

Designation: D 5832 - 98 (Reapproved 2003)

Standard Test Method for Volatile Matter Content of Activated Carbon Samples¹

This standard is issued under the fixed designation D 5832; 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 the percentage of gaseous products, exclusive of moisture vapor, present in virgin and used activated carbons which are released under specific conditions of the test.
- 1.2 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:
- D 2652 Terminology Relating to Activated Carbon²
- D 2867 Test Method for Moisture in Activated Carbon²
- D 3175 Test Method for Volatile Matter in the Analysis Sample of Coal and Coke³

3. Terminology

3.1 *Definitions*—For definitions of terms used in this test method relating to activated carbon, refer to Terminology D 2652.

4. Summary of Test Method talog/standards/sist/e26d66f

4.1 Volatile matter is determined by establishing the loss in mass resulting from heating an activated carbon sample under rigidly controlled conditions. The measured mass loss, corrected for moisture as determined in Test Method D 2867, establishes the volatile matter content of the activated carbon sample.

5. Significance and Use

5.1 Volatile matter, when determined as herein described, may be used as a relative measure of the extent of carbonization in an activated carbon and the extent of loading of volatile material on an activated carbon that has been used in an adsorption application.

- ¹ This test method is under the jurisdiction of ASTM Committee D28 on Activated Carbon and is the direct responsibility of Subcommittee D28.04 on Gas Phase Evaluation Tests.
- Current edition approved March 10, 2003. Published July 2003. Originally approved in 1995. Last previous edition approved in 1998 as D 5832 98.
 - ² Annual Book of ASTM Standards, Vol 15.01.
 - ³ Annual Book of ASTM Standards, Vol 05.05.

- 5.2 Combined with other information, the volatile matter of an activated carbon may be useful in evaluating its performance in an adsorption application.
- 5.3 Other automated methods for the determination of the volatile content of solids, such as using a thermogravimetric analyzer (TGA), can be used in place of this test method with equally reliable results.

6. Apparatus

- 6.1 Crucible and Cover, high temperature porcelain, high form, 30 cc capacity.
- 6.2 *Oven*, forced-air circulation, capable of temperature regulation up to 250°C.
- 6.3 Moisture Determination Apparatus, as described in Test Method D 2867.
- 6.4 Muffle Furnace, gravity circulation, capable of temperature regulation at 950 \pm 25°C. An electric furnace similar to the one described in Test Method D 3175 is suitable for use in this test method.
 - 6.5 Desiccator, glass with indicating type desiccant.
 - 6.6 Balance, analytical, capable of 0.1 mg sensitivity.

7. Hazards

- 7.1 The furnace used in this test method should be located in a well ventilated area to eliminate exposure to possible toxic vapors that may evolve from the carbon sample during the high temperature heating.
- 7.2 Exercise care when working with the high temperature furnace to eliminate the possibility of burns.

8. Procedure

- 8.1 Determine the moisture content of an as-received representative portion of the sample using the Xylene-Extraction Test Method described in D 2867. If the as-received sample is wet, drain it of all free liquid before the representative sample is taken.
- 8.2 Weigh to 0.1 mg accuracy a crucible and cover that have been ignited in a muffle furnace regulated at 950°C for 30 min and cooled in a desiccator. Record the weight.
- 8.3 Using a spoon or spatula, dip from the sample bottle approximately 1 g of the as-received sample and place it in the pre-dried and tared crucible. Cover it with a lid and immediately weigh it to the nearest 0.1 mg.