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Iron ores for blast furnace feedstocks — Determination of the reducibility by the rate of reduction index

*Minerais de fer pour charges de hauts fourneaux — Détermination de
la réductibilité à partir de la vitesse de réduction*

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ISO copyright office
CP 401 • Ch. de Blandonnet 8
CH-1214 Vernier, Geneva
Phone: +41 22 749 01 11
Email: copyright@iso.org
Website: www.iso.org

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Foreword

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This document was prepared by Technical Committee ISO/TC 201, *Iron ore and direct reduced iron*, Subcommittee SC 3, *Physical testing*.

This fifth edition cancels and replaces the fourth edition (ISO 4695:2015), of which it constitutes a minor revision to correct [Formula B.3](#) in [Annex B](#).

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

This document concerns one of a number of physical test methods that have been developed to measure various physical parameters and to evaluate the behaviour of iron ores, including reducibility, disintegration, crushing strength, apparent density, etc. This method was developed to provide a uniform procedure, validated by collaborative testing, to facilitate comparisons of tests made in different laboratories.

The results of this test have to be considered in conjunction with other tests used to evaluate the quality of iron ores as feedstocks for blast furnace processes.

This document can be used to provide test results as part of a production quality control system, as a basis of a contract or as part of a research project.

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Iron ores for blast furnace feedstocks — Determination of the reducibility by the rate of reduction index

CAUTION — This document can involve hazardous operations and equipment. This document does not purport to address all of the safety issues associated with its use. It is the responsibility of the user of this document to establish appropriate safety and health practices and determine the applicability of regulatory limitations prior to its use.

1 Scope

This document specifies a method to provide a relative measure for evaluating the extent to and ease with which oxygen can be removed from iron ores, when reduced under conditions resembling those prevailing in the reduction zone of a blast furnace.

This document is applicable to lump ores, sinters and hot-bonded pellets.

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 2597-1, *Iron ores — Determination of total iron content — Part 1: Titrimetric method after tin(II) chloride reduction*

[ISO/FDIS 4695](#)

ISO 2597-2, *Iron ores — Determination of total iron content — Part 2: Titrimetric methods after titanium(III) chloride reduction*

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ISO 3082, *Iron ores — Sampling and sample preparation procedures*

ISO 9035, *Iron ores — Determination of acid-soluble iron(II) content — Titrimetric method*

ISO 11323, *Iron ore and direct reduced iron — Vocabulary*

3 Terms and definitions

For the purposes of this document, the terms and definitions given in ISO 11323 apply.

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

The test portion is isothermally reduced in a fixed bed, at 950 °C, using a reducing gas consisting of CO and N₂, and the mass loss of the test portion is recorded continuously or at specified time intervals until its degree of reduction reaches 65 %. The rate of reduction is calculated at the oxygen/iron ratio of 0,9.

5 Sampling, sample preparation and preparation of test portions

5.1 Sampling and sample preparation

Sampling of a lot and preparation of a test sample shall be in accordance with ISO 3082.

The size range for pellets, sinters and lump ores shall be $-12,5 \text{ mm} + 10,0 \text{ mm}$.

A test sample of at least 2,5 kg, on a dry basis, of the sized material shall be obtained.

Oven-dry the test sample to constant mass at $105 \text{ °C} \pm 5 \text{ °C}$ and cool it to room temperature before preparation of the test portions.

NOTE Constant mass is achieved when the difference in mass between two subsequent measurements becomes less than 0,05 % of the initial mass of the test sample.

5.2 Preparation of test portions

Collect each test portion by taking ore particles at random.

NOTE Manual methods of division recommended in ISO 3082, such as riffing, can be applied to obtain the test portions.

At least five test portions, each of approximately 500 g (\pm the mass of one particle) shall be prepared from the test sample: four test portions for testing and one for chemical analysis.

Weigh the test portions to the nearest 1 g and register the mass of each test portion on its recipient label.

