EU DECLARATION OF CONFORMITY

EU Konformitätserklärung

MANUFACTURER:	**Red Milawa Pty Ltd T/A Magic Mobility [Australia]** 3 International Court, Scoresby VIC 3179 Australia
MANUFACTURER: **SRN/ EUDAMED ACTOR ID**	AU-MF-000034907
AUTHORIZED REPRESENTATIVE:	**Sunrise Medical GmbH** Kahlbachring 2-4 D-69254 Malsch / Heidelberg
AUTHORIZED REPRESENTATIVE: **SRN/ EUDAMED ACTOR ID**	DE-AR-000006321
PRODUCT NAME	Extreme X8
TRADE NAMES	Magic Extreme X8
EUDAMED REFERENCE/CATALOGUE NUMBER	Extreme X8
EUDAMED DEVICE NAME:(Basic UDI)	Power Wheelchair Magic Mobility
BASIC UDI-DI:	9356399pwcmagicT3
UDI-DI	09356399059695
CLASSIFICATION	Class I
REGULATION	MDR (REGULATION (EU) 2017/745 on medical devices)
NOMENCLATURE CODES	Y122127 ELECTRIC WHEELCHAIRS

INTENDED PURPOSE:
Magic Mobility Extreme X8 Power wheelchairs are exclusively for a user who is unable to walk or has limited mobility, for their own personal use indoor and outdoor. When an Attendant Control Module is fitted, the Power Wheelchair may be operated by an assistant on behalf of the user. When a Dual Control Module is fitted the Power Wheelchair may be operated by the user, or control may be switched to an assistant to operate on behalf of the user. When a folding back is fitted the wheelchair back maybe folded for transport.

We, Red Milawa Pty Ltd T/A Magic Mobility, declare under our sole responsibility that the product(s) to which this declaration relates, is a class 1 device, and is in conformity with the requirements of the MDR (REGULATION (EU) 2017/745 on medical devices).

Name:	Clare Daff	Terence Walsh
Title:	R&D Manager	National Quality and Regulatory Manager
Function:	R&D Manager	Person Responsible for Regulatory Compliance
Place and date of issue:	Australia 24/08/2023	Australia 24/08/2023
Signature: