



Product Service

EC Certificate

Production Quality Assurance System

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

No. G3 14 04 35504 007

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 test 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.: 713039919

Valid from: 2014-05-23
Valid until: 2019-05-22

Date, 2014-05-26

Hans-Heiner Junker



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

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