Issued Date: 2011-10-12 Page 1 of 26 Report Reference E252331-V2-S3

Revised Date: 2013-10-31

## UL TEST REPORT AND PROCEDURE

Standard: ANSI/AAMI ES60601-1:2005 (Medical Electrical Equipment – Part 1:

General Requirements for Basic Safety and Essential Performance); CSA C22.2 No. 60601-1:08 (Medical Electrical Equipment – Part 1: General Requirements for Basic Safety and Essential Performance)

Certification Type: Medical Electrical Equipment

CCN: QQHM2, QQHM8

Product: Switching Power Supply

Model: CFM351MXXX (XXX can be 050,120, 240,

480 to denote different output rating)

Rating: I/P: 100-240Vac, 50-60Hz, 5.0-2.0A

O/P:

Model Name Output Rating

CFM351M050 DC 5V/60A, 5V/0.3A, 12V/0.3A; CFM351M120 DC 12V/29.2A, 5V/0.3A, 12V/0.3A; CFM351M240 DC 24V/14.6A, 5V/0.3A, 12V/0.3A; CFM351M480 DC 48V/7.3A, 5V/0.3A, 12V/0.3A;

CFM351M120-02 DC 12V/29.2A, 5V/2A

Applicant Name and Address: CINCON ELECTRONICS CO LTD

8-1 FU KUNG RD FU HSING PARK FU HSING HSIANG

CHANGHUA HSIEN, 506 TAIWAN

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 Underwriters Laboratories Inc. ('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.

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