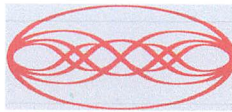


MaRhyThe



Matrix-Rhythmus-Therapie

## EC-Declaration of Conformity

**CE** 0123 marking

We, Manufacturer

**MaRhyThe®-Systems GmbH & Co. KG**

Industriestrasse 29

82194 Gröbenzell / München

Germany

Declare under our sole responsibility, that the product

Matrixmobil®

MaRhyThe® Applikator & Steuergerät

(Medical device class IIa (MDD93/42/EC, Annex IX))

Lot. No (Reference- and Serial Numbers) G1/A1 - 10.001 bis 10.600 und A2 - 01.001 bis 01.100

is in conformity with

Council Directive 93/42/EEC (MDD) as amended by 2007/47/EC

The product fulfills the essential requirements of Annex I of the MDD

The conformity assessment procedure was performed according to Annex VI

The product is classified as an class IIa product acc. to rule 9 of annex IX MDD 93/42 EEC as amended by 2007/47/EC

The product is manufactured in the EC.

Date: 20.08.2018

Signature

Managing Director

Ref. No.: G3 14 04 35504 007

Date: May 23<sup>rd</sup> 2014

Notified Body:  
**TÜV Süd Product Service GmbH**  
Ridlerstr. 65,  
80339 München

Notified by: **ZLG**  
Registration-Nr:  
ZLG-BS-244.10.08