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## DECLARATION OF CONFORMITY

FOR THE PRODUCT(S)

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### Freevent XtraCare

**Basic UDI-DI: 7331791-HME-0-000-0004-XM**

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 XtraCare and Freevent XtraCare Mini are single use Heat and Moisture Exchangers with electrostatic filters (HMEF) that condition and filter inhaled air in patients spontaneously breathing through a tracheostoma.

Hörby, Sweden, date as stated on last page

A handwritten signature in blue ink, appearing to read 'Henrik Heringslack', written over a dotted line.

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

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**SRN number:** SE-MF-00000725

**Competent Authority** Medical Products Agency  
Sweden

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FOR THE PRODUCT(S)

## 7331791-HME-0-000-0004-XM

REF	Device name	Class*	GMDN code
7767	Freevent XtraCare White	I	58705
7768	Freevent XtraCare Blue	I	58705
7788	Freevent XtraCare Blue	I	58705
7789	Freevent XtraCare White	I	58705
8004	Freevent XtraCare Mini White 30 pcs	I	58705
8005	Freevent XtraCare Mini Blue 30 pcs	I	58705
8006	Freevent XtraCare Mini Pink 30 pcs	I	58705
8008	Freevent XtraCare Mini White 5 pcs	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:

Coloplast Limited  
Nene Hall  
Peterborough Business Park  
Peterborough  
Cambridgeshire PE2 6FX

### 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: 2026-06-01

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