Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

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 023228 0013 Rev. 00

Manufacturer: **Dr. Hönle Medizintechnik GmbH**
Dornierstr. 4
82205 Gilching
GERMANY

Facility(ies): Dr. Hönle Medizintechnik GmbH
Thura Mark 8+10, 06780 Zörbig, GERMANY

Product Category(ies): **Light therapy devices and tap-water iontophoresis units including accessories for treatment of hyperhidrosis**

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.: 713136709

Valid from: 2019-01-31
Valid until: 2023-07-23

Date, 2019-01-31

Stefan Preiß

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

A4 / 07.17

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

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