Description

UL TEST REPORT AND PROCEDURE

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	Standby power (Optional and only applicable to y=blank): 5Vdc, 2A or 5.25Vdc, 1.9A; See enclosed Miscellaneous-Model different list for detail; Input: 100-240 Vac, 6.0 A or 4.0A, 50-60 Hz Output: - CUS500M1-12, CME500A-12: 12 Vdc (10.8Vdc-12.9Vdc), 41.7A - CUS500M1-19, CME500A-19: 19 Vdc (17.1Vdc-20.5Vdc), 26.4A - CUS500M1-24, CME500A-24: 24 Vdc (21.6Vdc-25.9Vdc), 20.9A - CUS500M1-28, CME500A-28: 28 Vdc (25.2Vdc-30.2Vdc), 17.9A - CUS500M1-32, CME500A-32: 32 Vdc (28.8Vdc-34.5Vdc), 15.7A - CUS500M1-36, CME500A-36: 36 Vdc (32.4Vdc-38.8Vdc), 13.9A - CUS500M1-48, CME500A-48: 48 Vdc (43.2Vdc-51.8Vdc), 10.5A
	See enclosed Miscellaneous-Model different list for detail.
Applicant Name and Address:	TDK-Lambda (China) Electronics Co Ltd No.95, Zhujiang Rd, Xinwu District, Wuxi JiangSu, CHINA

This is to certify that representative samples of the products covered by this Test Report have been investigated in accordance with the above referenced Standards. The products have been found to comply with the requirements covering the category and the products are judged to be eligible for Follow-Up Service under the indicated Test Procedure. The manufacturer is authorized to use the UL Mark on such products which comply with this Test Report and any other applicable requirements of UL LLC ('UL') in accordance with the Follow-Up Service Agreement. Only those products which properly bear the UL Mark are considered as being covered by UL's Follow-Up Service under the indicated Test Procedure.

The applicant is authorized to reproduce the referenced Test Report provided it is reproduced in its entirety.

UL authorizes the applicant to reproduce the latest pages of the referenced Test Report consisting of the first page of the Specific Technical Criteria through to the end of the Conditions of Acceptability as applicable.

Any information and documentation involving UL Mark services are provided on behalf of UL LLC (UL) or any authorized licensee of UL.

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Prepared by:
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Paul Zhang, Handler

Reviewed by: Jay Lu, Reviewer

Supporting Documentation

The following documents located at the beginning of this Procedure supplement the requirements of this Test Report:

- A. Authorization The Authorization page may include additional Factory Identification Code markings.
- B. Generic Inspection Instructions
 - i. **Part AC** details important information which may be applicable to products covered by this Procedure. Products described in this Test Report must comply with any applicable items listed unless otherwise stated in the body of this Test Report.
 - ii. **Part AE** details any requirements which may be applicable to all products covered by this Procedure. Products described in this Test Report must comply with any applicable items listed unless otherwise stated in the body of each Test Report.
 - iii. **Part AF** details the requirements for the UL Certification Mark which is not controlled by the technical standard used to investigate these products. Products are permitted to bear only the Certification Mark(s) corresponding to the countries for which it is certified, as indicated in each Test Report.

Product Description

The PSU is a component type switching mode power supplies intended for the earthed construction or nonearthed construction of medical equipment.

- For earthed construction (Class I), the PSU need to be reliably earthed and professionally installed and fixed with metal screws.

- For non-earthed construction (Class II), no earthing connection is required. The PSU need to be fixed so, that it is insulated from any unearthed accessible conductive part by reinforced insulation.

Refer to the Report Modifications page for any modifications made to this report.

Model Differences

Model CME600Ay-zxxxxxx is identical to model CUS600My-zxxxxxx except for model name. All models are identical, except for the optional chassis, cover, turns of Transformer and the rating of some components which results in different output ratings; y means blank and y means 1 have different T2 and relevant circuit construction;

For models: CUS500M1-zxxxxxx, CME500A-zxxxxxx; CUS500M1 & CME500A series is the modified models base on CUS600M1 & CME600A1 series, CUS500M1 & CME500A series Schematic is as same as CUS600M1 & CME600A1 series; CUS500M1 & CME500A series are without the standby power circuit; Model CME500A-zxxxxxx is identical to model CUS500M1-zxxxxxx except for model name.

See Enclosed Miscellaneous - Model Different List for details.

