



# **UK Declaration of Conformity**

For the following equipment :

Product Name: Switching Power Supply

Model Designation:MFM-30-X(X=3.3,5,12,15,24,48); MPM-30-XY (X=3.3, 5, 12, 15, 24, 48 Y=Blank or ST)

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 : TA50388097		
BS EN 60601-1-2:2015+A1:2021			

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

Conducted emission / Radiated emission

	BS EN 55011:2016+A11:20	120	Class B			
			Class D			
Harmonic current	BS EN IEC 61000-3-2:201	9				
Voltage flicker	BS EN 61000-3-3:2013+A	1:2019				
EMS (Electro-Magnetic Susceptibility)						
BS EN 60601-1-2:2015+A1:2021						
ESD air	BS EN 61000-4-2:2009	Level 4	15KV			
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 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	BS EN IEC 61000-4-11:2020	ų – 1	es,0% residual voltage for 1 cycles, ycles , 0% residual voltage for 250 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 numberSC3xxxxxxx

#### Person responsible for marking this declaration :

MEAN WELL Enterprises Co	o., Ltd.				
(Manufacturer Name)					
No.28, Wuquan 3rd Rd., Wugu Dist., New Taipei City 24891, Taiwan					
(Manufacturer Address)	$\wedge$ -				
Aries Jian/ Director, Group R&D :	Trips	Alex Tsai/ Director, Product Strategy Center :	$\mathcal{C}$		
(Name / Position)	(Signature)	(Name / Position)	(Signature)		
Taiwan	Jun. 14th, 2023	_			
(Place)	(Date)				