EC Certificate

For the Quality Assurance System according the directive 93/42/EEC, Annex II excluding section (4)

DEKRA Certification

As a notified body of the European Union, DEKRA Certification GmbH certifies, that the company

Rudolf Riester GmbH

Bruckstraße 31 • 72417 Jungingen, Germany

applies a quality assurance system for the medical devices listed in the annex according to the directive 93/42/EEC annex II. The approval is based on the result of the re-certification audit report no. 50828-Z3-00, the decision dated 07.11.2011 and is only valid in connection with the successful performance of the annual surveillance audits.

Date of the first certification:	24.10.1997	Date of the last recertification:	14.11.2011
This certificate is valid until:	13.11.2016	Certificates registration No.:	50828-16-04 English version

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

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

Stuttgart, 07.11.2011
DEKRA Certification GmbH
Handwerkstraße 15, 70565 Stuttgart, Germany
Notified Body ID-number: 0124

Annex to the EC Certificate 50828-16-04 dated 07.11.2011

English version

Revision status: 0 Date: 14.11.2011 Page 1 of 1

Devices/device categories included in the certificate

Non-invasive sphygmomanometers and equipment

Class II b:

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

- ri-vital®

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