



Designation: D1552 – 16

Standard Test Method for Sulfur in Petroleum Products by High Temperature Combustion and Infrared (IR) Detection or Thermal Conductivity Detection (TCD)¹

This standard is issued under the fixed designation D1552; 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 U.S. Department of Defense.

1. Scope*

1.1 This test method covers procedures for the determination of total sulfur in petroleum products including lubricating oils containing additives, and in additive concentrates. This test method is applicable to samples boiling above 177 °C (350 °F) and containing a mass fraction of sulfur between 0.22 % and 24.2 %. Other sulfur concentrations may be analyzed, but the precision stated may or may not apply. These procedures use IR detection or TCD following combustion in a furnace.

1.2 Petroleum coke containing a mass fraction of sulfur between 2.53 % to 3.79 % sulfur may be analyzed. Other sulfur concentrations may be analyzed, but the precision stated may or may not apply.

NOTE 1—The D1552–08 (2014) version of this standard contained two other procedures using iodate titrations. Since these procedures are no longer being used in the industry laboratories based on a survey of D02.SC 3 laboratories conducted in September 2014, they are being deleted. For earlier information on the deleted procedures, D1552–08 (2014) may be perused.

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

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

¹ This test method is under the jurisdiction of ASTM Committee D02 on Petroleum Products, Liquid Fuels, and Lubricants and is the direct responsibility of Subcommittee D02.03 on Elemental Analysis.

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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.

- D4057 Practice for Manual Sampling of Petroleum and Petroleum Products
- D4177 Practice for Automatic Sampling of Petroleum and Petroleum Products
- D6299 Practice for Applying Statistical Quality Assurance and Control Charting Techniques to Evaluate Analytical Measurement System Performance
- D6300 Practice for Determination of Precision and Bias Data for Use in Test Methods for Petroleum Products and Lubricants
- D6792 Practice for Quality System in Petroleum Products and Lubricants Testing Laboratories

3. Summary of Test Method

3.1 The sample is weighed and placed into a furnace having an oxygen atmosphere sufficient to combust the entire sample and a temperature between 1150 °C and 1450 °C for Procedure A or 1150 °C for Procedure B. Most sulfur present is combusted to SO₂, which is then measured with a detector after moisture and dust are removed by traps. The instrument calculates the mass percent sulfur from the sample mass, the integrated detector signal, and a predetermined calibration factor. The calibration factor is determined using standards approximating the material to be analyzed.

3.1.1 *Procedure A*—After combustion of the sample and subsequent moisture/dust removal, SO₂ is measured using infrared (IR) detection.

3.1.2 *Procedure B*—After combustion of the sample and subsequent moisture/dust removal, SO₂ is measured using thermal conductivity detection. An apparatus utilizing TCD may require the sample gas to pass an oxygen scrubber and adsorption/desorption traps to allow passing of contaminants prior to measuring SO₂.

4. Significance and Use

4.1 This test method provides a means of monitoring the sulfur level of various petroleum products and additives. This knowledge can be used to predict performance, handling, or processing properties. In some cases the presence of sulfur compounds is beneficial to the product and monitoring the

*A Summary of Changes section appears at the end of this standard

depletion of sulfur can provide useful information. In other cases the presence of sulfur compounds is detrimental to the processing or use of the product.

5. Apparatus

5.1 Combustion Furnace:

5.1.1 *Procedure A*—Furnace capable of maintaining a constant temperature (between 1150 °C minimum and 1450 °C maximum) sufficient to ensure quantitative recovery of sulfur as its corresponding gas SO₂.

5.1.2 *Procedure B*—Furnace capable of maintaining a temperature (1150 °C) sufficient to ensure quantitative recovery of sulfur as its corresponding gas SO₂.

5.2 *Combustion and Sulfur Detection System*, comprised of automatic balance, gas flow controls, drying tubes, oxygen scrubber, adsorption/desorption traps as required, combustion furnace, combustion boats or tin (Sn) containers as required and either an IR (Procedure A) or TCD (Procedure B) SO₂ detector.

5.3 *Sieve*, 60 mesh (250 μm).

