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Hårdmetall – Bestämning av kisel i koboltmetallpulver – Fotometrisk metod (ISO 11877:2008, IDT)

Hardmetals – Determination of silicon in cobalt metal powders – Photometric method (ISO 11877:2008, IDT)

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The International Standard ISO 11877:2008 has the status of a Swedish Standard. This document contains the official version of ISO 11877:2008.

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Denna standard är framtagen av kommittén för Pulvermetallurgi, SIS/TK 133.

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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 11877 was prepared by Technical Committee ISO/TC 119, *Powder metallurgy*, Subcommittee SC 4, *Sampling and testing methods for hardmetals*.

Hardmetals — Determination of silicon in cobalt metal powders — Photometric method

1 Scope

This International Standard specifies a photometric method to be used for the determination of the mass fraction of silicon in the range of 20 µg/g to 300 µg/g in cobalt metal powders.

2 Principle

After treatment of the sample with a mixture of acids, the so-formed silicon tetrafluoride is distilled into a mixture of boric acid and sodium molybdate. The obtained molybdatesilicate complex is extracted with methyl-isobutylketone and the absorbance of the solution is measured.

3 Reagents

Reagents of the highest purity and only double-distilled water or water of equivalent purity shall be used.

- 3.1 **Hydrofluoric acid**, 40 % by mass.
- 3.2 **Nitric acid**, 65 % by mass.
- 3.3 **Hydrochloric acid**, 30 % by mass.
- 3.4 **Sulfuric acid**, 96 % by mass.
- 3.5 **Perchloric acid**, 70 % by mass.
- 3.6 **Sodium hydroxide solution**, 30 % by mass.
- 3.7 **Boric acid**.
- 3.8 **Sodium molybdate**.
- 3.9 **Amidosulfuric acid**.
- 3.10 **Methyl-isobutylketone**.
- 3.11 **Acetone**.
- 3.12 **Water**.
- 3.13 **Nitrogen**.
- 3.14 **Silicon standard solution**, 100 µg/ml.

4 Apparatus and equipment

All apparatus and equipment in contact with reagents shall be made of silicon-free material like PTFE (polytetrafluoroethylene), PP (polypropylene), PE (polyethylene) and TPX (methylpenten-polymer).

4.1 Apparatus for acid treatment and distillation, for separation of silicon tetrafluoride.

4.2 Test-tube shaker.

4.3 Refrigerator.

4.4 Platinum thimbles, for weighing the sample.

4.5 Graduated flasks, stock bottles and pipettes.

5 Preparation of acid mixture for sample treatment

Mix 30 ml of hydrofluoric acid, 20 ml of nitric acid, 20 ml of perchloric acid and 30 ml of water in a PTFE flask.

6 Preparation of solutions for receiver

6.1 Solution I

Dissolve 37,2 g of boric acid and 14,4 g of sodium molybdate in water in a graduated PP flask and fill up to 1 000 ml with water.

6.2 Solution II

Dissolve 19,4 g of amidosulfuric acid in a graduated PP flask and fill up to 100 ml with water.

7 Procedure

7.1 Pour 30 ml of solution I and 1 ml of solution II into a graduated 100 ml flask.

7.2 Pour 3 ml of the acid mixture into the distilling flask, and add 100 mg of the sample weighed to a platinum thimble.

7.3 Connect the receiver and distilling flask and start the nitrogen flow.

7.4 Heat the apparatus to 170 ± 10 °C, within approximately 25 min, and keep it at this temperature for 50 min. Control the purge gas during distillation.

7.5 When the distillation time is over, remove the receiver flask and rinse the inlet tube with water.

7.6 Adjust the pH of the receiver solution to 1,3 to 1,4 with sodium hydroxide (approximately 800 µl to 1 000 µl) and leave it for 1 h.

7.7 Add 2,0 ml of hydrochloric acid and 20 ml of methyl-isobutylketone and shake for 1 min. Fill to the stopcock with water and close the flask. Keep the flask at + 5 °C for 1 h.