

# CERTIFICATE

for the  
Quality Assurance System



As a notified body of the European Union (Reg. No. 0124), DEKRA Certification GmbH hereby approves the Quality Assurance System applied for final inspection by the company

**Bosch + Sohn GmbH u. Co. KG**  
Bahnhofstraße 64 • 72417 Jungingen, Germany

Approval is based on the decision dated 23.04.2009 and the result of the report no. 50539-Z4-00 and is performed in accordance with the stipulations of

## Annex VI, Section 3 of the Directive 93/42/EEC

of the Council dated June 14, 1993 governing medical devices. The certification is applicable to the devices specified in the Annex. The devices in question are subjected to testing and examination in accordance with Annex VI, Section 3 of the Directive 93/42/EEC. The listed devices may be affixed with the CE marking indicated below.



Date of the first certification: 13.01.1995

Date of the last recertification: 29.04.2009

This certificate is valid until: 28.04.2014

Certificate-registration No.: 50539-18-02  
English version

DEKRA Certification GmbH  
Stuttgart, den 23.04.2009



Akkreditiert durch  
Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln  
und Medizinprodukten  
ZLG-ZQ-992.94.16

# Annex to the Certificate 50539-18-02 dated 23.04.2009

English version

Revision status: 0

Date: 29.04.2009

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## Devices/device categories included in the certificate

### Class II a:

- 24-hour ambulatory blood pressure monitor: boson TM-2430
- non invasive blood pressure units, electronic
- Communication aid, speak box
- thermometer, electronic
- thermometer, infrared, skin
- thermometer, infrared, ear



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