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

Product Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex VI
(Devices in Class I with measuring function)

No. G3M 020267 0020 Rev. 00

Manufacturer: **Friedrich Bosch GmbH & Co. KG**
Hohenlaienstr. 30
72406 Bisingen
GERMANY

Facility(ies): Friedrich Bosch GmbH & Co. KG
Hohenlaienstr. 30, 72406 Bisingen, GERMANY

Product Category(ies): **Mechanical Sphygmomanometers and Pressure Infusion Cuffs**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for final inspection and testing of the respective devices / device categories in accordance with MDD Annex VI. This quality assurance system covers those aspects of manufacture concerned with the metrological requirements of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report no.: 713157603

Valid from: 2019-06-25
Valid until: 2024-05-26

Date, 2019-06-25

Stefan Preiß
Head of Certification/Notified Body

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT
TÜV SÜD
A4 / 07.17