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Suggested revisions are invited and should be submitted to the Standards Department, API, 1220 L Street, NW, Washington, DC 20005, standards@api.org.
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INTRODUCTION

The previous version of the automatic sampling practice described the design, installation, testing, and operation of automated equipment for the extraction of representative samples from the flowing stream and storing mainly for crude oil.

This practice is a performance-based standard. It still includes the design, installation, testing, and operation of automated equipment for extraction of representative samples. It also includes the testing and proving of a sampling system in the field under actual operating conditions to ensure that the equipment, installation, and operating procedures produce representative samples. The acceptance criteria for custody transfer are covered in this practice. This practice does not address how to sample crude at temperatures below the freezing point of water. Extensive revisions have been made to the prior version of D4177 (API MPMS Chapter 8.2).

This practice also provides guidance for periodic verification of the sampling system.

This practice is separated into three parts:

General—Sections 5 – 17 (Part I) are currently applicable to crude oil and refined products. Review this section before designing or installing any automatic sampling system.

Crude Oil Sampling—Section 18 (Part II) contains additional information required to complete the design, testing, and monitoring of a crude oil sampling system.

Refined Product Sampling—Section 19 (Part III) contains additional information required to complete the design of a refined product sampling system.

A representative sample is “A portion extracted from the total volume that contains the constituents in the same proportions that are present in that total volume.” Representative samples are required for the determination of chemical and physical properties that are used to establish standard volumes, prices, and compliance with commercial and regulatory specifications.

The process of obtaining a representative sample consists of the following: the physical equipment, the correct matching of that equipment to the application, the adherence to procedures by the operator(s) of that equipment, and the proper handling and analysis.

1. Scope*

1.1 This practice describes general procedures and equipment for automatically obtaining samples of liquid petroleum and petroleum products, crude oils, and intermediate products from the sample point into the primary container. This practice also provides additional specific information about sample container selection, preparation, and sample handling. If sampling is for the precise determination of volatility, use Practice D5842 (API MPMS Chapter 8.4) in conjunction with this practice. For sample mixing and handling, refer to Practice D5854 (API MPMS Chapter 8.3). This practice does not cover sampling of electrical insulating oils and hydraulic fluids.

*A Summary of Changes section appears at the end of this standard
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1.3 Units—The values stated in either SI units or US Customary (USC) units are to be regarded separately as standard. The values stated in each system may not be exact equivalents; therefore, each system shall be used independently of the other. Combining values from the two systems may result in non-conformance with the standard. Except where there is no direct SI equivalent, such as for National Pipe Threads/diameters, or tubing.

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 of regulatory limitations prior to use.

2. Referenced Documents

2.1 ASTM Standards:

- D4007 Test Method for Water and Sediment in Crude Oil by the Centrifuge Method (Laboratory Procedure)
- D4840 Guide for Sample Chain-of-Custody Procedures
- D4928 Test Method for Water in Crude Oils by Coulometric Karl Fischer Titration
- D5842 Practice for Sampling and Handling of Fuels for Volatility Measurement

2.2 API Standards:

- MPMS Chapter 3 Tank Gauging
- MPMS Chapter 4 Proving Systems
- MPMS Chapter 5 Metering
- MPMS Chapter 8.3 Practice for Mixing and Handling of Liquid Samples of Petroleum and Petroleum Products (ASTM Practice D5854)
- MPMS Chapter 8.4 Practice for Manual Sampling and Handling of Fuels for Volatility Measurement (ASTM Practice D5842)
- MPMS Chapter 10 Sediment and Water
- MPMS Chapter 13 Statistical Aspects of Measuring and Sampling
- MPMS Chapter 20 Production Allocation Measurement for High Water Content Crude Oil Sampling
- MPMS Chapter 21 Flow Measurement Using Electronic Metering Systems

2.3 ISO Standards:


NOTE 1—See the Bibliography at the end of this standard for important historical references.

3. Terminology

3.1 Definitions of Terms Specific to This Standard:

3.1.1 automatic sampling system, n—fluid sampling system that consists of: (a) flowing fluid stream conditioning, if required; (b) a means of automatically extracting a representative sample; (c) pacing of the sample extraction in a flow or time proportional manner; and (d) delivering of each extracted sample to a sample container or an analyzer.

3.1.1.1 Discussion—The system consists of a sample extractor with an associated controller and flow-measuring or timing device, collectively referred to as an automatic sampler or auto-sampler. In addition, the system may include a flow conditioner, slipstream, sample probe, and sample conditioning.

3.1.1.2 Discussion—Systems may deliver the sample directly to an analytical device or may accumulate a composite sample for offline analysis, in which case, the system includes sample mixing and handling and a primary sample container.

3.1.1.3 Discussion—Automatic sampling systems may be used for liquids.

3.1.2 batch, n—discrete shipment of commodity defined by a specified quantity, a time interval, or quality.

3.1.3 component testing, n—process of individually testing the components of a system.

3.1.4 dead volume, n—in sampling, the volume trapped between the extraction point and the primary sample container.

3.1.4.1 Discussion—This represents potential for contamination between batches.