



Designation: ~~D 3373—93 (Reapproved 1998)~~ **3373 – 03**

Standard Test Method for Vanadium in Water¹

This standard is issued under the fixed designation D 3373; 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 Department of Defense.

1. Scope*

1.1 This test method covers the determination of dissolved and total recoverable vanadium in most waters and wastewater by graphite furnace atomic absorption spectrophotometry.

1.2 The optimum range of this test method is 10 to 200 $\mu\text{g/L}$ of vanadium based on a 20- μL sample size. Concentrations higher than 200 $\mu\text{g/L}$ may be determined by proper dilution of sample. A detection level as low as 4 $\mu\text{g/L}$ of vanadium has been reported.

1.3 This test method has been used successfully with reagent water, lake water, tap water, river water, condensate from a medium Btu coal gasification process, and well water. It is the user's responsibility to ensure the validity of this test method for waters of untested matrices.

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 or regulatory limitations prior to use.*

1.5 Former Test Method A (Catalytic) was discontinued. Refer to Appendix X1 for historical information.

2. Referenced Documents

2.1 *ASTM Standards:*

¹ This test method is under the jurisdiction of ASTM Committee D-19 on Water and is the direct responsibility of Subcommittee D19.05 on Inorganic Constituents in Water. Current edition approved ~~Sept 15, 1993~~, June 10, 2003. Published ~~November 1993~~, July 2003. Originally published as ~~D 3373—75~~, approved in 1975. Last previous edition approved in 1998 as D 3373 – 93 (19988).

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



- D 1066 Practice for Sampling Steam²
- D 1129 Terminology Relating to Water²
- D 1192 Specification for Equipment for Sampling Water and Steam in Closed Conduits²
- D 1193 Specification for Reagent Water²
- D 2777 Practice for Determination of Precision and Bias of Applicable Methods of Committee D-19 on Water²
- D 3370 Practices for Sampling Water from Closed Conduits²
- D 3919 Practice for Measuring Trace Elements in Water by Graphite Furnace Atomic Absorption Spectrophotometry²
- D 4841 Practice for Estimation of Holding Time for Water Samples Containing Organic and Inorganic Constituents²
- D 5810 Guide for Spiking into Aqueous Samples²
- D 5847 Practice for Writing Quality Control Specifications for Standard Test Methods for Water Analysis³

3. Terminology

3.1 *Definitions*—For definitions of terms used in this test method refer to Terminology D 1129.

3.2 *Definitions of Terms Specific to This Standard:*

3.2.1 *total recoverable vanadium*—dissolved vanadium plus that solubilized by acid digestion of particulates and organics in the sample.

4. Summary of Test Method

4.1 Vanadium is determined by an atomic absorption spectrophotometer utilizing a graphite furnace for sample atomization.

4.2 A sample volume of several microlitres, depending upon the concentration of the analyte, is transferred to a graphite tube housed within an electrical furnace and the system is heated in an inert or reducing atmosphere. The sample is evaporated to dryness, charred (pyrolyzed or ashed) and atomized.

4.3 Ground state atoms, produced in atomization, absorb the light of their spectrum emanating from a source and passing through the sample vapor. The amount of light absorbed is proportional to the concentration of the analyte in the sample.

4.4 Since the graphite furnace uses the sample much more efficiently than does flame atomization, the detection of low concentrations of the elements of interest in small volumes is possible.

4.5 Finally, the absorption signal produced during atomization is recorded and compared to those of standards, taken through the same process, by means of an analytical curve.

4.6 A general guide for graphite furnace applications is given in Practice D 3919.

4.7 Dissolved vanadium is determined after filtration and addition of HNO₃ to a pH of ≤ 2 .

4.8 Total recoverable vanadium is determined following acid digestion and filtration.

5. Significance and Use

5.1 Vanadium can be found in waste that result from chemical cleaning of components in which the metal is alloyed.

5.2 National Pollutant Discharge Elimination Systems permits or other standards, or both, require monitoring pollutants in waste discharged onto the water shed of, or into, navigable waters, and those disposed of in such a manner that eventual contamination of underground water could result.

5.3 This test method affords an accurate and sensitive means of determining compliance or noncompliance, or both, with those permits.

6. Interferences

6.1 For a complete discussion on general interferences with furnace procedures, the analyst is referred to Practice D 3919.

7. Apparatus

7.1 *Atomic Absorption Spectrophotometer*, for use at 318.4 nm with background correction.

