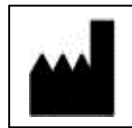




## EC DECLARATION of CONFORMITY

### 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**



Hereby declare under our sole responsibility as a legal manufacture 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

**Double-folded ramp, DF-Series DF-180, DF-240, DF-300**

Harmonized standards used during the validation process:  
PN EN ISO 14971:2012; EN 12182:2012; PN EN 1041:2001

Skanderborg, 2020-03-11, Thomas N. Christensen, Managing Director