Additional Information

Project 4789613276:

1. Add additional models CUS500M1-zxxxxxx, CME500A-zxxxxxxx (z = 12, 19, 24, 28, 32, 36 or 48; xxxxxx = /T, /J, /M, /C, /C2, /SF, /G, /EF, other alphanumeric character, symbol or blank), which are similar to original models CUS600M1-zxxxxxx, CME600A1-zxxxxxx with following differences: - Rated input current, output ratings;

- Add alternate heatsink combination 2 that is different from the original combination 1 on CUS600M1 series; see enclosed miscellaneous-heatsink combination 1&2 for details. Combination 2 assmebled CA922-32-01x on KFA1 (Pri, side) only, other heatsinks were removed;

- Component parameter adjustment for MOSFET (Q1), Diode (D1), Primary Electrolytic Capacitor (C6) and Resistor (R108).

2. update the definition of variables on model designations.

3. Add new alternate component of L4 PQ35-671/x (the x can be 00-99).

Project 4789581080:

1. Add new model CUS600M-12/GEKxxxx, that is identical with CUS600M-12xxxxxxx expect for:

- standby power rated O/P from 5Vdc/2A to 5.25V/1.9A;

- parts components were changed for gaining the changed standby power rating, but no any critical component were changed;

2. Correct the typo of Y-Capacitor in CCL;

3. Correct the file No from E508585-D1000-0 to E508584-D1000-1.

Project 4789290688:

For models: CUS600My-zxxxxxx, CME600Ay-zxxxxxx (y=blank);

1. Add additional new factory TDK-Lambda (China) Electronics Co., Ltd. (No.95, Zhujiang Road, Xinwu District, Wuxi Jiangsu P. R. China);

2. Add additional description of peak power in supplementary information of table 11.1.3;

3. Update critical components list for alternate sources including revising the typo on X-Cap (C1) and adding alternative source (RA series/Murata Mfg. Co. Ltd.) for Y-Cap;

4. Slightly increase 1.5 mm and 1 mm for case and studs respectively for model with suffix /EF;

5. The DC fan wire moves from the outside of the insulator to the inside of the insulator for model with suffix /EF;

6. Minor change in the circuitry as well as PCB layout;

7. Update the definition of variables(s) of models name;

8. Change Applicant and Manufacturer from WUXI TDK-LAMBDA ELECTRONICS CO LTD to TDK-Lambda (China) Electronics Co., Ltd.

9. Add new models serials: CUS600M1-zxxxxxx, CME600A1-zxxxxxx (y=1) which are the modified models based on CUS600My-zxxxxxx, CME600Ay-zxxxxxx (y=blank) except for minor variations in the circuitry and markings as below. Model CME600Ay-zxxxxxx is identical to model CUS600My-zxxxxxx except for model name designation;

10. Update the insulation diagram Table.

For models: CUS600M1-zxxxxxx, CME600A1-zxxxxxx (y=1);

1. All of derating curve are as same as CUS600M & CME600A series;

2. Cause the schematic has some differences which didn't impact the product safety, CUS600M1 &

CME600A1 series' PCB has minor differences from CUS600M & CME600A series';

3. CUS600M1 & CME600A1 series are no standby power circuit;

4. T2 is different model from the transformer in CUS600My-zxxxxxx, CME600Ay-zxxxxxxx, and it is located on the primary circuit. The output of T2 is intended to supply the internal chip, and its 12Vdc output can supply the Fan which is specific to model with suffix /EF. The Fan EFB0412HHD is identical with EFB0405HHD of CUS600My-zxxxxxx, CME600Ay-zxxxxxxx (y=blank) in CFM.

Technical Considerations

- The product was investigated to the following additional standards: N/A
- The following additional investigations were conducted: N/A
- The product was not investigated to the following standards or clauses: Electromagnetic Compatibility (IEC 60601-1-2), Clause 14 Programmable Electronic Systems, Biocompatibility (ISO 10993-1), Risk Management (ISO 14971)
- The following accessories were investigated for use with the product: N/A
- This PSU subject to this evaluation is not a medical device or system on its own right, but a component intended for building into such. Risk assessment was therefore not subject of this investigation. It shall be carried out for final medical electrical equipment or system. The insulation system of the PSU was evaluated for compliance with the MEANS OF PATIENT PROTECTION (MOPP). Compliance with IEC / EN 60601-1-2 shall be evaluated during the end system evaluation. The product is for building-in equipment, the overall compliance shall be investigated in the complete medical electrical equipment system, particular: or in Fire enclosure Mechanical enclosure Electrical enclosure Some components are pre-certified, which have been evaluated according to the relevant requirements of IEC 60601-1, employed in this product. are The equipment does not have circuits for direct connection to the patient and not is intended for use in the presence of flammable anesthetic mixtures with air, oxygen or nitrous oxide. The input circuit includes one fuse (F1A) in the Line conductor and the other fuse (F1B) is optional in neutral conductor. Consideration shall be given in the end-use product regarding addition of the second fuse having the same or better characteristics in order to comply with fusing requirements of Clause 8.11.5 of the standard. The metal enclosure of Class II equipment should be evaluated by end system.

