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**Brown coals and lignites —  
Determination of true relative density  
and apparent relative density**

*Charbons bruns et lignites — Détermination de la densité relative  
vraie et de la densité relative apparente*

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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.

International Standards are drafted in accordance with the rules given in the ISO/IEC Directives, Part 2.

The main task of technical committees is to prepare International Standards. Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

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.

ISO 5072 was prepared by Technical Committee ISO/TC 27, *Solid mineral fuels*, Subcommittee SC 5, *Methods of analysis*.

This second edition cancels and replaces the first edition (ISO 5072:1997), of which this constitutes a minor revision.

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# Brown coals and lignites — Determination of true relative density and apparent relative density

## 1 Scope

This International Standard describes methods for the determination of true relative density and the apparent relative density of brown coals and lignites.

## 2 Normative references

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

ISO 5068-2, *Brown coals and lignites — Determination of moisture — Part 2: Indirect gravimetric method for moisture in the analysis sample*

ISO 5069-2, *Brown coals and lignites — Principles of sampling — Part 2: Sample preparation for determination of moisture content and for general analysis*

## 3 Terms and definitions

### 3.1

#### **true relative density**

ratio of the mass of a sample of dry coal ground to pass through a 212  $\mu\text{m}$  sieve to the mass of an equal volume of water at a specified temperature

### 3.2

#### **apparent relative density**

ratio of the mass of a dry coal to the mass of a volume of water equal to the apparent volume of the coal at a specified temperature

## 4 Determination of the true relative density by the water method

### 4.1 Principle

The true relative density is determined pycnometrically by water displacement, with the inclusion of a wetting agent to ensure dispersion of the solid in the displacement medium.

**NOTE** The use of other displacement media such as methanol is not recommended because of possible swelling of some brown coals and lignites in such liquids.

### 4.2 Reagents

**4.2.1 Distilled or deionized water**, freshly boiled.

**4.2.2 Wetting agent**, 5 % (volume fraction) solution of detergent (such as sodium dodecyl sulfate) is suitable.

**4.2.3 Potassium dichromate-sulfuric acid mixture**, for cleaning pycnometers.

### 4.3 Apparatus

**4.3.1 Analytical balance**, sensitive to 0,1 mg.

**4.3.2 Camel hair brush**, of such a diameter that bristles can pass completely through the stem of the funnel (4.3.4).

**4.3.3 Filter paper**, for drying the necks of the pycnometers.

**4.3.4 Funnel**, with a stem of sufficient length to reach the middle of the pycnometer flask.

**4.3.5 Glass cloth**, fibre free for polishing the pycnometers prior to weighing.

**4.3.6 Glass syringe**, with a needle to bring the pycnometer liquid to the mark of the pycnometer.

**4.3.7 Earthing point**, to remove static charge from pycnometers.

**4.3.8 Mercury thermometer**, capable of measuring the temperature in the range of 20 °C to 30 °C with a minimum scale spacing of 0,1 °C.

**4.3.9 Pycnometers**, of capacity 50 ml, with capillary-bored ground stoppers, internal diameter of the neck no greater than 5 mm.

**4.3.10 Vacuum desiccator**, with protective cage.

NOTE The use of a vacuum desiccator is preferable for the purpose of degassing samples.

**4.3.11 Vacuum pump**, capable of attaining a vacuum of residual pressure from 0 kPa to 5 kPa.

**4.3.12 Water bath**, thermostatically controlled, the temperature of which shall be maintained at 25 °C ± 0,1 °C as measured by a thermometer (4.3.8) permanently placed in the water bath.

**4.3.13 Weighing boat**, with a capacity of not less than 2 g.

### 4.4 Sample

The sample for the determination of true relative density shall be the general analysis sample (ground to pass through a 212 µm sieve) prepared in accordance with ISO 5069-2.

### 4.5 Procedure

#### 4.5.1 Calibration of the mass of the pycnometer(s)

Pipette 10 ml of wetting agent (4.2.2) into the pycnometer(s) (4.3.9). Pipette 10 ml of water (4.2.1) into the pycnometer(s). Swirl to mix the solution. Place the pycnometer(s) in the vacuum desiccator (4.3.10) and evacuate to 0 Pa to 500 Pa for 15 min. Release the vacuum and fill the pycnometer(s) with water (4.2.1) using a glass syringe (4.3.6) to 3 mm from the top of the neck.

Place the pycnometer(s) in a water bath (4.3.12) thermostatically controlled at 25 °C ± 0,1 °C. Allow to equilibrate for at least 1 h 45 min. Top up the pycnometer(s) to the meniscus using a glass syringe without removing them from the water bath. Leave the pycnometer(s) in the water bath for a further 5 min.

