EC Certificate

For the Quality Assurance System according the directive 93/42/EEC, Annex VI

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 VI. The approval is based on the result of the re-certification audit report no. 50828-Z3-00, the decision dated 07.11.2011 is only valid in connection with the successful performance of the annual surveillance audits.

Date of the first certification:	21.02.1995	Date of the last recertification:	14.11.2011
This certificate is valid until:	13.11.2016	Certificates registration No.:	50828-18-04

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 50828-18-04 dated 07.11.2011

English version

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Devices/device categories included in the certificate

Class I devices with measurement function

For the products listed below, the review of the Quality System refers exclusively to the manufacturing steps associated with product conformity and metrological requirements.

Aneroid sphygmomanometers:	minimus®
	exacta®
	precisa® N
	sphygmotensiophone
	babyphon®
	big ben®
	ri-med®
	ri-mega®
	ri-san®
	sanaphon®
	R1 shock-proof
	e-mega®
Mercury sphygmomanometers:	diplomat-presameter®
	nova-presameter®
	empire® N
	global de luxe®
Eye tonometer:	schiötz

Class II a:

Digital sphygmomanometers:	ri-champion® N
Infrared-thermometer:	ri-thermo® N
Digital-thermometer:	ri-gital®
Pulsoxymeter:	ri-fox N
ENT-Endoscopy:	Sinuscopes