CE MD

COBI REHAB
The Positioning Company

Declaration of Conformity

Manufacturer:

COBI REHAB A/S
Silovej 16-18
DK-2690 Karlslunde, Danmark
Tel. +45 7025 2522
E-mail: cobi@cobi.dk

We hereby declare in sole and 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.

Risk analysis has been executed in accordance with EN ISO 14971:2019 Medical devices - Application of risk management to medical devices.

The devices comply to the following standards:

EN 12182 - Assistive products for persons with disability. General requirements and test methods
AS/NZS 3973:2009
AS/NZS 4688.2-2000
AS/NZS 4688.3-2000

Product name:	Vastus
Item No.:	0142-061-000, 0142-071-000
Risk Class:	I
Intended use:	Assistive devices for persons with special needs for a reinforced toilet or a toilet nearby or chair in a bath situation
Basic UDI-DI:	5740000100003MY
SRN:	DK-MF-000001370

Karlslunde, 08-01-2025

Michael Kock, CEO