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 5** **KMExx0-x-x**
Intended Purpose	Mobile Lifter - Patient/Resident
Basic UDI-DI	**5060693520112VK**
Additional Information	Also complies with the following EU Legislation: Machinery Regulation 2023/1230/EU RoHS Directive 2011/65/EU as amended
Risk Class and Rule	CE Class I, Rule 13

APPROVED BY		
Title: Regulatory Affairs Specialist	Signature:	*Electronically signed by: Champa Patel* *Reason: I am signing as approver of this document* *Date: Jan 21, 2025 16:28 GMT*
Name: Champa Patel	Date: Champa Patel	

On behalf of ArjoHuntleigh AB: Place: Magog

Form: SOP-047-B rev.7