



Declaration of Conformity

Manufacturer: COBI REHAB A/S
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The undersigned declares and takes the full responsibility that the product named and listed on this certificate is constructed and manufactured in accordance with the essential requirements of The Regulation (EU) 2017/745 of the European Parliament and of the Council of 5. April 2017 relating to Medical Devices Class 1. The product is suitable for reuse by other people after disinfection.

Risk analysis has been executed in accordance with EN ISO 14971:2012 Medical devices
- Application of risk management to medical devices.

The devices comply with the following standards:

EN 12182 - Assistive products for persons with disability. General requirements and test methods
ISO 11199-2:2005

Product name:	Rollator King
Item No.:	0172-050-000, 0172-071-000
Risk Class:	I
Intended use:	Assistive devices for use by individuals with mobility impairments
Basic UDI-DI:	5740000100007N8
SRN:	DK-MF-000001370

Karlslunde, 08-01-2025



Michael Kock, CEO