

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	Tenor, Patient Lift Models: KHA10x0-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	 Class I, Rule 13

APPROVED BY		
Title: Quality Director	Signature:	<i>Electronically signed by: Anna Nowotna Reason: I am signing as approver of this document Date: Jul 23, 2024 13:58 GMT+2</i>
Name: Anna Nowotna	Date: 	
Title: Senior Regulatory Compliance Manager	Signature:	<i>Electronically signed by: david moynham Reason: I am signing as approver of this document Date: Jul 23, 2024 12:04 GMT+1</i>
Name: David Moynham	Date: 	

On behalf of ArjoHuntleigh AB: Place: Poznan