



Designation: D7573 – 09 (Reapproved 2017)

Standard Test Method for Total Carbon and Organic Carbon in Water by High Temperature Catalytic Combustion and Infrared Detection¹

This standard is issued under the fixed designation D7573; 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 covers the determination of total carbon (TC), inorganic carbon (IC), total organic carbon (TOC), dissolved organic carbon (DOC), and non-purgable organic carbon (NPOC) in water, wastewater, and seawater in the range from 0.5 mg/L to 4000 mg/L of carbon. Higher levels may be determined by sample dilution. The sample is injected onto a quartz bed heated at 680°C. The sample converts into a gaseous phase and forced through a layer of catalyst ensuring conversion of all carbon containing compounds to CO₂. A non-dispersive infrared (NDIR) detector measures the resulting CO₂.

1.2 For TOC and DOC analysis a portion of the sample is injected to determine TC or dissolved carbon (DC). A portion of the sample is then acidified and purged to remove the IC. The purged inorganic carbon is measured as TIC, or DIC. TOC or DOC is calculated by subtracting the inorganic fraction from the total carbon:

$$\text{TOC} = \text{TC} - \text{IC}$$

1.3 For NPOC analysis a portion of sample is acidified and purged to remove IC. The purged sample is then injected to determine NPOC.

1.4 This test method was used successfully with reagent water spiked with potassium hydrogen phthalate, sucrose, nicotinic acid, benzoquinone, sodium dodecyl benzene sulfonate, urea, acetic acid, and humic acid. It is the user's responsibility to ensure the validity of this test method for waters of untested matrices.

1.5 This test method is applicable only to carbonaceous matter in the sample that can be introduced into the reaction zone. The syringe needle or injector opening size generally limits the maximum size of particles that can be so introduced.

1.6 In addition to laboratory analyses, this test method may be applied to stream monitoring.

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

1.8 *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

- D1129 Terminology Relating to Water
- D1192 Guide for Equipment for Sampling Water and Steam in Closed Conduits (Withdrawn 2003)³
- D1193 Specification for Reagent Water
- D2777 Practice for Determination of Precision and Bias of Applicable Test Methods of Committee D19 on Water
- D3370 Practices for Sampling Water from Closed Conduits
- D4129 Test Method for Total and Organic Carbon in Water by High Temperature Oxidation and by Coulometric Detection
- D5847 Practice for Writing Quality Control Specifications for Standard Test Methods for Water Analysis

3. Terminology

3.1 *Definitions:*

3.1.1 For definitions of terms used in this standard, refer to Terminology D1129.

3.2 *Definitions of Terms Specific to This Standard:*

3.2.1 *inorganic carbon (IC)*, *n*—carbon in the form of carbon dioxide, carbonate ion, or bicarbonate ion.

3.2.2 *total organic carbon (TOC)*, *n*—carbon in the form of organic compounds.

3.2.3 *non-purgable organic carbon (NPOC)*, *n*—carbon measured in a sample after acidification and sparging to remove inorganic carbon.

¹ This test method is under the jurisdiction of ASTM Committee D19 on Water and is the direct responsibility of Subcommittee D19.06 on Methods for Analysis for Organic Substances 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.

3.2.4 *total carbon (TC)*, *n*—the sum of IC and TOC.

3.2.5 *dissolved organic carbon (DOC)*, *n*—carbon determined on filtered samples.

3.2.6 *purgable organic carbon (POC)*, *n*—carbon that purges from acidified samples, also known as volatile organic compounds (VOC).

3.2.7 *refractory material*, *n*—that which cannot be oxidized completely under the test method conditions.

4. Summary of Test Method

4.1 *Fundamentals*—Carbon can occur in water as an inorganic and organic compound. This test method can be used to make independent measurements of IC, NPOC, and TC, and can also determine OC by the difference of TC and IC. DOC is determined on samples that have been filtered through a 0.45- μm filter.

4.2 TOC and DOC procedures require that IC has been removed from the sample before it is analyzed for organic carbon content. The sample free of IC is injected into the TOC instrument where all carbon is converted to CO_2 and measured by the detector. Failure of the method to remove all IC prior to analysis for organic carbon will result in significant error. A diagram of suitable apparatus is given in Fig. 1.

