Oven*, mechanical convection type for controlled circulation of air, with adjustable air intake and exhaust facilities, and designed for air velocities around  $\pm 250$  1000 mm/s [ $197 \pm 49$  ft/min].<sup>4</sup>

6.1.1 The oven shall be equipped with a temperature-control system designed to maintain the test temperature range from  $150 \pm 1^\circ\text{C}$  [ $302 \pm 1.8^\circ\text{F}$ ] and a device to prevent temperature override. With the oven adjusted to the nominal test temperature of 150°C, the override shall be set at 154°C [ $309^\circ\text{F}$ ]. A bimetallic-strip temperature switch has been found satisfactory.

6.2 *Oven Temperature Measurement System*, consisting of a thermocouple, thermometer, or resistance thermometer as the sensor, together with its associated conditions and readout instrumentation covering at least the temperature range from 0 to 200°C [ $32$  to  $372^\circ\text{F}$ ].

6.2.1 The thermometer must cover the range in one-degree subdivisions. It must be tested for bulb stability and standardized, in accordance with Test Method E 77.

6.2.2 The secondary standard shall be ASTM Thermometer 67C of Specification E 1.

6.2.3 Thermocouples shall comply with the requirements of Specification E 608 and shall be calibrated in accordance with Method E 220.

6.2.4 Resistance thermometers shall comply with the requirements of Test Methods E 644 and Specification E 1137, and be calibrated in accordance with NIST Special Publication 250-22.<sup>5,6</sup>

6.3 *Molding Press*, designed to operate at  $200 \pm 5^\circ\text{C}$  [ $392 \pm 9^\circ\text{F}$ ].

6.4 *Injection Molding Unit*, meeting the requirements of Practice D 3641.

6.5 *Mold*:

6.5.1 *Compression Mold*, comprised of the following:

6.5.1.1 *Compression Molding Chase*, having a blanked-out area of suitable size (Note 3) and capable of producing a plaque  $1.00 \pm 0.05$  mm [ $0.039 \pm 0.002$  in.] thick.

NOTE 3—A 152.4 by 152.4-mm [6 by 6-in.] blanked-out section has been found satisfactory.

6.5.1.2 *Backing Plates*, large enough to cover this chase and strong enough to resist warping or distortion, under the molding conditions. Polished steel plates, 3 mm [0.1 in.] thick, are satisfactory.

6.5.2 *Injection Mold*, designed using the guidelines specified in Practice D 3641. The mold shall be capable of producing either a plaque having a thickness of  $1.00 \pm 0.05$  mm [ $0.039 \pm 0.002$  in.] from which 50 by 10 by 1.00-mm [2 by 0.4 by 0.039-in.] specimens may be die cut, or mold a standard test specimen, 50 by 10 by 1.00 mm.

6.6 *Parting Sheets*—Fluoropolymer, polyester, or other film that will not affect the long-term thermal stability of polypropylene, 0.05 to 0.20 mm [0.002 to 0.008 in.] thick. The film must be free of wrinkles and foreign matter, such as lubricants and oils.

6.7 *Cutting Die*, to produce 50 by 10-mm [2 by 0.4-in.] specimens from either a compression-molded or injection-molded plaque. The die must be sharp and free of nicks.

6.8 *Specimen Holder*—The specimen holder may be biaxially rotated or uniaxially rotated provided that the test specimens are in a stream of air having a relative velocity about 1000 mm/s [ $197$  ft/min]. Illustrations of suitable apparatus for biaxially and uniaxially rotated specimen holders are shown in Figs. 1 and 2, respectively. Biaxial rotation increases the probability that all specimens will be exposed similarly. In the case of dispute, the use of biaxial rotation shall be the reference method.

6.8.1 *Biaxially Rotated Specimen Holder* (see Fig. 1)<sup>7</sup>. The frequency of rotation about the horizontal and vertical axes shall be 1 to 3  $\text{min}^{-1}$ .

6.8.2 *Uniaxially Rotated Specimen Holder* (see Fig. 2). The drum peripheral velocity shall be such that the air stream impinges on the flat section of the test specimens at about 1 m/s.

