



SLOVENSKI STANDARD

SIST EN ISO 80601-2-13:2013/A2:2020

01-februar-2020

Medicinska električna oprema - 2-13: del: Posebne zahteve za osnovno varnost in bistvene lastnosti delovnega mesta za anestezijo - Dopolnilo A2 (ISO 80601-2-13:2011/Amd 2:2018)

Medical electrical equipment - Part 2-13: Particular requirements for basic safety and essential performance of an anaesthetic workstation - Amendment 2 (ISO 80601-2-13:2011/Amd 2:2018)

Medizinische elektrische Geräte - Teil 2-13: Besondere Festlegungen für die Sicherheit einschließlich der wesentlichen Leistungsmerkmale für Anästhesie-Arbeitsplätzen - Änderung 2 (ISO 80601-2-13:2011/Amd 2:2018)

Appareils électromédicaux - Partie 2-13: Exigences particulières de sécurité de base et de performances essentielles pour les postes de travail d'anesthésie - Amendement 2 (ISO 80601-2-13:2011/Amd 2:2018)

Ta slovenski standard je istoveten z: EN ISO 80601-2-13:2012/A2:2019

ICS:

11.040.10	Anestzijska, respiratorna in reanimacijska oprema	Anaesthetic, respiratory and reanimation equipment
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SIST EN ISO 80601-2-13:2013/A2:2020 en

iTeh STANDARD PREVIEW
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Full standard:
<https://standards.itech.ai/catalog/standards/sist/e582a258-3efc-47b0-99fc-3bc41fcee671/sist-en-iso-80601-2-13-2013-a2-2020>

EUROPEAN STANDARD
NORME EUROPÉENNE
EUROPÄISCHE NORM

**EN ISO 80601-2-
13:2012/A2**

November 2019

ICS 11.040.10

English Version

**Medical electrical equipment - Part 2-13: Particular
requirements for basic safety and essential performance of
an anaesthetic workstation - Amendment 2 (ISO 80601-2-
13:2011/Amd 2:2018)**

Appareils électromédicaux - Partie 2-13: Exigences
particulières de sécurité de base et de performances
essentiels pour les postes de travail d'anesthésie -
Amendement 2 (ISO 80601-2-13:2011/Amd 2:2018)

Medizinische elektrische Geräte - Teil 2-13: Besondere
Festlegungen für die Sicherheit einschließlich der
wesentlichen Leistungsmerkmale für Anästhesie-
Arbeitsplätzen - Änderung 2 (ISO 80601-2-
13:2011/Amd 2:2018)

This amendment A2 modifies the European Standard EN ISO 80601-2-13:2012, it was approved by CEN on 4 November 2019.

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

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EUROPEAN COMMITTEE FOR STANDARDIZATION
COMITÉ EUROPÉEN DE NORMALISATION
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European foreword

This document (EN ISO 80601-2-13:2012/A2:2019) 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 Amendment to the European Standard EN ISO 80601-2-13:2012 shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by May 2020, and conflicting national standards shall be withdrawn at the latest by May 2020.

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

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, Republic of North Macedonia, Romania, Serbia, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey and the United Kingdom.

Endorsement notice

The text of ISO 80601-2-13:2011/Amd 2:2018 has been approved by CEN as EN ISO 80601-2-13:2012/A2:2019 without any modification.

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INTERNATIONAL STANDARD

ISO 80601-2-13

First edition
2011-08-01
AMENDMENT 2
2018-07

Medical electrical equipment — Part 2-13: Particular requirements for basic safety and essential performance of an anaesthetic workstation

AMENDMENT 2

Appareils électromédicaux —

*Partie 2-13: Exigences particulières de sécurité de base et de
performances essentielles pour les postes de travail d'anesthésie*

AMENDEMENT 2



Reference number
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Foreword

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The procedures used to develop this document and those intended for its further maintenance are described in the ISO/IEC Directives, Part 1. In particular the different approval criteria needed for the different types of ISO documents should be noted. This document was drafted in accordance with the editorial rules of the ISO/IEC Directives, Part 2 (see www.iso.org/directives).

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This document was prepared by ISO/TC 121, *Anaesthetic and respiratory equipment*, Subcommittee SC 1, *Breathing attachments and anaesthetic machines* and Technical Committee IEC/TC 62, *Electrical equipment in medical practice*, Subcommittee SC 62 D, *Electromedical equipment*.