

Designation: D 1384 – 01

Standard Test Method for Corrosion Test for Engine Coolants in Glassware¹

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1. Scope

1.1 This test method covers a simple beaker-type procedure for evaluating the effects of engine coolants on metal specimens under controlled laboratory conditions (see Appendix X1).

NOTE 1-For more information on engine coolants, see Refs (1-8).²

1.2 The values stated in SI units are to be regarded as the standard. The values given in parentheses are for information only.

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 hazards statements are given in 10.1.7.2, 10.1.7.3, and 10.1.7.4.

2. Referenced Documents

- 2.1 ASTM Standards:
- B 32 Specification for Solder Metal³
- B 36/B36M Specification for Brass Plate, Sheet, Strip, and Rolled Bar⁴
- D 1176 Test Method for Sampling and Preparing Aqueous Solutions of Engine Coolants or Antirusts for Testing Purposes⁵
- E 1 Specification for ASTM Thermometers⁶
- E 178 Practice for Dealing with Outlying Observations⁷

3. Summary of Test Method

3.1 In this test method, specimens of metals typical of those present in engine cooling systems are totally immersed in aerated engine coolant solutions for 336 h at 88°C (190°F) for high-boiling engine coolant or corrosion inhibitors and 71°C

(160°F) for low-boiling engine coolant. The corrosioninhibitive properties of the test solution are evaluated on the basis of the weight changes incurred by the specimens. Each test is run in triplicate, and the average weight change is determined for each metal. A single test may occasionally be completely out of line (see 11.2).

4. Significance and Use

4.1 This test method will generally distinguish between coolants that are definitely deleterious from the corrosion standpoint and those that are suitable for further evaluation. However, the results of this test method cannot stand alone as evidence of satisfactory corrosion inhibition. The actual service value of an engine coolant formulation can be determined only by more comprehensive bench, dynamometer, and field tests.

5. Apparatus Ch. 21

5.1 Container—A 1000-mL, tall-form, spoutless beaker, made of heat-resistant glass, for containing the engine coolant solution and test specimens. The beaker shall be tightly closed with a No. 15 rubber stopper, having drill holes to accommodate a water condenser, an aerator tube, and a thermometer as shown in Fig. $1.^{8}$

5.2 *Condenser*—A water condenser of the reflux, glass-tube type, having a 400-mm (16-in.) condenser jacket.

5.3 *Aerator Tube*— A gas-dispersion tube, porosity size 12-C,⁹ to assure continuous aeration without plugging.

5.4 *Thermometer*— An ASTM Partial Immersion Thermometer having a range from -20 to 150° C (0 to 302° F) and conforming to the requirements for Thermometer 1C (1F), as prescribed in Specification E 1.

5.5 *Heater*—A constant-temperature bath containing a highboiling liquid (see Note 2) that is capable of giving continuous service with the specified temperature control.¹⁰ The size of the

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² The boldface numbers in parentheses refer to the list of references at the end of this standard.

³ Annual Book of ASTM Standards, Vol 02.04.

⁴ Annual Book of ASTM Standards, Vol 02.01.

⁵ Annual Book of ASTM Standards, Vol 15.05.

⁶ Annual Book of ASTM Standards, Vol 14.03.

⁷ Annual Book of ASTM Standards, Vol 14.02.

⁸ Optionally, an all-glass apparatus may be used. Contact ASTM Headquarters for details. Request Adjunct No. 12-413841-12.

⁹ Gas-dispersion tube No. 39533, manufactured by the Corning Glass Works, 44-5 Crystal St., Corning, NY, generally has been found satisfactory for this purpose. Optionally, a capillary tip bleed tube with 0.28-in. (7-mm) bore and 11.2-in. (280-mm) length may be used when consistent early plugging of gas dispersion tubes occurs. The tube, catalog No. 7815-19, may be obtained from the Corning Glass Works, Corning, NY 14830.

