

ILNAS

Institut luxembourgeois de la normalisation
de l'accréditation, de la sécurité et qualité
des produits et services

ILNAS-EN ISO 10993-13:2010

Biological evaluation of medical devices - Part 13: Identification and quantification of degradation products from polymeric medical devices (ISO)

Biologische Beurteilung von
Medizinprodukten - Teil 13: Qualitativer
und quantitativer Nachweis von
Abbauprodukten in Medizinprodukten

Évaluation biologique des dispositifs
médicaux - Partie 13: Identification et
quantification de produits de
dégradation de dispositifs médicaux à

National Foreword

This European Standard EN ISO 10993-13:2010 was adopted as Luxembourgish Standard ILNAS-EN ISO 10993-13:2010.

Every interested party, which is member of an organization based in Luxembourg, can participate for FREE in the development of Luxembourgish (ILNAS), European (CEN, CENELEC) and International (ISO, IEC) standards:

- Participate in the design of standards
- Foresee future developments
- Participate in technical committee meetings

<https://portail-qualite.public.lu/fr/normes-normalisation/participer-normalisation.html>

THIS PUBLICATION IS COPYRIGHT PROTECTED

Nothing from this publication may be reproduced or utilized in any form or by any mean - electronic, mechanical, photocopying or any other data carries without prior permission!

English Version

Biological evaluation of medical devices - Part 13: Identification and quantification of degradation products from polymeric medical devices (ISO 10993-13:2010)

Évaluation biologique des dispositifs médicaux - Partie 13:
Identification et quantification de produits de dégradation
de dispositifs médicaux à base de polymères (ISO 10993-
13:2010)

Biologische Beurteilung von Medizinprodukten - Teil 13:
Qualitativer und quantitativer Nachweis von
Abbauprodukten in Medizinprodukten aus Polymeren (ISO
10993-13:2010)

This European Standard was approved by CEN on 5 June 2010.

CEN members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for giving this European Standard the status of a national standard without any alteration. Up-to-date lists and bibliographical references concerning such national standards may be obtained on application to the CEN Management Centre or to any CEN member.

This European Standard exists in three official versions (English, French, German). A version in any other language made by translation under the responsibility of a CEN member into its own language and notified to the CEN Management Centre has the same status as the official versions.

CEN members are the national standards bodies of Austria, Belgium, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland and United Kingdom.



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

Management Centre: Avenue Marnix 17, B-1000 Brussels