



Designation: D97 – 17b (Reapproved 2022)



Designation: 15/22

Standard Test Method for Pour Point of Petroleum Products^{1,2}

This standard is issued under the fixed designation D97; 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 and is intended for use on any petroleum product.³ A procedure suitable for black specimens, cylinder stock, and nondistillate fuel oil is described in 8.8. The cloud point procedure formerly part of this test method now appears as Test Method **D2500**.

1.2 Currently there is no ASTM test method for automated Test Method D97 pour point measurements.

1.3 Several ASTM test methods offering alternative procedures for determining pour points using automatic apparatus are available. None of them share the same designation number as Test Method D97. When an automatic instrument is used, the ASTM test method designation number specific to the technique shall be reported with the results. A procedure for testing the pour point of crude oils is described in Test Method **D5853**.

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

1.5 **WARNING**—Mercury has been designated by many regulatory agencies as a hazardous substance that can cause serious medical issues. Mercury, or its vapor, has been demonstrated to be hazardous to health and corrosive to materials. Use caution when handling mercury and mercury-containing products. See the applicable product Safety Data Sheet (SDS) for additional information. The potential exists that selling

mercury or mercury-containing products, or both, is prohibited by local or national law. Users must determine legality of sales in their location.

1.6 *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, health, and environmental practices and determine the applicability of regulatory limitations prior to use.*

1.7 *This international standard was developed in accordance with internationally recognized principles on standardization established in the Decision on Principles for the Development of International Standards, Guides and Recommendations issued by the World Trade Organization Technical Barriers to Trade (TBT) Committee.*

2. Referenced Documents

2.1 ASTM Standards:⁴

- D117** Guide for Sampling, Test Methods, and Specifications for Electrical Insulating Liquids
- D396** Specification for Fuel Oils
- D2500** Test Method for Cloud Point of Petroleum Products and Liquid Fuels
- D5853** Test Method for Pour Point of Crude Oils
- D6300** Practice for Determination of Precision and Bias Data for Use in Test Methods for Petroleum Products, Liquid Fuels, and Lubricants
- D7962** Practice for Determination of Minimum Immersion Depth and Assessment of Temperature Sensor Measurement Drift
- E1** Specification for ASTM Liquid-in-Glass Thermometers
- E644** Test Methods for Testing Industrial Resistance Thermometers
- E1137** Specification for Industrial Platinum Resistance Thermometers
- E2877** Guide for Digital Contact Thermometers

¹ This test method is under the jurisdiction of ASTM International Committee D02 on Petroleum Products, Liquid Fuels, and Lubricants and is the direct responsibility of ASTM Subcommittee D02.07 on Flow Properties. The technically equivalent standard as referenced is under the jurisdiction of the Energy Institute Subcommittee SC-C-4.

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This test method was adopted as a joint ASTM-IP Standard in 1965.

² This test method has been developed through the cooperative effort between ASTM and the Energy Institute, London. ASTM and IP standards were approved by ASTM and EI technical committees as being technically equivalent but that does not imply both standards are identical.

³ Statements defining this test and its significance when applied to electrical insulating oils of mineral origin will be found in Guide **D117**.

⁴ 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.

2.2 Energy Institute Standards:⁵
Specifications for IP Standard Thermometers

3. Terminology

3.1 Definitions:

3.1.1 *black oil, n*—lubricant containing asphaltic materials. Black oils are used in heavy-duty equipment applications, such as mining and quarrying, where extra adhesiveness is desired.

3.1.2 *cylinder stock, n*—lubricant for independently lubricated engine cylinders, such as those of steam engines and air compressors. Cylinder stock are also used for lubrication of valves and other elements in the cylinder area.

3.1.3 *digital contact thermometer (DCT), n*—an electronic device consisting of a digital display and associated temperature sensing probe.

3.1.3.1 *Discussion*—This device consists of a temperature sensor connected to a measuring instrument; this instrument measures the temperature-dependent quantity of the sensor, computes the temperature from the measured quantity, and provides a digital output. This digital output goes to a digital display and/or recording device that may be internal or external to the device. These devices are sometimes referred to as “digital thermometers.”

