

Declaration of Conformity (CE)

(Vaatimustenmukaisuusvakuutus)

Mobilex A/S, Grønlandsvej 5, DK-8660 Skanderborg, Denmark,

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

Arthritis Rollator Tiger - item no. 312201+312211

Arthritis Rollator Tiger with adjustable arm pads- item no. 312202

ACCESSORIES LIST

Item no.	Accessories item no.
312201+11	312315, 312320, 31233D, 312325, 312347, 312362, 312363, 312368, 31236A, 31236B, 312321

Item no.	Accessories item no.
312202	312315, 312320, 31233D, 312325, 312347, 312362, 312363, 312368, 31236A, 31236B, 312321

Harmonized norms used during conformity estimation:

PN-EN ISO 11199-2:2008; EN ISO 14971:2011, EN 1041:2009



EU2017/745 for Medical Equipment

In case of unintended incidents or suspected faults, failures or defects of the product, the dealer or Mobilex A/S must be contacted /

Thomas Nygaard Christensen
Managing Director/
Mobilex A/S

Skanderborg, 14-09-2021



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Signature