



Standard Test Method for Gross Calorific Value of Coal and Coke¹

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1. Scope

1.1 This test method pertains to the determination of the gross calorific value of coal and coke by either an isoperibol or adiabatic bomb calorimeter.

1.2 The values stated in SI units are regarded as 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.* Specific hazard statements are given in Section 8.

2. Referenced Documents

2.1 ASTM Standards:

- D 121 Terminology of Coal and Coke²
- D 346 Practice for Collection and Preparation of Coke Samples for Laboratory Analysis²
- D 388 Classification of Coals by Rank²
- D 1193 Specification for Reagent Water³
- D 2013 Method of Preparing Coal Samples for Analysis²
- D 3173 Test Method for Moisture in the Analysis Sample of Coal and Coke²
- D 3177 Test Method for Total Sulfur in the Analysis Sample of Coal and Coke²
- D 3180 Practice for Calculating Coal and Coke Analyses from As-Determined to Different Bases²
- D 4239 Test Method for Sulfur in the Analysis Sample of Coal and Coke Using High Temperature Tube Furnace Combustion Methods²
- D 5142 Test Methods for Proximate Analysis of the Analysis Sample of Coal and Coke by Instrumental Procedures²
- E 1 Specification for ASTM Thermometers⁴
- E 144 Practice for Safe Use of Oxygen Combustion Bombs⁵
- E 178 Practice for Dealing with Outlying Observations⁶

¹ This test method is under the jurisdiction of ASTM Committee D05 on Coal and Coke and is the direct responsibility of Subcommittee D05.21 on Methods of Analysis.

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² *Annual Book of ASTM Standards*, Vol 05.06.

³ *Annual Book of ASTM Standards*, Vol 11.01.

⁴ *Annual Book of ASTM Standards*, Vol 14.03.

⁵ *Annual Book of ASTM Standards*, Vol 14.04.

⁶ *Annual Book of ASTM Standards*, Vol 14.02.

3. Terminology

3.1 Definitions:

3.1.1 *calorific value*—the heat produced by combustion of a unit quantity of a substance under specified conditions.

3.1.2 *calorimeter*—a device for measuring calorific value consisting of a bomb, its contents, a vessel for holding the bomb, temperature measuring devices, ignition leads, water, stirrer, and a jacket maintained at specified temperature conditions.

3.1.3 *adiabatic calorimeter*—a calorimeter which has a jacket temperature adjusted to follow the calorimeter temperature so as to maintain zero thermal head.

3.1.4 *isoperibol calorimeter*—a calorimeter which has a jacket of uniform and constant temperature.

3.1.5 *gross calorific value (gross heat of combustion at constant volume), Q_v (gross)*—the heat produced by complete combustion of a substance at constant volume with all water formed condensed to a liquid.

3.1.6 *heat of formation*—the change in heat content resulting from the formation of 1 mole of a substance from its elements at constant pressure.

3.1.7 *net calorific value (net heat of combustion at constant pressure), Q_p (net)*—the heat produced by combustion of a substance at a constant pressure of 0.1 MPa (1 atm), with any water formed remaining as vapor.

3.2 Definitions of Terms Specific to This Standard:

3.2.1 *corrected temperature rise*—the calorimeter temperature change caused by the process that occurs inside the bomb corrected for various effects.

3.2.2 *heat capacity*—the energy required to raise the temperature of the calorimeter one arbitrary unit.

3.2.2.1 *Discussion*—The heat capacity can also be referred to as the energy equivalent or water equivalent of the calorimeter.

4. Summary of Test Method

4.1 The heat capacity of the calorimeter is determined by burning a specified mass of benzoic acid in oxygen. A comparable amount of the analysis sample is burned under the same conditions in the calorimeter. The calorific value of the analysis sample is computed by multiplying the corrected temperature rise, adjusted for extraneous heat effects, by the heat capacity and dividing by the mass of the sample.

NOTE 1—Oxidation of coal after sampling can result in a reduction of calorific value. This effect increases with decreasing rank. Unnecessary

exposure of the samples to air from the time of sampling or delay in analysis should be avoided.

5. Significance and Use

5.1 The gross calorific value can be used to compute the total calorific content of the quantity of coal or coke represented by the sample for payment purposes.

5.2 The gross calorific value can be used for computing the calorific value versus sulfur content to determine whether the coal meets regulatory requirements for industrial fuels.

5.3 The gross calorific value can be used to evaluate the effectiveness of beneficiation processes.

5.4 The gross calorific value can be required to classify coals according to Classification D 388.

6. Apparatus and Facilities

6.1 *Test Area*—An area free from drafts, shielded from direct sunlight and other radiation sources. Thermostatic control of room temperature and controlled relative humidity are desirable.

6.2 *Combustion Bomb*—Constructed of materials that are not affected by the combustion process or the products formed to introduce measurable heat input or alteration of end products. The bomb shall be designed so that all liquid combustion products can be completely recovered by washing the inner surfaces. There shall be no gas leakage. The bomb shall be capable of withstanding a hydrostatic pressure test to 20 MPa (3000 psig) at room temperature without stressing any part beyond its specified elastic limit.

6.3 *Balance*—A laboratory balance capable of weighing the analysis sample to the nearest 0.0001 g. The balance shall be checked weekly, at a minimum, for accuracy.

6.4 *Calorimeter Vessel*—Made of metal with a tarnish-resistant coating, with all outer surfaces highly polished. Its size shall be such that the bomb is completely immersed in water during a determination. A stirrer shall be provided for uniform mixing of the water. The immersed portion of the stirrer shall be accessible to the outside through a coupler of low thermal conductivity. The stirrer speed shall remain constant to minimize any temperature variations due to stirring. Continuous stirring for 10 min shall not raise the calorimeter temperature more than 0.01°C when starting with identical temperatures in the calorimeter, test area and jacket. For calorimeters having a bucket it can be a separate component or integral component of the bomb. The vessel shall be of such construction that the environment of the calorimeter's entire outer boundaries can be maintained at a uniform temperature.

6.5 *Jacket*—A container with the inner perimeter maintained at constant temperature $\pm 0.1^\circ\text{C}$ (isoperibol) or at the same temperature $\pm 0.1^\circ\text{C}$ as the calorimeter vessel (adiabatic) during the test. To minimize convection, the sides, top and bottom of the calorimeter vessel shall not be more than 10 mm from the inner surface of the jacket. Mechanical supports for the calorimeter vessel shall be of low thermal conductivity.

