



Standard Test Method for Electrical Conductivity of Liquid Hydrocarbons by Precision Meter¹

This standard is issued under the fixed designation D 4308; 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 Department of Defense.

^{ε1} NOTE—Warning notes were placed in the text editorially in April 2001.

1. Scope

1.1 This test method applies to the determination of the “rest” electrical conductivity of aviation fuels and other similar low-conductivity hydrocarbon liquids in the range from 0.1 to 2000 pS/m (see 3.2). This test method can be used in the laboratory or in the field.

1.2 The values stated in SI units are to be regarded as the 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.* For specific hazard statements, see 7.1.1 and 7.2.

2. Referenced Documents

2.1 ASTM Standards:

D 150 Test Methods for A-C Loss Characteristics and Permittivity (Dielectric Constant) of Solid Electrical Insulating Materials²

D 2624 Test Methods for Electrical Conductivity of Aviation and Distillate Fuels³

D 4306 Practice for Aviation Fuel Sample Containers for Tests Affected by Trace Contamination⁴

E 1 Specification for ASTM Thermometers⁵

3. Terminology

3.1 *picosiemens per metre*—the unit of electrical conductivity is also called a conductivity unit (CU). A siemen is the SI definition of reciprocal ohm sometimes called mho.

$$1 \text{ pS/m} = 1 \times 10^{-12} \Omega^{-1} \text{ m}^{-1} = 1 \text{ cu} = 1 \text{ picomho/m} \quad (1)$$

¹ This test method is under the jurisdiction of ASTM Committee D02 on Petroleum Products and Lubricants and is the direct responsibility of Subcommittee D02.J0 on Aviation Fuels.

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² *Annual Book of ASTM Standards*, Vol 10.01.

³ *Annual Book of ASTM Standards*, Vol 05.01.

⁴ *Annual Book of ASTM Standards*, Vol 05.02.

⁵ *Annual Book of ASTM Standards*, Vol 14.03.

3.2 *rest conductivity*—the reciprocal of the resistance of uncharged fuel in the absence of ionic depletion or polarization. It is the electrical conductivity at the initial instant of current measurement after a d-c voltage is impressed between electrodes.

4. Summary of Test Method

4.1 A sample of liquid hydrocarbon is introduced into a clean conductivity cell which is connected in series to a battery voltage source and a sensitive dc ammeter. The conductivity, automatically calculated from the observed peak current reading dc voltage and cell constant using Ohm’s law, appears as a digital value in either a manual or automatic mode of meter operation.

5. Significance and Use

5.1 The generation and dissipation of electrostatic charge in fuel due to handling depend largely on the ionic species present which may be characterized by the rest or equilibrium electrical conductivity. The time for static charge to dissipate is inversely related to conductivity. This test method can supplement Test Method D 2624 which is limited to fuels containing static dissipator additive.

NOTE 1—For low-conductivity fluids below 1 pS/m in conductivity, an a-c measurement technique is preferable to a d-c test method for sensing the electrical conductivity of bulk fluid. This d-c test method can be used at conductivities from 0.1 to 1 pS/m if precautions are observed in cell cleaning and sample handling. A waiting period of 15 min is required after filling the cell before measuring d-c conductivities below 1 pS/m. A single-laboratory program was conducted comparing this test method with a-c Test Method D 150. The results are on file at ASTM Headquarters. Request RR: D02-1241.

6. Apparatus

6.1 *Conductivity Apparatus*—Components of the dc conductivity apparatus are shown in Fig. 1.⁶

⁶ The KSLA Cell and Precision Conductivity Meter System, Emcee Model #1154 are available from Emcee Electronics, Inc., 520 Cypress Ave., Venice, FL 34292.

