

# Declaration of Conformity

According to ISO/IEC Guide 22 and EN 45014.

Mascot Electronics A/S

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We, **Mascot Electronics A/S**

declare under our sole responsibility that the products:

**Battery Charger for Lead-Acid or Li-Ion-Batteries (also for use with Medical Equipment)**

**Type: 2240** (with detachable mains cord / 2-pins IEC 60320 inlet or with non-detachable mains cord)  
(Version protected against ingress according to IP67, standard EN 60529, is available)

**Type: 2241** (with exchangeable mains plug-adapters or detachable mains cord)

**Data: Input: max.0.35A 100-240VAC 50-60 Hz, Class II**

**Versions for Lead-Acid Batteries:**

<b>Output: 6V-version:</b>	<b>7.35VDC</b>	<b>max. 1.3A</b>
<b>12V-version:</b>	<b>14.7VDC</b>	<b>max. 1.0A</b>
<b>24V-version:</b>	<b>29.4VDC</b>	<b>max. 0.5A</b>
<b>36V-version:</b>	<b>44.1VDC</b>	<b>max. 0.35A</b>

**Versions for Li-Ion Batteries:**

<b>Output: 1-cells version:</b>	<b>4.1 or 4.2VDC</b>	<b>max. 1.3A</b>
<b>2-cells version:</b>	<b>8.2 or 8.4VDC</b>	<b>max. 1.3A</b>
<b>3-cells version:</b>	<b>12.3 or 12.6VDC</b>	<b>max. 1.2A</b>
<b>4-cells version:</b>	<b>16.4 or 16.8VDC</b>	<b>max. 0.9A</b>
<b>5-cells version:</b>	<b>20.5 or 21.0VDC</b>	<b>max. 0.75A</b>
<b>6-cells version:</b>	<b>24.6 or 25.2VDC</b>	<b>max. 0.65A</b>
<b>7-cells version:</b>	<b>28.7 or 29.4VDC</b>	<b>max. 0.56A</b>

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

**Electrical Safety:**

<b>EN 60950-1</b>	(EN 60950-1:2006 +/A2:2013)	(IT-equipment)
<b>EN 60601-1</b>	(EN 60601-1:2006 +/AC:2010)	(Medical electrical equipment, 3 <sup>rd</sup> Ed.)

**Electromagnetic Compatibility (EMC):**

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

following the provisions of EU-Directives:

<b>2014/35/EU</b>	(repealing 2006/95/EC & 73/23/EEC)	(Low Voltage Directive, LVD, recast)
<b>2014/30/EU</b>	(repealing 2004/108/EC & 89/336/EEC)	(EMC Directive, recast)
<b>93/42/EEC</b>		(General Medical Devices Directive)
<b>2009/125/EC</b>	(repealing 2005/32/EC & 2008/28/EC)	(Energy-related Products Directive, ErP)
<b>2011/65/EU</b>	(repealing 2002/95/EC & 2008/35/EC)	(Restriction on use of Hazardous Substances in EEE, RoHS2)

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

Place of issue:  
**Fredrikstad, Norway**

Date of issue:  
**03 October 2014**

  
**Finn-Erik Wallin**  
Product Compliance Manager