6.6 Thermometers:

6.6.1 *Automated Calorimeters*—Platinum resistance or linear thermistor thermometers shall be capable of measuring to the nearest 0.0001°C. Thermometer calibration shall be traceable to a recognized certifying agency.

6.6.2 Manual Calorimeters:

6.6.2.1 *Platinum Resistance or Linear Thermistor Thermometers* shall be capable of measuring to the nearest 0.0001°C. Thermometer calibration shall be traceable to a recognized certifying agency.

6.6.2.2 *Liquid-in-Glass Thermometers*—Conforming to the requirements for thermometers 56C, 116C, or 117C as prescribed in Specification E 1. Thermometers 56C shall be calibrated at intervals no larger than 2.0°C over the entire graduated scale. The maximum difference in correction between any two calibration points shall be no more than 0.02°C. Thermometers 116C and 117C shall be calibrated at intervals no larger than 0.5°C over the entire graduated scale. The maximum difference in correction between any two calibration points shall not be more than 0.02°C.

6.6.2.3 *Beckman Differential Thermometer*—(Glass enclosed scale, adjustable), having a range of approximately 6°C in 0.01°C subdivisions reading upward and conforming to the requirements for Thermometer 115C, as prescribed in Specification E 1. The thermometer shall be calibrated at intervals no larger than 1°C over the entire graduated scale. The maximum difference in the correction between any two calibration points shall be less than 0.02°C.

6.6.2.4 *Thermometer Accessories*—A magnifier is required for reading liquid-in-glass thermometers to one tenth of the smallest scale division. The magnifier shall have a lens and holder designed so as to minimize errors as a result of parallax.

6.7 *Sample Holder*—An open crucible of platinum, quartz, or base metal alloy. Before use in the calorimeter, heat treat base metal crucibles for a minimum of 4 h at 500°C to ensure the crucible surface is completely oxidized. Base metal alloy crucibles are acceptable, if after three preliminary firings, the weight does not change by more than 0.0001 g.

6.8 *Ignition Fuse*—Ignition fuse of 100-mm length and 0.16-mm (No. 34 B&S gage) diameter or smaller. Nickel-chromium alloy (Chromel C) alloy, cotton thread, or iron wire are acceptable. Platinum or palladium wire, 0.10-mm diameter (No. 38 B&S gage), can be used provided constant ignition energy is supplied. Use the same type and length (or mass) of ignition fuse for calorific value determinations as used for standardization.

6.9 *Ignition Circuit*—A 6- to 30-V alternating or direct current is required for ignition purposes. A step-down transformer connected to an alternating current circuit, capacitors, or batteries can be used. For manually operated calorimeters, the ignition circuit switch shall be of the momentary double-contact type, normally open except when held closed by the operator. An ammeter or pilot light can be used in the circuit to indicate when current is flowing.

6.10 *Controller*—For automated calorimeters, capable of charging the bomb; filling the calorimeter vessel; firing the ignition circuit; recording calorimeter temperatures before, during, and after the test; recording the balance weights; and carrying out all necessary corrections and calculations.

6.11 *Crucible Liner*—Quartz fiber or alundum for lining the crucible to promote complete combustion of samples that do not burn completely during the determination of the calorific value.

7. Reagents

7.1 *Reagent Water*—Conforming to conductivity requirements for Type II of Specification D 1193 for preparation of reagents and washing of the bomb interior.

7.2 *Purity of Reagents*—Use reagent grade chemicals conforming to the specification of the Committee on Analytical Reagents of the American Chemical Society in all tests.⁷

7.3 *Benzoic Acid—Standard (C₆H₅COOH)*—Pellets made from benzoic acid available from the National Institute of Standards and Technology (NIST) or benzoic acid calibrated against NIST standard material. The calorific value of benzoic acid, for use in the calibration calculations, shall be traceable to a recognized certificate value.

7.4 *Oxygen*—Manufactured from liquid air, guaranteed to be greater than 99.5 % pure, and free of combustible matter. Oxygen made by the electrolytic process contains small amounts of hydrogen rendering it unfit unless purified by passage over copper oxide at 500°C.

7.5 *Titration Indicator*—Methyl orange, methyl red, or methyl purple for indicating the end point when titrating the acid formed during combustion. The same indicator shall be used for both calibration and calorific value determinations.

7.6 *Standard Solution*—Sodium carbonate (Na₂CO₃) or other suitable standard solution. Dissolve 3.76 g of sodium carbonate, dried for 24 h at 105°C in water, and dilute to 1 L. One millilitre of this solution is equivalent to 4.2 J (1.0 calorie) in the acid titration.

8. Hazards

8.1 The following precautions are recommended for safe calorimeter operation. Additional precautions are noted in Practice E 144. Also consult the calorimeter equipment manufacturer's installation and operating instructions before using the calorimeter.

8.1.1 The mass of sample and any combustion aid as well as the pressure of the oxygen admitted to the bomb shall not exceed the bomb manufacturer's specifications.

8.1.2 Inspect the bomb parts carefully after each use. Replace cracked or significantly worn parts. Replace O-rings and valve seats in accordance with manufacturer's instruction. For more details, consult the manufacturer.

8.1.3 Equip the oxygen supply cylinder with an approved type of safety device, such as a relief valve, in addition to the needle valve and pressure gage used in regulating the oxygen feed to the bomb. Valves, gages, and gaskets shall meet industry safety codes. Suitable reducing valves and adapters for 3- to 4-MPa (300- to 500-psig) discharge pressure can be obtained from commercial sources of compressed gas equipment. Check the pressure gage annually for accuracy or after any accidental over pressures that reach maximum gage pressure.

8.1.4 During ignition of a sample, the operator shall not extend any portion of the body over the calorimeter.

8.1.5 Do not fire the bomb if the bomb has been dropped or turned over after loading.

8.1.6 Do not fire the bomb if there is evidence of gas leakage when the bomb is submerged in the calorimeter vessel.

8.1.7 For manually operated calorimeters, the ignition switch shall be depressed only long enough to fire the charge.

9. Sample

9.1 The analysis sample is the material pulverized to pass 250- μ m (No. 60) sieve, prepared in accordance with either Practice D 346 for coke or Method D 2013 for coal.

10. Determination of the Heat Capacity of the Calorimeter

10.1 *Sample*—Weigh 0.8 to 1.2 g of benzoic acid into a sample holder. Record sample weight to the nearest 0.0001 g.