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6 Apparatus

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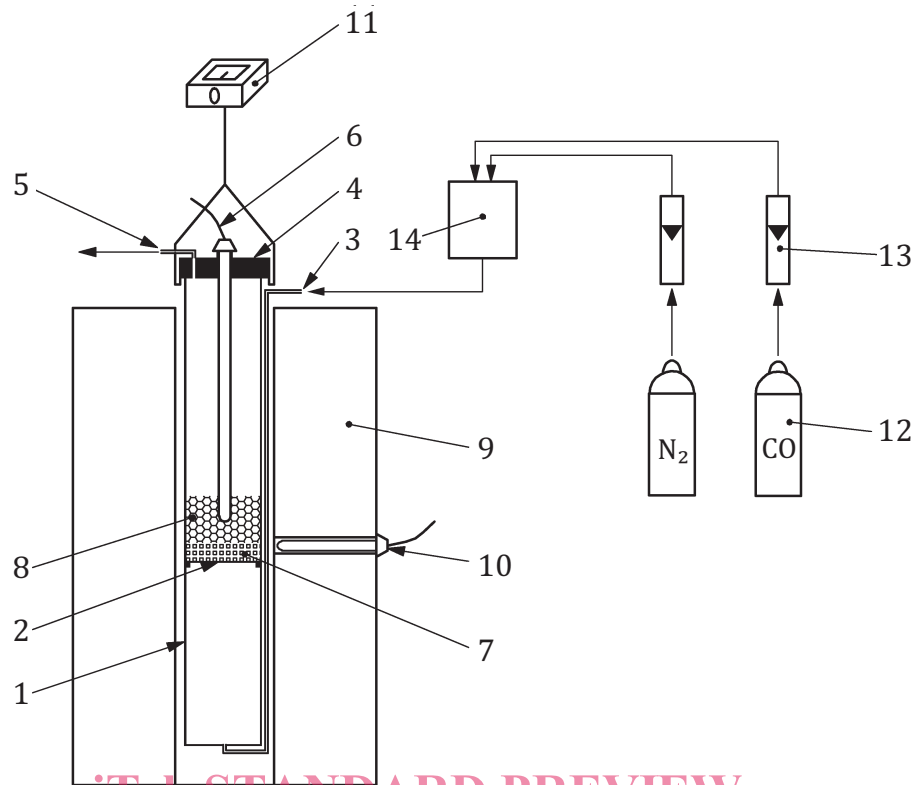
6.1 The test apparatus shall comprise the following:

- a) ordinary laboratory equipment, such as an oven, hand tools, a time-control device and safety equipment;
- b) reduction tube assembly;
- c) furnace, equipped with a balance for permitting the mass loss of the test portion to be read at any time during the test;
- d) system to supply the gases and regulate the flow rates;
- e) weighing device.

[Figure 1](#) shows an example of the test apparatus.

6.2 Reduction tube, made of non-scaling, heat-resistant metal to withstand temperatures higher than 950 °C and resistant to deformation. The internal diameter shall be $75 \text{ mm} \pm 1 \text{ mm}$. A removable perforated plate, made of non-scaling, heat-resistant metal to withstand temperatures higher than 950 °C , shall be mounted in the reduction tube to support the test portion and to ensure uniform gas flow through it. The perforated plate shall be 4 mm thick, with its diameter 1 mm less than the internal diameter of the tube. The holes in the plate shall be 2 mm to 3 mm in diameter at a pitch centre distance of 4 mm to 5 mm.

[Figure 2](#) shows an example of a reduction tube.

**Key****Reduction tube**

- 1 reduction tube
- 2 perforated plate
- 3 gas inlet
- 4 lid
- 5 gas outlet
- 6 thermocouple for measuring the reduction temperature
- 7 porcelain ball layer
- 8 test portion

Furnace

- 9 electrically heated furnace
- 10 thermocouple for temperature regulation of furnace
- 11 balance

