



## Standard Test Method for Ash in a Graphite Sample<sup>1</sup>

This standard is issued under the fixed designation C561; 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 ( $\epsilon$ ) indicates an editorial change since the last revision or reapproval.

*This standard has been approved for use by agencies of the Department of Defense.*

<sup>e1</sup> NOTE—Updated units of measure throughout the standard editorially in May 2010.

### 1. Scope

1.1 This test method provides a practical determination for the ash content in a graphite sample.

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

1.3 *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:*<sup>2</sup>

C562 Test Method for Moisture in a Graphite Sample

### 3. Significance and Use

3.1 This test method provides a practical estimate of non-burnable residues in commercially available graphite materials. The ash values determined by this test method are of use in comparing the relative purity of various grades of graphite. To facilitate use, this test method institutes simplifications that preclude the ability to determine absolutely the ash values of the test graphite material due to uncontrolled sources of trace contamination.

3.2 This test method is not intended for use in determining the ash content of purified graphites, for example, nuclear materials. The relationship between the mineral content of a graphite sample and the ash content of that sample is unknown and is not determined by the application of this test method.

<sup>1</sup> This test method is under the jurisdiction of ASTM Committee D02 on Petroleum Products and Lubricants and is the direct responsibility of Subcommittee D02.F0 on Manufactured Carbon and Graphite Products.

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<sup>2</sup> 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.

### 4. Interferences

4.1 Although permitted within the scope of this test method, the use of alumina ceramic crucibles may affect results due to difficulties in obtaining repeatable or proper weights, or both, because of (1) the hygroscopic nature of some ceramic crucibles, and (2) the possible chemical combination of trace elements with the ceramic crucible.

4.2 Any ash or trace elements introduced to the sample will influence results. Contamination can occur during drilling to obtain the sample and during pulverization. (See 6.1.)

### 5. Apparatus

5.1 *Alumina Ceramic or Platinum Crucible or Dish*, suitable for holding sample (subsequently called sample holder).

5.2 *Analytical Balance*, capable of weighing to  $\pm 0.0002$  g.

5.3 *Muffle Furnace*, capable of reaching 950°C with controller capable of maintaining a temperature of  $950^\circ \pm 20^\circ\text{C}$ .

5.4 *Platinum or Stainless Steel Wire*.

5.5 *Desiccator*, charged with indicating desiccant.

5.6 *Drying Oven*, air convection type, capable of being controlled to  $110 \pm 2^\circ\text{C}$ .

### 6. Sampling

6.1 Samples may be solid or particulate. Solid bodies may be sampled by removing one or more solid pieces from the body by, for example, sawing, turning, milling, or fracturing. Particulate samples may be generated from solid bodies by drilling, using a carbide drill to minimize contamination, or by other crushing and grinding methods.

### 7. Procedure

7.1 Dry the sample in accordance with Test Method C562, or for a minimum of 16 h in a drying oven at  $110 \pm 2^\circ\text{C}$ , and allow the sample to cool to room temperature in the desiccator.

7.2 Tare a dried sample holder using an analytical balance to  $\pm 0.002$  g. As soon as the sample has cooled to room temperature, remove it from the desiccator and weigh a 25- to