DECLARATION OF CONFORMITY TO COUNCIL DIRECTIVE 93/42/EEC (INCLUDING DIRECTIVE 2007/47/EEC) CONCERNING MEDICAL DEVICES

MANUFACTURER: Shenzhen Creative Industry Co., Ltd.

Floor 5, BLD 9, Baiwangxin High-Tech Industrial Park,

Songbai Road, Xili Street, Nanshan District,

518110 Shenzhen, PEOPLE'S REPUBLIC OF CHINA

MEDICAL DEVICE: Patient Monitor MODEL: K10, K12, K15

CLASSIFICATION - ANNEX IX: Class IIb, Rule 10

CONFORMITY ASSESSMENT ROUTE: Annex II .3

WE, **Shenzhen Creative Industry Co., Ltd.**, Herewith declare that the stated medical devices meet the transposition into national law, the provisions of Council Directive 93/42/EEC of 14 June 1993 (Amended by Directive 2007/47/EEC) concerning medical devices; All supporting documentation are retained under the premises of the manufacturer. We are exclusively responsible for the declaration of conformity.

STANDARDS APPLIED:

ISO 13485: 2016	EN/ISO 14971: 2012	IEC 60601-1: 2005+A1: 2012
IEC 60601-1-2: 2014	IEC 60601-1-6: 2010+A1:2013	EN 60601-1-8: 2007+A1: 2013
IEC 60601-2-49: 2011	IEC 60601-2-27: 2011	IEC 80601-2-30: 2009+A1: 2013
EN/ ISO 80601-2-61: 2017	EN/ISO 80601-2-56: 2017	ISO 80601-2-55: 2011
IEC 62304: 2006+A1: 2015	ISO 10993-1: 2009	ISO 10993-5: 2009
ISO 10993-10: 2010	EN/ISO 15233-1: 2012	EN 1041: 2008+A1: 2013

NOTIFIED BODY: TÜV SÜD PRODUCT SERVICE GMBH

RIDLERSTR 65, D-80339 MÜNCHEN, GERMANY

IDENTIFICATION NUMBER 0123

(EC) CERTIFICATE(S): G1 049076 0016 Rev .02

EC REP

EUROPEAN REPRESENTATIVE: Shanghai International Holding Corp. GmbH (Europe)

Eiffestraβe 80, 20537 Hamburg, Germany

START OF CE-MARKING: OCT.15, 2010

PLACE, DATE OF DECLARATION: Floor 5, BLD 9, Baiwangxin High-Tech Industrial Park,

Songbai Road, Xili Street, Nanshan District,

518110 Shenzhen, PEOPLE'S REPUBLIC OF CHINA,

Apr. 20, 2019

SIGNATURE:

NAME: Apr. 20, 2019
POSITION: Management Representative