



# **UK Declaration of Conformity**

For the following	equipment	:
-------------------	-----------	---

Product Name: Medical Type Switching Power Supply

Model Designation: RPS-300-x-y (x=12,15,24,27,48);y=C or blank

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 TUV certificate No : TA50236914

BS EN 60601-1-2:2015+A1:2021

## **EMI (Electro-Magnetic Interference)**

Conducted emission / Radiated emission

	BS EN 55011:2016+A11:2020	Class B
Harmonic current	BS EN IEC 61000-3-2:2019	
Voltage flicker	BS EN 61000-3-3:2013+A1:2019	

#### **EMS (Electro-Magnetic Susceptibility)**

Lino (Liectro-inagnetic Susceptibility)					
ESD air	BS EN 61000-4-2:2009	Level 4	15KV		
ESD contact	BS EN 61000-4-2:2009	Level 4	8KV		
RF field susceptibility	BS EN IEC 61000-4-3:2020	Level 3	10V/m(80MHz-2.7GHz)		
RF field susceptibility	BS EN IEC 61000-4-3:2020	Table 9	9~28V/m (385MHz~5.78GHz)		
EFT bursts	BS EN 61000-4-4:2012	Level 3	2KV/100KHz		
Surge susceptibility	BS EN 61000-4-5:2014+A1:2017	Level 4	2KV/Line-Line		
Surge susceptibility	BS EN 61000-4-5:2014+A1:2017	Level 4	4KV/Line-Earth		
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	BS EN IEC 61000-4-11:2020 100% dip 1 periods 30% dip 25 periods 100% interruptions 250 periods				

## 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 SC3xxxxxxx

# Person responsible for marking this declaration:

MEAN WELL Enterprises Co., Ltd.

(Manufacturer Name)

No.28, Wuquan 3rd Rd., Wugu Dist., New Taipei City 24891, Taiwan

(Manufacturer Address)

Aries Jian/ Director, Group R&D: (Name / Position)

(Name / Position) (Signature) Taiwan Nov. 29th, 2023

(Place) (Date)

Alex Tsai/ Director, Product Strategy Center:
(Name / Position)
(Signature)

Version: 3