



UK Declaration of Conformity

For the following equipment :

Product Name: Switching Power Supply

Model Designation: NMP650-aaaa-xx(a=C, D, E, H, K or #: x=0~9) C=NMS-240-05, E=NMS-240-12, H=NMS-240-24, K=NMS-240-48, #=Blank, D=NMD-240, #=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 : TA50393537 (for NMP650)

TUV certificate No : TA50393494 (for NMS-240&NMD-240)

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

Electrical Equipment (Safety) Regulations 2016 :

BS EN 62368-1:2014+A11:2017

TUV certificate No : R50417981

Electrical Compatibility Regulations 2016 :

EMI (Electro-Magnetic Interference)

Conducted emission / Radiated emission

BS EN 55032:2015+A11:2020

Class B

BS EN 55011:2016+A2:2021

Harmonic current

BS EN IEC 61000-3-2:2019

Voltage flicker

BS EN 61000-3-3:2013+A1:2019

EMS (Electro-Magnetic Susceptibility)

BS EN 55024:2010+A1:2015 BS EN 60601-1-2:2015+A1:2021 BS EN 55035:2017+A11:2020

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 2

3V

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 Regulations 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 TC3xxxxxxx

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 :

(Signature)

Alex Tsai/ Director, Product Strategy Center :

(Name / Position)

(Signature)

Taiwan

Nov. 13th, 2023

(Place)

(Date)