SLOVENSKI STANDARD
SIST EN ISO 7396-1:2007/oprA2:2009

01-april-2009

Sistemi napeljav za medicinske pline - 1. del: Napeljave za stisnjene medicinske pline in podtlak - Dopolnilo A2 (ISO 7396-1:2007/DAM 2:2009)

Medical gas pipeline systems - Part 1: Pipeline systems for compressed medical gases and vacuum - Amendment 2 (ISO 7396-1:2007/DAM 2:2009)

Rohrleitungssysteme für medizinische Gase - Teil 1: Rohrleitungenssyteme für medizinische Druckgase und Vakuum - Änderung 2 (ISO 7396-1:2007/DAM 2:2009)

Systèmes de distribution de gaz médicaux - Partie 1: Systèmes de distribution de gaz médicaux comprimés et de vide - Amendement 2 (ISO 7396-1:2007/DAM 2:2009)

Ta slovenski standard je istoveten z: **EN ISO 7396-1:2007/prA2**

ICS:

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

SIST EN ISO 7396-1:2007/oprA2:2009 **en**

EUROPEAN STANDARD

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Medical gas pipeline systems - Part 1: Pipeline systems for compressed medical gases and vacuum - Amendment 2 (ISO 7396-1:2007/DAM 2:2009)

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This draft amendment is submitted to CEN members for parallel enquiry. It has been drawn up by the Technical Committee CEN/TC 215.

This draft amendment A2, if approved, will modify the European Standard EN ISO 7396-1:2007. If this draft becomes an amendment, 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.

This draft amendment was established by CEN 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 Management Centre has the same status as the official versions.

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Recipients of this draft are invited to submit, with their comments, notification of any relevant patent rights of which they are aware and to provide supporting documentation.

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

Management Centre: Avenue Marnix 17, B-1000 Brussels

Contents

Page

Foreword

This document (EN ISO 7396-1:2007/prA2:2009) 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 document is currently submitted to the parallel Enquiry.

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

Endorsement notice

The text of ISO 7396-1:2007/DAM 2:2009 has been approved by CEN as a EN ISO 7396-1:2007/prA2:2009 without any modification.

DRAFT AMENDMENT ISO 7396-1:2007/DAmd 2

ISO/TC **121**/SC **6** Secretariat: **ANSI**

Voting begins on: **2009-02-12** Voting terminates on: **2009-07-12**

Medical gas pipeline systems —

Part 1:
Pipeline systems for compressed medical gases and vacuum

AMENDMENT 2

Systèmes de distribution de gaz médicaux —

Partie 1: Systèmes de distribution de gaz médicaux comprimés et de vide

AMENDEMENT 2

ICS 11.040.10

ISO/CEN PARALLEL PROCESSING

This draft has been developed within the International Organization for Standardization (ISO), and processed under the **ISO-lead** mode of collaboration as defined in the Vienna Agreement.

This draft is hereby submitted to the ISO member bodies and to the CEN member bodies for a parallel five-month enquiry.

Should this draft be accepted, a final draft, established on the basis of comments received, will be submitted to a parallel two-month approval vote in ISO and formal vote in CEN.

To expedite distribution, this document is circulated as received from the committee secretariat. ISO Central Secretariat work of editing and text composition will be undertaken at publication stage.

Pour accélérer la distribution, le présent document est distribué tel qu'il est parvenu du secrétariat du comité. Le travail de rédaction et de composition de texte sera effectué au Secrétariat central de l'ISO au stade de publication.

Contents

Page