



Standard Test Method for Trace Metals in Gas Turbine Fuels by Atomic Absorption and Flame Emission Spectroscopy¹

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1. Scope

1.1 This test method covers the determination of sodium, lead, calcium, and vanadium in Specification **D2880** Grade Nos. 1-GT and 2-GT fuels in the range from 0.1 to 2.0 mg/L. This test method is intended for the determination of oil-soluble metals and not waterborne contaminants in oil-water mixtures.

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.*

2. Referenced Documents

2.1 *ASTM Standards:*²

D2880 Specification for Gas Turbine Fuel Oils

D4057 Practice for Manual Sampling of Petroleum and Petroleum Products

3. Summary of Test Method

3.1 The samples are prepared to conform with the requirements of the method of standard additions, which is selected to obviate problems encountered with the direct analysis of typical gas turbine fuels that exhibit significant variations in physical properties. Different, but known, amounts of analyte are added to two portions of sample. These, together with the unaltered sample, are burned in the flame of an atomic absorption instrument that measures light absorption by the atomized metals. The analysis of the sample portions with

added analyte provides the calibration information necessary to calculate the analyte content of the unaltered sample.

3.2 Lead is determined by atomic absorption in a premixed air-acetylene flame, and sodium is determined by atomic absorption or atomic emission in a premixed air-acetylene flame. Calcium and vanadium are determined by atomic absorption or atomic emission in a premixed nitrous oxide-acetylene flame.

3.3 Most experience with this test method has been in the atomic absorption mode, although flame emission has been used successfully. Details in the subsequent sections are written for the atomic absorption mode. If the flame emission mode is used, minor details in the subsequent sections must be altered to conform to standard practice for flame emission spectroscopy. The precision statement applies only to the atomic absorption mode.

NOTE 1—Some GT fuel users may wish to determine potassium in addition to other metals included in this method. Potassium can be determined in a manner similar to that for sodium using a potassium hollow cathode lamp, (unless flame emission mode is used) a wavelength of 766.4 nm, and an appropriate organo-potassium standard. Precision data for potassium have not been determined.

4. Significance and Use

4.1 Knowledge of the presence of trace metals in gas turbine fuels enables the user to predict performance and, when necessary, to take appropriate action to prevent corrosion.

5. Apparatus

5.1 *Atomic Absorption Spectrophotometer*, capable of measuring radiation over the wavelength range from 280 to 600 nm. The instrument must be capable of measuring low-level signals (approximately 1 % absorption or 0.004 absorbance unit per mg/L vanadium). The instrument should also be equipped as follows.

5.1.1 *Burner*, with variable nebulizer and auxiliary oxidant supply to reduce non-atomic absorption from unburned hydrocarbons which cause interferences.

5.1.1.1 *Burner Head*, capable of supporting a nitrous oxide-acetylene flame.

5.1.1.2 *Burner Head*, single- or multiple-slot, capable of supporting an air-acetylene flame.

¹ This test method is under the jurisdiction of ASTM Committee **D02** on Petroleum Products and Lubricants and is the direct responsibility of Subcommittee **D02.03** 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.

5.1.2 *Electronic Detection System*, capable of reading to the nearest 0.1 % absorption or 0.0004 absorbance.

5.1.2.1 The text describes the measurement of absorption signals that is, either percent absorption or absorbance. For instruments reading in percent absorption, absorption signals of 0.1 % absorption must be measurable. For instruments reading in absorbance, signals of 0.0004 absorbance must be measurable.

5.1.3 *Hollow Cathode Lamp Power Supply*, regulated to minimize drift.

5.1.4 *Monochromator*, capable of resolving the 318.34–318.40-nm vanadium doublet from the 318.54-nm vanadium line.

5.1.5 *Hollow Cathode Lamps*, one each for calcium, sodium, lead, and vanadium.

NOTE 2—Electrodeless-discharge lamps can be an acceptable alternative, but the precision of this method was determined with hollow cathode lamps.

5.1.6 When the instrument has flame-emission capability, the emission technique can be used for the analyses of sodium, calcium, and vanadium.

5.2 *Volumetric Flasks*, 25-mL.

5.3 *Glass Vials*, 40-mL, screw-cap type, polyethylene-lined caps.

5.4 *Syringe*, 100- μ L, Hamilton type or equivalent.

6. Reagents

6.1 *Purity of Reagents*—Reagent grade chemicals shall be used in 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.³ Other grades may be used, provided it is ascertained that the reagent is of sufficiently high purity to permit its use without lessening the accuracy of the determination.

6.2 *1,2,3,4-tetrahydronaphthalene*,⁴ practical grade, analyte-sterile.

