



Standard Test Methods for Polyurethane Raw Materials: Determination of Viscosity of Crude or Modified Isocyanates¹

This standard is issued under the fixed designation D4889; 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 These test methods (A and B) determine the viscosity of crude or modified *isocyanates*. They are applicable to products derived from toluene diisocyanate, methylene di(phenylisocyanate), and polymeric (methylene phenylisocyanate) (see [Note 1](#)).

NOTE 1—Test method A includes a procedure for measuring dynamic viscosity using a rotational instrument. Test method B is simply a reference to a general procedure for measuring kinematic viscosity, [D445](#).

1.2 The values stated in SI units are to be regarded as standard.

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.* For specific hazards statement, see [Warning](#) at the end of [5.1](#).

NOTE 2—This standard is equivalent to ISO 3219 and ISO 3104.

2. Referenced Documents

2.1 ASTM Standards:²

[D445 Test Method for Kinematic Viscosity of Transparent and Opaque Liquids \(and Calculation of Dynamic Viscosity\)](#)

[D883 Terminology Relating to Plastics](#)

[E2251 Specification for Liquid-in-Glass ASTM Thermometers with Low-Hazard Precision Liquids](#)

2.2 ISO Standards:³

[ISO 3104 Petroleum Products—Transparent and Opaque Liquids—Determination of Kinematic Viscosity and Cal-](#)

[culation of Dynamic Viscosity](#)
[ISO 3219 Plastics—Polymers/Resins in the Liquid State or as Emulsions or Dispersions—Determination of Viscosity Using a Rotational Viscometer with Defined Shear Rate](#)

3. Terminology

3.1 *Definitions*—For definitions of terms used in these test methods see Terminology [D883](#).

4. Significance and Use

4.1 These test methods can be used for research, quality control, or specification tests to characterize *isocyanates* used in polyurethane products.

4.2 Viscosity measures the resistance of a fluid to uniform continuous flow without turbulence or other forces.

4.3 Some isocyanates exhibit non-Newtonian behavior under certain conditions. Whenever possible, generate results for comparison under the same conditions, that is, the same spindle/speed combination for rotational viscosity and the same tube size for kinematic viscosity.

5. Sampling

5.1 Since organic *isocyanates* react with atmospheric moisture, take special precautions in sampling. Usual sampling methods, even when conducted rapidly, can cause contamination of the sample with insoluble urea. Therefore, blanket the sample with dry air or nitrogen at all times. (**Warning**—*Diisocyanates* are eye, skin and respiratory irritants at concentrations above the occupational exposure limit (TLV or PEL). *Diisocyanates* can cause skin and respiratory sensitization (asthma) in some people. Once sensitized, it is essential to limit further exposure to diisocyanates. Use a combination of engineering controls and personal protective equipment, including respiratory, skin and eye protection, to prevent over-exposure to diisocyanates. Consult the product suppliers' Safety Data Sheet (SDS) for more detailed information about potential health effects and other specific safety and handling instructions for the product.)

6. Test Conditions

6.1 Since isocyanates react with moisture, keep laboratory humidity low, preferably about 50 % relative humidity. See [Warning](#) in [5.1](#).

¹ These test methods are under the jurisdiction of ASTM Committee D20 on Plastics and are the direct responsibility of Subcommittee D20.22 on Cellular Materials - Plastics and Elastomers.

Current edition approved May 1, 2015. Published March 2011. Originally approved in 1988. Last previous edition approved in 2011 as D4889 - 04(2011). DOI: 10.1520/D4889-15.

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.

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

TEST METHOD A—ROTATIONAL VISCOSITY

7. Summary of Test Method

7.1 The viscosity is measured by determining the torque on a spindle rotating at constant speed in the liquid sample which is adjusted to $25 \pm 0.1^\circ\text{C}$. Generation of comparative data using this method requires agreement on the speed, spindle, temperature, time of rotation and torque range of the instrument used.

8. Apparatus

8.1 *Constant-Temperature Bath*, capable of maintaining a temperature of $25 \pm 0.1^\circ\text{C}$ is to be used. Water, water and glycerin, or oil is used as the heating medium and the bath is to be provided with heating, circulating, and thermostating devices.

8.2 *Bath and Sample Thermometers*, graduated in 0.1°C subdivisions and standardized for the range of use to the nearest 0.01°C . ASTM Saybolt Viscosity Thermometers having ranges from 19 to 27°C and 49 to 57°C , as specified, and conforming to the requirements for Thermometers S117C and S64C, respectively, as prescribed in Specification **E2251** are recommended. Any other thermometric device of equal or better accuracy is also acceptable.

8.3 *Rotational Viscometer*, capable of user defined speed and spindle combinations. An instrument that is capable of providing the shear rate is recommended. The calibration of the instrument is to be checked periodically by measuring the viscosity of NIST traceable standard fluids.

9. Solvent

9.1 *Cleaning Solvent—dichloromethane or acetone*, reagent grade. Any solvent in which the isocyanate is completely miscible is acceptable.

