



Designation: D4326 – 11

# Standard Test Method for Major and Minor Elements in Coal and Coke Ash By X-Ray Fluorescence<sup>1</sup>

This standard is issued under the fixed designation D4326; 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.

## 1. Scope

1.1 This test method covers the analysis of the commonly determined major and minor elements in ash from coal or coke using X-ray fluorescence (XRF) techniques.

NOTE 1—Test Method D1757 is used for determination of sulfur.

NOTE 2—Although not included in the present method, the determination of barium, strontium, and manganese may be required to yield adequate totals.

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>

D121 Terminology of Coal and Coke

D1757 Test Method for Sulfate Sulfur in Ash from Coal and Coke (Withdrawn 2009)<sup>3</sup>

D3174 Test Method for Ash in the Analysis Sample of Coal and Coke from Coal

E2 Methods of Preparation of Micrographs of Metals and Alloys (Including Recommended Practice for Photography As Applied to Metallography); Replaced by E 883 (Withdrawn 1983)<sup>3</sup>

E11 Specification for Woven Wire Test Sieve Cloth and Test Sieves

<sup>1</sup> This test method is under the jurisdiction of ASTM Committee D05 on Coal and Coke and is the direct responsibility of Subcommittee D05.29 on Major Elements in Ash and Trace Elements of Coal.

Current edition approved April 1, 2011. Published April 2011. Originally approved in 1984. Last previous edition approved 2004 as D4326–04. DOI: 10.1520/D4326-11.

<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.

<sup>3</sup> The last approved version of this historical standard is referenced on www.astm.org.

## 3. Terminology

3.1 *Definitions*—Definitions applicable to this test method are listed in D121, Terminology of Coal and Coke.

## 4. Summary of Test Method

4.1 The coal or coke to be analyzed is ashed under standard conditions and ignited to constant weight. Previously ashed materials are ignited to constant weight under standard conditions. The ash is fused with lithium tetraborate ( $\text{Li}_2\text{B}_4\text{O}_7$ ) or other suitable flux and either ground and pressed into a pellet or cast into a glass disk. The pellet or disk is then irradiated by an X-ray beam of short wavelength (high energy). The characteristic X-rays of the atom that are emitted or fluoresced upon absorption of the primary or incident X-rays are dispersed and intensities at selected wavelengths are measured by sensitive detectors. Detector output is related to concentration by calibration curves or by computerized data-handling equipment.

4.2 The K spectral lines are used for all of the elements determined by this procedure.

4.3 All elements are determined as the element and reported as the oxide and include Si, Al, Fe, Ca, Mg, Na, K, P, Ti, Mn, Sr, and Ba.

## 5. Significance and Use

5.1 A compositional analysis of ash is used in describing the quality of coal for its complete characterization. Ash composition is useful in predicting slagging and fouling characteristics of combusted materials as well as the potential utilization of ash by-products.

5.2 The chemical composition of laboratory prepared coal or coke ash is rarely, if ever, representative of the composition of the mineral matter in the coal because the ashing process can alter some minerals. However, it can approximate the composition of the fly ash and slag resulting from commercial combustion of coal or coke.

## 6. Apparatus

6.1 *Ashing Furnace*, with air circulation as specified in Test Method D3174 and capable of having its temperature regulated between 700 and 750°C.

6.2 *Fusion Furnace or Fluxing Device*, with an operating temperature of at least 1000°C.

6.3 *Fusion Crucibles*, either high-purity graphite (22 mm high and 19 mm wide, inside diameter) or platinum-gold alloy of a similar or larger capacity.

6.4 *Pulverizers*, including agate, mullite or tungsten carbide mortar, and pestle, minimum capacity 25 mL.

6.5 *Sieves*, 250- $\mu\text{m}$  (No. 60) and 75- $\mu\text{m}$  (No. 200) U.S.A standard sieve as specified in Specification E11.

6.6 *Compactor*—A press equipped with a gage enabling reproducible pressures (exceeding  $1.72 \times 10^8$  Pa (25 000 psi)) if pressed pellets are utilized.

6.7 *Excitation Source*, with a stable electrical power supply ( $\pm 1$  %) and a high-intensity, short-wavelength X-ray capability.

6.8 *Spectrometer*—A wavelength or energy dispersive system equipped with a vacuum sample chamber.

6.8.1 *Analyzing Crystal (Wavelength Units)*—The choice of the analyzing crystal is made on the basis of the element to be determined. An attempt should be made to use the crystal that yields the maximum sensitivity with minimum interferences. The same crystal must be used for standards and unknowns.

6.8.2 *Detectors*—Scintillation and gas-flow counters are used with wavelength systems while lithium-drifted diodes are used for energy dispersive systems.

