ILNAS

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

ILNAS-EN ISO 7864:2016

Sterile hypodermic needles for single use - Requirements and test methods (ISO 7864:2016)

Aiguilles hypodermiques stériles, non réutilisables - Exigences et méthodes d'essai (ISO 7864:2016)

Sterile Injektionskanülen für den Einmalgebrauch - Anforderungen und Prüfverfahren (ISO 7864:2016)

08/2016

National Foreword

This European Standard EN ISO 7864:2016 was adopted as Luxembourgish Standard ILNAS-EN ISO 7864:2016.

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- Participate in the design of standards
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EUROPEAN STANDARD
NORME EUROPÉENNE
EUROPÄISCHE NORM

EN ISO 7864

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English Version

Sterile hypodermic needles for single use - Requirements and test methods (ISO 7864:2016)

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This European Standard was approved by CEN on 15 July 2016.

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 CEN-CENELEC Management Centre 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 CEN-CENELEC Management Centre has the same status as the official versions.

CEN members are the national standards bodies of Austria, Belgium, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, Former Yugoslav Republic of Macedonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey and United Kingdom.

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

CEN-CENELEC Management Centre: Avenue Marnix 17, B-1000 Brussels

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European foreword

This document (EN ISO 7864:2016) has been prepared by Technical Committee ISO/TC 76 "Transfusion, infusion and injection, and blood processing equipment for medical and pharmaceutical use" in collaboration with Technical Committee CEN/TC 205 “Non-active medical devices” the secretariat of which is held by DIN.

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 February 2017, and conflicting national standards shall be withdrawn at the latest by February 2017.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. CEN [and/or CENELEC] shall not be held responsible for identifying any or all such patent rights.

This document supersedes EN ISO 7864:1995.

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.

The following referenced documents are indispensable for the application of this document. For undated references, the latest edition of the referenced document (including any amendments) applies. For dated references, only the edition cited applies. However, for any use of this standard ‘within the meaning of Annex ZA’, the user should always check that any referenced document has not been superseded and that its relevant contents can still be considered the generally acknowledged state-of-art.

When an IEC or ISO standard is referred to in the ISO standard text, this shall be understood as a normative reference to the corresponding EN standard, if available, and otherwise to the dated ISO or IEC standard, as listed in Table 1.

NOTE The way in which these references documents are cited in normative requirements determines the extent (in whole or in part) to which they apply.

Table 1— Correlations between undated normative references and dated EN and ISO standards

Normative references as listed in Clause 2 of the ISO standard	Equivalent dated standard	
	EN	ISO or IEC
ISO 594-1	EN 20594-1: 1993/AC: 1996/ A1: 1997	ISO 594-1:1986
ISO 594-2	EN 1707:1996	ISO 594-2:1998
ISO 3696	EN ISO 3696:1995	ISO 3696:1987
ISO 6009	EN ISO 6009:1994/AC:2008	ISO 6009:2016
ISO 8601	--	ISO 8601: 2004
ISO 9626	EN ISO 9626:1995/A1:2001	ISO 9626:2016
ISO 10993-1	EN ISO 10993-1:2009	ISO 10993-1:2009
ISO 14971	EN ISO 14971:2012	ISO 14971:2007
ISO 23908	EN ISO 23908::2013	ISO 23908:2011
ISO 80369-1	EN ISO 80369-1:2010	ISO 80369-1:2010
ISO 15223-1	EN ISO 15223-1:2012	ISO 15223-1:2012

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

Endorsement notice

The text of ISO 7864:2016 has been approved by CEN as EN ISO 7864:2016 without any modification.

Annex ZA
(informative)

Relationship between this European Standard and the Essential Requirements of EU Directive 93/42/EEC [OJ L 169] aimed to be covered

This European Standard has been prepared under a Commission's standardization request M/295 concerning the development of European Standards related to medical devices to provide one voluntary means of conforming to essential requirements of Council Directive 93/42/EEC of 14 June 1993 concerning medical devices [OJ L 160].

Once this standard is cited in the Official Journal of the European Union under that Directive, 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.

NOTE 1 Where a reference from a clause of this standard to the risk management process is made, the risk management process needs to be in compliance with Directive 93/42/EEC as amended by 2007/47/EC. This means that risks have to be reduced 'as far as possible', 'to a minimum', 'to the lowest possible level', 'minimized' or 'removed', according to the wording of the corresponding essential requirement.

NOTE 2 The manufacturer's policy for determining acceptable risk must be in compliance with Essential Requirements 1, 2, 5, 6, 7, 8, 9, 11 and 12 of the Directive.

NOTE 3 This Annex ZA is based on normative references according to the table of references in the European foreword, replacing the references in the core text.

NOTE 4 When an Essential Requirement does not appear in Table ZA.1, it means that it is not addressed by this European Standard.

Table ZA.1 — Correspondence between this European Standard and Annex I of Directive 93/42/EEC [OJ L 169]

Essential Requirements of Directive 93/42/EEC	Clause(s)/sub-clause(s) of this EN	Remarks/Notes
7.1	4.9.1, 4.9.4, 4.14	
7.3	4.4, 4.5, 4.9.1, 4.9.4, 4.14	
7.5	4.4, 4.5, 4.9.1, 4.9.4, 4.14	The part of ER 7.5 related to phthalates is not explicitly covered.
7.6	4.8.1	
8.1	4.14, 5	The part of ER 8.1 relating to easy handling is not addressed.
8.3	5	
8.4	4.14.1, 5.1	
9.1	4.8.1	
9.2	4.6, 4. 9.3, 4.9.4, 4.10	

13.1	6.1	
13.2	4.7, 4.8.2, 4.9, 6.2, 6.3	
13.3(a)	6.2(e), 6.3(g), 6.4(e)	
13.3(b)	6.2(a), 6.3(a), 6.4(a)	
13.3 (c)	6.2(b), 6.3(b), 6.4(c)	
13.3(d)	6.2(c), 6.3(e),6.4(b)	
13.3(e)	6.2 (f), 6.3(f), 6.4(d)	
13.3 (f)	6.2(d), 6.3(c)	
13.3(i)	6.3(h), 6.4(f)	
13.3(k)	6.1, 6.3(d)	

WARNING 1 — Presumption of conformity stays valid only as long as a reference to this European Standard is maintained in the list published in the Official Journal of the European Union. Users of this standard should consult frequently the latest list published in the Official Journal of the European Union.

WARNING 2 — Other Union legislation may be applicable to the products falling within the scope of this standard.