Remove the pycnometer(s) from the water bath and insert the capillary-bored stopper(s) such that no air is trapped in the pycnometer(s). Wipe the outside of the pycnometer(s) gently using filter paper (4.3.3) to

remove excess water. Polish the pycnometer(s) using fibre-free glass cloth (4.3.5) to remove any visible film. Touch the pycnometer(s) to the earthing point (4.3.7). Weigh the pycnometer(s) and record the mass(es) to the nearest 0,1 mg.

The mean of three replicate determinations of the mass of each pycnometer which fall within the range of 1 mg is taken as the calibration mass of the pycnometer.

#### 4.5.2 Determination of true relative density

Weigh  $2 \text{ g} \pm 0,1 \text{ g}$  of sample, to the nearest 0,1 mg, into a weighing boat (4.3.13). Transfer the sample to a pycnometer using a funnel (4.3.4) and a camel hair brush (4.3.2). Ensure that no material is lost during the transfer.

Pipette 10 ml of wetting agent (4.2.2) into the pycnometer allowing the wetting agent to run down the inner surface of the pycnometer. Pipette 10 ml of water (4.2.1) into the pycnometer, allowing the water to run down the inner surface of the pycnometer. Swirl gently to wet the sample with the minimum production of bubbles. Place the pycnometer in the vacuum desiccator and evacuate to 0 Pa to 500 Pa for 15 min. To ensure that no sample containing froth is expelled from the pycnometer, regulate the vacuum supply to remove any froth produced below the neck of the pycnometer. Release the vacuum and fill the pycnometer with water (4.2.1) to 3 mm from the top of the neck using a glass syringe.

Place the pycnometer in a water bath thermostatically controlled at  $25 \text{ }^{\circ}\text{C} \pm 0,1 \text{ }^{\circ}\text{C}$ . Allow to equilibrate for at least 1 h 45 min. Top up the pycnometer to the meniscus using a glass syringe, without removing from the water bath. Leave the pycnometer in the water bath for a further 5 min.

Remove the pycnometer from the water bath and insert the capillary bored stopper such that no air is trapped in the pycnometer. Wipe the outside of the pycnometer gently using filter paper to remove excess water. Polish the pycnometer using fibre-free glass cloth to remove any visible film. Touch the pycnometer to the earthing point. Weigh the pycnometer and record the mass to the nearest 0,1 mg.

Determine the moisture content,  $M$ , on a separate test portion of the sample (4.4) in accordance with ISO 5068-2.

#### 4.6 Calculation of results

Calculate the true relative density of the dry coal,  $TRD_d$ , according to the formula:

$$TRD_d = \frac{m \times (100 - M_{ad})}{m \times (100 - M_{ad}) + 100 \times (m_1 - m_2)}$$

where

$m$  is the mass, in grams, of the analysis sample;

$m_1$  is the mass, in grams, of the pycnometer and water;

$m_2$  is the mass, in grams, of the pycnometer, sample and water;

$M_{ad}$  is the moisture, as a percentage by mass, of the analysis sample.

#### 4.7 Precision of the method

##### 4.7.1 Repeatability limit

The results of duplicate determinations, carried out in the same laboratory by the same operator using the same apparatus on the same sample within short intervals of time, shall not differ by more than the value given in Table 1.

#### 4.7.2 Reproducibility limit

The means of the results of duplicate determinations, carried out in each of two laboratories on representative test portions taken from the same sample, shall not differ by more than the value given in [Table 1](#).

**Table 1 — Precision of the method**

| Repeatability | Reproducibility |
|---------------|-----------------|
| 0,02          | 0,04            |

#### 4.8 Test report

The test report shall contain the following information:

- an identification of the sample;
- a reference to this International Standard including year of publication;
- the result and the method of expression used;
- any unusual features noted during the determination.

### 5 Determination of the apparent relative density

#### 5.1 Principle

The apparent relative density is determined by weighing a sample suspended in water, allowing the sample to drain to remove surface liquid and then reweighing the sample in air.

