



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

Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V

(Devices in Class IIa, IIb or III)

No. G2 016512 0046 Rev. 01

Manufacturer:

NIHON SEIMITSU SOKKI CO., LTD.

2508-13 Nakago, Shibukawa

Gunma

377-0293 JAPAN

**Product
Category(ies):**

**Non-invasive Digital Blood Pressure Monitors,
Oximeters Pulse, Non-contact Thermometer**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.:

JAQ235040622

Valid from:

2020-05-14

Valid until:

2024-05-26

Date, 2020-05-14

Christoph Dicks

Head of Certification/Notified Body