

<b>IEC 60601-1</b> <b>Medical electrical equipment</b> <b>Part 1: General requirements for basic safety and essential performance</b>	
Report Reference No.....:	RL/2016/A0046
Tested by.....:	Betty Tsai Engineer
Approved by.....:	Charles Chen Reviewer
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Total number of pages.....:	157
Testing Laboratory.....:	SGS Taiwan Ltd., Safety Laboratory
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Applicant's name.....:	Phihong Technology Co., Ltd.
Address.....:	No. 568, Fuxing 3rd Rd., Guishan District, Taoyuan City, Taiwan
<b>Test specification:</b>	
Standard.....:	IEC 60601-1: 2005 + CORR. 1 (2006) + CORR. 2 (2007) + AM1 (2012) + CORR.1 (2014) or IEC 60601-1: 2012 + CORR.1 (2014), EN 60601-1: 2006 + A1: 2013 + A12: 2014 ANSI/AAMI ES60601-1:2005 (R2012) + C1(2009) +A1(2012) +A2(2010), CAN/CSA C22.2 No. 60601-1:14.
Test procedure.....:	V.o.C Scheme
Non-standard test method.....:	None
Test Report Form No.....:	IEC60601_1K
Test Report Form Originator.....:	UL(US)
Master TRF.....:	2015-11
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Test item description.....:	Switching Power Supply
Trade Mark.....:	
Manufacturer.....:	Same as applicant
Model/Type reference.....:	PMA10R-050A, PMA10R-120A, PMA10R-240A, PMA09R-060A, PMA09R-090A

Unless otherwise stated the results shown in this test report refer only to the sample(s) tested and such sample(s) are retained for 90 days only.

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