NOTE 1—The manufacturer's instructions should be followed for all instrumental parameters.

7.2 *Vanadium Hollow Cathode Lamp*.

7.3 *Graphite Furnace* capable of reaching temperature sufficient to atomize the element of interest.

7.4 *Graphite Tubes*, pyrolytically coated, compatible with furnace device are recommended.

7.5 *Pipets*, microlitre with disposable tips, ranging in size from 1 to 100 μ L.

7.6 *Strip Chart Recorder*—A recorder is strongly recommended. The tracings can be used as permanent records and are of diagnostic value in the identification and analysis of instrument problems (such as drift, incomplete atomization, change in

² *Annual Book of ASTM Standards*, Vol 11.01.

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; ASTM Standards, Vol 11.02.

sensitivity, etc.). A fast response recorder (0.2 s or less for full-scale deflection) is recommended to ensure accuracy. Electronic peak-measuring devices have also been found useful.

7.7 *Automatic Sampling Accessory* is recommended, if available.

8. Reagents and Materials

8.1 *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 Committee on Analytical Reagents of the American Chemical Society, where such specifications are available.⁴ 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.

8.2 *Purity of Water*— Unless otherwise indicated, references to water shall be understood to mean reagent water conforming to Specification D 1193, Type I. Other reagent water types may be used provided it is first ascertained that the water is of sufficiently high purity to permit its use without adversely affecting the precision and bias of the test method.

8.3 *Nitric Acid* (sp gr 1.42)—Concentrated nitric acid (HNO₃).

8.4 *Vanadium Solution, Stock* (1.0 mL = 100 µg vanadium). Dissolve 0.2296 g of ammonium metavanadate (NH₄VO₃) in water and dilute to 1000 mL.

8.5 *Vanadium Solution, Intermediate* (1.0 mL = 1.0 µg vanadium)— Dilute 5 mL of vanadium stock solution to 500 mL with water.

8.6 *Vanadium Solution, Standard* (1.0 mL = 0.100 µg vanadium)—Dilute 10.0 mL of vanadium intermediate solution to 100 mL with water.

8.7 *Nitrogen Gas*—standard, welders grade, commercially available.

NOTE 2—It is probable that argon or some other inert gas can be used in place of nitrogen. It is the analyst's responsibility to verify suitability.

9. Sampling

9.1 Collect the sample in accordance with Practice D 1066, Specification D 1192, and Practices D 3370, as applicable. The holding time for the sample may be calculated in accordance with Practice D 4841.

9.2 Preserve samples with nitric acid (HNO₃, sp gr 1.42) to a pH of 2 or less immediately at the time of collection, normally about 2 mL/L. If only dissolved vanadium is to be determined, filter the sample through a 0.45-µm (No. 325) membrane filter before acidification.

10. Standardization

10.1 Initially, set the instrument in accordance with the manufacturer's specifications. Follow the general instructions as provided in Practice D 3919.

11. Procedure

11.1 Clean all glassware to be used for preparation of standard solutions or in the digestion step, or both, by rinsing first with HNO₃ (1 + 1) and then with water. Alternatively, soaking the glassware overnight in HNO₃ (1 + 1) is useful for low levels.

11.2 Measure 100.0 mL of each standard and well-mixed sample into a 125-mL beaker or flask. For total recoverable vanadium, add 5 mL of HNO₃ (sp gr 1.42) to each sample and proceed as directed in 11.4 through 11.6.

11.3 If only dissolved vanadium is to be determined, proceed to 11.6.

11.4 Heat the samples at 95°C on a steam bath or hotplate in a well-ventilated fume hood until the volume has been reduced to 15 to 20 mL making certain that the samples do not boil.

NOTE 3—When analyzing samples of brines or samples containing appreciable amounts of suspended matter or dissolved solids, the amount of reduction in volume is left to the discretion of the analyst.

11.5 Cool and filter the sample through a suitable filter (such as fine-textured, acid washed, ashless paper) into a 100- mL volumetric flask. Wash the filter paper 2 or 3 times with water and bring to volume. See Note 4. The acid concentration at this point should be 5 % HNO₃.

NOTE 4—If suspended material is not present, this filtration may be omitted, however, digested sample must still be diluted to 100 mL.

11.6 Inject a measured aliquot of sample into the furnace device following the directions as provided by the particular instrument manufacturer. Refer to Practice D 3919.

