



Designation: D7781 – 23

Standard Test Method for Nitrite-Nitrate in Water by Nitrate Reductase¹

This standard is issued under the fixed designation D7781; 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.

1. Scope

1.1 This test method is applicable to the determination of nitrate plus nitrite (as nitrogen) in drinking water, surface, saline, wastewater, and ground waters. The applicable range of this test method is from 0.05 to 5 mg/L of nitrogen. The range may be extended upward by dilution of an appropriate aliquot. The 40 CFR Part 136 Method Detection Limit (MDL) is 0.02 mg/L.

1.2 It is the user's responsibility to ensure the validity of this test method for waters of untested matrices. The quality control criteria in Section 17 for method blanks, laboratory control samples, matrix spikes and matrix duplicates must be met.

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

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, health, and environmental practices and determine the applicability of regulatory limitations prior to use.*

1.5 *This international standard was developed in accordance with internationally recognized principles on standardization established in the Decision on Principles for the Development of International Standards, Guides and Recommendations issued by the World Trade Organization Technical Barriers to Trade (TBT) Committee.*

2. Referenced Documents

2.1 *ASTM Standards:*²

D0992 Method of Test for Nitrate Ion in Water (Withdrawn 1983)³

D1129 Terminology Relating to Water

D1141 Practice for Preparation of Substitute Ocean Water

D1193 Specification for Reagent Water

D1254 Method of Test for Nitrite Ion in Water (Withdrawn 1980)³

D3867 Test Methods for Nitrite-Nitrate in Water

D5810 Guide for Spiking into Aqueous Samples

D6146 Guide for Monitoring Aqueous Nutrients in Watersheds

3. Terminology

3.1 *Definitions*—For definitions of terms used in these test methods, refer to Terminology D1129.

3.2 *Definitions of Terms Specific to This Standard:*

3.2.1 *bispecific NaR, n*—nitrate reductase that can use either NADH or NADPH as its electron donor (cofactor).

3.2.2 *discrete analyzer, n*—a programmable, computer-controlled instrument that automates wet-chemical analysis by using one or more robotic arms interfaced to high-precision volumetric dispensers to aspirate and dispense samples, standards, diluents, and reagents.

3.2.3 *Greiss reaction, n*—chemical formation of an azo dye by diazotization of nitrite ion with sulfanilamide and subsequent coupling with *N*-(1-naphthyl)ethylenediamine hydrochloride.

3.2.4 *NADH, n*—nicotinamide adenine dinucleotide, reduced form is a coenzyme found in all living cells.

3.2.5 *NADPH, n*—nicotinamide adenine dinucleotide phosphate, reduced form is a coenzyme found in all living cells; NADP⁺ is the oxidizing form and NADPH is the reducing form.

3.2.6 *nitrate reductase (NaR), n*—NADH:NaR (EC1.7.1.1 and CAS 9013-03-0) or bispecific NaR (EC 1.7.1.2 and CAS 9029-27-0) with 1 unit of enzyme activity defined as 1 micromol nitrite produced per minute at 30 °C, at pH 7 with NADH (refer to 3.2.4 and 10.2) as an electron donor.

4. Summary of Test Method

4.1 *Nitrite-Nitrate Nitrogen*—The sample is mixed with a buffered solution containing NAD(P)H: nitrate reductase (EC 1.7.1.3) and NADH or NADPH to reduce nitrate ion to nitrite ion. The combined nitrite-nitrate (expressed as mg/L NO₃+NO₂-N) is determined by diazotizing the total nitrite ion

¹ This test method is under the jurisdiction of ASTM Committee D19 on Water and is the direct responsibility of Subcommittee D19.05 on Inorganic Constituents in Water.

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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.

³ The last approved version of this historical standard is referenced on www.astm.org.

with sulfanilamide and coupling with *N*-(1-naphthyl) ethylenediamine dihydrochloride to form a highly colored azo dye that is measured spectrophotometrically at about 540 nm.

4.2 Nitrite Nitrogen—The nitrite ion (expressed as mg/L NO₂-N) originally present in the sample can be determined separately by carrying out the procedure and omitting the reduction step.

