

### SLOVENSKI STANDARD kSIST FprEN ISO 8871-5:2014

01-april-2014

Deli iz elastomera za parenteralne farmacevtske oblike - 5. del: Funkcionalne zahteve in preskušanje (ISO 8871-5:2005)

Elastomeric parts for parenterals and for devices for pharmaceutical use - Part 5: Functional requirements and testing (ISO 8871-5:2005)

Elastomere Teile für Parenteralia und für Geräte zur pharmazeutischen Verwendung - Teil 5: Funktionelle Anforderungen und Prüfung (ISO 8871-5:2005)

Éléments en élastomère pour administration parentérale et dispositifs à usage pharmaceutique - Partie 5: Exigences fonctionnelles et essais (ISO 8871-5:2005)

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**kSIST FprEN ISO 8871-5:2014** 

## EUROPEAN STANDARD NORME EUROPÉENNE EUROPÄISCHE NORM

### FINAL DRAFT FprEN ISO 8871-5

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#### **English Version**

# Elastomeric parts for parenterals and for devices for pharmaceutical use - Part 5: Functional requirements and testing (ISO 8871-5:2005)

Éléments en élastomère pour administration parentérale et dispositifs à usage pharmaceutique - Partie 5: Exigences fonctionnelles et essais (ISO 8871-5:2005)

Elastomere Teile für Parenteralia und für Geräte zur pharmazeutischen Verwendung - Teil 5: Funktionelle Anforderungen und Prüfung (ISO 8871-5:2005)

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#### **Foreword**

The text of ISO 8871-5:2005 has been prepared by Technical Committee ISO/TC 76 "Transfusion, infusion and injection, and blood processing equipment for medical and pharmaceutical use" of the International Organization for Standardization (ISO) and has been taken over as FprEN ISO 8871-5:2014 by Technical Committee CEN/TC 205 "Non-active medical devices" the secretariat of which is held by DIN.

This document is currently submitted to the Unique Acceptance Procedure.

#### **Endorsement notice**

The text of ISO 8871-5:2005 has been approved by CEN as FprEN ISO 8871-5:2014 without any modification.

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## INTERNATIONAL STANDARD

ISO 8871-5

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## Elastomeric parts for parenterals and for devices for pharmaceutical use —

Part 5:

Functional requirements and testing

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

Partie 5: Exigences fonctionnelles et essais



#### ISO 8871-5:2005(E)

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