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An American National Standard



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Standard Test Methods for Aniline Point and Mixed Aniline Point of Petroleum Products and Hydrocarbon Solvents¹

This standard is issued under the fixed designation D 611; 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.

These test methods were adopted as a joint ASTM-IP standard in 1964.

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

1. Scope

1.1 These test methods cover the determination of the aniline point of petroleum products and hydrocarbon solvents. Method A is suitable for transparent samples with an initial boiling point above room temperature and where the aniline point is below the bubble point and above the solidification point of the aniline-sample mixture. Method B, a thin-film method, is suitable for samples too dark for testing by Method A. Methods C and D are for samples that may vaporize appreciably at the aniline point. Method D is particularly suitable where only small quantities of sample are available. Method E describes a procedure using an automatic apparatus suitable for the range covered by Methods A and B.

1.2 These test methods also cover the determination of the mixed aniline point of petroleum products and hydrocarbon solvents having aniline points below the temperature at which aniline will crystallize from the aniline-sample mixture.

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.* Specific precautionary statements are given in Sections 7.1 and 7.3.

2. Referenced Documents

2.1 ASTM Standards:

D 1015 Test Method for Freezing Points of High-Purity Hydrocarbons²

D 1217 Test Method for Density and Relative Density (Specific Gravity) of Liquids by Bingham Pycnometer²

D 1218 Test Method for Refractive Index and Refractive Dispersion of Hydrocarbon Liquids²

¹ These test methods are under the jurisdiction of ASTM Committee D-2 on Petroleum Products and Lubricants and are the direct responsibility of D02.04 on Hydrocarbon Analysis.

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² *Annual Book of ASTM Standards*, Vol 05.01.

D 1500 Test Method for ASTM Color of Petroleum Products (ASTM Color Scale)²

D 2700 Test Method for Knock Characteristics of Motor and Aviation Fuels by the Motor Method³

E 1 Specification for ASTM Thermometers⁴

3. Terminology

3.1 Definitions:

3.1.1 *aniline point*—the minimum equilibrium solution temperature for equal volumes of aniline and sample.

3.1.2 *mixed aniline point*—the minimum equilibrium solution temperature of a mixture of two volumes of aniline, one volume of sample, and one volume of *n*-heptane of specified purity.

4. Summary of Test Methods

4.1 Specified volumes of aniline and sample, or aniline and sample plus *n*-heptane, are placed in a tube and mixed mechanically. The mixture is heated at a controlled rate until the two phases become miscible. The mixture is then cooled at a controlled rate and the temperature at which two phases separate is recorded as the aniline point or mixed aniline point.

5. Significance and Use

5.1 The aniline point (or mixed aniline point) is useful as an aid in the characterization of pure hydrocarbons and in the analysis of hydrocarbon mixtures. Aromatic hydrocarbons exhibit the lowest, and paraffins the highest values. Cycloparaffins and olefins exhibit values that lie between those for paraffins and aromatics. In homologous series the aniline points increase with increasing molecular weight. Although it occasionally is used in combination with other physical properties in correlative methods for hydrocarbon analysis, the aniline point is most often used to provide an estimate of the aromatic hydrocarbon content of mixtures.

³ *Annual Book of ASTM Standards*, Vol 05.04.

⁴ *Annual Book of ASTM Standards*, Vol 14.03.

6. Apparatus

6.1 For details of the aniline point apparatus required for each method see:

Annex A1 for Method A
Annex A2 for Method B
Annex A3 for Method C
Annex A4 for Method D
Annex A5 for Method E

NOTE 1—Alternative apparatus may be used, such as the U-tube method for dark oils, provided it has been shown to give results of the same precision and accuracy as those described in the Annexes.

6.2 *Heating and Cooling Bath*—A suitable air bath, a nonvolatile, transparent liquid bath, or an infrared lamp (250 to 375 W), provided with means for controlling the rate of heating.

NOTE 2—Water should not be used as either a heating or cooling medium since aniline is hygroscopic and moist aniline will give erroneous test results. For example, the aniline point of the *n*-heptane reagent as measured with aniline containing 0.1 volume % water is approximately 0.5°C (0.9°F) higher than that measured with dry aniline. If the aniline point is below the dew point of the atmosphere, pass a slow stream of dry inert gas into the aniline point tube to blanket the aniline-sample mixture.

