

Declaration of Conformity to Council Directive 93/42/EEC concerning Medical Devices

Manufacturer: Beijing Choice Electronic Technology Co., Ltd.
Room 4104, No. A12 Yuquan Road, Haidian District,
100143 Beijing, PEOPLE'S REPUBLIC OF CHINA.

European Representative: Shanghai International Holding Corp. GmbH (Europe)
Eiffestraße 80 20537 Hamburg GERMANY

Product Name: Fingertip Pulse Oximeter

Product Model: MD300C23

UMDNS Code: 17148

Classification: Class IIa, rule 10 to Annex IX of the MDD

Conformity assessment Route: Annex II excluding (4)

We, the manufacturer, herewith declare that the stated medical devices
meet the transposition into national law, the provisions of Council Directive
93/42/EEC concerning medical devices.

All supporting documentation is retained at the premises of the manufacturer.

Standards applied:

EN ISO 13485:2016/AC:2016 Medical devices- Quality management systems-
Requirements for regulatory purposes

EN ISO14971:2012 Medical devices - Application of risk management to medical devices

EN 60601-1:2006/A1:2013 Medical electrical equipment-Part 1: General requirements for
safety

EN 60601-1-2:2015 Medical electrical equipment Part 1-2: General requirements for basic
safety and essential performance - Collateral Standard: Electromagnetic disturbances -
Requirements and tests

EN 60601-1-6:2010 Medical electrical equipment -- Part 1-6: General requirements for
basic safety and essential performance - Collateral standard: Usability

EN 60601-1-11:2010 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.

ISO 80601-2-61:2011 Medical electrical equipment —Part 2-61: Particular requirements for basic safety and essential performance of pulse oximeter equipment

EN ISO10993-1:2009/AC:2010 Biological evaluation of medical devices - Part 1: Evaluation and testing within a risk management process

EN ISO10993-5:2009 Biological evaluation of medical devices - Part 5: Tests for in vitro cytotoxicity

ISO10993-10:2010 Biological evaluation of medical devices - Part 10: Tests for irritation and skin sensitization

EN1041:2008 Information supplied by the manufacture of medical devices

EN ISO15223-1:2016 Medical devices - Symbols to be used with medical device labels, labelling and information to be supplied - Part 1: General requirements

EN 62304:2006/AC:2008 Medical device software-Software life-cycle processes

MEDDEV 2.7/1: 2016 CLINICAL EVALUATION: A GUIDE FOR MANUFACTURERS AND NOTIFIED BODIES UNDER DIRECTIVES 93/42/EEC and 90/385/EEC

Notified Body: TÜV SÜD Product service GmbH
Ridlerstr 65, D-80339 München, Germany

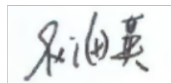
Identification Number:  0123

(EC) Certificate(s): No. G1 078179 0032 Rev.01

Start of CE-marking: 2016-05-06

Place, Date of Declaration: Beijing, 2019-05-22

Signature:



Name: Haiying Zhao

Position: Quality Director