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 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: designed to increase the mobility of persons who have difficulties standing or walking

PRODUCT LIST

Panther rollator:

REF / item no.	**318001**	**318002**
UDI-DI	**5740001403442**	**5740001403510**
BACIC-UDI-DI	**57400014PANTHERGX**	

ACCESSORIES LIST

Item nr.	**Accessories item nr.**
318011	Tray for Panther rollator

Harmonized norms used during conformity estimation:
PN-EN ISO 11199-2:2005; EN ISO 14971:2012, EN 1041:2009

Skanderborg, 2022-03-28, Thomas N. Christensen, Managing Director

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