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**Järnmalm – Bestämning av magnesium – Flamatomär
absorptionsspektrometrisk metod (ISO 10204:2017, IDT)**

**Iron ores – Determination of magnesium – Flame atomic
absorption spectrometric method (ISO 10204:2017, IDT)**



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Den internationella standarden ISO 10204:2017 gäller som svensk standard. Detta dokument innehåller den officiella engelska versionen av ISO 10204:2017.

Denna standard ersätter SS-ISO 10204:2006, utgåva 1.

The International Standard ISO 10204:2017 has the status of a Swedish Standard. This document contains the official version of ISO 10204:2017.

This standard supersedes the Swedish Standard SS-ISO 10204:2006, edition 1.

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Information about the content of the standard is available from the Swedish Standards Institute (SIS), telephone +46 8 555 520 00. Standards may be ordered from SIS, who can also provide general information about Swedish and foreign standards.

Denna standard är framtagen av kommittén för Järnmalm, SIS/TK 149.

Har du synpunkter på innehållet i den här standarden, vill du delta i ett kommande revideringsarbete eller vara med och ta fram andra standarder inom området? Gå in på www.sis.se - där hittar du mer information.

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SS-ISO 10204:2018 (E)**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 on 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 the following URL: www.iso.org/iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 102, *Iron ore and direct reduced iron*, Subcommittee SC 2, *Chemical analysis*.

This fourth edition cancels and replaces the third edition (ISO 10204:2015), of which it constitutes a minor revision with the following changes:

- in [7.2](#), reference to ISO 2596 has been incorporated and [7.2](#) has been rearranged in a more appropriate layout/format;
- in [8.2](#), reference to ISO 2596 has been incorporated;
- “predried” has been deleted where it is inappropriate in [8.1](#), [8.2](#), and [8.3.2](#);
- in [Table 1](#), some minor changes have been made;
- in [9.2.4](#), [Formula \(7\)](#) and relevant descriptions have been modified to harmonize this subclause across all documents for which ISO/TC 102/SC 2 is responsible.

Iron ores — Determination of magnesium — Flame atomic absorption spectrometric method

WARNING — This document might involve hazardous materials, operations, and equipment. This document does not purport to address all of the safety problems associated with its use. It is the responsibility of the user of this document to establish appropriate health and safety practices and determine the applicability of any limitations prior to use.

1 Scope

This document specifies a flame atomic absorption spectrometric method for the determination of the mass fraction of magnesium in iron ores.

This method is applicable to mass fractions of magnesium between 0,010 % and 2,00 % in natural iron ores, iron ore concentrates, and agglomerates, including sinter products.

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 648, *Laboratory glassware — Single-volume pipettes*

ISO 1042, *Laboratory glassware — One-mark volumetric flasks*

ISO 2596, *Iron ores — Determination of hygroscopic moisture in analytical samples — Gravimetric, Karl Fischer and mass-loss methods*

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

ISO 3696, *Water for analytical laboratory use — Specification and test methods*

ISO 7764, *Iron ores — Preparation of predried test samples for chemical analysis*

ISO 9516-1:2003, *Iron ores — Determination of various elements by X-ray fluorescence spectrometry — Part 1: Comprehensive procedure*

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:

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

4 Principle

The test portion is decomposed by the treatment with hydrochloric acid and a small amount of nitric acid, and then evaporated to dehydrate silica, followed by dilution and filtration.

The residue is ignited and silica is removed by evaporation with hydrofluoric and sulfuric acids. The residue is fused with sodium carbonate and the cooled melt is dissolved in the filtrate.

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The solution obtained is aspirated into the flame of an atomic absorption spectrometer using an air-acetylene burner.

The absorbance values obtained for magnesium are compared with those obtained from calibration solutions.

NOTE A dinitrogen oxide/acetylene flame can be used for the determination, in which case, the sensitivity is decreased by a factor of about 3.

5 Reagents

During the analysis, use only reagents of recognized analytical grade and only water that complies with Grade 2 of ISO 3696.

5.1 Sodium carbonate, (Na_2CO_3), anhydrous.

5.2 Hydrochloric acid, ρ 1,19 g/ml.

5.3 Hydrochloric acid, ρ 1,19 g/ml, diluted 1 + 9.

5.4 Nitric acid, ρ 1,4 g/ml.

5.5 Hydrofluoric acid, ρ 1,13 g/ml, 40 % (mass fraction) or ρ 1,185 g/ml, 48 % (mass fraction).

5.6 Sulfuric acid, ρ 1,84 g/ml, diluted 1 + 1.

5.7 Pure iron, minimum purity 99,9 % (mass fraction), of mass fraction of magnesium less than 0,0002 %.

5.8 Background solution.

Dissolve 5,0 g pure iron (5.7) in 50 ml of hydrochloric acid (5.2) and oxidize by adding nitric acid (5.4) drop by drop. Evaporate until a syrupy consistency is obtained. Add 20 ml of hydrochloric acid (5.2) and dilute to 200 ml with water. Dissolve 17 g of sodium carbonate (5.1) in water, add carefully to the iron solution, and heat to remove carbon dioxide. Transfer the cooled solution to a 1 000 ml one-mark volumetric flask, dilute to volume with water, and mix.

