

# CERTIFICATE OF COMPLIANCE

**Certificate Number** 20190814-E308218  
**Report Reference** E308218-20171211  
**Issue Date** 2019-AUGUST-14

**Issued to:** ARCH ELECTRONICS CORP  
3F No 79 Sec 1 Hsin Tai Wu Rd  
Hsi Chih District, New Taipei  
221 TAIWAN

**This certificate confirms that  
representative samples of**

COMPONENT - POWER SUPPLIES, MEDICAL AND  
DENTAL

Switching Power Supply  
Model: MQF500O-xSYYYYYYY, MQF500U-xSYYYYYYY,  
MQF500E-xSYYYYYYY  
(where "x" can be 9 to 12.48, 10 to 15.9, 18 to 24.96, 21 to  
29.4, 36 to 49.92; where "Y" can be 0-9, a-z, A-Z, - or blank  
for marketing purpose and not related to safety)

Have been investigated by UL in accordance with the  
component requirements in the Standard(s) indicated on  
this Certificate. UL Recognized components are incomplete  
in certain constructional features or restricted in  
performance capabilities and are intended for installation in  
complete equipment submitted for investigation to UL LLC.

**Standard(s) for Safety:**  
**Additional Information:**

See Addendum Page  
See the UL Online Certifications Directory at  
<https://iq.ulprospector.com> for additional information.

This *Certificate of Compliance* does not provide authorization to apply the UL Recognized Component Mark. Only the UL Follow-Up Services Procedure provides authorization to apply the UL Mark.

Only those products bearing the UL Recognized Component Mark should be considered as being UL Certified and covered under UL's Follow-Up Services.

Look for the UL Recognized Component Mark on the product.



Bruce Mahrenholz, Director North American Certification Program

UL LLC

Any information and documentation involving UL Mark services are provided on behalf of UL LLC (UL) or any authorized licensee of UL. For questions, please contact a local UL Customer Service Representative at <http://ul.com/aboutul/locations/>



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This is to certify that representative samples of the product as specified on this certificate were tested according to the current UL requirements.

**Standard(s) for Safety:-**

Medical Electrical Equipment, Part 1: General requirements for basic safety and essential performance, ANSI/AAMI ES60601-1 (2005/(R)2012 + A1:2012", C1:2009/(R)2012 + A2:2010/(R)2012) - Amendment 1 and CAN/CSA-C22.2 No. 60601-1:14.  
ANSI/AAMI ES 60601-1 can be representative for CAN/CSA-C22.2 No. 60601-1 (2014) since there's no difference between these two standards.



Bruce Mahrenholz, Director North American Certification Program

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