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	**Enterprise 8000X** 8Xxxxxxxxxxxx
Intended Purpose	Medical Bed
Basic UDI-DI	**5060693520143VW**
Additional Information	Also complies with the following EU Legislation: Machinery Regulation 2023/1230/EU RoHS Directive 2011/65/EU as amended
Risk Class and Rule	CE Class I, Rule 13

APPROVED BY		
Title: Local Quality Manager	Signature: Nowotna Anna	*Electronically signed by: Anna Nowotna* *Reason: I am signing as approver of this document* *Date: Dec 4, 2024 10:13 GMT+1*
Name: Anna Nowotna	Date: 04-Dec-2024	
Title: Regulatory Compliance Manager	Signature: D. Moynham	*Electronically signed by: david moynham* *Reason: I am signing as approver of this document* *Date: Dec 4, 2024 09:19 GMT*
Name: David Moynham	Date: 04-Dec-2024	

On behalf of ArjoHuntleigh AB: Place: Poznan

Form: SOP-047-B rev.7