



# Standard Test Method for Sulfur in Petroleum Products by Hydrogenolysis and Rateometric Colorimetry<sup>1</sup>

This standard is issued under the fixed designation D4045; 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 reappraisal. A superscript epsilon ( $\epsilon$ ) indicates an editorial change since the last revision or reappraisal.

## 1. Scope\*

1.1 This test method covers the determination of sulfur in petroleum products in the range from 0.02 mg/kg to 10.00 mg/kg.

1.2 This test method may be extended to higher concentration by dilution.

1.3 This test method is applicable to liquids whose boiling points are between 30 °C and 371 °C (86 °F and 700 °F). Materials that can be analyzed include naphtha, kerosine, alcohol, steam condensate, various distillates, jet fuel, benzene, and toluene.

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

1.4.1 Certain specifications for the recorder (see 5.5) are excepted.

1.5 *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:*<sup>2</sup>

[D1193 Specification for Reagent Water](#)

[D6299 Practice for Applying Statistical Quality Assurance and Control Charting Techniques to Evaluate Analytical Measurement System Performance](#)

## 3. Summary of Test Method

3.1 The sample is injected at a constant rate into a flowing hydrogen stream in a hydrogenolysis apparatus. The sample

and hydrogen are pyrolyzed at a temperature of 1300 °C, or above, to convert sulfur compounds to hydrogen sulfide (H<sub>2</sub>S). Readout is by the rateometric detection of the colorimetric reaction of H<sub>2</sub>S with lead acetate. Condensable components are converted to gaseous products, such as methane, during hydrogenolysis.

## 4. Significance and Use

4.1 In many petroleum refining processes, low levels of sulfur in feed stocks may poison expensive catalysts. This test method can be used to monitor the amount of sulfur in such petroleum fractions.

4.2 This test method may also be used as a quality-control tool for sulfur determination in finished products.

## 5. Apparatus<sup>3,4</sup>

5.1 *Pyrolysis Furnace*—A furnace that can provide an adjustable temperature from 900 °C to 1400 °C in a 5 mm inside diameter or larger tube is required to pyrolyze the sample. The furnace entry temperature shall allow insertion of the hypodermic tip to a depth at which the temperature is 550 °C to provide sample vaporization at the injection syringe tip. This temperature shall be above the boiling point of the sample and of the sulfur compounds in the sample (see Fig. 1). The pyrolyzer tube may be of quartz; however, the lifetime is limited above 1250 °C. Ceramic may be used at any temperature.

5.2 *Rateometric H<sub>2</sub>S Readout*—Hydrogenolysis products contain H<sub>2</sub>S in proportion to sulfur in the sample. The H<sub>2</sub>S is measured by measuring rate of change of reflectance caused by darkening when lead sulfide is formed. Rateometric electronics, adapted to provide a first derivative output, allows sufficient sensitivity to measure below 0.1 mg/L (see Fig. 2).

<sup>1</sup> 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.

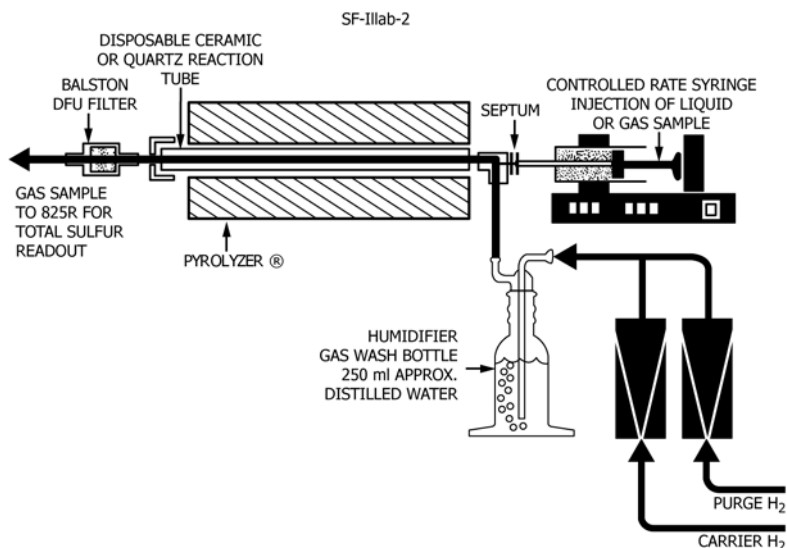
Current edition approved April 1, 2015. Published April 2015. Originally approved in 1987. Last previous edition approved in 2010 as D4045 – 04 (2010). DOI: 10.1520/D4045-15.

