

Declaration of Conformity

Manufacturer Guangdong Biolight Meditech Co., Ltd.
Address No.2 Innovation First Road, Technical Innovation Coast, Hi-tech Zone, Zhuhai, P.R. China
European Representative Shanghai International Holding Corp. GmbH (Europe)
Eiffestraße 80, 20537 Hamburg Germany
Product Electronic Sphygmomanometer
UMDNS Code 16157
Model Code WBP301/WBP302

Classification: Class IIa , rule 10 of Annex IX of the MDD 93/42/EEC

Conformity Assessment Route: Annex II without chapter 4 of the MDD 93/42/EEC

We herewith declare that the above mentioned products meet the transposition into national law, the provisions of the following EC Council Directives and Standards. We are exclusively responsible for this DoC. All supporting documentations are retained under the premises of the manufacturer.

DIRECTIVES

General applicable directives:

Medical Device Directive: COUNCIL DIRECTIVE 93/42/EEC concerning medical devices (MDD 93/42/EEC).

Standard applied:

See the appendix.

Notified Body: TÜV SÜD Product Service GmbH, Ridlerstr. 65, 80339 München, Germany

Identification number: 0123

(EC) Certificate(s): G10499570033 Rev.02

Expire date of the Certificate: 2024-05-26

Start of CE marking: 2017-11-09

Place, Date of Issue: Zhuhai, China. 2022-04-15

Signature Jin Liang

Name Jin Liang

Position Chief Engineer

APPENDIX

Standards for WBP301/WBP302 Electronic Sphygmomanometer

Item	Scope	Number of standard	Name of standard
1	General, Safety	IEC 60601-1:2005+A1:2012	Medical electrical equipment – Part 1: General requirements for basic safety and essential performance.
2	General, EMC	IEC 60601-1-2:2014	Medical electrical equipment – Part 1-2: General requirements for basic safety and essential performance- Collateral standard: Electromagnetic disturbances - Requirements and tests
3	General, Usability	IEC 60601-1-6:2010+A1:2013	Medical electrical equipment – Part 1-6: General requirements for basic safety and essential performance – Collateral standard: Usability.
4	General, home healthcare	IEC 60601-1-11:2015	Medical electrical equipment – Part1-11: 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
5	General, Software	IEC 62304:2006+A1:2015	Medical device software – Software life cycle processes.
6	Risk management	EN ISO 14971:2012	Medical devices- Application of risk management to medical devices
7	Labeling	ISO 15223-1:2016	Medical devices-Symbols to be used with medical device labels, labeling and information to be supplied--Part 1: General requirements
8	Biological evaluation	ISO 10993-1: 2009	Biological evaluation of medical devices – Part 1: Evaluation and testing within a risk management process.
9	Biological evaluation	ISO 10993-5:2009	Biological evaluation of medical devices—Part 5: Tests for in vitro cytotoxicity
10	Biological evaluation	ISO 10993-10:2010	Biological evaluation of medical devices--Part 10:Tests for irritation and skin sensitization
11	General, NIBP	IEC 80601-2-30:2009 +A1:2013	Medical electrical equipment – Part 2-30: Particular requirements for the basic safety and essential performance of automated non-invasive sphygmomanometers

12	General, NIBP	ISO 81060-2:2013	Non-invasive sphygmomanometers — Part 2: Clinical investigation of automated measurement type
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