EC DECLARATION of CONFORMITY

Regulation (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL on medical devices

We, **MOBILEX A/S**
Registered place of business
Grønlandsvej 5
8660 Skanderborg
Denmark

SRN: DK-MF-000021885

Hereby declare under our sole responsibility as a legal manufacturer that the product specified on the product list below, meet the essential health and safety requirements and is in conformance with the provisions of the Regulation (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL on medical devices.
The product is classified as Class I, medical device. The classification is based on the requirements of Rule 1 of annex VIII, of the Regulation (EU) 2017/745.

The CE marking has been affixed on the product according to Annex V of the Regulation (EU) 2017/745.

Intended purpose: To be used to assist patients in wheelchairs or scooters enable the passage of elevated points (e.g. Stairs).

PRODUCT LIST

Single-folded ramp:

REF / item no.	DF-180	DF-240	DF-300
UDI-DI	5740001415278	5740001415285	5740001415292
BASIC-UDI-DI	57400014DF-RAMPS6N		

ACCESSORIES LIST

Item nr.	Accessories item nr.
DF-180+240+300	NO

Harmonized norms used during conformity estimation:

EN 12182:2012, PN-EN ISO 14971:2012, EN 1041:2009

Skanderborg, 2022-04-12, Thomas N. Christensen, Managing Director

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