¹⁰ If a water bath is used, a significant reduction in evaporation rate is achieved by addition of floating plastic chips on the water surface.

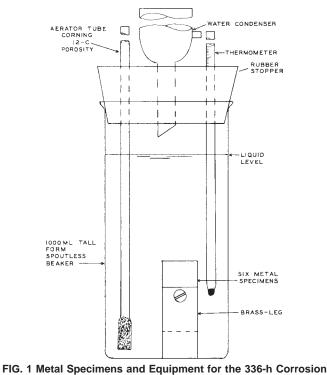


FIG. 1 Metal Specimens and Equipment for the 336-h Corrosion Test

bath will be determined by the number of corrosion tests that are to be run concurrently.

6. Metal Test Specimens

NOTE 2—The specimens prescribed in this test method have been accepted by automobile manufacturers, but their composition may not be the same as that of alloys currently used for engine cooling system components. Therefore, specimens other than those designated in this test method may be used by mutual agreement of the parties involved.

6.1 *Type*—The following metal test specimens,¹¹ representative of cooling system metals, shall be used:

6.1.1 *Steel*, UNS G10200 (SAE 1020),¹² cut from 1.59-mm ($\frac{1}{16}$ -in.) cold-rolled sheet stock to size 50.8 by 25.4 mm (2 by 1 in.). Chemical composition of the carbon steel is as follows: carbon, 0.17 to 0.23 %; manganese, 0.30 to 0.60 %; phosphorus, 0.040 % maximum; sulfur, 0.050 % maximum.

6.1.2 *Copper*, conforming to UNS C11000 (SAE CA110)¹² or UNS C11300 (SAE CA113)¹². Cold-rolled, cut from 1.59-mm ($\frac{1}{16}$ -in.) sheet stock to size 50.8 by 25.4 mm (2 by 1 in.).

6.1.3 *Brass*, conforming to Alloy UNS C26000 (SAE CA 260).¹³ Half-hard, cut from 1.59-mm ($\frac{1}{16}$ -in.) sheet stock to size 50.8 by 25.4 mm (2 by 1 in.).

¹² UNIFIED numbering system for metals and alloys, SAE-ASTM, July 1995.

6.1.4 Solder—A brass specimen as described in 6.1.3, coated with solder conforming to Alloy Grade 30A (SAE 3A) of Specification B 32.¹³ Solder-coated specimens may be prepared, or used specimens recoated for reuse, by the procedure given in Annex A1. A solid solder specimen cut from 1.59-mm ($\frac{1}{16}$ -in.) sheet stock of Alloy Grade 30A (SAE 3A) to size 50.8 by 25.4 mm (2 by 1 in.) may be used subject to mutual agreement of the parties involved. The use of a solid solder specimen must be reported along with the metal specimen weight loss results.

6.1.4.1 When agreed upon between the supplier and the purchaser of engine coolants, the standard solder specimen may be replaced with one having a different alloy composition than standard Alloy Grade 30A or 30B. Use of specimens other than standard Alloy Grade 30A or 30B shall be noted in the test report.

NOTE 3—Where non-standard alloy is used, the standard flux shown in A1.1.5 may not be satisfactory. A low corrosive flux may be required.

6.1.5 *Cast Aluminum*, conforming to Alloy UNS A23190 (SAE 329).¹² Specimen size, 50.8 by 25.4 by 3.18 mm (2 by 1 by $\frac{1}{\sin}$).

6.1.6 *Cast Iron*, conforming to Alloy UNS F10007 (SAE G3500).¹¹ Specimen size, 50.8 by 25.4 by 3.18 mm (2 by 1 by $\frac{1}{\sin}$).

