

SLOVENSKI STANDARD

SIST EN ISO 10079-1:2016/A1:2019

01-april-2019

Medicinska sukcijška (aspiracijska) oprema - 1. del: Električna sukcijška (aspiracijska) oprema - Dopolnilo A1: Spremembe zahtev za delovanje pri ekstremnih temperaturah (ISO 10079-1:2015/Amd 1:2018)

Medical suction equipment - Part 1: Electrically powered suction equipment - Amendment 1: Changes to requirements for operating at extremes of temperature (ISO 10079-1:2015/Amd 1:2018)

Medizinische Absauggeräte — Teil 1: Elektrisch betriebene Absauggeräte - Änderung 1: Änderungen der Anforderungen an Grenzwerte für Betriebstemperaturen (ISO 10079-1:2015/Amd 1:2018)

Appareils d'aspiration médicale - Partie 1: Appareils électriques d'aspiration - Amendement 1: Modifications des exigences de fonctionnement à des valeurs extrêmes de température (ISO 10079-1:2015/Amd 1:2018)

Ta slovenski standard je istoveten z: EN ISO 10079-1:2015/A1:2019

ICS:

11.040.10	Anestezijska, respiratorna in reanimacijska oprema	Anaesthetic, respiratory and reanimation equipment
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EUROPEAN STANDARD
NORME EUROPÉENNE
EUROPÄISCHE NORM

EN ISO 10079-1:2015/A1

January 2019

ICS 11.040.10

English Version

**Medical suction equipment - Part 1: Electrically powered
suction equipment - Amendment 1: Changes to
requirements for operating at extremes of temperature
(ISO 10079-1:2015/Amd 1:2018)**

Appareils d'aspiration médicale - Partie 1: Appareils
électriques d'aspiration - Amendement 1:
Modifications des exigences de fonctionnement à des
valeurs extrêmes de température (ISO 10079-
1:2015/Amd 1:2018)

Medizinische Absauggeräte - Teil 1: Elektrisch
betriebene Absauggeräte - Änderung 1: Änderungen
der Anforderungen an Grenzwerte für
Betriebstemperaturen (ISO 10079-1:2015/Amd
1:2018)

This amendment A1 modifies the European Standard EN ISO 10079-1:2015; it was approved by CEN on 13 December 2018.

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
EUROPÄISCHES KOMITEE FÜR NORMUNG

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

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European foreword

This document (EN ISO 10079-1:2015/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 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 July 2019, and conflicting national standards shall be withdrawn at the latest by July 2019.

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, Former Yugoslav Republic of Macedonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Serbia, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey and the United Kingdom.

Endorsement notice

The text of ISO 10079-1:2015/Amd 1:2018 has been approved by CEN as EN ISO 10079-1:2015/A1:2019 without any modification.

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INTERNATIONAL
STANDARDISO
10079-1Third edition
2015-11-01**AMENDMENT 1**
2018-06

Medical suction equipment —

Part 1:

**Electrically powered suction
equipment**

AMENDMENT 1: Changes to

requirements for operating at extremes
of temperature
*(standards.iteh.ai)**Appareils d'aspiration médicale —**Partie 1: Appareils électriques d'aspiration**AMENDEMENT 1: Modifications des exigences de fonctionnement à
des valeurs extrêmes de température*Reference number
ISO 10079-1:2015/Amd.1:2018(E)

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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.

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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For an explanation on the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT) see the following URL: www.iso.org/iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 121, *Anaesthetic and respiratory equipment*, Subcommittee SC 8, *Suction devices for hospital and emergency care use*.

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