



Designation: D2883 – 95(Reapproved 2009)

Standard Test Method for Reaction Threshold Temperature of Liquid and Solid Materials¹

This standard is issued under the fixed designation D2883; 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 covers determination of the pre-flame, cool-flame, and hot-flame reaction threshold temperatures and the incipient reaction temperature of liquids and solids. Data may be obtained at pressures from low vacuum to 0.8 MPa (115 psia) for temperatures within the range from room temperature to 925 K (1200°F).

1.2 This test method may be applied to any substance that is a liquid or a solid at room temperature and atmospheric pressure and that, at room temperature, is compatible with glass and stainless steel. Air is the intended oxidizing medium; however, other media may be substituted provided appropriate precautions are taken for their safe use.

1.3 The values stated in SI units are to be regarded as the standard. In cases where materials, products, or equipment are available in inch-pound units only, SI units are omitted.

1.4 *This standard is used to measure and describe the response of materials, products, or assemblies to heat and flame under controlled conditions, but does not by itself incorporate all factors required for fire hazard or fire risk assessment of the materials, products, or assemblies under actual fire conditions.*

1.5 *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 warning statements, see 6.8, Sections 7 and 9.*

2. Referenced Documents

2.1 *ASTM Standards:*²

D1193 Specification for Reagent Water

¹ This test method is under the jurisdiction of ASTM Committee D02 on Petroleum Products, Liquid Fuels, and Lubricants and is the direct responsibility of Subcommittee D02.L0.07 on Engineering Sciences of High Performance Fluids and Solids (Formally D02.1100).

Current edition approved Oct. 1, 2009. Published November 2009. Originally approved in 1970. Last previous edition approved in 2005 as D2883–95(2005). DOI: 10.1520/D2883-95R09.

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.

D2021 Specification for Neutral Detergent, 40 Percent Alkylbenzene Sulfonate Type (Withdrawn 2000)³

E659 Test Method for Autoignition Temperature of Liquid Chemicals

2.2 *Military Standards:*⁴

MIL-C-81302 Trichlorotrifluoroethane

MIL-T-7003 Trichloroethylene

3. Terminology

3.1 *Definitions of Terms Specific to This Standard:*

3.1.1 *catalytic reaction, n*—a fast, self-sustaining, energetic, sometimes luminous, sometimes audible reaction that occurs as a result of catalytic action on the surface of the thermocouple or other solid surface within the combustion chamber.

3.1.2 *cool-flame reaction, n*—a relatively slow, self-sustaining, barely luminous reaction of the sample or its decomposition products with the atmosphere in the combustion chamber.

3.1.2.1 *Discussion*—This type of flame is visible only in a darkened area. Figs. 1-4 illustrate the type of temperature records obtained for cool-flames.

3.1.3 *hot-flame reaction, n*—a rapid, self-sustaining, luminous, sometimes audible reaction of the sample or its decomposition products with the atmosphere in the combustion chamber.

3.1.3.1 *Discussion*—A yellow or blue flame usually accompanies the reaction. Figs. 5-7 illustrate the type of temperature records obtained for hot-flames.

3.1.4 *incipient reaction temperature, n*—the temperature obtained by extrapolation of a plot of the reciprocal of the reaction delay time versus the initial reaction temperature to that value of the reciprocal which represents infinite delay ($t^{-1} = 0$) (Fig. 8).

3.1.4.1 *Discussion*—The detection of sample reactions and the distinctions between different reactions are based upon the records produced by the various sensing devices used in the apparatus. The temperature record is basic to the method and

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

⁴ Available from Standardization Documents Order Desk, Bldg 4, Section D, 700 Robbins Ave., Philadelphia, PA 19111-5094. Attn: NPODS.

