MOTOmed®

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 muvi 300.000**
Basic UDI-DI: 4260193710057V

REF	SN
300.000	xxx-VM3-000xxx

Product options: all (according to current pricelist)

Product classification: IIa (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 (registration no. of certificate: 50293-16-05) 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