6.3 *Thermometers*, having the following ranges and conforming to the requirements of the designated ASTM or IP specification:

Range	ASTM (Specification E 1)	IP
-38 to +42°C (-36.5 to +107.5°F)	33C, 33F	20C
25 to 105°C (77 to 221°F)	34C, 34F	21C
90 to 170°C (194 to 338°F)	35C, 35F	59C

6.4 *Pipets*, with capacities of 10 ± 0.04 mL, 5 ± 0.02 mL, the latter equipped with a long, fine tip. Provide a rubber suction bulb for use with pipets when measuring aniline.

6.5 *Balance*—A laboratory balance sensitive to 0.01 g, suitable for weighing the tube and sample when the sample cannot be pipetted conveniently.

6.6 *Safety Goggles*.

6.7 *Plastic Gloves*, impervious to aniline.

7. Reagents

7.1 *Aniline* (**Warning**—See Note 3.) Dry chemically pure aniline over potassium hydroxide pellets, decant, and distill fresh on the day of use, discarding the first and last 10 %. Aniline thus prepared when tested with *n*-heptane according to Section 9 shall give an aniline point of $69.3 \pm 0.2^\circ\text{C}$ ($156.7 \pm 0.4^\circ\text{F}$) as determined from the average of two independent tests having a difference of not more than 0.1°C (0.2°F).

NOTE 3—**Warning**: Aniline should not be pipetted directly by mouth because of its extreme toxicity. Aniline is also toxic by absorption through the skin even in *very small quantities*, and should be handled with great caution.

NOTE 4—For routine purposes the distillation process is not mandatory provided the aniline meets the requirements of the test with *n*-heptane.

NOTE 5—The aniline point of aniline and *n*-heptane determined with automatic apparatus (Method E) shall be $69.3 \pm 0.2^\circ\text{C}$ ($156.7^\circ\text{F} \pm 0.4^\circ\text{F}$) when corrected in accordance with the equation in Section A5.2.1

NOTE 6—As an alternative to distilling the aniline on the day of use, the aniline may be distilled as described in 7.1, collecting the distillate in ampoules, sealing the ampoules under vacuum or dry nitrogen, and storing in a cool dark place for future use. In either case, rigid precaution must be

taken to avoid contamination from atmospheric moisture (Note 2). It is believed that under these conditions the aniline will remain unchanged for a period exceeding 6 months.

7.2 *Calcium Sulfate*, anhydrous.

7.3 *n-Heptane* (**Warning**—See Note 7), conforming to the requirements listed in Table 1.⁵

NOTE 7—**Warning**: Flammable. Harmful if inhaled. See Annex A6.1.

8. Sample

8.1 Dry the sample by shaking vigorously for 3 to 5 min with about 10 volume % of a suitable drying agent such as anhydrous calcium sulfate or anhydrous sodium sulfate. Reduce the viscosity of viscous samples by warming to a temperature below that which would cause the loss of light ends or the dehydration of the drying agent. Remove any suspended drying agent by use of a centrifuge or by filtration. Heat samples containing separated wax until they are homogeneous and keep heated during filtration or centrifugation to ensure against separation of wax. When suspended water is visibly present and the sample material is known to dissolve less than 0.03 mass % of water, the use of a centrifuge for the removal of suspended water is an acceptable procedure.

9. Procedure for Aniline Point

9.1 The following methods, to be used as applicable, are covered as follows:

9.1.1 *Method A*, described in detail in Annex A1, is applicable to clear samples or to samples not darker than No. 6.5 ASTM color, as determined by Test Method D 1500, having initial boiling points well above the expected aniline point.

9.1.2 *Method B*, described in detail in Annex A2, is applicable to light-colored samples, moderately dark samples, and to very dark samples. It is suitable for samples that are too dark to be tested by Method A.

9.1.3 *Method C*, described in detail in Annex A3, is applicable to clear samples or to samples not darker than No. 6.5 ASTM color, as determined by Test Method D 1500, having initial boiling points sufficiently low as to give incorrect aniline point readings by Method A, for example, aviation gasoline.

9.1.4 *Method D*, described in detail in Annex A4, is applicable to the same type of sample as Method C. It is particularly useful when only limited quantities of sample are available.

⁵ These requirements for *n*-heptane are identical, except for tetraethyl lead, with those prescribed in the 1987 Annual Book of ASTM Standards, Vol 05.04.

TABLE 1 Requirements for *n*-Heptane

		ASTM Method
ASTM Motor Octane Number	0.0 ± 0.2	D 2700
Density at 20°C, g/mL	0.68380 ± 0.00015	D 1217
Refractive index, $n_D^{20^\circ\text{C}}$	1.38770 ± 0.00015	D 1218
Freezing point, °C	-90.710 min	D 1015
Distillation, 50 % recovered at 1.013 bar (760 mm Hg), °C	98.427 ± 0.025	A
Differential, 80 % recovered minus 20 % recovered, °C	0.020 max	

^AFor equipment and method used, see *Journal of Research*, National Institute of Standards and Technology, Vol 44, No. 3, 1950, pp. 309 and 310 (RP2079).

