

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	HUMIDO
Basic UDI-DI	405194846150HUMIDOHMEO22U
Class acc. to EC regulation 2017/745 Annex VIII	Class IIa
Classification Rule	2
REFs / Medical Devices	see annex on next page
Intended purpose	HMEs serve to humidify the airways and warm the air one breathes. They reduce the formation of viscous secretions in the lungs and prevent the trachea from drying out. HMEs filter the breathing air, thereby reducing intrusion of particles into the patient's airways. The use of an HME creates the respiratory resistance that is necessary to maintain physiological lung function. Variants with an O ₂ connection also allow for the supply of additional oxygen.
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 according to EC regulation 2017/745, Annex IX. Certificate No.: 1320GB448251015 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.

The manufacturer is subject to supervision by the notified body:
DNV MEDCERT GmbH (CE 0482), Pilatuspool 2, 20355 Hamburg, Germany

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, 15.10.2025

Andreas Fahl
Medizintechnik-Vertrieb GmbH
August-Horch-Straße 4a
51149 Köln, Germany
Christoph Bernads

Person Responsible for Regulatory Compliance

Annex to EC Declaration of Conformity: List of Medical Devices

REF	Medical Device
46150	HUMIDOSTOM® O2
46155	HUMIDOSTOM® O2 ISO
46450	HUMIDOTRACH®
46452	HUMIDOTRACH® sterile
46453	HUMIDOTRACH® PRO sterile
46454	HUMIDOTRACH® PRO ISO sterile
46456	HUMIDOTRACH® PRO
46457	HUMIDOTRACH® PRO ISO
46470	HUMIDUAL®