5.9 Lanthanum chloride solution.

Dissolve 50 g of lanthanum chloride ($\text{LaCl}_3 \cdot 7\text{H}_2\text{O}$) (of mass fraction of magnesium less than 0,002 %) in 50 ml of hydrochloric acid (5.2) and 300 ml of hot water. Cool and dilute to 1 l.

5.10 Magnesium standard solution, 25 μg Mg/ml.

Dissolve 0,500 0 g of oxide-free magnesium metal [minimum purity 99,9 % (mass fraction)] or 0,829 2 g of analytical grade magnesium oxide (dried and weighed in accordance with ISO 9516-1:2003, 4.9) by slowly adding 75 ml of hydrochloric acid (5.2), diluted 1 + 3. When dissolved, cool, transfer to a 1 000 ml one-mark volumetric flask, dilute to volume with water, and mix. Transfer 10 ml of this solution to a 200 ml one-mark volumetric flask, dilute to volume with water, and mix.

5.11 Magnesium calibration solutions.

Using pipettes, transfer 2,0 ml, 5,0 ml, 10,0 ml, 15,0 ml, and 20,0 ml portions of magnesium standard solution (5.10) to 200 ml volumetric flasks (see next paragraph). Add 6 ml of hydrochloric acid (5.2), 60 ml of background solution (5.8), and 40 ml of lanthanum chloride solution (5.9) to each flask. Dilute the solution to volume with water and mix. Prepare a zero magnesium calibration solution

by transferring 60 ml of background solution (5.8) to a 200 ml volumetric flask. Then add 6 ml of hydrochloric acid (5.2) and 40 ml of lanthanum chloride solution (5.9). Dilute the solution to volume with water and mix.

The range of magnesium that can be covered can vary from instrument to instrument. Attention should be paid to the minimum criteria given in 6.3. For instruments having high sensitivity, smaller portions of standard solution or a more diluted standard solution can be used.

6 Apparatus

Ordinary laboratory apparatus, including one-mark pipettes and one-mark volumetric flasks complying with the specifications of ISO 648 and ISO 1042, respectively, and the following.

6.1 Platinum crucible with a lid, of minimum capacity 30 ml.

6.2 Muffle furnace, capable of maintaining a temperature of approximately 1 100 °C.

6.3 Atomic absorption spectrometer, equipped with an air-acetylene burner.

WARNING — Follow the manufacturer's instructions for igniting and extinguishing the dinitrogen oxide/acetylene flame to avoid possible explosion hazards. Wear tinted safety glasses whenever the flame is burning.

The atomic absorption spectrometer used in this method shall meet the following criteria.

- a) Minimum sensitivity: the absorbance of the most concentrated magnesium calibration solution (see 5.11) shall be at least 0,3.
- b) Graph linearity: the ratio between the slope of the calibration graph covering the top 20 % of the concentration range (expressed as a change in absorbance) and the value of the slope for the bottom 20 % of the concentration range determined in the same way shall not be less than 0,7.
- c) Minimum stability: the standard deviation of the absorbance of the most concentrated calibration solution and that of the zero calibration solution, each being calculated from a sufficient number of repetitive measurements, shall be less than 1,5 % and 0,5 %, respectively, of the mean value of the absorbance of the most concentrated calibration solution.

The use of a strip chart recorder and/or digital readout device is recommended to evaluate criteria a), b), and c) and for all subsequent measurements.

NOTE Instrument parameters can vary with each instrument. The following parameters were successfully used in several laboratories and they can be used as guidelines. Solutions were aspirated into an air-acetylene burner.

In systems where the values shown below for gas flow rates do not apply, the ratio of the gas flow rates can still be a useful guideline.

— Hollow cathode lamp, mA	15
— Wavelength, nm	285,2
— Air flow rate, l/min	22
— Acetylene flow rate, l/min	4,2

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7 Sampling and samples

7.1 Laboratory sample

For analysis, use a laboratory sample of $-100\ \mu\text{m}$ particle size which has been taken and prepared in accordance with ISO 3082. In the case of ores having significant contents of combined water or oxidizable compounds, use a particle size of $-160\ \mu\text{m}$.

