# SLOVENSKI STANDARD
# kSIST FprEN ISO 8362-2:2010

**01-april-2010**

---

**Vsebniki za parenteralne farmacevtske oblike in dodatna oprema - 2. del: Zapirala za viale (ISO 8362-2:2008)**

Injection containers and accessories - Part 2: Closures for injection vials (ISO 8362-2:2008)

Injektionsbehältnisse und Zubehör - Teil 2: Stopfen für Injektionsflaschen (ISO 8362-2:2008)

Récipients et accessoires pour produits injectables - Partie 2: Bouchons pour flacons (ISO 8362-2:2008)

**Ta slovenski standard je istoveten z:** **FprEN ISO 8362-2**

---

**ICS:**

| | | |
|---|---|---|
| 11.040.20 | Transfuzijska, infuzijska in injekcijska oprema | Transfusion, infusion and injection equipment |

**kSIST FprEN ISO 8362-2:2010** **en**

EUROPEAN STANDARD

NORME EUROPÉENNE

EUROPÄISCHE NORM

# FINAL DRAFT
# FprEN ISO 8362-2

February 2010

ICS 11.040.20

Will supersede EN 28362-2:1993

English Version

## Injection containers and accessories - Part 2: Closures for injection vials (ISO 8362-2:2008)

Récipients et accessoires pour produits injectables - Partie 2: Bouchons pour flacons (ISO 8362-2:2008)

Injektionsbehältnisse und Zubehör - Teil 2: Stopfen für Injektionsflaschen (ISO 8362-2:2008)

This draft European Standard is submitted to CEN members for unique acceptance procedure. It has been drawn up by the Technical Committee CEN/SS S02.

If this draft becomes a European Standard, 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.

This draft European Standard was established by CEN 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.

**Warning** : This document is not a European Standard. It is distributed for review and comments. It is subject to change without notice and shall not be referred to as a European Standard.

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

**Management Centre: Avenue Marnix 17, B-1000 Brussels**



Ref. No. FprEN ISO 8362-2:2010: E

# Contents

Page

# Foreword

The text of ISO 8362-2:2008 has been prepared by Technical Committee ISO/TC 76 “Transfusion, infusion and injection equipment for medical and pharmaceutical use” of the International Organization for Standardization (ISO) and has been taken over as FprEN ISO 8362-2:2010.

This document is currently submitted to the Unique Acceptance Procedure.

This document will supersede EN 28362-2:1993.

**Endorsement notice**

The text of ISO 8362-2:2008 has been approved by CEN as a FprEN ISO 8362-2:2010 without any modification.

INTERNATIONAL STANDARD

**ISO 8362-2**

Second edition
2008-10-15

# Injection containers and accessories —

## Part 2: Closures for injection vials

*Récipients et accessoires pour produits injectables —*

*Partie 2: Bouchons pour flacons*

Reference number
ISO 8362-2:2008(E)

**PDF disclaimer**

This PDF file may contain embedded typefaces. In accordance with Adobe's licensing policy, this file may be printed or viewed but shall not be edited unless the typefaces which are embedded are licensed to and installed on the computer performing the editing. In downloading this file, parties accept therein the responsibility of not infringing Adobe's licensing policy. The ISO Central Secretariat accepts no liability in this area.

Adobe is a trademark of Adobe Systems Incorporated.

Details of the software products used to create this PDF file can be found in the General Info relative to the file; the PDF-creation parameters were optimized for printing. Every care has been taken to ensure that the file is suitable for use by ISO member bodies. In the unlikely event that a problem relating to it is found, please inform the Central Secretariat at the address given below.



ISO copyright office
Case postale 56 • CH-1211 Geneva 20
Tel. + 41 22 749 01 11
Fax + 41 22 749 09 47
E-mail copyright@iso.org
Web www.iso.org

Published in Switzerland

 