
Medical gas pipeline systems —
Part 1:
Pipeline systems for compressed
medical gases and vacuum
AMENDMENT 1

iTeh STANDARD PREVIEW
(standards.iteh.ai)

Systèmes de distribution de gaz médicaux —

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

ISO 7396-1:2016/Amd 1:2017

AMENDEMENT 1

<https://standards.iteh.ai/catalog/standards/sist/4b1ed067-49e9-4943-9f97-1aff927059d5/iso-7396-1-2016-amd-1-2017>



iTeh STANDARD PREVIEW
(standards.iteh.ai)

ISO 7396-1:2016/Amd 1:2017
<https://standards.iteh.ai/catalog/standards/sist/4b1ed067-49e9-4943-9f97-1aff927059d5/iso-7396-1-2016-amd-1-2017>



COPYRIGHT PROTECTED DOCUMENT

© ISO 2017, Published in Switzerland

All rights reserved. Unless otherwise specified, no part of this publication may be reproduced or utilized otherwise in any form or by any means, electronic or mechanical, including photocopying, or posting on the internet or an intranet, without prior written permission. Permission can be requested from either ISO at the address below or ISO's member body in the country of the requester.

ISO copyright office
Ch. de Blandonnet 8 • CP 401
CH-1214 Vernier, Geneva, Switzerland
Tel. +41 22 749 01 11
Fax +41 22 749 09 47
copyright@iso.org
www.iso.org

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).

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights. Details of any patent rights identified during the development of the document will be in the Introduction and/or on the ISO list of patent declarations received (see www.iso.org/patents).

Any trade name used in this document is information given for the convenience of users and does not constitute an endorsement.

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 6, *Medical gas systems*.

A list of all parts in the ISO 7396 series can be found on the ISO website.

iTeh STANDARD PREVIEW
(standards.iteh.ai)

ISO 7396-1:2016/Amd 1:2017
<https://standards.iteh.ai/catalog/standards/sist/4b1ed067-49e9-4943-9f97-1aff927059d5/iso-7396-1-2016-amd-1-2017>

Medical gas pipeline systems —

Part 1:

Pipeline systems for compressed medical gases and vacuum

AMENDMENT 1

5.8.2, fifth paragraph

Replace the following paragraph:

Locations of supply systems which contain gases other than medical air and in which the gas can accumulate shall be provided with an oxygen monitor with an indicator at the entry to warn of oxygen concentrations below 19.5% or above 23.5%. The monitor shall activate an alarm with an auditory and visual signal at the entrance to warn of oxygen concentrations below 19.5% or above 23.5%.

with:

Locations of supply systems which contain gases other than CO₂, medical air and air for driving surgical tools, and in which the gas can accumulate, shall be provided with an oxygen monitor with an indicator at the entry to warn of oxygen concentrations below 19,5 % or above 23,5 %. The monitor shall activate an alarm with audible and visual signals outside the entrance to warn of oxygen concentrations below 19,5 % or above 23,5 %. The O₂ sensor should be installed at a height of approximately 1 m. Labelling information on risks (anoxia, hypoxia) shall be placed on the outside of the entrance door. In case of supply system for CO₂, a CO₂ sensor with an indicator shall be installed at the entry to ensure the safety of personnel. In this case, the threshold level for CO₂ is 1,5 % in ambient air. The monitor shall activate an alarm with audible and visual signals outside the entrance, to warn of CO₂ concentrations equal to or above 1,5 %. The CO₂ sensor shall be installed at height of approximately 1 m.

For CO₂, complementary to these requirements labelling, information on risks (anoxia, hypoxia) shall be placed outside of the room in a position that is clearly visible to anyone entering the room.

iTeh STANDARD PREVIEW
(standards.iteh.ai)

ISO 7396-1:2016/Amd 1:2017
<https://standards.iteh.ai/catalog/standards/sist/4b1ed067-49e9-4943-9f97-1aff927059d5/iso-7396-1-2016-amd-1-2017>