



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

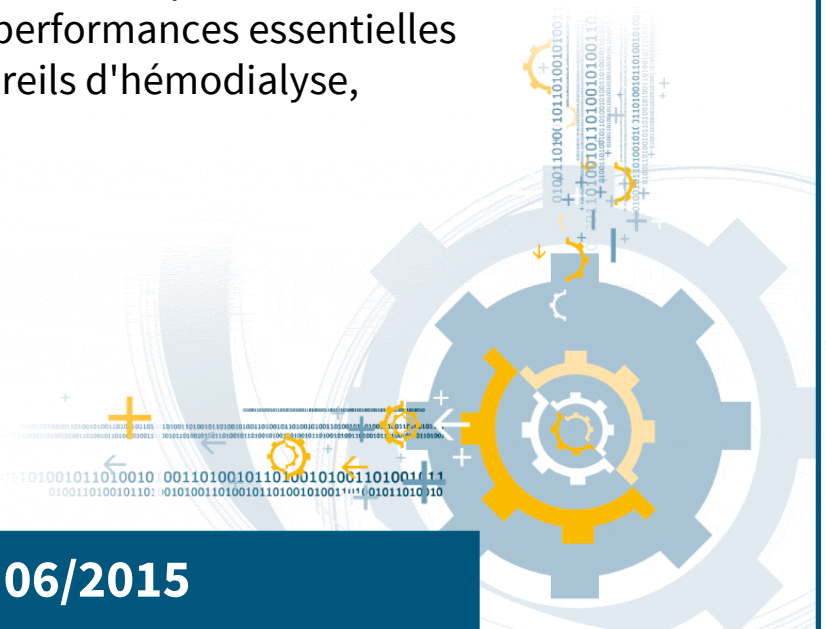
ILNAS-EN 60601-2-16:2015

**Medical electrical equipment - Part
2-16: Particular requirements for the
basic safety and essential performance
of haemodialysis, haemodiafiltration**

Medizinische elektrische Geräte - Teil
2-16: Besondere Festlegungen für die
Sicherheit einschließlich der
wesentlichen Leistungsmerkmale von

Appareils électromédicaux - Partie 2-16:
Exigences particulières pour la sécurité
de base et les performances essentielles
des appareils d'hémodialyse,

06/2015



National Foreword

This European Standard EN 60601-2-16:2015 was adopted as Luxembourgish Standard ILNAS-EN 60601-2-16:2015.

Every interested party, which is member of an organization based in Luxembourg, can participate for FREE in the development of Luxembourgish (ILNAS), European (CEN, CENELEC) and International (ISO, IEC) standards:

- Participate in the design of standards
- Foresee future developments
- Participate in technical committee meetings

<https://portail-qualite.public.lu/fr/normes-normalisation/participer-normalisation.html>

THIS PUBLICATION IS COPYRIGHT PROTECTED

Nothing from this publication may be reproduced or utilized in any form or by any mean - electronic, mechanical, photocopying or any other data carries without prior permission!

ILNAS-EN 60601-2-16:2015

EUROPEAN STANDARD **EN 60601-2-16**

NORME EUROPÉENNE

EUROPÄISCHE NORM

June 2015

ICS 11.040.20; 11.040.25

Supersedes EN 60601-2-16:1998

English Version

**Medical electrical equipment - Part 2-16: Particular requirements
for the basic safety and essential performance of haemodialysis,
haemodiafiltration and haemofiltration equipment
(IEC 60601-2-16:2012)**

Appareils électromédicaux - Partie 2-16: Exigences
particulières pour la sécurité de base et les performances
essentielles des appareils d'hémodialyse,
d'hémodiafiltration et d'hémofiltration
(IEC 60601-2-16:2012)

Medizinische elektrische Geräte - Teil 2-16: Besondere
Festlegungen für die Sicherheit einschließlich der
wesentlichen Leistungsmerkmale von Hämodialyse-,
Hämodiafiltrations- und Hämofiltrationsgeräten
(IEC 60601-2-16:2012)

This European Standard was approved by CENELEC on 2015-04-14. CENELEC 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 CENELEC 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 CENELEC member into its own language and notified to the CEN-CENELEC Management Centre has the same status as the official versions.

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



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

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

Foreword

The text of document 62D/972/FDIS, future edition 4 of IEC 60601-2-16, prepared by SC 62D, "Electromedical equipment", of IEC TC 62, "Electrical equipment in medical practice" was submitted to the IEC-CENELEC parallel vote and approved by CENELEC as EN 60601-2-16:2015.

The following dates are fixed:

- latest date by which the document has to be implemented at national level by publication of an identical national standard or by endorsement (dop) 2016-01-14
- latest date by which the national standards conflicting with the document have to be withdrawn (dow) 2018-04-14

This document supersedes EN 60601-2-16:1998.

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

This document has been prepared under a mandate given to CENELEC by the European Commission and the European Free Trade Association, and supports essential requirements of EU Directive(s).

For the relationship with EU Directive see informative Annex ZZ, which is an integral part of this document.

Endorsement notice

The text of the International Standard IEC 60601-2-16:2012 was approved by CENELEC as a European Standard without any modification.

In the official version, for Bibliography, the following notes have to be added for the standards indicated:

IEC 60601-2-16:1998	NOTE Harmonized as EN 60601-2-16:1998 (not modified).
IEC 60601-2-39	NOTE Harmonized as EN 60601-2-39.
ISO 11197	NOTE Harmonized as EN ISO 11197.

Annex ZA (normative)

Normative references to international publications with their corresponding European publications

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

NOTE 1 When an International Publication has been modified by common modifications, indicated by (mod), the relevant EN/HD applies.