The others see Enclosed Miscellaneous-Recommend by manufacturer

Engineering Conditions of Acceptability

For use only in or with complete equipment where the acceptability of the combination is determined by UL LLC. When installed in an end-product, consideration must be given to the following:

 Scope of Power Supply evaluation defers the following clauses to be determined as part of the end product investigation:

Clause 7.2.7 ELECTRICAL INPUT POWER FROM THE SUPPLY MINS,

Clause 7.5 SAFETY SIGNS,

Clause 7.6 SYMBOLS,

Clause 7.9 ACCOMPANYING DOCUMENTS,

Clause 9 PROTECTION AGAINST MECHANICAL HAZARDS OF ME EQUIPMENT AND ME SYSTEMS,

Clause 10 PROTECTION AGAINST UNWANTED AND EXCESSIVE RADIATION HAZARDS, Clause 12 ACCURACY OF CONTROLS AND INSTRUMENTS AND PROTECTION AGAINST HAZARDOUS OUTPUTS,

Clause 14 PROGRAMMABLE ELECTRICAL MEDICAL SYSTEMS (PEMS),

Clause 16 ME SYSTEMS,

Risk Management was excluded from this investigation

Risk Controls/ Engineering Considerations for component power supply:

For use only in or with complete equipment where the acceptability of the combination is determined by the CB Testing Laboratory, when installed in an end-product, consideration must be given to the following:For Power Supplies with No RM: End product Risk Management Process to include consideration of requirements specific to the Power Supply.

For Power Supplies with No RM: End product Risk Management Process to consider the acceptability of risk for the following components that were identified as High-Integrity Component: i.e. Fuse (F1A).

For Power Supplies with No RM: End product Risk Management Process to consider the need for simultaneous fault condition testing.

For Power Supplies with No RM: End product Risk Management Process to consider the need for different orientations of installation during testing.

For Power Supplies with No RM with Exposure Condition outside of Humidity Range: Power Supply tested in 40°C, 95%RH. End product Risk Management Process to determine risk acceptability criteria.

For Power Supplies with No RM and Insulating Materials: End product to determine the acceptability of risk in conjunction to insulation to resistance to heat, moisture, and dielectric strength. For Power Supplies with No RM: End product to determine the acceptability of risk in

conjunction to the movement of components as part of the power supply.For Power Supplies with No RM: End product to determine the acceptability of risk in conjunction to the movement of conductors as part of the power supply.

For Power Supplies with No RM: End product to determine the acceptability of risk in

conjunction to the routing of wires away from moving parts and sharp edges as part of the power supply.

For Power Supplies with No RM and Not tested with Test Corner: Temperature Test was conducted without Test Corner. End product to determine the acceptability of risk in conjunction to temperature testing without test corner as part of the power supply.For Power Supplies with No RM or Units without Cleaning/Disinfection Methods: End product to determine the acceptability of risk in conjunction to the Cleaning and Disinfection Methods as part of the power supply.

For Power Supplies with No RM or Units with Liquids: End product to determine the acceptability of risk in conjunction to the Leakage of Liquids as part of the power supply.

For Power Supplies with No RM or Units with Indicators: End product to determine the acceptability of risk in conjunction to the Arrangement of Indicators as part of the power supply.

For Power Supplies with No RM or Units with Enclosures: End product to determine the acceptability of risk in conjunction to the results of Mechanical Testing conducted as part of the power supply. For Power Supplies with No RM: End product to determine the acceptability of risk in

conjunction to the selection of components as it pertains to the intended use, essential performance, transport, storage conditions as part of the power supply.

For Power Supplies with Thermal Cut-off and No RM: End product to determine the acceptability of

risk in conjunction to the use of Thermal Cut-off and Overcurrent releases as part of the power supply.

For Power Supplies with Pre-set components and No RM: End product to determine the acceptability of risk in conjunction to the use of Pre-set controls as part of the power supply.