

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