EC CERTIFICATE

for the Quality Assurance System

according the Directive 93/42/EEC, Annex II excluding section (4)

As a Notified Body of the European Union, DEKRA Certification GmbH certifies, that the company

Rudolf Riester GmbH

Bruckstraße 31, 72417 Jungingen, Germany

Certified location:

Bruckstraße 31, 72417 Jungingen, Germany

applies a quality assurance system according to the Directive 93/42/EEC Annex II for the medical devices listed in the annex. The approval is based on the result of the re-certification audit report no. 50828-Z5-00, the decision dated 2019-11-11 and is only valid in connection with the successful performance of the annual surveillance audits.

This certificate is valid from 2019-11-14 to 2024-05-26

Registration No.: 50828-16-06

DEKRA Certification GmbH Stuttgart, Handwerkstraße 15, D-70565 Stuttgart

Ruth Delbeck-Bayer
DEKRA Certification GmbH Stuttgart; 2019-11-11
Notified Body ID-number: 0124

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-295.10.02

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Annex to the EC Certificate No. 50828-16-06

Valid from 2019-11-14 to 2024-05-26

Revision status of the annex: 0 dated 2019-11-14

Devices/device categories included in the certificate:

Class II b:

Measuring device indicating the oxygen saturation of pulse and non-invasive blood pressure

- ri-vital®

Vital Signs Monitoring

- RVS-100

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Ruth Delbeck-Bayer
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