

# Atos

## DECLARATION OF CONFORMITY FOR THE PRODUCT(S)

### Provox FreeHands HME

**Basic UDI-DI: 7331791-HME-0-000-0003-XJ**

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 FreeHands HME Cassette/Moist/Flow is intended for single use for spontaneously breathing laryngectomized patients, utilizing a voice prosthesis and must be used in combination with a Provox FreeHands speaking valve, a Provox cap or DigiTop O2. Provox FreeHands combines pulmonary rehabilitation with its Heat and Moisture Exchanging functionality with voice rehabilitation using an Automatic Speaking Valve or Manual Occlusion. The HME conditions inhaled air by retaining heat and moisture from the exhaled air. The device also partially restores lost breathing resistance.

Hörby, Sweden, date as stated on last page



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

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**SRN number:** SE-MF-000000725

**Competent Authority** Medical Products Agency  
Sweden

# DECLARATION OF CONFORMITY

FOR THE PRODUCT(S)

**7331791-HME-0-000-0003-XJ**

REF	Device name	Class*	GMDN code
8220	Provox FreeHands HME Moist (30 pcs)	I	58705
8221	Provox FreeHands HME Flow (30 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:

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: 2024-11-22

Approval Task Verdict: Approve	MARRIC Martin Richardson, VP, International Manufacturing, Sweden, Germany & China (martin.richardson-atosmedical@coloplast.com) Management 22-Nov-2024 08:03:12 GMT+0000
Approval Task Verdict: Approve	SEHRBHNU Ulrika Svensson, Regulatory Affairs Professional (ulrika.svensson-atosmedical@coloplast.com) Issuer 22-Nov-2024 09:12:09 GMT+0000
Approval Task Verdict: Approve	KARNIL Karolina Nilsson, Head of Regulatory Affairs (karolina.nilsson-atosmedical@coloplast.com) Regulatory 22-Nov-2024 14:07:55 GMT+0000