

## EU DECLARATION OF CONFORMITY

### *EU Konformitätserklärung*

<b>MANUFACTURER:</b>	<b>Red Milawa Pty Ltd T/A Magic Mobility [Australia]</b> 3 International Court, Scoresby VIC 3179 Australia
<b>MANUFACTURER: SRN/ EUDAMED ACTOR ID</b>	AU-MF-000034907
<b>AUTHORIZED REPRESENTATIVE:</b>	<b>Sunrise Medical GmbH</b> Kahlbachring 2-4 D-69254 Malsch / Heidelberg
<b>AUTHORIZED REPRESENTATIVE: SRN/ EUDAMED ACTOR ID</b>	DE-AR-000006321
<b>PRODUCT NAME</b>	XT2
<b>TRADE NAMES</b>	XT2
<b>EUDAMED REFERENCE/CATALOGUE NUMBER</b>	XT2
<b>EUDAMED DEVICE NAME:(Basic UDI)</b>	Power Wheelchair Magic Mobility
<b>BASIC UDI-DI:</b>	9356399pwcmagicT3
<b>UDI-DI</b>	<b>09356399060974</b>
<b>CLASSIFICATION</b>	Class I
<b>REGULATION</b>	MDR (REGULATION (EU) 2017/745 on medical devices)
<b>NOMENCLATURE CODES</b>	Y122127 ELECTRIC WHEELCHAIRS
<b>INTENDED PURPOSE:</b> 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

**Title:** Engineering Manager

**Function:** R&D Engineering

**Place and date of issue:** Australia 30 Jan 2025

Ed Tan

National Quality and Regulatory Manager

Person Responsible for Regulatory Compliance

Australia 30 Jan 2025

**Signature:**



*Ed Tan*