

## 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 15 11 20267 018

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 test 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 .:

713069052

Valid from: Valid until: 2015-11-16 2020-11-15



Date, 2015-09-29

Hans-Heiner Junker

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

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