
**Extenders for paints — Specifications and
methods of test —**

**Part 14:
Cristobalite**

*Matières de charge pour peintures — Spécifications et méthodes d'essai —
Partie 14: Cristobalite*
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ISO 3262-14:2000

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Reference number
ISO 3262-14:2000(E)

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Printed in Switzerland

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 3.

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 part of ISO 3262 may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights.

International Standard ISO 3262-14 was prepared by Technical Committee ISO/TC 35, *Paints and varnishes*, Subcommittee SC 2, *Pigments and extenders*.

Together with the other parts (see below), this part of ISO 3262 cancels and replaces ISO 3262:1975, which has been technically revised. Part 1 comprises the definition of the term extender and a number of test methods that are applicable to most extenders, whilst part 2 and the following parts specify requirements and, where appropriate, particular test methods for individual extenders.

ISO 3262 consists of the following parts, under the general title *Extenders for paints — Specifications and methods of test*:

- Part 1: *Introduction and general test methods*
- Part 2: *Barytes (natural barium sulfate)*
- Part 3: *Blanc fixe*
- Part 4: *Whiting*
- Part 5: *Natural crystalline calcium carbonate*
- Part 6: *Precipitated calcium carbonate*
- Part 7: *Dolomite*
- Part 8: *Natural clay*
- Part 9: *Calcined clay*
- Part 10: *Natural talc/chlorite in lamellar form*
- Part 11: *Natural talc, in lamellar form, containing carbonates*
- Part 12: *Muscovite-type mica*
- Part 13: *Natural quartz (ground)*
- Part 14: *Cristobalite*
- Part 15: *Vitreous silica*
- Part 16: *Aluminium hydroxides*
- Part 17: *Precipitated calcium silicate*
- Part 18: *Precipitated sodium aluminium silicate*

- *Part 19: Precipitated silica*
- *Part 20: Fumed silica*
- *Part 21: Silica sand (unground natural quartz)*
- *Part 22: Flux-calcined kieselguhr*

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Extenders for paints — Specifications and methods of test —

Part 14: Cristobalite

1 Scope

This part of ISO 3262 specifies requirements and corresponding methods of test for cristobalite.

2 Normative references

The following normative documents contain provisions which, through reference in this text, constitute provisions of this part of ISO 3262. For dated references, subsequent amendments to, or revisions of, any of these publications do not apply. However, parties to agreements based on this part of ISO 3262 are encouraged to investigate the possibility of applying the most recent editions of the normative documents indicated below. For undated references, the latest edition of the normative document referred to applies. Members of ISO and IEC maintain registers of currently valid International Standards.

ISO 787-2:1981, *General methods of test for pigments and extenders — Part 2: Determination of matter volatile at 105 °C.*

ISO 787-3:—¹⁾, *General methods of test for pigments and extenders — Part 3: Determination of matter soluble in water — Hot extraction method*

ISO 787-5:1980, *General methods of test for pigments and extenders — Part 5: Determination of oil absorption value.*

ISO 787-7:1981, *General methods of test for pigments and extenders — Part 7: Determination of residue on sieve — Water method — Manual procedure.*

ISO 787-9:1981, *General methods of test for pigments and extenders — Part 9: Determination of pH value of an aqueous suspension.*

ISO 787-14:1973, *General methods of test for pigments — Part 14: Determination of resistivity of aqueous extract.*

ISO 3262-1:1997, *Extenders for paints — Specifications and methods of test — Part 1: Introduction and general test methods.*

ISO 3696:1987, *Water for analytical laboratory use — Specification and test methods.*

ISO 15528:—²⁾, *Paints, varnishes and raw materials for paints and varnishes — Sampling.*

3 Term and definition

For the purposes of this part of ISO 3262, the following term and definition applies.

1) To be published. (Revision of ISO 787-3:1979)

2) To be published. (Revision of ISO 842:1984 and ISO 1512:1991)

3.1

crystalite

a low-temperature modification of crystalline silicon dioxide with a theoretical density of 2,35 g/cm³

4 Requirements and test methods

For cristobalite complying with this part of ISO 3262, the essential requirements are specified in Table 1 and the conditional requirements are listed in Table 2.