Gas supply system

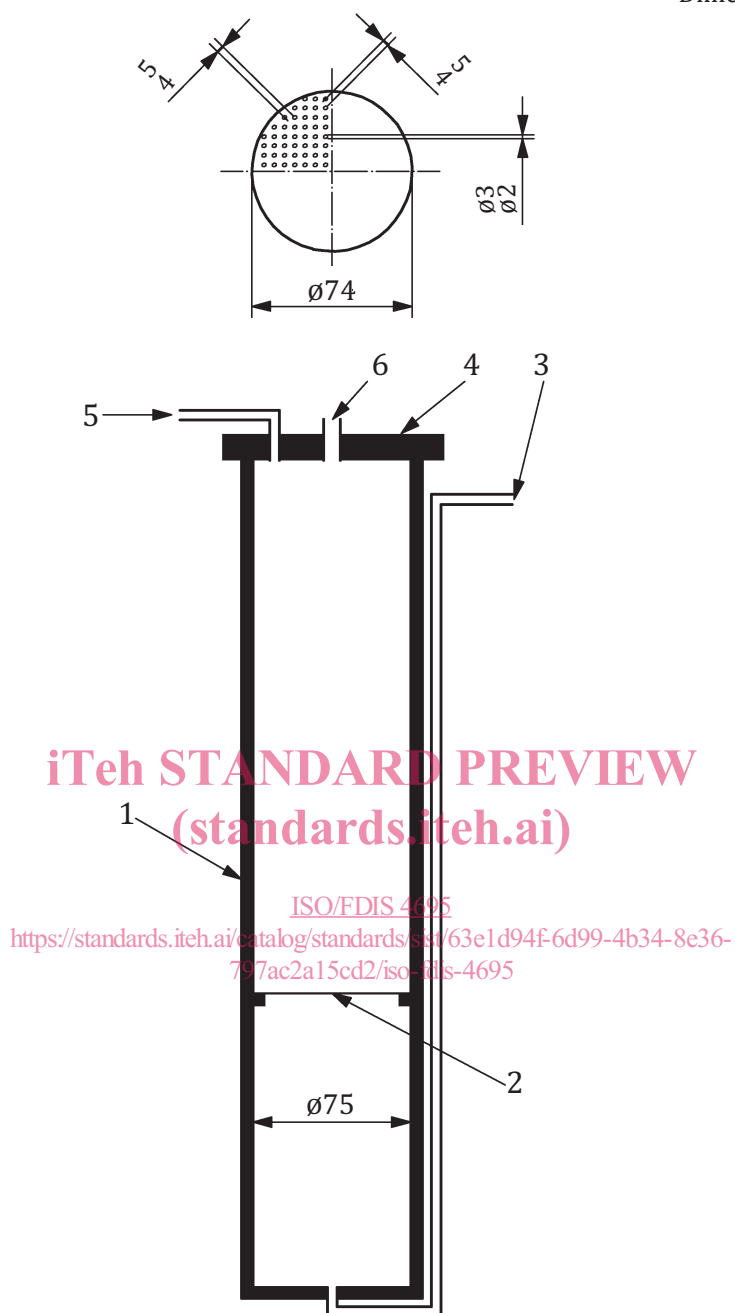
- 12 gas cylinder
- 13 gas flow meters
- 14 mixing vessel

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Figure 1 — Example of test apparatus (schematic diagram)



Key

- 1 reduction tube
- 2 perforated plate
- 3 opening for gas inlet
- 4 lid
- 5 opening for gas outlet
- 6 opening for thermocouple insertion

NOTE Dimensions not specified in [Clause 6](#) are shown for information only.

Figure 2 — Example of reduction tube (schematic diagram)

6.3 Furnace, having a heating capacity and temperature control able to maintain the entire test portion, as well as the gas entering the bed, at $950\text{ °C} \pm 10\text{ °C}$.

6.4 Balance, capable of weighing the reduction tube assembly, including the test portion, to an accuracy of 1 g. The balance shall have an appropriate device to suspend the reduction tube assembly.

6.5 Gas-supply system, capable of supplying the gases and regulating gas flow rates. It shall be ensured that a frictionless connection between the gas-supply system and the reduction tube does not affect the weight loss determination during reduction.

6.6 Weighing device, capable of weighing the test sample and test portions to an accuracy of 1 g.

7 Test conditions

7.1 General

Volumes and flow rates of gases used are as measured at a reference temperature of 0 °C and at a reference atmospheric pressure of $101,325\text{ kPa}$ ($1,013\ 25\text{ bar}$).

7.2 Reducing gas

7.2.1 Composition

The reducing gas shall consist of

CO $40,0\% \pm 0,5\%$ (volume fraction)

N₂ $60,0\% \pm 0,5\%$ (volume fraction)

7.2.2 Purity

Impurities in the reducing gas shall not exceed the following:

H₂ $0,2\%$ (volume fraction)

CO₂ $0,2\%$ (volume fraction)

O₂ $0,1\%$ (volume fraction)

H₂O $0,2\%$ (volume fraction)

7.2.3 Flow rate

The flow rate of the reducing gas, during the entire reducing period, shall be maintained at $50\text{ l/min} \pm 0,5\text{ l/min}$.

7.3 Heating and cooling gas

Nitrogen (N₂) shall be used as the heating and cooling gas. Impurities shall not exceed $0,1\%$ (volume fraction).

The flow rate of N₂ shall be maintained at 25 l/min until the test portion reaches 950 °C , and at 50 l/min during the temperature-equilibration period. During cooling, it shall be maintained at 5 l/min .