

DECLARATION OF CONFORMITY

FOR THE PRODUCT(S)

Provox Life LP Kit

Basic UDI-DI: 7331791-KIT-0-000-0006-J6

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:

Provox Life LP Kit is an assortment of devices and accessories, intended for rehabilitation for users breathing through a tracheostoma after a total laryngectomy.

Hörby, Sweden, date as stated on last page

Martin Richardson, Senior Vice President Operations & Quality on behalf of the CEO of Atos Medical AB

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Competent Authority Medical Products Agency

Sweden

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REF	Device name	Class*	GMDN code
6147	Provox Life LP Kit 18 - ADH	1	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

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^{**}The choice of GMDN code is based on the primary product in the kit, the HME.

Document Approvals Approved Date: 2025-09-18

Approval Task Verdict: Approve	HIND.ABUTALEB Hind Palm, Regulatory Affairs Professional (hind.abutaleb-atosmedical@coloplast.com) Issuer 15-Sep-2025 15:14:48 GMT+0000	
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