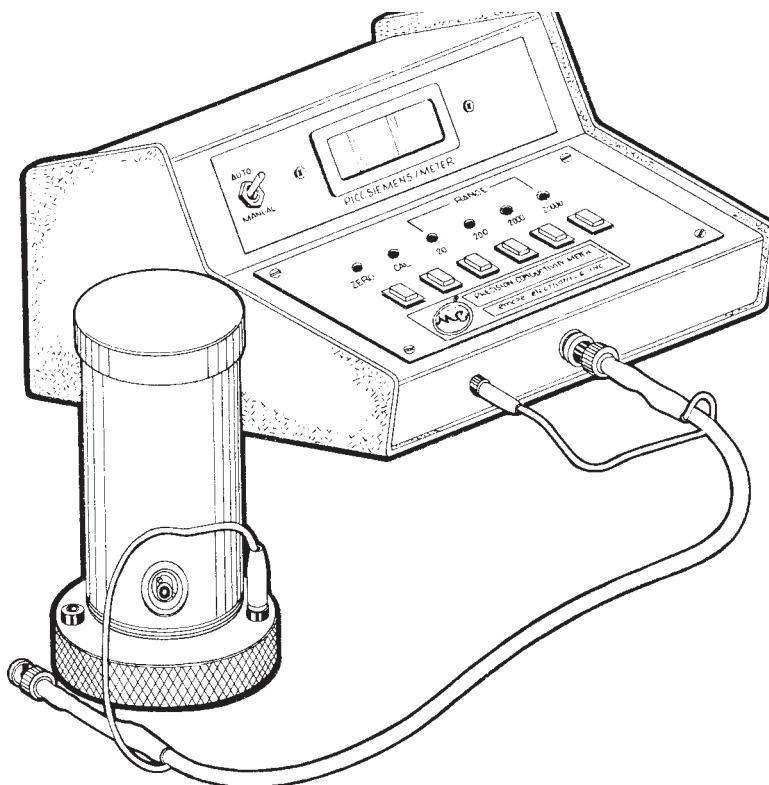


FIG. 1 Precision Conductivity Meter

6.1.1 The conductivity cell shown in Fig. 1 consists of an inner electrode and an outer electrode separated by an insulator. The outer electrode and cap provide a ground path and electrostatic (Faraday) shield.

6.1.2 The electrometer shown in Fig. 1 contains a battery which supplies a voltage to the cell and a bridge circuit which senses the flow of current and converts the output signal directly into conductivity units, that is, pS/m. A pushbutton selector allows selection of zero reading, calibration, and four range selections.

6.1.3 The cell and electrometer are connected by a triaxial cable as shown in Fig. 1.

6.2 *Thermometer*, calibrated to 0.5°C and conforming to Specification E 1.

7. Reagents

7.1 *Cleaning Solvent*—The following may be used:

7.1.1 *Toluene-Isopropyl Alcohol Mixture*—(**Warning**—Flammable. Vapor harmful. See Annex A1.1.) Mix two volumes of toluene and three volumes of isopropyl alcohol both of reagent grade and distill. Discard the first 20 % and last 5 % fractions.

7.2 *n-Heptane*—(**Warning**—Flammable. Harmful if inhaled. See Annex A1.2.) Prepare by percolating ASTM reference fuel grade *n*-heptane through silica gel⁷ as follows:

7.2.1 Activate approximately 2000 g of 100 to 200 mesh silica gel by heating at 180°C for 24 h. Allow it to cool in a

desiccator under nitrogen or in vacuum. Soak approximately 0.5 g of glass wool⁸ for 24 h in clean *n*-heptane.

7.2.2 Take a tube of borosilicate glass having an inside diameter of 60 to 70 mm, a length 1500 mm, with a conically shaped lower end provided with a glass cock. Place a perforated porcelain disk (diameter 25 mm) in the lower end of the tube and put the soaked glass wool on top of the disk. Fill the tube with the activated silica gel while tapping to achieve homogeneous filling. The silica gel layer will be approximately 1250 mm high. Wrap the column in black paper to exclude light.

7.2.3 Percolate *n*-heptane through the column at a rate of about 2 to 3 L/h. Discard the first 3 L. Never allow the column to run dry. The silica gel charge is sufficient for the percolation of 1000 L of *n*-heptane, provided the conductivity of the untreated *n*-heptane is below 1 pS/m.

NOTE 2—If the conductivity of the *n*-heptane after treatment, measured in accordance with Section 11 in a thoroughly cleaned cell, is higher than 0.1 pS/m, the treatment should be repeated.

