



UK Declaration of Conformity

For the following equipment :

Product Name: AC/DC Medical Adaptor

Model Designation:GSM120Bx (x=12,15,20,24,48)

The designated product(s) is(are) in conformity with the relevant legislation:

The Restriction of the Use of Certain Hazardous Substances in Electrical and Electronic Equipment Regulations 2012: SI 2012 No. 3032

Medical Devices Regulations 2002 (SI 2002 No 618, as amended) (UK MDR 2002) BS EN 60601-1:2006+A1+A12+A2

BS EN60601-1-11:2015+A1

TUV certificate No : TA50293271

BS EN60601-1-2:2015+A1:2021

EMI (Electro-Magnetic Interference)

Conducted emission / Radiated emission

| | BS EN 55011:2016+A2:2021 BS EN IEC 61024-3:2018 | Class B | |
|------------------|--|---------|--|
| Harmonic current | BS EN IEC 61000-3-2:2019+A1:2021 | | |
| Voltage flicker | BS EN 61000-3-3:2013+A1:2019+A2:2021 | | |

EMS (Electro-Magnetic Susceptibility)

| BS EN60601-1-2:2015+A1:2021 BS EN IEC 61204-3:2018 | | | | | | |
|--|----------------------------|---|---|--|--|--|
| ESD air | BS EN 61000-4-2:2009 | Level 4 | 15KV | | | |
| RF field susceptibility | BS IEC EN61000-4-3:2020 | Level 3 | 10V/m(80MHz-2.7GHz) | | | |
| RF field susceptibility | BS IEC EN 61000-4-3:2020 | Table 9 | 9~28V/m (385MHz~5.78GHz) | | | |
| EFT bursts | BS EN 61000-4-4:2012 | Level 3 | 2KV/5KHz | | | |
| Surge susceptibility | BS EN 61000-4-5:2014+A1:20 |)17 Level 3 | 1KV/Line-Line | | | |
| Conducted susceptibility | BS EN 61000-4-6:2014 | Level 3 | 10V | | | |
| Magnetic field immunity | BS EN 61000-4-8:2010 | Level 4 | 30A/m | | | |
| Voltage dip, interruption | | 0% residual voltage for 1 cy 0% residual voltage for 250 | cles, 70% residual voltage for 25 cycles,) cycles | | | |

Note:

The power supply is considered as a component that will be operated in combination with final equipment. Since EMC performance will be affected by the complete system, the final equipment manufacturers must re-qualify EMC Directive on the complete system again.

For guidance on how to perform these EMC tests, please refer to TDF (Technical Documentation File). The product under declaration is just a unit without medical function. Complete MDR should only be verified when it is used together with particular medical device(s).

This Declaration is effective from serial number SC3xxxxxx

Person responsible for marking this declaration :

| MEAN WELL Enterprises Co | o., Ltd. | | |
|-----------------------------------|-------------------------|--|---------------|
| (Manufacturer Name) | | | |
| No.28, Wuquan 3rd Rd., Wu | igu Dist., New Taipei (| City 24891, Taiwan | |
| (Manufacturer Address) | \wedge - | | |
| Aries Jian/ Director, Group R&D : | Tries | Alex Tsai/ Director, Product Strategy Center : | \mathcal{C} |
| (Name / Position) | (Signature) | (Name / Position) | (Signature) |
| Taiwan | Sep 15th, 2023 | | |
| (Place) | (Date) | | |

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