Declaration of Conformity

Manufacturer	**ArjoHuntleigh AB** Hans Michelsensgatan 10 211 20 Malmö, Sweden
Single Registration Number	**SE-MF-000000696**
Declaration	ArjoHuntleigh AB as the manufacturer of the following medical devices, takes sole responsibility and declares conformity with the applicable provisions of Medical Device Regulation (EU) 2017/745 concerning medical devices, by Annex IX.
Device Family Name	Maxi Move – KMCxAx-xx Where x depends of options and country
Intended Purpose	Mobile Lifter - Patient/Resident
Basic UDI-DI	**5060693520082W3**
Risk Class and Rule	Class IM, Rule 13
Additional Information	Also complies with the following EU Legislation: Machinery Directive 2006/42/EC RoHS Directive 2011/65/EU as amend Non-Automatic Weighing Instruments Directive 2014/31/EU
Notified Body Name and Number	CE 2797 BSI Group The Netherlands B.V. Number: 2797 CE Certificate Number MDR 718928

APPROVED BY		
Title: Senior Regulatory Affairs Specialist Name: Champa Patel	Signature: Date:	*Electronically signed by: Champa Patel* *Reason: I am signing as reviewer of this document* *Date: Apr 3, 2024 09:21 GMT+1*
Title: Local Quality Manager Name: Mélanie Chassé	Signature: Date:	*Electronically signed by: Melanie Chasse* *Reason: I am signing as approver of this document* *Date: Apr 3, 2024 06:46 EDT*

On behalf of ArjoHuntleigh AB: Place: Magog

Form: SOP-047-C rev.6

Declaration of Conformity

Manufacturer	**ArjoHuntleigh AB** Hans Michelsensgatan 10 211 20 Malmö, Sweden
Single Registration Number	**SE-MF-000000696**
Declaration	ArjoHuntleigh AB as the manufacturer of the following medical devices, takes sole responsibility and declares conformity with the applicable provisions of Medical Device Regulation (EU) 2017/745 concerning medical devices, by Annex IX.
Device Family Name	**Maxi Move – KMCxUx-xx** Where X depends of options and country
Intended Purpose	Mobile Lifter - Patient/Resident
Basic UDI-DI	**5060693520082W3**
Risk Class and Rule	Class IM, Rule 13
Additional Information	Also complies with the following EU Legislation: Machinery Directive 2006/42/EC RoHS Directive 2011/65/EU as amend
Notified Body Name and Number	CE 2797 BSI Group The Netherlands B.V. Number: 2797 CE Certificate Number MDR 718928

APPROVED BY	
Title: Regulatory Affairs Specialist	Signature: Champa Patel
Name: Champa Patel	Date: 01-Nov-2023
Title: Local Quality Manager	Signature:
Name: Mélanie Chassé	Date: 09-Nov-2023

On behalf of ArjoHuntleigh AB: Place: Magog

Declaration of Conformity

Manufacturer	**ArjoHuntleigh AB** Hans Michelsensgatan 10 211 20 Malmö, Sweden
Single Registration Number	**SE-MF-000000696**
Declaration	ArjoHuntleigh AB as the manufacturer of the following medical devices, takes sole responsibility and declares conformity with the applicable provisions of Medical Device Regulation (EU) 2017/745 concerning medical devices.
Device Family Name	Maxi Move – KMCxXx-xx Where X depends of options and country
Intended Purpose	Mobile Lifter - Patient/Resident
Basic UDI-DI	**5060693520112VK**
Additional Information	Also complies with the following EU Legislation: Machinery Directive 2006/42/EC RoHS Directive 2011/65/EU
Risk Class and Rule	CE Class I, Rule 13

APPROVED BY	
Title: Regulatory Affairs Specialist	Signature: Champa Patel
Name: Champa Patel	Date: 2022-Nov-04
Title: Local Quality Manager	Signature:
Name: Mélanie Chassé	Date: 11-Nov-2022

On behalf of ArjoHuntleigh AB: Place: Magog

Form: SOP-047-B rev.6