



Designation: D2273 – 08 (Reapproved 2016)

Standard Test Method for Trace Sediment in Lubricating Oils¹

This standard is issued under the fixed designation D2273; 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.

This standard has been approved for use by agencies of the U.S. Department of Defense.

1. Scope

1.1 This test method covers the determination of trace amounts (less than 0.05 % by volume) of sediment in lubricating oils. Since oil-soluble material precipitated by the specified solvent is not intended as part of the measured sediment, the test method is not applicable in cases where precipitated oil-soluble components will appreciably contribute to the sediment readings.

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

2. Referenced Documents

2.1 ASTM Standards:²

D4057 Practice for Manual Sampling of Petroleum and Petroleum Products

D4177 Practice for Automatic Sampling of Petroleum and Petroleum Products

3. Terminology

3.1 Definitions of Terms Specific to This Standard:

3.1.1 *trace sediment, n*—the number of millilitres of sediment precipitated from 100 mL of oil sample (volume percent) when equal parts of the oil sample and the specified solvent are mixed and centrifuged under the prescribed conditions.

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 D02.06 on Analysis of Liquid Fuels and Lubricants.

Current edition approved Oct. 1, 2016. Published November 2016. Originally approved in 1964. Last previous edition approved in 2012 as D2273 – 08 (2012). DOI: 10.1520/D2273-08R16.

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.

4. Significance and Use

4.1 This test measures the trace level amount of sediment that is naphtha-insoluble and can be separated by centrifuging. Excessive amounts of sediment in oil could lead to system malfunction in critical applications.

5. Apparatus

5.1 *Centrifuge*, meeting all the safety requirements for normal use and capable of whirling two or more filled centrifuge tubes at a speed which can be controlled to give a relative centrifugal force (rcf) between 600 and 700 at the tip of the tubes. The revolving head, trunnion rings, and trunnion cups, including the rubber cushion, shall be soundly constructed to withstand the maximum centrifugal force capable of being delivered by the power source. The trunnion cups and cushions shall firmly support the tubes when the centrifuge is in motion. The centrifuge shall be enclosed by a metal shield or case strong enough to eliminate danger if any breakage occurs. Calculate the speed of the rotating head as follows:

$$\text{rpm} = 1337 \sqrt{\text{rcf}/d} \quad (1)$$

where:

rcf = relative centrifugal force, and

d = diameter of swing, in millimetres, measured between tips of opposite tubes when in rotating position.

The relationship between the diameter swing, relative centrifugal force, and revolutions per minute is given in Table 1.

5.2 *Centrifuge Tube*, cone-shaped, conforming to the dimensions given in Fig. 1, and made of thoroughly annealed glass. The graduations, numbered as shown in Fig. 1, shall be clear and distinct, and the mouth shall be constructed in a shape suitable for closure with a cork. Scale-error tolerances and smallest graduations between various calibration marks are given in Table 2. Calibrated centrifuge tubes shall be purchased from the manufacturer.

6. Reagents

6.1 *Hexanes*, reagent grade, minimum purity. (**Warning**—Extremely flammable. Harmful if inhaled.) See also Note 1.

NOTE 1—Reagent grade minimum purity hexanes are sometimes referred to or sold by other names such as precipitation naphtha,

TABLE 1 Rotation Speeds for Centrifuges of Various Diameters

Diameter of Swing, mm ^A	RPM at 600 rcf	RPM at 700 rcf
483	1490	1610
508	1450	1570
533	1420	1530
559	1390	1500

^A Measured in mm between tips of opposite tubes when rotating-position.

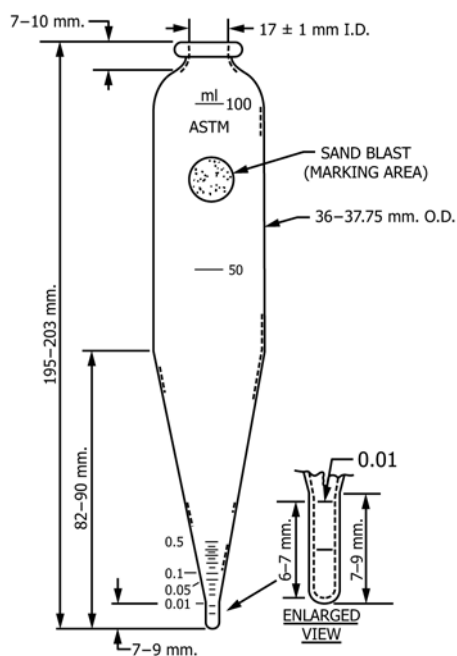


FIG. 1 Trace Sediment Tube

TABLE 2 Trace Sediment Tube Calibration Tolerances

Range, mL	Smallest Scale Division, mL	Scale Error, max, mL
0 to 0.01	0.005	±0.001 at 0.01
0.01 to 0.05	0.01	±0.005
0.05 to 0.15	0.05	±0.01
0.15 to 0.30	0.05	±0.02
0.30 to 0.50	0.05	±0.03
0.50 to 50	none	±1.0
50 to 100	none	±1.0

petroleum naphtha, petroleum ether, ligroine, petroleum benzin, or industrial naphthas.