<sup>5</sup> Mangum, B. W., "Platinum Resistance Thermometer Calibration," NBS Special Publication 250-22 (1987).

<sup>6</sup> Available from National Institute of Standards and Technology (NIST), 100 Bureau Dr., Stop 3460, Gaithersburg, MD 20899-3460.

<sup>7</sup> Standard Scientific Supply Company, Model CS191, or equivalent, can be used.

<sup>4</sup> A Precision Scientific Freas Model 835 B, or Blue M POM-206C-1, or equivalent, can be used.

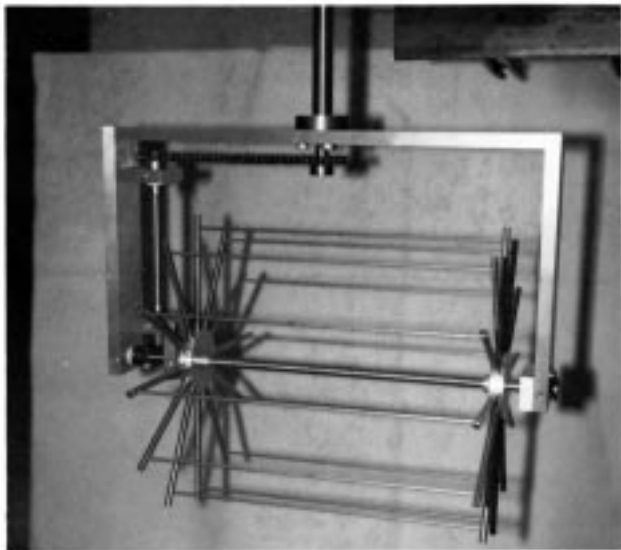


FIG. 1 Biaxial Ferris-Wheel-Type Rotator

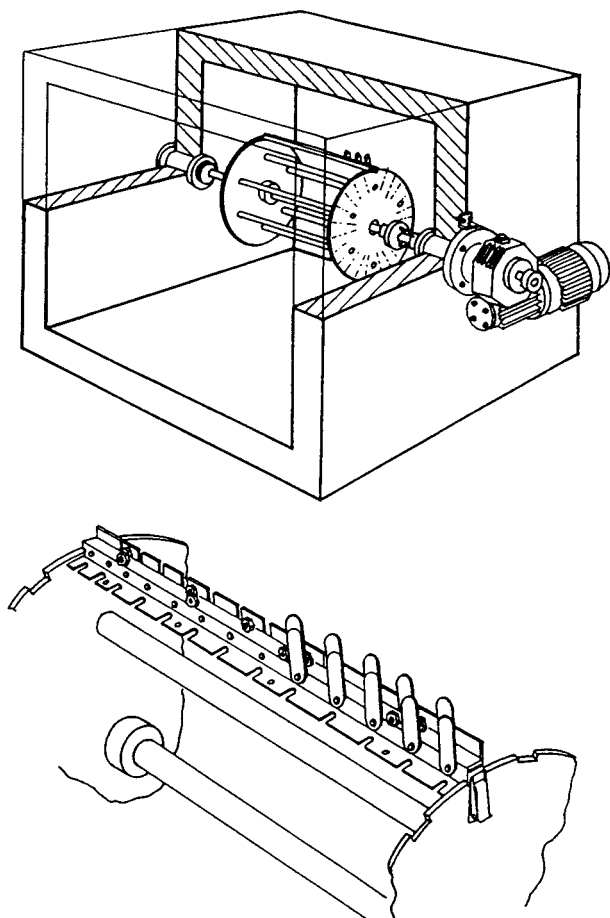


FIG. 2 Uniaxially Rotated Specimen Holder—Drum Rotator

6.9 *Air Velocity Meter*, nondirectional resistance wire type for measuring the air velocity in the oven.

6.10 *Anemometer*, with a circular vane, for determining the frequency of air changes in the oven. The anemometer shall be positioned directly in front of the oven outlet. Based on the

design of the outlet and the anemometer, every effort shall be made to position the anemometer's electronics in a way as to minimize the degree of exposure to the heated air exiting the oven.

6.11 *Metal Clips*, lined with fluorocarbon film or other materials that have no adverse effect on the oxidative thermal stability of polypropylene.

## 7. Specimen Preparation

7.1 The test specimens shall be cut from either a compression-molded plaque or injection-molded plaque prepared from granules or other homogeneous molding material. Plaques shall be prepared as follows:

### 7.2 Compression Molding:

7.2.1 Adjust the temperature of the platens to  $200 \pm 5^\circ\text{C}$  [ $392 \pm 9^\circ\text{F}$ ].

7.2.2 Place a smooth, clean parting sheet on a backing plate and center the chase on it. Put enough of the sample into the cavity to fill it completely when molded. A slight excess of material is desirable. Cover the loaded chase first with a clean parting sheet and then a backing plate. Finally, put the assembled mold on the lower platen and close the press carefully until both platens are in contact with the assembly. When the material has melted, apply sufficient pressure to form a void-free plaque in the  $1.00 \pm 0.5 \text{ mm}$  [ $0.039 \pm 0.002 \text{ in.}$ ] thick and record this pressure. Leave the polypropylene in the heated press under pressure for 3 to 4 min at  $200 \pm 5^\circ\text{C}$  [ $392 \pm 9^\circ\text{F}$ ]. Flash cool the mold assembly by transferring to a water-cooled press or by water quenching.

### 7.3 Injection Molding:

7.3.1 Plaques or test specimens shall be injection-molded in accordance with the requirements of Specification D 4101.

7.4 Prepare a minimum of five specimens per material sample by die-cutting specimens from the plaque or directly molding the standard test specimen. The standard specimen shall be 10 mm wide, 50 mm long, and  $1.00 \pm 0.05 \text{ mm}$  thick. Edges shall be smoothed, if necessary, to remove imperfections introduced by cutting.

7.5 Test specimens prepared directly by injection molding without die cutting or specimens cut from polypropylene products can be used as agreed upon between the interested parties. In all cases the referee method will be based on compression molded die cut specimens.

NOTE 4—Failure test times for compression-molded and injection-molded specimens may vary due to the skin surface effect and the distinctness or sharpness of the specimen edge.

7.6 Specimens of other thicknesses can be used as agreed upon between the interested parties.

7.7 In cases of dispute, the referee specimens shall be die-cut only from compression-molded specimens, unless the test specimen is otherwise agreed upon.

7.8 In handling the plaques and cut specimens, use clean gloves or tongs to prevent contamination of the test specimens.

## 8. Conditioning

8.1 Condition the test specimens in accordance with Specification D 4101. When unsure of the material formulation, condition the test specimen at  $23 \pm 2^\circ\text{C}$  [ $73.4 \pm 3.6^\circ\text{F}$ ] and 50

$\pm 5\%$  relative humidity for not less than 40 h prior to test in accordance with Procedure A of Practice D 618. In cases of disagreement, the tolerances shall be  $\pm 1^\circ\text{C}$  [ $\pm 1.8^\circ\text{F}$ ] and  $\pm 2\%$  relative humidity.

## 9. Oven Parameters

9.1 Unless otherwise specified, the oven temperature shall be  $150^\circ\text{C}$  [ $302^\circ\text{F}$ ] and shall not vary more than  $\pm 1^\circ\text{C}$ , measured at a single location within the oven, or more than  $\pm 1.5^\circ\text{C}$  throughout the oven.

9.1.1 To monitor temperature, the oven temperature measurement sensor should be conveniently located about 80 mm [3 in.] from the top liner of the oven, in the vicinity of the exhaust port.

9.2 In the event that the average time to failure at  $150^\circ\text{C}$  is less than seven days, the test report shall indicate “less than seven days at  $150^\circ\text{C}$ ” and the test shall be repeated at an oven temperature of  $140 \pm 1^\circ\text{C}$ , and, if necessary, at lower temperatures at intervals of  $10^\circ\text{C}$  until time to failure of seven days is achieved.

NOTE 5—If it is preferred to perform the test at  $140^\circ\text{C}$ , this is permitted provided that this is stated in the test report. This is applicable even for materials lasting more than seven days at  $150^\circ\text{C}$ .

9.3 The air velocity shall be  $1000 \pm 250$  mm/s [ $197 \pm 49$  ft/mm] during the course of the test as measured at the center of the oven.

