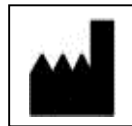




## EC DECLARATION of CONFORMITY

### 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~~ 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**

#### **Telescopic Ramps TR & LR-Series**

Item number: **TR-150, TR-210, TR-300, LR-150, LR-210**

#### **ACCESSORIES LIST**

Harmonized norms used during conformity estimation:

EN 12182:1999, PN-EN ISO 14971:2004, EN 1041:2001

Skanderborg, 2020-03-24, Thomas N. Christensen, Managing Director

