# **EU Declaration of Conformity**



We, the responsible manufacturer;				
Company Name:	Mascot Electronics AS			
Postal Address:	P.O.Box 177, N-1601 Fredrikstad, NORWAY			
Visiting Address:	Mosseveien 109, N-1624 Gressvik, NORWAY			
Telephone:	(+47) 69 36 43 00 E-mail: sales@mascot.com WEB: www.mascot.com			
declare that this Declaration is issued under our sole responsibility and belongs to the following product(s):Product andBattery Charger for 1 cell Li-Ion Batteriesintended purpose:				
Brand(s):	and/or managed (may also carry additional customer name, logo or trade mark)			
Type(s)/Model(s)/ UDI-DI:	3845 (may also carry additional customer model name or part number)			
Batch / Serial No./ UDI-PI:	all CE-marked products produced from the date indicated below (for production date: see marking on the product)			
Description:	Input: max. 0.25 A, 100-240 VAC 50-60 Hz, Class II Output: 4.2 VDC, max. 1.5 A / 6.3 W			

The product(s) desci legislation:	e product(s) described above are in conformity with the relevant European Union harmonisation gislation:		
2014/35/EU	EU Directive - Safety of electrical equipment ("Low-Voltage Directive") (LVD) recast, repealing Directives 2006/95/EC & 73/23/EEC		
2014/30/EU	EU Directive - Electromagnetic Compatibility (EMC) recast, repealing Directives 2004/108/EC & 89/336/EEC		
93/42/EEC	EU Directive - General Medical Devices (MDD), Risk Class I Device will from 05.05.2020 be repealed by Regulation (EU) 2017/745		
2009/125/EC	EU Directive - Energy Related Products, Ecodesign (ERP) recast, repealing Directive 2005/32/EC (EUP)		
2015/863/EU	EU Directive - Restriction on use of Hazardous Substances in EEE ("RoHS3") recast, repealing Directives 2002/95/EC, 2008/35/EC & 2011/65/EU		

The following harmonised standards and technical specifications have been applied: (International editions and comments indicated in brackets)

## Electrical Safety (to LVD- & MDD-Directives):

EN 60950-1	EN 60950-1:2006 + /A1:2010, + /A11:2009, + /AC:2011, + /A12:2011 + /A2:2 (IEC 60950-1:2005 modified + /A1:2009 modified + /A2:2013 modified, Editi (will from 20.06.2019 be replaced by standard EN 62368-1:2014 + /AC:2015, (IEC 62368-1:2014, Edition 2.0)	ion 2.2)
EN 60335-1	EN 60335-1:2012 + /AC:2014 + /A11:2014 Household and similar ap (IEC 60335-1:2010 modified, Edition 5.0)(also IEC 60335-1:2010 modified + /	pliances-General requirements, Edition 5.0 /A1:2013 + /A2:2016, Edition 5.2)
EN 60335-2-29	EN 60335-2-29:2004 + /A2:2010 Household and similar appliances-Req (IEC 60335-2-29:2002 + /A1:2004 + /A2:2009, Edition 4.2) (also IEC 60335-2-	uirements for battery chargers, Edition 4.2 -29:2016, Edition 5.0)
EN 60601-1	EN 60601-1:2006 + /AC:2010 +/A1:2013 (IEC 60601-1:2005 + /A1:2012)	Medical electrical equipment, Edition 3.1

# **EU Declaration of Conformity**



## Electromagnetic Compatibility (to EMC- & MDD-Directives):

EN 61000-6-1	EN 61000-6-1:2007 Immunity-residential, comm. & light-industrial environment, Edition (IEC 61000-6-1:2005, Edition 2.0) (also IEC 61000-6-1:2016, Edition 3.0, not yet an EN-norm)
EN 61000-6-3	EN 61000-6-3:2007 + /A1:2011 & /AC:2012 Emission-residential, comm. & light-industrial environment, Edition (IEC 61000-6-3:2007 + /A1:2010)
EN 55014-1	EN 55014-1:2006 + /A1:2009 & /A2:2011 Emission-household appliances, Edition (CISPR 14-1:2005 + /A1:2008 & /A2:2011, Edition 5.2) (also CISPR 14-1:2016, Edition 6.0, but not yet an EN-norm)
EN 55014-2	EN 55014-2:1997 + /AC:1997, /A1:2001, /A2:2008 Immunity-household appliances, Edition (CISPR 14-2:1997 + /A1:2001 & /A2:2008, Edition 1.2) ( <i>also CISPR 14-2:2015, Edition 2.0, but not yet an EN-norm</i> )
EN 55022	EN 55022:2010 + /AC:2011 Emission-IT-Equipment, Edition (CISPR 22:2008 modified, Edition 6.0)( <i>Note: CISPR 22 is now replaced by CISPR 32:2012</i> )
EN 55024	EN 55024:2010 Immunity-IT-Equipment, Edition (CISPR 24:2010, Edition 2.0) (also CISPR 24:2010 + /Corr.1:2011 + /A1:2015, Edition 2.1, but not yet an EN-norm)
EN 55032	EN 55032:2012 + /AC:2013 (CISPR 32:2012 + /Corr.1:2012 + /Corr 2:2012, Edition 1.0) (also CISPR 32:2015, Edition 2.0, but not yet an EN-norm
EN 60601-1-2	EN 60601-1-2:2007 Medical equipment, EMC - Requirements and tests, Edition from 31/12/2018: EN 60601-1-2:2015 Medical equipment, EMC - Requirements and tests, Edition (IEC 60601-1-2:2007 modified, Edition 3.0)( <i>Note: for IEC: Edition 3.0 is replaced by IEC 60601-1-2:2014, Edition 4.0</i> )

#### Ecodesign (to ERP-Directive):

Commission Regulation (EC) No 278/2009

implementing Directive 2005/32/EC with regard to ecodesign requirements for noload condition electric power consumption and average active efficiency of external power supplies (*Note: not applicable to Battery Chargers, ref. Article 1.2 item c*) )

### Additional Information:

Compliance with harmonised standards and technical specifications may have been verified by the manufacturer, by third party testing or by a Certification Body (NCB).

The products are considered Risk Class I devices according to the General Medical Devices Directive.

The product(s) may be produced at production sites (for specific product: see "Made in"-marking on the product):

	the free model and free managements and free model and free model and free model and free model.					
	Mascot Electronics AS	Mascot Baltic OÜ	Mascot Power Supplies (Ningbo) Co.,Ltd			
	P.O.Box 177,	Taevakivi 15	No.128 Jinchuan Road, Zhenhai			
	N-1601 Fredrikstad,	EE-13619 Tallinn	Ningbo 315221			
	NORWAY	ESTONIA	CHINA			
The production sites are certified to standard EN 29001:2015 (ISO 9001:2015):						
	Mascot Electronics AS:	Mascot Baltic OÜ:	Mascot Power Supplies (Ningbo) Co.,Ltd:			
	Kiwa Teknologisk Institutt	Metrosert	DNV-GL			
	certificate ref. 044	certificate ref. K-144	certificate ref. 179027-2015			

The most recent issue of this Declaration is available at www.mascot.com.

Fredrikstad, Norway

2018-06-25

Place of issue

Date of issue

## Signed on behalf of Mascot Electronics AS

Finn-Erik Wallin, Compliance Manager

Name, function, signature