NOTE 3—Analyte-sterile 1,2,3,4-tetrahydronaphthalene can be prepared by extracting a portion of tetralin with an equal amount of hydrochloric acid in a covered screw-cap vial. Heat the vial on a steam bath for 1 h and shake the vial for 1 h. If the acid extracted 1,2,3,4-tetrahydronaphthalene and unextracted 1,2,3,4-tetrahydronaphthalene give indistinguishable absorption signals for each of the analytes under optimal experimental conditions, the unextracted 1,2,3,4-tetrahydronaphthalene can be used throughout this method.

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 *Analar Standards for Laboratory Chemicals*, BDH Ltd., Poole, Dorset, U.K., and the *United States Pharmacopeia and National Formulary*, U.S. Pharmaceutical Convention, Inc. (USPC), Rockville, MD.

⁴ Tetralin (1,2,3,4-tetrahydronaphthalene), manufactured by E. I. duPont de Nemours and Co., has been found satisfactory. 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 committee¹, which you may attend.

TABLE 1 Experimental Conditions

Element	Mode	Wavelength, nm	Fuel	Oxidant
Na	Absorption	589.6	C ₂ H ₂	air
Na	Emission	589.6	C ₂ H ₂	air
Pb	Absorption	283.3	C ₂ H ₂	air
Ca	Absorption	422.7	C ₂ H ₂	N ₂ O
Ca	Emission	422.7	C ₂ H ₂	N ₂ O
V	Absorption	318.34–318.40	C ₂ H ₂	N ₂ O
V	Emission	437.9	C ₂ H ₂	N ₂ O

6.3 *Organometallic Standards*—Oil-soluble salts of sodium, lead, calcium, and vanadium of known concentration.⁵

6.4 *Mixed Standard*—Prepare a mixed standard containing 250 mg/L each of sodium, lead, calcium, and vanadium by dissolving the appropriate amounts of organometallic standards in 1,2,3,4-tetrahydronaphthalene and making the required dilutions. Prepare fresh daily, as needed.

7. Sampling

7.1 Samples shall be taken in accordance with the instructions in Practice [D4057](#).

8. Procedure

8.1 Fill two clean 25-mL volumetric flasks to the line with sample. With the microlitre syringe add 50 μ L of mixed standard to one flask and 100 μ L to the other. Touch the needle of the syringe to the inner wall of the flask to ensure quantitative transfer of the standard. Invert and mix the contents. (The two flasks are now spiked with 0.5 mg/L and 1.0 mg/L of sodium, lead, calcium, and vanadium). Alternatively, weigh 25.0 g of sample into each of two clean disposable glass vials and add the standard in the same manner. (The two vials are now spiked with 0.5 mg/kg and 1.0 mg/kg of sodium, lead, calcium, and vanadium.)

8.2 Prepare a third spiked sample by adding approximately 1 mL of the mixed standard to approximately 25 mL of sample. This solution serves only to aid in establishing satisfactory operating conditions for the atomic absorption instrument.

8.3 Establish the atomic absorption instrument operating conditions, which are recommended by the manufacturer, and consider the following special points. Select the mode, flame gases, and spectral lines from the information presented in [Table 1](#).

8.4 *Analysis*:

8.4.1 With the atomic absorption instrument in operation for monitoring lead absorption and with 1,2,3,4-tetrahydronaphthalene nebulizing, zero the instrument. Aspirate into the flame the third spiked sample described in [8.2](#) and note the net lead absorption signal. Optimize experimental conditions by adjusting the burner position (relative to the hollow cathode beam), the flow rates of the fuel and oxidant gases, and the sample aspiration rate until the net lead

⁵ Conostan standards, available from Conostan Division, Continental Oil Co., Ponca City, OK 74601, were used in determining the precision quoted in this method. Other standards are available from the Office of Standard Reference Materials, Room B314, Chemistry Bldg., National Institute of Standards and Technology, Washington, DC 20234, and from Angstrom, Inc., P. O. Box 252, Belleville, MI 48111, but the precision statement may or may not apply to results obtained with these standards.

absorption signal maximizes. Re-zero the instrument with 1,2,3,4-tetrahydronaphthalene nebulizing, and consecutively introduce into the flame the unaltered sample and the two spiked samples, with 1,2,3,4-tetrahydronaphthalene nebulizing between each sample. Record the absorption signal of each sample and of each blank between samples.

8.4.1.1 At optimal experimental conditions, the analyte concentration that accounts for 1% absorption should be approximately 1 mg/L. The percent absorption to concentration ratio must be near unity in order to achieve the lower limits of detection that are required. This note does not apply to flame emission measurements.

