

SLOVENSKI STANDARD SIST EN ISO 5832-3:2012

01-oktober-2012

Vsadki (implantati) za kirurgijo - Kovinski materiali - 3. del: Kovana zlitina (ISO 5832-3:1996)

Implants for surgery - Metallic materials - Part 3: Wrought titanium 6-aluminium 4-vanadium alloy (ISO 5832-3:1996)

Chirurgische Implantate - Metallische Werkstoffe - Teil 3: Titan 6-Aluminium 4-Vanadium Knetlegierungen (ISO 5832-3:1996) NDARD PREVIEW

Implants chirurgicaux - Produits à base de métaux - Partie 3: Alliage à forger à base de titane, d'aluminium 6 et de vanadium 4 (ISQ 5832-3;1996)

https://standards.iteh.ai/catalog/standards/sist/74c6e81a-817c-411b-aa52-

Ta slovenski standard je istoveten z: EN ISO 5832-3-2012

ICS:

11.040.40 Implantanti za kirurgijo, protetiko in ortetiko Implants for surgery, prosthetics and orthotics
77.120.50 Titan in titanove zlitine Titanium and titanium alloys

SIST EN ISO 5832-3:2012 en

SIST EN ISO 5832-3:2012

iTeh STANDARD PREVIEW (standards.iteh.ai)

SIST EN ISO 5832-3:2012

https://standards.iteh.ai/catalog/standards/sist/74c6e81a-817c-411b-aa52-b1f2e5af3c59/sist-en-iso-5832-3-2012

EUROPEAN STANDARD

EN ISO 5832-3

NORME EUROPÉENNE

EUROPÄISCHE NORM

May 2012

ICS 11.040.40

English Version

Implants for surgery - Metallic materials - Part 3: Wrought titanium 6-aluminium 4-vanadium alloy (ISO 5832-3:1996)

Implants chirurgicaux - Produits à base de métaux - Partie 3: Alliage à forger à base de titane, d'aluminium 6 et de vanadium 4 (ISO 5832-3:1996) Chirurgische Implantate - Metallische Werkstoffe - Teil 3: Titan 6-Aluminium 4-Vanadium Knetlegierungen (ISO 5832-3:1996)

This European Standard was approved by CEN on 28 April 2012.

CEN members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for giving this European Standard the status of a 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 European Standard 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, 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/74c6e81a-817c-411b-aa52-b1f2e5af3c59/sist-en-iso-5832-3-2012



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

Management Centre: Avenue Marnix 17, B-1000 Brussels

EN ISO 5832-3:2012 (E)

Contents	Pag
Foreword	

iTeh STANDARD PREVIEW (standards.iteh.ai)

SIST EN ISO 5832-3:2012 https://standards.iteh.ai/catalog/standards/sist/74c6e81a-817c-411b-aa52-b1f2e5af3c59/sist-en-iso-5832-3-2012

EN ISO 5832-3:2012 (E)

Foreword

The text of ISO 5832-3:1996 has been prepared by Technical Committee ISO/TC 150 "Implants for surgery" of the International Organization for Standardization (ISO) and has been taken over as EN ISO 5832-3:2012 by 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 November 2012, and conflicting national standards shall be withdrawn at the latest by November 2012.

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, 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.

iTeh STANEndersement notice VIEW

The text of ISO 5832-3:1996 has been approved by CEN as a EN ISO 5832-3:2012 without any modification.

<u>SIST EN ISO 5832-3:2012</u> https://standards.iteh.ai/catalog/standards/sist/74c6e81a-817c-411b-aa52-b1f2e5af3c59/sist-en-iso-5832-3-2012 **SIST EN ISO 5832-3:2012**

iTeh STANDARD PREVIEW (standards.iteh.ai)

SIST EN ISO 5832-3:2012

https://standards.iteh.ai/catalog/standards/sist/74c6e81a-817c-411b-aa52-b1f2e5af3c59/sist-en-iso-5832-3-2012

SIST EN ISO 5832-3:2012

INTERNATIONAL STANDARD

ISO 5832-3

> Third edition 1996-07-01

Implants for surgery — Metallic materials —

Part 3:

Wrought titanium 6-aluminium 4-vanadium alloy

iTen S Implants chirurgicaux — Produits à base de métaux —

Partie 3: Alliage à forger à base de titane, d'aluminium 6 et de vanadium 4

SIST EN ISO 5832-3:2012

https://standards.iteh.ai/catalog/standards/sist/74c6e81a-817c-411b-aa52-b1f2e5af3c59/sist-en-iso-5832-3-2012



ISO 5832-3:1996(E)

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.

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.

International Standard ISO 5832-3 was prepared by Technical Committee ISO/TC 150, Implants for surgery, Subcommittee SC 1, Materials.

SIST EN ISO 5832-3:2012

This third edition cancels ps/and-darteplaces at a the stander of the standard edition 817c-411b-aa52-(ISO 5832-3:1990), which has been technically revised 9/sist-en-iso-5832-3-2012

ISO 5832 consists of the following parts, under the general title *Implants* for surgery — Metallic materials:

- Part 1: Wrought stainless steel
- Part 2: Unalloyed titanium
- Part 3: Wrought titanium 6-aluminium 4-vanadium alloy
- Part 4: Cobalt-chromium-molybdenum casting alloy
- Part 5: Wrought cobalt-chromium-tungsten-nickel alloy
- Part 6: Wrought cobalt-nickel-chromium-molybdenum alloy
- Part 7: Forgeable and cold-formed cobalt-chromium-nickel-molybdenum-iron alloy
- Part 8: Wrought cobalt-nickel-chromium-molybdenum-tungsten-iron alloy

© ISO 1996

All rights reserved. Unless otherwise specified, no part of this publication may be reproduced or utilized in any form or by any means, electronic or mechanical, including photocopying and microfilm, without permission in writing from the publisher.

International Organization for Standardization Case postale 56 • CH-1211 Genève 20 • Switzerland

Printed in Switzerland

ISO 5832-3:1996(E)

- © ISO
- Part 9: Wrought high nitrogen stainless steel
- Part 10: Wrought titanium 5-aluminium 2,5-iron alloy
- Part 11: Wrought titanium 6-aluminium 7-niobium alloy
- Part 12: Wrought cobalt-chromium-molybdenum alloy

iTeh STANDARD PREVIEW (standards.iteh.ai)

<u>SIST EN ISO 5832-3:2012</u> https://standards.iteh.ai/catalog/standards/sist/74c6e81a-817c-411b-aa52-b1f2e5af3c59/sist-en-iso-5832-3-2012