





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	XT2
TRADE NAMES	XT2
EUDAMED REFERENCE/CATALOGUE NUMBER	XT2
EUDAMED DEVICE NAME:(Basic UDI)	Power Wheelchair Magic Mobility
BASIC UDI-DI:	9356399pwcmagicT3
UDI-DI	09356399060974
CLASSIFICATION	Class I
REGULATION	MDR (REGULATION (EU) 2017/745 on medical devices)
NOMENCLATURE CODES	Y122127 ELECTRIC WHEELCHAIRS
INTENDED PURPOSE:	Magic Mobility XT2 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 may be 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	Ed Tan
Title:	Engineering Manager	National Quality and Regulatory Manager
Function:	R&D Engineering	Person Responsible for Regulatory Compliance
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Signature:		

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