NOTE 6—The air velocity in the oven is dependent on the design of the oven and will not vary a great deal. It should be checked occasionally to determine if the blowers and other moving parts are functioning properly.

NOTE 7—The air velocity should be measured at  $+23^\circ\text{C}$  for non-thermally corrected anemometers. A thermally-corrected anemometer can be used in the range from  $+20$  to  $+80^\circ\text{C}$ . In no case should an anemometer be used to measure air flow at the test temperature. Typically an anemometer will have a 25 % error for each  $35^\circ\text{C}$  above its maximum calibrated temperature point.

9.4 The air intake and exhaust ports shall be adjusted to allow approximately  $1 \pm 15\%$  air volume change/min.

9.4.1 The number of air changes that take place in the oven should be determined by the power consumption technique described in Test Methods D 5374.

9.4.2 An anemometer can be used to measure the number of air exchanges also. This method is described in Appendix X1. If used, this method should be verified with the power consumption technique, and the anemometer value must be within 15 % of the number of air changes observed with the power consumption method. The advantages of the anemometer method is that it is quick, easy, and can be conducted on a more frequent basis to ensure that no changes have occurred in the settings or conditions in the oven. It is not an absolute measure.

## 10. Procedure

NOTE 8—**Warning:** Handle specimens with caution to avoid contamination.

10.1 The metal clips with fluoropolymer lining shall be cleaned beforehand with trichloroethylene or any other suitable solvent to remove all traces of oil.

10.2 Attach the five specimens to the biaxial rotator by suitable metal clips lined with fluoropolymer film. Space the specimens 30 to 40 mm [1.2 to 1.6 in.] from each other on the rack.

10.2.1 Avoid direct contact of the specimens with the metal clips or metal parts. Certain metals, such as aluminum, are known to affect with the long-term thermal endurance of some polypropylenes.

10.3 Inspect the specimens at least once a day, turning off the biaxial rotator during inspection.

10.4 Failure by this test method for polypropylene is visual evidence of localized discoloration and crumbling, crazing or pin holes that open when sample is flexed on any part of the specimen directly exposed to the air flow. The area within 5 mm of the clips shall not be included in this evaluation.

NOTE 9—When testing polypropylene containing recycled material localized discoloration often appears on the surface of the specimen within a few days of exposure to the heat. This should not be interpreted as a failure unless the criteria of failure specified in 10.4 are met.

NOTE 10—Visual inspection of the deterioration of the specimen surface, as a reliable evaluation of oxidative aging endurance, is justified by the fact that oxidation of propylene thermoplastics usually takes place on the surface of the material before propagating inward.

10.5 Record the time to failure of each specimen in days.

## 11. Report

11.1 The report shall include the following information:

11.1.1 A complete identification of the sample,

11.1.2 The average time to failure, in days at  $150^\circ\text{C}$ <sup>8</sup> of the five specimens,

11.1.3 The range of times to failure in days at  $150^\circ\text{C}$ <sup>8</sup> of the five specimens,

11.1.4 The preparation or history of the specimen,

11.1.5 The thickness of the specimen,

11.1.6 The type of rotator,

11.1.7 The test temperature, if other than  $150^\circ\text{C}$ ,

11.1.8 Any departures from the specified test conditions, and

11.1.9 Details of conditioning (temperature and humidity) if appropriate.

## 12. Precision and Bias

12.1 Table 1 is based on a round robin conducted in 1970 in accordance with Practice E 691, involving three materials tested by seven laboratories. Each material was supplied from one source, but the individual specimens were prepared at the laboratory that tested the specimens. Test specimens were 1.25 mm [0.50 in.] in thickness. Each of two test results that was obtained by each laboratory was the average of five individual determinations. **Warning**—The following explanations of *r* and *R* (12.2-12.2.3) are only intended to present a meaningful way of considering the approximate precision of this test method. The data in Table 1 should not be rigorously applied to acceptance or rejection of material, as those data are specific to

<sup>8</sup> If the time to failure is less than seven days at  $150^\circ\text{C}$ , report as “less than seven days at  $150^\circ\text{C}$ .” In addition, report the time (range of times) to failure at  $140^\circ\text{C}$  (or at other temperatures used).



