MOTOmed®
move differently

EC-Declaration of Conformity on Medical Devices

According to annex II of Medical Device Directive 93/42/EEC (2007/47/EG)

Manufacturer: **RECK-Technik GmbH & Co. KG** Engineering Division Sector
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Name of Product:
MOTOmed viva2, Item no. 200.003
MOTOmed viva2 ATAP, Item no. 200.003 + 250.000
MOTOmed viva2 stativ, Item no. 200.012
MOTOmed viva2 light, Item no. 200.004
MOTOmed viva2 light ATAP, Item no. 200.004 + 250.000
MOTOmed viva2 light stativ, Item no. 200.021
MOTOmed viva2 Parkinson, Item no. 200.008
MOTOmed viva2 Parkinson ATAP, Item no. 200.008 + 251.000
MOTOmed letto2, Item no. 279.003
MOTOmed letto2, Item no. 279.008
MOTOmed letto2, Item no. 279.024
MOTOmed letto2, Item no. 279.016
MOTOmed gracile 12, Item no. 594.003
MOTOmed gracile 12 ATAP, Item no. 594.003 + 599.000
MOTOmed muvi, Item no. 300.000

Product options: all (according to current pricelist)

Product classification: IIa (rule 9, Medical Device Directive 93/42/EEC)

The manufacturer hereby declares under his sole responsibility that the products identified above are in conformity with the MDD 93/42/EEC (2007/47/EC) concerning medical devices, annex II, paragraph 3.

This Declaration of Conformity is valid until 02.08.2023 or until a revised declaration is made due to product modifications.

Following party was involved in the evaluation of the conformity:
DEKRA Certification GmbH
Handwerkstrasse 15, 70565 Stuttgart, GERMANY
Notified Body No. 0124

Betzenweiler, 03.08.2018

Andreas Reck Executive Director

RECK
Medizintechnik