EU Declaration of Conformity

This EU declaration of conformity is issued under the sole responsibility of the manufacturer. The device that is covered by the present declaration is in conformity with EC Council Directive for Medical Devices 93/42/EEC of 14 June1993 & Medical Device Regulation (EU) 2017/745.

Product Name:	Oxford Professional Advance
Description:	The Oxford Professional Advance range of hoists are ergonomically designed mobile devices for the safe lifting and handling of patients for the transfer or moving a person over short distances.
Product Codes:	OXF-ADVANCE-AU-S OXF-ADVANCE-EM OXF-ADVANCE-S
Manufacturer SRN:	CA010152
Basic UDI-DI (GMN):	50556184OXF-PROSD **GMDN:** 12330
Classification Type:	Class I [Rule 1 of MDR 2017/745, Annex VIII, Chapter III, Section 4.1]
Authorised Representative:	Joerns Healthcare BV, Zeilschip 26,3991 CT Houten, Netherlands
Registered place of business:	Joerns Healthcare BV, Zeilschip 26, 3991 CT Houten, Netherlands
Responsible Person:	Chris Murray
Registered place of business:	Joerns Healthcare LLC, 2100 Design Road, Arlington. TX76014, United States

This device was tested and is in conformity with all relevant clauses within the following standards:

Standard	Description
BS EN ISO 10535:2006	Hoists for the transfer of disabled persons
BS EN ISO 12182:2012	Assistive products for persons with disability. General requirements & test methods
BS EN ISO 14971:2019	Application of Risk Management to medical devices

Issued By:	Stan Snijders – Managing Director of Joerns Healthcare, Europe
Issue Date:	07/01/2021
Location:	UK / Netherlands
On behalf of:	Joerns Healthcare Limited, UK
Signature:	