10.2 *Preparation of Bomb:*

10.2.1 Rinse the bomb with water to wet internal seals and surface areas of the bomb or precondition the calorimeter according to the manufacturer's instructions. Add 1.0 mL of water to the bomb before assembly.

10.2.2 Connect a measured fuse in accordance with manufacturer's guidelines.

10.2.3 Assemble the bomb. Admit oxygen to the bomb to a consistent pressure of between 2 and 3 MPa (20 and 30 atm). The same pressure is used for each heat capacity run. Control oxygen flow to the bomb so as not to blow material from the sample holder. If the pressure exceeds the specified pressure, detach the filling connection and exhaust the bomb. Discard the sample.

10.3 *Preparation of Calorimeter:*

10.3.1 Fill the calorimeter vessel with water at a temperature not more than 2°C below room temperature and place the assembled bomb in the calorimeter. Check that no oxygen bubbles are leaking from the bomb. If there is evidence of leakage, remove and exhaust the bomb. Discard the sample.

10.3.2 The mass of water used for each test run shall be $M \pm 0.5$ g where M is a fixed mass of water. Devices used to supply the required mass of water on a volumetric basis shall be adjusted when necessary to compensate for change in the density of water with temperature.

10.3.3 With the calorimeter vessel positioned in the jacket start the stirrers.

10.4 *Temperature Observations Automated Calorimeters:*

10.4.1 *Stabilization*—The calorimeter vessel's temperature shall remain stable over a period of 30 s before firing. The stability shall be $\pm 0.001^\circ\text{C}$ for an adiabatic calorimeters and $\pm 0.001^\circ\text{C/s}$ or less for an isoperibol calorimeter.

10.4.2 *Extrapolation Method*—Fire the charge, record the temperature rise. The test can be terminated when the observed thermal curve matches a thermal curve which allows extrapolation to a final temperature with a maximum uncertainty of $\pm 0.002^\circ\text{C}$.

10.4.3 *Full Development Method*—Fire the charge and record the temperature rise until the temperature has stabilized for a period of 30 s in accordance with the stability requirements specified in 10.4.1.

10.5 *Temperature Observations Manual Calorimeters:*

10.5.1 When using ASTM Thermometers 56C, estimate all

⁷ *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. Pharmacopeial Convention, Inc. (USPC), Rockville, MD.

readings to the nearest 0.002°C. When using ASTM Thermometers 115C, 116C, or 117C, estimate readings to 0.001°C and 25-Ω resistance thermometer readings to the nearest 0.0001 Ω. Tap or vibrate mercury thermometers just before reading to avoid errors caused by mercury sticking to the walls of the capillary.

10.5.2 Allow 5 min for the temperature of the calorimeter vessel to stabilize. Adjust the jacket temperature to match the calorimeter vessel temperature within 0.01°C and maintain for 3 min.

10.5.3 Fire the charge. Record the time as *a* and the temperature as *t_a*.

10.5.4 For adiabatic calorimeters adjust the jacket temperature to match that of the calorimeter vessel temperature during the period of the rise. Keep the two temperatures as equal as possible during the period of rapid rise. Adjust to within 0.01°C when approaching the final stabilization temperature. Record subsequent readings at intervals no greater than 1 min until three successive readings do not differ by more than ±0.001°C. Record the first reading after the rate of change has stabilized as the final temperature *t_c* and the time of this reading as *c*. For isoperibol calorimeters, when approaching the final stabilization temperature, record readings until three successive readings do not differ by more than 0.001°C per min. Record the first reading after the rate of change has stabilized as the final temperature as *t_c* and the time of this reading as *c*.

10.5.5 Open the calorimeter and remove the bomb. Release the pressure at a uniform rate such that the operation will not be less than 1 min. Open the bomb and examine the bomb interior. Discard the test if unburned sample or sooty deposits are found.

10.6 Thermochemical Corrections (see Appendix X1):

10.6.1 Acid Correction (see X1.1)—One may use either the titration (10.6.1.1) or calculated titration (10.6.1.2) procedure for coal and coke samples.

10.6.1.1 Titration Method—Wash the interior of the bomb with distilled water containing the titration indicator (see 7.5) until the washings are free of acid and combine with the rinse of the capsule. Titrate the washings with the standard solutions (see 7.6) using a titration indicator, or a pH or millivolt meter. The number of millilitres of standard Na₂CO₃ used in the titration shall be taken as *e1*.

10.6.1.2 Calculated Titration Method—Each calorimeter system shall be tested at several energy levels with benzoic acid pellets weighing 0.8, 1.0, and 1.2 g. This range corresponds to the optimum energy levels of 5000 through 8000 calories. Two runs shall be made at each weight. Plot millilitres of titrant (*y*) versus temperature rise, degree C (*x*), for each calibration and use linear regression to determine the formula for the line $y = m(x) + b$ (see Fig. 1). The resulting formula for a line is the equation for determining the calculated millilitres of titrant (*e1*). The calculated titrant = $m(x) + b$ where *m* and *b* have been determined by linear regression. The temperature rise (*x*) for each test shall be plotted on the graph to determine the calculated acid correction (*e1*) or determined from the equation $y = mx + b$.

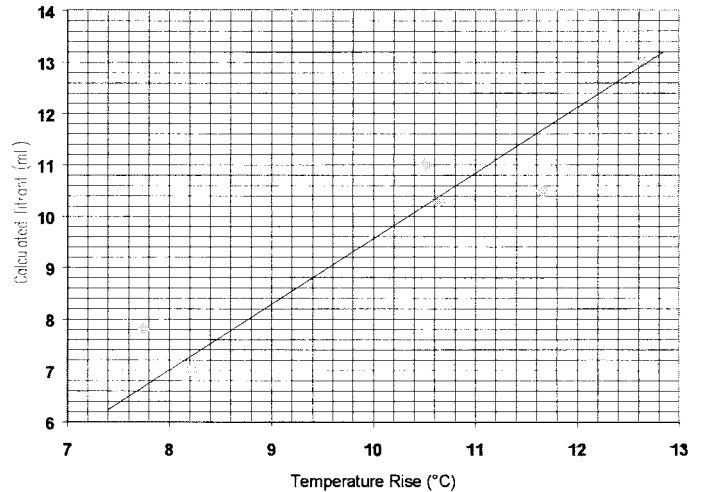


FIG. 1 Titration Versus Temperature Rise

Example:

Sample wt., g	Measured Titration (y)	Temperature Rise (x)
0.7643	7.8	7.7443
0.8104	7.0	8.2188
1.0392	11.0	10.5114
1.0506	10.3	10.6420
1.1539	10.5	11.6584
1.2562	13.0	12.6491

$$y = m(x) + b \tag{1}$$

Using regression analysis, the above data yield the following data: slope = 1.0826, intercept = -1.1496, and the equation for the millilitres of titrant = $y = 1.0826x - 1.1496$.

