

Stainless steel needle tubing for the manufacture of medical devices

The European Standard EN ISO 9626:1995, with the incorporation of amendment A1:2001, has the status of a British Standard

Committees responsible for this British Standard

The preparation of this British Standard was entrusted by the Health Care Standards Policy Committee (HCC/-) to Technical Committee HCC/2, upon which the following bodies were represented:

- Association of British Health-care Industries
- Association of Clinical Pathologists
- 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 Care Standards Policy Committee, was published under the authority of the Standards Board and comes into effect on 20 December 1991

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The following BSI references relate to the work on this standard:
Committee reference HCC/2
Draft for comment 88/55196 DC

ISBN 0 580 20414 6

Amendments issued since publication

Amd. No.	Date	Comments
8750	July 1995	
13264	21 December 2001	Indicated by a sideline
13672 Incorp. Corr. No. 1	31 January 2002	Incorporating annex ZA

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National foreword

This British Standard is the official English language version of EN ISO 9626:1995, including amendment A1:2001. It is identical with ISO 9626:1991, including amendment 1:2001.

Cross-references

The British Standards which implement international or European publications referred to in this document 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, pages 2 to 11, the annex ZA page, an inside back cover and a back cover.

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ICS 11.040.20

Descriptors: Medical equipment, hypodermic needles, stainless steels, steel tubes, specifications, dimensions, tests

English version

Stainless steel needle tubing for the manufacture of medical devices

(includes amendment A1:2001)

(ISO 9626:1991 + amd. 1:2000)

Tubes d'aiguilles en acier inoxydable pour la
fabrication de matériel médical
(inclut l'amendement A1:2001)
(ISO 9626:1991 + amd. 1:2001)

Kanülenrohre aus nichtrostendem Stahl zur
Herstellung von Medizinprodukten
(enthält Änderung A1:2001)
(ISO 9626:1991 + amd. 1:2001)

This European Standard was approved by CEN on 1994-11-14. Amendment A1:2001 was approved by CEN on 2001-06-01. 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.

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CEN

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

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

Foreword

This European Standard has been taken over by Technical Committee CEN/TC 205, Non-active medical devices, from the work of ISO/TC 84, Medical devices for injections, of the International Organization for Standardization (ISO).

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 August 1995, conflicting national standards shall be withdrawn at the latest by August 1995.

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

CORRECTED 2001-12-12

Foreword to amendment A1

The text of this Amendment EN ISO 9626:1995/A1:2001 to the EN ISO 9626:1995 has been prepared by Technical Committee ISO/TC 84, Medical devices for injections, in collaboration with 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 December 2001, and conflicting national standards shall be withdrawn at the latest by December 2001.

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 United Kingdom.

Endorsement notice

The text of the Amendment to the International Standard ISO 9626:1991/AM 1:2001 has been approved by CEN as an Amendment to the European Standard without any modification.

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