ATTESTATION OF CONFORMITY

Directive(s):	*2004/108/EC & 2014/30/EU*
Attestation No.:	*SECM1601003*
Applicant / Holder:	*GlobTek, Inc.*
Address:	*186 Veterans Dr. Northvale, NJ 07647, USA*
Product / Test Item:	*MEDICIAL POWER SUPPLY*
Model / Type Reference:	*GT*41078-*05-USB（The 1st "*" part can be 'M' or '-' or 'H' for market identification and not related to safety. The 2nd "*" part denotes the rated output wattage designation, which can be "01" to "06", with interval of 1.）*

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 60601-1-2 : 2007 (EN 55011: 2009+ A1: 2010)	EN 60601-1-2 : 2007
EN 61000-3-2 : 2014	IEC 61000-4-2 : 2008
EN 61000-3-3 : 2013	IEC 61000-4-3 : 2006+A1:2007+A2:2010
	IEC 61000-4-4 : 2012
	IEC 61000-4-5 : 2005
	IEC 61000-4-6 : 2008
	IEC 61000-4-8 : 2009
	IEC 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

Miro Chueh / Manager
2016-01-19

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