EU Declaration of Conformity

Manufacturers Name:	Ki Mobility
Manufacturers Address:	5201 Woodward Dr. Stevens Point, WI 54481
SRN (Single Registration Number):	US-MF-000012633
Authorized Representative Name (if applicable):	Etac Supply Center AB
Authorized Representative Address (if applicable):	Langgatan 12 33233 Anderstorp, Sweden
Authorized Representative SRN:	SE-AR-000001601
Basic UDI-DI:	0850013379SEATINGAG
UDI-DI:	00850013379163
Name of the Device(s):	Axiom LB
GMDN product code:	36254
Device Classification:	Class I, Rule 1
Intended Purpose:	A wheelchair component is a device intended for medical purposes that is generally sold as an integral part of a wheelchair but may also be sold separately as a replacement part.
Notified Body name:	Not Applicable
Notified Body Address:	Not Applicable
Notified Body Identification number:	Not Applicable
Conformity assessment route:	Ki Mobility uses the following procedures for the CE-labeling of their products according to the Regulation MDR 2017/745: Class 1: EU conformity declaration according to annex VIII

This declaration of conformity is issued under the sole responsibility of Ki Mobility. We hereby declare that the medical device(s) specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices. This declaration is supported by the Quality System certification to ISO 13485:2016 and on assessment of technical documentation.
All supporting documentation is retained at the premises of Ki Mobility.

Name:	Mark Murphy
Title:	Vice President of Operations, PRRC
Signature:	Mark Murphy Digitally signed by Mark Murphy Date: 2023.11.29 13:59:04 -05'00'
Date (YYYY-MM-DD) of issue:	2023-11-29