ATTESTATION OF CONFORMITY

Directive(s):	*2014/30/EU & 93/42/EEC*
Attestation No.:	*SECM1707216*
Product / Test Item:	*Medical Type Switching Power Supply*
Model / Type Reference:	*RPTG-160X(X=A,B,C,D)-C* *RPT-160X(X=A,B,C,D)-C*

The submitted sample(s) have been tested with the following standard(s) and found to be in compliance with the essential requirements of the Directive(s):

Standard(s)	
EN 55011 : 2009+A1:2010 (Class B)	*EN 60601-1-2 : 2015*
(CISPR 11: 2009+ A1: 2010)	*EN 61000-4-2 : 2009*
EN 61000-3-2 : 2014	*EN 61000-4-3 : 2006+A1:2008+A2:2010*
EN 61000-3-3 : 2013	*EN 61000-4-4 : 2012*
	EN 61000-4-5 : 2014
	EN 61000-4-6 : 2014
	EN 61000-4-8 : 2010
	EN 61000-4-11 : 2004

The referred test report(s) show that the product fulfills the essential requirements set out in the Directive(s). On this basis, together with the manufacturer's own documented production control, the manufacturer or his European authorized representative can in his EC Declaration of Conformity verify compliance with the Directive(s). The CE marking could be affixed only when all the relevant and effective EC Directives are complied with.

CE

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