

Oxygen Sensor OOM201

Use the advantages:

- Compliant with European MDD (CE certification)
- Meets ISO 80601-2-55
- Designed and manufactured according to EN ISO 13485
- Accurate and reliable fast response
- Resistant to N₂O
- Excellent signal stability
- High product quality
- Short delivery times
- Technical support
- · Made in Germany
- FDA cleared



From standard sensors to customized sensors

Experienced EnviteC engineers analyze customer requirements. This input is used for different standard and OEM applications, and ongoing support is provided right up to the final integration in the solution. EnviteC designs customized sensors characterized by a maximum possible degree of precision, for example with different signal levels or temperature compensation elements.

Intendend use

The EnviteC Medical Oxygen Sensors are intended as oxygensensing component of an oxygen analyzer that measures oxygen concentration in breathing gas mixtures in the following applications:

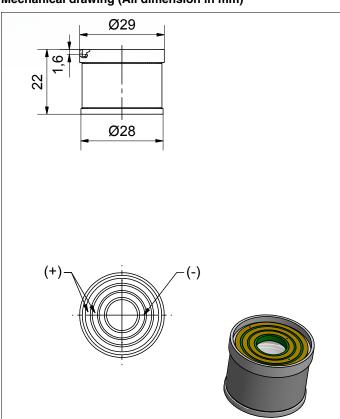
Sensing device for oxygen in

- control device of oxygen concentrators
- medical ventilators
- anaesthesia equipment
- incubators.

The use is limited to system monitoring. The sensors are not suited for breath by breath analysis of breath gases. Please refer to the Instructions for Use! If the sensor is intended to replace the original oxygen-sensing component of an oxygen analyzer, consult the EnviteC XRL Cross Reference List for selecting the appropriate sensor.



Mechanical drawing (All dimension in mm)



General tolerances ISO 2768-c

Additional information

The Instructions for Use as well as the EnviteC XRL Cross Reference List are available under www.EnviteC.com and in the Apple App Store under EnviteC XRL as free download.

For more information please contact us!

We look forward to assisting you either on the phone or in a personal talk.

Technical Specifications OOM201

Measurement range	0 % 100 % oxygen (at atmospheric pressure)
Nominal sensor lifetime	≥ 500 000 % volume oxygen hours
Output in ambient air	14 mV 20.7 mV (Dual Cathode), load 600 Ohms
Electrical interface	Gold plated slip rings
Accuracy	meets ISO 80601-2-55 requirements
Repeatability	$<$ 1 % volume O_2 at constant temperature and pressure
Linearity error	< 3 % relative
Response time	< 12 s to 90 % of final value
Zero offset voltage	< 200 µV in 100 % nitrogen, applied for 5 min
Cross interference	meets ISO 80601-2-55 requirements
Influence of humidity	-0.03 % rel. per % RH at 25 °C
Pressure range	0.6 bar 2 bar (ppO $_2$ 0 1250 mbar O $_2$)
Influence of pressure	proportional to change in oxygen partial pressure
Influence of mechanical shock	< 1 % relative after a fall from 1 m
Operating temperature	0 °C +50 °C
Temperature compensation	no temperature compensation
Operating humidity	0 % 99 % RH non-condensing
Long term output drift	< 1 % volume oxygen per month
	typically < -15 % relative over lifetime
Storage temperature	-20 °C +50 °C
Recommended storage	+5 °C +15 °C
Recommended load	≥ 10 kOhms
Warm-up time	< 30 minutes, after replacement of sensor
Weight	approximately 28 grams
Part number	01-00-0014





All specifications are applicable at standard conditions: 1013 hPa, 25 °C dry ambient air

For suitable accessories and sensors please refer to the EnviteC Cross Reference List under www.EnviteC.com and in the Apple App Store unter EnviteC XRL as free download.

EnviteC-Wismar GmbH a Honeywell Company

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Doc. No. 001-33-Datasheet_OOM201-0 March 2016 Technical information is subject to change without notice! © 2016 Honeywell International Inc.

EnviteC by Honeywell reserves the right to make changes in product specifications and adjust its production at any time and without notice.





