

## EU Declaration of Conformity



Otto Bock Mobility Solutions GmbH  
Lindenstraße 13  
07426 Königsee, Germany  
SRN: DE-MF-000005061

<b>Product</b>	Kimba see reference list
<b>Basic UDI-DI (GMN)</b>	40644060000000470G71RW
<b>Intended Purpose</b>	The product is intended for everyday indoor and outdoor use, by an attendant, of children and up with temporary or permanent limitations of the ability to walk, inability to walk or difficulty standing up.
<b>Classification</b>	Class I
<b>Conformity assessment procedure</b>	Annex II and Annex III of the Regulation (EU) 2017/745

Otto Bock Mobility Solutions GmbH hereby declares in sole responsibility that the medical device mentioned above complies with the current version of the Regulation (EU) 2017/745 of the European Parliament and of the Council on medical devices.

Applicable specifications are considered.  
The declaration is valid as of the date of signatures.

Königsee, October 18, 2023

Ralf Theisen  
Managing Director

Björn Kölle  
Person Responsible for  
Regulatory Compliance

**Reference list of EU Declaration of Conformity  
of Kimba from October 18, 2023**

<b>Item-N°</b>	<b>Description</b>
470G71=5_C	Kimba
HR3255=0000_K	Kimba Inline
HR3401=0000_K	Kimba Cross
HR2233=0000_K	Kimba Indoor mobility base
HR22330000-011	Kimba Indoor mobility base Sz 1
HR22430000-011	Kimba Indoor mobility base Sz 2