Declaration of Conformity

Manufacturer:

COBI Rehab ApS
Fuglebækvej 1 D
DK-2770 Kastrup, Danmark
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The undersigned declares 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 14971:2012 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
EN 11199-2:2005

Product name:	Rollator Support
Item No.:	0173-050-000, 0173-071-000
Risk Class:	I
Intended use:	Assistive devices for persons with special needs
Basic UDI-DI:	5740000100007N8
SRN:	DK-MF-000001370

Kastrup, 17-02-2021

Claus Nymark, Director