

Sterile hypodermic syringes for single use —

Part 2: Syringes for use with power-driven syringe pumps

The European Standard EN ISO 7886-2:1997 has the status of a
British Standard

ICS 11.040.20

Committees responsible for this British Standard

The preparation of this British Standard was entrusted to Technical Committee CH/2, Hypodermic equipment, upon which the following bodies were represented:

- Association of British Healthcare Industries
- British Diabetic Association
- British Glass Manufacturers' Confederation
- British Surgical Trades' Association
- Department of Health
- Disposable Hypodermic and Allied Equipment Manufacturers' Association (UK)
- Medical Sterile Products Association
- Royal Pharmaceutical Society of Great Britain

This British Standard, having been prepared under the direction of the Health and Environment Sector Board, was published under the authority of the Standards Board and comes into effect on 15 September 1996

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The following BSI references relate to the work on this standard:
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National foreword

This British Standard is the English language version of EN ISO 7886-2:1997. |

This British Standard is published under the direction of the Health and Environment Sector Board whose Technical Committee CH/2 has the responsibility to:

- aid enquirers to understand the text;
- present to the responsible international committee any enquiries on interpretation, or proposals for change, and keep UK interests informed;
- monitor related international and European developments and promulgate them in the UK.

NOTE International and European Standard, as well as overseas standards, are available from Customer Services, BSI, 389 Chiswick High Road, London W4 4AL.

A British Standard does not purport to include all the necessary provisions of a contract. Users of British Standards are responsible for their correct application.

Compliance with a British Standard does not of itself confer immunity from legal obligations.

Summary of pages

This document comprises a front cover, an inside front cover, pages i and ii, the EN ISO title page, page 2, the ISO title page, pages ii to iv, pages 1 to 12, an inside back cover and a back cover.

This standard has been updated (see copyright date) and may have had amendments incorporated. This will be indicated in the amendment table on the inside front cover.

English version

Sterile hypodermic syringes for single use — Part 2: Syringes for use with power-driven syringe pumps

(ISO 7886-2:1996)

Seringues hypodermiques stériles, non
réutilisables —
Partie 2: Seringues pour pousse-seringues mûs
par un moteur
(ISO 7886-2:1996)

Sterile Einmalspritzen für medizinische
Zwecke —
Teil 2: Spritzen zur Verwendung mit
Spritzenpumpen
(ISO 7886-2:1996)

This European Standard was approved by CEN on 1997-08-23. 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.

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 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 84, Medical devices for injections, of the International Organization for Standardization (ISO) has been taken over as a European Standard by Technical Committee CEN/TC 205 Non-active medical devices, the secretariat of which is held by BSI.

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

According to 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 7886-2:1996 has been approved by CEN as a European Standard without any modification.

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

INTERNATIONAL STANDARD

**ISO
7886-2**

First edition
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BS EN ISO

Sterile hypodermic syringes for single use —

Part 2:

Syringes for use with power-driven syringe
pumps

Seringues hypodermiques stériles, non réutilisables —

Partie 2: Seringues pour pousse-seringues mûs par un moteur



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