

EC Declaration of Conformity

Manufacturer:

DeVilbiss Healthcare LLC
100 DeVilbiss Drive
Somerset, PA 15501, USA

EC Authorized Representative:

DeVilbiss Healthcare GmbH
Kamenzerstraße 3, 68309
Mannheim, Germany

1. Oxygen Concentrators (UMDNS 12-873)

Catalogue nos.: 1025KS, 1025UK
Classification (MDD Annex IX): IIa (Rule 11)
Conformity Assessment Procedure: MDD 93/42/EEC, Annex II excluding Section 4

2. Accessories:

Product Description (Catalogue no.):

Caster, 4 pkg.-DFT	501DZ-603
Caster, Locking, 2 pk.	525DS-603
EXT Life Intake Bacteria Filter-DFT	MC44D-605
Cabinet Air Filter	303DZ-605
Intake Bacteria Filter	1025D-605
Compressor Filter	1025D-682
230V Compressor SSP	1025K-625
Final Bacteria Filter	PV5LD-651
Oxygen outlet connector (plastic, 1/pack)	CN100
Transfiller Caddy	525DD-650

Applied standards: All applicable harmonized standards as adopted under the EC Directive 93/42/EEC and published in the Official Journal of the European Communities (See attached listing).

This Declaration of Conformity is based on the EC Directive 93/42/EEC, Annex II.3 and supported by the TÜV NORD CERT GmbH (0044) certificate, with reference to articles 1 and 3 of the directive 93/42/EEC.

Notified Body:

TÜV NORD CERT GmbH
Langemarckstrasse 20, 45141 Essen, Germany

Identification No.:

0044

EC Certificate No.:

44 232 117803

Start of EC Marking:

2017-11-15

We, DeVilbiss Healthcare LLC., hereby declare under our sole responsibility that the above-mentioned products comply with the essential requirements and provisions of Council Directive 93/42/EEC.

Validity of this Declaration:

2019-08-07 – 2024-05-26

Somerset, PA,

Dec. 21, 2020

Place, Date


Tejash Naik

Senior Director, Regulatory Affairs and Quality Assurance
Name and Position

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Applied standards:

1025KS / 1025UK
<i>AAMI / ANSI ES60601-1:2005/(R) 2012 Ed 3.1 and A1:2012, C1:2009/(R) 2012 and A2:2010/(R) 2012 (Consolidated Text) Medical Electrical Equipment—Part 1: General Requirements for Basic Safety and Essential Performance (IEC 60601-1:2005, Mod). (General I (QS/RM))</i>
<i>AAMI / ANSI / IEC 60601–1–2:2014, Ed. 4.0, Medical Electrical Equipment—Part 1-2: General Requirements for Basic Safety And Essential Performance – Collateral Standard; Electromagnetic Disturbances - Requirements and Tests (associated with IEC 60601-1 Ed. 3.0)</i>
<i>IEC 60601-1-8 Ed. 2.1 b.2012 Medical electrical equipment - Part 1-8: General requirements for basic safety and essential performance - Collateral Standard: General requirements, tests and guidance for alarm systems in medical electrical equipment and medical electrical systems (associated with IEC 60601-1 3rd Edition)</i>
<i>IEC 60601-1-11:2010 Medical electrical equipment - Part 1-11: General requirements for basic safety and essential performance - Collateral Standard: Requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment (associated with IEC 60601-1 3rd Edition, referenced by ISO 80601-2-69:2014)</i>
<i>AAMI / ANSI / IEC 62304:2006, Medical Device Software – Software Life Cycle Process (Software/Informatics)</i>
<i>ISO 80601-2-69:2014 Medical electrical equipment – Part 2-69: Particular requirements for basic safety and essential performance of oxygen concentrator equipment</i>
<i>ISTA Procedure 6A FedEx Packaged-Products for Parcel Delivery System Shipments 70kg (150 lb) or Less (standard)</i>
<i>AMMI / ANSI / ISO 10993-1:2009, Biological Evaluation Of Medical Devices – Part 1: Evaluation And Testing Within A Risk Management Process (Biocompatibility)</i>