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

TÜV SÜD Product Service

EC Certificate

Product Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex VI
(Devices in class IIa or IIb)

No. G3 035504 0011 Rev. 00

Manufacturer:
MaRhyThe - Systems GmbH & Co. KG
Industrie Str. 29
82194 Gröbenzell/München
GERMANY

Facility(ies):
MaRhyThe - Systems GmbH & Co. KG
Industrie Str. 29, 82194 Gröbenzell/München, GERMANY

Product Category(ies):
Active therapeutic devices in the area of the Matrix-Rhythm-Therapy

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

Report no.: 713156270

Valid from: 2019-06-24
Valid until: 2024-05-22

Date, 2019-06-24

Stefan Preiß
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