EC Declaration of Conformity

Manufacturer:	**Xuzhou Yongkang Electronic Science Technology Co., Ltd** **4F Building C8, 40 Jingshan Road, Economic and Technological Development Zone, Xuzhou, China**
European Representative:	**Prolinx GmbH** **Brehmstr. 56, 40239, Düsseldorf,GERMANY**
Product Name:	Braun Fingertip Pulse Oximeter
Models:	YK-81CEU
GMDN Code:	**17148**
UMDNS Code:	**17148**

Classification (MDD, Annex IX): **IIa，Rule 10**

Conformity Assessment Route: **Annex II(excluding section 4) and Annex VII of Directive 93/42/EEC**

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

DIRECTIVES

General applicable directives:

Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment

Medical Device Directive: COUNCIL DIRECTIVE 93/42/EEC of 14 June 1993 concerning medical devices (MDD 93/42/EEC). Amended by DIRECTIVE 2007/47/EC of 5 September 2007

Notified Body:	TÜV SÜD Product Service GmbH, Ridlerstr. 65, 80339 München, Germany
NB Identification number:	0123
(EC) Certificate(s):	G1 092582 0009 Rev.00
Expire date of the Certificate:	2024-05-26
Start of CE Marking:	2019-11-26
Place, Date of Issue:	Xuzhou, 2022-04-12

Signature:

Name: Zhao Xuecheng

Position: **General Manager**