

EC Declaration of Conformity

The Company:

Andreas Fahl Medizintechnik-Vertrieb GmbH
August-Horch-Straße 4a
51149 Köln, Germany
SRN: DE-MF-000006665

hereby account for the medical devices listed in the annex on which this declaration is based, that they conform to the following EC legislation:

Regulation (EC) 2017/745 of 05 April 2017, Medical Device Regulation (MDR)

Product group	X25900 – LARYVOX® Inserter
Basic UDI-DI	405194825900LVINSERT+VADS
Class acc. to EC regulation 2017/745 Annex VIII	Class I
Classification Rule	5
REFs / Medical Devices	see annex on next page
Intended purpose	LARYVOX® INSERTER devices are used for the anterograde placement of a voice prosthesis when replacing the voice prosthesis.
Conformity assessment procedure	Medical devices listed in the annex conform to "general safety and performance requirements" according to EC regulation 2017/745, Annex I. The conformity of medical devices is declared based on technical documentation according to EC regulation 2017/745, Annex II and III. The manufacturer assumes sole responsibility for compliance with the requirements of EC regulation 2017/745 and all other Union legislation applicable to the medical devices listed in the annex below.

This Declaration of Conformity applies to all devices supplied after the production date until further notice. In the event of changes for the conformity assessment procedure, device alterations and changes to normative or regulatory requirements, conformity will be re-evaluated and declared again.

Cologne, 15.01.2025

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Annex to EC Declaration of Conformity: List of Medical Devices

REF	Medical Device
X25900	LARYVOX® INSERTER
X25910	LARYVOX® INSERTER EXPERT
X25915	LARYVOX® INSERTER EXPERT SMALL
X25920	LARYVOX® INSERTER SPECIAL
X25925	LARYVOX® INSERTER UNIEK
X25930	LARYVOX® INSERTER UNIEK PRO