

SLOVENSKI STANDARD SIST EN ISO 15002:2008/A1:2020

01-februar-2020

Pretočni merilniki za priključitev na končne dele napeljav za medicinske pline - Dopolnilo A1 (ISO 15002:2008/Amd 1:2018)

Flow-metering devices for connection to terminal units of medical gas pipeline systems - Amendment 1 (ISO 15002:2008/Amd 1:2018)

Durchflussmesseinrichtungen zum Anschluss an Entnahmestellen von Rohrleitungssystemen für medizinische Gase - Änderung 1 (ISO 15002:2008/Amd 1:2018)

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Dispositifs de mesure de débit pour raccordement aux prises murales des systèmes de distribution de gaz médicaux - Amendement 1 (ISO 15002:2008/Amd 1:2018)

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Ta slovenski standard je istoveten z: EN ISO 15002:2008/A1:2019

ICS:

11.040.10 Anestezijska, respiratorna in Anaesthetic, respiratory and reanimacijska oprema reanimation equipment

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EUROPEAN STANDARD NORME EUROPÉENNE EUROPÄISCHE NORM

EN ISO 15002:2008/A1

November 2019

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English Version

Flow-metering devices for connection to terminal units of medical gas pipeline systems - Amendment 1 (ISO 15002:2008/Amd 1:2018)

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This amendment A1 modifies the European Standard EN ISO 15002:2008; 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.

SIST EN ISO 15002:2008/A1:2020

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

CEN-CENELEC Management Centre: Rue de la Science 23, B-1040 Brussels

EN ISO 15002:2008/A1:2019 (E)

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EN ISO 15002:2008/A1:2019 (E)

European foreword

This document (EN ISO 15002:2008/A1:2019) 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 15002:2008 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.

iTeh STANDARD PREVIEW

The text of ISO 15002:2008/Amd 1:2018 has been approved by CEN as EN ISO 15002:2008/A1:2019 without any modification.

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INTERNATIONAL STANDARD

ISO 15002

Second edition 2008-07-01 **AMENDMENT 1** 2018-01

Flow-metering devices for connection to terminal units of medical gas pipeline systems

AMENDMENT 1

Dispositifs de mesure de débit pour raccordement aux prises murales des systèmes de distribution de gaz médicaux

AMENDARD PRE AMENDEMENT 1 (standards.iteh.ai)

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ISO 15002:2008/Amd.1:2018(E)

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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