NOTE 2—Before use, the hexanes should be free of any extraneous material that might affect the final test readings. For this purpose, it should either be filtered through a membrane filter or centrifuged several times and decanted just prior to its use.

7. Sampling

7.1 Refer to Practice D4057 (manual) or D4177 (automatic) for recommended practices for obtaining samples.

7.2 The sample shall be thoroughly representative of the material in question and the portion used for the test shall be thoroughly representative of the sample itself. This requires vigorous agitation of the sample immediately before transferring the sample to the tube. The difficulties in obtaining representative samples for this determination are unusually great; hence, the importance of sampling cannot be too strongly emphasized.

8. Procedure

8.1 Measure 50 mL ± 1 mL of hexanes (see 6.1) into each of two clean, dry centrifuge tubes at room temperature. (**Warning**—Extremely flammable.) Then fill each tube to the 100 mL mark with the oil sample and close tightly with a softened cork covered with a thin pliable plastic film that is resistant to petroleum products (not a rubber stopper). Shake sample well to ensure complete mixing, then invert each tube at least 20 times allowing the liquid to drain thoroughly from the tip of the tube at each inversion. If the liquid refuses to drain upon inversion, gently tap the inverted tube against the palm of the hand to jolt the liquid out of the tip. Place the tubes in a water bath at 32 °C to 35 °C for 5 min ± 1 min. Momentarily remove the corks to relieve any pressure, and invert each tube again at least 20 times exactly as before. The success of this method depends to a large degree upon having a thoroughly homogeneous mixture which will drain completely as possible from the tip when the tube is inverted.

8.2 Balance the two centrifuge tubes or pairs of tubes with their respective trunnion cups and place them on opposite sides of the centrifuge head. Then whirl them for 10 min ± 1 min at a rate sufficient to produce a relative centrifuge force (rcf) between 600 and 700 at the tips of the whirling tubes (see 5.1). At the end of the 10 min whirling period, decant the mixture carefully, allowing the sediment to remain in the tubes. Measure another 50 mL ± 1 mL of the hexanes into each of the two tubes, then add the oil sample until the total volume reaches the 100 mL mark. Then stopper the tubes, repeatedly invert, heat, and again invert as described in 8.1. Whirl the tubes in the centrifuge for 10 min ± 1 min as before, repeating the 10 min whirling operation until the volume of sediment in each tube remains constant for three consecutive readings. In general, not more than four whirlings will be required for oils having low sediment values. Record the final reading for sediment in each tube.

9. Calculation

9.1 If sediment is present in one or both tubes, as recorded in 8.2, average the final readings for sediment in the two tubes containing the sample to obtain the average volume of sediment per 100 mL sample.

10. Report

10.1 If no sediment is present in either tube, as determined and recorded in 8.2, report the sample result as 0.

10.2 If the average result determined in 9.1 is >0 % and <0.01 %, report the result to the nearest 0.001 %.

10.3 If the average result determined in 9.1 is within the range of 0.01 % to 0.05 %, report the result to the nearest 0.01 %.

11. Precision and Bias

11.1 The precision of this test method as determined by statistical examination of interlaboratory results is as follows:

11.1.1 *Repeatability*—The difference between two test results, obtained by the same operator with the same apparatus under constant operating conditions on identical test material,

would in the long run, in the normal and correct operation of the test method, exceed the following values only in one case in twenty:

Sediment, percent volume	Repeatability
0.000 to 0.002	0.001
0.003 to 0.005	0.001
0.006 to 0.01	0.002

11.1.2 *Reproducibility*—The difference between two single and independent results obtained by different operators working in different laboratories on identical test material would, in the long run, in the normal and correct operation of the test method, exceed the following values only in one case in twenty:

Sediment, percent volume	Reproducibility
0.000 to 0.002	0.001
0.003 to 0.005	0.002
0.006 to 0.01	0.003

11.1.3 *Bias*—Test Method D2273 is empirical and no statement of bias can be made.

12. Keywords

12.1 lubricating oils; trace sediment

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/>