Oxygen Sensor OOM202

Use the advantages:

- Compliant with European MDD (CE certification)
- Meets ISO 80601-2-55
- Designed and manufactured according to EN ISO 13485
- Accurate and reliable fast response
- Resistant to N₂O
- Excellent signal stability
- High product quality
- Short delivery times
- Technical support
- · Made in Germany
- FDA cleared



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Intendend use

The EnviteC Medical Oxygen Sensors are intended as oxygensensing component of an oxygen analyzer that measures oxygen concentration in breathing gas mixtures in the following applications:

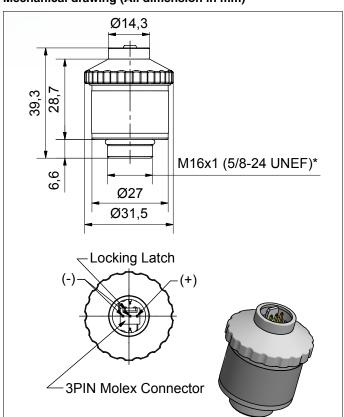
Sensing device for oxygen in

- control device of oxygen concentrators
- medical ventilators
- anaesthesia equipment
- incubators.

The use is limited to system monitoring. The sensors are not suited for breath by breath analysis of breath gases. Please refer to the Instructions for Use! If the sensor is intended to replace the original oxygen-sensing component of an oxygen analyzer, consult the EnviteC XRL Cross Reference List for selecting the appropriate sensor.



Mechanical drawing (All dimension in mm)



General tolerances ISO 2768-c

*Intermediate thread: Metric / Unified Extra Fine

Additional information

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Technical Specifications OOM202

Measurement range	0 % 100 % oxygen (at atmospheric pressure)
Nominal sensor lifetime	≥ 1 000 000 % volume oxygen hours
Output in ambient air	13 mV 16 mV
Electrical interface	3 pin (Molex® 22-11-1031)
Accuracy	meets ISO 80601-2-55 requirements
Repeatability	$<$ 1 % volume O_2 at constant temperature and pressure
Linearity error	< 3 % relative
Response time	< 12 s to 90 % of final value
Zero offset voltage	< 200 μV in 100 % nitrogen, applied for 5 min
Cross interference	meets ISO 80601-2-55 requirements
Influence of humidity	-0.03 % rel. per % RH at 25 °C
Pressure range	0.6 bar 2 bar (ppO ₂ 0 1250 mbar O ₂)
Influence of pressure	proportional to change in oxygen partial pressure
Influence of mechanical shock	< 1 % relative after a fall from 1 m
Operating temperature	0 °C +50 °C
Temperature compensation	built-in NTC compensation
Effect of temperature compensation (steady	between +25 °C and +40 °C: 3 % relative error
state)	between 0 °C and +50 °C: 8 % relative error
Operating humidity	0 % 99 % RH non-condensing
Long term output drift	< 1 % volume oxygen per month
	typically < -15 % relative over lifetime
Storage temperature	-20 °C +50 °C
Recommended storage	+5 °C +15 °C
Recommended load	≥ 10 kOhms
Warm-up time	< 30 minutes, after replacement of sensor
Weight	approximately 28 grams
Part number	01-00-0047





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Flow Sensor SpiroQuant A+

Use the advantages:

- Proven constant temperature hot-wire anemometry
- Fast response time
- High product quality
- RoHS conform
- Biocompatible components
- Short delivery times
- Technical support
- Made in Germany
- Certified quality management system according EN ISO 13485



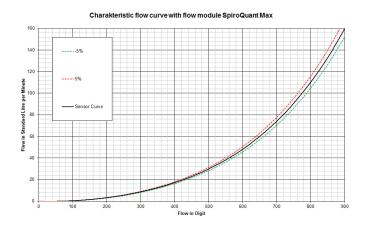
"We keep your devices operating at their best."

EnviteC has been developing and manufacturing highly specialized products for medical applications. EnviteC's research and development activities are consistently aligned to customer and market needs – identification and optimization of sustainable product solutions are the main concerns of the company.

