BS EN ISO 18113-3:2011

BSI Standards Publication

In vitro diagnostic medical devices — Information supplied by the manufacturer (labelling)

Part 3: In vitro diagnostic instruments for professional use (ISO 18113-3:2009)

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This British Standard is the UK implementation of EN ISO 18113-3:2011. It is identical to ISO 18113-3:2009. It supersedes BS EN ISO 18113-3:2009 which is withdrawn.

The UK participation in its preparation was entrusted to Technical Committee CH/212, IVDs.

A list of organizations represented on this committee can be obtained on request to its secretary.

ISBN 978 0 580 76942 9

ICS 11.100.10

This British Standard was published under the authority of the Standards Policy and Strategy Committee on 30 November 2011.

Amendments issued since publication

Date	Text affected

EUROPÄISCHE NORM

October 2011

ICS 11.100.10

Supersedes EN ISO 18113-3:2009

English Version

In vitro diagnostic medical devices - Information supplied by the manufacturer (labelling) - Part 3: In vitro diagnostic instruments for professional use (ISO 18113-3:2009)

Dispositifs médicaux de diagnostic in vitro - Informations fournies par le fabricant (étiquetage) - Partie 3: Instruments de diagnostic in vitro à usage professionnel (ISO 18113-3:2009)

In-vitro-Diagnostika - Bereitstellung von Informationen durch den Hersteller - Teil 3: Geräte für in-vitro-diagnostische Untersuchungen zum Gebrauch durch Fachpersonal (ISO 18113-3:2009)

This European Standard was approved by CEN on 20 September 2011.

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

Management Centre: Avenue Marnix 17, B-1000 Brussels

Ref. No. EN ISO 18113-3:2011: E

Foreword

This document (EN ISO 18113-3:2011) has been prepared by Technical Committee ISO/TC 212 "Clinical laboratory testing and in vitro diagnostic test systems" in collaboration with Technical Committee CEN/TC 140 “In vitro diagnostic 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of an identical text or by endorsement, at the latest by April 2012, and conflicting national standards shall be withdrawn at the latest by October 2014.

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 18113-3:2009.

This new edition contains a revised Annex ZA.

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.

For relationship with EU Directive, 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, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland and United Kingdom.

Endorsement notice

The text of ISO 18113-3:2009 has been approved by CEN as EN ISO 18113-3:2011 without any modification.

Annex ZA
(informative)

Relationship between this European Standard and the Essential Requirements of the EU Directive 98/79/EC on “in vitro Diagnostic Medical Devices”

This European Standard has been prepared under a mandate given to CEN by the European Commission to provide a means of conforming to the Essential Requirements of the New Approach Directive 98/79/EC on “*in vitro* Diagnostic Medical Devices”.

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

Table ZA.1 — Correspondence between this European Standard and European Directive 98/79/EC

Clauses of this European Standard	Essential Requirements (ERs) of Directive 98/79/EC	Qualifying comments/Notes
5, 6, 7	B.8.1	Presumption of conformity with ER B.8.1 also requires compliance with clauses 4.1, 4.2.1 and 4.6 of EN ISO 18113-1.
7.1	B.8.4(a)	**NOTE 1**
5.2.1	B.8.4(b)	
5.2.2	B.8.4(d)	Full compliance with ER B.8.4(d) requires the use of EN 980, clause 5.4, symbol [LOT] or EN 980, clause 5.5, symbol [SN], as applicable.
5.2.3	B.8.4(g)	
7.3	B. 8.5	
7.1, 7.2.1, 7.3, 7.4, 7.5, 7.12	B.8.7(a)	Presumption of conformity with ER B.8.7(a) requires compliance also with clause 4.5 of EN ISO 18113-1. **NOTE 1** **NOTE 2**
7.9.	B.8.7(d)	
7.2.2, 7.11, 7.12	B.8.7(e)	
7.11, 7.12	B.8.7(f)	
7.12, 7.15, 7.17	B.8.7(g)	
7.7, 7.8, 7.9, 7.10, 7.11, 7.12	B.8.7(h)	
7.14	B.8.7(i)	
7.20	B.8.7(j)	
7.13	B.8.7(k)	
7.6, 7.11, 7.12	B.8.7(m)	
7.6, 7.11, 7.12, 7.13, 7.18, 7.19	B.8.7(n)	
7.11	B.8.7(o)	

Clauses of this European Standard	Essential Requirements (ERs) of Directive 98/79/EC	Qualifying comments/Notes
7.19	B.8.7(p)	
7.19	B.8.7(q)	This clause covers only partially ER B.8.7(q), namely only the information about cleaning, decontamination or disinfection. Any other information related to reuse and restrictions on the number of reuse does not apply to this kind of IVD medical device.
7.5, 7.6.3, 7.11	B.8.7(r)	**NOTE 2**
7.5, 7.18	B.8.7(s)	**NOTE 2**

NOTE 1 In the European Union, the name and address of the manufacturer's "EC Authorized representative" is required on the outer container label or in the instructions for use, if the legal manufacturer is not located within the European Union.

