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 Sky 440 - LExxxxx-xx** **Voyager Portable 200 – JExxxxx-xx** Where x depends on options and country
Intended Purpose	Electric hoist – patient/resident
Basic UDI-DI	**5060693520105VN**
Additional Information	Also complies with the following EU Legislation: Machinery Directive 2006/42/EC RoHS directive 2011/65/EU as amended by 2015/863
Risk Class and Rule	CE Class I, Rule 13

APPROVED BY		
Title: Regulatory Affairs Specialist	Signature: Champa Patel	*Electronically signed by: Champa Patel* *Reason: I am signing as approver of this document* *Date: Sep 25, 2024 10:42 GMT+1*
Name: Champa Patel	Date: 25-Sep-2024	
Title: Local Quality Manager	Signature:	*Electronically signed by: Melanie Chasse* *Reason: I am signing as approver of this document* *Date: Sep 27, 2024 14:38 EDT*
Name: Mélanie Chassé	Date: 27-Sep-2024	

On behalf of ArjoHuntleigh AB: Place: Magog