SLOVENSKI STANDARD
SIST EN 27740:2000/A1:2000

01-januar-2000

Kirurški instrumenti, skalpeli s snemnimi rezili, mere nastavkov (ISO 7740:1985)

Instruments for surgery, scalpels with detachable blades, fitting dimensions (ISO 7740:1985)

Chirurgische Instrumente, Skalpelle mit auswechselbaren Klingen, Paßmaße (ISO 7740:1985)

Instruments chirurgicaux, bistouris a lames détachables, dimensions d'assemblage (ISO 7740:1985)

Ta slovenski standard je istoveten z: **EN 27740:1992/A1:1997**

ICS:

11.040.30	Operacijski instrumenti in materiali	Surgical instruments and materials

SIST EN 27740:2000/A1:2000 **en**

EUROPEAN STANDARD

NORME EUROPÉENNE

EUROPÄISCHE NORM

EN 27740:1992/A1

November 1997

ICS 11.040.30

Descriptors: medical equipment, surgical equipment, scalpels, handles, blades, joining, dimensions

English version

Instruments for surgery, scalpels with detachable blades, fitting dimensions (ISO 7740:1985)

Instruments chirurgicaux, bistouris à lames détachables, dimensions d'assemblage (ISO 7740:1985)

Chirurgische Instrumente, Skalpelle mit auswechselbaren Klingen, Paßmaße (ISO 7740:1985)

This amendment A1 modifies the European Standard EN 27740:1992; it was approved by CEN on 30 October 1997.

CEN members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for inclusion of this amendment into the relevant 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 amendment 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.

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

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

Foreword

This Amendment EN 27740:1992/A1:1997 to EN 27740:1992 has been prepared 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 27740:1992 shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by May 1998, and conflicting national standards shall be withdrawn at the latest by May 1998.

This Amendment consists of annex ZA to EN 27740:1992.

For relationship with EU Directive(s), see informative Annex ZA, which is 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.

Annex ZA (informative)

Clauses of this European Standard addressing essential requirements or other provisions of EU Directives

This European Standard has been prepared under a mandate given to CEN by the European Commission and the European Free Trade Association and supports essential requirements of EU Directive 93/42/EEC.

> WARNING: Other requirements and other EU Directives may be applicable to the product(s) falling within the scope of this standard.

The following clauses of this standard are likely to support requirements of Directive 93/42/EEC.

Compliance with these clauses of this standard provides one means of conforming with the specific essential requirements of the Directive concerned and associated EFTA regulations.

TABLE ZA.1 : Correspondence between this European Standard and EU Directives

Clause/subclause of this European Standard	Corresponding Essential Requirement of Directive 93/42/EEC	Comments
2	2, 3, 9.1	