With any given temperature rise (*x*), the value *y* (*e1*) may be determined.

NOTE 2—*m* above represents the slope of the line, whereas in other references in this method *m* represents mass.

NOTE 3—Regression analysis to determine the equation for the millilitres of titrant (*e1*) is to be done without forcing the data through zero.

10.6.2 Fuse Correction (see X1.3)—Determine the fuse correction using one of the two alternatives:

10.6.2.1 Measure the combined pieces of unburned ignition fuse and subtract from the original length to determine the fuse consumed in firing according to Eq 2.

$$e2 = K_f \times l \tag{2}$$

where:

- e2* = the correction for the heat of combustion of the firing fuse,
- l* = the length of fuse consumed during combustion,
- K_f* = 0.96 J/mm (0.23 cal/mm) for No. 34 B&S gage Chromel C,
- K_f* = 1.13 J/mm (0.27 cal/mm) for No. 34 B&S gage iron wire, and
- K_f* = 0.00 J/mm for platinum or palladium wire provided the ignition energy is constant.

or;

10.6.2.2 Weigh the combined pieces of unburned fuse and subtract from the original weight to determine the weight in

milligrams of the fuse consumed in firing (m). Remove any ball of oxidized metal from the ends before weighing.

$$e2 = K_m \times m \quad (3)$$

where:

- $e2$ = the correction for the heat of combustion of the firing fuse,
- m = the weight in mg of fuse consumed during combustion,
- K_m = 5.9 J/mg (1.4 cal/mg) for No. 34 B&S gage Chromel C,
- K_m = 7.5 J/mg (1.8 cal/mg) for No. 34 B&S gage iron wire, and
- K_m = 0.00 J/mg for platinum or palladium wire provided the ignition energy is constant.

When cotton thread is used, employ the correction in J recommended by the instrument manufacturer.

10.7 Calculation of the Corrected Temperature Rise—Compute the corrected temperature rise, t , as follows:

$$t = t_c - t_a + C_e + C_r + C_s \quad (4)$$

where:

- t = corrected temperature rise, °C;
- t_a = initial temperature reading at time of firing;
- t_c = final temperature reading;
- C_e = thermometer, emergent stem correction (see Eq A1.4);
- C_r = radiation correction (see Eq A1.2); and
- C_s = thermometer setting correction (see Eq A1.3).

10.7.1 The temperature rise in isoperibol calorimeters require a radiation correction.

10.7.2 Beckman differential thermometers require a setting correction and an emergent stem correction.

10.7.3 Solid-stem ASTM Thermometers 56C do not require emergent stem corrections if all tests are performed within the same 5.5°C interval. If the operating temperature range is beyond this limit, a differential emergent stem correction shall be applied.

10.8 Calculation of the Heat Capacity—Calculate the heat capacity (E) of the calorimeter using the following equation:

$$E = [(H_c \times m) + e1 + e2]/t \quad (5)$$

where:

- E = the calorimeter heat capacity, J/°C;
- H_c = heat of combustion of benzoic acid, as stated in the certificate, J/g;
- m = mass of benzoic acid, g;
- $e1$ = acid correction from 10.6.1 from either the titration method (10.6.1.1) or the calculated titration (10.6.1.2);
- $e2$ = fuse correction from 10.6.2, J; and
- t = corrected temperature rise from 10.7, °C.

10.8.1 Using the procedures described in 10.1-10.8 complete a total of ten acceptable test runs. An individual test shall be rejected only if there is evidence of incomplete combustion.

10.8.2 The relative standard deviation of the heat capacity of ten acceptable test runs shall be 0.15 % or less of the average energy equivalent. If after considering the possibility of outliers using criterion established in Practice E 178, this limit is not met, one should review operation of the calorimeter for any assignable cause which should be corrected before repeating

the series. Table 1 summarizes a series of heat capacity runs. This table would be applicable regardless of the unit of measure for the heat capacity.

11. Heat Capacity Checks

11.1 The heat capacity value shall be checked a minimum of once a month, after changing any part of the calorimeter, or after changing the oxygen supply. Two procedures are available for heat capacity check: Standard Method and Rolling Average Method.

NOTE 4—Although it is only required to check the heat capacity once a month, this may be inadequate. A more frequent check of heat capacity values is recommended for laboratories making a large number of tests on a daily basis. The frequency of the heat capacity check should be determined to minimize the number of tests that would be affected by an undetected shift in the heat capacity values.

11.2 Standard Method:

11.2.1 A single new heat capacity test value shall not differ from the existing heat capacity value by more than ± 0.17 %. If this requirement is met, the existing heat capacity value is acceptable. For example: existing heat capacity value is 2402 cal/°C. $2402 \text{ cal/°C} \times 0.0017 = 4.1 \text{ cal/°C}$. If single test value is within 4.1 cal/°C of the 2402 cal/°C value, then the existing heat capacity value is still acceptable.

11.2.2 If the requirements given in 11.2.1 are not met, an additional heat capacity test shall be run. The difference between the two new heat capacity values shall not exceed 0.21 % of the existing heat capacity value. The average of the two new heat capacity values shall not differ from the existing heat capacity value by more than ± 0.13 %. If this requirement is met, the existing heat capacity value is acceptable.

11.2.3 If the requirements given in 11.2.2 are not met, two more heat capacity tests shall be run. The range of the four new test values shall not exceed 0.33 % of the existing heat capacity value. The average of the four new heat capacity values shall not differ from the existing heat capacity value by more than ± 0.08 %. If this requirement is met, the existing heat capacity value is acceptable.

11.2.4 If the requirements given in 11.2.3 are not met, a fifth and sixth heat capacity test shall be run. The range of the six new test values shall not exceed 0.42 % of the existing heat

TABLE 1 Heat Capacity Runs

Note—Variance = $s^2 = \{(\text{Sum Column C} - [(\text{Sum Column B})^2/10])\}/9 = 89.51$.
Standard Deviation = $s = \sqrt{s^2} = 9.46$.
Relative Standard Deviation = $(s/\text{Average}) \times 100 = 0.09$ %.

