

SLOVENSKI STANDARD SIST EN ISO 13017:2012/A1:2016

01-februar-2016

Zobozdravstvo - Magnetni priključki - Dopolnilo A1 (ISO 13017:2012/Amd 1:2015)

Dentistry - Magnetic attachments - Amendment 1 (ISO 13017:2012/Amd 1:2015)

Zahnheilkunde - Magnetische Retentionselemente - Änderung 1 (ISO 13017:2012/Amd 1:2015)

Médecine bucco-dentaire - Attaches magnétiques - Amendement 1 (ISO 13017:2012/Amd 1:2015) (standards.iteh.ai)

Ta slovenski standard je istoveten za EN ISO 13017:2012/A1:2015

976f8d9a537b/sist-en-iso-13017-2012-a1-2016

ICS:

11.060.10 Zobotehnični materiali Dental materials

SIST EN ISO 13017:2012/A1:2016 en

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EUROPEAN STANDARD NORME EUROPÉENNE EUROPÄISCHE NORM EN ISO 13017:2012/A1

December 2015

ICS 11.060.10

English Version

Dentistry - Magnetic attachments - Amendment 1 (ISO 13017:2012/Amd 1:2015)

Médecine bucco-dentaire - Attaches magnétiques - Amendement 1 (ISO 13017:2012/Amd 1:2015)

Zahnheilkunde - Magnetische Retentionselemente - Änderung 1 (ISO 13017:2012/Amd 1:2015)

This amendment A1 modifies the European Standard EN ISO 13017:2012; it was approved by CEN on 3 October 2015.

CEN members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for inclusion of this amendment into the relevant national standard without any alteration. Up-to-date lists and bibliographical references concerning such national standards may be obtained on application to the CEN-CENELEC Management Centre or to any CEN member.

This amendment exists in three official versions (English, French, German). A version in any other language made by translation under the responsibility of a CEN member into its own language and notified to the CEN-CENELEC Management Centre has the same status as the official versions.

CEN members are the national standards bodies of Austria, Belgium, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, Former Yugoslav Republic of Macedonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey and United Kingdom.

https://standards.iteh.ai/catalog/standards/sist/a45925fc-5018-4963-8642-976f8d9a537b/sist-en-iso-13017-2012-a1-2016



EUROPEAN COMMITTEE FOR STANDARDIZATION COMITÉ EUROPÉEN DE NORMALISATION EUROPÄISCHES KOMITEE FÜR NORMUNG

CEN-CENELEC Management Centre: Avenue Marnix 17, B-1000 Brussels

EN ISO 13017:2012/A1:2015 (E)

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EN ISO 13017:2012/A1:2015 (E)

European foreword

This document (EN ISO 13017:2012/A1:2015) 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 Amendment to the European Standard EN ISO 13017:2012 shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by June 2016, and conflicting national standards shall be withdrawn at the latest by June 2016.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. CEN [and/or CENELEC] shall not be held responsible for identifying any or all such patent rights.

According to the CEN-CENELEC Internal Regulations, the national standards organizations of the following countries are bound to implement this European Standard: Austria, Belgium, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, Former Yugoslav Republic of Macedonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey and the United Kingdom.

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The text of ISO 13017:2012/Amd 1:2015 has been approved by CEN as EN ISO 13017:2012/A1:2015 without any modification.

SIST EN ISO 13017:2012/A1:2016

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INTERNATIONAL STANDARD

ISO 13017

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Dentistry — Magnetic attachmentsAMENDMENT 1

Médecine bucco-dentaire — Attaches magnétiques AMENDEMENT 1

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ISO 13017:2012/Amd.1:2015(E)

Foreword

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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. www.iso.org/directives

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For an explanation on the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the WTO principles in the Technical Barriers to Trade (TBT), see the following URL: Foreword - Supplementary information

The committee responsible for this document is ISO/TC 106, Dentistry, Subcommittee SC 2, Prosthodontic materials.