Underwriters Laboratories (UL LLC)
Safety Certification (Manufacturing Factory) Report

Model:	CFM21Mx-Y (where x can be 050, 090, 120, 150, 240; -Y can be blank, -E, -T or -S)
Device Description:	Component Medical Power Supply
Applicant:	CINCON ELECTRONICS CO LTD 8-1 FU KUNG RD, FU HSING PARK, FU HSING HSIANG, CHANGHUA HSIEN, 506, TAIWAN
Manufacturer:	Same as Applicant
Manufacturing Facility(ies):	DONGGUAN CINCON ELECTRONICS LTD DONGGUAN CINCON ELECTRONICS LTD, 1 JING XIANG RD DONGCHENG, FOREIGN TRADE INDUSTRIAL PARK, ZHUSHAN DONGCHENG DISTRICT, DONGGUAN, GUANGDONG 523128, CHINA CINCON ELECTRONICS CO LTD 8-1 FU KUNG RD, FU HSING PARK, FU HSING HSIANG, CHANGHUA HSIEN, 506, TAIWAN
Report No.:	E252331-D1005-1/A0/C0-(M)
Report (Re)Issue Date:	2017-09-20
Base Standard(s):	ANSI/AAMI ES60601-1 (2005/(R)2012 + A1:2012, C1:2009/(R)2012 + A2:2010/(R)2012) - Amendment 1 - Revision Date 2012/08/21 CAN/CSA-C22.2 No. 60601-1:14 - Edition 3 - Revision Date 2014/03
Additional Standards:	N/A
Report Types:	This report consists of the following report types: [Yes] US Certification (UL Recognition) [Yes] CAN Certification (cUL Recognition)

This report covers the Safety evaluation of the referenced model(s) according to the standard(s) specified above.

This is the Manufacturing Factory report only, which is used as part of the factory FUS inspections.