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## DECLARATION OF CONFORMITY FOR THE PRODUCT(S)

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### Freevent HME 22

**Basic UDI-DI: 7331791-HME-0-000-0011-XH**

This Declaration of Conformity is issued under the sole responsibility of Atos Medical AB. Atos Medical AB declares that the devices covered by the present declaration are in conformity with the Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices and the requirements specified herein are fulfilled and, clause 6.6 of Schedule 3 to the Australian Therapeutic Goods (Medical Devices) Regulations 2002 including amendments in effect at the issuance date.

**Intended use/purpose:**

Freevent HME 22 Regular is a Heat and Moisture Exchanger (HME) intended for spontaneously breathing tracheostomized patients using a tracheostomy tube with a Ø22mm connector. The HME conditions inhaled air by retaining heat and moisture from the exhaled air. The device also partially restores lost breathing resistance.

The HME is used in combination with Freevent DualCare Speaking Valve/Freevent DualCare Speaking Valve Blue, with Freevent HME DigiTop/Freevent HME DigiTop Blue, or with HME DigiTop O2.

The HME is for single use, i.e. it has to be exchanged at least every 24 hours.

Hörby, Sweden, date as stated on last page

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Henrik Heringslack, Atos Medical Site Manager  
on behalf of the CEO of Atos Medical AB.

<b>Manufacturer:</b>	<b>Atos Medical AB</b> Kraftgatan 8, SE-242 35 Hörby Sweden	<b>Telephone:</b> Email: Web:	+46 (0)415 198 00 Info@atosmedical.com www.atosmedical.com
<b>SRN number:</b>	<b>SE-MF-000000725</b>		
<b>Competent Authority</b>	<b>Medical Products Agency</b> Sweden		

# DECLARATION OF CONFORMITY

FOR THE PRODUCT(S)

**7331791-HME-0-000-0011-XH**

REF	Device name	Class*	GMDN code
7747	Freevent HME 22 Regular (30pcs)	I	58705

\*Product Classification according to Annex VIII EU Medical Device Regulation 2017/745

## In compliance with Therapeutic Goods (Medical Devices) Regulations 2002

Each kind of medical device to which the system has been applied complies with the applicable provisions of the essential principles, the classification rules, and the full quality assurance procedures, at each stage, from the design of the device until its final inspection before being supplied.

## In compliance with UK Medical Devices Regulations 2002 as amended

Each kind of medical device to which the system has been applied fulfils the applicable obligations imposed by Annex VII and the device meets the provisions of the applicable section of the Medical Devices Regulations.

Authorized representative/UK Responsible Person:

Atos Medical UK Ltd  
Tottle Road  
Cartwright House  
Nottingham  
Nottinghamshire NG2 1RT  
England, United Kingdom

## Common Specification(s) as per Article 9, and other Union Legislation(s)

- No relevant Common Specifications as defined in Article 9 to list.
- No relevant Union Legislations to list
- No European Representative

Document Approvals

Approved Date: 2026-02-13

Approval Task Verdict: Approve	SEHRBHNU Ulrika Svensson, Regulatory Affairs Professional (ulrika.svensson-atosmedical@coloplast.com) Issuer 13-Feb-2026 07:39:47 GMT+0000
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Approval Task Verdict: Approve	HENRIK.HERINGSLACK Henrik Heringslack, Site Director (henrik.heringslack-atosmedical@coloplast.com) Management 13-Feb-2026 14:43:30 GMT+0000
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Approval Task Verdict: Approve	SEHRBPNH Håkan Persson, Quality Manager (hakan.persson-atosmedical@coloplast.com) Quality 13-Feb-2026 14:47:26 GMT+0000
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