3.1.3.2 *Discussion*—PET is an acronym for portable electronic thermometers, a subset of digital contact thermometers (DCT).

3.1.4 *pour point, n—in petroleum products*, the lowest temperature at which movement of the test specimen is observed under prescribed conditions of test.

3.1.5 *residual fuel, n*—a liquid fuel containing bottoms remaining from crude distillation or thermal cracking; sometimes referred to as heavy fuel oil.

3.1.5.1 *Discussion*—Residual fuels comprise Grades 4, 5, and 6 fuel oils, as defined in Specification D396.

4. Summary of Test Method

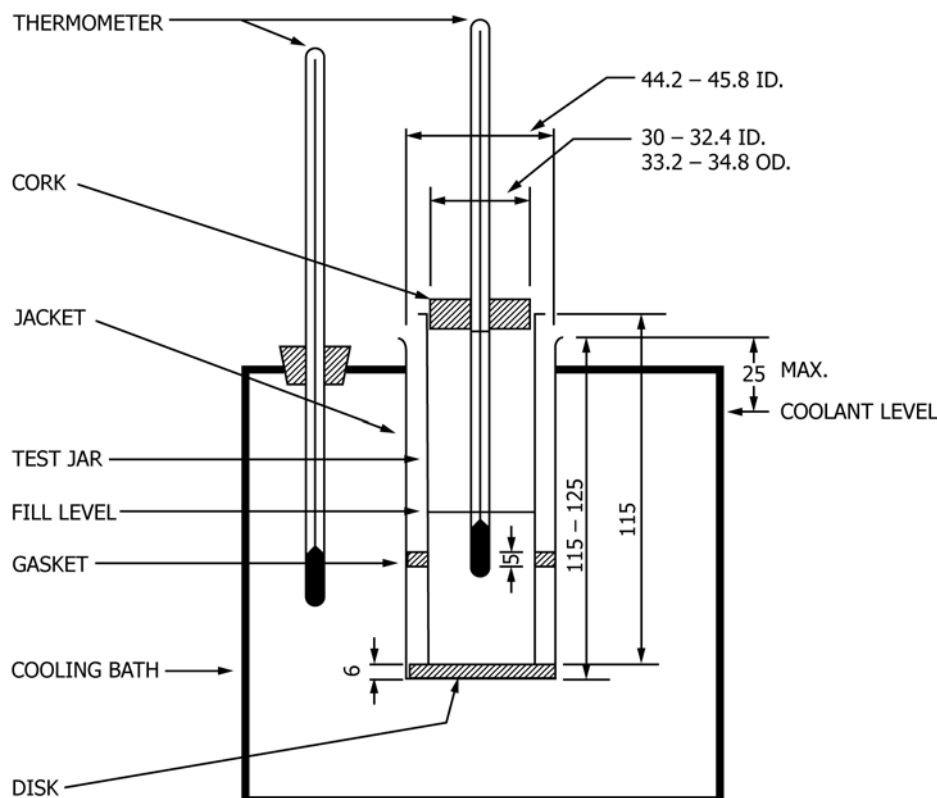
4.1 After preliminary heating, the sample is cooled at a specified rate and examined at intervals of 3 °C for flow characteristics. The lowest temperature at which movement of the specimen is observed is recorded as the pour point.

5. Significance and Use

5.1 The pour point of a petroleum specimen is an index of the lowest temperature of its utility for certain applications.

6. Apparatus

6.1 *Test Jar*, cylindrical, of clear glass, flat bottom, 33.2 mm to 34.8 mm outside diameter, and 115 mm to 125 mm in height. The inside diameter of the jar can range from 30.0 mm



NOTE 1—Dimensions are in millimetres (not to scale).

FIG. 1 Apparatus for Pour Point Test

to 32.4 mm, within the constraint that the wall thickness be no greater than 1.6 mm. The jar shall have a line to indicate a sample height 54 mm \pm 3 mm above the inside bottom. See Fig. 1.

