

## Standard Practice for Preparation of Calcined Petroleum Coke Samples for Analysis<sup>1</sup>

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### 1. Scope

1.1 This practice covers the preparation procedures necessary for the reduction and division of calcined petroleum coke samples in order to generate appropriate analytical samples upon which physical and chemical analytical tests will be performed.

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>

- D1552 Test Method for Sulfur in Petroleum Products (High-Temperature Method)
- D2638 Test Method for Real Density of Calcined Petroleum Coke by Helium Pycnometer
- D4292 Test Method for Determination of Vibrated Bulk Density of Calcined Petroleum Coke
- D4422 Test Method for Ash in Analysis of Petroleum Coke
- D4930 Test Method for Dust Control Material on Calcined Petroleum Coke
- D4931 Test Method for Gross Moisture in Green Petroleum Coke

- D5004 Test Method for Real Density of Calcined Petroleum Coke by Xylene Displacement
- D5056 Test Method for Trace Metals in Petroleum Coke by Atomic Absorption
- D5187 Test Method for Determination of Crystallite Size  $(L_c \text{ of Calcined Petroleum Coke by X-Ray Diffraction})$
- D5600 Test Method for Trace Metals in Petroleum Coke by Inductively Coupled Plasma Atomic Emission Spectrometry (ICP-AES)
- D5709 Test Method for Sieve Analysis of Petroleum Coke
- D6376 Test Method for Determination of Trace Metals in Petroleum Coke by Wavelength Dispersive X-ray Fluorescence Spectroscopy

### 3. Terminology

3.1 Definitions:

3.1.1 *analysis sample*—the reduced and divided representative portion of the bulk sample, prepared for use in the laboratory.

3.1.2 *composite sample*—a sample, representative of an entire consignment of calcined petroleum coke, generated by mixing portions of gross samples from different lots together in mass fractions proportioned to the consignment.

3.1.3 *gross sample*—the original, uncrushed representative portion taken from a shipment or lot of coke.

3.1.4 *intermediate sample*—a sample, representative of a gross sample, upon which no analysis is to be performed, yet required for generation of analysis samples after undergoing further division and reduction.

3.1.5 *lot*—a quantity of calcined petroleum coke to be represented by a gross sample.

3.1.6 *riffle*—a manual sample divider which splits the sample stream into a number of alternate elements.

3.1.7 *sample division*—the process whereby a sample is reduced in mass without change in particle size.

3.1.8 *sample preparation*—the process that may include drying, crushing, division, and mixing of a gross sample for the purpose of obtaining an unbiased analysis sample.

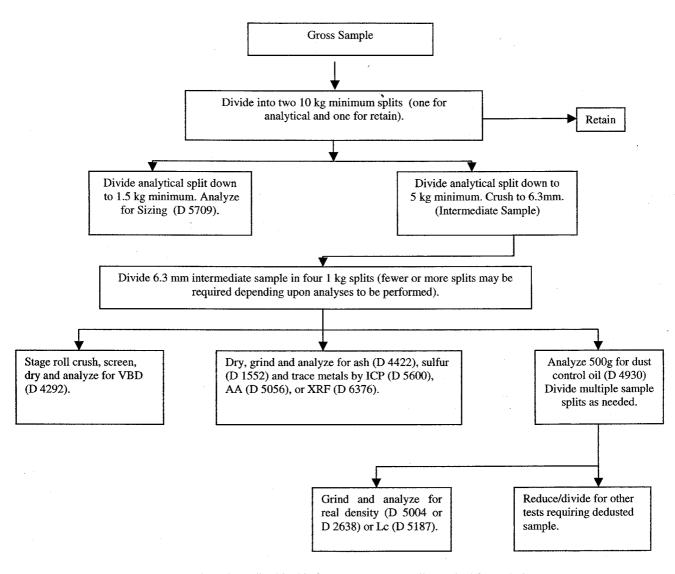
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<sup>&</sup>lt;sup>1</sup> This practice is under the jurisdiction of ASTM Committee D02 on Petroleum Products and Lubricants and is the direct responsibility of Subcommittee D02.05 on Properties of Fuels, Petroleum Coke and Carbon Material

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<sup>&</sup>lt;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.

