
**Coal — Determination of caking power
— Gray-King coke test**

Charbon — Détermination du pouvoir agglutinant — Essai Gray-King

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Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

The procedures used to develop this document and those intended for its further maintenance are described in the ISO/IEC Directives, Part 1. In particular the different approval criteria needed for the different types of ISO documents should be noted. This document was drafted in accordance with the editorial rules of the ISO/IEC Directives, Part 2 (see www.iso.org/directives).

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights. Details of any patent rights identified during the development of the document will be in the Introduction and/or on the ISO list of patent declarations received (see www.iso.org/patents).

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For an explanation on the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the WTO principles in the Technical Barriers to Trade (TBT) see the following URL: [Foreword - Supplementary information](#)

The committee responsible for this document is ISO/TC 27, *Solid mineral fuels*, Subcommittee SC 05, *Methods of analysis*.

This third edition cancels and replaces the second edition (ISO 502:1982), which has been technically revised.

Introduction

The purpose of the Gray-King coke test, which is one of the parameters adopted for the International Classification of Hard Coal by Type by the United Nations Economic Commission for Europe, is to assess the caking properties of a type of coal or a blend of coals by carbonizing under standard conditions.

Although the Gray-King test and the Roga test both assess the caking properties of a coal, they do not measure precisely the same parameters and are not recommended to be regarded as alternative methods.

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Coal — Determination of caking power — Gray-King coke test

1 Scope

This International Standard specifies a method of assessing the caking power of coal under standard conditions.

2 Normative references

The following documents, in whole or in part, are normatively referenced in this document and are indispensable for its application. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

ISO 1014, *Coke — Determination of true relative density, apparent relative density and porosity*

3 Principle

The sample is heated under standard conditions to a final temperature of 600 °C. The coke residue obtained is classified by reference to a series of standard residues. If the coke residue produced is so swollen that it fills the cross-section of the retort tube, the determination is repeated with the coal admixed with a suitable quantity of electrode carbon or equivalent material. For these highly swelling coals, the Gray-King coke type is defined by the minimum amount of electrode carbon required to produce a strong hard coke residue of the same volume as the original coal and electrode carbon mixture.

4 Reagent

4.1 Standard electrode carbon (see [10.1](#))

High temperature electrode carbon:

Moisture	less than 1 %
Volatile matter	less than 1,5 %
Ash	less than 5 %
Bulk density at 25 °C (see Annex A)	1,00 g/cm ³ to 1,05 g/cm ³
Relative density at 25 °C (see 10.2)	2,05 to 2,09

Size analysis:

Retained on 212 µm test sieve	nil
Through 212 µm test sieve, retained on 125 µm test sieve	less than 26 %
Through 125 µm test sieve, retained on 63 µm test sieve	10 % to 40 %
Through 63 µm test sieve	50 % to 85 %

It is recommended that a dust mask is used while using the inert carbonaceous material which can contain undesirable trace elements from the original processing of this material.

5 Apparatus

5.1 Furnace, horizontal electric, 50 mm internal diameter and 300 mm long, with one end closed and the other carrying a plug of insulating material which is bored centrally with a hole 25 mm in diameter. The winding of the furnace shall be such that the middle 200 mm is at a uniform temperature within ± 5 °C at both 300 °C and 600 °C. Alternatively, the furnace may be constructed from an electrically-heated aluminium-bronze block, with one or several, bores of 25 mm diameter. The furnace shall be insulated and located in a cover of metal or other suitable material, and shall be equipped with a suitable thermocouple, lying above the retort tube when the latter is in position and with the junction at the centre of the furnace. An indicator shall be provided for showing the furnace temperature with an accuracy of ± 5 °C. A suitable means of controlling the energy input shall also be provided to permit an increase in temperature at a rate of 5 °C/min. A multiple tube furnace to allow simultaneous determinations is convenient. The furnace may be of the fixed type or mounted on rails. Suitable furnaces are shown in [Figure 2](#) and [Figure 3](#).

5.2 Retort tube (see [Figure 4](#)), a heat-resistant glass or transparent silica tube, 20 mm internal diameter and 300 mm long, closed at one end, with a side arm, 8 mm internal diameter and 50 mm long, sealed in at a distance of about 20 mm from the open end. The tube shall be smooth and either of uniform bore or with a slight taper (19 mm to 21 mm) such that the open end is the larger.

5.3 Distance rod, with a flat disk at one end to assist in the packing of the coal and to indicate the free end of the coal sample in the retort tube.

5.4 Receiver and outlet tube

A glass vessel of adequate size, suitably supported and attached to the side arm of the retort tube, fitted with an outlet tube leading to atmosphere or to a piece of small bore silica tubing at the end of which the gas leaving the receiver can be burned through Bunsen burner (in a fume cabinet) to ensure toxic fumes are burnt before venting to the atmosphere through the fume cabinet.

The receiver may conveniently be a U-tube which can be immersed in water.

The outlet shall be open to the atmosphere to prevent pressure build-up.

Fumes are toxic and it is recommended that the process is carried out in a fume cupboard.

The tar receiver should be cleared on a regular basis, by placing tar in a heat resistant crucible and burning off in a furnace.

WARNING — THE FUMES ARE TOXIC AND DUE CARE SHOULD BE EXERCISED IN THEIR DISPOSAL.

