



DECLARATION OF CONFORMITY

FOR THE PRODUCT(S)

Provox LaryClip

Basic UDI-DI: 7331791-LTU-A-000-0001-JT

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:

The Provox LaryClip is used for extra support for LaryButton and LaryTube. The product consists of two parts, one that is attached to the patients' skin on each side of the stoma and the other part is attached to the LaryButton or the LaryTube. The two parts are then connected by Velcro.

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, AtosMedical 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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REF	Device name	Class*	GMDN code
7669	Provox LaryClip	I	35752

*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: 2026-06-15

Approval Task Verdict: Approve	SOFTHO Sofia Thomasson, Regulatory Affairs Specialist (sofia.thomasson-atosmedical@coloplast.com) Issuer 12-Jun-2026 12:25:48 GMT+0000
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