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Designation: D 3117 – 03

Standard Test Method for Wax Appearance Point of Distillate Fuels¹

This standard is issued under the fixed designation D 3117; 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.

 ϵ^1 Note— Figure was corrected editorially in October 1997.

¹ This test method is under the jurisdiction of ASTM Committee D-2 D02 on Petroleum Products and Lubricants and is the direct responsibility of Subcommittee D02.07 on Flow Properties.

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1. Scope *

1.1 This test method covers the detection of wax in burner fuels, diesel fuels, and turbine engine fuels in the range from -26 to $+2^{\circ}$ C. It is applicable to a dark-colored oil if the stirrer can be seen under illumination.

1.2 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 hazard statements, see Notes 2 through 5. Section 7.

2. Referenced Documents

2.1 ASTM Standards:

D-2386 Test 445 Test Method for Freezing Point Kinematic Viscosity of Aviation Fuels Transparent and Opaque Liquids² D-2500 Test 2386 Test Method for Freezing Point of Aviation Fuels²

D 2500 Test Method for Cloud Point of Petroleum-Oils Products²

E 1 Specification for ASTM Thermometers³

3. Terminology

3.1 Definitions of Terms Specific to This Standard:

3.1.1 *wax appearance point*—the temperature at which wax first begins to separate from the liquid when it is cooled under prescribed conditions.

4. Summary of Test Method

4.1 A specimen of distillate fuel is cooled under prescribed conditions while being stirred. The temperature at which wax first appears is recorded as the wax appearance point.

5. Significance and Use

5.1 Wax appearance point is the temperature at which wax crystals begin to precipitate out of a fuel under specified cooling conditions. The presence of wax crystals in the fuel may restrict flow or plug the fuel filter. In critical fuel systems, wax appearance point may define the lower limit of acceptable operability.

6. Apparatus (Fig. 1) Apparatus

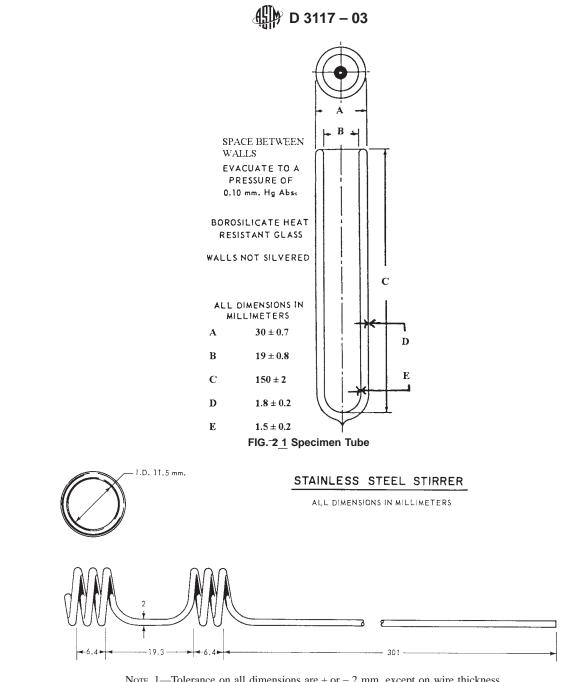
6.1 Specimen Tube—A double-walled (Dewar-type) jacketed tube with dimensions shown in Fig.-2_1.

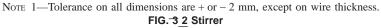
6.2 *Temperature Measuring Device*—Liquid-in-glass thermometer—C_conforming to specifications for ASTM Thermometer 62C in accordance with Specification E 1, or any other temperature measuring device with equal or better accuracy and equal temperature response.

6.3 *Stirrer Assembly*—A stainless steel wire configured in the manner described in Fig. <u>3.2</u> and manipulated by a motor or other suitable device in a vertical direction. The frequency of movement shall be 55 ± 5 cycles/min with an amplitude of 50 ± 5 mm. The stirrer shall be concentric with the <u>thermometer temperature measuring device</u> and shall be fitted with the moisture proof collar specified in Test Method D 2386. A No. 3, two-hole neoprene rubber stopper shall be used to seal the top of the specimen tube.

² Annual Book of ASTM Standards, Vol 05.01.

³ Annual Book of ASTM Standards, Vol 14.03.





6.4 *Cooling Bath*—Use an unsilvered vacuum flask having minimum dimensions of 200-mm depth and 65-mm internal diameter. The bath temperature, below –45°C, may be maintained by refrigeration or suitable freezing mixtures (Note 27.1). Bath temperature is monitored with an appropriate thermometer temperature measuring device such as ASTM Thermometer 6C/IP2C 6C.

NOTE 1—Solid carbon dioxide chips (dry ice) and isopropanol is a recommended mixture for coolant. An excess of dry ice should be avoided to prevent obscuring the sample tube in a continuous stream of bubbles. Isopropanol should be replaced daily or when low temperature viscosity is noticeably higher than a fresh bath. Liquid nitrogen may also be used as coolant instead of liquids cooled with solid carbon dioxide.

6.5 *Illumination*—A 150 to 230-mm long, 5 to 8-W fluorescent tube shall be mounted behind the specimen to illuminate it with transmitted light. Observations shall be made with the sample tube between the observer's eye and the lamp.

6.6 *Clock*—Use a clock or other timing device readable to 10 s to monitor the cooling rate.

7. Materials

7.1 *Carbon Dioxide (Solid) or Dry Ice*—(Warning—see Note 2).— (Warning—Extremely cold, -78°C.) A commercial grade of dry ice is suitable for use in the cooling bath (Note-6).