7.3 *Hydrocarbon*, for calibration. The dielectric constant must be known to ±5 % at the temperature of calibration.⁹

8. Sampling

8.1 The sample volume should be at least 0.7 L.

⁷ A suitable grade is available from W. R. Grace & Co., Davison Chemical Division, Baltimore, MD 21202 by specifying Code 923.

⁸ A suitable product is filtering fiber Pyrex Wool. Catalogue No. 3950, supplied by Owens-Corning Fiber Glass Corp.

⁹ A standard, such as cyclohexane, with certified dielectric constant, may be obtained from the National Bureau of Standards, Washington, DC 20234.

8.2 Use a clean epoxy-lined can, or a new glass bottle that has been rinsed successively with hot water, distilled water, acetone, and cleaning solvent then flush with dry nitrogen. Use only non-contaminating caps.

NOTE 3—Test method results are known to be sensitive to trace contamination from sampling containers. For recommended sampling containers refer to Practice D 4306.

NOTE 4—Bottle samples should be *tested immediately*, since the glass surface tends to absorb from the fuel the conductive substances that the test method is intended to measure.

8.3 Rinse the container several times with portions of the aviation turbine fuel to be sampled (Jet A or A-1, (**Warning**—Combustible. Vapor harmful. See Annex A1.3.), Jet B (**Warning**—Extremely flammable. Harmful if inhaled. Vapors can cause flash fire. See Annex A1.4). If possible fill the container, let stand, then empty and refill. Avoid taking the sample for test by pouring from the container; pipet instead. The sample should be clean and bright when tested.

9. Preparation of Apparatus

9.1 *Cleaning the Cell*—The cleaning procedure to be used depends on the estimated conductivity of the sample to be tested.

9.1.1 For samples that are expected to exhibit conductivities below 1 pS/m, the KSLA cell should be completely disassembled, the parts cleaned and the cell reassembled using protective gloves.

9.1.1.1 Dismantle the cell by removing the loose battery cap, the outer electrode electrical connector and the bottom screw-on cap. Press the inner electrode towards the bottom of the outer electrode and remove the inner electrode TFE-fluorocarbon insulator assembly.

9.1.1.2 Each part of the cell should be rinsed thoroughly five times with cleaning solvent followed by rinsing with treated *n*-heptane. The parts should be dried with a stream of nitrogen gas.

9.1.1.3 After reassembly, the cell should be rinsed with treated *n*-heptane.

9.1.1.4 After cleaning, check the cleanliness of the cell by measuring the conductivity of treated *n*-heptane in accordance with Section 11. The corrected value should be lower than 0.05 pS/m.

9.1.2 For samples that are expected to exhibit conductivities above 1 pS/m, the KSLA cell still assembled should be rinsed five times with cleaning solvent, followed by rinsing with treated *n*-heptane. The cell should be dried with a stream of nitrogen gas.

9.1.2.1 After cleaning, check the cleanliness of the cell by measuring the conductivity of treated *n*-heptane in accordance with Section 11. The corrected value should be lower than 0.1 pS/m.

9.2 *Cleaning of Auxiliary Equipment:*

9.2.1 Pipets used to transfer samples should be rinsed inside and outside with cleaning solvent using a non-contaminating squeeze bottle, then blown dry with clean, dry nitrogen. Thermometers should be similarly rinsed and maintained.

NOTE 5—If a cell has been used to test samples of high-conductivity, that is, more than 1000 pS/m, it should be disassembled for thorough

cleaning. Very thorough cleaning may also be accomplished by placing the disassembled cell in a Soxhlet apparatus containing boiling toluene/isopropyl alcohol for several hours.

NOTE 6—If testing is to be done on both low-conductivity (<1 pS/m) and high-conductivity (>1000 pS/m) samples, separate cells are recommended.

10. Calibration and Standardization

10.1 *Checking the Test Equipment:*

10.1.1 Remove cell and cable from the meter.

10.1.2 Depress the 20-pS/m switch. The digital reading should indicate 0.00 ± 0.01 pS/m after 3 s. If readings exceed ± 0.01 either adjust zero or record the zero error for calculating final report value.

