

STN	<p>Kardiovaskulárne implantáty a umelé orgány Výmenníky pre krv-plyn (oxygenátory) (ISO 7199: 2016/Amd 1: 2020) Zmena A1</p>	<p>STN EN ISO 7199/A1</p>
		85 6250

Cardiovascular implants and artificial organs - Blood-gas exchangers (oxygenators) (ISO 7199:2016)

Táto norma obsahuje anglickú verziu európskej normy.
This standard includes the English version of the European Standard.

Táto norma bola označená vo Vestníku ÚNMS SR č. 06/20

Obsahuje: EN ISO 7199:2017/A1:2020, ISO 7199:2016/Amd 1:2020

130956

EUROPEAN STANDARD
NORME EUROPÉENNE
EUROPÄISCHE NORM

EN ISO 7199:2017/A1

April 2020

ICS 11.040.40

English Version

**Cardiovascular implants and artificial organs - Blood-gas
exchangers (oxygenators) - Amendment 1: Connectors
(ISO 7199:2016/Amd 1:2020)**

Implants cardiovasculaires et organes artificiels -
Échangeurs gaz/sang extracorporels (oxygénateurs) -
Amendment 1 (ISO 7199:2016/Amd 1:2020)

Kardiovaskuläre Implantate und künstliche Organe -
Blut-Gas-Austauscher (Oxygenatoren) - Änderung 1:
Anschlüsse (ISO 7199:2016/Amd 1:2020)

This amendment A1 modifies the European Standard EN ISO 7199:2017; it was approved by CEN on 4 March 2020.

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.

CEN members are the national standards bodies of 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 United Kingdom.



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

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

Contents

	Page
European foreword.....	3

European foreword

This document (EN ISO 7199:2017/A1:2020) has been prepared by Technical Committee ISO/TC 150 "Implants for surgery" in collaboration with Technical Committee CEN/TC 205 "Non-active 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 October 2020, and conflicting national standards shall be withdrawn at the latest by October 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 7199:2016/Amd 1:2020 has been approved by CEN as EN ISO 7199:2017/A1:2020 without any modification.

koniec náhľadu – text d'alej pokračuje v platenej verzii STN