<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.

<sup>3</sup> The apparatus described in 5.1 – 5.4 inclusive is similar in specification to the equipment available from Analytical Systems Keco, 9215 Solon Rd., Suite A4, Houston, TX 77064. For further information see Drushel, H. V., "Trace Sulfur Determination Petroleum Fractions," *Analytical Chemistry*, Vol 50, 1978, p. 76.

<sup>4</sup> The sole source of supply of the apparatus known to the committee at this time is Analytical Systems Keco. 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,<sup>1</sup> which you may attend.

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



NOTE 1—The humidifier gas wash bottle is optional.

FIG. 1 Hydrogenolysis Flow Diagram

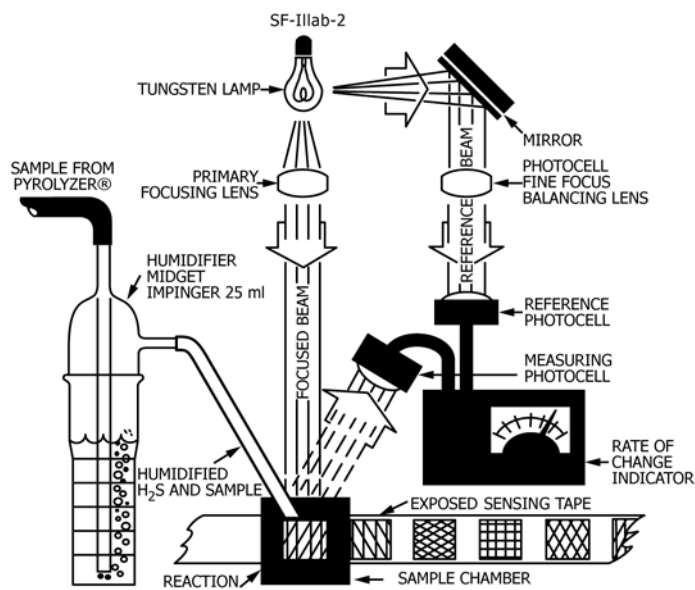


FIG. 2 Photorateometry H<sub>2</sub>S Readout

5.3 *Hypodermic Syringe*—A hypodermic having a needle long enough to reach the 550 °C zone is required. A side port is convenient for vacuum filling and for flushing the syringe. A 100 µL syringe is satisfactory for injection rates down to 3 µL/min and a 25 µL syringe for lower rates. (**Warning**—Exercise caution as hypodermics can cause accidental injury.)

5.4 *Syringe Injection Drive*—The drive shall provide uniform, continuous sample injections. Variation in drive injection rate caused by mechanical irregularities of gears will cause noise. The adjustable drive shall be capable of injection from 6 µL/min to 0.06 µL/min over a 6 min interval.

5.5 *Recorder*—A chart recorder with 10 V full scale and 10 000-Ω input or greater and having a chart speed of 0.2 in./min to 1 in./min (approximately 0.5 cm/min to 3 cm-

/min) is required. An attenuator can be used for more sensitive recorders. Newer instruments that display the results are acceptable in lieu of a chart recorder.

5.6 *Thermocouple*—A thermocouple suitable for use at 500 °C to 1400 °C, 250 mm long with readout is required. Type K, 1/16 in. (1.6 mm) diameter, Type 316 stainless steel sheath is suitable.

## 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 conform to the specifications of the Committee on Analytical Reagents of the American Chemical Society where such specifications are available.<sup>5</sup> 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 *Purity of Water*—Unless otherwise indicated, references to water shall be understood to mean Type II reagent grade water conforming to Specification D1193.

6.3 *Sensing Tape*—Lead acetate impregnated paper of chromatographic quality shall be used. (**Warning**—Lead is a cumulative poison.)

6.4 *Hydrogen*—As no commercial grade of hydrogen has a sulfur specification sufficiently low, each new source of supply must be tested. A change in the zero base line of 5 % of full scale from no flow to full flow indicates impure hydrogen. (**Warning**—Extremely flammable gas under pressure. Hydrogen is a flammable gas. Test all flow systems for leaks and

<sup>5</sup> *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.

purge with inert gas before introducing hydrogen and after removing hydrogen. Keep all flow systems as small in volume as practical and provide protective screening for containers other than sample flow lines. Dispose of exhaust gases in a fume hood or by vacuuming to a safe area. If gas cylinders are used handle carefully as rupture of the valve or cylinder is dangerous.)

