



Declaration of Conformity

For the following equipment :

Product Name: 100W GaN Fast Charger

Model Designation: NGE100xyyyy (x=Blank ,U ; yyy= maybe blank ,-,0-9 , A-Z or a-z for market purpose 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

Low Voltage Directive (2014/35/EU) :

EN 62368-1:2014+A11:2017

Dekra Certificate: 35-147458

Electromagnetic Compatibility Directive (2014/30/EU) :

EMI (Electro-Magnetic Interference)

| | | |
|--------------------|---------------------------------------|----------|
| Conducted emission | EN 55032:2015+A1:2020 | |
| Radiated emission | EN 55032:2015+A11:2020 | Class B |
| Harmonic current | EN IEC 61000-3-2:2019+A1:2021 | Class A |
| Voltage flicker | EN IEC 61000-3-3:2013+A1:2019+A2:2021 | Clause 5 |

EMS (Electro-Magnetic Susceptibility)

EN 55035:2017+A11:2020

| | | | |
|---------------------------|---------------------------|--|--------------------|
| ESD air | EN 61000-4-2:2009 | Level3 | 8KV |
| RF field susceptibility | EN IEC 61000-4-3:2020 | Level 2 | 3V/m(80MHz~2.7GHz) |
| EFT bursts | EN 61000-4-4:2012 | Level 2 | 1KV |
| Surge susceptibility | EN 61000-4-5:2014+A1:2017 | Level 3 | 1KV/Line-Line |
| Conducted susceptibility | EN 61000-4-6:2014 | Level 2 | 3V |
| Magnetic field immunity | EN 61000-4-8:2010 | Level 1 | 1A/m |
| Voltage dip, interruption | EN IEC 61000-4-11:2020 | <5% residual voltage for 0,5cycle, 70% residual voltage for 25cycles, <5% residual voltage for 250cycles | |

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 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).

Energy-Related Products Directive (2009/125/EC) :

Ecodesign requirements for no-load condition electric power consumption and average active efficiency of external power supplies (EU)2019/1782

This Declaration is effective from serial number SC4xxxxxxx

Person responsible for marking this declaration :

MEAN WELL Enterprises Co., Ltd.

(Manufacturer Name)

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(Manufacturer Address)

Aries Jian/ Director, Group R&D :

(Name / Position)

Aries
(Signature)

Alex Tsai/Director, Product Strategy Center :

(Name / Position)

[Signature]
(Signature)

Taiwan

(Place)

Feb. 18th, 2024

(Date)