8.4.2 With the atomic absorption instrument in operation for monitoring vanadium absorption and with 1,2,3,4-tetrahydronaphthalene nebulizing, zero the instrument. Introduce into the flame the third spiked sample described in 8.2. Record the net vanadium absorption signal. Optimize the experimental conditions as in 8.4.1. Re-zero the instrument with 1,2,3,4-tetrahydronaphthalene nebulizing and consecutively introduce into the flame the unaltered sample and the two spiked samples. Record the absorption signal of each sample and of each 1,2,3,4-tetrahydronaphthalene blank between samples.

8.4.3 With the atomic absorption instrument in operation for monitoring sodium absorption, zero the instrument with 1,2,3,4-tetrahydronaphthalene nebulizing, and consecutively introduce into the flame the unaltered sample and the two spiked samples. Record the absorption signal of each sample and of each 1,2,3,4-tetrahydronaphthalene blank between samples.

NOTE 4—For determining sodium and calcium, the maximization of absorption or emission signals is not critical.

8.4.4 With the atomic absorption instrument in operation for monitoring calcium absorption, zero the instrument with 1,2,3,4-tetrahydronaphthalene nebulizing, and consecutively introduce into the flame the unaltered sample and the two spiked samples. Record the absorption signal of each sample and of each 1,2,3,4-tetrahydronaphthalene blank between samples.

9. Calculation

9.1 For each absorption signal, calculate the net absorption signal as follows:

$$a = A - [(b_1 + b_2)/2] \quad (1)$$

where:

- a = net absorption signal,
- A = observed absorption signal,
- b_1 = blank signal before the sample, and
- b_2 = blank signal after the sample.

NOTE 5—Blank corrections are usually small. If a large drift in blank signals is observed during a series of measurements, some experimental parameter is out of control. The cause of the variation should be corrected, and the measurements repeated.

9.2 For each element in turn, calculate sensitivities as follows:

$$S_{0.5} = 2(a_1 - a_2) \quad (2)$$

$$S_{1.0} = (a_3 - a_2) \quad (3)$$

$$S = (S_{0.5} + S_{1.0})/2 \quad (4)$$

where:

- $S_{0.5}$ = sensitivity for the 0.5-mg/L spiked sample,
- $S_{1.0}$ = sensitivity for the 1.0-mg/L spiked sample,
- S = average sensitivity,
- a_1 = net absorption signal for the 0.5-mg/L spiked sample,
- a_2 = net absorption signal for the unaltered sample, and
- a_3 = net absorption signal for the 1.0-mg/L spiked sample.

9.3 Calculate the ratio of sensitivities, R , as follows:

$$R = S_{1.0}/S_{0.5} \quad (5)$$

If R falls outside the range of $0.90 \leq R \leq 1.10$, the data are nonlinear. Readjust experimental conditions and repeat the analysis.

9.4 Calculate the concentration of each element as follows:

$$\text{mg/L} = a_2/S \quad (6)$$

$$\text{mg/kg} = a_2/(S \times d) \quad (7)$$

where d = density of the sample in g/ml.

10. Quality Control Checks (QA/QC)

10.1 When QA/QC protocols are already established in the testing facility, these may be used to confirm the reliability of the test method.

10.2 Since reference materials for the various matrices are not available and the test method utilizes the method of standard additions, the only practical quality control check is to verify the accuracy of the organometallic standard in 6.3. A source independent of this standard should be acquired and a sample carried through the analysis using the second-source standard as the known addition. Results of this test can be compared to the original sample result using the original standard material to verify accuracy.

11. Report

11.1 Report results for each element to the nearest 0.1 mg/L.

12. Precision and Bias

12.1 The precision of this test method as obtained by statistical examination of interlaboratory test results is as follows:

12.1.1 *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 the test method exceed the following values only in one case in twenty:

Element	Repeatability
V	0.452 (concentration) ^{1/2}
Pb	0.244 (concentration) ^{1/2}
Ca	0.202 (concentration) ^{1/2}
Na	0.232 (concentration) ^{1/2}

12.1.2 *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, in the normal and correct operation of the test method exceed the following values only in one case in twenty:

Element	Reproducibility
V	0.616 (concentration) ^{1/2}
Pb	0.900 (concentration) ^{1/2}
Ca	0.402 (concentration) ^{1/2}
Na	0.738 (concentration) ^{1/2}

NOTE 6—The above repeatability and reproducibility are based on a concentration range of 0.1 to 0.5 mg/L and apply only to the atomic absorption mode.

12.2 *Bias*—The bias of this test method cannot be determined since appropriate reference materials containing known levels of vanadium, lead calcium and sodium are not available.

13. Keywords

13.1 atomic absorption; calcium; flame emission; gas turbine fuels; lead; sodium; vanadium

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