10.1.3 Depress the calibrate switch. The digital reading should indicate 1000 ± 3 pS/m.

10.1.4 If low battery indicator is displayed during measure or calibration, the internal batteries should be replaced.

10.2 *Checking the Cell Constant:*

10.2.1 A check on the cell constant is necessary only if the cell has been damaged. Two capacitance measurements are required with a precision a-c bridge. Make a rigid two-terminal connection between the cell assembly and the bridge. Measure the total capacitance, C_E (picofarad) of the empty assembly. Without disturbing the connection, add 100 mL of a hydrocarbon standard and measure the new total capacitance, C_S (picofarad) and the temperature in the cell. Alternatively, the cell can be sent to the manufacturer for recalibration.

10.2.2 Calculate the actual capacitance, C_A , of the empty cell as follows:

$$C_A = (C_S - C_E)/(k-1) \quad (2)$$

where:

k = dielectric constant of the hydrocarbon at test temperatures.

10.2.3 Calculate the cell constant as follows:

$$K = 8.854/C_A \quad (3)$$

10.2.4 The cell constant of a new KSLA cell is 1.0. Because of its configuration, the cell constant cannot change significantly unless there is gross damage.

11. Procedure

11.1 Attach the cleaned cell to the meter as shown in Fig. 2.

11.1.1 Depress the ZERO switch, the digital reading should indicate the same value recorded in 10.1.2.

11.1.2 Depress the CAL switch, the digital reading should indicate 1000 ± 5 .

11.1.3 Rinse the cleaned cell three times with the sample, empty completely, then fill the outer chamber until sample overflows into the center receptacle. (Alternatively, the outer chamber can be filled by pipet with 100 mL of sample).

NOTE 7—Static charges generated by handling the sample must be allowed to dissipate. If the sample is expected to have a conductivity below 1 pS/m, 15-min waiting time must be allowed before reading.

11.1.4 If the sample conductivity is known, select the corresponding range position. When the conductivity is unknown, first check the fuel on a 2000 pS/m range position, then read in a lower scale if appropriate.



FIG. 2 Cleaned Cell Attached to Meter

11.1.5 Using the AUTO mode, depress the appropriate RANGE switch. In the AUTO mode, the reading is stabilized after 3 s and held on display for 9 s. Record the pS/m value.

11.1.6 Repeat readings can be taken after a 1-min delay.

11.1.7 Remove the cell cover and measure the temperature of the test samples to the nearest 1°C with a clean thermometer.

12. Calculation

12.1 If the zero error in 10.1.2 was greater than ± 0.01 and the zero of the meter was not adjusted, the readings should be corrected by subtracting any positive zero error or adding any negative zero error. For example:

Sample reading (11.1.5)	1.67	
Uncorrected zero error (10.1.2)	+ 0.03	
Calculated reading	1.64	

13. Report

13.1 The report is to include the following:

13.1.1 The calculated conductivity of the sample in pS/m after correcting for the zero reading. Note MANUAL or AUTO mode.

13.1.2 Temperature of sample.

NOTE 8—It is recognized that the electrical conductivity of a fuel varies significantly with temperature and that the relationship differs for various types of aviation and distillate fuel. If it is necessary to correct conductivity readings to a particular temperature, each laboratory would have to establish this relationship for the fuels and temperature range of interest.

14. Precision and Bias ¹⁰

14.1 The precision of this test method obtained by statistical examination of test results in the range between 0.1 and 2000

pS/m by operator/instrument pairs at a common test site is described in 14.2 and 14.3.

NOTE 9—The data used to determine the precision of this test method were obtained using the auto mode.

14.2 *Repeatability*—The difference between successive test results obtained by the same operator with the same apparatus under constant operating conditions on identical test samples would, in the long run, in the normal and correct operation of the test method, exceed the values in Fig. 3 in only one case in twenty.

14.3 *Reproducibility*—The difference between two single and independent results obtained by different operators working at the same location on identical test materials would, in the long run, exceed the values in Fig. 3 only in one case in twenty.

