
**Carbonaceous materials for the
production of aluminium — Cathode
block materials —**

**Part 1:
Determination of the expansion due to
sodium penetration with application
of pressure**

*Produits carbonés utilisés pour la production de l'aluminium — Blocs
cathodiques —*

*Partie 1: Détermination de l'expansion due à la pénétration du
sodium avec application de pression*



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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).

Any trade name used in this document is information given for the convenience of users and does not constitute an endorsement.

For an explanation of the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT) see www.iso.org/iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 226, *Materials for the production of primary aluminium*.

This third edition cancels and replaces the second edition (ISO 15379-1:2015), which has been technically revised. The main changes to the previous edition are as follows:

- ASTM E 220 has been removed from [Clause 2](#) and [5.3](#);
- minor changes have been made in [6.2](#), [6.3](#), [6.4](#) and [6.5](#);
- [Clause 10](#) has been aligned with test reports in other ISO/TC 226 standards.

A list of all parts in the ISO 15379 series can be found on the ISO website.

Any feedback or questions on this document should be directed to the user's national standards body. A complete listing of these bodies can be found at www.iso.org/members.html.

Introduction

Expansion due to sodium penetration is an important property of carbon cathode blocks. As soon as alumina electrolysis starts, sodium penetrates into the carbon cathode blocks causing swelling of these blocks. This increase in volume creates mechanical stresses within the blocks and/or bulging of the bottom block plate. This can lead to cracks through which liquid aluminium and/or liquid electrolyte can flow, reaching the thermal insulation beneath the blocks and destroying these ceramic materials. In such cases, the electrolysis cell has to be relined, resulting in loss of aluminium production and high expenses. Therefore, cathode blocks produced with materials allowing only low sodium penetration and with the lowest possible expansion due to sodium penetration are preferred.

The study can be supplemented by measuring electrical resistivity (see ISO 11713) before and after the test.

Due to thermal and sodium expansion when heating up the cathode blocks in the electrolysis cells to operational temperature, the block will be subject to pressure from the steel shell; therefore, in the present method, 5 MPa is chosen as a realistic maximum pressure.

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Carbonaceous materials for the production of aluminium — Cathode block materials —

Part 1:

Determination of the expansion due to sodium penetration with application of pressure

1 Scope

This document specifies a method that covers the determination of linear expansion under external pressure due to sodium penetration in cathode-block materials used in the production of aluminium.

The linear expansion of the blocks depends on the sampling direction due to anisotropy.

2 Normative references

The following documents are referred to in the text in such a way that some or all of their content constitutes requirements 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 8007-1, *Carbonaceous materials used in the production of aluminium — Sampling plans and sampling from individual units — Part 1: Cathode blocks*

3 Terms and definitions

No terms and definitions are listed in this document.

ISO and IEC maintain terminological databases for use in standardization at the following addresses:

- ISO Online browsing platform: available at <https://www.iso.org/obp>
- IEC Electropedia: available at <http://www.electropedia.org/>

4 Principle

An anodic graphite crucible containing a cryolitic bath with an initial cryolite ratio of 4,0 and with a cathode carbon sample placed in the crucible, as shown in [Figure 1](#), is used for this experiment. The whole sample is immersed in a cryolitic bath with a graphite cylinder fitted as an extension from the sample. The crucible is placed in a crucible support, which is connected to a hydraulic power cylinder. A constant pressure of 5 MPa is applied to the sample by pressing the graphite cylinder against a stop rod in the top of the furnace by the hydraulic power cylinder. The whole assembly is heated in a tubular furnace to $(980 \pm 5) ^\circ\text{C}$ and is then electrolyzed for 2 h with a current density of approximately 0,7 A/cm² (see [Clause 7](#) for precise setting). The expansion is measured by a probe, which is fastened to the frame of the furnace and measures the position of the crucible support.

5 Apparatus

The principle of the apparatus for measuring sodium expansion is shown in [Figure 1](#).

5.1 Furnace, capable of maintaining a temperature of 980 °C, with a temperature gradient over the melt of less than 10 °C.

5.2 Furnace control device, suitable for holding the temperature at (980 ± 5) °C.

5.3 Temperature-measuring device, i.e. a thermocouple, preferably of type K or S, capable of determining the temperature to ± 5 °C at 980 °C.

5.4 Crucible, made of graphite, with an inner diameter of 90 mm and an inner height of 90 mm.

The crucible acts as an anode.

5.5 Lid, made of graphite, which shall have a hole in the centre to allow the sample/graphite extension to reach above the lid.

5.6 Insulating ring, made of a ceramic material, which resists the temperature and the fluoride environment.

The ring shall be placed in the hole of the graphite lid and act as an electrical insulation between the lid and the sample/graphite extension. The ring shall be large enough to allow the sample/graphite extension to move freely in the vertical direction.

