CE Declaration of Conformity

Manufacturer:
Hankamp Rehab BV, Buurserstraat 198-200, 7544 RG Enschede; The Netherlands
SRN | NL-MF-000006108

In accordance with the following Regulation:

EU 2017/745	European Medical Device Regulation And its amending regulation

Hereby declare under sole responsibility that:

Equipment:	ReStep
Classification:	Class I (In accordance with Annex VIII, 2017/745)
Intended Purpose:	Ceiling-mounted track system for fall prevention
UDI-DI:	8 72089 2136442

Is in conformity with applicable requirements of the following standards:

Ref. No.	Title	Edition
NEN-EN-ISO 13485	Medical devices — Quality management systems — Requirements for regulatory purposes	2016
EN-ISO 20417	Medical devices — Information to be supplied by the manufacturer	2021
EN ISO 15223-1	Medical Devices – Symbols to be used with medical device labels, labelling and information to be supplied - Part 1 – General Requirements	2021
EN ISO 14971	Medical Devices – Application of risk management to medical devices	2019

And is in conformity with the following Common Specifications: -

I hereby declare that the equipment named above has been designed to comply with the relevant sections of the above reference specifications. The unit complies with all applicable Essential Requirements of the Regulation.

Signed (date): 01-05-2026

Name:	Frederik Tönis, CEO
Place of Issuance:	Hankamp Rehab BV, Enschede, the Netherlands
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Conformity assessment procedure: Annex IV of EU 2017/745 considering the amendment by the regulation: EU 2020/561