6.2 Temperature Measuring Device—Either liquid-in-glass thermometer as described in 6.2.1 or Digital Contact Thermometer (DCT) meeting the requirements described in 6.2.2.⁶

6.2.1 Liquid-in-Glass Thermometers, having the following ranges and conforming to the requirements prescribed in Specification E1 or Specifications for IP Standard Thermometers:

Thermometer	Temperature	Thermometer Number	
	Range	ASTM	IP
High cloud and pour	–38 °C to +50 °C	5C	1C
Low cloud and pour	–80 °C to +20 °C	6C	2C
Melting point	+32 °C to +127 °C	61C	63C

6.2.1.1 Since separation of liquid column thermometers occasionally occurs and may escape detection, thermometers should be checked immediately prior to the test and used only if they prove accurate within ± 1 °C (for example ice point).

6.2.2 Digital Contact Thermometer Requirements:

Parameter	Requirement
DCT	Guide E2877 Class F or better
Nominal temperature range ^A	High Pour: –38 °C to 50 °C Low Pour: –80 °C to 20 °C Melt Point: 32 °C to 127 °C
Display resolution	0.1 °C minimum
Accuracy ^B	± 500 mK (± 0.5 °C)
Sensor type	Platinum resistance thermometer (PRT), thermistor, thermocouple
Sensor sheath ^C	4.2 mm OD maximum
Sensor length ^D	Less than 10 mm
Immersion depth ^E	Less than 40 mm per Practice D7962
Sample immersion depth	Tip of sheath between 10 mm and 15 mm below sample meniscus Fig. 1
Measurement drift ^E	Less than 500 mK (0.5 °C) per year
Response time ^F	Less than or equal to 4 s per footnote F
Calibration error	Less than 500 mK (0.5 °C) over the range of intended use
Calibration range	Consistent with temperature range of use
Calibration data	Four data points evenly distributed over the calibration range that is consistent with the range of use. The calibration data is to be included in the calibration report.
Calibration report	From a calibration laboratory with demonstrated competency in temperature calibration which is traceable to a national calibration laboratory or metrology standards body

^A The nominal temperature range may be different from the values shown provided the calibration and accuracy criteria are met.

^B Accuracy is the combined accuracy of the DCT unit, which is the display and sensor.

^C Sensor sheath is the tube that holds the sensing element. The value is the outside diameter of the sheath segment containing the sensor element.

^D The physical length of the temperature sensing element.

^E As determined by Practice D7962 or an equivalent procedure.

^F **Response Time**—The time for a DCT to respond to a step change in temperature. The response time is 63.2 % of the step change time as determined per Section 9 of Test Methods E644. The step change evaluation begins at 20 °C \pm 5 °C air to 77 °C \pm 5 °C with water circulating at 0.9 m/s \pm 0.09 m/s past the sensor.

NOTE 1—When making measurements below –40 °C with a PRT, it

may be necessary to use a 1000 Ω sensor in order to obtain accurate measurements.

NOTE 2—When the DCT display is mounted on the end to the probe's sheath, the test jar with the probe inserted will be unstable. To resolve this, it is recommended that the probe be less than 30 cm in length but no less than 15 cm. A 5 cm long stopper, that has a low thermal conductivity, with approximately half of it inserted in the sample tube will improve stability.

6.2.2.1 The DCT calibration drift shall be checked at least annually by either measuring the ice point or against a reference thermometer in a constant temperature bath at the prescribed immersion depth to ensure compliance with 6.2.2. See Practice D7962.

NOTE 3—When a DCT's calibration drifts in one direction over several calibration checks, that is, ice point, it may be an indication of deterioration of the DCT.

6.3 Cork, to fit the test jar, bored centrally for the test temperature measuring device.

6.4 Jacket, watertight, cylindrical, metal, flat-bottomed, 115 mm \pm 3 mm depth, with inside diameter of 44.2 mm to 45.8 mm. It shall be supported in a vertical position in the cooling bath (see 6.7) so that not more than 25 mm projects out of the cooling medium, and shall be capable of being cleaned.

6.5 Disk, cork or felt, 6 mm thick to fit loosely inside the jacket.

6.6 Gasket Ring Form, about 5 mm in thickness, to fit snugly around the outside of the test jar and loosely inside the jacket. The gasket may be made of rubber, leather, or other material that is elastic enough to cling to the test jar and hard enough to hold its shape. Its purpose is to prevent the test jar from touching the jacket.

