Designation: E1953 – 14

Standard Practice for Description of Thermal Analysis and Rheology Apparatus[1]

This standard is issued under the fixed designation E1953; 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 practice covers generic descriptions of apparatus used for thermal analysis or rheometry measurements and its purpose is to achieve uniformity in description of thermal analysis, rheometry, and viscometer instrumentation throughout standard test methods. These descriptions are intended to be used as templates for inclusion in any test method where the thermal analysis instrumentation described herein is cited.

1.2 Each description contains quantifiable instrument performance requirements to be specified for each test method.

1.3 The values stated in SI units are to be regarded as standard. No other units of measurement are included in this standard.

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.*

2. Referenced Documents

2.1 *ASTM Standards:*[2]

E473 Terminology Relating to Thermal Analysis and Rheology

E1142 Terminology Relating to Thermophysical Properties

IEEE/ASTM SI 10 Standard for Use of the International System of Units (SI): The Modern Metric System

3. Terminology

3.1 Technical terms used in this document are found in Terminologies E473 and E1142 and Standard IEEE/ASTM SI 10.

4. Significance and Use

4.1 Section 5 identifies essential instrumentation and accessories required to perform thermal analysis, rheometry, or viscometry for a variety of different instruments. The appropriate generic instrument description should be included in any test method describing use or application of the thermal analysis, rheometry, or viscometry instrumentation described herein.

4.2 Units included in these descriptions are used to identify needed performance criteria and are considered typical. Other units may be used when including these descriptions in a specific test method. Items underlined constitute required inputs specifically established for each test method (for example, sensitivity of temperature sensor).

4.3 Additional components and accessories may be added as needed, with the appropriate performance requirements specified. Items listed in these descriptions but not used in a test method (for example, vacuum system) may be deleted.

5. Apparatus

5.1 *Differential Scanning Calorimeter* (DSC)—The essential instrumentation required to provide the minimum differential scanning calorimetric capability for this method includes:

5.1.1 *DSC Test Chamber* composed of:

5.1.1.1 A furnace(s) to provide uniform controlled heating or cooling of a specimen and reference to a constant temperature or at a constant rate within the applicable temperature range of this method.

5.1.1.2 A temperature sensor to provide an indication of the specimen temperature to ± _____ K.

5.1.1.3 Differential sensors to detect a heat flow (power) difference between the specimen and reference with a range of _____ mW and a sensitivity of ± _____ μW.

5.1.1.4 A means of sustaining a *test chamber environment* of _____ at a purge rate of mL/min ± _____ mL/min.

NOTE 1—Typically, _____ % pure nitrogen, argon, or helium is employed when oxidation in air is a concern. Unless effects of moisture are to be studied, use of dry purge gas is recommended and is essential for operation at subambient temperatures.

5.1.2 A *temperature controller*, capable of executing a specific temperature program by operating the furnace(s) between selected temperature limits at a rate of temperature change of _____ K/min constant to ± _____ K/min (list

[1] This practice is under the jurisdiction of Committee E37 on Thermal Measurements and is the direct responsibility of Subcommittee E37.10 on Fundamental, Statistical and Mechanical Properties.

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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.

cooling requirements separately if different) or at an isothermal temperature constant to ± ______ K.

5.1.3 A *data collection device*, to provide a means of acquiring, storing, and displaying measured or calculated signals, or both. The minimum output signals required for DSC are heat flow, temperature, and time.

5.1.4 *Containers* (pans, crucibles, vials, lids, closures, seals, etc.) that are inert to the specimen and reference materials and that are of suitable structural shape and integrity to contain the specimen and reference in accordance with the specific requirements of this test method including:

5.1.5 *Pressure/Vacuum System* consisting of:

5.1.5.1 A pressure vessel or similar means of sealing the test chamber at any applied pressure within the pressure limits required for this method.

5.1.5.2 A source of pressurized gas or vacuum capable of sustaining a regulated gas pressure in the test chamber of between ______ Pa and ______ Pa.

5.1.5.3 A pressure transducer or similar device to measure the pressure inside the test chamber to ±_______ %, including any temperature dependence of the transducer.

NOTE 2—The link between test chamber and pressure transducer should allow fast pressure equilibration to ensure accurate recording of the pressure above the specimen during testing.