**TABLE 1 Precision Data for Oxidative Stability of Polypropylene**

Material	Average Time to Failure, days	$S_r^A$	$S_R^B$	$r^C$	$R^D$
L	14.0	0.765	2.63	2.16	7.44
M	34.9	3.08	6.95	8.72	19.6
H	63.4	5.42	18.7	15.3	52.9

<sup>A</sup>  $S_r$  = within-laboratory standard deviation of the average.

<sup>B</sup>  $S_R$  = between-laboratory standard deviation of the average.

<sup>C</sup>  $r = 2.8 \times S_r$ .

<sup>D</sup>  $R = 2.8 \times S_R$ .

the round robin and may not be representative of the other lots, conditions, materials, or laboratories. Users of this test method should apply the principles outlined in Practice E 691 to generate data specific to their laboratory and materials, or between specific laboratories. The principles of 12.2-12.2.3 would then be valid for such data.

12.2 *Concept of r and R*—If  $S_r$  and  $S_R$  have been calculated from a large enough body of data, and for test results that were averages from testing five test specimens:

12.2.1 *Repeatability (r)*—In comparing two test results for the same material, obtained by the same operator using the same equipment on the same day, the two test results should be judged not equivalent if they differ by more than the  $I_r$  value for that material.

12.2.2 *Reproducibility (R)*—In comparing two test results for the same material, obtained by different operators using different equipment on different days, the two test results should be judged not equivalent if they differ by more than the  $I_R$  value for that material.

12.2.3 Any judgment in accordance with 12.2.1 and 12.2.2 would have an approximate 95 % (0.95) probability of being correct.

12.3 *Bias*—There are no recognized standards on which to base an estimate of bias for this test method.

### 13. Keywords

13.1 aging; biaxial rotator; polypropylene; thermal aging; thermal oxidative stability

## APPENDIX

### (Nonmandatory Information)

#### X1. SECONDARY METHOD FOR DETERMINING THE NUMBER OF AIR EXCHANGES

##### X1.1 Scope

X1.1.1 The purpose of this appendix is to provide a secondary method for determining the number of air exchanges that occur within the forced draft oven used for measurement of thermal oxidative stability.

##### X1.2 Summary of Test Method

X1.2.1 This method uses an anemometer to measure the linear velocity of the air exiting the heated oven through the oven exhaust outlet. Knowing the volume of the interior of the oven and the cross-sectional area of the exhaust outlet and linear velocity of the air, the number of air volume changes occurring in the oven over a 1-h period can be calculated.

##### X1.3 Apparatus

X1.3.1 *Anemometer*, with circular vane. The vane shall be of equal or smaller diameter than the inside diameter of the exhaust outlet. The electronics within the anemometer shall be below the level of the heated exhaust to prevent the electronics from being heated and errors in calibration occurring. The unit should have a digital display of the air velocity.

X1.3.2 *Temperature Measuring Devices*, to determine the temperature in the room and in the oven.

##### X1.4 Procedure

X1.4.1 Ensure that the anemometer has been properly calibrated before using.

X1.4.2 Position the vane of the anemometer at the exhaust outlet, ensuring that it is centered on the outlet.

X1.4.3 Monitor the digital display of the anemometer until the reading stabilizes. Record this value as the linear air velocity in metres per minute. Move the vane away from the exhaust port and allow it to cool. Once cool, repeat the air velocity measurement. Ensure that the anemometer is cooled between each measurement to prevent heat buildup that might effect the electronics. Make a minimum of three measurements of the air velocity.

X1.4.4 Determine the volume of the interior of the oven in cubic metres.

X1.4.5 Determine the inside cross-sectional area of the exhaust outlet, in square metres.

X1.4.6 Measure the air temperature within the laboratory to the nearest degree, in degree centigrade.

X1.4.7 Measure the air temperature within the oven, in degree centigrade.

##### X1.5 Calculation

X1.5.1 Determine the average linear air velocity from data collected in X1.4.3.

