



Standard Test Method for Determination of Moisture in New and In-Service Lubricating Oils and Additives by Relative Humidity Sensor¹

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1. Scope

1.1 This test method covers the quantitative determination of water in new and in-service lubricating oils and additives in the range of 10 to 100 000 mg/kg (0.001 to 10% wt./wt.) using a relative humidity (RH) sensor. Methanol, acetonitrile, and other compounds are known to interfere with this test method.

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

1.3 **Warning**—Samples tested in this test method can be flammable, explosive, and toxic. Use caution when handling them before and after testing.

1.4 *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.*

2. Referenced Documents

2.1 *ASTM Standards:*²

[D4175 Terminology Relating to Petroleum, Petroleum Products, and Lubricants](#)

3. Terminology

3.1 For definitions of terms used in this test method, refer to Terminology [D4175](#).

4. Summary of Test Method

4.1 An aliquot of sample is injected onto a heated stainless steel coil. The temperature of the coil is programmable in 1°C increments from 25°C to 200°C. The coil is maintained at a constant temperature for the duration of the test. As the sample travels down the coil, an opposing dry inert gas carries the

thermally evolved moisture past a relative humidity sensor. The sensor signal is integrated over time to provide a measurement of total mass of water in the sample.

4.2 The sample injection may be done either by mass or by volume.

4.3 This test method utilizes anhydrous compressed gas or ambient air passed through a desiccant to prevent contamination from moisture present in the atmosphere.

4.4 Viscous samples can be analyzed by preheating them to place them in a more fluid state allowing them to be drawn into a syringe, or by dissolving them in a compatible anhydrous solvent. Care should be taken to minimize time spent preheating samples to prevent moisture loss.

5. Significance and Use

5.1 Knowledge of the water content of lubricating oils, additives, and similar products is important in the manufacture, purchase, sale, transfer, or use of such petroleum products to help in predicting their quality and performance characteristics.

5.2 For lubricating oils, the presence of water can lead to premature corrosion and wear, an increase in the debris load resulting in diminished lubrication and premature plugging of filters, impedance to the effect of additives, and undesirable support of deleterious bacterial growth.

6. Interferences

6.1 Methanol and acetonitrile are known to interfere with the determination of moisture by this test method. These substances contribute to a high bias in the final results. More generally, some short-chained polar molecules mimic the effect of water at the RH sensor resulting in a positive interference. Strong polar solvents, such as n-methyl-pyrrolidone, can severely damage the RH sensor.

7. Apparatus

7.1 *Moisture Analyzer*,³ an apparatus that consists of:

³ The sole source of supply of the apparatus known to the committee at this time is Arizona Instrument, 3375 N Delaware St., Chandler, AZ 85225. If you are aware of alternative suppliers, please provide this information to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee,¹ which you may attend.

¹ This test method is under the jurisdiction of ASTM Committee D02 on Petroleum Products and Lubricants and is the direct responsibility of Subcommittee D02.96.02 on Chemistry for the Evaluation of In-Service Lubricants.

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² 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.

TABLE 1 Suggested Injection Volume Based on Expected Water Content

Expected Water Concentration, %	Sample Volume, mL
Less than 0.02	5
0.02 to 0.025	4
0.025 to 0.035	3
0.035 to 0.050	2
Greater than 0.05	1

7.1.1 *Flow Regulator*, capable of maintaining the carrier gas flow rate within the manufacturer’s specified conditions.

7.1.2 *Flow Meter*, capable of measuring the carrier gas flow rate within the manufacturer’s specified conditions.

7.1.3 *Stainless Steel Sample Coil*, for heating the sample as it is transported from the sample inlet to the sump.

7.1.4 *Sample Coil Heater*, capable of maintaining the sample coil temperature within 1°C of the programmed temperature between 25°C and 200°C.

7.1.5 *Sample Delivery System*, provides programmable variable speed injection of the sample into the coil.

7.1.6 *Sump*, allows for collection of the sample at the bottom of the sample coil during testing and discharge of the sample to a waste container after testing is completed.

7.1.7 *Manifold*, which provides:

7.1.7.1 A thermally stable port for mounting and operation of the relative humidity sensor.

7.1.7.2 Inlet and outlet ports for the carrier gas.

7.1.8 *Relative Humidity (RH) Sensor*, a capacitive sensing element that measures the relative humidity of the carrier gas.

7.1.9 *Microcontroller*, which provides:

7.1.9.1 Capability of integrating and converting the RH sensor signal.

7.1.9.2 Capability of controlling the temperature of the coil heater, and the sensor manifold.

7.1.9.3 Capability of controlling the speed of the sample delivery system.

7.1.10 Balance, external, with 1 mg precision for weighing sample.

8. Reagents and Materials

8.1 *Traceable⁴ Syringe*, traceable at the desired total mass of water, typically 500 µg (0.5 µL).

8.2 *Glass or Plastic Syringe*, 1 or 5 mL capacity.

