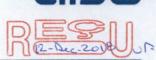


Declaration of Conformity



Manufacturer	ArjoHuntleigh AB Hans Michelsensgatan 10 211 20 Malmö, Sweden	
Declaration	ArjoHuntleigh AB as the manufacturer of the following medical devices, declares conformity with the applicable provisions of Directive 93/42/EEC of 14 June 1993, concerning medical devices, by Annex VII.	
Device Family Name	Maxi 500 KM56xxxx Where x depends on options	
GMDN Number and Term	12330 Mobile patient lifting system, battery powered	
Risk Class and Rule	CE Class I, Rule 12	

APPROVED BY		
Title: Senior RA Specialist	Date: 23-NOV-ZOI8	
Name: Maria Van Loon	Date: 23-NOV-Z018	
Title: Engineering Director	Signature: Signature:	
Name: Isabelle Bissonnette	Date: 21 NOV. 2018	
Title: LQM	Signature:	
Name: Mélanie Chassé	Date: 28-No.J- 70:8	