

# **SLOVENSKI STANDARD**

## **SIST EN ISO 8872:2003**

**01-september-2003**

**BUXca Yý U**  
**SIST EN 28872:2000**

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### **Aluminijeve zaporke za transfuzijske, infuzijske in injekcijske steklenice - Splošne zahteve in preskusne metode (ISO 8872:2003)**

Aluminium caps for transfusion, infusion and injection bottles - General requirements and test methods (ISO 8872:2003)

Aluminium-Bördekappen für Transfusions-, Infusions- und Injektionsflaschen - Allgemeine Anforderungen und Prüfverfahren (ISO 8872:2003)

Capsules en aluminium pour flacons de transfusion, perfusion et injection - Spécifications générales et méthodes d'essai (ISO 8872:2003)

**Ta slovenski standard je istoveten z: EN ISO 8872:2003**

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#### **ICS:**

11.040.20	Transfuzijska, infuzijska in injekcijska oprema	Transfusion, infusion and injection equipment
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**SIST EN ISO 8872:2003**

**en**

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EUROPEAN STANDARD  
NORME EUROPÉENNE  
EUROPÄISCHE NORM

**EN ISO 8872**

March 2003

ICS 11.040.20

Supersedes EN 28872:1993

English version

**Aluminium caps for transfusion, infusion and injection bottles -  
General requirements and test methods (ISO 8872:2003)**

Capsules en aluminium pour flacons de transfusion,  
perfusion et injection - Spécifications générales et  
méthodes d'essai (ISO 8872:2003)

Aluminium-Bördelkappen für Transfusions-, Infusions- und  
Injektionsflaschen - Allgemeine Anforderungen und  
Prüfverfahren (ISO 8872:2003)

This European Standard was approved by CEN on 11 March 2003.

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 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 Management Centre has the same status as the official versions.

CEN members are the national standards bodies of Austria, Belgium, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Luxembourg, Malta, Netherlands, Norway, Portugal, Slovakia, Spain, Sweden, Switzerland and United Kingdom.

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

**Management Centre: rue de Stassart, 36 B-1050 Brussels**

## EN ISO 8872:2003 (E)

<b>CORRECTED 2003-05-14</b>
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**Foreword**

This document (EN ISO 8872:2003) 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 September 2003, and conflicting national standards shall be withdrawn at the latest by September 2003.

This document supersedes EN ISO 28872:1993.

According to the CEN/CENELEC Internal Regulations, the national standards organizations of the following countries are bound to implement this European Standard: Austria, Belgium, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Luxembourg, Malta, Netherlands, Norway, Portugal, Slovakia, Spain, Sweden, Switzerland and the United Kingdom.

**Endorsement notice**

The text of ISO 8872:2003 has been approved by CEN as EN ISO 8872:2003 without any modifications.

NOTE Normative references to International Standards are listed in Annex ZA (normative).

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## Annex ZA (normative)

### Normative references to international publications with their relevant European publications

This European Standard incorporates by dated or undated reference, provisions from other publications. These normative references are cited at the appropriate places in the text and the publications are listed hereafter. For dated references, subsequent amendments to or revisions of any of these publications apply to this European Standard only when incorporated in it by amendment or revision. For undated references the latest edition of the publication referred to applies (including amendments).

NOTE Where an International Publication has been modified by common modifications, indicated by (mod.), the relevant EN/HD applies.

<u>Publication</u>	<u>Year</u>	<u>Title</u>	<u>EN</u>	<u>Year</u>
ISO 7500-1	1999	Metallic materials - Verification of static uniaxial testing machines - Part 1: Tension/compression testing machines	EN ISO 7500-1	1999
ISO 8362-3	2001	Injection containers and accessories - Part 3: Aluminium caps for injection vials	EN ISO 8362-3	2003
ISO 8536-3	1999	Infusion equipment for medical use - Part 3: Aluminium caps for infusion bottles	EN ISO 8536-3	1999

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# INTERNATIONAL STANDARD

**ISO  
8872**

Second edition  
2003-03-15

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## Aluminium caps for transfusion, infusion and injection bottles — General requirements and test methods

*Capsules en aluminium pour flacons de transfusion, perfusion et  
injection — Spécifications générales et méthodes d'essai*

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Reference number  
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## Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

International Standards are drafted in accordance with the rules given in the ISO/IEC Directives, Part 2.

The main task of technical committees is to prepare International Standards. Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights.

ISO 8872 was prepared by Technical Committee ISO/TC 76, *Transfusion, infusion and injection equipment for medical and pharmaceutical use*.

This second edition cancels and replaces the first edition (ISO 8872:1988), which has been technically revised.

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