5.7 Crucible support, made of heat-resistant steel to conduct anodic current from the power supply to the crucible.

5.8 Stop rod, made of heat-resistant steel to conduct cathodic current to the top of the sample.

The stop rod acts as a fixed reference point for the expansion measurements.

The material for the steel support and the steel stop rod should have minimal deformation at a pressure of 5 MPa.

The steel quality Sanicro 31HT¹⁾, X 10 NiCrAlTi32.20 is suitable for the crucible support rod and the stop rod. If other steel grades are used, it shall be verified that they do not deform significantly versus the length change of the sample.

5.9 Alumina disk, capable of covering the bottom of the crucible and capable of acting as electrical insulation between the crucible and the sample. The disk should preferably have a centring slot to help position the sample in the centre of the crucible.

5.10 Graphite cylinder, which shall be used as an extension to the sample and shall have a diameter of $(30,0 \pm 0,1)$ mm and a length of (40 ± 1) mm.

5.11 Hydraulic power cylinder, capable of applying a constant pressure of $5 \text{ MPa} \pm 0,1 \text{ MPa}$ during the whole experiment independent of the expansion of the sample.

5.12 Extensometer, connected to a computer or data recorder, with a measuring range of 10 mm and an accuracy of 1 μm over the whole range, in order to observe the expansion due to sodium penetration.

5.13 Power supply, capable of supplying 39,6 A DC, with a current density of the cathode which shall be 0,7 A/cm².

1) Sanicro 31HT is an example of a suitable product available commercially. This information is given for the convenience of users of this document and does not constitute an endorsement by ISO of this product.

6 Reagents

6.1 Argon, welding-grade quality.

6.2 Cryolite, Na_3AlF_6 , natural (≥ 99 % by mass) or synthetic (≥ 97 % by mass).

6.3 Sodium fluoride, NaF , purum, ≥ 99 % by mass.

6.4 Calcium fluoride, CaF_2 , precipitated pure, ≥ 97 % by mass.

6.5 Alumina, Al_2O_3 , extra pure, ≥ 98 % by mass.

6.6 Bath composition, which shall have a cryolite ratio of 4,0 and consist of the following: 71,5 % Na_3AlF_6 , 14,5 % NaF , 5,0 % CaF_2 , 9,0 % Al_2O_3 . The bath is crushed to < 2 mm using a jaw crusher and shall have a mass of 765 g.

7 Samples

Sample the material in accordance with ISO 8007-1. The diameter of the sample shall be $(30,0 \pm 0,1)$ mm. The length shall be (60 ± 1) mm.

8 Procedure

Assemble the crucible support (5.7) with the crucible (5.4) and the sample so they can be pressed together with a hydraulic power cylinder against a heat-resistant steel stop rod (5.8) conducting the cathodic current to the top of the sample (see Figure 1). The stop rod (5.8) acts as a fixed reference point for the expansion measurements. Place the alumina disk (5.9) in the bottom of the crucible (5.4). Measure the length, l_0 , of the sample at room temperature with an accuracy of 0,1 mm. Place the sample on the alumina disk in the centre of the crucible. Place the graphite cylinder (5.10) on top of the sample. Fill the crucible with the bath prepared in accordance with 6.6. Place the lid (5.5), with an insulating ring (5.6) around the graphite extension, on top of the crucible.

Lift the crucible by the hydraulic power cylinder (5.11) until the sample touches the stop rod (5.8) inside the furnace (5.1). Adjust the hydraulic pressure to 5 MPa.

Place the thermocouple close to the crucible near the centre of the melt height.

Place the extensometer (5.12) underneath the base of the crucible holder outside the furnace.

Heat the furnace to (980 ± 5) °C with an argon (6.1) flush, and measure the change in the length of the sample and the apparatus. Wait until no further thermally induced movement can be detected in the sample and apparatus. Determine the position of the crucible support and take this reading as the zero reference for subsequent measurements of the change in length, $\Delta l_{\text{meas}, t}$.

Connect the power supply (5.13) to the crucible support and the stop rod. Electrolyze the system for 2 h with a constant current of 39,6 A. Record the change in length, $\Delta l_{\text{meas}, t}$, every minute, then turn off the electrolysis.

Release the pressure, but keep the crucible inside the furnace. Allow the furnace to cool down to room temperature.

9 Results

9.1 Calculation

Calculate the relative expansion of the sample for each recording using [Formula \(1\)](#):

$$\Delta L_t = \frac{\Delta l_{\text{meas}, t}}{l_0} \times 100 \quad (1)$$

where

ΔL_t is the relative expansion, in percent, at time, t ;

$\Delta l_{\text{meas}, t}$ is the measured change in length, in millimetres, at time, t ;

l_0 is the initial sample length, in millimetres.

Plot the relative expansion, ΔL_t , versus time. Determine the maximum relative expansion, ΔL_{max} . Round the results to the second decimal place.

9.2 Precision

The precision was calculated in accordance with ISO 5725-2. The degrees of freedom, i.e. the number of laboratories (e.g. four) times the number of samples (e.g. three), was equal to 12.

