Declaration of Conformity

Manufacturer	**ArjoHuntleigh AB** Hans Michelsensgatan 10 211 20 Malmö, Sweden
Single Registration Number	Not available at date of signing.
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	Passive Clip Sling **Model: MAA2040M, MAA2070M, MAA4050M, MAA2000M, MAA4000M, MAA4000MA, MAA4100M, MAA4060M, MAA4061M, MAA4160M, MFA1000M**
Basic UDI-DI	5060693520150
GMDN Number and Term	GMDN Number: 37480 General – Purpose patient lifting system Sling/harness.
Additional Information	Also complies with the following EU Legislation: Machinery Directive 2006/42/EC
Risk Class and Rule	CE Class I, Rule 1

APPROVED BY		
Title: Senior Regulatory Compliance Manager	Signature:	D. Moynham
Name: David Moynham	Date:	04-Nov-2021
Title: Local Quality Manager	Signature:	Yacaida Almonte
Name:Yacaida Almonte	Date:	18/oct/2021

On behalf of ArjoHuntleigh AB: Place: Dominican Replublic

Form: SOP-047-B rev.5