Designation: D1657 - 12 (Reapproved 2017)

# AMERICAN PETROLEUM INSTITUTE

# Manual of Petroleum Measurement Standards (MPMS), Chapter 9.2

# Standard Test Method for Density or Relative Density of Light Hydrocarbons by Pressure Hydrometer<sup>1</sup>

This standard is issued under the fixed designation D1657; 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  $(\varepsilon)$  indicates an editorial change since the last revision or reapproval.

## 1. Scope

- 1.1 This test method covers the determination of the density or relative density of light hydrocarbons including liquefied petroleum gases (LPG) having Reid vapor pressures exceeding 101.325 kPa (14.696 psi).
- 1.2 The prescribed apparatus should not be used for materials having vapor pressures higher than 1.4 MPa (200 psi) at the test temperature. This pressure limit is dictated by the type of equipment. Higher pressures can apply to other equipment designs.
- 1.3 The initial pressure hydrometer readings obtained are uncorrected hydrometer readings and not density measurements. Readings are measured on a hydrometer at either the reference temperature or at another convenient temperature, and readings are corrected for the meniscus effect, the thermal glass expansion effect, alternate calibration temperature effects and to the reference temperature by means of calculations and Adjunct to D1250 Guide for Petroleum Measurement Tables (API MPMS Chapter 11.1) or API MPMS Chapter 11.2.4 (GPA TP-27), as applicable.
- 1.4 Values determined as density or relative density can be converted to equivalent values in the other units or alternative reference temperatures by means of Interconversion Procedures API *MPMS* Chapter 11.5, or Adjunct to D1250 Guide for Petroleum Measurement Tables (API *MPMS* Chapter 11.1) or API *MPMS* Chapter 11.2.4 (GPA TP-27), as applicable.
- 1.5 The calculations required in Section 11 shall be applied to the initial pressure hydrometer reading with observations

Current edition approved July 15, 2017. Published July 2017. Originally approved in 1939. Last previous edition approved in 2012 as D1657 – 12. DOI: 10.1520/D1657-12R17.

and results reported as required by Section 11 prior to use in a subsequent calculation procedure (measurement ticket calculation, meter factor calculation, or base prover volume determination).

- 1.6 Annex A1 contains a procedure for verifying or certifying the equipment for this test method.
- 1.7 The values in SI units are to be regarded as the standard. US Customary values shown in adjacent parentheses are for information only and may not be exactly equivalent. Both SI and customary units have been rounded so that they may not be exactly equivalent.
- 1.8 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.
- 1.9 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:<sup>2</sup>
- D1250 Guide for Use of the Petroleum Measurement Tables
  D1265 Practice for Sampling Liquefied Petroleum (LP)
  Gases, Manual Method
- D1298 Test Method for Density, Relative Density, and API Gravity of Crude Petroleum and Liquid Petroleum Products by Hydrometer Method (API *MPMS* Chapter 9.1)
- E1 Specification for ASTM Liquid-in-Glass Thermometers

<sup>&</sup>lt;sup>1</sup> This test method is under the jurisdiction of ASTM Committee D02 on Petroleum Products, Liquid Fuels, and Lubricants and the API Committee on Petroleum Measurement, and is the direct responsibility of Subcommittee D02.02 /COMQ, the joint ASTM-API Committee on Hydrocarbon Measurement for Custody Transfer (Joint ASTM-API). This test method has been approved by the sponsoring committees and accepted by the Cooperating Societies in accordance with established procedures.

<sup>&</sup>lt;sup>2</sup> 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

E100 Specification for ASTM Hydrometers 2.2 *API Standards*:<sup>3</sup>

MPMS Chapter 9.1 Test Method for Density, Relative Density, and API Gravity of Crude Petroleum and Liquid Petroleum Products by Hydrometer Method (ASTM Test Method D1298)

MPMS Chapter 11.1 Temperature and Pressure Volume Correction Factors for Generalized Crude Oils, Refined Products and Lubricating Oils (Adjunct to ASTM D1250)

MPMS Chapter 11.2.2 Compressibility Factors for Hydrocarbons: 0.350-0.0637 Relative Density (60°F/60°F) and -50°F to 140°F Metering Temperature

MPMS Chapter 11.2.2M Compressibility Factors for Hydrocarbons: 350-637 Kilograms per Cubic Meter Density 15°C and -46°C to 60°C Metering Temperatures

MPMSChapter 11.2.4 Temperature Correction for the Volume of NGL and LPG Tables 23E, 24E, 53E, 54E, 59E and 60E (joint standard with GPA TP-27)

