Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 016389 0017 Rev. 01

Manufacturer:
VBM Medizintechnik GmbH
Einsteinstr. 1
72172 Sulz a. N.
GERMANY

Facility(ies):
VBM Medizintechnik GmbH
Einsteinstr. 1, 72172 Sulz a. N., GERMANY

Product Category(ies):
Tourniquet Systems and Pressure Infusors with cuffs, Medical Devices for Trans Tracheal Ventilations, Laryngeal Tubes, Respiration Sets, Respiration accessories (except class I), Rectal Tampon by Roche (RTR)

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: 713137731_1

Valid from: 2018-10-21
Valid until: 2023-10-20

Date, 2018-10-18

Digitally signed by Rotari Vladimir
Date: 2023.06.20 20:44:52 EEST
Reason: MoldSign Signature
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Ștefan Preiß

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

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