6.7 Bath or Baths, maintained at prescribed temperatures with a firm support to hold the jacket vertical. The required bath temperatures may be obtained by refrigeration if available, otherwise by suitable cooling mixtures. Cooling mixtures commonly used for bath temperatures down to those shown are in Table 1.

7. Reagents and Materials

7.1 The following solvents of technical grade are appropriate for low-temperature bath media.

7.1.1 Acetone, (**Warning**—Extremely flammable).

7.1.2 Alcohol, Ethanol (**Warning**—Flammable).

7.1.3 Alcohol, Methanol (**Warning**—Flammable. Vapor harmful).

7.1.4 Petroleum Naphtha, (**Warning**—Combustible. Vapor harmful).

7.1.5 Solid Carbon Dioxide, (**Warning**—Extremely cold –78.5 °C).

8. Procedure

8.1 Pour the specimen into the test jar to the level mark. When necessary, heat the specimen in a bath until it is just sufficiently fluid to pour into the test jar.

NOTE 4—It is known that some materials, when heated to a temperature higher than 45 °C during the preceding 24 h, do not yield the same pour point results as when they are kept at room temperature for 24 h prior to testing. Examples of materials which are known to show sensitivity to thermal history are residual fuels, black oils, and cylinder stocks.

⁶ Supporting data have been filed at ASTM International Headquarters and may be obtained by requesting Research Report RR:D02-1826. Contact ASTM Customer Service at service@astm.org.

TABLE 1 Cooling Mixtures and Bath Temperatures

Cooling Mixture	Bath Temperature
Ice and water	0 °C ± 1.5 °C
Crushed ice and sodium chloride crystals or Acetone or petroleum naphtha, or methanol or ethanol (see Section 7) with solid carbon dioxide added to give the desired temperature	–18 °C ± 1.5 °C
Acetone or petroleum naphtha or methanol or ethanol (see Section 7) with solid carbon dioxide added to give the desired temperature	–33 °C ± 1.5 °C
Acetone or petroleum naphtha or methanol or ethanol (see Section 7) with solid carbon dioxide added to give the desired temperature	–51 °C ± 1.5 °C
Acetone or petroleum naphtha or methanol or ethanol (see Section 7) with solid carbon dioxide added to give the desired temperature	–69 °C ± 1.5 °C

8.1.1 Samples of residual fuels, black oils, and cylinder stocks which have been heated to a temperature higher than 45 °C during the preceding 24 h, or when the thermal history of these sample types is not known, shall be kept at room temperature for 24 h before testing. Samples which are known by the operator not to be sensitive to thermal history need not be kept at room temperature for 24 h before testing.

8.1.2 Experimental evidence supporting elimination of the 24 h waiting period for some sample types is contained in a research report.⁷

8.2 In the case of pour points above 36 °C, use a higher range temperature measuring device (6.2) such as IP 63C or ASTM 61C, or a digital contact thermometer. Close the test jar with the cork carrying the test temperature measuring device (6.2). Adjust the position of the cork and temperature measuring device so the cork fits tightly, the temperature measuring device and the jar are coaxial, and the temperature measuring device is immersed to the correct depth.

8.2.1 For liquid-in-glass, the thermometer bulb should be immersed so the beginning of the capillary is 3 mm below the surface of the specimen.

8.2.2 For digital contact thermometers, the probe should be immersed so the end of the probe is 10 mm to 15 mm below the surface of the specimen.

8.3 For the measurement of pour point, subject the specimen in the test jar to the following preliminary treatment:

8.3.1 *Specimens Having Pour Points Above –33 °C*—Heat the specimen without stirring to 9 °C above the expected pour point, but to at least 45 °C, in a bath maintained at 12 °C above the expected pour point, but at least 48 °C. Transfer the test jar to a bath maintained at 24 °C ± 1.5 °C and commence observations for pour point. When using a liquid bath, ensure that the liquid level is between the fill mark on the test jar and the top of the test jar.