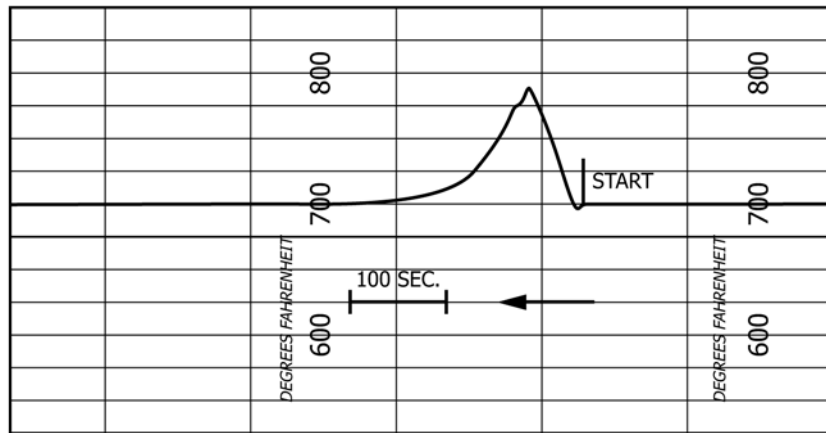


FIG. 1 Time-Temperature Profile for Typical Cool-Flame Reactions

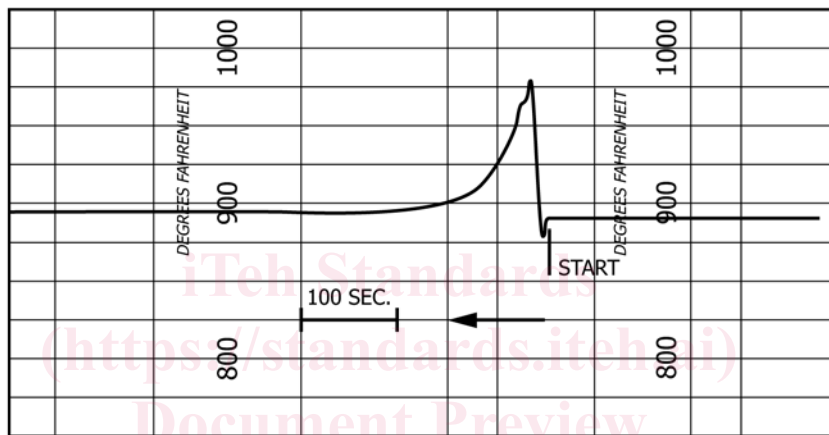


FIG. 2 Time-Temperature Profile for Typical Cool-Flame Reactions

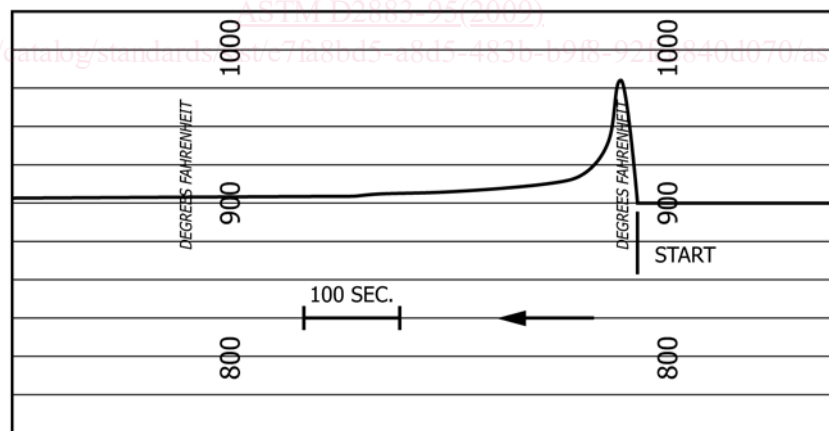


FIG. 3 Time-Temperature Profile for Typical Cool-Flame Reactions

shall be regarded as the primary source of such data. The use of a pressure record and the records of other sensing devices to obtain additional data is optional.

3.1.5 *noncombustive reaction, n*—a reaction other than combustion or thermal degradation undergone by certain materials upon exposure to elevated temperatures.

3.1.5.1 *Discussion*—Thermal polymerization is an example

of the type of reaction that may occur when dealing with potentially reactive substances.

3.1.6 *pre-flame reaction, n*—a slow, nonluminous reaction of the sample or its decomposition products with the atmosphere in the combustion chamber.

3.1.6.1 *Discussion*—Fig. 9 and Fig. 10 illustrate the type of temperature records obtained for preflame reactions.