6 Preparation of sample

The coal used for the determination of the Gray-King coke type is the analysis sample ground to pass a sieve of 212 μm aperture. If necessary, expose the sample in a thin layer for the minimum time required for the moisture content to reach approximate equilibrium with the laboratory atmosphere.

Before commencing the determination, mix the air-dried sample thoroughly for at least 1 min, preferably by mechanical means. The sample shall be prepared on the same day as the determination is carried out.

7 Procedure

7.1 Coals with a Gray-King coke type within the range A to G₂ (see 10.3)

Raise the temperature of the furnace until it is steady at 325 °C.

Weigh on a scoop 20,0 g ± 0,1g of the sample and transfer it to the retort tube (5.2), held in such a manner that the coal cannot enter the side arm. Complete the transfer with a soft brush and allow the coal to fall to the far end of the retort tube. Hold the tube horizontally, insert the distance rod (5.3) so that the face of the disk is 150 mm from the closed end of the retort tube and spread the coal into a layer of uniform depth by shaking and turning. Withdraw the distance rod and insert a flattened pad of asbestos wool or a notched asbestos disk to retain the coal in position. Without disturbing the position of the coal, close the open end of the retort tube with a heat-resisting stopper. Connect the receiver (5.4) to the side arm and insert the retort tube in position in the furnace (5.1) so that the centre of the coal layer coincides with the centre of the furnace. If the furnace is mounted on rails, clamp the retort tube in a horizontal position and run the furnace into position.

Raise the energy input to the furnace in such a manner that the temperature of 325 °C is regained in 3 min to 7 min and maintain a uniform rate of rise of 5 °C/min thereafter until a temperature of 590 °C is reached. At this point, regulate the energy input to the furnace so that a temperature of 600 °C is reached, and maintain this temperature for 15 min.

Withdraw the retort tube (or retract the furnace) and allow it to cool. Detach the receiver, remove the stopper and slide the coke residue out for examination.

NOTE If the coal has an ash greater than 10 %, the Gray-King result might be affected.

7.2 Coals with a Gray-King coke type greater than G₂ (see 10.3)

Weigh X g of the electrode carbon (4.1), where X is always an integer, into a weighing bottle and add (20 – X) g of the coal sample. Insert the stopper and mix the contents thoroughly.

Transfer the mixture to the retort tube and proceed exactly as specified in 7.1.

Repeat the determination if necessary, varying the amount of electrode carbon in 20 g of the mixture, until a coke residue of type G is obtained using the minimum mass of electrode carbon.

8 Expression of results

Report the Gray-King coke type of a coal by reference to Figure 1 and Table 3, where the appearance of typical coke residues is described and illustrated. For coals giving a coke type with an index greater than G₂, the subscript defines the minimum number of grams of electrode carbon added to produce a standard G type coke residue.

9 Precision of the method

9.1 General

Table 1

Type of coke	Maximum acceptable difference between results	
	Same laboratory (Repeatability)	Different laboratory (Reproducibility)
A to G ₁ Greater than G ₁	One letter One unit in the subscript	One letter One unit in the subscript

9.2 Repeatability

The results of duplicate determinations, carried out at different times in the same laboratory by the same operator using the same apparatus on the same analysis sample, shall not differ by more than the above value.

9.3 Reproducibility

The means of the results of duplicate determinations, carried out in each of two different laboratories on representative portions taken from the same analysis sample after the last stage of sample preparation, shall not differ by more than the above value.

10 Notes on procedure

10.1 It has been shown that anthracite may be used as an alternative to electrode carbon. Any material which has been found by experiment to produce results equivalent to those obtained when using standard electrode carbon may be used.

If anthracite is to be used, it should meet the following parameters:

Retained on 212 μm test sieve	nil
Through 212 μm , retained on 125 μm	5 % to 10 %
Through 125 μm , retained on 63 μm	20 % to 25 %
Through 63 μm	65 % to 75 %

10.2 Determine the true relative density using a density bottle (see ISO 1014). To ensure complete wetting of the electrode carbon, use a 1 % solution of a wetting agent and evacuate the density bottle containing the electrode carbon and wetting agent solution to a pressure of 8 kPa¹⁾ in a vacuum desiccator. Maintain this vacuum for 10 min before transferring the density bottle to a water-bath thermostatically controlled at 25 °C.

10.3 Although there is no precise relationship between the crucible swelling number and the Gray-King coke type, the following table shows the broad relationship to be expected. This can be found useful for indicating the necessity for blending with electrode carbon and the probable amount required.

Table 2

Crucible swelling number	Gray-King coke type
0 to 1/2	A to B
1 to 4	C to G ₂
4 1/2 to 6	F to G ₄
6 1/2 to 8	G ₃ to G ₉
8 1/2 to 9	G ₇ or above

The above relationship is known to be applicable to United Kingdom coals and is intended only as a general guide. Each country should determine the correlation applicable to its own coals.

1) 8 kPa = 80 mbar.

11 Test report

The test report shall include the following particulars:

- a) an identification of the product tested;
- b) the reference of the method used;
- c) the results and the method of expression used;
- d) any unusual features noted during the determination;
- e) any operation not included in this International Standard or regarded as optional.

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