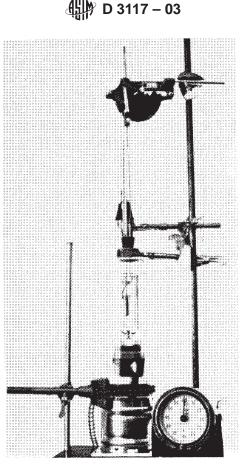


FIG. 1 3 Complete Assembly

Note 2—Warning: Extremely cold, - 78°C. 2).

7.2 Isopropanol or Isopropyl Alcohol—(Warning—see Note 3).— (Warning—Extremely flammable.) A commercial or technical grade of isopropanol is suitable for the cooling bath. Note 3—Warning: Extremely flammable. 7.3 Liquid

<u>7.3 Liquid Nitrogen (Warning—see Note 4).</u> (Warning—Extremely cold, –196°C.) A commercial or technical grade of liquid nitrogen is suitable for the cooling bath. See Note-6.

Note 4-Warning: Extremely cold, - 196°C.

Note 5—Warning: Carbon 2. (Warning— Carbon dioxide (solid) and liquid nitrogen liberate gases that can cause suffocation. Contact with skin causes burns, freezing, or both.)

8. Procedure

8.1 Check

<u>8.1 Ensure that</u> the thermometer each day the test temperature measuring device is run by immersing it in an ice-water slurry. An acceptable thermometer will show $0 \pm 0.1^{\circ}$ C. calibration. For liquid-in-glass temperature measuring devices, see the Annex on Calibration in Test Method D 445.

Note 62—Because ambient temperatures are well above the range of <u>ASTM</u> Thermometer 62C, the <u>mercury liquid-in-glass</u> thread will extend upward into the expansion chamber. During cooling, the <u>thermometer liquid-in-glass</u> temperature measuring device must be examined to make certain that no mercury separation occurs. If a low reading is obtained <u>observed</u> in the ice-water slurry, it may be due to separation. Warm the thermometer, tap until the mercury no longer shows any separation, and retest. The thermometer should be stored in a vertical position with the mercury bulb at the bottom. liquid.

8.2 Sample temperature shall not be lower than 10°C when starting the measurement. Dry the specimen by filtration through a lintless filter paper.

8.3 Assemble the unit as shown in Fig.-1.-3.

8.4 With the oil at 10°C or above, introduce a 25 \pm 1-mL specimen into the tube. Add three drops of anhydrous isopropanol. 8.5 Adjust the stopper, thermometer, temperature measuring device, and stirrer so that the bottom of the thermometer bulb temperature measuring device is 25 mm above the bottom of the specimen tube and the stirrer does not break the surface at the upper end of its stroke. 8.6 When the stopper has been firmly seated, start the stirrer, raise the cooling bath around the specimen tube, and cool rapidly to 2° C.

8.7 Once the thermometer temperature reading is within the graduated region, below 2° C, adjust the bath height to keep the cooling rate between 1 and 2° C/min. The preferred cooling rate is 1.5° C/min.

8.8 Continue cooling until wax distinctly appears when the specimen tube is examined by transmitted light. This is manifested by the appearance of very small wax crystals which make the stirring pattern quite obvious. An example of the change observed is shown in Fig. 4 and Fig. 5. Read. Observe the thermometer temperature reading immediately to the nearest 0.2°C.

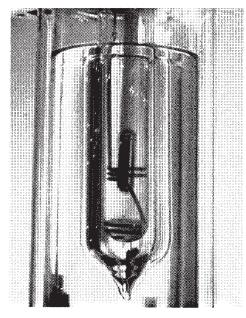


FIG. 5 Tube After Wax Appearance

NOTE 73—The wax appearance point should not be misinterpreted to be the appearance of the very first crystals. The temperature at which a distinct swirl of wax particles around the stirrer occurs is the correct interpretation.

9. Report

9.1 The temperature at which the wax crystals are obviously present is the wax appearance point and is reported to the nearest 0.2°C.

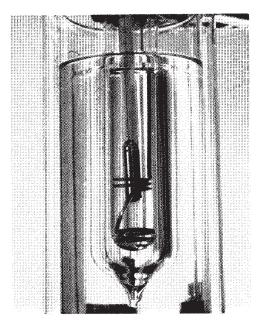


FIG. 4 Tube Before Wax Appearance

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10. Precision and Bias

10.1 *Precision*—The precision of this test method obtained by the statistical examination of interlaboratory test results is as follows:

10.1.1 *Repeatability*—The difference between two test results obtained by the same operator with the same apparatus under constant operating conditions on the same fuel sample would, in the long run, in the normal and correct operation of the test method, exceed 0.8° C only in one case in twenty.

10.1.2 *Reproducibility*—The difference between two single and independent results obtained by different operators working in different laboratories on the same fuel sample would, in the long run, in the normal and correct operation of the test method, exceed 2.2°C only in one case in twenty.

10.2 *Bias*—Because liquid hydrocarbon standards of known wax appearance point are not available, no statement of bias can be made.

Note 84—These precision data were obtained in a cooperative program among eight laboratories testing eight fuels for both wax appearance point and cloud point by Test Method D 2500. None of the fuels contained wax crystal modifying additives.

11. Keywords

11.1 distillate fuels; wax

SUMMARY OF CHANGES

Subcommittee D02.07 has identified the location of selected changes to this standard since the last issue (D 3117–96^{ϵ 1}) that may impact the use of this standard.

(1) Added Test Method D 445 to Section 2, Referenced Documents. (2) Added "temperature measuring device" throughout.

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