Declaration of Conformity

RESU 12-Dec-2018

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	Signature:
Name: Maria Van Loon	Date: 23-NOV-2018
Title: Engineering Director	Signature:
Name: Isabelle Bissonnette	Date: 27 NOV. 2018
Title: LQM	Signature:
Name: Mélanie Chassé	Date: 28-Nov-2018

Form: SOP-047-B rev.3