



Declaration of Conformity

For the fo	llowing	equipment	
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Product Name: Switching Power Supply

Model Designation: MPM-X-YZ (X=45 65 90; Y=5, 12, 15, 24, 48; Z=Blank or ST

is herewith confirmed to comply with the requirements set out in the Council Directive, the following standards were applied:

RoHS Directive (2011/65/EU), (EU)2015/863

MDR Directive (EU) 2017/745

EN 60601-1:2006+A1+A12+A2 TUV certificate No: TA50457911

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

EMI (Electro-Magnetic Interference)

Conducted emission / Radiated emission

	EN 55011:2016+A2:2021	Class B
Harmonic current	EN IEC 61000-3-2:2019+A1:2021	
Voltage flicker	EN 61000-3-3:2013+A1:2019+A2:2021	
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EMS (Electro-Magnetic Susceptibility)

EN 0601-1-2:2015+A1:202	FN	0601	-1-2·20°	15+A1	1.202
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EN 61000-4-2:2009	Level 4	15KV
EN IEC 61000-4-3:2020	Level 3	10V/m(80MHz-2.7GHz)
EN IEC 61000-4-3:2020	Table 9	9~28V/m (385MHz~5.78GHz)
EN 61000-4-4:2012	Level 3	2KV/100KHz
EN 61000-4-5:2014+A1:20	17 Level 3	1KV/Line-Line
EN 61000-4-6:2014	Level 3	10V
EN 61000-4-8:2010	Level 4	30A/m
EN IEC 61000-4-11:2020	0% residual voltage for 0.5 cycl 70% residual voltage for 25 cycl	es, 0% residual voltage for 1 cycles, les , 0% residual voltage for 250 cycles
	EN IEC 61000-4-3:2020 EN IEC 61000-4-3:2020 EN 61000-4-4:2012 EN 61000-4-5:2014+A1:20 EN 61000-4-6:2014 EN 61000-4-8:2010	EN IEC 61000-4-3:2020 Level 3 EN IEC 61000-4-3:2020 Table 9 EN 61000-4-4:2012 Level 3 EN 61000-4-5:2014+A1:2017 Level 3 EN 61000-4-6:2014 Level 3 EN 61000-4-8:2010 Level 4 EN IEC 61000-4-11:2020 0% residual voltage for 0.5 cycl

Note:

A component power supply will be installed into final equipment. Since EMC performance will be affected by the complete installation, the final equipment manufacturers must re-qualify EMC Directive on the complete installation 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)

(Signature)

Alex Tsai/Director, Product Strategy Center:

(Name / Position)

(Signature)

Taiwan

Oct. 19th, 2023

(Place) (Date)