ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 14 10 30178 012

Manufacturer:

RECO Medizintechnik
Wolfgang Rentsch e.K.
Altjessen 2
01796 Pirna
GERMANY

Facility(ies):

RECO Medizintechnik Wolfgang Rentsch e.K.
Altjessen 2, 01796 Pirna, GERMANY

Product Category(ies):

Equipment for Diathermy, Microwave Therapy, Uroflow Recorders

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: 713050056

Valid from: 2014-10-20
Valid until: 2019-10-13

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Date, 2014-10-23

Hans-Heiner Junker

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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