

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	Auralis Mattress & Seat Cushion
Intended Purpose	For the prevention and/or management of pressure injuries
Basic UDI-DI	5060693520136VZ
Additional Information	Also complies with the following EU Legislation: None.
Risk Class and Rule	 Class I, Rule 1

APPROVED BY	
Title:Local Quality Manager	Signature: 
Name: Angela Jiang	Date: 1st Nov 2022
Title:Senior Regulatory Compliance Manager	Signature: 
Name:David Moynham	Date: 01-NOV 2022

On behalf of ArjoHuntleigh AB: Place: Suzhou

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Product/Model Numbers	Description
636T01S	Auralis Mattress Standard 110
636T01N	Auralis Mattress Narrow 110
636M02S	Auralis Mattress Standard 200
636M02N	Auralis Mattress Narrow 200
636M05S	Auralis Mattress Standard 175
636M05SUS	US labelled variant
636M05SCA	CA labelled variant
636M05N	Auralis Mattress Narrow 175
636M05NUS	US labelled variant
636M05NCA	CA labelled variant
636B02	Auralis Plus Mattress
636C01S	Auralis Seat Cushion
636T01S-P	Auralis 110 mattress STD
636T01N-P	Auralis 110 mattress NAR
636M05S-P	Auralis 175 mattress STD
636M05N-P	Auralis 175 mattress NAR
636M02S-P	Auralis 200 mattress STD
636M02N-P	Auralis 200 mattress NAR
636B02-P	Auralis Plus Mattress