## 7. Reagents

7.1 *Purity of Reagents*—Reagent grade chemicals shall be used in all tests. Unless otherwise indicated, it is intended that all reagents shall conform to the specifications of the Committee on Analytical Reagents of the American Chemical Society, where such specifications are available.<sup>4</sup> Other grades may be used, provided it is first ascertained that the reagent is of sufficiently high purity to permit its use without lessening the accuracy of the determination.

7.2 *Detector Gas*—The usual gas composition of the gas used in the flow-proportional counters is 90 % argon, 10 % methane, although other compositions may be used.

7.3 *Heavy Absorber*—Where heavy absorbers, such as lanthanum oxide or barium oxide, are used they shall be a minimum of 99.99 % purity.

7.4 *Fluxes*—Lithium or sodium borates or carbonates, or combination thereof, are often used for sample fusion. Lithium or ammonium iodide used as a nonwetting agent and potassium or ammonium nitrate used as an oxidizing agent may be used provided they do not contribute to spectral interference.

7.5 *Binders*—Where pressed pellets are used for analysis, the binder used shall contribute no spectral interferences during the determination.

<sup>4</sup> *Reagent Chemicals, American Chemical Society Specifications*, American Chemical Society, Washington, DC. For suggestions on the testing of reagents not listed by the American Chemical Society, see *Analar Standards for Laboratory Chemicals*, BDH Ltd., Poole, Dorset, U.K., and the *United States Pharmacopeia and National Formulary*, U.S. Pharmaceutical Convention, Inc. (USPC), Rockville, MD.

## 8. Preparation of Coal Ash and Coke Ash

8.1 Prepare the ash from a thoroughly mixed representative analysis sample of air-dried coal or coke that has been ground to pass a 250- $\mu\text{m}$  (No. 60) U.S.A standard sieve. Spread the coal or coke in a layer not over 6 mm ( $\frac{1}{4}$  in.) in depth in a fireclay, porcelain, or platinum roasting dish. Place the ash in a cold muffle furnace and heat gradually so that the temperature reaches 500°C in 1 h and 750°C in 2 h. Ignite at 750°C to constant weight, cool, grind to pass a 74- $\mu\text{m}$  (No. 200) U.S.A standard sieve, and reignite at 750°C for 1 h. Immediately, prepare the analysis sample or store the dry ash in a vacuum desiccator. If samples are stored and the absorption of moisture or CO<sub>2</sub>, or both, is in question, reignite the ash using the 500-750°C staged combustion before use. Alternatively, determine loss on ignition using the 500-750°C staged combustion on a separate sample weighed out at the same time as the test portion and make the necessary corrections. Thoroughly mix each sample before weighing.

8.2 Materials previously ashed, fly ash, or bottom ash must be ignited to constant weight at 750°C and cooled in a desiccator before analysis sample preparation, or alternatively, weight loss or gain must be determined on a second sample at 750°C taken at the same time as the analysis sample so that analysis determined on an as-received basis can be reported on an ignited ash basis.

## 9. Preparation of Analytical Sample

9.1 Mix the powdered sample, weigh a portion of the sample, and mix with a suitable amount of flux (2 to 10 g of flux per gram of sample) (Note 3). When a heavy absorber is used (Note 4), it is added at this point in an amount equal to the amount of sample and thoroughly blended with the mix. A portion of the flux used as a cap on the mix ensures washing down any of the material from the sides of the crucible.

NOTE 3—The amount of sample and flux used are determined by the necessity of forming a disk or pellet of 2-mm thickness to fit the sample holder of the spectrometer used. All fluxes and other additives should be added in an appropriate manner at the time of sample preparation.

9.1.1 Fluxing materials commonly available are not consistent and can vary in volatile losses upon fusing. In order to eliminate errors caused by this loss, one of three methods must be employed. First, the entire bottle of flux may be heated to the fused state, then cooled, reground, and stored in a desiccator. Second, a weighed sample from each bottle is fused and a loss on fusion is determined, which is then applied as a correction for each sample prepared from that bottle. Third, the entire mass of each sample prepared (flux, sample, and heavy absorber, if used) may be weighed and an independent fusion loss calculated.

NOTE 4—Use of a heavy absorber has the advantage of allowing the use of a much smaller sample weight to reach infinite thickness and allows for calculation of concentrations using a simple linear regression coefficient. It does, however, prevent determination of a number of trace elements and may be impractical for energy dispersive systems where its addition may cause increased detector dead time and complicate correction procedures. The use of a heavy absorber is recommended only when its absence is impractical or inconvenient.

9.2 The sample mix is fused at approximately 1000°C, in a fusion furnace or fluxing device, for a length of time sufficient