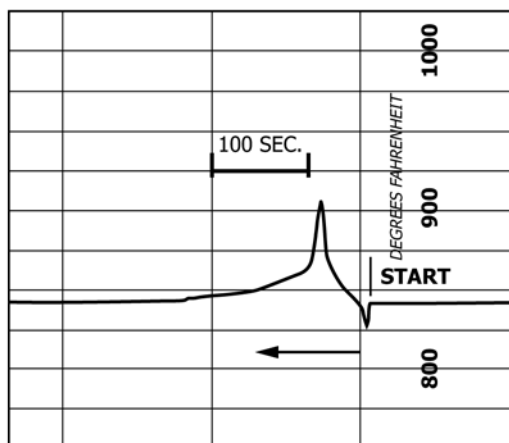


FIG. 4 Time-Temperature Profile for Typical Cool-Flame Reactions

3.1.7 *reaction delay time, n*—the time, measured in seconds, that elapses between the introduction of the sample into the reaction chamber and the attainment of maximum response from one of the sensors used to follow the reaction (Fig. 3).

3.1.8 *reaction threshold temperature, n*—the lowest temperature at which any reaction of the sample or its decomposition products may be detected by a thermocouple or other sensing devices.

3.2 Symbols:

- T = temperature, °C
- T_i = incipient reaction temperature, °C
- t = delay time, s
- P = pressure, Pa or torr
- h = hot-flame reaction
- c = cool-flame reaction
- p = pre-flame reaction
- c = catalytic reaction
- n = noncombustion reaction

4. Summary of Test Method

4.1 A small measured amount of the sample contained in a glass ampoule is introduced into a stainless steel reaction chamber maintained at preselected temperature and pressure. The reactions of the sample subsequent to its introduction into the reaction chamber are followed by monitoring the temperature or the temperature and pressure (optional) of the reaction chamber as a function of time.

4.2 At any selected system pressure, the minimum temperature (reaction threshold temperature) required to produce a given reaction is determined as a function of the size of the sample employed. The delay times for cool-flame and hot-flame reactions are also measured. A plot of the reciprocal of the reaction delay time versus the initial reaction chamber temperature for a cool-flame or hot-flame reaction may be used to determine the incipient reaction temperature by extrapolation to infinite delay time ($t^{-1} = 0$).

NOTE 1—The hot-flame reaction, cool-flame reaction, and reaction threshold temperature obtained by this test method may approximate those temperatures obtained by Test Method E659, for hot-flame autoignition, cool-flame autoignition, and reaction threshold temperature, respectively.

5. Significance and Use

5.1 The reaction thresholds of a material are a measure of the tendency of the material or its decomposition products to undergo gas phase reactions of various types. Hot-flame and cool-flame thresholds relate directly to reactions which are involved in autoignition phenomena. Pre-flame, catalytic and thermal polymerization thresholds also relate to autoignition in that they represent reactions which can be under some conditions the precursors of ignition reactions.

6. Apparatus^{5,6}

6.1 *Oven*, fan-assisted air-circulating, with a 300 by 300 by 300 mm minimum volume, centrally located in the oven work space within which temperature uniformity shall be ± 3 K over the temperature range from room temperature to 925 K. A typical oven is detailed in the Appendix.

6.1.1 Temperature uniformity shall be determined by the following test: Eight thermocouples shall be arranged within the oven in such a manner that they shall be located at positions designated by the corners of a 300 mm cube. Awg No. 30, calibrated, bare junction iron-constantan or Chromel-Alumel thermocouples are suitable. Lead-in ports may be plugged with insulating material during the uniformity test.

6.2 *Reaction Temperature Apparatus*, assembly as shown in Fig. 11, comprising the following:

6.2.1 *Combustion Chamber*, 1000 mL round-bottom, long-neck, AISI Type 316 stainless steel flask with 25.4 mm (1-in.) Corning ferrule, as shown in Fig. 12.