#### 5.2 Reagents

**5.2.1 Water**, distilled or deionized.

#### 5.3 Apparatus

**5.3.1 Beam balance**, sensitive to 10 mg, modified as in [Figure 1](#).

**5.3.2 Evaporating dishes**, 160 mm diameter, 60 mm deep.

**5.3.3 Paper towels or filter papers**.

**5.3.4 Top pan balance**, capacity of 1 kg, sensitive to 10 mg.

**5.3.5 Vacuum pump**, capable of attaining 250 Pa pressure.

**5.3.6 Vacuum desiccator**, 200 mm diameter.

#### 5.4 Sample

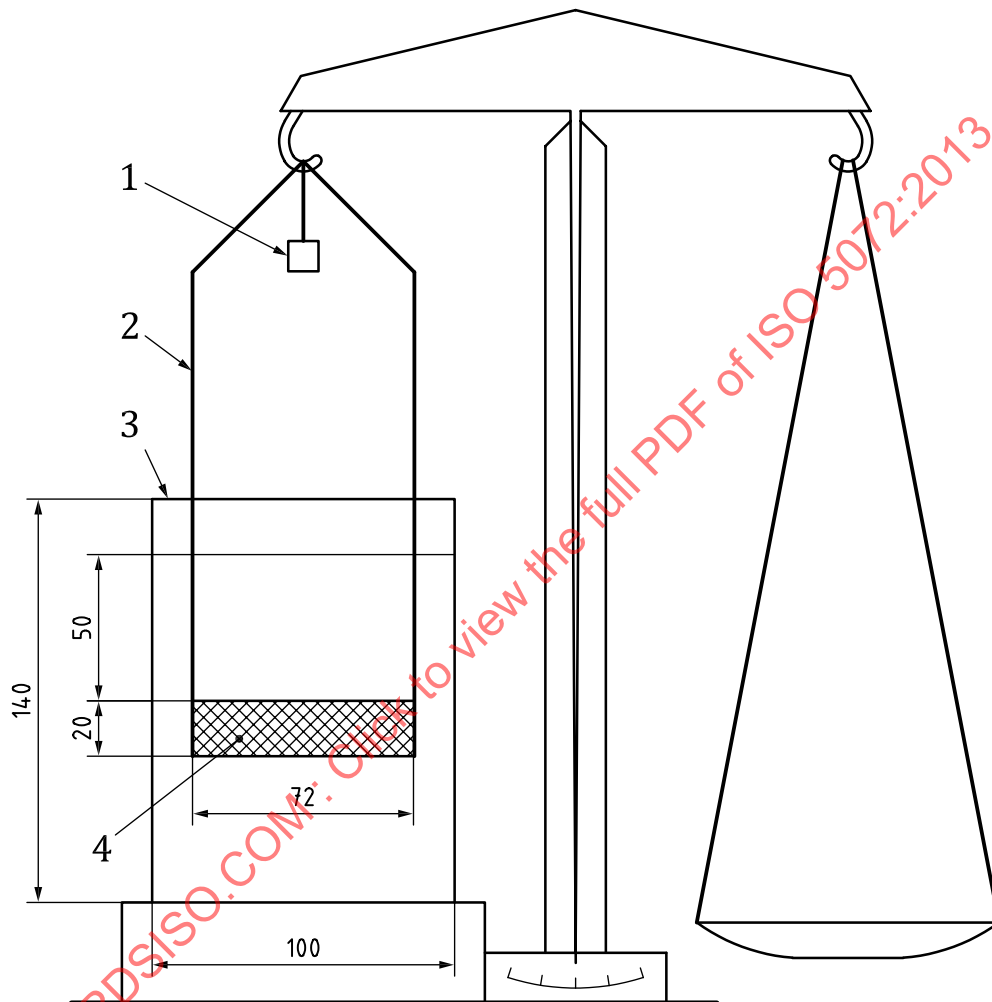
The determination of apparent relative density shall be carried out on an air-dried sample of particle size 10 mm to 30 mm.

## 5.5 Procedure

Set up the beam balance (5.3.1) as shown in Figure 1, with a minimum of 50 mm of water (5.2.1) covering the gauge basket. Tare the beam balance so that at the balance point approximately 5 g is added to the weight pan. Record the tare mass.

NOTE Check the beam balance tare at least once every 5 to 10 determinations.

Dimensions in millimetres



### Key

- 1 tare weight, approximately 5 g
- 2 copper wire
- 3 beaker, 800 ml
- 4 gauze basket (phosphor-bronze 12 mm aperture)

**Figure 1 — Modified beam balance**

Weigh from 30 g to 35 g of sample (5.4) into an evaporating dish (5.3.2). Add water (5.2.1) to the dish to completely immerse the sample. Place the evaporating dish plus sample in a vacuum desiccator (5.3.6). Evacuate the desiccator to 0 Pa to 250 Pa for 5 min. Release the vacuum and allow to stand for a further 5 min.

Transfer the sample to a pad of paper towels (5.3.3), using the paper towels to remove excess water. Immediately place the sample in the gauze basket of the beam balance ensuring that the sample is totally covered with water. Allow the balance to attain equilibrium and record the mass of the sample in water to the nearest 10 mg.