Run Number	Column A Heat Capacity, J/°C	Column B Difference From Average	Column C (Column B) ²
1	10 257.7	+4.2	17.6
2	10 249.3	-4.2	17.6
3	10 270.2	+16.7	278.9
4	10 253.5	0.0	0
5	10 245.1	-8.4	70.6
6	10 249.3	-4.2	17.6
7	10 240.9	-12.6	158.8
8	10 266.0	+12.5	156.3
9	10 257.7	+4.2	17.6
10	10 245.1	-8.4	70.6
SUM		-0.2	805.6
AVERAGE	E = 10 253.5		

capacity value. The average of the six new heat capacity values shall not differ from the existing heat capacity value by more than ±0.08 %. If these requirements are met, do not change the existing heat capacity value.

11.2.5 If the requirements given in 11.2.4 are not met, four more heat capacity tests shall be run. The range of the ten new test values shall not exceed 0.50 % of the existing heat capacity value. The average of the ten new heat capacity values shall not differ from the existing heat capacity value by more than ±0.04 %. If this requirement is met, the existing heat capacity value is acceptable.

11.2.6 If requirements given in 11.2.5 are not met, the average value from the ten new test values shall be used to determine a new heat capacity value provided the relative standard deviation of the ten values does not exceed 0.15 %.

11.2.7 The summary of the numerical requirements at each step in checking the heat capacity is given in Table 2.

11.3 Rolling Average Method:

11.3.1 A single new heat capacity value shall not differ from the existing heat capacity value by more than ±0.17 %.

11.3.2 Values that serve to confirm existing heat capacity values will be included with the original 10 calibration tests until a total of 20 tests are made. These tests will comprise a database for calculating the mean heat capacity value provided the relative standard deviation of the 20 values does not exceed 0.15 %. Any new calibration check beyond the 20 tests will replace the oldest value in the heat capacity database of 20 tests. A maximum relative standard deviation of 0.15 % shall be maintained for the heat capacity database of 20 tests.

11.3.3 When using a rolling average, the heat capacity data must be continually evaluated for four indications of out-of-control conditions: first, seven consecutive results are higher than the mean; second, seven consecutive results are lower than the mean; third, seven consecutive results are continually increasing; and fourth, seven consecutive results are continually decreasing.

11.3.4 Out-of-control trends indicate that the calorimeter operation is suspect and causes should be identified. Whether or not causes are identified, the calorimeter should be recalibrated according to the procedure in Section 10.

12. Procedure for Coal and Coke Samples

12.1 Weigh 0.8 to 1.2 g of sample into a sample holder. Record the weight to the nearest 0.0001 g (see 12.6.3).

12.2 Follow the procedures as described in 10.2-10.5 for determination of heat capacity. For the calorific value of coke, it is necessary to use 3-MPa (30-atm) pressure for both

standardization and analysis. The starting temperature for determinations shall be within ±0.5°C of that used in the determination of the heat capacity.

12.3 Carry out a moisture determination in accordance with Test Method D 3173 or Test Methods D 5142 on a separate portion of the analysis sample preferably on the same day but not more than 24 h apart from the calorific value determination so that reliable corrections to other bases can be made.

12.4 Conduct the sulfur analysis in accordance with Test Methods D 3177 or D 4239. From the weight % sulfur, calculate the sulfur corrections (see X1.2):

$$e3 = 55.2 \text{ J/g} \times S \times m \text{ (13.18 cal/g} \times S \times m) \tag{6}$$

where:

e3 = a correction for the difference between the heat of formation of H₂SO₄ from SO₂ with respect to the formation of HNO₃, J;

S = wt % sulfur in the sample; and

m = mass of sample from 12.1, g.

12.4.1 When titration method is used (see 10.6.1.1), the sulfur correction is

$$e3 = 55.18 \text{ J/g} \times S \times m \text{ or (13.18 cal/g} \times S \times m) \tag{7}$$

12.4.2 When the calculated titration method is used (see 10.6.1.2), the sulfur correction is

$$e3 = 94.51 \text{ J/g} \times S \times m \text{ or (22.57 cal/g} \times S \times m) \tag{8}$$

12.5 For eight mesh samples, analyze coals susceptible to oxidation within 24 h of preparation.

12.6 Coal or coke that do not burn completely can be treated as follows:

12.6.1 Use a crucible liner of the type recommended in 6.11.

12.6.2 Use a combustion aid such as benzoic acid, ethylene glycol, mineral oil or a gelatin capsule. A minimum of 0.4 g of combustion aid shall be used. Record the weight to the nearest 0.0001 g. Calculate the correction for use of a combustion aid using the following:

$$e4 = Ha \times ma \tag{9}$$

where:

e4 = correction for use of a combustion aid,

Ha = heat of combustion of the combustion aid J/g (cal/g), and

ma = mass of combustion aid, g.

12.6.3 Vary the mass of the sample to obtain good ignition and so that the total heat generated is the same as the heat generated during calibration.

13. Calculation

13.1 Gross Calorific Value—Calculate the gross calorific value *Q_{vad}*(gross) using the following equation:

$$Q_{vad}(\text{gross}) = [(tE_e) - e1 - e2 - e3 - e4]/m \tag{10}$$

where:

Q_{vad}(gross) = gross calorific value at constant volume as determined, J/g (cal/g);

E_e = the heat capacity of the calorimeter, J/°C (cal/°C);

t = corrected temperature rise according to 10.7, °C;

TABLE 2 Numerical Requirements

Number of Check Runs	Maximum Range ((<i>E_{nmax}</i> - <i>E_{nmin}</i>)/ <i>E_e</i>) × 100	Maximum Difference ((<i>E_{nnav}</i> - <i>E_e</i>)/ <i>E_e</i>) × 100
1	...	±0.17
2	0.21	±0.13
4	0.33	±0.08
6	0.42	±0.08
10	0.50	±0.04

E_e is the existing heat capacity value.

E_{nmin} is the minimum reading in group of heat capacity check runs.

E_{nmax} is the maximum reading in a group of heat capacity check runs.

E_{nnav} is the average of the group of heat capacity check runs.

- e1 = acid correction according to 10.6.1, J;
- e2 = fuse correction according to 10.6.2, J;
- e3 = sulfur correction determined according to 12.4, J;
- e4 = combustion aid correction determined according to 12.7, J; and
- m = mass of the sample, g.