6.2 Arrangement (See Fig. 2):

6.2.1 *Metal Specimen Arrangement*—None of the hardware used in metal specimen arrangement (metal specimen, screws, washers, metal spacers, insulating sleeves, insulating spacers and nuts) can be reused for a test. The metal test specimens shall be drilled through the center with a 6.75-mm ($^{17}_{64}$ -in.) drill to accommodate a 50.8-mm (2-in.) 10–24 brass machine screw covered with a thin-walled insulating sleeve. Tetrafluoroethylene tubing with a 6.35-mm ($^{14}_{-10}$.) outside diameter 1.59-mm ($^{14}_{-10}$ -in.) wide and a wall thickness of 0.4 mm ($^{164}_{-10}$.) is satisfactory. Two half-hard brass legs shall be cut from 1.59-mm ($^{14}_{-10}$.) sheet stock to size 50.8 by 25.4 mm (2 by 1 in.). A 6.35-mm ($^{14}_{-10}$.) diameter hole shall be drilled in each leg with the center 6.35 mm ($^{14}_{-10}$.) from the top and 12.7 mm ($^{14}_{-10}$.) from each side. The test "bundle" shall be made up

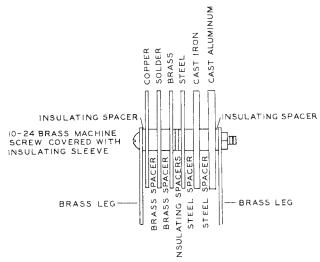


FIG. 2 Metal Specimen Arrangement

¹¹ Complete sets or individual metal test specimens are available from (*a*) Chemical Specialties Manufacturers Association, Inc., Suite 1120, 1001 Connecticut Ave., N.W., Washington, DC 20036; (*b*) Astro-Mechanics, Inc., 8500 Research Blvd., Austin, TX 78766; (*c*) The Metaspec Company, P.O. Box 27707, San Antonio, TX 78227; or (*d*) Metal Samples Co. Inc., P.O. Box 8, Munford, AL 36268.

¹³ Round-robin evaluation of coated solder report is available from ASTM Headquarters. Request RR:D15-0132.

on the insulated screw with the specimens in the following order: brass leg, copper, solder, brass, steel, cast iron, cast aluminum, and brass leg. The specimens shall be separated by 4.76-mm ($\frac{3}{16}\text{-in.}$) thick solid metal spacers having a 6.75-mm ($\frac{17}{64}\text{-in.}$) inside diameter and a 11.11-mm ($\frac{7}{16}\text{-in.}$) outside diameter. Insulating spacers made from tetrafluoroethylene shall be used between the brass legs and the specimen "bundle," and between the brass and steel specimens. Brass spacers shall be used between the brass, solder, and copper specimens, and steel spacers between the cast iron, steel, and cast aluminum specimens. The nut shall be tightened firmly to ensure good electrical contact between the test specimens in each section of the "bundle."

6.2.2 Alternate Metal Specimen Arrangement—When agreed upon between the supplier and the purchaser, an alternate metal specimen arrangement may be used to evaluate multiple solder alloys, such as high lead Alloy Grade L50113¹³ consisting of 97 % lead, 2.5 % tin, 0.3 % silver, concurrently with Standard Alloy Grade 30A or 30B. It is recommended that the metal specimen arrangement be modified by replacing the copper specimen with the high lead solder specimen and arranging specimens in the bundle as follows:

High Lead	Brass	Alloy Grade	Steel	Cast Iron	Cast Alumi-
Solder		30A or 30B			num

Use of alternate specimens and metal specimens arrangements shall be noted in the test report.

7. Preparation of Test Specimens

7.1 Sand the cast iron and cast aluminum specimens on the 25.4 by 50.8-mm (1 by 2-in.) cut surfaces with "coarse" grade (No. 1) emery cloth. Remove any burrs from coupon edges and hole. Scrub all specimens vigorously, using a moistened bristle brush and ground pumice powder or fine silicon carbide grit until the entire metal area is bright, shiny, and free from any visible oxide film or tarnish.

