Declaration of Conformity

According to ISO/IEC Guide 22 and EN 45014.

Mascot Electronics A/S
Postal address: P.O. Box 177
N-1601 Fredrikstad - Norway

Visiting Address: Mosseveien 109
N-1624 Gressvik - Norway

Phone: +47 69 36 43 00
Fax: +47 69 36 43 01
E-mail: sales@mascot.no
Internet: www.mascot.no

We, ***Mascot Electronics A/S***
declare under our sole responsibility that the products:

Desktop Power Supply ***(also for Medical Equipment)***

Type: 2924 ***(with 2-pins IEC 60320/C8 AC-inlet for detachable mains cord set)***

Type: 2925 ***(with non-detachable mains cord) (May also be delivered in a special version protected against ingress of objects and water according to IP67, standard IEC 60529)***

Direct Plug-In Power Supply ***(also for Medical Equipment)***

Type: 2926 ***(with 2-pins IEC 60320/C8 AC-inlet and exchangeable mains plugs (EU/UK/US/AUS))***

Data: Input:	***0.35A 100 - 240VAC 50-60 Hz, Cl. II***		
Output:	***6V-version:***	***4.5 - 7.5VDC***	***max. 1.5A/9W***
	9V-version:	***7.5 - 11VDC***	***max. 1.5A/13.5W***
	12V-version:	***11 - 15VDC***	***max. 1.2A/15W***
	16V-version:	***15 - 18VDC***	***max. 1.0A/16W***
	24V-version:	***20 - 30VDC***	***max. 650mA/16W***
	48V-version:	***40 - 52VDC***	***max. 330mA/16W***

are in conformity with the following standards or other normative documents:

Electrical Safety:

EN 60950-1	***(EN 60950-1:2006 +/A11:2009, /A1:2010, /A12:2011 & /A2:2013)***	***(IT-equipment)***
EN 60601-1	***(EN 60601-1:2006 +/AC:2010)***	***(Medical electrical equipment, 3rd Ed.)***

Electromagnetic Compatibility (EMC):

EN 61000-6-1	***(EN 61000-6-1:2007)***	***(Immunity-residential, commercial & light-industrial environment)***
EN 61000-6-3	***(EN 61000-6-3:2007 & /A1:2011)***	***(Emission-residential, commercial & light-industrial environment)***
EN 55022	***(EN 55022:2010 & /AC:2011)***	***(Emission-IT-Equipment)***
EN 55024	***(EN 55024:2010)***	***(Immunity-IT-Equipment)***
EN 60601-1-2	***(EN 60601-1-2:2007 +/AC:2010)***	***(Medical equipment, EMC - Requirements and tests)***

following the provisions of EU-Directives:

2006/95/EC	***(repealing 73/23/EEC)***	***(Low Voltage Directive, LVD)***
2004/108/EC	***(repealing 89/336/EEC)***	***(EMC Directive)***
93/42/EEC		***(General Medical Devices Directive)***
2009/125/EC	***(repealing 2005/32/EC & 2008/28/EC)***	***(Energy-related Products Directive, ErP)***
2011/65/EU	***(repealing 2002/95/EC & 2008/35/EC)***	***(Restriction on use of Hazardous Substances in EEE, RoHS2)***

and are produced under a quality system certified to standard EN 29001:2008 (ISO 9001:2008).

Place of issue:
Fredrikstad, Norway

Date of issue:
25 January 2013

Finn-Erik Wallin
Compliance Manager