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 Twin, Non-AC-powered patient lift** Models: KTBBxxxxx-xx
Basic UDI-DI	**5060693520112VK**
GMDN Number and Term	**GMDN Number: 12330** Mobile patient lifting system, battery-powered
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: Quality Director	Signature:	*Electronically signed by: Anna Nowotna* *Reason: I am signing as approver of this document* *Date: Jul 23, 2024 13:58 GMT+2*
Name: Anna Nowotna	Date: Nowotna Anna	
Title: Senior Regulatory Compliance Manager	Signature:	*Electronically signed by: david moynham* *Reason: I am signing as approver of this document* *Date: Jul 23, 2024 12:04 GMT+1*
Name: David Moynham	Date: D. Moynham	

On behalf of ArjoHuntleigh AB: Place: Poznan

Form: SOP-047-B rev.5