

EU
declaration of conformity following directive 93/42 EEC

We

emotion fitness GmbH & Co. KG

(Name of producer)

Trippstadter Str. 68, D-67691 Hochspeyer, Germany

(Address)

declare under our sole responsibility that the product

cardiovascular and muscular training device

motion cycle 900 med item number F-MED-MC-900

from the serial number 9MER000000121 (or 9MEB0000000121) up to the machines build until May 26th, 2024;

motion body 900 med item number F-MED-MO-900 and

motion body 900 med (wall mounted version) item number F-MED-MO-901

from the serial number 9MUR000000121 (or 9MUB0000000121) up to the machines build until May 26th, 2024;

motion relax 900 med item number F-MED-MR-900

from the serial number 9MRR000000121 (or 9MRB0000000121) up to the machines build until May 26th, 2024;

motion cross 900 med item number F-MED-CR-900

from the serial number 9MCR000000121 (or 9MCB0000000121) up to the machines build until May 26th, 2024;

motion stair 900 med item number F-MED-MS-900

from the serial number 9MSR000000121 up to the machines build until May 26th, 2024;

(name of type or model, batch-, or serial number)

do meet the requirements, following the provisions of the medical device directive 93/42 EEC annex VI and VII (class IIa).

Additional specifications with the used standards

DIN EN ISO 20957-1 2014, DIN EN ISO 20957-5 2015 (cycle, body, relax), DIN EN ISO 20957-9 2015 (cross), DIN EN ISO 20957-8 (stair)

DIN EN 60601-1-2 2015, DIN EN 60601-1 2013, DIN EN 62366 2008

(name and/or numbers, as well as the date)

Involved notified body at this procedure of conformity CE 0633
Berlin Cert - Prüf- und Zertifizierstelle für Medizinprodukte GmbH
an der Technischen Universität Berlin
Dovestraße 6
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Managing Director

Hochspeyer, May 20th, 2021