8.3.2 *Specimens Having Pour Points of –33 °C and Below*—Heat the specimen without stirring to at least 45 °C in a bath maintained at 48 °C ± 1.5 °C. Transfer the test jar to a bath maintained at 24 °C ± 1.5 °C. When using a liquid bath, ensure that the liquid level is between the fill mark on the test jar and the top of the test jar. When the specimen temperature reaches 27 °C, and if using liquid-in-glass thermometers, remove the high cloud and pour thermometer, and place the low cloud and pour thermometer in position. Transfer the test jar to the cooling bath (see 8.6.1).

8.4 See that the disk, gasket, and the inside of the jacket are clean and dry. Place the disk in the bottom of the jacket. Place the gasket around the test jar, 25 mm from the bottom. Insert the test jar in the jacket. Never place a jar directly into the cooling medium.

8.5 After the specimen has cooled to allow the formation of paraffin wax crystals, take great care not to disturb the mass of specimen nor permit the thermometer to shift in the specimen; any disturbance of the spongy network of wax crystals will lead to low and erroneous results.

8.6 Pour points are expressed in integers that are positive or negative multiples of 3 °C. Begin to examine the appearance of the specimen when the temperature of the specimen is 9 °C above the expected pour point (estimated as a multiple of 3 °C). At each test temperature that is a multiple of 3 °C below the starting temperature remove the test jar from the jacket. To remove condensed moisture that limits visibility wipe the surface with a clean cloth moistened in alcohol (ethanol or methanol). Tilt the jar just enough to ascertain whether there is a movement of the specimen in the test jar. If movement of specimen in the test jar is noted, then replace the test jar immediately in the jacket and repeat a test for flow at the next temperature, 3 °C lower. Typically, the complete operation of removal, wiping, and replacement shall require not more than 3 s.

8.6.1 If the specimen has not ceased to flow when its temperature has reached 27 °C, transfer the test jar to a jacket in a cooling bath maintained at 0 °C ± 1.5 °C. As the specimen

⁷ Supporting data have been filed at ASTM International Headquarters and may be obtained by requesting Research Report RR:D02-1377. Contact ASTM Customer Service at service@astm.org.

continues to get colder, transfer the test jar to a jacket in the next lower temperature cooling bath in accordance with [Table 2](#).

8.6.2 If the specimen in the jar does not show movement when tilted, hold the jar in a horizontal position for 5 s, as noted by an accurate timing device, and observe the specimen carefully. If the specimen shows any signs of movement before 5 s has passed, replace the test jar immediately in the jacket and repeat a test for flow at the next temperature, 3 °C lower.

8.7 Continue in this manner until a point is reached at which the specimen shows no movement when the test jar is held in a horizontal position for 5 s. Record the observed reading of the test thermometer.

8.8 For black specimen, cylinder stock, and residual fuel specimen, the result obtained by the procedure described in [8.1](#) through [8.7](#) is the upper (maximum) pour point. If required, determine the lower (minimum) pour point by heating the sample while stirring, to 105 °C, pouring it into the jar, and determining the pour point as described in [8.4](#) through [8.7](#).

8.9 Some specifications allow for a pass/fail test or have pour point limits at temperatures not divisible by 3 °C. In these cases, it is acceptable practice to conduct the pour point measurement according to the following schedule: Begin to examine the appearance of the specimen when the temperature of the specimen is 9 °C above the specification pour point. Continue observations at 3 °C intervals as described in [8.6](#) and [8.7](#) until the specification temperature is reached. Report the sample as passing or failing the specification limit.

9. Calculation and Report

9.1 Add 3 °C to the temperature recorded in [8.7](#) and report the result as the Pour Point, ASTM D97. For black oil, and so forth, add 3 °C to the temperature recorded in [8.7](#) and report the result as Upper Pour Point, ASTM D97, or Lower Pour Point, ASTM D97, as required.

10. Precision and Bias

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

NOTE 5—The precision statements were developed using liquid-in-glass thermometers corresponding to those in Specification [E1](#) or IP Specifications for IP Standard Thermometers.

