# INTERNATIONAL STANDARD

ISO 591-1

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# **Titanium dioxide pigments for paints** — Part 1:

# Specifications and methods of test

Pigments de dioxyde de titane pour peintures —

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<u>ISO 591-1:2000</u> https://standards.iteh.ai/catalog/standards/sist/4fc18d4d-2935-4d48-a649-4b5721b6673e/iso-591-1-2000



Reference number ISO 591-1:2000(E)

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

International Standard ISO 591-1 was prepared by Technical Committee ISO/TC 35, *Paints and varnishes*, Subcommittee SC 2, *Pigments and extenders*.

This part of ISO 591 cancels and replaces ISO 591:1977, which has been technically revised. In contrast to ISO 591:1977, which specified the Nakazono reductor method for the determination of the titanium dioxide content and allowed other methods to be used by agreement between the interested parties, ISO 591-1 contains two methods (see clause 7).

ISO 591 consists of the following parts, under the general title *Titanium dioxide pigments for paints*: https://standards.iteh.ai/catalog/standards/sist/4fc18d4d-2935-4d48-a649-

- Part 1: Specifications and methods of test  $\frac{4b5721b6673e}{iso-591-1-2000}$
- ,
- Part 2: Determination of the content of secondary constituents

## Titanium dioxide pigments for paints —

# Part 1: **Specifications and methods of test**

## 1 Scope

This part of ISO 591 specifies the requirements and corresponding methods of test for titanium dioxide pigments for paints.

## 2 Normative references

The following normative documents contain provisions which, through reference in this text, constitute provisions of this part of ISO 591. 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 591 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 385-1:1984, Laboratory glassware — Burettes 5 Part 20 General requirements. https://standards.iteh.ai/catalog/standards/sist/4fc18d4d-2935-4d48-a649

ISO 648:1977, Laboratory glassware — One-mark pipettes.591-1-2000

ISO 787-1:1982, General methods of test for pigments and extenders — Part 1: Comparison of colour of pigments.

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

ISO 787-3:2000, 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 787-9:1981, 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 787-18:1983, General methods of test for pigments and extenders — Part 18: Determination of residue on sieve — Mechanical flushing procedure.

ISO 787-24:1985, General methods of test for pigments and extenders — Part 24: Determination of relative tinting strength of coloured pigments and relative scattering power of white pigments — Photometric methods.

ISO 787-25:1993, General methods of test for pigments and extenders — Part 25: Comparison of the colour, in full-shade systems, of white, black and coloured pigments – Colorimetric method.

ISO 1042:1998, Laboratory glassware — One-mark volumetric flasks.

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

ISO 15528:2000, Paints, varnishes and raw materials for paints and varnishes — Sampling.

## 3 Term and definition

For the purposes of this part of ISO 591, the following term and definition apply.

## 3.1

## titanium dioxide pigment

pigment consisting essentially of titanium dioxide (TiO<sub>2</sub>) of the anatase or the rutile crystal structure, as determined by X-ray examination

## 4 Classification

## 4.1 Types

This part of ISO 591 covers two types of titanium dioxide pigment, as follows:

Type A: Anatase type

Type R: Rutile type

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## 4.2 Grades

ISO 591-1:2000 The pigments are further classified into the following grades: ds/sist/4fc18d4d-2935-4d48-a649-4b5721b6673e/iso-591-1-2000

Grade A1	)
Grade A2	Fype A
	,
Grade R1	)
Grade R2	Type R
Grade R3	

## 5 Required characteristics and associated tolerances

## 5.1 Appearance

The material shall be in the form of a soft, dry powder or in such a condition that it may be readily reduced thereto by crushing under a palette knife, without any grinding action.

## 5.2 Other characteristics

**5.2.1** For titanium dioxide pigments complying with this part of ISO 591, the essential requirements are specified in Table 1 and the conditional requirements are listed in Table 2. The conditional requirements shall be specified by agreement between the interested parties.

The requirement for matter volatile at 105 °C after preconditioning in Table 2 shall only apply if this characteristic is explicitly specified by the interested parties or in a contract.

#### 5.2.2 The agreed reference pigment referred to in Table 2 shall comply with the requirements of Table 1.

	Re					
Characteristic	Туре А			Type R	Method of test	
	A1	A2	R1	R2	R3	
Titanium dioxide content % (by mass), min.	98	92	97	90	80	See clause 7
Matter volatile at 105 °C at point of acceptance % (by mass), max.	0,5	0,8	0,5	To be agreed between the interested parties		ISO 787-2
Matter soluble in water % (by mass), max.	0,6	0,5	0,6	0,5	0,7	ISO 787-3ª
Residue on sieve (45 μm) % (by mass), max.	0,1	0,1	0,1	0,1	0,1	ISO 787-18 or other agreed method

Table 1 — Essential requirements

# iTeh STANDARD PREVIEW Table 2 — Conditional requirements

	Requirement depending on type and grade					
Characteristic https://sta	ndards.iteh.ai/c	pe ASO 591-1:2 atalog/standards/s 721b6673e/iso-5	ist/4fc18d4d	<b>Type R</b> -2935-4d48-a R2	1649- R3	Method of test
Colour Similar to that of the agreed reference pigment (see 5.2.2)						ISO 787-1 or ISO 787-25 <sup>a</sup>
Scattering power	To be agreed between the interested parties				ISO 787-24	
Matter volatile at 105 °C after 24 h preconditioning at $(23 \pm 2)$ °C and $(50 \pm 5)$ % relative humidity % (by mass), max. <sup>b</sup>	0,5	0,8	0,5	1,5	2,5	ISO 787-2
pH of aqueous suspension	Similar to that of the agreed reference pigment (see 5.2.2)					ISO 787-9
Oil absorption value						ISO 787-5
Resistivity of aqueous extract	_	Similar to that of the agreed reference pigment (see 5.2.2) Similar to that of the agreed reference pigment (see 5.2.2)		ISO 787-14		
a By agreement only.	•					
<sup>b</sup> See 5.2.1.						