NOTE A guideline on significant contents of combined water and oxidizable compounds is incorporated in ISO 7764.

7.2 Preparation of test samples

7.2.1 General

Depending on the ore type, proceed in accordance with either [7.2.2](#) or [7.2.3](#).

7.2.2 Method specified in ISO 2596

This method is applicable to all types of ores.

Thoroughly mix the laboratory sample and, taking multiple increments, extract a test sample in such a way that it is representative of the whole contents of the container. Bring the test sample into equilibrium with the laboratory atmosphere in accordance with ISO 2596.

7.2.3 Method specified in ISO 7764

This method is not applicable to the following types of ores:

- a) processed ores containing metallic iron (direct-reduces iron);
- b) natural or processed ores in which the sulfur content is higher than 0,2 %;
- c) natural or processed ores in which the content of combined water is higher than 2,5 % by mass.

Thoroughly mix the laboratory sample and, taking multiple increments, extract a test sample in such a way that it is representative of the whole contents of the container.

8 Procedure

8.1 Number of determinations

Carry out the analysis at least in duplicate in accordance with [Annex A](#), independently, on one test sample (see [7.2](#)).

NOTE The expression "independently" means that the second and any subsequent result is not affected by the previous result(s). For this particular analytical method, this condition implies that the repetition of the procedure is carried out either by the same operator at a different time or by a different operator including, in either case, appropriate recalibration.

8.2 Test portion

Taking several increments, weigh, to the nearest 0,000 2 g, approximately 0,5 g of the air-equilibrated test sample obtained in accordance with [7.2.2](#). Concurrently, determine the hygroscopic moisture content in accordance with ISO 2596 and calculate the dried mass of the test portion.

Where the test sample is obtained in accordance with [7.2.3](#), transfer 0,5 g of the test portion to a weighing bottle, then dry at $105\text{ °C} \pm 2\text{ °C}$ and determine the dried mass of the test portion to the nearest 0,000 2 g in accordance with ISO 7764.

8.3 Blank test and check test

8.3.1 Blank test

8.3.1.1 Preparation of blank test solution

Using pure iron ([5.7](#)) with a 60 % mass of the test portion in place of the test portion, carry out the same procedure described from [8.4.1](#) to [8.4.3](#) as the test portion in parallel with it.

8.3.1.2 Measurement of blank value

Carry out in accordance with [8.4.5](#).

Where the analysis is carried out on several samples at the same time, the blank value may be represented by one test, provided that the procedure is the same and that the reagents used are from the same reagent bottles.

8.3.2 Check test

In each run, one analysis of a certified reference material (CRM) of the same type of ore shall be carried out in parallel with the analysis of the ore sample(s) under the same conditions. A test sample of the CRM shall be prepared as specified in [7.2](#). The CRM should be of the same type as the sample to be analysed and the properties of the two materials should be sufficiently similar to ensure that, in either case, no significant changes in the analytical procedure will become necessary.

Where the analysis is carried out on several samples of the same type of ore at the same time, the analytical value of one CRM may be used.

8.4 Determination

8.4.1 Decomposition of the test portion

Transfer the test portion ([8.2](#)) to a 250 ml beaker. Moisten with a few millilitres of water, add 25 ml of hydrochloric acid ([5.2](#)), cover with a watch-glass, and heat gently. Increase the heat and digest just below boiling, until no further attack is apparent. Add 2 ml of nitric acid ([5.4](#)) and digest for several minutes. Remove the watch-glass and evaporate the solution until dry. Heat the salts on a hot-plate at 105 °C to 110 °C for 30 min. Add 5 ml of hydrochloric acid ([5.2](#)), cover the beaker with a watch-glass, and warm for several minutes. Add 50 ml of water, stirring to avoid the hydrolysis of titanium, and heat to boiling. Wash the watch-glass and the walls of the beaker and filter the solution through a medium-texture paper containing some filter pulp into a 250 ml beaker. Carefully remove all adhering particles with a rubber-tipped glass rod or moistened filter paper and transfer to the filter, washing three times with diluted hydrochloric acid ([5.3](#)), then with hot water until the filter paper is free of iron. Transfer the paper and residue to a platinum crucible ([6.1](#)).

Evaporate the filtrate to about 100 ml and retain it.

8.4.2 Treatment of the residue

Ignite the filter paper and residue in a platinum crucible ([6.1](#)) at a low temperature (500 °C to 800 °C). Cool, moisten with a few drops of water, and then add three or four drops of sulfuric acid ([5.6](#)) and 10 ml of hydrofluoric acid ([5.5](#)).