6.2.2 *Pressure Control Manifold*, as shown in Fig. 13, consisting of:

6.2.2.1 *Thermocouple Gland*.^{6,7}

6.2.2.2 *Insertion Rod Guide*, AISI Type 316 tube, 6.4 mm (¼ in.) in outside diameter by 1.245 mm (0.049 in.) wall.

6.2.2.3 *Chamber Head* with 25.4 mm (1-in.) Corning ferrule.

6.2.2.4 *Pressure Transducer* (optional).

6.2.2.5 *Pressure Relief Valve*, capable of relieving pressure at 40 % above test equilibrium pressure.

6.2.2.6 *Air Inlet Valve*, AISI Type 316 stainless steel.

6.2.3 *Electromagnet*, capable of propelling insertion rod against anvil with sufficient force to break sample ampoule.

6.2.4 *Insertion Rod*, AISI Type 316 tubing, 3.2 mm (⅛ in.) in outside diameter by 673 mm (26.5 in.) in length with attached armature.

6.2.5 *Flange Assembly*, Corning Style 1 for 25.4 mm (1-in.) pipe.

6.2.6 *Thermocouples*, calibrated precision grade iron-constantan (ISA-Type and Chromel-Alumel (ISA-Type K), Awg No. 30 glass-fiber insulated, bare-junction (welded) in 5-mm borosilicate glass tube.

⁵ The sole source of supply of the apparatus known to the committee at this time is W.A. Sales Ltd., 419 Harvester Ct., Wheeling, IL 60090.

⁶ If you are aware of alternative suppliers, please provide this information to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee,¹ which you may attend.

⁷ Reducer, Swagelok 200-R-4-316BT with silicone rubber disk insert to form seal around wire has been found satisfactory.

