





## Declaration of Conformity

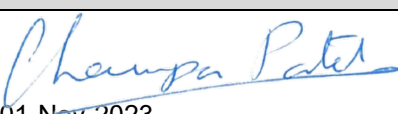
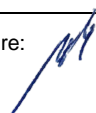
Manufacturer	<b>ArjoHuntleigh AB</b> Hans Michelsensgatan 10 211 20 Malmö, Sweden
Single Registration Number	<b>SE-MF-000000696</b>
Declaration	ArjoHuntleigh AB as the manufacturer of the following medical devices, takes sole responsibility and declares conformity with the applicable provisions of Medical 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	<b>5060693520082W3</b>
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	 BSI Group The Netherlands B.V. Number: 2797 2797 CE Certificate Number MDR 718928

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

On behalf of ArjoHuntleigh AB: Place: Magog


## Declaration of Conformity



Manufacturer	<b>ArjoHuntleigh AB</b> Hans Michelsensgatan 10 211 20 Malmö, Sweden
Single Registration Number	<b>SE-MF-000000696</b>
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	<b>Maxi Move – KMCxUx-xx</b> Where X depends of options and country
Intended Purpose	Mobile Lifter - Patient/Resident
Basic UDI-DI	<b>5060693520082W3</b>
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	 <b>2797</b> BSI Group The Netherlands B.V. Number: 2797 CE Certificate Number MDR 718928

APPROVED BY	
Title: Regulatory Affairs Specialist	Signature: 
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

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Manufacturer	<b>ArjoHuntleigh AB</b> Hans Michelsensgatan 10 211 20 Malmö, Sweden
Single Registration Number	<b>SE-MF-000000696</b>
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	<b>5060693520112VK</b>
Additional Information	Also complies with the following EU Legislation: Machinery Directive 2006/42/EC RoHS Directive 2011/65/EU
Risk Class and Rule	 Class I, Rule 13

APPROVED BY	
Title: Regulatory Affairs Specialist	Signature: 
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