MPMS Chapter 11.5 Density/Weight/Volume Intraconversion

2.3 GPA Standards:<sup>4</sup>

GPA Technical Publication TP-27 Temperature Correction for the Volume of NGL and LPG, Tables 23E, 24E, 53E, 54E, 59E and 60E (joint standard with API *MPMS* Chapter 11.2.4)

2.4 ASTM Adjuncts:

Adjunct to D1250 Guide for Petroleum Measurement Tables (API MPMS Chapter 11.1)<sup>5</sup>

## 3. Terminology

- 3.1 Definitions:
- 3.1.1 *density, n*—the mass of liquid per unit volume at 15 °C and its saturation pressure with the standard unit of measurement being kilograms per cubic metre.
- 3.1.1.1 *Discussion*—Other reference temperatures, such as 20 °C may be used for some products or in some locations. Less preferred units of measurement; for example, kg/L or g/mL, are still in use.
- 3.1.2 relative density (specific gravity), n—the ratio of the mass of a given volume of liquid at a specific temperature to the mass of an equal volume of pure water at the same or different temperature. Both reference temperatures shall be explicitly stated.
- 3.1.2.1 *Discussion*—Common reference temperatures include 60/60 °F, 20/20 °C, 20/4 °C. The historic deprecated term "specific gravity" may still be found.
- 3.1.3 *thermohydrometer*, *n*—a glass hydrometer with a self-contained mercury thermometer.

#### 4. Summary of Test Method

4.1 The apparatus is purged with a portion of the sample before filling with the portion to be used for testing. The

pressure cylinder is filled to a level at which the enclosed hydrometer floats freely, and the cylinder is then placed in a constant-temperature bath (if needed). When the temperature has reached equilibrium, the hydrometer reading and the temperature of the sample are read. The observed hydrometer reading is corrected for the meniscus effect, alternate calibration temperature effects and reduced to the reference temperature by means of the volume correction factors or tables as applicable by use of the appropriate Adjunct to D1250 Guide for Petroleum Measurement Tables (API MPMS Chapter 11.1) or API MPMS Chapter 11.2.4 (GPA TP-27), as applicable, and observed temperature from the thermometer.

## 5. Significance and Use

5.1 The density or relative density of light hydrocarbons and liquefied petroleum gases is used in custody transfer quantity calculations or to satisfy transportation, storage, and regulatory requirements.

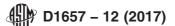
#### 6. Apparatus

- 6.1 *Hydrometers*, graduated in density with a range from 500 kg/m<sup>3</sup> to 650 kg/m<sup>3</sup>, or in relative density with a range from 0.500 to 0.650, and conforming to the dimensions in Specification E100.
- 6.1.1 *Thermohydrometers*, for field applications, thermohydrometers may be more convenient than hydrometers with separate thermometers. They shall conform to Specification E100, Thermohydrometer Nos. 101H or 310H.
- 6.1.1.1 Thermohydrometers shall be of suitable range and have dimensions to float freely within the pressure hydrometer cylinder with clearances of 5 mm at the wall and 25 mm at the top and bottom.
- 6.1.1.2 The test report shall state that a thermohydrometer was used.
- 6.1.1.3 The user should ascertain that the instruments used for this procedure conform to the requirements set out above with respect to materials, dimensions, and scale errors. In cases where the instrument is provided with a calibration certificate issued by a recognized standardizing body, the instrument is classed as certified and the appropriate corrections for the meniscus effect, the thermal glass expansion effect, and alternative calibration temperature effects shall be applied to the observed readings prior to corrections. Instruments that satisfy the requirements of this test method, but are not provided with a recognized calibration certificate, are classed as uncertified and the appropriate corrections for the meniscus effect, the thermal glass expansion effect, and alternative calibration temperature effects shall be applied to the observed readings prior to corrections.
- 6.2 Hydrometer Cylinder, constructed of transparent plastic; for example, poly(methyl methacrylate) or equivalent material, conforming to the design and recommended dimensions given in Fig. 1. The cylinder shall be of such dimensions that the hydrometer shall float freely within it. The ends shall be tightly sealed with neoprene gaskets and metal end plates as shown in Fig. 1. (Warning—A protective shield shall be placed around the cylinder. Replace any cylinders that show signs of fogging, crazing, cracking, or etching.)

<sup>&</sup>lt;sup>3</sup> Available from American Petroleum Institute (API), 1220 L. St., NW, Washington, DC 20005-4070, http://www.api.org.

<sup>&</sup>lt;sup>4</sup> Available from Gas Processors Association (GPA), 6526 E. 60th St., Tulsa, OK 74145, http://www.gpaglobal.org.

