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	**Maxi Sky 1000 - LFxxxxx-xx** Where x depends on options and country
Intended Purpose	**Electric hoist – Patient/Resident**
Basic UDI-DI	**5060693520105VN**
Additional Information	Also complies with the following EU Legislation: Machinery Directive 2006/42/EC RoHS Directive 2011/65/EU as amended by 2015-863
Risk Class and Rule	CE Class I, Rule 13

APPROVED BY	
Title: Regulatory Affairs Specialist	Signature: Champa Patel
Name: Champa Patel	Date: 2022-Sep-01
Title: Local Quality Manager	Signature:
Name: Mélanie Chassé	Date: 07-Sep-2022

On behalf of ArjoHuntleigh AB: Place: Magog

Form: SOP-047-B rev.6