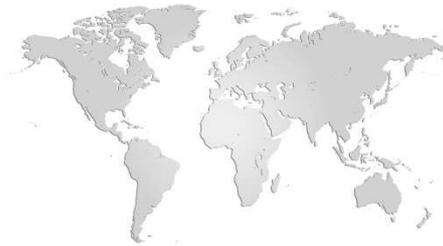


# EC CERTIFICATE

## for the Quality Assurance System



### according the Directive 93/42/EEC, Annex V

As a Notified Body of the European Union, DEKRA Certification GmbH certifies, that the company  
**Bosch + Sohn GmbH & Co. KG**

Bahnhofstraße 64, 72417 Jungingen, Germany

**Certified location:**

Bahnhofstraße 64, 72417 Jungingen, Germany

applies a quality assurance system according to the Directive 93/42/EEC Annex V for the medical devices listed in the annex. The approval is based on the result of the re-certification audit report no. 50539-Z7-00, the decision dated 2020-04-23 and is only valid in connection with the successful performance of the annual surveillance audits.

This certificate is valid from 2020-04-29 to 2024-05-26

Registration No.: 50539-17-05



Ruth Delbeck-Bayer  
DEKRA Certification GmbH Stuttgart; 2020-04-23  
Notified Body ID-number: 0124



Benannt durch/Designated by  
Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln und  
Medizinprodukten  
**ZLG-BS-295.10.02**

# Annex to the EC Certificate No. 50539-17-05

Valid from 2020-04-29 to 2024-05-26

Revision status of the annex: 0 dated 2020-04-29

Devices/device categories included in the certificate:

## Class I m:

For the products listed below, the review of the Quality System refers exclusively to the aspects of manufacture concerned with the conformity of the products with the metrological requirements.

- Non active medical devices with measuring function
  - non invasive aneroid sphygmomanometers

## Class II a:

- Devices for monitoring vital physiological parameters
  - Noninvasive blood pressure units and systems
  - Thermometer, electronic
  - Thermometer, infrared
  - 24-hour ambulatory blood pressure monitor
- Evaluation software for blood pressure units and systems



Ruth Delbeck-Bayer  
DEKRA Certification GmbH, Stuttgart, 2020-04-23  
Notified Body ID-number: 0124

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