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	**Loop comfort repositioning sling** Model: MAA7000X
Intended Purpose	**Patient/Resident - Non-Rigid support for Lifters**
Basic UDI-DI	**5060693520150VT**
Additional Information	Also complies with the following EU Legislation: Not applicable
Risk Class and Rule	CE Class I, Rule 1

APPROVED BY	
Title: Local Quality Manager	Signature: Anna Nowotna
Name: Anna Nowotna	Date: 20-DEC-2022
Title:Senior Regulatory Standards and Test Specialist	Signature: Małgorzata Kózka
Name: Malgorzata Kózka	Date: 20-DEC-2022

On behalf of ArjoHuntleigh AB: Place: Poznan

Form: SOP-047-B rev.6