4.3 Nitrate Nitrogen—The nitrate ion as nitrogen can be calculated as the difference between the combined nitrate plus nitrite (NO₃+NO₂-N) and the nitrite (NO₂-N):

$$\text{mg/LNO}_3 - \text{N} = \text{mg/L}(\text{NO}_3 + \text{NO}_2 - \text{N}) - \text{mg/L}(\text{NO}_2 - \text{N}) \quad (1)$$

5. Significance and Use

5.1 This test method replaces Methods **D1254** (Nitrite) and **D992** (Nitrate). The nitrite test method (Method **D1254**) used a reagent that is considered to be a potential carcinogen. The nitrate test method (Method **D992**) has been shown to have relatively large errors when used in wastewaters and also has greater manipulative difficulties than the test method described herein.

5.2 This test method can be used in place of Test Methods **D3867** (Nitrite-Nitrate). Test Methods **D3867** uses cadmium for the reduction of nitrate to nitrite. Cadmium is considered a toxic metal. Also, the heterogeneous cadmium reductant creates greater difficulty than the reduction described in this test method.

6. Interferences

6.1 Turbid samples should be filtered prior to analysis to eliminate particulate interference.

6.2 Sample color that absorbs at wavelengths between 520 and 540 nm interferes with the absorbance measurements. When color is suspect, analyze a sample blank, omitting the *N*-(1-naphthyl)ethylenediamine dihydrochloride from the color reagent.

NOTE 1—The instrumentation described in this standard may automatically correct for some turbidity and sample color. See the instrument manual for further information.

6.3 Certain ions may cause interferences. See **Table 1**.

7. Purity of Reagents

7.1 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, when such specifications are available.⁴ Other grades may be used, provided it is first ascertained that the reagent is of sufficient high purity to permit its use without lessening the accuracy of the determination.

⁴ ACS Reagent Chemicals, Specifications and Procedures for Reagents and Standard-Grade Reference Materials, 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. Pharmacopeial Convention, Inc. (USPC), Rockville, MD.

TABLE 1 Determination of Nitrate in the Presence of Potential Interferences

Species	Concentration Added (mg/L)	Unspiked Sample Result (mg/L)	Spiked Sample Results (mg/L)	Spike Added (mg/L)	% Recovery
Cl ⁻	500	0.02	0.23	0.200	105
		0.17	2.54	2.50	95
F ⁻	500	0.01	0.22	0.200	105
Br ⁻	500	<0.01	0.21	0.200	100
		0.15	2.65	2.50	100
PO ₄ ⁻³	500	0.01	0.22	0.200	105
		0.14	2.54	2.50	96
SO ₄ ⁻²	500	<0.01	0.21	0.200	105
		0.14	2.53	2.50	96
Fe	500	0.17	2.60	2.50	97
	1.0	<0.01	0.21	0.200	105
		0.168	2.59	2.50	96
Zn	1.0	<0.01	0.22	0.200	110
		0.14	2.64	2.50	100
Al	1.0	<0.01	0.21	0.200	105
		0.14	2.53	2.50	96
BrO ₃ ⁻	1.0	<0.01	0.22	0.200	110
		0.17	2.64	2.50	99
ClO ₂ ⁻	1.0	0.01	0.22	0.200	110
		0.14	2.54	2.50	96
ClO ₃ ⁻	1.0	0.23	2.45	2.50	89
CHCl ₃	>Miscibility	<0.01	0.21	0.200	105

7.2 Purity of Water—Unless otherwise indicated, references to water shall be understood to mean reagent water conforming to Specification **D1193**, Type I or Type II. 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 bias and precision of these test methods.

8. Sampling and Sample Preservation

8.1 Collect the sample in accordance with Guide **D6146**.

8.2 When nitrite ion is to be determined separately, analyze within 48 hours after sampling. Even when sterile bottles are used, bacteria naturally present in the water may cause conversion of all or part of nitrite ion to other forms such as nitrate or ammonia. Ammonia and natural amines, which are frequently present in natural waters, may react with nitrites to form nitrogen. If samples are to be stored for 48 h or less, preserve the sample by refrigeration at 2–6 °C. If the sample must be stored for more than 48 h, preserve it by the addition of sulfuric acid to pH 2 in addition to refrigeration at 2–6 °C.

NOTE 2—Use sulfuric acid for preservation of nitrite-nitrate nitrogen only. Samples for nitrite must be analyzed within 48 hours.

NOTE 3—Sulfuric acid does not necessarily inhibit oxidation and mercury compounds should be avoided to prevent environmental pollution.

NOTE 4—Residual chlorine does not interfere, however, attempts to remove residual chlorine (such as addition of ascorbic acid) interfere by