

## **Declaration of Conformity**

ArjoHuntleigh AB, Hans Michelsensgatan 10, 211 20 Malmö, Sweden, 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 VII.

This declaration covers the devices listed below:-

Device Family Name:	Maxi Twin, Non-AC-powered patient lift
	Models: KTBBxxxxx-xx
GMDN Number and Term:	GMDN Number: 12330 Mobile patient lifting system, battery powered
Risk Class and Rule:	Class I, Rule 12

Name: Paweł Cisowski

Signature:

Plant Manager

Date:

Name: Katarzyna Bobrow

Signature

Quality & Regulatory Manager

Date: 2014-12-18

2014-12-18