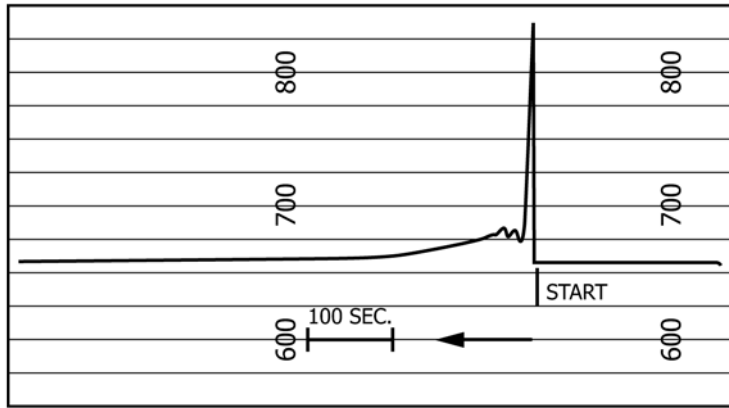


FIG. 5 Time-Temperature Profile and Delay Time for Typical Hot-Flame Reactions

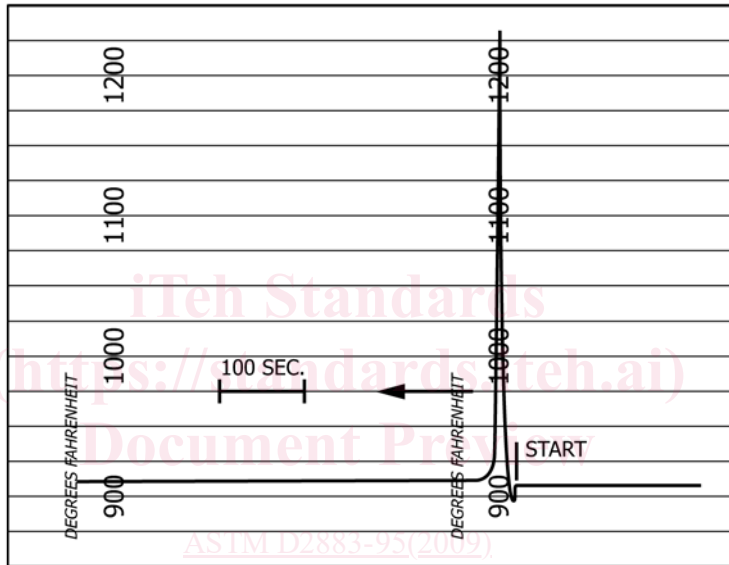


FIG. 6 Time-Temperature Profile and Delay Time for Typical Hot-Flame Reactions

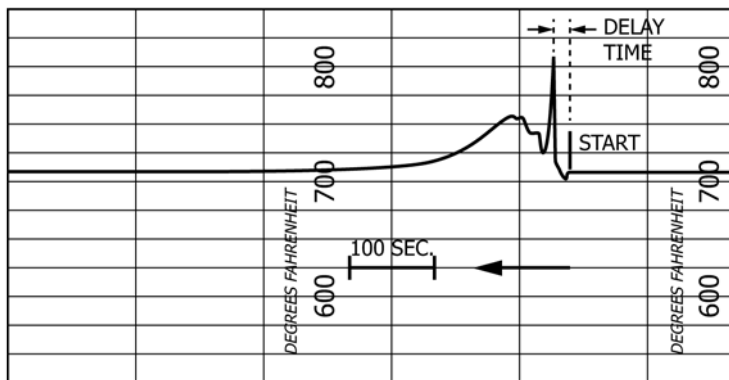


FIG. 7 Time-Temperature Profile and Delay Time for Typical Hot-Flame Reactions

6.3 *Sample Ampoule*, illustrated for solids and liquids in Fig. 14.

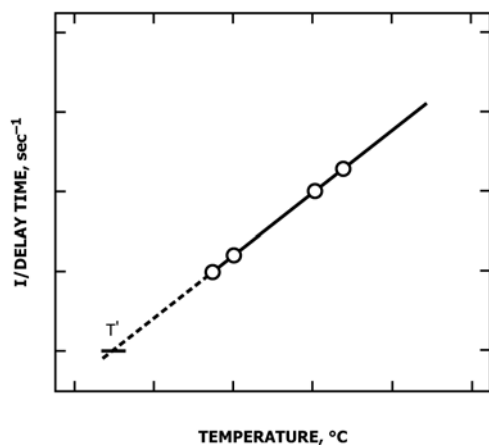


FIG. 8 Effect of the Reaction Temperature on the Delay Time to Determine Incipient Reaction Temperature

6.4 *Temperature Recorder*,^{6,8} 475 to 1025 K, (400 to 1400°F) range, ¼ s full-scale deflection, thermocouple actuation with range changes for iron-constantan and Chromel-Alumel elements.

6.5 *Hypodermic Syringe*,^{6,9} fluorocarbon-tipped 0.5 mL and 2.5 mL with hypodermic needles suitable for introduction into sample ampoules.

6.6 *Pressure Gauges*, capable of determining the appropriate pressure with an accuracy of ±1 %.

6.7 *Vacuum System*, capable of attaining and maintaining the required pressure with mechanical vacuum pump^{6,10} and cold trap inserted in the line between the reaction temperature apparatus and the pump.

6.8 *Gas Supply System*,^{6,11} capable of supplying the appropriate gas, dewpoint 215 K (−70°F) maximum and essentially free of contaminants at pressures up to 0.8 MPa. (**Warning**—Compressed gas under high pressure. Gas reduces oxygen available for breathing. See A2.1.)

6.9 *Exhaust Hood*, capable of disposing of fumes vented from the reaction temperature apparatus.

7. Reagents and Materials

7.1 *Trichloroethylene*, conforming to MIL-T-7003. Trichlorotrifluoroethane conforming to MIL-C-81302 or high flash point naphtha may be used as alternatives. (**Warning**—Harmful if inhaled. High concentrations may cause unconsciousness or death. Contact can cause skin irritation and dermatitis. See A2.2.)

7.2 *Cleaning Compounds*.^{6,12}

⁸ Honeywell Model KSY 153 × 116—(PH)-II-III-163-66-D has been found satisfactory.

⁹ The sole source of supply of Models 1750 and 1002 with Luer Lok known to the committee at this time is Hamilton Co., Whittier, CA.

¹⁰ 125 W (½ hp) carbon vane pump has been found satisfactory for most applications.

¹¹ The sole source of supply of dry air known to the committee at this time is Matheson Co., East Rutherford, NJ.

