




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, and by Annex IX Chapter I and III.
Device Family Name	Citadel Bed Frame System CXxxxxxxxxxxxxx
Intended Purpose	Medical Bed
Basic UDI-DI	5060693520075W6
Risk Class and Rule	Class IM, Rule 13 + metrology function
Additional Information	Also complies with the following EU Legislation: Machinery Regulation 2023/1230/EU RoHS Directive 2011/65/EU as amended Non-Automatic Weighing Instruments Directive 2014/31/EU
Notified Body Name and Number	 2797 BSI Group The Netherlands B.V. Number: 2797 CE Certificate Number MDR 718928

APPROVED BY		
Title: Local Quality Manager	Signature: 	Electronically signed by: Anna Nowotna Reason: I am signing as approver of this document Date: Nov 22, 2024 09:35 GMT+1
Name: Anna Nowotna	Date: 22-Nov-2024	
Title: Senior Regulatory Compliance Manager	Signature: 	Electronically signed by: david moynham Reason: I am signing as approver of this document Date: Nov 22, 2024 09:51 GMT
Name: David Moynham	Date: 22-Nov-2024	

On behalf of ArjoHuntleigh AB: Place: Poznan