ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Product Service

EC Certificate

Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V (Devices in Class IIa, IIb or III)

No. G2 17 12 31998 026

Digitally signed by Rotari Vladimir
Date: 2022.06.18 13:26:55 EEST
Reason: MoldSign Signature
Location: Moldova

Manufacturer:

DEAS s.r.l.
Via dell'Industria 49
48014 Castel Bolognese (RA)
ITALY

Facility(ies):

DEAS s.r.l.
Via dell'Industria 49, 48014 Castel Bolognese (RA), ITALY

Product Category(ies):

Sterile and non sterile medical devices: breathing circuits and connectors, humidification chambers, catheters mount, masks, filters, spirometry filters and mouthpieces, bags and manual breathing units, monitor lines, extraglottic, tracheal, tracheal-bronchial, tracheostomy tubes, introducers, tracheostomy kits, aerosol and oxygen therapy devices, PEEP valves, suction devices

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

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Valid from: 2018-03-21
Valid until: 2023-03-20

Date, 2018-02-19

Stefan Preiß

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123