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Bellman & Symfon®
DESIGN FOR EARS

EC DECLARATION OF CONFORMITY

The undersigned, representing the following manufacturer.

Name:	**Bellman & Symfon AB**
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Herewith declares that the products

Type of equipment	Brand name	Model	Conformity with the following EC/EU directives. Table 1.	Standard used to prove conformity. Table 2.
Light My Way System	Bellman & Symfon	BE8108	A, B, C, D, E, F, I	1,2,3,4,5,6,7,8,9,13,14,15,16,17, 18,19,20,21,22
Light My Way System	Bellman & Symfon	BE8107	A, B, C, D, E, F, I	1,2,3,4,5,6,7,8,9,13,14,15,16,17,18, 19,20,21,22
Bed Exit Sensor	Bellman & Symfon	BE2500	B, C, E, F, I	1,2,3,4,5
Bedroom LED Light Bulb	Bellman & Symfon	BE2510	A, B, C, E, F, G, H, I	1,2,3,4,5,6,7,8,9
Hall LED Light Bulb	Bellman & Symfon	BE2511	A, B, C, E, F, G, H, I	1,2,3,4,5,6,7,8,9
Bathroom LED Light Bulb	Bellman & Symfon	BE2520	A, B, C, E, F, G, H, I	1,2,3,4,5,6,7,8,9
Bedroom Switch	Bellman & Symfon	BE2530	B, C, E, F, I	1,2,3,4,5
Bathroom Switch	Bellman & Symfon	BE2540	B, C, E, F, I	1,2,3,4,5
Bulb Socket Adaptor E14-E27	Bellman & Symfon	BE9278	A, B, E, F, I	10,11,12
Bulb Socket Adaptor B22d-E27	Bellman & Symfon	BE9279	A, B, E, F, I	10,11,12

Is in conformity with the provisions of the following EEC/EU directives and applicable essential requirements set out in related directives.

No	Reference No:	Title:
A	2014/35/EU	Low Voltage Directive (LVD)
B	2014/30/EU	Electromagnetic Compatibility (EMC Directive)
C	2014/53/EU	Radio Equipment Directive (RED)
D	93/42/EEC	Medical Devices Directive (MDD)
E	2015/863/EU	Restriction of the use of certain hazardous substances in electrical and electronic equipment (RoHS Directive)

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F	**EU Regulation (EC) No 1907/2006**	**Registration, Evaluation, Authorization and Restriction of Chemicals (REACH)**
G	**Directive 2009/125/EC**	**ERP ECODESIGN** **Regulation EC 244/2009, ecodesign requirements for non-directional household lamps;** **Regulation EC 859/2009, ecodesign requirements on ultraviolet radiation of non-directional household lamps;** **Regulation EU 1194/2012, ecodesign requirements for directional lamps, light emitting diode lamps and related equipment**
H	**Directive 2010/30/EC**	**Energy labelling of household** **Regulation 874/2012, energy labelling of electrical lamps and luminaires**
I	**Directive 2012/19/EU**	**WEEE Waste electrical and electronic equipment**

Table 1. List of EC directives

This declaration of conformity is issued under the sole responsibility of the manufacturer, and that related applicable harmonized standards and/or technical specifications referenced overleaf have been applied.

Bellman & Symfon AB declares herby that the products above conform to applicable requirements in the Swedish Act (1993:584), Ordinance (SFS 1993:876), and regulation, LVFS 2003:11 regarding medical device provision form Medical Products Agency MPA - Läkemedelsverket.

Gothenburg, Sweden; Jul 21 ,2019

Vicky 2019.7.21

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CTO, Bellman & Symfon AB

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Relevant harmonised standards used or specifications in relation to conformity:

No.	Standard No. and edition/year	Title
1	EN 301 489-1 V2.2.0;	Electromagnetic compatibility and Radio spectrum Matters (ERM); ElectroMagnetic Compatibility (EMC) standard for radio equipment and services; Part 1: Common technical requirements
2	EN 301489-17 V3.2.0	ElectroMagnetic Compatibility (EMC) standard for radio equipment and services; Part 17: Specific conditions for Broadband Data Transmission Systems; Harmonised Standard covering the essential requirements of article 3.1(b) of the Directive 2014/53/EU
3	EN 300 328 V2.1.1	Wideband transmission systems; Data transmission equipment operating in the 2,4 GHz ISM band and using wide band modulation techniques; Harmonised Standard covering the essential requirements of article 3.2 of Directive 2014/53/EU
4	EN 62479:2010	Assessment of the compliance of low power electronic and electrical equipment with the basic restrictions related to human exposure to electromagnetic fields (10 MHz to 300 GHz)
5	EN 60950-1:2006 + A11:2009 + A1:2010 + A12:2011 + A2:2013	Information technology equipment - Safety Part 1: General requirements
6	EN 62560:2012 + A1:2015 + AC:2015	Self-ballasted LED-lamps for general lighting services by voltage > 50 V — Safety specifications
7	EN 62493:2015	Assessment of lighting equipment related to human exposure to electromagnetic fields
8	EN 62031:2008 + A1:2013 + A2:2015	LED modules for general lighting — Safety specifications
9	EN 62471:2008	Photobiological safety of lamps and lamp systems
10	EN 60238:2004	Edison screw lampholders
11	EN 60061-1:2017	Lamp caps and holders together with gauges for the control of interchangeability and safety - Part 1: Lamp caps
12	EN 60061-3:2017	Lamp caps and holders together with gauges for the control of interchangeability and safety - Part 3: Gauges
13	EN 61547:2009	Equipment for general lighting purposes - EMC immunity requirements
14	EN 61000-3-2:2014	Electromagnetic compatibility (EMC) - Part 3-2: Limits - Limits for harmonic current emissions (equipment input current ≤ 16 A per phase)
15	EN 61000-3-3:2013	Electromagnetic compatibility (EMC) - Part 3-3: Limits - Limitation of voltage changes, voltage fluctuations and flicker in public low-voltage supply systems, for equipment with rated current <= 16 A per phase and not subject to conditional connection
16	EN 55015:2013+A1:2015	Limits and methods of measurement of radio disturbance characteristics of electrical lighting and similar equipment
17	EN 60601-1:2006 + A1:2013	Medical electrical equipment – Part 1: General requirements for basic safety and essential performance
18	EN 60601-1-2:2015	Medical electrical equipment – Part 1-11: General requirements for basic safety and essential performance – Collateral Standard: Requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment
19	EN ISO 13485:2016	Medical devices — Quality management systems — Requirements for regulatory purposes
20	EN ISO 15223-1:2016	Medical devices — Symbols to be used with medical device labels, labelling and information to be supplied — Part 1: General requirements
21	EN 1041:2008	Information supplied by the manufacturer of medical devices
22	EN ISO 14971:2012	Medical devices — Application of risk management to medical devices

Table 2. List of standards