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- an Serial Numbers) G1/A1 – 11.001 bis 11.500 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: 15.02.2024

Signature _____
Managing Director

Ref. No.: G3 035504 0011 Rev.00
Date: June 24th 2019

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

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