

# **SLOVENSKI STANDARD**

## **SIST EN ISO 18618:2018**

**01-september-2018**

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### **Zobozdravstvo - Skladno delovanje sistemov CAD/CAM (ISO 16818:2018)**

Dentistry - Interoperability of CAD/CAM-systems (ISO 16818:2018)

Zahnheilkunde - Interoperabilität der CAD/CAM-Systeme (ISO 16818:2018)

Médecine bucco-dentaire - Interopérabilité des systèmes de CFAO (ISO 18618:2018)

**Ta slovenski standard je istoveten z: EN ISO 18618:2018**

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#### **ICS:**

11.060.01	Zobozdravstvo na splošno	Dentistry in general
35.240.80	Uporabniške rešitve IT v zdravstveni tehniki	IT applications in health care technology

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EUROPEAN STANDARD  
NORME EUROPÉENNE  
EUROPÄISCHE NORM

**EN ISO 18618**

June 2018

ICS 11.060.01

English Version

**Dentistry - Interoperability of CAD/CAM systems (ISO 18618:2018)**

Médecine bucco-dentaire - Interopérabilité des systèmes de CFAO (ISO 18618:2018)

Zahnheilkunde - Interoperabilität der CAD/CAM-Systeme (ISO 18618:2018)

This European Standard was approved by CEN on 3 May 2018.

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## European foreword

This document (EN ISO 18618:2018) has been prepared by Technical Committee ISO/TC 106 “Dentistry” in collaboration with Technical Committee CEN/TC 55 “Dentistry” the secretariat of which is held by DIN.

This European Standard shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by December 2018, and conflicting national standards shall be withdrawn at the latest by December 2018.

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## Endorsement notice

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# INTERNATIONAL STANDARD

**ISO  
18618**

First edition  
2018-05

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## Dentistry — Interoperability of CAD/ CAM systems

*Médecine bucco-dentaire — Interopérabilité des systèmes de CFAO*

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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.

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](http://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](http://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](http://www.iso.org/iso/foreword.html). (standards.iteh.ai)

This document was prepared by Technical Committee ISO/TC 106, *Dentistry*, Subcommittee SC 9, *CAD/CAM Systems*.

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## Introduction

Manufacturers of dental CAD/CAM systems differ in how they exchange manufacturing information and three dimensional data. This causes difficulty in data processing, design processes, and manufacturing processes for users of those systems. In order to overcome these interoperability issues, this document has been prepared to facilitate open interoperability between CAD/CAM systems in dentistry.

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# Dentistry — Interoperability of CAD/CAM systems

## 1 Scope

This document specifies an extensible markup language (XML) format to facilitate the transfer of dental case data and CAD/CAM data between software systems.

## 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 1942, *Dentistry — Vocabulary*

ISO 3166-1, *Codes for the representation of names of countries and their subdivisions — Part 1: Country codes*

ISO 3950, *Dentistry — Designation system for teeth and areas of the oral cavity*

ISO 16443, *Dentistry — Vocabulary for dental implants systems and related procedure*

ISO 18739, *Dentistry — Vocabulary of process chain for CAD/CAM systems*

ISO 19429:2015, *Dentistry — Designation system for dental implants' after 'ISO 18739, Dentistry — Vocabulary of process chain for CAD/CAM systems'*

W3C — Extensible Markup Language (XML) 1.0 (Fifth Edition), November 2008

W3C XML Schema Definition Language (XSD) 1.1, 2012

## 3 Terms and definitions

For the purposes of this document, the terms and definitions given in ISO 1942, ISO 16443, ISO 18739, W3C XML1.0, W3C XSD 1.1 and the following apply.

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 <https://www.iso.org/obp>

**NOTE 1** Throughout the IDS (interface for dental CAD/CAM systems) schema there are terms that have special meaning or definition. Understanding the use of these terms is the key to well-defined IDS documents that all parties can understand universally.

**NOTE 2** The IDS schema defines several peer level nodes immediately within the enveloping root <IDS> element that organizes the IDS document into structures for specific transactions. They represent a submission, a query, an update of a previous submission, a notification of an event or status change and a series of catalogs. A single IDS document can contain a combination of different transaction nodes or consist of only a single transactional node.