### 6.5 Reference Standards:

6.5.1 *2,2,4 Trimethyl Pentane (Isooctane)*—(**Warning**—Extremely flammable.)

6.5.1.1 ASTM Knock Test Reference Fuel<sup>6</sup> may be used as the solvent. However, when this material is used, each new lot shall be tested for sulfur by this procedure because the specifications are not rigorous enough for this application.

NOTE 1—Heptane or equivalent material may also be used but precision data is based on the use of *isooctane*.

6.6 *Acetic Acid Solution*—Mix glacial acetic acid 1 part by volume into 19 parts water (see 6.2). (**Warning**—Corrosive.)

6.7 *Di-n-Butyl Sulfide*—( $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2$ )<sub>2</sub>S) is used to prepare standards. Equivalent sulfur compound may be used if care is exercised to prevent more volatile compounds from evaporating during preparation or use of standards.

6.8 *Helium or Nitrogen Purge Gas*. (**Warning**—Compressed gas under high pressure.)

## 7. Preparation of Apparatus

7.1 Turn on the furnace with temperature controls at minimum. Gradually increase furnace control over a 3 h period to approximately 1300 °C to minimize thermal shock. Reverse the procedure when preparing for long-term storage. For shutdown at night and over weekends, reduce temperature to about 900 °C but do not turn off the furnace. Furnace and quartz tubing life are extended by not cooling to room temperature.

7.2 Connect all tubing and fill prehumidifier outside the cabinet with water if this apparatus is being used, and final humidifier inside the cabinet with 5 % by volume acetic acid solution. Purge with inert gas, then close valve. Check all connections and repair any leaks that are found. (**Warning**—On instruments where electronics are in close proximity with the flow systems, exercise care if leak checking with a soap solution. A suitable method is to block the line that goes to the final humidifier and observe the flow meter. If the flow drops to zero, the flow systems are adequately leak-free.) Set hydrogen flow at 200 mL/min, and allow temperature to stabilize. (**Warning**—Extremely flammable gas under pressure.) Make final temperature adjustment to 1315 °C ± 15 °C. Use a standard thermocouple to verify temperature by inserting through a septum with hydrogen flowing at the rate noted above. Determine depth of insertion required, and always insert the hypodermic tip to the 550 °C point (see 7.6). (**Warning**—The use of a humidifier gas wash bottle filled with approximately 250 mL of water is a potential safety hazard as hydrogen pressure may build up inside the container.

The user of this test method should take appropriate safety measures to prevent an accidental injury, if the humidifier gas wash bottle is used in the analysis.)

7.3 Prepare the sample injection drive. Check to be sure desired injection rate is obtained at various settings. Verify that erratic pulses of fast drive do not occur when the drive range is switched. Pulses of high sample flow above 15 µL/min will cause carboning and spurious readings.

7.4 Install sensing tape, and turn H<sub>2</sub>S readout analyzer on.

7.5 Connect the recorder and adjust the zero to desired position with hydrogen flowing.

7.6 Fill syringe with blank reference standard solution, typically *isooctane*, insert the needle through the septum to the 550 °C temperature zone, and clamp to the syringe drive. At high temperature the hot needle may absorb sulfur and at lower temperature heavy compounds may not evaporate. Set the syringe drive rate desired, normally 6 µL/min, maximum with 200 mL/min hydrogen flow. Drive rate may be increased for increased sensitivity up to the point at which carbon is formed. (Hydrogen flow at 500 mL/min allows injection of 15 µL/min; however, dibenzothiophene conversion will be low.)

## 8. Calibration Standard

8.1 Prepare a reference standard-solution or solutions of strength near that expected in the unknown. Measurements can be made by weight or by volume for carrier liquid.