¹² The sole source of supply of “Transpo” cleaner known to the committee at this time is Turco Products, Inc., Rockdale, IL.

7.3 *Chloroform*, technical grade. (**Warning**—Can be fatal if swallowed. Harmful if inhaled. Can produce toxic vapors if burned. See A2.5.)

7.4 *Detergent*, free-rinsing, conforming to Specification D2021.

7.5 *Reagent Water*, conforming to Specification D1193, either grade.

7.6 *Acetic Acid*, ACS reagent grade.

7.7 *Hydroiodic Acid*, 57 % reagent grade.

8. Test Specimen

8.1 *Liquid Sample*—200 mL is sufficient for the determination of the various reaction threshold temperatures at a single pressure.

8.2 *Solid Sample*—150 g is sufficient for the determination of the various reaction threshold temperatures at a single pressure.

9. Procedure

9.1 *Method A for Liquids*:

9.1.1 Clean a combustion chamber as follows: rinse with trichloroethylene (**Warning**—Harmful if inhaled. High concentrations may cause unconsciousness or death. Contact can cause skin irritation and dermatitis. See A2.2.), immerse in cleaning compound for a minimum of 15 min at 293 to 308 K, wash with detergent, and rinse thoroughly with reagent water. Dry in an oven at 393 to 423 K. Discard the chamber if not free of deposits. Clean the thermocouple internal supports, anvil, and insertion rod in the same manner.

NOTE 2—A chloroform (see 7.3) rinse prior to washing can prove useful in some cases. Difficult deposits may be removed by walnut shell abrasion. Discard the chamber if deposits can not be removed or if inspection reveals that the interior of the chamber has been damaged or corroded.

NOTE 3—Stainless steel, Type 316 combustion chambers and an air atmosphere have been specified with this test method. However, chambers of other materials and of different sizes, and atmospheres of other gases may be used when required.

NOTE 4—Chambers may be tested for the presence of peroxide or free radical residues by purging them with nitrogen and rinsing with a freshly prepared mixture of 3 mL each of chloroform, glacial acetic acid, (**Warning**—Poison. Corrosive. Combustible. Can be fatal if swallowed. Causes severe burns. Harmful if inhaled. See A2.6) and hydroiodic acid (57 % analytical grade, not stabilized) (**Warning**—Poison. Corrosive. Can be fatal if swallowed. Liquid and vapor cause severe burns. Harmful if inhaled. See A2.7). When this mixture is poured into a clean glass vessel and 10 drops of stabilized starch solution added, a trace of the starch-iodine color reaction indicates no peroxides or free radicals are present.

9.1.2 Assemble the reaction temperature apparatus in accordance with Fig. 11, using a clean combustion chamber. Insert the assembly horizontally in the oven to the middle of the temperature-controlled zone to a minimum inside depth of 21 cm.

9.1.3 Connect the vacuum system, gas supply system (**Warning**—Compressed gas under high pressure. Gas reduces oxygen available for breathing. See A2.1.), pressure gauges, and temperature recorder to the apparatus in accordance with Fig. 13.

