

# 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. Suction Units (UMDNS 13-846):**

Catalogue nos.: 7325D-AP, 7325D-D, 7325D-D-EXF, 7325D-I, 7325D-LA, 7325D-U,  
7325P-AP, 7325P-D, 7325P-D-EXF, 7325P-I, 7325P-LA, 7325P-T, 7325P-U  
Classification (MDD Annex IX): Ila (Rule 11)  
Conformity Assessment Procedure: MDD 93/42/EEC, Annex II excluding Section 4

**2. Accessories:**

Product Description (Catalogue no.):

Universal Suction Tubing and Filter Kit	18600-KITN
6' (1.8m) Patient Tubing, PVC	6305D-611
10" (25.4cm) Connection Tubing, PVC	SUCP TUBING 10
DC Power Cord, 1 each	7304D-619
800 ml Disposable Container w/ internal filter cartridge, splash guard, 4 3/8" tubing, 48 each	7305D-632
Collection Container Kit (internal filter cartridge, splash guard, 800 ml container, 4 3/8" and 6" tubing package)	7305D-633
Filter Cartridge, 12 pk. (for disposable container use)	7305D-635
4.5" (11.4cm) Connection Tubing, silicon	7305D-639
Collection Container Kit (1200 ml reusable container, external bacteria filter, elbow, 4 3/8" tubing)	7314D-603
1200 ml Reusable Container (external bacteria filter, elbow, 4 3/8" tubing) 6 pk.	7314D-604
AC to DC Power Adapter/Charger	7314P-613
External Bacteria Filter (non-sterile), 12 pk. (for reusable container use)	7305D-608
Battery, replacement assy.	7325P-614
Carry Case	7325D-635
Power Cord, US	DV51D-606
Power Cord, EU	DV51D-607
Power Cord, UK	DV51D-608
Power Cord, AU	DV51D-609

**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



# EC Declaration of Conformity

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, June 14, 2022

Place, Date

Roberto Munoz Director, Regulatory Affairs and Audit  
Name and Position

**Applied Standards:**

<b>7325 series</b>
BS EN ISO 10079-1:2015 + AMD 1:2019 (Ed 3.0) - Medical Suction Equipment
IEC 60601-1:2005+A1:2012 (Ed 3.0) - Medical Electrical Equipment—Part 1: General Requirements for Safety; Safety Requirements for Medical Electrical Systems [FDA Recognized Consensus Standard]
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 compatibility – Requirements and tests (FDA Recognition Number 19-8)
IEC 60601-1-6:2010 + AMD 1:2013 (Ed. 3.1) Medical electrical equipment - Part 1-6: General requirements for basic safety and essential performance - Collateral standard: Usability (associated with IEC 60601-1 Ed. 3.0) (FDA Recognition Number 5-89)
IEC 60601-1-9:2007 + A1:2013 (Ed. 1.1) Medical electrical equipment - Part 1-9: General requirements for basic safety and essential performance - Collateral standard: Requirements for environmentally conscious design (associated with IEC 60601-1 Ed. 3.0)
IEC 60601-1-11:2010 (Ed 1.0), 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 (FDA Recognition Number 19-14)
IEC 62366-1:2015 (Ed. 1.0) – Medical devices - Application of usability engineering to medical devices (FDA Recognition Number 5-114)
ISO 14971:2019 (Third Ed), Medical devices - Application of risk management to medical devices [FDA Recognized Consensus Standard Number 5-125]
ISTA Procedure 3A: Packaged-Products for Parcel Delivery System Shipments 70kg (150 lb) or less (standard).