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

Mobilex Transfer board:

REF / item no.	240501	240502	240505	240507	240701
UDI-DI	5740001403459	5740001403466	5740001439014	5740001403473	5740001403480
BASIC-UDI-DI	57400014TRASFERBOARDMU				

PN-EN 12182:2012, PN-EN ISO14971:2012, PN-EN 1041:2009

Skanderborg, 2022-04-005, Thomas N. Christensen, Managing Director

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