



Institut luxembourgeois de la normalisation
de l'accréditation, de la sécurité et qualité
des produits et services

ILNAS-EN ISO 5356-1:2004

Anaesthetic and respiratory equipment - Conical connectors - Part 1: Cones and sockets (ISO 5356-1:2004)

Matériel d'anesthésie et de réanimation
respiratoire - Raccords coniques - Partie
1: Raccords mâles et femelles (ISO
5356-1:2004)

Anästhesie- und Beatmungsgeräte -
Konische Konnektoren - Teil 1: Männliche
und weibliche Konen (ISO 5356-1:2004)

05/2004



National Foreword

This European Standard EN ISO 5356-1:2004 was adopted as Luxembourgish Standard ILNAS-EN ISO 5356-1:2004.

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ILNAS-EN ISO 5356-1:2004

EUROPEAN STANDARD **EN ISO 5356-1**

NORME EUROPÉENNE

EUROPÄISCHE NORM

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Supersedes EN 1281-1:1997

English version

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Matériel d'anesthésie et de réanimation respiratoire -
Raccords coniques - Partie 1: Raccords mâles et femelles
(ISO 5356-1:2004)

Anästhesie- und Beatmungsgeräte - Konische Konnektoren
- Teil 1: Männliche und weibliche Konen (ISO 5356-1:2004)

This European Standard was approved by CEN on 1 April 2004.

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, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Slovakia, Slovenia, Spain, Sweden, Switzerland and United Kingdom.



EUROPEAN COMMITTEE FOR STANDARDIZATION
COMITÉ EUROPÉEN DE NORMALISATION
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Foreword

This document (EN ISO 5356-1:2004) has been prepared by Technical Committee ISO/TC 121 "Anaesthetic and respiratory equipment" in collaboration with Technical Committee CEN/TC 215 "Respiratory and anaesthetic equipment", 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 November 2004, and conflicting national standards shall be withdrawn at the latest by November 2004.

This document supersedes EN 1281-1:1997.

This document 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(s).

For relationship with EU Directive(s), see informative Annex ZA, which is an integral part of this document.

According to the CEN/CENELEC Internal Regulations, the national standards organizations of the following countries are bound to implement this European Standard: Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Slovakia, Slovenia, Spain, Sweden, Switzerland and United Kingdom.

Endorsement notice

The text of ISO 5356-1:2004 has been approved by CEN as EN ISO 5356-1:2004 without any modifications.

ANNEX ZA

(informative)

Relationship between this European Standard and the Essential Requirements of EU Directive 93/42 EEC

This European Standard has been prepared under a mandate given to CEN by the European Commission and the European Free Trade Association to provide a means of conforming to Essential Requirements of the New Approach Directive 93/42 EEC.

Once this standard is cited in the Official Journal of the European Communities under that Directive and has been implemented as a national standard in at least one Member State, compliance with the normative clauses of this standard given in Table ZA.1 confers, within the limits of the scope of this standard, a presumption of conformity with the corresponding Essential Requirements of that Directive and associated EFTA regulations.

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

Table ZA.1 – Correspondence between this European Standard and EU Directives

Clause/subclause of this European Standard	Corresponding essential requirements of Directive 93/42/EEC	Comments
4	1, 2, 7.5, 7.6, 9.1	
5	1, 2, 7.5, 7.6, 9.1	
6	1, 2, 4, 7.5, 7.6, 9.1	

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Conical connectors —**

**Part 1:
Cones and sockets**

*Matériel d'anesthésie et de réanimation respiratoire — Raccords
coniques —*

Partie 1: Raccords mâles et femelles



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