

EU Certificate

for the assessment of the
quality management system



according to Medical Device Regulation (EU) 2017/745 Annex IX Chapter I+III

As a Notified Body of the European Union DEKRA Certification GmbH certifies, that the
manufacturer

RECK-Technik GmbH & Co. KG

Single Registration Number (SRN): DE-MF-000005730

Reckstraße 1-5, 88422 Betzenweiler, Germany

applies a quality management system according to Annex IX Chapter I+III of the Medical Device Regulation (EU) 2017/745 for the medical devices listed in the annex. This certificate is based on the assessments listed in CNo50293-00 and is only valid in conjunction with the successful completion of the annual surveillance audits.

EU Certificate no.: 50293-60-00-01

Certificate valid from:

2026-01-20

Certificate valid to:

2028-02-12

Previous certificate no. 50293-60-00-00, issued on 2025-08-20



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de

BS-MDR-092

DEKRA Certification GmbH, Stuttgart 2026-01-20
Notified Body ID number: 0124

Annex to the EU Certificate no. 50293-60-00-01

Following devices/device categories are included in this certificate:

Class IIa

- Movement therapy devices
 - MOTOmed loop
Basis-UDI-DI: 4260193710027P
 - MOTOmed layson
Basis-UDI-DI: 4260193710037R
 - MOTOmed gracile
Basis-UDI-DI: 4260193710588J
 - MOTOmed muvi
Basis-UDI-DI: 4260193710057V

Change(s) to previous certificate: Product MOTOmed muvi has been added to the certificate.