



Designation: D6060 – 17

Standard Test Method for Sampling of Process Vents with a Portable Gas Chromatograph¹

This standard is issued under the fixed designation D6060; 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 describes a method for direct sampling and analysis of process vents for volatile organic compound (VOC) vapors and permanent gases using a portable gas chromatograph (GC).

1.2 This test method is applicable to analysis of permanent gases such as oxygen (O_2), carbon dioxide (CO_2) and nitrogen (N_2), as well as vapors from organic compounds with boiling points up to 125°C.

1.3 The detection limits obtained will depend on the portable gas chromatograph and detector used. Detectors available include but are not limited to thermal conductivity, photoionization, argon ionization, and electron capture. For instruments equipped with thermal conductivity detectors, typical detection limits are one to two parts per million by volume (ppm(v)) with an applicable concentration range to high percent by volume levels. For instruments with photoionization detectors detection limit of one to ten parts per billion by volume (ppb(v)) are obtainable with a concentration range from 1000 to 2000 ppm(v). The argon ionization detector has an achievable detection limit of one (ppb(v)), while the electron capture detector has an achievable detection limit of one part per trillion by volume (ppt(v)) for chlorinated compounds.

1.4 The applicability of this test method should be evaluated for each VOC by determining stability, reproducibility, and linearity.

1.5 The appropriate concentration range must also be determined for each VOC, as the range will depend on the vapor pressure of the particular VOC.

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

1.7 *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. Refer to Section 8 on Hazards for additional safety precautions.*

1.8 *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

D1356 Terminology Relating to Sampling and Analysis of Atmospheres

D3154 Test Method for Average Velocity in a Duct (Pitot Tube Method)

D3464 Test Method for Average Velocity in a Duct Using a Thermal Anemometer

E355 Practice for Gas Chromatography Terms and Relationships

2.2 Other Document:

NFPA 70 National Electrical Code³

NFPA 496 Standard for Purged and Pressurized Enclosures for Electrical Equipment³

3. Terminology

3.1 *Definitions*—For the definition of terms used in this test method, refer to Terminology **D1356** and Practice **E355**.

3.2 Definitions of Terms Specific to This Standard:

3.2.1 *portable, n*—refers to gas chromatograph with internal battery, internal sample pump, and internal/rechargeable carrier gas supply cylinder.

¹ This test method is under the jurisdiction of ASTM Committee **D22** on Air Quality and is the direct responsibility of Subcommittee **D22.03** on Ambient Atmospheres and Source Emissions.

Current edition approved Oct. 1, 2017. Published October 2017. Originally approved in 1996. Last previous edition approved in 2009 as D6060 – 96 (2009). DOI: 10.1520/D6060-17.

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 Available from National Fire Protection Association (NFPA), 1 Batterymarch Park, Quincy, MA 02169-7471, <http://www.nfpa.org>.

4. Summary of Test Method

4.1 One end of a sampling line (typically 6 mm outside diameter TFE-fluorocarbon tubing) is connected to a tee in a process vent and the other end to a condensation trap (see 6.1), which is connected to a gas sampling bulb. The outlet of the gas sampling bulb is connected to a sampling pump set at a flow rate of 0.5 to 2 L/min. The sample line from the portable gas chromatograph is inserted through the septum port of the gas sampling bulb. At user selected intervals, the internal pump of the portable gas chromatograph is activated and process vapors drawn through the injection valve of the gas chromatograph and analyzed.

5. Significance and Use

5.1 This test method has been widely used to obtain mass balance data for process scrubbers, to determine the efficiency of VOC emission control equipment, and to obtain data to support air permit applications.

5.2 This test method will have applications to the Maximum Achievable Control Technology (MACT) Rule and may have applications to Compliance Assurance Monitoring verification required by the 1990 Clean Air Act Title III Amendments.

5.3 This test method, when used with Test Methods D3464 or D3154 or on-line process flow meter data, can be used to calculate detailed emission rate profiles for VOCs from process vents.

5.4 This test method provides nearly real time results that can detect process changes or upsets that may be missed using conventional sorbent tube or integrated gas sampling bag sampling.

6. Interferences

6.1 Water or liquid in the process line will plug the sample line of the gas chromatograph, since the injection valve of most portable GCs is not heated. The condensation trap is designed to protect the portable gas chromatograph if liquids are present or occur during process upset.

6.2 Interferences sometimes result from analytes having similar retention times during gas chromatography.

6.3 General approaches which can be followed to resolve such interferences are given below:

6.3.1 Change the type of column, length of column, or operating conditions.

6.3.2 Analyze using a nonpolar methyl silicone column which separates according to boiling point of the compounds and a polar column whose separations are influenced by the polarity of the compounds.⁴

6.3.3 Use a mass spectrometer to verify the identity of peaks.

⁴ The columns in most portable gas chromatographs are easily interchanged. One manufacturer has an instrument that simultaneously injects onto two user selected column modules.

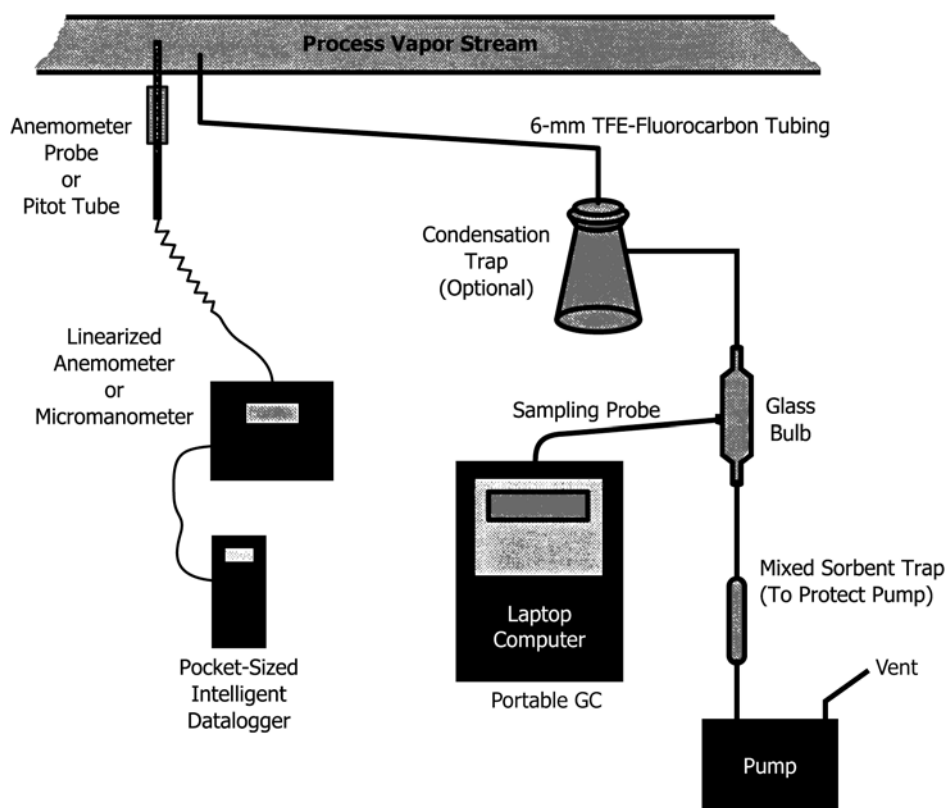


FIG. 1 Schematic of Process Sampling Equipment