The repeatability is calculated from [Formula \(2\)](#):

$$r = 0,20 \times \Delta \bar{L}_{\text{max}} + 0,02 \quad (2)$$

where

$\Delta \bar{L}_{\text{max}}$ is the maximum average expansion;

0,02 is in absolute %.

EXAMPLE Material A has a maximum average expansion equal to 0,56 %. The repeatability is then $0,20 \times 0,56 + 0,03 = 0,15$, in % absolute.

The reproducibility is calculated from [Formula \(3\)](#):

$$R = 0,24 \times \Delta \bar{L}_{\text{max}} + 0,03 \quad (3)$$

where the parameters are the same as for [Formula \(2\)](#).

10 Test report

The report shall include the following information:

- an identification of the sample;
- the International Standard used (including its year of publication), i.e. ISO 15379-1:2019;
- the method used (if the standard includes several);
- the date of the test;
- the result(s), including a reference to the clause which explains how the results were calculated;
- any deviations from the procedure;

g) any unusual features observed.

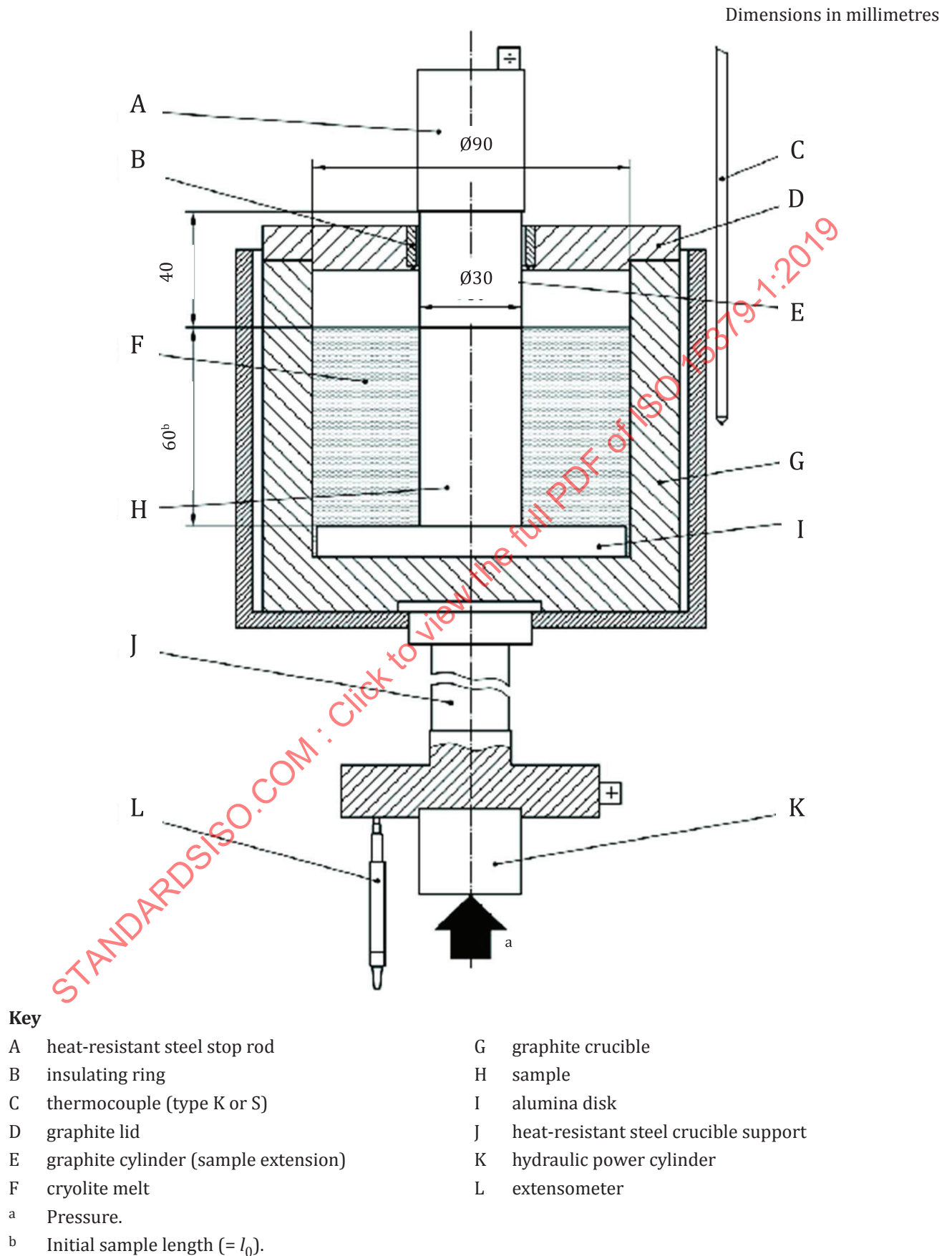


Figure 1 — Apparatus for measuring expansion due to sodium penetration