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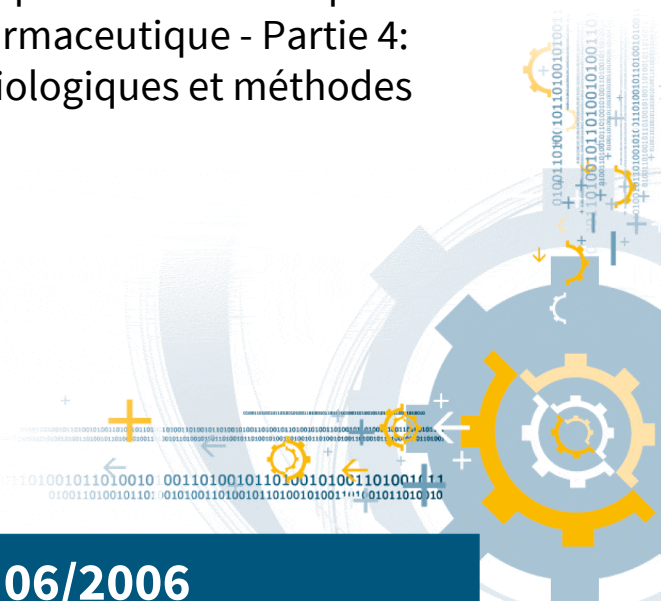
ILNAS-EN ISO 8871-4:2006

Elastomeric parts for parenterals and for devices for pharmaceutical use - Part 4: Biological requirements and test methods (ISO 8871-4:2006)

Elastomere Teile für Parenteralia und für
Geräte zur pharmazeutischen
Verwendung - Teil 4: Biologische
Anforderungen und Prüfverfahren (ISO

Éléments en élastomère pour
administration parentérale et dispositifs
à usage pharmaceutique - Partie 4:
Exigences biologiques et méthodes

06/2006



National Foreword

This European Standard EN ISO 8871-4:2006 was adopted as Luxembourgish Standard ILNAS-EN ISO 8871-4:2006.

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- Participate in the design of standards
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ILNAS-EN ISO 8871-4:2006

EUROPEAN STANDARD **EN ISO 8871-4**

NORME EUROPÉENNE

EUROPÄISCHE NORM

June 2006

ICS 11.040.20

Supersedes EN ISO 8871:1997

English Version

**Elastomeric parts for parenterals and for devices for
pharmaceutical use - Part 4: Biological requirements and test
methods (ISO 8871-4:2006)**

Éléments en élastomère pour administration parentérale et
dispositifs à usage pharmaceutique - Partie 4: Exigences
biologiques et méthodes d'essais (ISO 8871-4:2006)

Elastomere Teile für Parenteralia und für Geräte zur
pharmazeutischen Verwendung - Teil 4: Biologische
Anforderungen und Prüfverfahren (ISO 8871-4:2006)

This European Standard was approved by CEN on 5 June 2006.

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 Central Secretariat 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 Central Secretariat has the same status as the official versions.

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

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Foreword

This document (EN ISO 8871-4:2006) has been prepared by Technical Committee ISO/TC 76 "Transfusion, infusion and injection equipment for medical and pharmaceutical use" in collaboration with CMC.

This European Standard shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by December 2006, and conflicting national standards shall be withdrawn at the latest by December 2006.

This document supersedes EN ISO 8871:1997.

According to the CEN/CENELEC Internal Regulations, the national standards organizations of the following countries are bound to implement this European Standard: Austria, Belgium, 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.

Endorsement notice

The text of ISO 8871-4:2006 has been approved by CEN as EN ISO 8871-4:2006 without any modifications.

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2006-06-15

Elastomeric parts for parenterals and for devices for pharmaceutical use —

Part 4: Biological requirements and test methods

*Éléments en élastomère pour administration parentérale et dispositifs à
usage pharmaceutique —*

Partie 4: Exigences biologiques et méthodes d'essai



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