8.3 *Needle*, 18- or 22-gauge.

8.4 *Water*, deionized.

8.5 *Carrier Gas*, any dry inert gas including, but not limited to, dry air, nitrogen, helium, or argon.

9. Sampling

9.1 Laboratory sample shall be thoroughly homogeneous before drawing a test specimen.

9.2 Select test specimen size as indicated in [Table 1](#) based on the expected water concentration.

⁴ Traceable to national or international standards like NIST, ISO, etc., as defined in Terminology [D4175](#).

10. Preparation of Apparatus

10.1 Establish carrier gas flow to the analyzer by either opening the source regulator or turning on the dry air generator.

10.2 Turn on analyzer and allow equilibration for at least 15 min.

11. Calibration and Standardization

11.1 To ensure the integrity of the test results, the RH sensor shall be verified and calibrated using a traceable syringe. Alternatively, the RH sensor may be verified using a traceable standard solution of water in a compatible solvent. Other suitable instrument calibration methods and standards may be used as specified by the instrument manufacturer.

NOTE 1—Examples of suitable water standards include water in propylene carbonate or water in xylenes.

11.2 Perform the coil heater calibration in accordance with the manufacturer’s instructions. After calibration, the coil shall verifiably maintain an arbitrary set temperature from 25 to 200°C within ±1°C. Coil heater calibration should be performed at least once annually.

11.2.1 If the results are not within the acceptable range, contact the analyzer manufacturer.

11.3 To perform the RH sensor verification, set the instrument to calibration/verification mode and inject 0.5 µL water directly into the sensor chamber of the instrument using the calibrated syringe. Acceptable results are 475 to 525 µg water detected.

11.3.1 If the RH sensor verification is not within acceptable range:

11.3.2 Perform at least five injections using the procedure outlined in [11.3](#). If the coefficient of variation between the five injections is < 2%, use the mean result of the five injections to perform a single-point recalibration of the instrument.

11.3.3 Repeat step [11.3](#) to verify RH sensor calibration.

11.3.4 If results are not within acceptable range, contact the analyzer manufacturer.

12. Procedure

12.1 *Sample Analysis*:

12.1.1 Program the analyzer with appropriate test conditions.

12.1.2 Flush clean syringe and needle 2 to 5 times with the material to be tested.

12.1.3 Place empty syringe on the balance, and tare it.

12.1.4 Draw desired amount of sample into the syringe.

NOTE 2—Suggested test specimen sizes are listed in [Table 1](#).

12.1.5 Place syringe with test specimen back onto balance and reweigh.

12.1.6 Record the sample weight.

12.1.7 Load the syringe with the test specimen in the testing portion of the analyzer.

12.1.8 Begin the program and follow the instrument provided prompts for starting analysis.

12.1.9 Record result displayed at the end of the test.

12.1.10 Repeat [12.1.4-12.1.9](#) for subsequent tests.

TABLE 2 Repeatability Limits for Instrument Utilizing Relative Humidity Sensor

NOTE—Repeatability data based on 30 tests per material/single operator/two instruments

Material =	Hydraulic Fluid
Sample Size =	4 mL
Test Temperature =	120°C
Syringe Speed =	5.0 min
Ending Criteria =	Rate = 0.10 µg/s
Mean =	0.0277% (wt./wt.)
Standard Deviation =	0.0017
Repeatability Limit (2.77 *S.D.) =	0.0046
<hr/>	
Material =	Turbine Oil
Sample Size =	5 mL
Test Temperature =	120°C
Syringe Speed =	5.0 min
Ending Criteria =	Rate = 0.10 µg/s
Mean =	0.0015% (wt./wt.)
Standard Deviation =	0.0002
Repeatability Limit (2.77 *S.D.) =	0.0007

13. Calculation or Interpretation of Results

13.1 Calculate the water concentration in mass or volume % of the sample as follows:

$$\text{water, mass \%} = \frac{R}{10^4 \times W} \text{ or} \quad (1)$$

$$\text{volume \%} = \frac{R}{10^4 \times V}$$

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where:

R = total water result for test specimen, µg,

W = sample weight, g, and

V = volume of sample used, mL.

13.2 If results are displayed as percent of water present and conversion to mg/kg is desired, calculate as follows:

$$\text{mg/kg} = \text{water (\%)} \times 10^4 \quad (2)$$

13.3 No further calculation or interpretation is necessary.

14. Report

14.1 Report the water concentration to the nearest whole mg/kg the nearest 0.01 mass %, the nearest whole µL/mL, or the nearest 0.01 volume %.

15. Precision and Bias ⁵

15.1 Precision and bias statements will be established following completion of the round robin analyses in accordance with ASTM requirements. See Table 2 for information on repeatability of this test method.

16. Keywords

16.1 additives; lubricating oils; moisture determination; relative humidity sensor

⁵ Supporting data have been filed at ASTM International Headquarters and may be obtained by requesting Research Report RR: D02-1674.