



DECLARATION OF CONFORMITY

FOR THE PRODUCT(S)

Freevent Neckband

Basic UDI-DI: 7331791-GEN-A-000-0005-EM

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 Neckband is used for holding a tube or button that connects to the trachea in place. It is a single use device intended for adult and pediatric patients. Not for neonatal use.

Hörby, Sweden, date as stated on last page

A handwritten signature in blue ink, appearing to read "Henrik Heringslack".

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

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Competent Authority Medical Products Agency
Sweden

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REF	Device name	Class*	GMDN code
1651	Freeevent Neckband, one-piece, small	I	63438
1652	Freeevent Neckband, two-piece, small	I	63438
1661	Freeevent Neckband, one-piece, large	I	63438
1662	Freeevent Neckband, two-piece, large	I	63438

*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

Approval Task Verdict: Approve	SOFTHO Sofia Thomasson, Regulatory Affairs Specialist (sofia.thomasson-atosmedical@coloplast.com) Issuer 01-Jun-2026 08:58:12 GMT+0000
Approval Task Verdict: Approve	HENRIK.HERINGSLACK Henrik Heringslack, Site Director (henrik.heringslack-atosmedical@coloplast.com) Management 01-Jun-2026 09:20:21 GMT+0000
Approval Task Verdict: Approve	KARNIL Karolina Nilsson, Head of RA, Voice & Respiratory Care (karolina.nilsson-atosmedical@coloplast.com) Regulatory 01-Jun-2026 13:16:40 GMT+0000