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Note—The tasks outlined in this figure are not necessarily required for analysis. FIG. 1 Preparation of Calcined Petroleum Coke

3.1.9 *sample reduction*—the process whereby a sample is reduced in particle size by crushing or grinding without significant change in chemical properties.

3.1.10 *top size*—the size of the smallest opening of one sieve of a series upon which is cumulatively retained a total of less than 5 % of the sample. This defined top size is not to be confused with the size of the largest particle in a lot.

### 4. Significance and Use

4.1 This practice provides field personnel and laboratories with standard procedures for dividing, reducing, and mixing gross samples and intermediate samples, such that the resulting prepared analysis samples are more uniform from laboratory to laboratory. Adherence to these guidelines is expected to provide significant reduction in interlaboratory variability.

### 5. Organization of Sample Preparation

5.1 Initial Division of Gross Sample:

5.1.1 Determine the required analyses for the lot and manage the division steps accordingly to achieve enough analytical sample to perform all required analyses in duplicate (see flowchart in Fig. 1 as an example). Divide the gross sample by use of a riffle or rotary sample divider to an intermediate sample of sufficient mass.

5.1.2 Exercise care in the division operation to preserve the particle size distribution of the gross sample. Riffle openings must be at least three times the top size of the calcined petroleum coke being divided. A feed hopper or vibratory feeder, or both, are recommended to feed the coke into the riffle. Enclosed riffles are recommended to minimize dusting and loss of sample.

5.2 Subsequent Reduction and Division:

5.2.1 Crushing processes are to be carried out such that the apparatus does not contribute significant impurities into the analysis sample. For example, if trace metal analyses are to be performed on a number 60 mesh analysis sample, a ring and

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TABLE 1	Example of Mixing	a Gross Samples to	o Generate a Weighted Avera	ge Vessel Composite Sample

	Lot 1 Hold 1	Lot 2 Hold 2	Lot 3 Hold 3	Lot 4 Hold 4	Lot 5 Hold 5	Lot 6 Hold 6	Lot 7 Hold 7	Total
Lot tonnage (short tons) Target sample mass mixed into	7800 1426 ± 29	7500 1371 ± 27	8200 1499 ± 30	7500 1371 ± 27	8000 1463 ± 29	7800 1426 ± 29	7900 1444 ± 29	54 700 10 000 ± 200
composite (grams)	1420 ± 29	1371 ± 27	1499 ± 30	1371 ± 27	1403 ± 29	1420 ± 29	1444 ± 29	10 000 ± 200

puck mill with tungsten carbide grinding components is recommended to minimize the metallic impurities of analytical interest that may be added to sample.

5.2.2 If trace metals are not required for testing on a number 60 mesh analysis sample, a bench-top hammer mill or ball mill are adequate for performing the grinding operation.

5.3 Refer to actual test methods for size and mass of samples required for analysis.

5.4 Removal of dedusting oil may be required for analytical purposes and for subsequent analyses such as real density or porosity. Oil may be extracted and quantified by Test Method D4931 or thermally removed.

5.5 Mixing of gross samples to generate a composite sample representing more than one lot is frequently required. Mixing must be planned such that the final composite sample has a mass of no less than 10 kg so that all the required analyses may be performed.

5.5.1 Divide each gross sample down into an intermediate sample weighing no less than 1.5 kg such that each intermediate sample is still representative for sizing determination by Test Method D5709.

5.5.2 Each lot sample will be proportionately represented in the composite, as the lot tonnage was representative of the total cargo. An example of a large seven-hold vessel cargo represented by a 10 kg vessel composite sample is illustrated in Table 1. To calculate the individual target sample mass, use the following formula:

target sample mass 
$$= \frac{A}{B} \times C$$
 (1)

where:

A = mass of individual lot in tons,

B = mass of total vessel in tons, and

C = mass of composite sample in grams.

5.5.3 Adjust the mass of the intermediate sample to within 2 % of the target mass contribution to the composite sample by removing material from the intermediate sample. Use appropriate means to preserve the particle size distribution of the original gross sample.

### 6. Keywords

6.1 calcined petroleum coke; sample division; sample preparation; sample reduction

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