MOTOmed®

[EN] EU Declaration of Conformity for Medical Devices

in accordance with Annex IV of the EU MDR 2017/745 Regulation

[1]	Manufacturer:	**RECK-Technik GmbH & Co. KG** Reckstraße 1–5, 88422 Betzenweiler, GERMANY Tel. 07374 18-84, Fax 0 73 74 18-80 SRN: DE-MF-000005730 E-Mail: vigilance@motomed.com
[2]	Product name	**MOTOmed loop** REF **260.xxx** UDI Basic UDI-DI: 4260193710027P
[3]	Intended purpose:	The MOTOmed loop is intended exclusively for passive, assistive and active movement of the lower and upper extremities of seated persons. During use, the MOTOmed loop can be controlled via an operating panel. The MOTOmed loop is mobile and can therefore be used at different locations. The MOTOmed loop helps wheelchair users and people with limited mobility maintain the necessary daily movement through passive and active movement of the extremities. Treatment goal: to prevent, reduce, and improve damage (and subsequent damage) caused by loss of mobility or lack of exercise.
[4]	Product variants:	

	REF	SN
MOTOmed loop.l	260.010	xxx-LP-010xxx
MOTOmed loop.a	260.020	xxx-LP-020xxx
MOTOmed loop.la	260.030	xxx-LP-030xxx
MOTOmed loop.la prof	260.039	xxx-LP-039xxx
MOTOmed loop p.l	260.040	xxx-LP-040xxx
MOTOmed loop p.la	260.060	xxx-LP-060xxx
MOTOmed loop light.l	260.100	xxx-LP-100xxx
MOTOmed loop light.a	260.110	xxx-LP-110xxx
MOTOmed loop light.la	260.120	xxx-LP-120xxx
MOTOmed loop kidz.l	260.070	xxx-LP-070xxx
MOTOmed loop kidz.a	260.080	xxx-LP-080xxx
MOTOmed loop kidz.la	260.090	xxx-LP-090xxx

[5]	Classification of the risk class:	IIa, Annex VIII, Rule 9
[6]	Applicable common specifications	-
[7]	Other EU legislations met:	REACH Regulation (EC) No. 1907/2006 RoHS Directive 2011/65/EU Machinery Directive 2006/42/EC
[8]	Conformity assessment procedure according to MDR (EU) 2017/745	Conformity assessment according to Annex IX Chapter I
[9]	Certificate number	50293-60-00-00
[10]	Valid from: Date of Expiry	20.08.2025 12.02.2028

We hereby assure under our sole responsibility that the above-mentioned products comply with the relevant provisions of the EU MDR 2017/745 Regulation for Medical Devices and, if applicable, other relevant legal provisions of the European Union.

This declaration of conformity is valid until 12.02.2028 (certificate number: 50293-60-00-00) or until a revised declaration of conformity is issued after the product has been modified.

The following notified body was involved in the conformity assessment procedure: DEKRA Certification GmbH, Handwerkstrasse 15, 70565 Stuttgart, Germany, ID number 0124.

Betzenweiler, 21.08.2025

Christine Reck
CEO

i.A. Andreas Brauch
PRRC

RECK-Technik GmbH & Co. KG
Reckstraße 1-5, 88422 Betzenweiler
RECK