



### Declaration of Conformity

For the following equipment :

Product Name: Medical Type Switching Power Supply

Model Designation: RPXY-160Z (X=D, T) (Y=G or blank) (Z=-5, -12, -15, -24, -48, A, B, C, D)

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+A11+A1+A12

TUV certificate No : TA50147896

**MDR Directive (EU) 2017/745**

EN 60601-1-2:2015

**EMI (Electro-Magnetic Interference)**

Conducted emission / Radiated emission

EN 55011:2016+A2:2021

Class B

Harmonic current

EN IEC 61000-3-2:2019

Voltage flicker

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

**EMS (Electro-Magnetic Susceptibility)**

ESD air

EN 61000-4-2:2009

Level 4

15KV

ESD contact

EN 61000-4-2:2009

Level 4

8KV

RF field susceptibility

EN IEC 61000-4-3:2020

Level 3

10V/m(80MHz-2.7GHz)

RF field susceptibility

EN IEC 61000-4-3:2020

Table 9

9~28V/m (385MHz~5.78GHz)

EFT bursts

EN 61000-4-4:2012

Level 3

2KV/100KHz

Surge susceptibility

EN 61000-4-5:2014+A1:2017

Level 3

1KV/Line-Line

Surge susceptibility

EN 61000-4-5:2014+A1:2017

Level 3

2KV/Line-Earth

Conducted susceptibility

EN 61000-4-6:2014

Level 3

10V

Magnetic field immunity

EN 61000-4-8:2010

Level 4

30A/m

Voltage dip, interruption

EN IEC 61000-4-11:2020 0% residual voltage for 1 cycles, 70% residual voltage for 25 cycles, 0% residual voltage for 250 cycles

**Note:**

A component power supply with load will be installed into final equipment which consists of an electronically shielded metal enclosure. Since EMC performance will be affected by the complete installation, the final equipment manufacturers must re-qualify EMC Directive on the complete installation again.

The EMC tests mentioned above are performed using a well defined metal plate to simulate said metal enclosure.

For guidance on how to perform these EMC tests, please refer to "EMI testing of component power supplies".(as available on <http://www.meanwell.com>)".

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 SC1xxxxxx

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 :

*Aries*  
(Signature)

Alex Tsai/Director, Product Strategy Center :

*[Signature]*  
(Signature)

(Name / Position)

(Name / Position)

(Signature)

Taiwan

Oct. 29th, 2021

(Place)

(Date)