5. Significance and Use

5.1 This test method is used for determination of the carbon content of water from a variety of natural, domestic, and industrial sources. In its most common form, this test method is used to measure organic carbon as a means of monitoring organic pollutants in industrial wastewater. These measurements are also used in monitoring waste treatment processes.

5.2 The relationship of TOC to other water quality parameters such as chemical oxygen demand (COD) and total oxygen demand (TOD) is described in the literature.⁴

6. Interferences and Limitations

6.1 The oxidation of dissolved carbon to CO_2 is brought about at high temperatures (680°C) in the presence of oxygen. A catalyst promotes the oxidation process and the resulting carbon dioxide is measured by a non-dispersive infrared detector (NDIR). Suspended and refractory materials are completely oxidized under these conditions.

6.2 Acid preservation can precipitate some compounds, such as humic acids, removing them from solution and causing erroneously low results.

6.3 Homogenizing or sparging of a sample, or both, may cause loss of purgable organic compounds, thus yielding a value lower than the true TOC level. (For this reason, such measurements are sometimes known as NPOC). The extent and significance of such losses must be evaluated on an individual basis. Comparison of the difference, if any, between NPOC and TOC by subtraction represents POC lost during sparging.

6.4 If POC is important then TOC must be measured by subtraction:

$$\text{TOC} = \text{TC} - \text{TIC}$$

6.5 Note that error will be introduced when the method of difference is used to derive a relatively small level from two large levels. For example, a ground water high in IC and low in TOC will give a poorer TOC value as $(\text{TC} - \text{IC})$ than by direct measurement as NPOC.

6.6 Samples containing high levels (>1 ppm) of surfactant may lose TOC by foaming.

6.7 Elemental carbon may not be completely combusted at 680°C ; however, it is not generally found in water samples. Elemental carbon does not form during the catalytic oxidation of water samples.

6.8 Inorganics dissolved in the sample are not volatilized into gas and remain on the catalyst or quartz shard surfaces. High amounts of solids eventually react with the quartz surfaces causing devitrification, or solidify in the catalyst bed decreasing flow rates. Limit sample volume injected to reduce the amount of soluble salts and to reduce cooling of the reaction chamber. Buildup of salts; reduction of flow rate, or large injection volumes could result in peak splitting.

6.9 Carbon in reagent water and reagent blanks can be reduced to a minimum, and consistent value, but cannot be completely eliminated. Analyzing low-level TOC (less than 1.0 mg/L) bears special consideration requiring that the same water used to set the baseline be used to prepare the calibration standards.

6.10 Atmospheric carbon dioxide absorbs into reagent water increasing its inorganic carbon content with time. The small

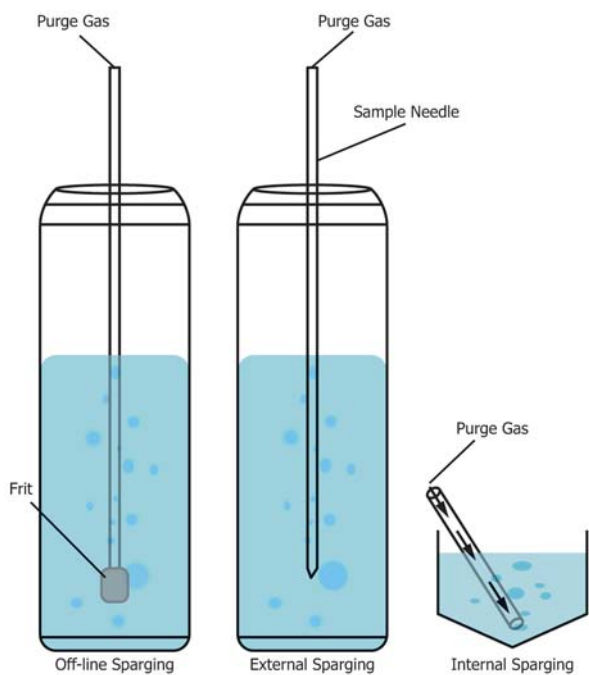


FIG. 1 TIC Removal

⁴ Handbook for Monitoring Industrial Wastewater, Section 5.3, U.S. Environmental Protection Agency, August 1973, pp. 5–12.

levels of CO₂ absorbed into reagent water can cause considerable inaccuracies in low-level TIC analysis. For instance, a 40-milliliter vial of reagent water containing no detectable TIC was analyzed to contain 160 µg/LTIC after 1 hour of exposure to ambient air.