TABLE 2 Bath and Sample Temperature Ranges

Bath Temperature Setting, °C	Sample Temperature Range, °C
48 ± 1.5 or 12 above expected pour point	Preheat to at least 45 or 9 above expected pour point
24 ± 1.5	Start to 27
0 ± 1.5	27 to 9
−18 ± 1.5	9 to −6
−33 ± 1.5	−6 to −24
−51 ± 1.5	−24 to −42
−69 ± 1.5	−42 to −60

10.1.1 *Lubricating Oil*:⁸

10.1.1.1 *Repeatability*—The difference between successive test results, obtained by the same operator using the same apparatus under constant operating conditions on identical test material would, in the long run, in the normal and correct operation of this test method, exceed 6 °C only in one case in twenty. Differences greater than this should be considered suspect.

10.1.1.2 *Reproducibility*—The difference between two single and independent test 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 this test method, exceed 9 °C only in one case in twenty. Differences greater than this should be considered suspect.

10.1.1.3 The precision statements⁸ were derived from a 1998 interlaboratory test program using Practice [D6300](#). Participants analyzed five sets of duplicate base oils, three sets of duplicate multigrade lubricating oils, and one set each of duplicate hydraulic oils and automatic transmission fluid in the temperature range of −51 °C to −11 °C. Seven laboratories participated with the manual Test Method D97. Information on the type of samples and their average pour points are in Research Report RR:D02-1499.⁸

NOTE 6—The precision statements are the derived values rounded up to the next testing interval value. The actual derived precision values appear in [Table X1.1](#).

10.1.2 *Middle Distillate and Residual Fuel*:⁹

10.1.2.1 *Repeatability*—The difference between successive test results obtained by the same operator using the same apparatus under constant operation conditions on identical test material would, in the long run, in the normal and correct operation of this test method, exceed 3 °C only in one case in twenty. Differences greater than this should be considered suspect.

10.1.2.2 *Reproducibility*—The difference between two single and independent test 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 this test method, exceed 9 °C only in one case in twenty. Differences greater than this should be considered suspect.

10.1.2.3 The precision statements⁹ were prepared with data on sixteen middle distillate and residual fuels tested by twelve cooperators. The fuels had pour points ranging from −33 °C to +51 °C.

NOTE 7—The precision statements are the derived values rounded up to the next testing interval value. The actual derived precision values can be seen in [Table X1.1](#).

NOTE 8—The precisions in [10.1.2](#) are not known to have been derived using Practice [D6300](#).

10.2 *Bias*—There being no criteria for measuring bias in these test-product combinations, no statement of bias can be made.

⁸ Supporting data (the results of the 1998 interlaboratory cooperative test program) have been filed at ASTM International Headquarters and may be obtained by requesting Research Report RR:D02-1499. Contact ASTM Customer Service at service@astm.org.

⁹ Based on the results of the 1983 interlaboratory cooperative test program.

11. Keywords

11.1 petroleum products; pour point

APPENDIXES

(Nonmandatory Information)

X1. ACTUAL DERIVED PRECISION VALUES

X1.1 See [Table X1.1](#).

TABLE X1.1 Actual Derived Precision Values

95 % Confidence	1998 Research Program Lubricating Oil, °C	1983 Research Program Middle Distillate and Residual Fuels, °C
Repeatability	5.3	2.5
Reproducibility	8.0	6.6

X2. THERMOMETER SPECIFICATIONS

X2.1 See [Table X2.1](#).

TABLE X2.1 Thermometer Specifications

		Low cloud and pour	High cloud and pour	Melting point
Range	°C	–80 to +20	–38 to +50	32 to 127
Immersion	mm	76	108	79
Graduation at each	°C	1	1	0.2
Longer lines at each	°C	5	5	1
Figured at each	°C	10	10	2
Scale error, max	°C	1 down to –33 2 below –33	0.5	0.2
Expansion chamber to permit heating to	°C	60	100	150
Overall length	mm	230 ± 5	230 ± 5	380 ± 5
Stem diameter	mm	6 to 8	6 to 8	6 to 8
Bulb length	mm	7.0 to 10	7.0 to 10	18 to 28
Bulb diameter	mm	5.0 to stem	5.5 to stem	5.0 to 6.0
Distance from bottom of bulb to line at	°C	–70	–38	32
	mm	100 to 120	120 to 130	105 to 115
Length of scale/range	mm	70 to 100	65 to 85	200 to 240

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