

**SLOVENSKI STANDARD
SIST EN ISO 5832-6:2022****01-junij-2022****Nadomešča:****SIST EN ISO 5832-6:2019**

Vsadki (implantati) za kirurgijo - Kovinski materiali - 6. del: Kobalt-nikelj-krom-molibdenova kovana zlitina (ISO 5832-6:2022)

Implants for surgery - Metallic materials - Part 6: Wrought cobalt-nickel-chromium-molybdenum alloy (ISO 5832-6:2022)

Chirurgische Implantate - Metallische Werkstoffe - Teil 6: Kobalt-Nickel-Chrom-Molybdän-Schmiedelegerung (ISO 5832-6:2022)

Implants chirurgicaux - Matériaux métalliques - Partie 6: Alliage corroyé à base de cobalt, de nickel, de chrome et de molybdène (ISO 5832-6:2022)

[SIST EN ISO 5832-6:2022](https://standards.iteh.ai/catalog/standards/sist/e4339523-3d508222-5832-6-2022)[https://standards.iteh.ai/catalog/standards/sist/e4339523-](https://standards.iteh.ai/catalog/standards/sist/e4339523-3d508222-5832-6-2022)**Ta slovenski standard je istoveten z: EN ISO 5832-6:2022****ICS:**

11.040.40	Implantanti za kirurgijo, protetiko in ortetiko	Implants for surgery, prosthetics and orthotics
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SIST EN ISO 5832-6:2022**en,fr,de**

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EUROPEAN STANDARD

EN ISO 5832-6

NORME EUROPÉENNE

EUROPÄISCHE NORM

March 2022

ICS 11.040.40

Supersedes EN ISO 5832-6:2019

English Version

Implants for surgery - Metallic materials - Part 6: Wrought cobalt-nickel-chromium-molybdenum alloy (ISO 5832-6:2022)

Implants chirurgicaux - Matériaux métalliques - Partie 6: Alliage corroyé à base de cobalt, de nickel, de chrome et de molybdène (ISO 5832-6:2022)

Chirurgische Implantate - Metallische Werkstoffe - Teil 6: Kobalt-Nickel-Chrom-Molybdän-Schmiedelegerung (ISO 5832-6:2022)

This European Standard was approved by CEN on 20 March 2022.

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EUROPEAN COMMITTEE FOR STANDARDIZATION
COMITÉ EUROPÉEN DE NORMALISATION
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CEN-CENELEC Management Centre: Rue de la Science 23, B-1040 Brussels

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European foreword

This document (EN ISO 5832-6:2022) has been prepared by Technical Committee ISO/TC 150 "Implants for surgery" in collaboration with Technical Committee CEN/TC 285 "Non-active surgical implants" 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 September 2022, and conflicting national standards shall be withdrawn at the latest by September 2022.

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

This document supersedes EN ISO 5832-6:2019.

Any feedback and questions on this document should be directed to the users' national standards body/national committee. A complete listing of these bodies can be found on the CEN website.

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INTERNATIONAL
STANDARD

ISO
5832-6

Third edition
2022-03

**Implants for surgery — Metallic
materials —**

Part 6:
**Wrought cobalt-nickel-chromium-
molybdenum alloy**

Implants chirurgicaux — Matériaux métalliques —

*Partie 6: Alliage corroyé à base de cobalt, de nickel, de chrome et de
molybdène*

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ISO 5832-6:2022(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.

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).

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For an explanation of 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 www.iso.org/iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 150, *Implants for surgery*, Subcommittee SC 1, *Materials*, in collaboration with the European Committee for Standardization (CEN) Technical Committee CEN/TC 285, *Non-active surgical implants*, in accordance with the Agreement on technical cooperation between ISO and CEN (Vienna Agreement).

This third edition cancels and replaces the second edition (ISO 5832-6:1997) which has been technically revised.

The main changes compared to the previous edition are as follows:

- the introduction has been updated;
- requirements for boron in [Table 1](#) has been added;
- information on grain size in [5.2](#) has been added;
- requirements for tensile properties in [Table 2](#) have been updated and harmonized with the ISO 5832 series.

A list of all parts in the ISO 5832 series can be found on the ISO website.

Any feedback or questions on this document should be directed to the user's national standards body. A complete listing of these bodies can be found at www.iso.org/members.html.