9.1.5 *Method E* is applicable when using automatic apparatus in accordance with the instructions in Annex A5.

10. Procedure for Mixed Aniline Point

10.1 This procedure is applicable to samples having aniline points below the temperature at which aniline crystallizes from the mixture. Pipet 10 mL of aniline (**Warning:** See Note 3), 5 mL of sample, and 5 mL of *n*-heptane into a clean, dry apparatus. Determine the aniline point of the mixture by Method A or B as described in Annex A1 or Annex A2.

11. Report

11.1 If the range of three successive observations of the aniline point temperature is not greater than 0.1°C (0.2°F) for light-colored samples or 0.2°C (0.4°F) for dark samples, report the average temperature of these observations, corrected for thermometer calibration errors, to the nearest 0.05°C (0.1°F) as the aniline point.

11.2 If such a range is not obtained after five observations, repeat the test using fresh quantities of aniline and sample in a clean, dry apparatus, and if consecutive temperature observations show a progressive change, or if the range of observations is greater than the repeatability given in 12.1, report the method as being inapplicable.

12. Precision and Bias

12.1 The precision of these test methods as obtained by statistical examination of interlaboratory test results is as follows:

12.1.1 *Repeatability*—The difference between successive test results (two average temperatures obtained in a series of observations as described in Section 11 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:

Repeatability		
Aniline point of:		
Clear, light-colored samples	0.16°C	(0.3°F)
Moderately dark to very dark samples	0.3°C	(0.6°F) ^A
Mixed aniline point of:		
Clear, light-colored samples	0.16°C	(0.3°F) ^A
Moderately dark to very dark samples	0.3°C	(0.6°F) ^A

^ANot determined from recent cooperative tests; however, the ratios with those given in the 1953 version are believed to apply.

12.1.2 *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:

12.2 *Bias*—A statement of bias is now being developed by the subcommittee.

Reproducibility	
Aniline point of:	
Clear, light-colored samples	0.5°C (0.9°F)
Moderately dark to very dark samples	1.0°C (1.8°F) ^A
Mixed aniline point of:	
Clear, light-colored samples	0.7°C (1.3°F) ^A
Moderately dark to very dark samples	1.0°C (1.8°F) ^A

^ANot determined from recent cooperative tests; however, the ratios with those given in the 1953 version are believed to apply.

12.3 The precision of this test was not obtained in accordance with Committee D-2 Research Report RR:D02-1007, "Manual on Determining Precision Data for ASTM Methods on Petroleum Products and Lubricants."

13. Keywords

13.1 aniline point; aromatics; mixed aniline point

ANNEXES

(Mandatory Information)

A1. METHOD A

A1.1 Apparatus

A1.1.1 The apparatus shown in Fig. A1.1 shall consist of the following:

A1.1.1.1 *Test Tube*, approximately 25 mm in diameter and 150 mm in length, made of heat-resistant glass.

A1.1.1.2 *Jacket*, approximately 37 to 42 mm in diameter and 175 mm in length, made of heat-resistant glass.

A1.1.1.3 *Stirrer*, manually operated, metal, approximately 2 mm in diameter (14 B&S gage) metal wire as shown in Fig. A1.1. A concentric ring shall be at the bottom, having a diameter of approximately 19 mm. The length of the stirrer to a right-angle bend shall be approximately 200 mm. The right-angle bend shall be approximately 55 mm long. A glass sleeve approximately 65 mm in length of 3-mm inside diameter shall be used as a guide for the stirrer. Any suitable mechanical device for operating the stirrer as specified is an approved

alternative for the manual operation.

A1.2 Procedure

A1.2.1 Clean and dry the apparatus. Pipet 10 mL of aniline (**Warning**— see 7.1) and 10 mL of the dried sample (8.1) into the test tube fitted with stirrer and thermometer. If the material is too viscous for pipetting, weigh to the nearest 0.01 g a quantity of the sample corresponding to 10 mL at room temperature. Center the thermometer in the test tube so that the immersion mark is at the liquid level, making sure that the thermometer bulb does not touch the side of the tube. Center the test tube in the jacket tube. Stir the mixture rapidly using a 50-mm (2-in.) stroke, avoiding the introduction of air bubbles.

A1.2.2 If the aniline-sample mixture is not miscible at room temperature, apply heat directly to the jacket tube so that the temperature rises at a rate of 1 to 3°C (2 to 5°F)/min by