7.2 Rinse the specimens thoroughly with tap water; then rinse with acetone, dry, and weigh to the nearest 1 mg. Cast aluminum specimens should be dried in a 100°C oven for 1 h, to a constant weight, prior to recording the weight.

NOTE 4—If the test specimens are not to be used immediately, keep them in a desiccator until required.

8. Test Solutions

8.1 The concentration of the engine coolant to be tested shall be as follows:

8.1.1 *Engine Coolant*—The engine coolant, EG or PG based, shall be mixed with the proper quantity of corrosive water to give a 33¹/₃volume % coolant test solution.

8.1.2 *Corrosive Water* (Note 4)—The corrosive water shall contain 100 ppm each of sulfate, chloride, and bicarbonate ions introduced as sodium salts.

8.2 *Preparation of Sample*—The preparation of the sample shall be done in accordance with the section on Preparation of Solutions Requiring Inclusion of Separated Solids and Liquids in Test Method D 1176, except that the corrosive water shall be used for dilution instead of distilled water. Thus, any insoluble materials will be included in the representative sample.

NOTE 5-The specified corrosive water can be prepared by dissolving

the following amounts of anhydrous sodium salts in a quanity of distilled or deionized water.

sodium sulfate	148 mg
sodium chloride	165 mg
sodium bicarbonate	138 mg

The resulting solution should be made up to a volume of 1 L with distilled or deionized water at 20°C.

If relatively large amounts of corrosive water are needed for testing, a concentrate may be prepared by dissolving ten times the above amounts of the three chemicals, in distilled or deionized water, and adjusting the total volume to 1 L by further additions of distilled or deionized water. When needed, the water concentrate is diluted to the ratio of one part by volume of concentrate to nine parts of distilled or deionized water.

9. Test Conditions

9.1 *Beaker Assembly*—The arrangement of the assembled metal specimens with relation to the aerator tube and other components is shown in Fig. 1. Note that the tip of the condenser just emerges from the bottom of the rubber stopper.

9.2 *Test Temperature*—The test solution shall be maintained at a temperature of $88 \pm 2^{\circ}$ C (190 $\pm 5^{\circ}$ F) for high-boiling engine coolants.

9.3 Aeration Rate—The aeration rate shall be 100 ± 10 mL/min. The aerator tube should be located at least 12.7 mm (¹/₂in.) away from the test "bundle" to avoid direct contact with the metal specimens.

9.4 *Test Duration*—The test shall be run continuously for 2 weeks (336 h).

10. Procedure

10.1 Make triplicate tests concurrently on each engine coolant solution in accordance with the following procedure:

10.1.1 Carefully clean the test beaker, condenser, rubber stopper, and aerator tube, and thoroughly rinse with water.

10.1.2 Bolt the specimens together in the order given in 6.2 and place the "bundle" in the test beaker as shown in Fig. 1.

 $10.1.3\,$ Pour 750 mL of the prepared test solution into the 1000-mL beaker.

10.1.4 Fit the condenser and aeration tube to the beaker, and set the aeration rate at 100 mL/min, using a flowmeter or other suitable device.

10.1.5 Raise the temperature of the test solution to 88° C (190°F) for high-boiling engine coolants. Pass water through the condenser at a rate sufficient to maintain adequate cooling.

10.1.6 Check the tests once each working day to ensure proper solution temperature, aeration rate, and solution level. The tests may operate unattended on weekends and holidays. Make up evaporation losses during the corrosion tests by addition of distilled or deionized water.

10.1.7 At the end of the test, *immediately* disassemble specimens and brush very lightly with a soft bristle brush and water to remove loosely held corrosion products. To remove the more tenacious corrosion products and films, the individual specimens shall then be subjected to additional cleaning treatments as follows:

10.1.7.1 *Iron and Steel*—Remove adherent deposits by means of a brass scraper or brass bristle brush, followed by scrubbing with a wet bristle brush and fine pumice to clean the specimen completely.