14.3.1 In the event of a dispute or concern regarding shipped sample conductivity, it is recommended that operators come to the bulk hydrocarbon storage site to measure conductivity on freshly obtained samples according to the cited procedure. This ensures that samples identical to bulk storage are tested by either or both parties and the precision data in Fig. 3 shall apply (Note 10).

NOTE 10—In 1987, a test program using Test Method D 2624 was carried out to investigate reproducibility of results when samples are shipped between laboratories.¹¹ While repeatability values were similar to those from common site testing, it was concluded that adequate reproducibility values were not obtained due to changes in conductivity of samples during shipment and storage. The same possibility of conductivity change would occur for samples tested according to Test Method D 4308.

¹⁰ A report on statistical analysis of interlaboratory test results is on file in ASTM Headquarters as RR: D02-1170.

¹¹ A report of data and conclusions from this program is on file at ASTM Headquarters; request RR:D02-1235.

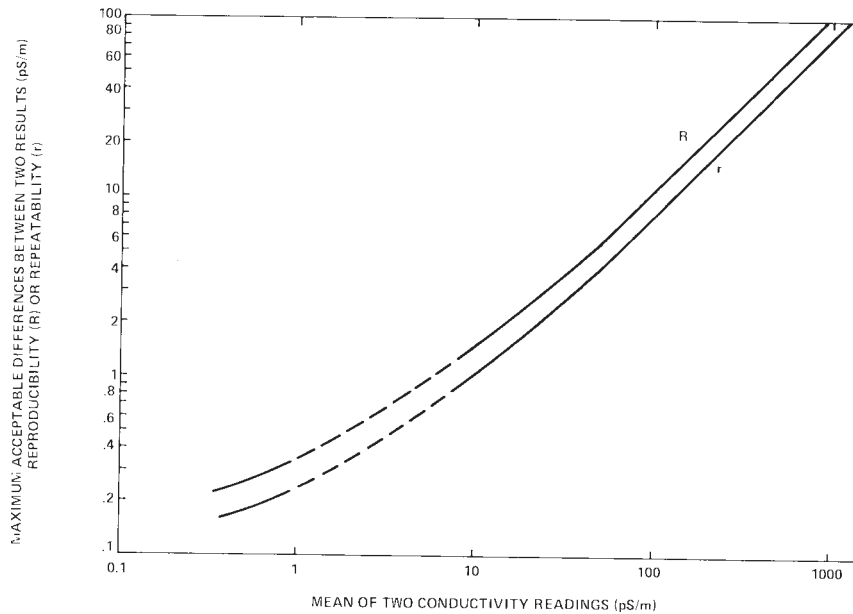


FIG. 3 Variation of Repeatability and Reproducibility with Conductivity Level

14.4 *Bias*—Since there is no accepted reference material suitable for determining the bias of the procedure in Test Method D 4308 for measuring conductivity, bias cannot be determined.

ANNEX

(Mandatory Information)

A1. PRECAUTIONARY STATEMENTS

A1.1 Toluene

Keep away from heat, sparks, and open flame.
 Keep container closed.
 Use with adequate ventilation.
 Avoid breathing of vapor or spray mist.
 Avoid prolonged or repeated contact with skin.

A1.2 n-Heptane

Keep away from heat, sparks, and open flame.
 Keep container closed.
 Use with adequate ventilation.
 Avoid prolonged breathing of vapor or spray mist.
 Avoid prolonged or repeated skin contact.


A1.3 Aviation Turbine Fuel (Jet A or A-1)

Keep away from heat, sparks, and open flame.
 Keep container closed.

Use with adequate ventilation.
 Avoid breathing vapor or spray mist.
 Avoid prolonged or repeated contact with skin.

A1.4 Aviation Turbine Fuel (Jet B)

Keep away from heat, sparks, and open flame.
 Keep container closed.
 Use with adequate ventilation.
 Avoid buildup of vapors and eliminate all sources of ignition, especially non-explosionproof electrical apparatus and heaters.
 Avoid breathing vapor or spray mist.
 Avoid prolonged or repeated contact with skin.

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