
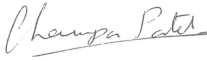



Declaration of Conformity

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	Sara Stedy – NTB2000-xx Sara Stedy Compact – NTB3000-xx Where x depends of country
Intended Purpose	Mobile Lifter - Patient/Resident
Basic UDI-DI	5060693520235W4
Additional Information	Also complies with the following EU Legislation: Machinery Directive 2006/42/EC
Risk Class and Rule	 Class I, Rule 1

APPROVED BY		
Title: Regulatory Affairs Specialist	Signature: 	<i>Electronically signed by: Champa Patel Reason: I am signing as approver of this document Date: Oct 2, 2024 08:21 GMT+1</i>
Name: Champa Patel	Date: 02-Oct-2024	
Title: Local Quality Manager	Signature: 	<i>Electronically signed by: Melanie Chasse Reason: I am signing as approver of this document Date: Oct 2, 2024 07:13 EDT</i>
Name: Mélanie Chassé	Date: 02-Oct-2024	

On behalf of ArjoHuntleigh AB: Place: Magog










100084266 - 01-Oct-2024


Final Audit Report

2024-10-02

Created:	2024-10-01 (Central European Summer Time)
By:	Joanie Masse (joanie.masse@arjo.com)
Status:	Signed
Transaction ID:	CBJCHBCAABAATFqeCEQPQAD5m5Povt-rZSf9imFZZIMf


"100084266 - 01-Oct-2024" History

-  Document created by Joanie Masse (joanie.masse@arjo.com)
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