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 REPUBILIC OF CHINA.
European Representative:	Shanghai International Holding Corp. GmbH (Europe) Eiffestraβe 80 20537 Hamburg GERMANY
Product Name：	Fingertip Pulse Oximeter
Product Model:	See attached list
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 13485:2012 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
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
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 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

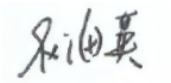
EN ISO10993-10:2010 Biological evaluation of medical devices - Part 10: Tests for irritation and skin sensitization

EN1041:2008 Information supplied by the manufacture of medical devices

EN 980:2008 Symbols for use in the labelling of medical devices

EN 62304:2006 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：	CE 0123
(EC) Certificate(s):	No. G1 17 11 78179 032
Start of CE-marking:	2016-05-06
Place, Date of Declaration:	Beijing, 2018-04-16
Signature:	
	Name: Haiying Zhao
	Position: Quality Director

Attached list

MD300CN310, MD300CN330, MD300CN340, MD300CN350, MD300CN356, MD300CN360, MD300CN130, MD300CN150, MD300CN160

MD300C1, MD300C11, MD300C12, MD300C13, MD300C15, MD300C16, MD300C17, MD300C18, MD300C19, MD300C1B, MD300C1C, MD300C1D, MD300C1E, MD300C1F

MD300C2, MD300C20, MD300C201, MD300C203, MD300C204, MD300C21, MD300C21C, MD300C22, MD300C221, MD300C23, MD300C25, MD300C26, MD300C29, MD300C2A, MD300C2B, MD300C2D, MD300C2E, MD300C2F

MD300C4, MD300C41

MD300C5, MD300C52, MD300C53, MD300C54

MD300C63, MD300C634, MD300CF3, MD300CH3