



# SLOVENSKI STANDARD SIST EN ISO 8871:2000

01-januar-2000

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## Deli iz elastomera za parenteralne farmacevtske oblike (ISO 8871:1990)

Elastomeric parts for aqueous parenteral preparations (ISO 8871:1990)

Elastomere Teile für wäßrige parenterale Zubereitungen (ISO 8871:1990)

Éléments en élastomère pour préparations aqueuses parentérales (ISO 8871:1990)

Ta slovenski standard je istoveten z: **EN ISO 8871:1997**

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### **ICS:**

|           |  |  |
|-----------|--|--|
| 11.040.20 | Transfuzijska, infuzijska in<br>injekcijska oprema | Transfusion, infusion and<br>injection equipment |
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EUROPEAN STANDARD

EN ISO 8871

NORME EUROPÉENNE

EUROPÄISCHE NORM

July 1997

ICS 11.040.20

Supersedes EN 28871:1993

Descriptors: see ISO document

English version

**Elastomeric parts for aqueous parenteral  
preparations (ISO 8871:1990)**

Eléments en élastomère pour préparations aqueuses parentérales (ISO 8871:1990)      Elastomere Teile für wässrige parenterale Zubereitungen (ISO 8871:1990)

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This European Standard was approved by CEN on 1997-06-19. 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.

The European Standards exist 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.

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

**CEN**

European Committee for Standardization  
Comité Européen de Normalisation  
Europäisches Komitee für Normung

Central Secretariat: rue de Stassart, 36 B-1050 Brussels

**Foreword**

The text of the International Standard from Technical Committee ISO/TC 76 "Transfusion, infusion and injection equipment for medical use" of the International Organization for Standardization (ISO) has been taken over as an European Standard by Technical Committee CEN/TC 205 "Non-active medical devices", the secretariat of which is held by BSI.

This European Standard replaces EN 28871:1993.

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 January 1998, and conflicting national standards shall be withdrawn at the latest by January 1998.

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, Iceland, Ireland, Italy, Luxembourg, Netherlands, Norway, Portugal, Spain, Sweden, Switzerland and the United Kingdom.

**Endorsement notice**

The text of the International Standard ISO 8871:1990 has been approved by CEN as a European Standard without any modification.

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.

| <u>Publication</u> | <u>Year</u> | <u>Title</u>   | <u>EN</u>   | <u>Year</u> |
|--------------------|-------------|--|-------------|-------------|
| ISO 3696           | 1987        | Water for analytical laboratory use - Specification and test methods | EN ISO 3696 | 1995        |

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# INTERNATIONAL STANDARD

**ISO**  
**8871**

Second edition  
1990-08-01

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## Elastomeric parts for aqueous parenteral preparations

**iTeh** ~~STANDARD PREVIEW~~  
*Éléments en élastomère pour préparations aqueuses parentérales*  
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Reference number  
ISO 8871:1990(E)

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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.

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.

International Standard ISO 8871 was prepared by Technical Committee ISO/TC 76, *Transfusion, infusion and injection equipment for medical use*.

This second edition cancels and replaces the first edition (ISO 8871:1988); presentation has been modified and clauses D.3.2, E.3.2, F.3.2, G.3, J.3.2 and K.3.2 have been technically revised.

Annexes A, B, C, D, E, F, G, H, J, K, L and M form an integral part of this International Standard.