5.1.5.4 A pressure regulator or similar device to adjust the applied pressure in the test chamber to ± ______ % of the desired value.

5.1.5.5 A ballast or similar means to maintain the applied pressure in the test chamber constant to ± ______ Pa or ± ______ %.

5.1.5.6 Valves to control the gas or vacuum environment in the test chamber or to isolate components of the pressure/vacuum system, or both.

5.1.6 Auxiliary instrumentation considered necessary or useful for conducting this method includes:

5.1.6.1 A cooling capability to hasten cool down from elevated temperatures, to provide constant cooling rates, or to sustain an isothermal subambient temperature.

5.1.6.2 A balance to weigh specimens or containers (pans, crucibles, vials, etc.), or both, to ± ______ mg.

5.1.6.3 A means, tool, or device to close, encapsulate, or seal the container of choice.

5.2 *Thermomechanical Analyzer* (TMA)—The essential instrumentation required to provide the minimum thermomechanical analytical or thermodilatometric capability for this method includes:

5.2.1 A rigid specimen holder of inert low expansivity material ______ μm/(m-K) to center the specimen in the furnace and to fix the specimen to mechanical ground.

5.2.2 A rigid (expansion, compression, flexure, tensile, etc.) *probe* of inert low expansivity material ______ μ m/(m-K) which contacts the specimen with an applied compressive or tensile force.

5.2.3 Rigid specimen clamps of inert low expansivity material ______ μm/(m-K) that grip the specimen between the rigid specimen holder and the rigid probe without distortion ______ or slippage______ [for tensile or flexure mode only].

5.2.4 A sensing element linear over a minimum range of______ mm to measure the displacement of the rigid __ probe to ± __ μm resulting from changes in length/height of the specimen.

5.2.5 A weight or force transducer to generate a constant force of ______± ______ [or between _____ and ______± ______] that is applied through the rigid ______probe to the specimen.

5.2.6 A furnace to provide uniform controlled heating or cooling of a specimen to a constant temperature or at a constant rate within the applicable temperature range of this method.

5.2.7 A temperature controller capable of executing a specific temperature program by operating the furnace between selected temperature limits at a rate of temperature change of ______ K/min constant to ± ______ K/min [list cooling requirements separately if different] or at an isothermal temperature constant to ± ______ K.

5.2.8 A temperature sensor to provide an indication of the specimen/furnace temperature to ± ______ K.

5.2.9 A means of sustaining an environment around the specimen of ______ at a purge rate of ______ mL/min ± ______.

NOTE 3—Typically, ______ % pure nitrogen, argon, or helium is employed when oxidation in air is a concern. Unless effects of moisture are to be studied, use of dry purge gas is recommended and is essential for operation at subambient temperatures.

5.2.10 A *data collection device*, to provide a means of acquiring, storing, and displaying measured or calculated signals, or both. The minimum output signals required for TMA are a change in linear dimension, temperature, and time.

5.2.11 Auxiliary instrumentation considered necessary or useful in conducting this method includes:

5.2.11.1 A cooling capability to hasten cool down from elevated temperatures, to provide constant cooling rates or to sustain an isothermal subambient temperature.

5.2.11.2 Micrometer or other *measuring device* to determine specimen dimensions of _____ mm ± _____ mm.

5.2.11.3 A balance with a minimum capacity of __ mg to weigh specimens or clamps, or both, to ± ______ mg.

5.3 *Thermogravimetric Analyzer* (TGA)—The essential instrumentation required to provide the minimum thermogravimetric analytical capability for this method includes:

5.3.1 A thermobalance composed of:

5.3.1.1 A furnace to provide uniform controlled heating or cooling of a specimen to a constant temperature or at a constant rate within the applicable temperature range of this method.

5.3.1.2 A temperature sensor to provide an indication of the specimen/furnace temperature to ± ______ K.

5.3.1.3 A continuously recording balance to measure the specimen mass with a minimum capacity of _____ mg and a sensitivity of ± ______ μg.

5.3.1.4 A means of maintaining the specimen/container under atmospheric control of ______ of ______ % purity at a purge rate of ______ L/min ± _____.

NOTE 4—Excessive purge rates should be avoided as this may introduce interferences due to turbulance effects and temperature gradients.