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 032017 0012 Rev. 01

Manufacturer:

gbo Medizintechnik AG
Kleiststrasse 6
64668 Rimbach
GERMANY

Facility(ies):

gbo Medizintechnik AG
Kleiststrasse 6, 64668 Rimbach, GERMANY

Product Category(ies):

Stimulators for nerve and muscle, Ultrasound therapy devices, suction application aids, microwave therapy devices, shortwave therapy devices, cryo therapy devices, traction therapy devices, magnetic field therapy devices

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.: 713160427

Valid from: 2020-01-07
Valid until: 2024-05-25

Date, 2020-01-07

Christoph Dicks
Head of Certification/Notified Body

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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TÜV SÜD

A4 / 07.17