**NOTE 3** In addition to the transactional nodes mentioned above, the IDS schema also defines several nodes that provide traceability and source identification features as well as provide information on how to reply to a document transaction.

### 3.1 General terms

## ISO 18618:2018(E)

### 3.1.1

#### **broker**

entity that acts as a middleman or intermediary

Note 1 to entry: Such organizations may take multiple orders from multiple sources and consolidate them into a single order for a Provider or they may take single orders from an originator and split them among multiple providers or they may just pass orders through between originators and providers.

### 3.1.2

#### **originator**

entity (organization or person) that is responsible for creating the current document, order, submission, etc.

Note 1 to entry: As such, they are the “originator” of the data being exchanged.

Note 2 to entry: Most often an originator would be a dental practice. In some cases, an originator may be a dental laboratory that is outsourcing work to another lab.

### 3.1.3

#### **provider**

entity (company, lab, etc.) that is responsible for providing the services or products that are being requested in an order

Note 1 to entry: Most often, this would be a dental laboratory or manufacturer.

## 3.2 Terms and definitions relating to XML content

### 3.2.1

#### **CADDataCatalog**

collection of nodes describing CAD data associated with one or more of the orders and/or restorations

Note 1 to entry: It can include digital scan and/or design files, etc.  
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### 3.2.2

#### **case**

set of one or more orders for dental appliances, products or services, all of which are being submitted for a single patient

EXAMPLE A case might contain one order for a crown, and another order for a bridge.

### 3.2.3

#### **catalogs**

data that are referenced in other elements or areas

Note 1 to entry: The catalogs are subdivided by the data they are grouping, making it easier to manage and reference.

### 3.2.4

#### **DeliveryRequest**

information for the out-going, finished order, which will be sent to the originator (or an originator's agent) as a separate delivery

Note 1 to entry: A delivery may be physical, electronic, or both.

### 3.2.5

#### **dentist**

dentist or responsible clinician who requested the order

**3.2.6****DentistCatalog**

collection of *dentist* (3.2.5) nodes that provides attribute and elements to define the dentists being referenced within this document

Note 1 to entry: The definition can include billing information, license information, etc.

**3.2.7****ExtraInfo**

child node that can be used to extend the schema with undefined XML

**EXAMPLE** Many of the elements will contain child nodes with the suffix “ExtraInfo” (i.e. <DentistExtraInfo>, <OrderExtraInfo>, etc. These sections are intended to be areas that can be used to extend the defined schema with proprietary or undefined XML. For example, an implementation may use one (or more) of these sections to embed XML that is only of use to the implementer for an internal workflow. Another use could be two business partners using these sections to experiment with XML they intend to propose for future versions or to pass proprietary XML they have previously defined between themselves. The IDS schema and XSD will ignore the contents of these sections so they will not be validated as part of the IDS schema. It is highly recommended that if these sections are used, that any XML be enclosed within some proprietary element tag so that if the XML document passes through multiple handlers there are no conflicts:

```
<DentistExtraInfo>
```

```
  <MyCompanyData>
```

```
    data specific and of use only to “MyCompany”...
```

```
  </MyCompanyData>
```

```
</DentistExtraInfo>
```

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**3.2.8****FileCatalog**

collection of <IDSFile> nodes that describe files associated with the <Case>, <Order> or CAD data (scans, design files, etc.)

**3.2.9****host service**

system that receives the IDS document and processes the contents

**3.2.10****IdMapCatalog**

collection of <IdMapItem> nodes which provide a means of defining alternate identifiers for key elements within the IDS

**3.2.11****notification**

means for publishing or returning a defined status, event or message related to an order

Note 1 to entry: Within the notification node is an untyped element that can be defined according to the needs of the parties exchanging information.

**3.2.12****order**

request for a self-contained dental appliance, service or product that is being requested by an originator

Note 1 to entry: Each order in a case might be created or manufactured by a different provider. Each order contains its own delivery (or reply) instruction nodes.