

Elastomeric parts for aqueous parenteral preparations

The European Standard EN ISO 8871:1997 has the status of a
British Standard

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National foreword

This British Standard has been prepared by Technical Committee CH/11 and is the English language version of EN ISO 8871:1997 *Elastomeric parts for aqueous parenteral preparations* published by the European Committee for Standardization (CEN). It is identical with ISO 8871:1990, published by the International Organization for Standardization (ISO).

This British Standard contains an informative annex which is identical with ISO 8871:1990 Amendment 1:1995. At the time of publication of this British Standard, this ISO Amendment had not been adopted by CEN as an amendment to EN ISO 8871:1997 and it is for that reason that the amendment is included in this British Standard as an informative annex.

A list of organizations represented on this committee can be obtained on request to its secretary.

Cross-references

Attention is drawn to the fact that CEN and CENELEC standards normally include an annex which lists normative references to international publications with their corresponding European publications. The British Standards which implement these international or European publications may be found in the BSI Standards Catalogue under the section entitled “International Standards Correspondence Index”, or by using the “Find” facility of the BSI Standards Electronic Catalogue.

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Summary of pages

This document comprises a front cover, an inside front cover, pages i and ii, the EN ISO title page, the EN ISO foreword page, the ISO title page, the ISO pages ii to iv, pages 1 to 12, an inside back cover and a back cover.

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Contents

	Page
National foreword	Inside front cover
Foreword	2
ISO Foreword	iv
Text of EN ISO 8871	1

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)

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.

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CEN

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

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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.

Attention is drawn to annex ZB (informative), concerning A-deviations. Annexes N and P form an integral part of this standard.

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).

Foreword to EN ISO 8817:1997/A1:1999

The text of this amendment EN 8871:1997/A1:1999 to the EN 8871:1997 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 amendment to the European Standard by Technical Committee CEN/TC 205, Non-active medical devices, the Secretariat of which is held by BSI.

This amendment to the European Standard EN ISO 8871:1997 shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by September 1999, and conflicting national standards shall be withdrawn at the latest by September 1999.

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.

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The text of the amendment to the International Standard ISO 8871:1997/amendment 1:1999 has been approved by CEN as an amendment to the European Standard without any modification.

INTERNATIONAL STANDARD

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Elastomeric parts for aqueous parenteral preparations

Éléments en élastomère pour préparations aqueuses parentérales



Contents

	Page
Foreword	iv
Introduction	1
1 Scope	1
2 Normative references	1
3 Classification	1
4 Identification	1
5 Requirements	2
6 Testing	2
7 Packaging	3
8 Storage	3
9 Marking and labelling	3
Annex A (normative) Ultraviolet spectrometry of extracts	4
A.1 Principle	4
A.2 Procedure	4
A.3 Expression of results	4
Annex B (normative) Infra-red spectrometry of pyrolysates	4
B.1 Principle	4
B.2 Procedure	4
B.3 Expression of results	4
Annex C (normative) Determination of extracted reducing matter (oxidizables)	4
C.1 Principle	4
C.2 Reagents	4
C.3 Procedure	5
C.4 Expression of results	5
Annex D (normative) Determination of extracted heavy metals	5
D.1 Principle	5
D.2 Reagents	5
D.3 Procedure	5
D.4 Expression of results	5
Annex E (normative) Determination of extracted ammonia	5
E.1 Principle	5
E.2 Reagent	5
E.3 Procedure	6
E.4 Expression of results	6
Annex F (normative) Determination of extracted halides	6
F.1 Principle	6
F.2 Reagents	6
F.3 Procedure	6
F.4 Expression of results	6
Annex G (normative) Determination of acidity and alkalinity	6
G.1 Principle	6
G.2 Reagents	6
G.3 Procedure	6
G.4 Expression of results	7
Annex H (normative) Determination of extracted non-volatile solids	7
H.1 Principle	7
H.2 Procedure	7
H.3 Expression of results	7
Annex J (normative) Determination of volatile sulfides	7
J.1 Principle	7