NOTE 2 Essential Requirement B.8.7 of Directive 98/79/EC should be consulted for a comprehensive list of the information required.

Contents

Page

Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

International Standards are drafted in accordance with the rules given in the ISO/IEC Directives, Part 2.

The main task of technical committees is to prepare International Standards. Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

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

ISO 18113-3 was prepared by Technical Committee ISO/TC 212, *Clinical laboratory testing and* in vitro *diagnostic test systems*.

ISO 18113 consists of the following parts, under the general title In vitro *diagnostic medical devices — Information supplied by the manufacturer (labelling)*:

— *Part 1: Terms, definitions and general requirements*

— *Part 2:* In vitro *diagnostic reagents for professional use*

— *Part 3*: In vitro *diagnostic instruments for professional use*

— *Part 4:* In vitro *diagnostic reagents for self-testing*

— *Part 5:* In vitro *diagnostic instruments for self-testing*

Introduction

Manufacturers of *in vitro* diagnostic (IVD) instruments for professional use supply users with information to enable the safe use and expected performance of their devices. The type and level of detail varies according to the intended uses and country-specific regulations.

The Global Harmonization Task Force (GHTF) encourages convergence of the evolution of regulatory systems for medical devices at the global level. Eliminating differences among regulatory jurisdictions could allow patients earlier access to new technologies and treatments. See Reference [5]. This part of ISO 18113 provides a basis for harmonization of labelling requirements for IVD instruments for professional use.

This part of ISO 18113 is concerned solely with information supplied with IVD instruments and equipment intended for professional use. It is intended to be used in conjunction with ISO 18113-1, which contains the general requirements for information supplied by the manufacturer and definitions of general labelling concepts.

This part of ISO 18113 is based on EN 591[3]. The text has been modified to conform to Part 2 of the ISO/IEC Directives[2], but the requirements including those in ISO 18113-1, are substantially equivalent to the original European harmonized standard. This part of ISO 18113 is intended to support the essential labelling requirements of all the GHTF partners, as well as other countries that have or plan to enact labelling regulations for IVD medical devices.

For IVD instruments for professional use that are intended to be used as a system with reagents provided by the same manufacturer, this part of ISO 18113 is also intended to be used together with ISO 18113-1 and ISO 18113-2[1].

In vitro diagnostic medical devices — Information supplied by the manufacturer (labelling) —

Part 3: *In vitro* diagnostic instruments for professional use

1 Scope

This part of ISO 18113 specifies requirements for information supplied by the manufacturer of IVD instruments for professional use.

This part of ISO 18113 also applies to apparatus and equipment intended to be used with IVD instruments for professional use.

This part of ISO 18113 can also be applied to accessories.

This part of ISO 18113 does not apply to:

a) instructions for instrument servicing or repair,

b) IVD reagents, including calibrators and control materials for use in control of the reagent,

c) IVD instruments for self-testing.

2 Normative references

The following referenced documents are indispensable for the application of this document. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

ISO 14971, *Medical devices — Application of risk management to medical devices*

ISO 15223-1, *Medical devices — Symbols to be used with medical device labels, labelling and information to be supplied — Part 1: General requirements*

ISO 18113-1, In vitro *diagnostic medical devices — Information supplied by the manufacturer (labelling) — Part 1: Terms, definitions and general requirements*

IEC 61010-1, *Safety requirements for electrical equipment for measurement, control and laboratory use — Part 1: General requirements*

IEC 61010-2-101, *Safety requirements for electrical equipment for measurement, control and laboratory use — Part 2-101: Particular requirements for* in vitro *diagnostic (IVD) medical equipment*

IEC 61326-2-6, *Electrical equipment for measurement, control and laboratory use — EMC requirements — Part 2-6: Particular requirements —* In vitro *diagnostic (IVD) medical equipment*

IEC 62366, *Medical devices — Application of usability engineering to medical devices*

EN 980, *Symbols for use in the labelling of medical devices*