NOTE 2 Up-to-date information on the latest versions of the European Standards listed in this annex is available here: www.cenelec.eu.

Annex ZA of EN 60601-1:2006 applies with the following exceptions:

<u>Publication</u>	<u>Year</u>	<u>Title</u>	<u>EN/HD</u>	<u>Year</u>
<i>Replacement :</i>				
IEC 60601-1-2	2007	Medical electrical equipment - Part 1-2: General requirements for basic safety and essential performance - Collateral standard: Electromagnetic compatibility - Requirements and tests	EN 60601-1-2 + corr March	2007 2010
IEC 60601-1-6	2010	Medical electrical equipment - Part 1-6: General requirements for basic safety and essential performance - Collateral standard: Usability	EN 60601-1-6	2010
IEC 60601-1-8	2006	Medical electrical equipment - Part 1-8: General requirements for basic safety and essential performance - Collateral Standard: General requirements, tests and guidance for alarm systems in medical electrical equipment and medical electrical systems	EN 60601-1-8 + corr March	2007 2010
<i>Addition:</i>				
IEC 60601-1-10	2007	Medical electrical equipment - Part 1-10: General requirements for basic safety and essential performance - Collateral Standard: Requirements for the development of physiologic closed-loop controllers	EN 60601-1-10	2008
IEC 60601-1-11	2010	Medical electrical equipment - Part 1-11: General requirements for basic safety and essential performance - Collateral standard: Requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment	EN 60601-1-11	2010
IEC 62366	2007	Medical devices - Application of usability engineering to medical devices	EN 62366	2008

ISO 594-2	-	Conical fittings with 6 % (Luer) taper for syringes, needles and certain other medical equipment - Part 2: Lock fittings		
ISO 3744	-	Acoustics - Determination of sound power levels and sound energy levels of noise sources using sound pressure - Engineering methods for an essentially free field over a reflecting plane	EN ISO 3744	-
ISO 8638	-	Cardiovascular implants and extracorporeal systems - Extracorporeal blood circuit for haemodialysers, haemodiafilters and haemofilters	EN ISO 8638	-

Annex ZZ (informative)

Coverage of Essential Requirements of EU Directives

This European Standard has been prepared under a mandate given to CENELEC by the European Commission and the European Free Trade Association, and within its scope the Standard covers all relevant essential requirements given in Annex I of EU Directive 93/42/EEC of 14 June 1993 concerning medical devices.

Compliance with this standard provides one means of conformity with the specified essential requirements of the Directive concerned.

WARNING: Other requirements and other EU Directives can be applied to the products falling within the scope of this standard.



INTERNATIONAL STANDARD

NORME INTERNATIONALE

Medical electrical equipment –

Part 2-16: Particular requirements for the basic safety and essential performance of haemodialysis, haemodiafiltration and haemofiltration equipment

Appareils électromédicaux –

Partie 2-16: Exigences particulières pour la sécurité de base et les performances essentielles des appareils d'hémodialyse, d'hémodiafiltration et d'hémofiltration

CONTENTS

FOREWORD.....	3
INTRODUCTION.....	6
201.1 Scope, object and related standards	7
201.2 Normative references	9
201.3 Terms and definitions	10
201.4 General requirements	12
201.5 General requirements for testing of ME EQUIPMENT	15
201.6 Classification of ME EQUIPMENT and ME SYSTEMS	16
201.7 ME EQUIPMENT identification, marking and documents	16
201.8 Protection against electrical HAZARDS from ME EQUIPMENT	19
201.9 Protection against MECHANICAL HAZARDS of ME EQUIPMENT and ME SYSTEMS	20
201.10 Protection against unwanted and excessive radiation HAZARDS	21
201.11 Protection against excessive temperatures and other HAZARDS	21
201.12 * Accuracy of controls and instruments and protection against hazardous outputs.....	22
201.13 HAZARDOUS SITUATIONS and fault conditions	30
201.14 PROGRAMMABLE ELECTRICAL MEDICAL SYSTEMS (PEMS).....	31
201.15 Construction of ME EQUIPMENT.....	31
201.16 * ME SYSTEMS	32
201.17 Electromagnetic compatibility of ME EQUIPMENT and ME SYSTEMS	33
202 Electromagnetic compatibility – Requirements and tests.....	33
208 General requirements, tests and guidance for alarm systems in medical electrical equipment and medical electrical systems	33
210 Process requirements for the development of PHYSIOLOGIC CLOSED-LOOP CONTROLLERS.....	35
211 * Requirements for MEDICAL ELECTRICAL EQUIPMENT and MEDICAL ELECTRICAL SYSTEMS used in the HOME HEALTHCARE ENVIRONMENT	35
Annexes	35
Annex G (normative) Protection against HAZARDS of ignition of flammable anaesthetic mixtures	36
Annex AA (informative) Particular guidance and rationale	37
Annex BB (informative) Examples of HAZARDS, foreseeable sequences of events, and HAZARDOUS SITUATIONS in HAEMODIALYSIS EQUIPMENT	55
Bibliography	63
Index of defined terms used in this particular standard	64
Figure 201.101 – Continuous air infusion test set-up with example dimensions	28
Figure AA.1 – Example of the HAEMODIALYSIS ME SYSTEM	51
Table 201.101 – ESSENTIAL PERFORMANCE requirements	12
Table AA.1: Possible ALARM CONDITION priorities according to 6.1.2 of IEC 60601-1-8:2006,.....	53
Table BB.1 – Hazardous situation list following ISO 14971:2007, Annex E	55