INTERNATIONAL STANDARD

ISO 5832-7

Third edition 2016-11-15

Implants for surgery — Metallic materials —

Part 7:

Forgeable and cold-formed cobaltchromium-nickel-molybdenum-iron

iTeh STANDARD PREVIEW

Simplants chirurgicaux—Produits à base de métaux—
Partie 7: Alliage à forger mis en forme à froid à base de cobalt, de chrome, de nickel, de molybdène et de fer

https://standards.iteh.ai/catalog/standards/sist/501868f8-7a61-40ac-83f8-db61e3bdec47/iso-5832-7-2016



iTeh STANDARD PREVIEW (standards.iteh.ai)

ISO 5832-7:2016 https://standards.iteh.ai/catalog/standards/sist/501868f8-7a61-40ac-83f8-db61e3bdec47/iso-5832-7-2016



COPYRIGHT PROTECTED DOCUMENT

© ISO 2016, Published in Switzerland

All rights reserved. Unless otherwise specified, no part of this publication may be reproduced or utilized otherwise in any form or by any means, electronic or mechanical, including photocopying, or posting on the internet or an intranet, without prior written permission. Permission can be requested from either ISO at the address below or ISO's member body in the country of the requester.

ISO copyright office Ch. de Blandonnet 8 • CP 401 CH-1214 Vernier, Geneva, Switzerland Tel. +41 22 749 01 11 Fax +41 22 749 09 47 copyright@iso.org www.iso.org

Co	Contents				
Fore	eword		iv		
Intr	oductio	n	v		
1	Scop	e	1		
2	Nori	native references	1		
3	Tern	ns and definitions	1		
4	Cher	nical composition	1		
5	Micr	ostructure	2		
	5.1	Grain size			
	5.2	Inclusion content	2		
6	Mec	nanical properties	2		
7	Test	methods	3		

iTeh STANDARD PREVIEW (standards.iteh.ai)

ISO 5832-7:2016 https://standards.iteh.ai/catalog/standards/sist/501868f8-7a61-40ac-83f8-db61e3bdec47/iso-5832-7-2016

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).

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).

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 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.

The committee responsible for this document is ISO/TC 150, *Implants for surgery*, Subcommittee SC 1, *Materials*.

ISO 5832-7:2016

This third edition cancels and replaces the second edition (ISO 583247:1994), which has been technically revised. db61e3bdec47/iso-5832-7-2016

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

Introduction

No known surgical implant material has ever been shown to be completely free of adverse reactions in the human body. However, long-term clinical experience of the use of the material referred to in this document has shown that an acceptable level of biological response can be expected when the material is used in appropriate conditions.

iTeh STANDARD PREVIEW (standards.iteh.ai)

ISO 5832-7:2016 https://standards.iteh.ai/catalog/standards/sist/501868f8-7a61-40ac-83f8-db61e3bdec47/iso-5832-7-2016

iTeh STANDARD PREVIEW (standards.iteh.ai)

ISO 5832-7:2016 https://standards.iteh.ai/catalog/standards/sist/501868f8-7a61-40ac-83f8-db61e3bdec47/iso-5832-7-2016

Implants for surgery — Metallic materials —

Part 7:

Forgeable and cold-formed cobalt-chromium-nickel-molybdenum-iron alloy

1 Scope

This document specifies the characteristics of, and corresponding test methods for, forgeable and cold-formed cobalt-chromium-nickel-molybdenum-iron alloy for use in the manufacture of surgical implants.

NOTE The mechanical properties of a sample obtained from a finished product made of this alloy do not necessarily comply with those specified in this document.

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 643, Steels — Micrographic determination of the apparent grain size

ISO 6892-1, Metallic materials — Tensile testing 150 Part 1: Method of test at room temperature

3 Terms and definitions

No terms and definitions are listed in this document.

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 http://www.iso.org/obp

4 Chemical composition

The heat analysis of the alloy when determined as specified in <u>Clause 7</u> shall comply with the chemical composition specified in <u>Table 1</u>. The analysis of samples taken from products manufactured from the alloy shall also comply with <u>Table 1</u>.

Table 1 — Chemical composition

Element	Element compositional limits,	
Element	% (m/m)	
Cobalt	39 to 42	
Chromium	18,5 to 21,5	
Nickel	14 to 18	
Molybdenum	6,5 to 8,0	
Manganese	1,0 to 2,5	
Silicon	1 max.	
Carbon	0,15 max.	
Phosphorus	0,015 max.	
Sulfur	0,015 max.	
Beryllium	0,001 max.	
Iron	Balance	

5 Microstructure

5.1 Grain size

The microscopic structure shall be uniform. The grain size, determined as specified in <u>Clause 7</u>, shall be no coarser than grain size No. 5.

(standards.iteh.ai)

5.2 Inclusion content

ISO 5832-7:2016

The non-metallic inclusion content of the alloy determined as specified in Glause 7, shall not exceed the limits given in Table 2. db61e3bdec47/iso-5832-7-2016

Table 2 — Inclusion content limits

Type of inclusion	Inclusion content thina		
A – Sulfides	1		
B – Aluminates	3		
C – Silicates	1		
D – Oxides, globular	3		
a There shall be no thick inclusions.			

6 Mechanical properties

The mechanical properties, determined as specified in <u>Clause 7</u>, shall be in accordance with the requirements of <u>Table 3</u>.

Table 3 — Mechanical properties

	Tensile strength	Proof stress of non- proportional elongation	Percentage elongation		
Condition	min.	min.	min.		
	MPa	MPa	%		
Annealed	950	450	65		
Hot worked	950	600	20		
30 % cold-worked	1 450	1 300	8		
Spring tempera	1 650	1 400	1		
^a For specific applications.					

7 Test methods

The test methods to be used in determining compliance with the requirements of this document shall be those given in $\underline{\text{Table 4}}$.

Table 4 — Test methods

Requirement STANDA	Relevant clause or subclause	Test method
Chemical composition (standar	ds.it@ause4i)	Recognized analytical procedures (ISO methods where these exist)
Inclusion content	<u>5.2</u>	ISO 4967
Grain size ISO 58	32-7:2016 5.1	ISO 643
Mechanical properties db61e3bdec47	iso-5832-7-2016	Vac-8318-
Tensile strength	Clause 6	ISO 6892-1
Percentage elongation		
Proof stress of non-proportional elongation		