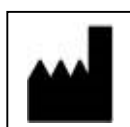




EC DECLARATION of CONFORMITY

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. | SF-040 | SF-090 | SF-120 | SF-150 |
|----------------|--------------------|-------------------|-------------------|-------------------|
| UDI-DI | 57400014 15230 | 57400014 15247 | 57400014 15254 | 57400014 15261 |
| BASIC-UDI-DI | 57400014SF-RAMPSFK | | | |

ACCESSORIES LIST

| Item nr. | Accessories item nr. |
|------------------------|----------------------|
| SF-045+060+090+120+150 | 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

Issued: 2022/04