6. Reagents and Materials

6.1 *Purity of Reagents*—Reagent grade chemicals shall be used in all tests. Unless otherwise indicated, it is intended that all reagents shall conform to the specifications of the Committee on Analytical Reagents of the American Chemical Society, where such specifications are available.³ Other grades may be used, provided it is first ascertained that the reagent is of sufficiently high purity to permit its use without lessening the accuracy of the determination.

6.2 *Combustion Promoter*—Magnesium oxide (MgO), tungsten trioxide (WO₃), or aluminum oxide (Al₂O₃). Alternatively, COM-CAT, a dual promoter that is both a fixing agent and oxidizing agent, may be used.⁴

6.3 *Drying Agent*, Magnesium perchlorate: anhydrous (Mg(ClO₄)₂) or phosphorus pentoxide (P₂O₅). (**Warning**—In addition to other precautions, handle magnesium perchlorate with care. Avoid contacting it with acid and organic materials. Reactions with fuel may be violent.)

6.4 *Oxygen (Extra Dry)*—The oxygen shall be at least 99.5 % pure and show no detectable sulfur by blank determination. (**Warning**—Oxygen vigorously accelerates combustion.)

6.5 *Inert Gas*—Helium or argon, high purity grade, 99.995 % minimum purity, as required by manufacturer's recommendations.

³ *Reagent Chemicals, American Chemical Society Specifications*, American Chemical Society, Washington, DC. For Suggestions on the testing of reagents not listed by the American Chemical Society, see *Annual Standards for Laboratory Chemicals*, BDH Ltd., Poole, Dorset, U.K., and the *United States Pharmacopeia and National Formulary*, U.S. Pharmacopeial Convention, Inc. (USPC), Rockville, MD.

⁴ The sole source of supply of COM-CAT known to the committee at this time is LECO Corporation, 3000 Lakeview Ave. St. Joseph, MI 49085 USA. 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.

6.6 *Quality Control (QC) Sample(s)*, preferably are portions of one or more petroleum products that are stable and representative of the samples of interest. These QC samples can be used to check the validity of the testing process and performance of the instrument as described in Section 11.

7. Sampling

7.1 Take samples in accordance with the instructions in Practice D4057 or D4177.

8. Preparation of Apparatus

8.1 Assemble and adjust apparatus according to manufacturer's instructions. Initialize instrument, check power supplies, set gas pressure and flows, and set furnace temperature.

8.1.1 Condition the instrument with samples that are representative or typical of the sample types to be analyzed. During the interlaboratory study, laboratories analyzed between one and five conditioning samples.

8.1.2 Calibrate the automatic balance according to manufacturer's instructions.

9. Standardization

9.1 Determination of Standardization Factor:

9.1.1 Because effects such as sample volatility can also affect the relative recovery as SO₂ of the sulfur originally present in the sample, it is necessary to determine a standardization factor. Proceed as described in Sections 9 through 12, using an oil sample of similar type to the unknown sample and of accurately known sulfur content.⁵

9.1.2 Determine the calibration factor for the particular type of sample to be analyzed (lubricating oil, petroleum coke, residual fuel) as recommended by the manufacturer.

9.2 *Quality Control*—Run a suitable analytical quality control sample several times daily. When the observed value lies between acceptable limits on a quality control chart, proceed with sample determinations.

10. Preparation of Coke

10.1 It is assumed that a representative sample has been received for analysis. If the sample of coke received is not dry, it is recommended that the sample be dried prior to grinding.

10.2 Grind and sieve the sample received so as to pass a 60 mesh (250 μm) sieve.

10.3 Dry the sieved material to constant weight at 105 °C to 110 °C.

11. Analysis of Quality Control Samples

11.1 A QC sample shall be analyzed each day samples are analyzed to verify the testing procedure and instrument performance. Additional QC samples may be analyzed. The QC samples shall be treated as outlined in Section 12, depending upon the type of furnace set-up used by the lab.

⁵ Residual fuel oil Standard Reference Materials may be obtained from the National Institute of Standards and Technology or other sources.