<sup>&</sup>lt;sup>5</sup> Available from ASTM International Headquarters. Order Adjunct No. ADJD1250. Original adjunct produced in 1983.



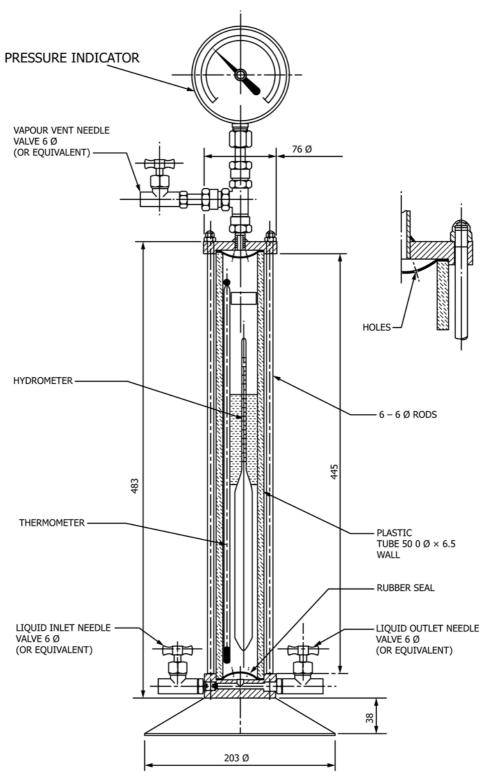


FIG. 1 Pressure Hydrometer Cylinder

Note 1—Certain compounds attack plastics and cloud the inner surface of the cylinder, making it difficult or impossible to read the hydrometer. Tests showed no attack by ethane, ethene (ethylene), propane, propylene, butane, methylpropane (isobutane), butenes (normal butylenes), methylpropene (isobutylene), pentane, and methylbutane (isopentane), and no attack is expected from butadiene and acetaldehyde. Users are cautioned,

however, to clean the cylinder thoroughly after each determination. Ketones and alcohols should not be used for cleaning as they attack and weaken plastics while aromatics also tend to attack the surface of plastics and should similarly not be used. A light aliphatic hydrocarbon is recommended for cleaning. (Warning—Extremely flammable. Harmful if inhaled. Vapors may cause flash fire.)

- 6.2.1 The liquid inlet valve and the liquid outlet valve shall be tightly connected to a base plate that shall be bored to give both valves a common inlet to the cylinder. The vapor vent valve shall be similarly connected to the top plate, which shall be bored to provide a vapor outlet from the pressure cylinder. All valves shall be 6 mm (1/4 in.) or equivalent needle valves.
- 6.2.2 The cylinder shall not be operated at a gage pressure greater than 1.4 MPa (200 psi). A hydrostatic test at 2.8 MPa (400 psi) shall be carried out at intervals no greater than 12 months.
- Note 2—For Safety—The cylinder should be equipped with appropriate safety devices to prevent over-pressure. Consideration should be given to pressure testing the cylinder at 1.25 to 1.5 times its normal maximum operating pressure on a periodic basis, not to exceed two years.
- 6.3 *Thermometers*, conforming to ASTM 12C or 12F in Specification E1. The thermometer shall be held firmly inside the hydrometer cylinder by a clip in such a manner that it does not interfere with the free movement of the hydrometer.
- 6.4 Constant-Temperature Bath, of dimensions such that it can accommodate the hydrometer cylinder with the test portion fully immersed below the test portion liquid surface, and a temperature control system capable of maintaining the bath temperature within 0.25 °C of the test temperature throughout the duration of the test.

## 7. Reference Liquids

- 7.1 The following reference liquids are required for standardization of the hydrometer:
- 7.1.1 *Propane*, pure grade, having a nominal density of 507.6 kg/m $^3$  at 15 °C or a relative density 60/60 °F of 0.50699. (**Warning**—Extremely flammable. Harmful if inhaled. Vapors may cause flash fire.) The density of the reference liquid used shall be known.
- 7.1.2 *Butane*, pure grade, having a nominal density of 584.1 kg/m<sup>3</sup> at 15 °C or a relative density 60/60 °F of 0.5840. (**Warning**—Extremely flammable. Harmful if inhaled. Vapors may cause flash fire.) The density of the reference liquid used shall be known.

