Declaration of Conformity

Manufacturer	**ArjoHuntleigh AB** Hans Michelsensgatan 10 211 20 Malmö, Sweden
Declaration	ArjoHuntleigh AB as the manufacturer of the following medical devices, declare conformity with the applicable provisions of Directive 93/42/EEC of 14 June 1993, concerning medical devices, by Annex II.
Additional information	Manufactured and distributed on behalf of ArjoHuntleigh AB by: Huntleigh Healthcare Ltd 35 Portmanmoor Road Cardiff CF24 5HN United Kingdom
Device Family Name	Ultrasound Transducer – US1 / US1L / US2-EUR
GMDN Number and Term	41917 – **Foetal Doppler system probe**
Risk Class and Rule	Class IIb, Rule 10
Notified Body Name and Number	CE 2797 BSI 2797 CE Certificate Number CE01945

APPROVED BY	
Title: QRE Compliance Director	Signature:
Name: Steve Monks	Date: 7th December 2020

Form: SOP-047-C rev.5