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 layson edition 261.xxx**
Basic UDI-DI: 4260193710037R

		REF	SN
Product variants:	MOTOmed layson.la	261.130	xxx-LY-130xxx
	MOTOmed layson.la	261.030	xxx-LY-030xxx
	MOTOmed layson.l	261.110	xxx-LY-110xxx
	MOTOmed layson.l	261.010	xxx-LY-010xxx
	MOTOmed layson kidz.la	261.330	xxx-LY-330xxx
	MOTOmed layson kidz.la	261.230	xxx-LY-230xxx
	MOTOmed layson kidz.l	261.310	xxx-LY-310xxx
	MOTOmed layson kidz.l	261.210	xxx-LY-210xxx
	MOTOmed layson.l dia	261.119	xxx-LY-119xxx
	MOTOmed layson.la prof	261.139	xxx-LY-139xxx

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, 20.07.2020 **Andreas Reck** Executive Director