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 Reckstraße 1–5, 88422 Betzenweiler, GERMANY phone +49 7374 18-85, fax +49 7374 18-480 email: contact@motomed.com
Name of Product:	**MOTOmed loop**, Item no. 260.xxx
Product variants:	loop l, loop a, loop la, loop p l, loop p a, loop p la, loop kidz l, loop kidz a, loop kidz la, loop light l, loop light a, loop light la
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 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, 24.01.2019

Andreas Reck Executive Director

RECK
Medizintechnik