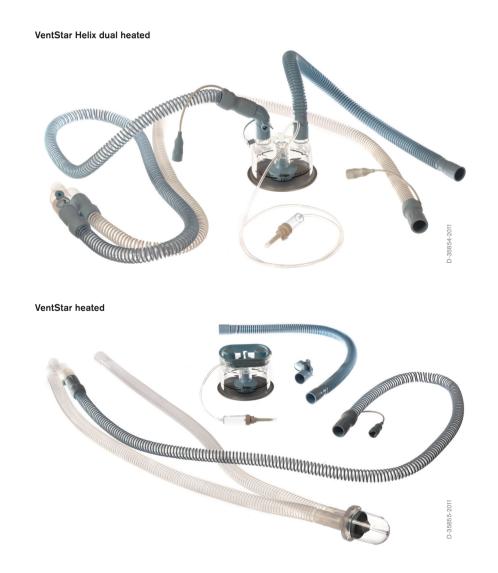


VentStar Helix - Heated Breathing Circuits Consumables and Accessories

Breathing gas humidification with heated hose systems in mechanically ventilated patients is an essential component in today's intensive care. Choosing the right hose system can support the therapy success.



Benefits

Clinical application

Heated hose systems are used to avoid damage to the lungs and airways of longterm ventilated patients due to dry and cool breathing gases. The patient is adequate ly hydrated when a breathing gas humidification system is used [1]. This maintains the mucociliary clearance and secretions can be transported. The risk of infection is reduced subsequently [2].

Ricard JD, Cook D, Griffith L, Brochard L, Dreyfuss D. Physicians' attitude to use heat and moisture exchangers or heated humidifiers: a Franco-Canadian survey. Intensive Care Med 2002; 28: 719-725.

[2] Williams T. Humidification in intensive care SAJCC: July 2005, Vol. 21, No. 1: 26-31

Innovative hose heater design

The innovative double helix ensures a more uniform heating of the breathing gas than conventional hose heaters. It supports the prevention of condensation and provides patients with the optimal level of hydration. No hose heater inside the hose - for optimised resistance values.

Versatile use

Removable Y-adapter for carrying out oxygen therapy in conjunction with the humidifier and the ventilator or to insert a drug nebuliser in the inspiratory branch of the hose system.

High quality standards

Due to the high Dräger quality standards, all respiratory gas-carrying components are tested for biocompatibility and do not contain any potentially hazardous plasticizers such as DEHP.

Wide product range

Dräger offers a wide range of humidi - fication products. VentStar Helix is part of the product range consisting of breathing hose systems and water chambers for active humidification in the ICU. All relevant respiratory gas conditioning accessories come from a single supplier.

Efficient workflows

Designed as a disposable product, the VentStar Helix provides heat for the optimal support of the clinical workflows. No disassembly of components is required to prepare for cleaning and sterilisation. After use with a patient, the entire system can be disposed of in one step. This means the medical staff has more time to focus on the care of the patient.

Technical Data

Length of breathing hose	1,63m (64,20 Inch)
Material	
Breathing hoses	PP (polypropylene)
Connectors	PP (polypropylene)
Y-adapter	PP (polypropylene)
U.midifiar abambar	DC SDC cilicana aluminium

Ordering Information

AutoFeed water chamber	MP02605
VentStar Helix dual heated	MP02606
VentStar Helix heated	MP02607

Notes

CORPORATE HEADQUARTERS

Drägerwerk AG & Co. KGaA Moislinger Allee 53–55 23558 Lübeck, Germany www.draeger.com

Manufacturer:

Dräger Medical GmbH Moislinger Allee 53-55 23558 Lübeck, Germany

As of August 2015

Dräger Medical GmbH changes to Drägerwerk AG & Co. KGaA

Locate your Regional Sales Representative at: www.draeger.com/contact



REGION EUROPE CENTRAL AND EUROPE NORTH

Dräger Medical GmbH Moislinger Allee 53-55 23558 Lübeck, Germany Tel +49 451 882 0 Fax +49 451 882 2080 info@draeger.com

REGION EUROPE SOUTH

Dräger Médical S.A.S.
Parc de Haute Technologie
d'Antony 2
25, rue Georges Besse
92182 Antony Cedex, France
Tel +33 1 46 11 56 00
Fax +33 1 40 96 97 20
dlmfr-contact@draeger.com

REGION MIDDLE EAST, AFRICA

Dräger Medical GmbH Branch Office P.O. Box 505108 Dubai, United Arab Emirates Tel +971 4 4294 600 Fax +971 4 4294 699 contactuae@draeger.com

REGION ASIA / PACIFIC

Draeger Medical
South East Asia Pte Ltd.
25 International Business Park
#04-27/29 German Centre
Singapore 609916, Singapore
Tel +65 6572 4388
Fax +65 6572 4399
asia.pacific@draeger.com

REGION CENTRAL AND SOUTH AMERICA

Dräger Panama Comercial S. de R.L.
Complejo Business Park, V tower, 10th floor
Panama City
Tel +507 377 9100
Fax +507 377 9130
contactcsa@draeger.com

The quality management system at Dräger Medical GmbH is certified according to ISO 13485, ISO 9001 and Annex II.3 of Directive 93/42/EEC (Medical devices).