EC-Declaration of Conformity on Medical Devices


Manufacturer: RECK-Technik GmbH & Co. KG Engineering Division Sector
Reckstraße 1–5, 88422 Betzenweiler, GERMANY
phone +49 7374 18-85, fax +49 7374 18-480
email: contact@motomed.com

Name of Product: MOTOmed muvi, Item no. 300.000

Product options: all (according to current pricelist)

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

We hereby declare under our sole responsibility that the above products comply with the relevant provisions of the Medical Devices Directive 93/42 / EEC (2007), Annex II, Section 3 and that we are solely responsible for issuing this declaration of conformity.

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