6.11 Trace organics in the atmosphere can be absorbed into reagent water increasing its organic carbon content with time. The small levels of organics absorbed into reagent water can cause considerable inaccuracies in low-level (<1 mg/L) TOC measurements.

7. Apparatus

NOTE 1—See also Fig. 2.

7.1 *Sampling Devices*—Manually operated or automatically operated sampling valves, or syringes are typically used with this method. Sampling devices with inside diameters as small as 0.15 mm may be used with samples containing little or no particulate matter. Larger inside dimensions, such as 0.2 mm, will be required for samples with particulate matter.

7.2 *Apparatus for Carbon Determination*—This instrument consists of reagent and sample introduction mechanism, a gas-sparged reaction vessel for TIC removal, the high temperature combustion chamber with catalyst, a gas demister or dryer and halogen trap, an optional CO₂ trap, a CO₂-specific infrared

detector, a control system, and a display. Fig. 1 shows a diagram of such an arrangement.⁵

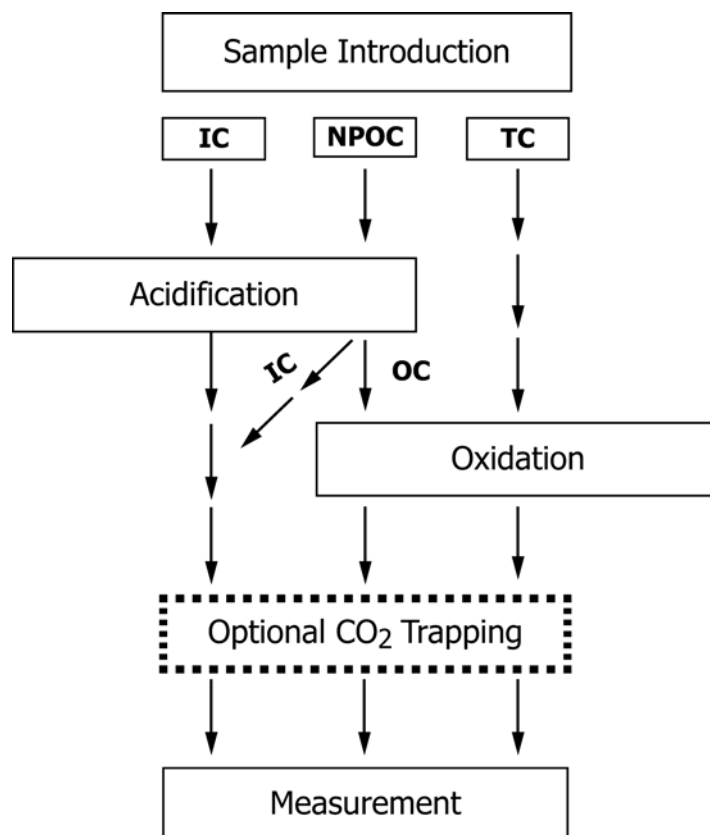
7.2.1 Reaction vessel consists of TIC removal and the combustion chamber.

7.2.1.1 *TIC Removal*—Sparging requires an inert vessel with a capacity of at least double the sample size with provision for sparging with 50 to 200 mL/min of carbon-free gas. This procedure should remove essentially all IC in 2 to 10 min, depending on design and can be at room temperature or at elevated temperatures (≤70°C) to promote CO₂ removal. Verify that heated sparging does not remove >5 % of the NPOC. Fig. 1 illustrates three different options for TIC removal.

7.2.1.2 *Combustion Chamber*—A heated catalyst contained in a quartz tube, may contain quartz wool, quartz shards, or other items to protect the catalyst from dissolved salts to extend its life.

7.2.2 *Gas Conditioning*—The gas passing from the reactor is dried, and the CO₂ produced is either trapped and later

⁵ The sole source of supply of the apparatus known to the committee at this time is the OI Analytical Aurora 1030C and 1020. 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.



IC—Inorganic Carbon
 NPOC—Non Purgable Organic Carbon
 TC—Total Carbon
 OC—Organic Carbon

FIG. 2 Diagram of Apparatus