Table 1 — Essential requirements

Characteristic	Unit	Requirement	Test method
Cristobalite content, min.	% (m/m)	60	X-ray diffraction
Silica content, min.	% (m/m)	98	See clause 6
Matter volatile at 105 °C, max.	% (m/m)	0,2	ISO 787-2 ^a
Loss on ignition, max.	% (m/m)	0,2 0,5 ^b	ISO 3262-1
Matter soluble in water, max.	% (m/m)	0,2	ISO 787-3
pH value of aqueous suspension	—	6 to 9 6 to 10 ^b	ISO 787-9

^a Another method may be agreed between the interested parties provided it gives the same results.

^b These values take into account the effect on the result if the product has undergone surface treatment.

Table 2 — Conditional requirements

Characteristic	Unit	Requirement	Test method
Residue on 45 µm sieve	% (m/m)	To be agreed between the interested parties	ISO 787-7
Particle size distribution (instrumental method)	% (m/m)	To be agreed between the interested parties ^a	
Colour	—	To be agreed between the interested parties	ISO 3262-1
Lightness	—		To be agreed between the interested parties ^b
Resistivity of aqueous extract	Ω · m		ISO 787-14
Oil absorption value	g/100 g		ISO 787-5

^a A general description of a sedimentation method using X-ray absorption is given in EN 725-5:1996, *Advanced technical ceramics — Methods of test for ceramic powders — Part 5: Determination of the particle size distribution*.

^b Test method in preparation.

5 Sampling

Take a representative sample of the product to be tested, as described in ISO 15528.

6 Determination of silica content

6.1 Reagents

During the analysis, use only reagents of recognized analytical grade and only water of at least grade 3 purity as defined in ISO 3696.

6.1.1 Sulfuric acid, diluted 1 + 1.

Add 1 part by volume of concentrated sulfuric acid, approximately 96 % (m/m), $\rho \approx 1,84$ g/ml, slowly to 1 part by volume of water.

6.1.2 Hydrofluoric acid, approximately 40 % (m/m), $\rho \approx 1,13$ g/ml.

6.2 Apparatus

Use ordinary laboratory apparatus and glassware, together with the following:

6.2.1 Platinum dish.

6.2.2 Muffle furnace.

6.3 Procedure

Weigh, to the nearest 1 mg, approximately 2 g of the test sample (clause 5), previously dried at 105 °C in accordance with ISO 787-2, into the tared platinum dish (6.2.1), ignite in the muffle furnace (6.2.2) at $(1\,000 \pm 25)$ °C to constant mass (m_1) and allow to cool in a desiccator containing phosphorus pentoxide.

Add approximately 1 ml of sulfuric acid (6.1.1). Heat the platinum dish gently until fuming ceases and then continue the heating at 900 °C for 15 min in the muffle furnace. Remove from the furnace, allow to cool in a desiccator and weigh to the nearest 0,1 mg (m_2).

Add to the residue in the platinum dish 15 ml of hydrofluoric acid (6.1.2) and 1 ml of sulfuric acid (6.1.1) and evaporate to a syrup, taking care to avoid loss by spitting. Cool the dish and wash the sides down with small quantities of water. Then add a further 10 ml of hydrofluoric acid and evaporate to dryness. Heat the residue on a hot-plate until white fumes are no longer evolved, then ignite in the muffle furnace at 900 °C for 15 min. Remove the dish from the furnace, cool in the desiccator and weigh to the nearest 0,1 mg (m_3).

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6.4 Expression of results

Calculate the silica content $w(\text{SiO}_2)$, expressed as a percentage by mass, using the equation:

$$w(\text{SiO}_2) = \frac{m_2 - m_3}{m_1} \times 100$$

where

m_1 is the mass, in grams, of the ignited residue;

m_2 is the mass, in grams, after treatment with sulfuric acid and igniting;

m_3 is the mass, in grams, after treatment with hydrofluoric acid and igniting.

7 Test report

The test report shall contain at least the following information:

- all details necessary to identify the product tested;
- a reference to this part of ISO 3262 (ISO 3262-14);
- the results of the tests and whether or not the product complies with the relevant specification limits;
- any deviation from the test methods specified;
- the dates of the tests.

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ICS 87.060.10

Price based on 3 pages