### 8. Sampling

- 8.1 Unless otherwise specified, samples of liquid hydrocarbons shall be obtained by the procedures described in Practice D1265. The procedure for sampling for verification of the apparatus and for subsequent testing is described as follows.
- 8.1.1 Connect the source of supply of the liquid to be tested to the inlet valve by suitable fittings so that a representative sample can be introduced into the cylinder. Ascertain that these connections are free of leaks. Open the outlet and vent valves and purge the sampling connections by opening the inlet valve slightly, permitting the product to flow through the outlet valve at the bottom of the cylinder and the vent valve at the top of the cylinder.
- 8.1.2 When the connections have been purged, close the outlet and vent valves and open the inlet valve, permitting the liquid to enter the cylinder until it is full. If necessary, the vent valve may be opened slightly to permit complete filling of the cylinder after which it shall be closed. At no time shall the

- pressure in the cylinder be allowed to rise above a gage pressure of 1.4 MPa (200 psi).
- 8.1.3 When the cylinder has been filled, close the inlet valve and open the outlet and vent valves, permitting the contents of the cylinder to be withdrawn completely and the pressure inside the cylinder to be returned to atmospheric pressure.
- 8.1.4 Close the outlet and vent valves and open the inlet valve, filling the cylinder to a level at which the enclosed hydrometer floats freely. If it is necessary to accomplish this filling by venting vapor through the vent valve, repeat the purging to cool the cylinder sufficiently to permit its being filled without venting.
- 8.1.5 With all valves closed, examine the apparatus for leaks. If leaks are detected, discard the sample, reduce the pressure to atmospheric and repair the leaks. Repeat the sampling procedure.

## 9. Verification of Apparatus

- 9.1 Carefully clean and dry the hydrometer and the inside wall of the pressure cylinder.
- 9.2 Insert the hydrometer in the pressure cylinder and attach the thermometer and cover plate. Connect the source of supply of the reference liquid to the inlet valve and ascertain that the connections are free of leaks. Fill the hydrometer with one of the reference fluids (see Section 7) by the procedure given in 8.1.2 8.1.5.
- 9.3 Disconnect the hydrometer cylinder from the source of supply of liquid and place it in the constant-temperature bath maintained within 0.2 °C of the test temperature until thermal equilibrium has been obtained. To accelerate thermal adjustment, occasionally remove the hydrometer cylinder from the constant-temperature bath, tilt to a horizontal position, rock gently a few times to ensure mixing, and replace in the constant-temperature bath. Exercise care to prevent damage to the hydrometer and thermometer.
- 9.4 When the hydrometer cylinder and thermometer have reached equilibrium, as shown by the thermometer inside the cylinder, remove the cylinder from the constant-temperature bath, stand it on a firm level surface, and while the hydrometer is floating freely, take the hydrometer reading promptly as described in 9.4.1 and 9.4.2.
- 9.4.1 Observe a point slightly below the plane of the liquid surface and then raise the line of vision until this surface, seen as an ellipse, becomes a straight line. Record the point where this line cuts the hydrometer scale, estimating to the nearest one fifth of a scale division the observed hydrometer reading. A white card held behind the cylinder just below the liquid level will improve the visibility of the surface.
- 9.4.2 Read and record the temperature of the liquid to the nearest  $0.2\,^{\circ}\text{C}$ , immediately before and after reading the hydrometer. If the temperature differs by more than  $0.4\,^{\circ}\text{C}$ , replace the hydrometer cylinder in the bath to restore equilibrium and repeat the readings.
- 9.5 Repeat the verification with a second portion of the same reference fluid.
- 9.6 If the temperatures measured in 9.4.2 are within 0.2 °C of the reference temperature and the two determinations are

within 0.7 kg/m³ or 0.0007 relative density, average the two determinations. If the temperatures are not within 0.2 °C of the reference temperature, correct the observed hydrometer readings to the reference temperature by use of the appropriate table from Adjunct to D1250 Guide for Petroleum Measurement Tables (API *MPMS* Chapter 11.1) or API *MPMS* Chapter 11.2.4 (GPA TP-27). Repeat the verification with two additional determinations.

- 9.7 Compare the average results with the known density of the reference fluids used (see 9.2).
- 9.7.1 If the results are within  $0.2 \text{ kg/m}^3$  (0.0002 relative density), accept the hydrometer scale as suitable for direct reading.
- 9.7.2 If the difference is between  $0.2 \text{ kg/m}^3$  and  $1.0 \text{ kg/m}^3$  (0.0002 and 0.001 relative density) from the known value, apply this correction to readings obtained on test portions of samples.
- 9.7.3 If the readings differ more than 1.0 kg/m<sup>3</sup> (0.001 relative density) from the known value, replace the hydrometer.
- 9.8 After each determination, empty the cylinder and vent to reduce the pressure to atmospheric in a safe place. Highly volatile liquids and liquefied petroleum gases shall not be left in the apparatus because, at high ambient temperatures, they might generate sufficient pressure to burst the cylinder.

