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	Intraoperative Probe Adapter (PA8XS, PA8-HG)
Intended Purpose	Surgically invasive device intended for transient use to monitor the central circulatory system
Basic UDI-DI	**5060693520365WJ**
Additional Information	Also complies with the following EU Legislation: RoHS Directive 2011/65/EU
Risk Class and Rule	CE Class I, Rule 13

APPROVED BY	
Title: QRE Compliance Director	Signature:
Name: Steve Monks	Date: 30/09/2024

On behalf of ArjoHuntleigh AB: Place: Cardiff