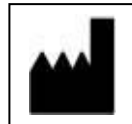




## EC DECLARATION of CONFORMITY

### 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 specified on the product list below is "technical aid for the disabled", 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.

#### **PRODUCT LIST**

Doorstep ramps:

REF / item no.	DS4025	DS4076	DS2176
UDI-DI	5740001415018	5740001415001	5740001415025
BASIC-UDI-DI	57400014DOORSTEPRAMPSZM		

Harmonized norms used during conformity estimation:

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

Skanderborg, 2022-03-30, Thomas N. Christensen, Managing Director

