



**SLOVENSKI STANDARD
SIST EN ISO 10993-1:2010/AC:2010**

Biološko ovrednotenje medicinskih pripomočkov - 1. del: Ocena in preskušanje znotraj procesa obvladovanja tveganja - Popravek 1 (ISO 10993-1:2009/Cor 1:2010)

Biological evaluation of medical devices - Part 1: Evaluation and testing within a risk management process - Technical Corrigendum 1 (ISO 10993-1:2009/Cor 1:2010)

Biologische Beurteilung von Medizinprodukten - Teil 1: Beurteilung und Prüfung im Rahmen eines Risikomanagementverfahrens (ISO 10993-1:2009/Cor 1:2010)

Évaluation biologique des dispositifs médicaux - Partie 1: Évaluation et essais au sein d'un processus de gestion du risque - Rectificatif technique 1 (ISO 10993-1:2009/Cor 1:2010)

Ta slovenski standard je istoveten z: EN ISO 10993-1:2009/AC:2010

ICS:

11.100.20 Biološko ovrednotenje medicinskih pripomočkov Biological evaluation of medical devices

SIST EN ISO 10993-1:2010/AC:2010 en

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EUROPEAN STANDARD

EN ISO 10993-1:2009/AC

NORME EUROPÉENNE

June 2010

EUROPÄISCHE NORM

Juin 2010

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ICS 11.100.20

English version
 Version Française
 Deutsche Fassung

Biological evaluation of medical devices - Part 1: Evaluation and testing
 within a risk management process - Technical Corrigendum 1 (ISO 10993-
 1:2009/Cor 1:2010)

Évaluation biologique des dispositifs
 médicaux - Partie 1: Évaluation et essais
 au sein d'un processus de gestion du
 risque - Rectificatif technique 1 (ISO 10993-
 1:2009/Cor 1:2010)

Biologische Beurteilung von
 Medizinprodukten - Teil 1: Beurteilung und
 Prüfung im Rahmen eines
 Risikomanagementverfahrens (ISO 10993-
 1:2009/Cor 1:2010)

This corrigendum becomes effective on 15 June 2010 for incorporation in the three official language
 versions of the EN.

Ce corrigendum prendra effet le 15 juin 2010 pour incorporation dans les trois versions linguistiques
 officielles de la EN.

Die Berichtigung tritt am 15.Juni 2010 zur Einarbeitung in die drei offiziellen Sprachfassungen der EN
 in Kraft.

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

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EN ISO 10993-1:2009/AC:2010 (E)

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Foreword

This document (EN ISO 10993-1:2009/AC:2010) has been prepared by Technical Committee ISO/TC 194 "Biological evaluation of medical devices" in collaboration with Technical Committee CEN/TC 206 "Biological evaluation of medical devices" the secretariat of which is held by NEN.

Endorsement notice

The text of ISO 10993-1:2009/Cor 1:2010 has been approved by CEN as a EN ISO 10993-1:2009/AC:2010 without any modification.

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INTERNATIONAL STANDARD ISO 10993-1:2009
TECHNICAL CORRIGENDUM 1

Published 2010-06-15

INTERNATIONAL ORGANIZATION FOR STANDARDIZATION • МЕЖДУНАРОДНА ОРГАНИЗАЦИЯ ПО СТАНДАРТИЗАЦИИ • ORGANISATION INTERNATIONALE DE NORMALISATION

Biological evaluation of medical devices —

Part 1: Evaluation and testing within a risk management process

TECHNICAL CORRIGENDUM 1

Évaluation biologique des dispositifs médicaux —

Partie 1: Évaluation et essais au sein d'un processus de gestion du risque

RECTIFICATIF TECHNIQUE 1

Technical Corrigendum 1 to ISO 10993-1:2009 was prepared by Technical Committee ISO/TC 194, *Biological evaluation of medical devices*.

*1st STANDARD PREVIEW
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 4674-ba81-5da03951b571ist-en-iso-10993-1-2010-2010*

Page 5, Figure 1

On the top left-hand side of the flowchart, and to the right of the rhombus indicating “Is there either direct or indirect contact?”, replace “1.0” with “Clause 1”.

On the lower left-hand side of the flowchart, and to the top right of the rectangle indicating “Perform further evaluation of device ... and type and duration of contact”, replace “7.0” with “Clause 7”.

On the lower right-hand side of the flowchart, replace the text in the bottom right rectangle with “Perform toxicological risk assessment (Annex B)”.