12. Calculation

12.1 Determine the concentration of vanadium in each sample in accordance with the Samples Analysis Procedure Section of Practice D 3919.

⁴Supporting data are available and

⁴ *Reagent Chemicals, American Chemical Society Specifications*, American Chemical Society, Washington, DC. For suggestions on loan from ASTM Headquarters, Request RR- D19-1114; 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.



13. Precision and Bias ⁵

13.1 Based on data from twelve participating laboratories, the overall precision of this test method and recoveries from a series of standards containing known amounts of vanadium, were as given in Table 1.

13.2 Because of the large number of metals analyzed in this study the requirement for replicate tests have been waived; therefore, single-operator precision is not available.

13.3 Each participating laboratory evaluated this test method in reagent water. Individual laboratories selected one water of choice as an additional matrix of interest in which to test recovery. Listed among those choices were: lake water, tap water, river water, well water, and condensate from a medium Btu coal gasification process.

13.4 These data may not apply to waters of other matrices.

13.5 This section on precision and bias conforms to Practice D 2777 – 77 which was in place at the time of collaborative testing. Under the allowances made in 1.4 of D 2777 – 98, these precision and bias data do meet existing requirements of interlaboratory studies of Committee D19 test methods.

14. Quality Control

14.1 In order to be certain that analytical values obtained using these test methods are valid and accurate within the confidence limits of the test, the following QC procedures must be followed when analyzing vanadium.

14.2 Calibration and Calibration Verification

14.2.1 Analyze at least three working standards containing concentrations of vanadium that bracket the expected sample concentration, prior to analysis of samples, to calibrate the instrument. The calibration correlation coefficient shall be equal to or greater than 0.990. In addition to the initial calibration blank, a calibration blank shall be analyzed at the end of the batch run to ensure contamination was not a problem during the batch analysis.

14.2.2 Verify instrument calibration after standardization by analyzing a standard at the concentration of one of the calibration standards. The concentration of a mid-range standard should fall within $\pm 15\%$ of the known concentration.

14.2.3 If calibration cannot be verified, recalibrate the instrument.

14.3 Initial Demonstration of Laboratory Capability

14.3.1 If a laboratory has not performed the test before, or if there has been a major change in the measurement system, for example, new analyst, new instrument, and so forth, a precision and bias study must be performed to demonstrate laboratory capability.

14.3.2 Analyze seven replicates of a standard solution prepared from an Independent Reference Material containing a midrange concentration of vanadium. The matrix and chemistry of the solution should be equivalent to the solution used in the collaborative study. Each replicate must be taken through the complete analytical test method including any sample preservation and pretreatment steps. The replicates may be interspersed with samples.

14.3.3 Calculate the mean and standard deviation of the seven values and compare to the acceptable ranges of bias in Table 1. This study should be repeated until the recoveries are within the limits given in Table 1. If a concentration other than the recommended concentration is used, refer to Practice D5847 for information on applying the F test and t test in evaluating the acceptability of the mean and standard deviation.

14.4 Laboratory Control Sample (LCS)

14.4.1 To ensure that the test method is in control, analyze a LCS containing a known concentration of vanadium with each batch or ten samples. If large numbers of samples are analyzed in the batch, analyze the LCS after every ten samples. The laboratory control samples for a large batch should cover the analytical range when possible. The LCS must be taken through all of the steps of the analytical method including sample preservation and pretreatment. The result obtained for a mid-range LCS shall fall within $\pm 15\%$ of the known concentration.

14.4.2 If the result is not within these limits, analysis of samples is halted until the problem is corrected, and either all the samples in the batch must be reanalyzed, or the results must be qualified with an indication that they do not fall within the performance criteria of the test method.

⁵ Supporting data have been filed at ASTM International Headquarters and may be obtained by requesting Research Report RR: D19-1114.

TABLE 1 Determination of Precision and Bias for Vanadium

Amount Added, $\mu\text{g V/L}^A$	Amount Found, $\mu\text{g V/L}^A$	S_T , $\mu\text{g/L}$	Bias, $\mu\text{g/L}$	% Bias	Statistically Significant
<i>Reagent Water Type II</i>					
25	24	2.6	-1.0	4.0	no
70	73.4	4.8	+ 3.4	4.9	no
240	232.7	25.5	-7.3	3.0	no
<i>Water of Choice</i>					
25	25.9	3.6	+ 0.9	3.6	no
70	70	7.6	0	0	no
240	247.3	27.7	+ 7.3	3.0	no

^A Editorially corrected.

