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Standard Practice for
Preparation of Calcined Petroleum Coke Samples for Analysis[1]

This standard is issued under the fixed designation D6969; 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~~ 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 either SI units or inch-pound units are to be regarded ~~as standard. No other units of measurement are included in this~~ 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.

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:*[2]
D1552 Test Method for Sulfur in Petroleum Products by High Temperature Combustion and IR Detection
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 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~~ sample, n—*the reduced and divided representative portion of the bulk sample, prepared for use in the laboratory.

3.1.2 *composite ~~sample~~ sample, n—*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~~ sample, n—*the original, uncrushed representative portion taken from a shipment or lot of coke.

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*A Summary of Changes section appears at the end of this standard.

3.1.4 *intermediate ~~sample~~—sample, n—*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~~—lot, n—*a quantity of calcined petroleum coke to be represented by a gross sample.

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

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

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

3.1.9 *sample ~~reduction~~—reduction, n—*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~~—size, n—*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 :~~ 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 :~~ 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 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~~ D4930 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:

$$\text{target sample mass} = \frac{A}{B} \times C \tag{1}$$

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