13.1.1 See X1.2.3 for an example calculation.

13.2 *Net Calorific Value*—Calculate the net calorific value $Q_p(\text{net})$ as follows:

$$Q_p(\text{net}) = Q_{\text{var}}(\text{gross}) - 215.5 \text{ J/g} \times H_{\text{ar}} \quad (11)$$

or:

$$(Q_p(\text{net}))_{\text{ar}} = Q_{\text{var}}(\text{gross}) - 92.67 \text{ Btu/lb} \times H_{\text{ar}} \quad (12)$$

where:

- $Q_p(\text{net})$ = net calorific value, at constant pressure;
- $Q_{\text{var}}(\text{gross})$ = gross calorific value, at constant volume, as-received basis; and
- H_{ar} = total hydrogen, %, as-received basis, where hydrogen includes hydrogen in the sample moisture.

Example:

Calorific value as determined $Q_{\text{vad}}(\text{gross}) = 31\,420 \text{ J/g}$
 Moisture, as determined $M_{\text{ad}} = 2.13 \text{ wt } \%$
 Moisture, as received $M_{\text{ar}} = 8.00 \text{ wt } \%$
 Hydrogen, as determined $H_{\text{ad}} = 5.00 \text{ wt } \%$
 $Q_{\text{var}}(\text{gross}) = Q_{\text{vad}}(\text{gross}) \times [(100 - M_{\text{ar}})/(100 - M_{\text{ad}})] = 31\,420 \text{ J/g} \times [(100 - 8.00)/(100 - 2.13)] = 29\,535 \text{ J/g}$
 $H_{\text{ar}} = [(H_{\text{ad}} - 0.1119 \times M_{\text{ad}}) \times \{(100 - M_{\text{ar}})/(100 - M_{\text{ad}})\}] + 0.1119 M_{\text{ar}} = [(5.00 - 0.1119 \times 2.13) \times \{(100 - 8.00)/(100 - 2.13)\}] + 0.1119 \times 8.0 = 5.37$
 $Q_p(\text{net}) = 29\,535 \text{ J/g} - (215.5 \times 5.37) = 29\,535 \text{ J/g} - 1153 \text{ J/g} = 28\,388 \text{ J/g}$

14. Report

14.1 Report the calorific value as $Q_{\text{vad}}(\text{gross})$ along with the moisture of the sample as determined M_{ad} from 12.3.

14.2 The results of the calorific value can be reported in any of a number of bases differing in the manner the moisture is treated. Procedures for converting the value obtained on an analysis sample to other bases are described in Practice D 3180.

15. Precision and Bias

15.1 Manual Calorimeters:

15.1.1 The relative precision of this test method for the gross calorific value covers the range from 29,535 to 33,721 J/g (12,700 to 14,500 Btu/lb) for bituminous coals and from 20,442 to 29,651 J/g (8,790 to 12,750 Btu/lb) for subbituminous and lignite coals.

15.1.2 *Repeatability*—The difference in absolute value between two test results calculated to a dry basis (Practice D 3180) performed on two separate test portions of the same analysis sample of 250- μm (No. 60) coal in the same laboratory, by the same operator, using the same equipment with the same heat capacity value shall not exceed the repeatability interval $I(r)$ of 115 J/g (50 Btu/lb) more than 5 % of such paired values (95 % confidence level). When such a difference

is found to exceed the repeatability interval, there is reason to question one or both of the test results.

15.1.3 *Repeatability Limit (r)*—the value below which the absolute difference between two test results of separate and consecutive test determinations, carried out on the same sample in the same laboratory by the same operator, using the same apparatus on a single test specimen of two separate 2.36-mm (No. 8) test units of coal reduced entirely to 250- μm (No. 60) on samples taken at random from a single quantity of homogeneous material, may be expected to occur with a probability of approximately 95 %. The repeatability limit for this test method on a dry basis is:

Bituminous coals	160 J/g (69 Btu/lb)
Subbituminous and lignite coals	140 J/g (60 Btu/lb)

15.1.4 *Reproducibility*—The difference in absolute value between test results calculated to a dry basis (Practice D 3180) performed in different laboratories on representative analysis samples of 250- μm (No. 60) coal shall not exceed the reproducibility interval $I(r)$ of 250 J/g (100 Btu/lb) more than 5 % of such paired values (95 % confidence level). When such a difference is found to exceed the reproducibility interval there is reason to question one or both of the test results.

15.1.5 *Reproducibility Limit (R)*—the value below which the absolute difference between two test results, carried out in different laboratories using single test samples of two separate 2.36-mm (No. 8) test units of coal reduced entirely to 250- μm (No. 60) and prepared from samples taken at random from a single quantity of material that is as nearly homogeneous as possible, may be expected to occur with a probability of 95 %. The reproducibility limit for this test method on a dry basis is:

Bituminous coals	249 J/g (107 Btu/lb)
Subbituminous and lignite coals	326 J/g (140 Btu/lb)

NOTE 5—Supporting data for 2.36-mm (No. 8) coal has been filed at ASTM Headquarters and may be obtained by requesting RR:DO5-11015.

15.1.6 *Bias*—Bias in the determination of the gross calorific value is eliminated provided samples are treated identically to the benzoic acid used in the determination of the calorimeter heat capacity.

15.2 Microprocessor Controlled Automated Calorimeters:

15.2.1 *Precision*—The relative precision of this test method for the determination of gross calorific value covers the range from 26 279 to 34 186 J/g (11 300 to 14 700 Btu/lb) for bituminous coals, 21 860 to 30 000 J/g (9 400 to 12 900 Btu/lb) for subbituminous and lignitic coals, and 30 628 J/g (13 170 Btu/lb) for the coke value as based upon the analysis of only one coke sample.

15.2.2 *Repeatability Limit (r)*—The value below which the absolute difference between two test results of separate and consecutive test determinations, carried out on the same sample in the same laboratory by the same operator using the same apparatus on samples taken at random from a 250- μm (No. 60 sieve) single sample of homogeneous material may be expected to occur with a probability of approximately 95 %. The repeatability limit for this test method on a dry basis is as follows:

Bituminous coals	149 J/g (64 Btu/lb)
Subbituminous and lignitic coals	193 J/g (83 Btu/lb)
Coke	479 J/g (206 Btu/lb)

15.2.3 *Reproducibility Limit (R)*—The value below which the absolute difference between two test results, carried out in different laboratories using samples taken at random from a 250-um (No. 60 sieve) single quantity of material that is as nearly homogeneous as possible, may be expected to occur with a probability of 95 %. The reproducibility limit for this test method on a dry basis is as follows (see Note 6):

Bituminous coals	256 J/g (110 Btu/lb)
Subbituminous and lignitic coals	381 J/g (164 Btu/lb)
Coke	928 J/g (399 Btu/lb)

NOTE 6—Calorific values have not been determined for anthracite coals.