8.2 Units of sulfur in milligrams per litre of sample are preferred as this is independent of the density of the carrier liquid. The following equation is used to calculate the volume of solvent required to dissolve a precise weight of sulfur compound, of known composition and purity to prepare a liquid standard:

$$z = \left( \frac{b}{c} \times d \times e \times 10^6 \right) / (a) \text{ or alternatively:} \quad (1)$$

$$a = \left( \frac{b}{c} \times d \times e \times 10^6 \right) / (z) \quad (2)$$

where:

- a* = desired concentration of sulfur, mg/L, of the standard solution of *z* millilitre of volume,
- b* = molecular weight of sulfur: 32.06,
- c* = molecular weight of the sulfur compound to be used to prepare the standard.
- d* = mass of the sulfur compound used to prepare the standard, g,
- e* = purity of sulfur compound expressed as a decimal, and
- z* = millilitres of standard solution required to give the desired concentration *a*.

*Example:*

Calculate the volume of sulfur free *isooctane* with volume of sulfur compound necessary to dissolve 0.5013 g of 98 % by weight di-*n*-butyl sulfide to obtain a standard containing 1000 mg/L of sulfur in a solution.

$$a = 1000 \text{ mg/L}$$

$$b = 32.06$$

$$c = 146.29 \text{ di-}n\text{-butyl sulfide } (\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2)_2\text{S}$$

<sup>6</sup> Available from Phillips Petroleum Co., P.O. Drawer O, Borger, TX 79071.

$$d = 0.5013 \text{ g}$$

$$e = 0.98$$

$$z = \frac{[32.06 / 146.29] \times 0.5013 \times 0.98 \times 10^6}{1000} = 107.66 \text{ mL}$$

Isooctane is added to bring the solution to a total volume of 107.66 mL. When results are to be reported in mass of parts per million mg/kg, the conversion from milligrams per litre should be done as the last step in the calculations.

8.3 To prepare a sulfur standard with a sulfur concentration of 1000 mg/L, as previously described, obtain a clean 125 mL glass container, a 100 mL flask, and a 20 mL graduated glass pipet. Rinse each thoroughly with 2,2,4-trimethyl pentane (isooctane). (**Warning**—Extremely flammable.) Pour approximately 90 mL of isooctane into the 100 mL flask. Weigh approximately 0.5 g of di-*n*-butyl sulfide directly into the flask and record the mass added, to  $\pm 50 \mu\text{g}$ . Add additional isooctane to the flask to 100 mL. Transfer the mixture to the 125 mL container and add isooctane equal to the difference of  $z$  minus 100 mL. Keep containers closed as much as possible. Do not open containers of pure sulfur compound in the vicinity of sulfur free stocks or low-level standards. Evaporation from containers of pure sulfur compounds can contaminate other nearby liquids. This is particularly troublesome when working below 1 mg/L near volatile sulfur compounds. Volumetrically dilute stock to prepare low-level standards.

## 9. Calibration and Standardization

9.1 With hydrogen flow at 200 mL/min, advance new tape and note baseline. Adjust the offset up scale about 5 % to be clear of the recorder stops. Record the stable reading average value as the zero sulfur reference and record as  $R_b$  in 11.1. There will be essentially no difference in reading with or without hydrogen flow and with or without blank injection, if blank and hydrogen have no sulfur.

9.2 Advance the tape and inject a reference material with a sulfur concentration near that expected in the unknown. After about 4 min injection time, adjust the recorder span for approximately 90 % of scale. Record the average reading as  $R_{\text{std}}$  in 10.1.

## 10. Procedure

10.1 Advance the tape and inject the unknown sample. After a stable reading is obtained record this average value as  $R_s$  in 11.1.

10.2 Proceed with additional samples. Every 2 h, or as needed, verify blank and span values.

10.3 To measure sulfur below 1 mg/L, inject the sample at the maximum rate, normally 6  $\mu\text{L}/\text{min}$ , that does not form carbon or gums to obtain the best signal to noise ratio. Samples above 1 mg/L require proportionally lower injection rates or span adjustment. A sharp fall in response at high sulfur levels indicates color saturation of the tape, which can be prevented by slower injection rates.

## 11. Calculations and Report

11.1 Calculate the concentration of the sulfur in the sample as follows:

$$S, \text{ mg/L} = C_{\text{std}} \times (R_s - R_b) / (R_{\text{std}} - R_b) \quad (3)$$

where:

- $C_{\text{std}}$  = concentration of sulfur in the standard, mg/L,
- $R_b$  = response for blank run using no sample or for solvent known to be sulfur-free,
- $R_s$  = response for unknown sample, and
- $R_{\text{std}}$  = response for standard reference material.

