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EU Declaration of Conformity

For MR Patient Transportation Wheelchairs

Basic UDI-DI: 506500603700GMDN40699LP

Regulation (EU) 2017/745 of the European Parliament and the Council of 5 April 2017 concerning Medical Devices.

The undersigned declares that the products described in this document meet the Council provisions that apply to them, and the CE Mark may be affixed. This declaration is issued under the sole responsibility of the manufacturer.

General Product Name:	MR Patient Transportation Wheelchairs
Legal Manufacturer: (Name on Label)	Wardray Premise Limited, Unit 2 Springvale Works, Elland Road, Brighouse, West Yorkshire, HD6 2RN United Kingdom
Manufacturers SRN:	GB-MF-000021675
Variants:	As per Appendix II (This document) – Product Listing/Schedule
Intended Purpose:	Patient Transport to and from MRI Department
MDR Classification:	Class I Classification rule 1 from Annex VIII of the MDR directive specifies that all non-invasive devices are classified as class 1 unless one of the other rules listed is applicable.
Notified Body:	Not Applicable for Class I
EC Certificate:	Not Applicable as we self-certify Class I devices.
EU Authorised Representative:	Advena Limited. Tower Business Centre, 2nd Flr., Tower Street, Swatar, BKR 4013 Malta.
EU Authorised Representative SRN:	MT-AR-000000234
Medical Device Regulation Assessment Route:	In conformity with Annex IX of the Medical Device Regulation. Issuing of the Declaration of Conformity in accordance with Article 19 after drawing up the technical documentation laid out in Annexes II and III of the EU MDR 2017/745

Name Justine Colquhoun **Position** Director **Place** Head Office, Sutton, Surrey

Signed Justine Colquhoun **Date** 25/03/2024

Who is the natural and legal person with responsibility for the design, manufacture, packaging and labelling before the device is placed on the market under this manufacturer's name regardless of whether these operations are carried out by the manufacturer or on his behalf by a third party.

Appendix I – Applicable Standards

This present declaration is also in conformity with the following European standards and Common Specifications:

Standard/Document Name	Description
2017/745	Regulation (EU) 2017/745 of the European Parliament and the Council of 5 April 2017 concerning Medical Devices
BS EN 12183:2022	Specification for Manually Propelled Wheelchairs
BS EN ISO 20417:2021	Information supplied by the manufacturer of medical devices
EN ISO 13485:2016	Medical Devices – Quality Management Systems – Requirements for Regulatory Purposes
EN ISO 14971:2019 + A11: 2021	Medical Devices – Application of Risk Management to Medical Devices
BS EN ISO 15223-1:2021	Medical Devices – Symbols to be used with Medical Device labels
PDCEN ISO/TR 24971:2020	Guidance on the application of ISO 14971

Appendix II – Product Listing/Schedule

Part/Catalogue Number	Device Name	EMDN Code
MR4500	MR Folding Portering Chair	41630
MR4501	MR Non Folding Portering Chair	41630
MR4588	MR Folding Bariatric Portering Chair	57979
FGPC-200	Ferrous Free Portering Chair	41630

Version History

Version	Compiled by	Date	Description
1.0	J C Colquhoun	30/10/2019	First issue
2.0	J C Colquhoun	27/05/2021	Amended to include basic UDI-DI
3.0	J C Colquhoun	02/07/2021	Removal of reference to Annex IX under Medical Device Regulation Assessment Route
4.0	J C Colquhoun	09/12/2021	Addition of FGPC-200 Portering Chair
5.0	J C Colquhoun	15/07/2022	Addition of Manufacturers SRN number
6.0	J C Colquhoun	25/03/2024	Update of British Standards