14.5 Method Blank

14.5.1 Analyze a reagent water test blank with each batch. The concentration of vanadium found in the blank should be less than 0.5 times the lowest calibration standard. If the concentration of vanadium is found above this level, analysis of samples is halted until the contamination is eliminated, and a blank shows no contamination at or above this level, or the results must be qualified with an indication that they do not fall within the performance criteria of the test method.

14.6 Matrix Spike (MS)

14.6.1 To check for interferences in the specific matrix being tested, perform a MS on at least one sample from each batch by spiking an aliquot of the sample with a known concentration of vanadium and taking it through the analytical method.

14.6.2 The spike concentration plus the background concentration of vanadium must not exceed the high calibration standard. The spike must produce a concentration in the spiked sample that is 2 to 5 times the analyte concentration in the unspiked sample, or 10 to 50 times the detection limit of the test method, whichever is greater.

14.6.3 Calculate the percent recovery of the spike (P) using the following formula:

$$P = 100 [A(V_s + V) - BV_s] / CV \quad (1)$$

where:

- A ≡ analyte concentration (mg/L) in spiked sample,
- B ≡ analyte concentration (mg/L) in unspiked sample,
- C ≡ concentration (mg/L) of analyte in spiking solution,
- V_s ≡ volume (mL) of sample used, and
- V ≡ volume (mL) added with spike.

14.6.4 The percent recovery of the spike shall fall within the limits, based on the analyte concentration, listed in Test Method D 5810, Table 1. If the percent recovery is not within these limits, a matrix interference may be present in the sample selected for spiking. Under these circumstances, one of the following remedies must be employed: the matrix interference must be removed, all samples in the batch must be analyzed by a test method not affected by the matrix interference, or the results must be qualified with an indication that they do not fall within the performance criteria of the test method.

NOTE 5—Acceptable spike recoveries are dependent on the concentration of the component of interest. See Test Method D 5810 for additional information.

14.7 Duplicate

14.7.1 To check the precision of sample analyses, analyze a sample in duplicate with each batch. If the concentration of the analyte is less than five times the detection limit for the analyte, a matrix spike duplicate (MSD) should be used.

14.7.2 Calculate the standard deviation of the duplicate values and compare to the precision in the collaborative study using an F test. Refer to 6.4.4 of Practice D 5847 for information on applying the F test.

14.7.3 If the result exceeds the precision limit, the batch must be reanalyzed or the results must be qualified with an indication that they do not fall within the performance criteria of the test method.

14.8 Independent Reference Material (IRM)

14.8.1 In order to verify the quantitative value produced by the test method, analyze an Independent Reference Material (IRM) submitted as a regular sample (if practical) to the laboratory at least once per quarter. The concentration of the IRM should be in the concentration mid-range for the method chosen. The value obtained must fall within the control limits established by the laboratory.

15. Keywords

14.5.1 atomic absorption; furnace; total recoverable vanadium; water



APPENDIX

(Nonmandatory Information)

X1. RATIONALE FOR DISCONTINUATION OF TEST METHODS

X1.1 Former Test Method A—Catalytic:

X1.1.1 This test method was discontinued in 1988. The test method may be found in its entirety in the 1988 *Annual Book of ASTM Standards*, Vol 11.01.

X1.1.2 This test method covers the determination of dissolved and total recoverable vanadium found in concentrations of 1.0 to 8.0 µg/L.

X1.1.3 Trace amounts of vanadium are determined by measuring the catalytic effect it exerts on the rate of oxidation of gallic acid by persulfate in acid solution. Under given conditions of reactant concentration, temperature, and reaction time, the extent of oxidation of gallic acid is proportional to the concentration of vanadium present. Depending on the amount of vanadium present, the reaction produces a yellow-to-red color. Vanadium is determined by measuring the absorbance of the sample at 415 nm and comparing it with standard solutions treated in an identical manner.

X1.1.4 This test method was discontinued because there were insufficient laboratories interested in participating in another collaborative study to obtain the necessary precision and bias as required by Practice D 2777.

SUMMARY OF CHANGES

Committee D19 has identified the location of selected changes to this standard since the last issue (D3373–93 (1998)) that may impact the use of this standard.

- (1) Section 8.2 was revised.
- (2) Section 13.5 was added.
- (3) The QC Section 14 was added to the test method.

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