
**Implants for surgery — Metallic
materials —**

**Part 7:
Forgeable and cold-formed cobalt-
chromium-nickel-molybdenum-iron
alloy**

Implants 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*

ISO 5832-7:2016

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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 (see www.iso.org/directives).

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

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

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.

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