#### 10. Procedure

- 10.1 Purge and empty the apparatus and draw a sample of the product to be tested as in Section 8.
- 10.2 Carry out the procedures and take the reading as described in Section 9. After each determination, empty the cylinder and vent to reduce the pressure to atmospheric in a safe place.

Note 3—For measurements in the field, this method may be used at ambient temperatures, with the realization that the accuracy is decreased. (**Warning**—If so used, the cylinder must be vented and the test discarded if the pressure rises above a gauge pressure of 1.4 MPa (200 psi).)

Note 4—ASTM Test Method D1298 (API MPMS Chapter 9.1) can provide additional assistance in how to perform these tests

### 11. Calculation and Report

- 11.1 Apply hydrometer corrections, if any, to the observed readings and report the corrected reading to the nearest  $0.5 \text{ kg/m}^3$  or 0.0005 relative density.
- 11.2 If the determinations were not carried out within 0.2 °C of the reference temperature, correct the results by use of the appropriate Adjunct to D1250 Guide for Petroleum Measurement Tables (API *MPMS* Chapter 11.1) or API *MPMS* Chapter 11.2.4 (GPA TP-27).
- 11.2.1 When a density-scaled hydrometer was used, use Table 53E for densities up to  $635 \text{ kg/m}^3$  to obtain density at 15 °C.
- 11.2.2 When a relative density hydrometer was used, use Table 23E for relative densities up to 0.655 to obtain relative density at  $60 \,^{\circ}$ F.
- 11.3 Report the final result as density to the nearest  $0.5~kg/m^3$  at  $15~^{\circ}C$  or as relative density to the nearest 0.0005 and specify the reference temperature.

- 11.3.1 The test report shall contain at least the following information:
  - 11.3.1.1 A reference to this standard.
- 11.3.1.2 The type and complete identification of the product tested.
  - 11.3.1.3 The result of the test.
  - 11.3.1.4 The date of the test.
  - 11.3.1.5 Any deviations from the test method.
  - 11.3.1.6 State whether a thermohydrometer was used.
- 11.4 The reporting values have no precision or bias determination. It is up to the user to determine whether this test method provides results of sufficient accuracy for the intended purpose.
- 11.5 Certified hydrometers from a recognized standardizing body, such as NIST, report the output density as 'Density in Vacuo.'

### 12. Precision and Bias

- 12.1 *Precision*—The precision of this test method as determined by statistical examination of interlaboratory test results is as follows:
- 12.1.1 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 value only one case in twenty.

Repeatability = 
$$1 \text{ kg/m}^3$$

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 value only in one case in twenty.

Reproducibility = 
$$3 \text{ kg/m}^3$$

Note 5—The precision of this test method has only been determined in the units of density.

Note 6-No precision data are available for thermohydrometers.

- 12.2 *Bias*—Bias for this test method has not been determined. However, to determine that the bias is within acceptable limits, verification of the hydrometer and thermometer using standards traceable to international standards shall occur before the thermohydrometer or hydrometer and thermometer are placed into service. Periodic re-verification may be required.
- 12.3 The repeatability and reproducibility values provided are not based on any interlaboratory round robin results. They should be considered historical numbers, the source of which cannot be verified by either ASTM or API and have been in this standard prior to the current slate of Light Hydrocarbons including liquefied petroleum gases (LPG). These values do not apply to the current calculation procedures, and it is up to the user to determine whether this test method provides results of sufficient accuracy for the intended purpose.

#### 13. Keywords

13.1 density; hydrocarbons; hydrometer; petroleum products; pressure hydrometer; relative density

## **ANNEX**

## A1. APPARATUS

(Mandatory Information)

## **A1.1 Apparatus Verification and Certification**

A1.1.1 *Hydrometers*, shall either be certified or verified. Verification shall be either by comparison with a certified hydrometer (see 6.1.1) or by the use of a certified reference material (CRM) specific to the reference temperature used.

A1.1.1.1 The hydrometer scale shall be correctly located within the hydrometer stem by reference to the datum mark. If the scale has moved, reject the hydrometer.

A1.1.1.2 Hydrometers shall be certified or verified at intervals of no more than 24 months.

A1.1.2 *Thermometers*, shall be verified at intervals of no more than six months for conformance with specifications. Either comparison with a referenced temperature measurement system traceable to an international standard, or a determination of ice point, is suitable.

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 jointly copyrighted by ASTM International (ASTM), 100 Barr Harbor Drive, PO Box C700, West Conshohocken, PA 19428-2959, United States, and the American Petroleum Institute (API), 1220 L Street NW, Washington DC 20005, 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/