11.2 Report mass of parts per million of sulfur as follows:

$$S, \text{ ppm} = (\text{mg/L}) / (\text{density}) = \text{mg/kg} \quad (4)$$

since density of sample is in grams per cubic centimetres.

## 12. Quality Assurance/Quality Control (QA/QC)

12.1 Confirm the performance of the instrument and the test procedure by analyzing a control (QC) sample.

12.1.1 When QA/QC protocols are already established in the testing facility, these may be used when they confirm the reliability of test result.

12.1.2 When there is no QA/QC protocol established in the testing facility, Appendix X1 may be used as the QA/QC system.

12.2 Users of this test method are advised that in contractual agreements, one or more of the contracting parties can and may make Appendix X1 a mandatory practice.

## 13. Precision and Bias

13.1 The precision of this test method as obtained by statistical analysis of interlaboratory test results is as follows:

13.1.1 *Repeatability*—The difference between successive 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 this test method, exceed the following values only in one case in twenty (see Table 1).

$$\text{Repeatability} = 0.16 \sqrt{X} \quad (5)$$

where

$X$  = average value of two results, mg/kg.

13.1.2 *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, exceed the following values only in one case in twenty (see Table 1).

$$\text{Reproducibility} = 0.26 \sqrt{X} \quad (6)$$

NOTE 2—One laboratory conducted a statistical evaluation by analyzing the same sample (with a nominal sulfur concentration of 0.2 ppm), using

TABLE 1 Repeatability and Reproducibility

Average value mg/kg of two results ( $x$ )	Repeatability, mg/kg	Reproducibility, mg/kg
0.02	0.02	0.04
0.10	0.05	0.08
0.50	0.11	0.18
1.00	0.16	0.26
2.50	0.25	0.41
10.00	0.50	0.82

multiple technicians and the same instrumentation, with and without the humidifier gas wash bottle installed, and determined that results were statistically equivalent for both precision and accuracy at the 95 % confidence interval. Results of the statistical evaluation are available from ASTM Headquarters.<sup>7</sup>

<sup>7</sup> Supporting data have been filed at ASTM International Headquarters and may be obtained by requesting Research Report RR:D02-1405.

13.2 *Bias*—The bias of this test method cannot be determined since an appropriate standard reference material containing trace sulfur levels in petroleum products is not available.

## 14. Keywords

14.1 rateometric colorimetry; sulfur

## APPENDIX

### (Nonmandatory Information)

#### X1. GENERIC QUALITY CONTROL STATEMENT FOR D02 TEST METHODS

X1.1 Confirm the performance of the instrument and the test procedure by analyzing a quality control (QC) sample.

X1.2 Prior to monitoring the measurement process, the user of this test method needs to determine the average value and control limits of the QC sample (see Practice **D6299** and MNL7<sup>8</sup>).

X1.3 Record the QC results and analyze by control charts or other statistically equivalent techniques to ascertain the statistical control status of the total testing process (see Practice **D6299** and MNL7<sup>8</sup>). Any out-of-control data should trigger investigation for root cause(s). The results of this investigation may, but not necessarily, result in instrument recalibration.

X1.4 In the absence of explicit requirements given in this test method, the frequency of QC testing is dependent on the

criticality of the quality being measured, the demonstrated stability of the testing process, and customer requirements. Generally, a QC sample should be analyzed on each day of testing routine samples. The QC frequency should be increased when a large number of samples are routinely analyzed. However, when it is demonstrated that the testing is under statistical control, the QC testing frequency may be reduced. The QC sample precision should be periodically checked against the ASTM test method precision to ensure data quality.

X1.5 It is recommended that, when possible, the type of QC sample that is regularly tested be representative of the samples routinely analyzed. An ample supply of QC sample material should be available for the intended period of use and must be homogeneous and stable under the anticipated storage conditions.

X1.6 See Practice **D6299** and MNL7<sup>8</sup> for further guidance on QC and control charting techniques.

<sup>8</sup> ASTM MNL7, *Manual of Presentation of Data Control Chart Analysis*, 6th ed., Section 3: Control Chart of Individuals, ASTM International, W. Conshohocken, PA.

## SUMMARY OF CHANGES

Subcommittee D02.03 has identified the location of selected changes to this standard since the last issue (D4045 – 04 (2010)) that may impact the use of this standard. (Approved April 1, 2015.)

(1) Updated footnotes 3 and 4.

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