
**Implants for surgery — Metallic
materials —**

**Part 9:
Wrought high nitrogen stainless steel**

Implants chirurgicaux — Matériaux métalliques —

Partie 9: Acier inoxydable corroyé à haute teneur en azote

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Contents

	Page
Foreword	iv
Introduction	v
1 Scope	1
2 Normative references	1
3 Terms and definitions	1
4 Chemical composition	1
4.1 Test samples.....	1
4.2 Cast analysis.....	2
5 Microstructure	2
5.1 Grain size.....	2
5.2 Absence of foreign phases.....	2
5.3 Inclusion content.....	2
6 Corrosion resistance	3
7 Mechanical properties	3
7.1 Test pieces.....	3
7.2 Tensile test.....	3
8 Test methods	4

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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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This document was prepared by Technical Committee ISO/TC 150, *Implants for surgery*, Subcommittee SC 1, *Materials*.

This third edition cancels and replaces the second edition (ISO 5832-9:2007), which has been technically revised.

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.

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 applications.

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