



Benannt durch/Designated by  
 Zentralstelle der Länder  
 für Gesundheitsschutz  
 bei Arzneimitteln und  
 Medizinprodukten  
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 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.

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**Valid until:** 2023-10-20

**Date,** 2018-10-18

Stefan Preis

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 ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ CERTIFICADO ◆ CERTIFICAT