

EU Declaration of Conformity (DoC)

in accordance with Regulation (EU) 2017/745 on medical devices (MDR)
in application of the transitional provisions MDR Art. 120 (3a)



We, the company

emotion fitness GmbH & Co KG

Trippstadter Str. 68

67691 Hochspeyer

SRN: DE-MF-000016584

declare under our sole responsibility
that the following listed and in the technical documentation "motion cardio line 900 & 100 med" as
of April 26th, 2024 described and specified

Cardiovascular and Muscular Training Devices

Product name	REF
motion cycle 900 med	F-MED-MC-900
motion body 900 med	F-MED-MO-900 F-MED-MO-901
motion relax 900 med	F-MED-MR-900
motion cross 900 med	F-MED-CR-900
motion stair 900 med	F-MED-MS-900
motion cycle 100 med	F-MED-MC-100

for carrying out physiotherapeutic training and generating reproducible loads during cardiological
diagnostics

comply with the Essential Requirements of
EC Directive 93/42/EEC on medical devices (MDD), Annex I

Basic UDI-DI

426054547935AD

Medical device class:

(according to MDD, Annex IX)

Ila

(Rule 10)

**applied conformity assessment
procedure:**

MDD, Annex VI and Annex VII

Notified body

Berlin Cert – Prüf- und Zertifizierstelle für
Medizinprodukte GmbH
Dovestrasse 6
10587 Berlin
Notified Body No.: 0633

Certificate number

Z-19-065-S-R VI-E-N1

Place, date

Michael Schmitt, PRRC

The declaration of conformity is valid until 31.12.2028 at the latest
Product changes or changes on the technical documentation requires a reissue of this DoC.