NOTE 7—Precision statements have not been determined for No. 8 sieve size coal.

16. Keywords

16.1 adiabatic calorimeter; bomb calorimeters; calorific value; calorimeter; coal; coke; isoperibol bomb calorimeter

ANNEX

(Mandatory Information)

A1. THERMOMETRIC CORRECTIONS

A1.1 *Thermometer Corrections*—The following corrections shall be made:

A1.1.1 *Calibration Correction*, shall be made in accordance with the calibration certificate furnished by the calibration authority.

A1.1.2 *Radiation Corrections*—Radiation corrections are required to calculate heat loss or gain to the isoperibol water jacket. They are based on the Dickinson formula,⁸ the Regnault-Pfaundler formula,⁹ or the U.S. Bureau of Mines method.¹⁰ The same method of determining the radiation correction shall be used consistently in the determination of heat capacity and sample measurements.

A1.1.2.1 *Dickinson Formula*:

$$C_r = -r_1 \times (b - a) - r_2 \times (c - b) \quad (\text{A1.1})$$

where:

C_r = radiation correction;

r_1 = rate of rise in temperature per minute in the preliminary period;

r_2 = rate of rise of temperature per minute in the final period (if temperature is falling, r_2 is negative);

t_a = firing temperature;

t_c = final temperature, being the first temperature after which the rate of change is constant;

a = time at temperature t_a , min;

b = time at temperature $t_a + 0.60 (t_c - t_a)$, min; and

c = time at temperature t_c , min.

A1.1.2.2 *Regnault-Pfaundler Formula*:

$$C_r = nr_1 + kS \quad (\text{A1.2})$$

where:

C_r = radiation correction,

n = number of minutes in the combustion period,

$$k = (r_1 - r_2) / (t'' - t'), \quad (\text{A1.3})$$

$$S = tn - 1 + (1/2)(ti + tf)nt' \quad (\text{A1.4})$$

t' = average temperature during the preliminary period,

t'' = average temperature during the final period,

r_1, r_2 see A1.1.2.1,

t_1, t_2, \dots, t_n = successive temperature recorded during the combustion period, at 1-min intervals, and

$$tn - 1 = \text{sum of } t_1, t_2, t_3 \dots t_{n-1} \quad (\text{A1.5})$$

A1.1.2.3 *Bureau of Mines Method*—A table of radiation corrections can be established so that only the initial and final readings are required to determine the calorific value of a fuel. This can be done by carrying out a series of tests using the procedure described in Section 10, using the following conditions. Regulate the amount of sample burned so that a series of determinations is made in which different temperature rises are obtained. For all determinations, keep the water jacket temperature constant, fire the bomb at the same initial temperature, and have the same time, $c - a$, elapse (± 2 s) between the initial and final readings. Determine the radiation corrections for each of the series of temperature rises using the Dickinson method (see A1.1.2.1), or the Regnault-Pfaundler method (see A1.1.2.2). These corrections are constant for a given temperature rise. From the series of readings, a table or graph is plotted to show radiation correction versus temperature rise. Once the table or graph is established, the radiation corrections can be obtained from it until there is a major change in the equipment.

A1.1.3 *Setting Correction*

This is necessary for the Beckman thermometer. It shall be made in accordance with the directions furnished by the calibration authority.

A1.1.4 *Differential Emergent Stem Correction*

The calculation of differential stem correction depends upon the way the thermometer was calibrated and how it is used. Two conditions are possible.

A1.1.4.1 *Thermometers Calibrated in Total Immersion and Used in Partial Immersion*

This emergent stem correction is made as follows:

$$C_e = K (tf - ti)(tf + ti - L - T) \quad (\text{A1.6})$$

⁸ Dickinson, H. C., *Bulletin*, U.S. Bureau of Standards, Vol. 11, 1951, p. 189.

⁹ Pfaundler, L., *Annalen der Physik* (Leipzig), ANPYA, Vol. 129, 1966, p. 102.

¹⁰ "Methods of Analyzing & Testing Coal and Coke," U.S. Bureau of Mines *Bulletin* 638, XMBUA, 1967, pp. 16–17.

where:

C_e = emergent stem correction,
 K = 0.000 16 for thermometers calibrated in °C,
 L = scale reading to which the thermometer was immersed,
 T = mean temperature of emergent stem,
 t_i = initial temperature reading, and
 t_f = final temperature reading.

Example:

A thermometer was immersed to 16°C; its initial reading, t_i , was 24.127°C; its final reading, t_f , was 27.876; the mean temperature of the emergent stem, T , was 26°C.

$$C_e = 0.000\ 16 \times (28 - 24) \times (28 + 24 - 16 - 26) = 0.0064^\circ\text{C} \quad (\text{A1.7})$$

A1.1.4.2 Thermometers Calibrated and Used in Partial Immersion, But at a Different Temperature Than the Calibrated Temperature:

$$C_e = K (t_f - t_i)(t_c - t_o) \quad (\text{A1.8})$$

where:

C_e = emergent stem correction,
 K = 0.000 16 for thermometers calibrated in °C,
 t_i = initial temperature reading,
 t_f = final temperature reading,
 t_o = observed stem temperature, and
 t_c = stem temperature at which the thermometer was calibrated.

Example:

A thermometer has an initial reading, t_i , 27°C; a final reading, t_f , 30°C; the observed stem temperature, t_o , 28°C; and the calibration temperature, t_c , 22°C.

$$C_e = 0.000\ 16 \times (30 - 27) \times (28 - 22) = 0.003^\circ\text{C} \quad (\text{A1.9})$$

APPENDIX

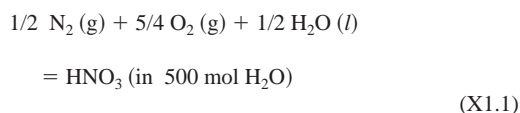
(Nonmandatory Information)

X1. THERMOCHEMICAL CORRECTIONS

X1.1 Energy of Formation of Nitric Acid (HNO_3):

X1.1.1 A correction, e_1 , (10.6.1) is applied for the formation of nitric acid.