## 6 Sampling

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

## 7 Determination of titanium dioxide content

## 7.1 General

For the determination of the titanium dioxide content, two methods (A and B) are provided. In cases of dispute, either of the methods may be used as the referee method, by agreement between the interested parties.

Method A uses aluminium metal as the reducing agent. It is suitable for individual determinations and for routine analysis.

Method B uses chromium(II) chloride and zinc amalgam as reducing agents and can be automated. It is particularly suitable as a routine method.

## 7.2 Method A: Aluminium reduction method

## 7.2.1 Principle

A test portion of the dried sample is dissolved in sulfuric acid, with the addition of ammonium sulfate. The titanium(IV) is then reduced to titanium(III) with aluminium under a carbon dioxide atmosphere. The solution is then titrated with ammonium iron(III) sulfate, using ammonium thiocyanate as indicator.

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## 7.2.2 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. 4b5721b6673e/iso-591-1-2000

### WARNING — Use the reagents in accordance with the appropriate health and safety regulations.

- **7.2.2.1 Hydrochloric acid**, concentrated, approximately 37 % (by mass),  $\rho \approx 1,19$  g/ml.
- **7.2.2.2** Sulfuric acid, concentrated, approximately 96 % (by mass),  $\rho \approx$  1,84 g/ml.
- 7.2.2.3 Ammonium sulfate.
- **7.2.2.4 Sodium hydrogen carbonate**, saturated solution.

Prepare this solution at the time of use. About 10 g of sodium hydrogen carbonate (NaHCO<sub>3</sub>) to 90 ml of water are required.

### 7.2.2.5 Ammonium thiocyanate indicator.

Dissolve 24,5 g of ammonium thiocyanate ( $NH_4SCN$ ) in 80 ml of hot water, filter, cool to room temperature and dilute to 100 ml. Keep in a well-stoppered, dark-coloured bottle.

**7.2.2.6** Ammonium iron(III) sulfate, standard volumetric solution, 1 ml equivalent to 0,0048 g of TiO<sub>2</sub>.

### 7.2.2.6.1 Preparation

Dissolve, in a 1000 ml one-mark volumetric flask, 30 g of freshly prepared ammonium iron(III) sulfate  $[(FeNH_4(SO_4)_2 \cdot 12H_2O]$  in 300 ml of water containing 15 ml of sulfuric acid (7.2.2.2). Add potassium permanganate solution (7.2.2.7), drop by drop, until the solution is pink coloured. Dilute to the mark with water and mix well. Filter if the solution is cloudy.

## 7.2.2.6.2 Standardization

Standardize the solution using 190 mg to 210 mg of titanium dioxide reference standard, dried to constant mass at  $(105 \pm 2)$  °C, following the procedure described in 7.2.4.3.

Calculate the titanium dioxide equivalent  $T_1$  of the solution, expressed in grams of TiO<sub>2</sub> per millilitre, using the equation

$$T_1 = \frac{m_1 \times P}{V_1 \times 100}$$

where

- $m_1$  is the mass, in grams, of reference standard used;
- *P* is the titanium dioxide content of the reference standard, expressed as a percentage by mass (i.e. 99,74 %);
- $V_1$  is the volume, in millilitres, of the ammonium iron(III) sulfate solution required in the titration.

## **7.2.2.7** Potassium permanganate, standard volumetric solution, $c(\frac{1}{5} \text{KMnO}_4) = 0,1 \text{ mol/l.}$

Dissolve 3,1607 g of potassium permanganate in 500 ml of water in a 1000 ml one-mark volumetric flask, dilute to the mark with water and mix well.

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7.2.2.8 Aluminium metal, electrolytic grade, for example in the form of foil, sheet or cut wire.

## 7.2.3 Apparatus

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Use ordinary laboratory apparatus, and burettes, pipettes, and one-mark volumetric flasks complying with the requirements of ISO 385-1, ISO 648 and ISO 1042, respectively, together with the following:

**7.2.3.1 Delivery tube:** an essentially U-shaped glass tube, 4 mm in internal diameter, comprising a horizontal section and two vertical arms, one of the vertical arms being 150 mm long and the other 75 mm long.

As an alternative, an absorption apparatus as shown in Figure 1 or a Contat-Göckel cap as shown in Figure 2 may be used.

**7.2.3.2** Weighing bottle, wide-mouth, with an external-fitting cap, and no larger than necessary for the test portion.

- **7.2.3.3 Oven**, capable of maintaining a temperature of  $(105 \pm 2)$  °C.
- **7.2.3.4 Desiccator**, containing a suitable desiccant, for example silica gel.

## 7.2.4 Procedure

7.2.4.1 General

Carry out the determination in duplicate.

## 7.2.4.2 Test portion

Take about 10 g of the sample (see clause 6) and dry it in the open weighing bottle (7.2.3.2) at  $(105 \pm 2)$  °C to constant mass. Cap the bottle and allow it to cool to room temperature in the desiccator (7.2.3.4).

Weigh out, to the nearest 0,1 mg, 190 mg to 210 mg of this test sample  $(m_2)$ .