10. Preparation of Sample

10.1 The preparation of a homogeneous sample is of primary importance in viscosity measurements. A non-uniform temperature distribution as well as the presence of air bubbles and traces of extraneous material are to be avoided. The sample must be thoroughly mixed and the temperature measured at several locations in the sample vessel before determining the viscosity.

11. Preparation of Apparatus

11.1 Follow the manufacturer's instructions to set up the instrument and ensure that the viscometer is level.

12. Choice of Temperature

12.1 Samples that are liquid and have a viscosity of less than 100 000 mPa·s(cP) at 25°C are to be measured at 25°C .

12.2 In cases of interlaboratory studies and higher viscosity samples, all parties are to agree upon a set measurement temperature.

13. Choice of Spindle and Rotational Speed

13.1 Rotational Viscometers offer a variety of spindle size and rotational speeds. In the case of non-Newtonian liquids,

changing these factors will cause variation in the results obtained. In general, the following recommendations provide guidance for choosing the spindle size and speed to be used for a specific sample.

13.1.1 The combination chosen shall generate a torque value between 15 and 90% of full scale, or that specified by the instrument manufacturer.

13.1.1.1 If more than one speed/spindle combination will fulfill the requirement of **13.1.1**, the combination with the higher speed will provide higher accuracy and the combination with the lower speed will minimize certain types of non-Newtonian behavior.

13.1.1.2 There must be agreement between the testing laboratory and the submitter on the spindle/speed selection.

14. Procedure

14.1 Using the smallest container recommended by the manufacturer, place sufficient sample to cover the immersion mark on the viscometer spindle. Cover the container and immerse it to the sample level in a constant temperature bath. Stir occasionally without trapping air bubbles. Check the temperature at several different locations in the container to make sure uniformity has been achieved.

14.2 After the desired temperature has been observed throughout the sample for 10 min, immerse the viscometer spindle (and the guard when recommended by the manufacturer) into a sample to the immersion line marked on the spindle. Exercise caution to avoid air bubbles gathering under the spindle during immersion. If bubbles are observed, detach the spindle, keeping it in the sample, and stir until the bubbles are released. Reattach the spindle.

14.3 Follow the manufacturer's instructions to measure the viscosity for the sample using a 15 second rotation time.

14.4 After the analysis, spindles are cleaned with a solvent appropriate for the isocyanate and equipment used, for example, dichloromethane or acetone.

15. Calculation

15.1 Multiply the reading by the factor provided by the manufacture for the speed/spindle combination used to convert the instrument reading to the viscosity in mPa·s (cP). Most instruments automatically perform this calculation.

16. Report

16.1 Report the following information:

16.1.1 Temperature of test,

16.1.2 Model of viscometer,

16.1.3 Speed of rotation,

16.1.4 Spindle number, and

16.1.5 Viscosity in millipascal seconds (centipoises) [mPa·s(cP)].

17. Precision and Bias

17.1 *Precision*—Attempts to develop a precision and bias statement for this test method have not been successful; however, the precision is expected to be equivalent to that reported by the instrument manufacturer. For this reason, data

on precision and bias cannot be given. Because this test method does not contain a numerical precision and bias statement, it shall not be used as a referee test method in case of dispute. Anyone wishing to participate in the development of precision and bias data should contact the Chairman, Subcommittee D 20.22 (Section D 20.22.01), ASTM, 100 Barr Harbor Drive, West Conshohocken, PA 19428.

17.2 *Bias*—The bias of this test method has not yet been determined.

TEST METHOD B—KINEMATIC VISCOSITY

18. General:

18.1 A general test method for kinematic viscosity which applies to *isocyanates* as well as other materials is published in Test Method **D445**.

19. Keywords

19.1 dynamic viscosity; isocyanates; kinematic viscosity; polyurethane raw materials; rotational viscosity; viscosity

SUMMARY OF CHANGES

Committee D20 has identified the location of selected changes to this standard since the last issue (D4889–04(2011)) that may impact the use of this standard. (May 1, 2015)

(1) Subsection 1.1: Scope—Changed naming of methylene di(phenylisocyanate) to be consistent with the other polyurethane raw material methods.

(2) Section 1, NOTE 1: Scope—Numerous changes throughout to remove specific instrument manufacturer names and make the method more applicable to all instruments.

(3) Section 1, NOTE 2: Scope—Updated ISO equivalency.

(4) Subsection 2.2: ISO Standards—Added equivalent ISO standards.

(5) Subsection 5.1: Sampling—The previous warning statement was edited to remove non-mandatory language and updated with input from the Center for the Polyurethanes Industry's (CPI) Product Stewardship Committee.

(6) Subsection 8.2: Apparatus—Changed reference from E1 to E2251 to allow for use of non-mercury thermometers.

(7) Section 14: Procedure—Removed parameter tables for specific instrument manufacturer and stated to use the parameters specified by the instrument manufacturer.

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). Permission rights to photocopy the standard may also be secured from the Copyright Clearance Center, 222 Rosewood Drive, Danvers, MA 01923, Tel: (978) 646-2600; <http://www.copyright.com/>