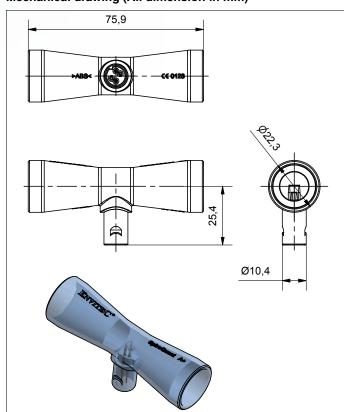
The company trusts in the quality of advanced production methods and processes. The result are flow measurement solutions for medical applications, which share the same extraordinary quality, excellent signal stability and reliable response for accurate readings. The flow sensor SpiroQuant A+ is a hot-wire sensor for measuring volume gas flow in anesthesia machines and ventilators. The flow sensor is not suitable for sterilization. For operation please refer to the instructions for use of the basic device to which this sensor is connected.

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Mechanical drawing (All dimension in mm)



General tolerances ISO 2768-c

Additional information

If the sensor is intended to replace the original flow-sensing component in anesthesia machines or ventilators, consult the EnviteC XRL Cross Reference List under www. EnviteC.com for selecting the appropriate sensor.

For more information please contact us!

We look forward to assisting you either on the phone or in a personal talk.

Technical Specifications SpiroQuant A+

Measuring principle	Constant temperature hot-wire anemometry	
Cross reference	Designed for EnviteC SpiroQuant MAX module; compatible with Dräger Spirolog® flow	
	sensor	
Sensor type	Disposable sensor for single patient use	
Flow range	0 160 slpm (standard liters per minute)	
Accuracy	± 7% from measured value between 0 10 l/min with flow module SpiroQuant MAX;	
	± 5% from measured value between 10 160 l/min with flow module SpiroQuant MAX;	
	(± 8 % reading used in Dräger ventilators)	
Flow resistance	< 2.5 mbar (added resistance)	
Operating temperature	+15 °C +40 °C	
Operating time	Single patient use, applicable as long as a sensor calibration is successful	
Storage temperature	-20 °C +70 °C	
Storage ambient pressure	570 1200 hPa	
Storage humidity	5 % 95 % RH in orginal packaging	
Storage time	Max. 5 years in original packaging	
Cleaning / disinfection	Single patient use	
Weight	Approx. 10 g	
Electrical connection	Gold plated 4 pin (compatible with Dräger Spirolog® sensor)	
Material	Housing parts: MABS;	
	Contact pins: gold-plated;	
	Hot-wires: platin	
Part number	1000470: Flow sensor SpiroQuant A+ (Box with 5 pieces);	
	1001487: Flow module SpiroQuant MAX	

All specifications are applicable at standard conditions: 1013 hPa, 25 $^{\circ}$ C dry ambient air



Certified Quality Management

EnviteC is maintaining a quality management system, which meets the requirements of EN ISO 13485 for medical devices.

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Doc. No. 007-05-SpiroQuant_A+_Spec-0 December 2017 Technical information is subject to change without notice! © 2017 Honeywell International Inc.

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EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 021697 0017 Rev. 01

Manufacturer:

EnviteC - Wismar GmbH

Alter Holzhafen 18 23966 Wismar GERMANY

Facility(ies):

EnviteC - Wismar GmbH

Alter Holzhafen 18, 23966 Wismar, GERMANY

Product Category(ies): Oxygen Saturation Sensors and Monitors,

Sensors and Control Units for Monitoring of Respiratory Parameters and Gas Exchange, Non-invasive Blood Pressure Equipment,

Temperature Sensors

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.:

713172795

Valid from:

2020-02-17

Valid until:

2024-05-26

Date,

2020-02-17

Christoph Dicks

Head of Certification/Notified Body









DAKKS CRT2 / 10.13



CERTIFICATE

No. Q5 17 12 21697 018

Holder of Certificate: EnviteC - Wismar GmbH

> Alter Holzhafen 18 23966 Wismar **GERMANY**

Facility(ies):

EnviteC - Wismar GmbH

Alter Holzhafen 18, 23966 Wismar, GERMANY

Certification Mark:





Scope of Certificate: Design and development, production and distribution

of sensors and control units for monitoring of vital physiological parameters, sensors and control units for monitoring of respiratory mechanics parameters and gas exchange, measurement devices and sensors

for alcohol blood concentration

Applied

EN ISO 13485:2016

Standard(s):

Medical devices - Quality management systems -

Requirements for regulatory purposes

(ISO 13485:2016) DIN EN ISO 13485:2016

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). See also notes overleaf.

Report No.:

713119332

Valid from:

2018-02-06

Valid until:

2021-01-30

Date, 2018-02-06

Stefan Preiß