X1.5.2 Convert the oven temperature and the room temperature from degree centigrade to degree Kelvin ( $^{\circ}\text{C} + 273 = \text{K}$ )

X1.5.3 Using the Perfect Gas Law,  $PV = nRT$ , and assuming pressure remains constant, the volume of air once heated in the oven that will exit the exhaust outlet can be determined as follows:

$$V_1/V_2 = T_1/T_2 \quad (\text{X1.1})$$

where:

- $V_1$  = volume of air in the oven at room temperature,  $m^3$ ,
- $V_2$  = volume of air in the oven at the set temperature,  $m^3$ ,
- $T_1$  = temperature of the room, K, and
- $T_2$  = temperature of the oven, K.

NOTE X1.1—Should  $V_1$  be more than  $3^\circ$  from  $+23^\circ\text{C}$ , it is required that you calculate  $V_1$  first using the following equation:

$$V_1 = (VO \times T_1)/296 \quad (\text{X1.2})$$

where:

- $VO$  = volume of oven calculated from dimensions at  $+23^\circ\text{C}$ ,  $m^3$ , and
- $296 = 296\text{K} = +23^\circ\text{C} + 273\text{K}$ .

X1.5.4 Calculate the air volume discharged from the oven as follows:

$$CMH = V_3 \times A \times (V_1/V_2) \times 60 \quad (\text{X1.3})$$

where:

- $CMH$  = air volume exiting the exhaust outlet per hour,  $m^3/h$ ,
- $V_3$  = air velocity of air exiting the exhaust outlet,  $m/min$ ,
- $A$  = cross-sectional area of exhaust outlet,  $m^2$ , and
- $60$  = conversion from minutes to hours.

X1.5.5 Calculate the number of air changes per hour as follows:

$$AC = CMH/VO \quad (\text{X1.4})$$

where:

- $AC$  = number of air changes per hour.

## SUMMARY OF CHANGES

This section identifies the location of selected changes to this test method. For the convenience of the user, Committee D20 has highlighted those changes that may impact the use of this test method. This section may also include descriptions of the changes or reasons for the changes, or both.

### *D 3012 – 04:*

- (1) Deleted D 5423 from 2.1.
- (2) Deleted last sentence of 6.1.
- (3) Revised 10.4 for expansion of failure criteria.
- (4) Added new Note 9 to address recycled material.
- (5) Renumbered old Note 9 to Note 10.

### *D 3012 – 00:*

- (1) Changed “propylene plastics” to “polypropylene” throughout.
- (2) Changed “°K” to “K” throughout.
- (3) Changed permissive language from “may be” to “can be” throughout.
- (4) Added thermal oxidative in 1.2.
- (5) Made editorial corrections throughout.

- (6) Revised text in 6.9.
- (7) Added 6.10, Anemometer, and made editorial corrections.
- (8) Made editorial corrections, elimination of redundancy, and added information on how to smooth edges of specimens with flash in 7.3.
- (9) Eliminated last sentence in 7.4, as it was redundant.
- (10) Added the sentence, “It is not an absolute measure.” and editorial correction in 9.2.
- (11) Added anemometer and referral to Appendix X1 in 9.4.2. Added tolerance on air changes, and eliminated word “approximately.”
- (12) Added Appendix X1 on how to use an anemometer to measure the number of air changes in the oven.

*ASTM International takes no position respecting the validity of any patent rights asserted in connection with any item mentioned in this standard. Users of this standard are expressly advised that determination of the validity of any such patent rights, and the risk of infringement of such rights, are entirely their own responsibility.*

*This standard is subject to revision at any time by the responsible technical committee and must be reviewed every five years and if not revised, either reapproved or withdrawn. Your comments are invited either for revision of this standard or for additional standards and should be addressed to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee, which you may attend. If you feel that your comments have not received a fair hearing you should make your views known to the ASTM Committee on Standards, at the address shown below.*

*This standard is copyrighted by ASTM International, 100 Barr Harbor Drive, PO Box C700, West Conshohocken, PA 19428-2959, United States. Individual reprints (single or multiple copies) of this standard may be obtained by contacting ASTM at the above address or at 610-832-9585 (phone), 610-832-9555 (fax), or service@astm.org (e-mail); or through the ASTM website (www.astm.org).*