X1.1.2 (1) HNO_3 is formed in the calorimeter by the following reaction:



X1.1.3 (2) the energy of formation of HNO_3 in approximately 500 mol of water under bomb conditions is minus 59.0 kJ/mol (14.09 Kcal/mole).¹¹

X1.1.4 Normal convention assigns a negative value for a heat of formation that is exothermic. By definition, heat released from combustion processes are expressed as positive values. Hence, the negative factors developed for nitric and sulfuric acid corrections are expressed as positive values in the calculations.

X1.1.5 A convenient concentration of Na_2CO_3 is 3.76-g $\text{Na}_2\text{CO}_3/\text{L}$ which gives $e_1 = V$ where V is the volume of Na_2CO_3 in millilitres. When H_2SO_4 is produced during the combustion of coal or coke, a part of the correction for H_2SO_4 is present in the e_1 correction. The remainder is in the e_3 correction (see X1.2).

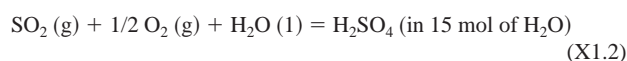
X1.2 Energy of Formation of Sulfuric Acid (H_2SO_4)—By definition (see Terminology D 121), the gross calorific value is obtained when the product of the combustion of sulfur in the sample is $\text{SO}_2(\text{g})$. However, in actual bomb combustion

processes, all the sulfur is found as H_2SO_4 in the bomb washings.

X1.2.1 A correction e_3 is applied for the sulfur that is converted to H_2SO_4 . This correction is based upon the energy of formation of H_2SO_4 in solutions, such as will be present in the bomb at the end of a combustion from SO_2 . This energy is taken as -297.2 kJ/mol.¹²

X1.2.2 When the bomb washings are titrated, a correction of 2×59.7 kJ/mole of sulfur is applied in the e_1 correction so that the additional correction that is necessary is the difference in the heats of formation for nitric and sulfuric acid and this correction is $-297.2 - (-2 \text{ times } 59.7) = -177.8$ kJ/mol, or -55.45 J/g (13.2 cal/g) of sulfur times the weight of sample in grams times percent sulfur in sample.

X1.2.3 If a 1-g sample is burned, the resulting H_2SO_4 condensed with water formed on the walls of the bomb will have a ratio of about 15 mol of water to 1 mol of H_2SO_4 . For this concentration, the energy of the reaction under the conditions of the bomb process is -303 kJ/mole.



X1.2.4 The values above are based on a sample containing approximately 5 % sulfur and approximately 5 % hydrogen. The assumption is also made that the H_2SO_4 is dissolved entirely in the water condensed during combustion of the sample.¹³

X1.2.5 For different sample weights or sulfur content, or

¹¹ Mott, R. A., and Parker, C., "Studies in Bomb Calorimetry IV—Corrections," Fuel, FUELB, Vol. 34, 1955, p. 303–316.

¹² The NBS Tables of Chemical Thermodynamic Properties, Selected Values for Inorganic and C_1 and C_2 Organic Substances in SI Units, *Journal of Phys. Chem. Ref. Data*, 11, Supplement No. 2, 1985, 392 pp.

¹³ Mott, R. A., and Parker, C., "Studies in Bomb Calorimetry IX—Formation of Sulfuric Acid," *Fuel, Fuel B*, Vol. 37, 1958, p. 371.

both, the resultant normality of acid formed can be different, and therefore, the normality of titrant must be adjusted accordingly. Basing the calculation upon a sample of comparatively large sulfur content reduces the possible overall errors, because, for small percentages of sulfur, the correction is smaller.

X1.3 Fuse Correction—The energy required to melt a platinum or palladium wire is constant for each experiment if the same amount of platinum or palladium wire is used. As the energy is small, its effect is essentially cancelled out in the relationship between the standardization experiments and the calorific value determinations, and it can be neglected.

X1.4 Reporting Results in Other Units:

X1.4.1 The gross calorific value can be expressed in joules per gram, calories per gram, or British thermal units per pound. The relationships between these units are given in the table below:

1 Btu = 1055.06 J	1 J/g = 0.430 Btu/lb
1 calorie IT ¹⁴ = 4.1868 J	1 J/g = 0.239 cal/g
	1 cal/g = 1.8 Btu/lb

X1.5 Sample Calculations:

X1.5.1 Heat Capacity:

$$E = [(Hc \times m) + e1 + e2]/t$$

$$Hc = 26\,435 \text{ J/g,}$$

$$m = 1.0047 \text{ g,}$$

$$e1 = 43\text{-J acid correction,}$$

$$e2 = 55\text{-J fuse correction,}$$

$$t = 2.6006^\circ\text{C,}$$

$$E = [(26\,435 \text{ J/g} \times 1.0047 \text{ g}) + 43 \text{ J} + 55 \text{ J}]/2.6006^\circ\text{C, and}$$

$$E = 10\,250.4 \text{ J}^\circ\text{C.}$$

X1.6 Heat of Combustion:

$$Q_{\text{vad}} (\text{gross}) = [(tE) - e1 - e2 - e3 - e4]/m;$$

$$E_e = 10\,250.4 \text{ J}^\circ\text{C;}$$

$$t = 2.417^\circ\text{C;}$$

$$e1 = 77\text{-J acid correction;}$$

$$e2 = 52\text{-J fuse correction;}$$

$$e3 = 58 \times 1.24 \% \times 0.7423 \text{ g, sulfur correction;}$$

$$e4 = 46\,025 \text{ J/g} \times 0.2043 \text{ g, combustion aid correction;}$$

$$m = 0.7423 \text{ g, mass of sample;}$$

$$Q_{\text{vad}} = [(10\,250.4 \text{ J}^\circ\text{C} \times 2.417^\circ\text{C}) - 77 \text{ J} - 52 \text{ J} - 53 \text{ J} - 9403 \text{ J}]/0.7423 \text{ g; and}$$

$$Q_{\text{vad}} = 20\,464 \text{ J/g.}$$

¹⁴ Cohen, E. R., and Taylor, B. N., "The 1986 CODATA Recommended Values for the Fundamental Physical Constants," *Journal of Phys. Chem. Ref. Data*, Vol 17, No. 4, 1988, pp. 1795–1803.

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