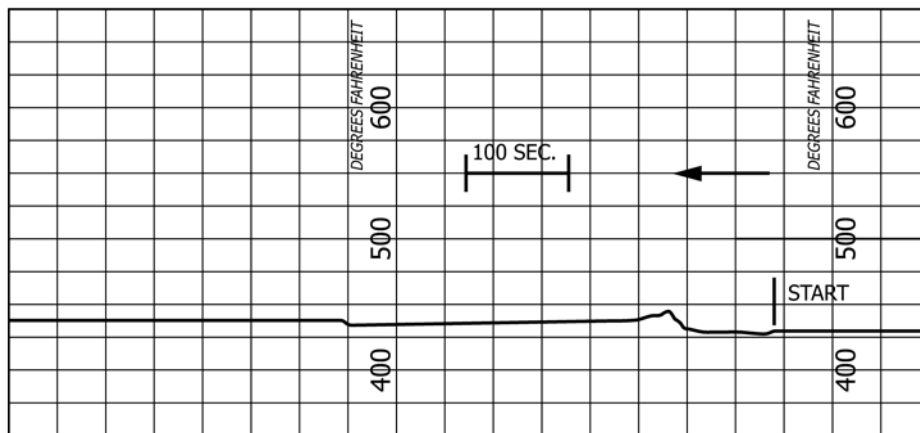


FIG. 9 Time-Temperature Profile for Typical Pre-Flame Reactions

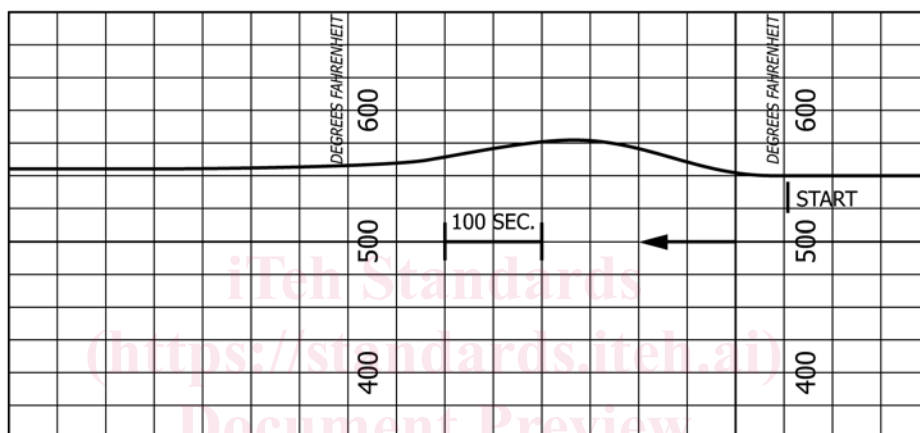


FIG. 10 Time-Temperature Profile for Typical Pre-Flame Reactions

9.1.4 Set the temperature controller of the oven at a temperature approximately 50 K below the expected threshold for the hot-flame reaction of the sample. Turn on the heat and blower switches of the oven. Increase the pressure in the gas supply system or decrease the pressure through the vacuum system as required.

9.1.5 Establish thermal and pressure equilibrium in the system. Record the temperature and pressure for at least 10 min to ensure a satisfactory equilibrium.

9.1.6 Fill the 0.5-mL syringe with a portion of the sample and transfer 0.2 mL to the thin-walled bulb portion of the liquid sample ampoule (Fig. 14). Place the cap on the ampoule and affix the covered ampoule to the end of the insertion rod.

9.1.6.1 If the sample is extremely volatile, seal the ampoule by drawing the open end closed in the flame of a torch. Use caution to prevent possible violent reaction between sample and flame.

9.1.6.2 Sample volatility is evaluated by placing a filled, weighed ampoule in the combustion chamber head with the system at test temperature and pressure for 10 min. A weight loss from the ampoule greater than 10 % after exposure indicates that the volatility is sufficiently great to require sealing.

9.1.7 Open the air inlet valve and allow the combustion chamber to attain atmospheric pressure. Remove the seal cap

on the insertion rod guide and place the ampoule-insertion rod inside the guide. Replace the seal cap.

9.1.8 Reestablish the temperature and pressure of the system at the appropriate level and record the temperature and pressure for at least 10 min to reaffirm that the system is free of leaks.

9.1.9 Activate the electromagnet and rapidly move the magnet forward to drive the insertion rod into the furnace and smash the ampoule against the anvil affixed to the thermocouple support. Mark the start of test on the recorder. Withdraw the insertion rod with magnet immediately after the sample has been delivered. Perform this entire operation as quickly as possible (no longer than 4 s).

9.1.10 Continue to record the temperature and other reaction parameters of the combustion chamber for a minimum of 10 min after sample introduction or until there is no evidence of continuing reaction within the system. Record the initial equilibrium temperature as the test temperature.

9.1.10.1 Reactions are detected by the evolution of heat that increases the temperature or pressure, etc., of the combustion chamber in a characteristic manner depending on the type of reaction. The measurement of temperature is presently the principal method for following the reactions of the sample. A pressure transducer, optical device, or other suitable detectors may also be used.