SUZHOU BEING MEDICAL DEVICE.CO.,LTD

DECLARATION OF CONFORMITY UE

In accordance with Annex II A - Directive 2006/42/CE
Annex IV - EMC Directive
and Annex VI - Directive 2011/65/UE (RoHS)



No. ISETC.002420200624

Manufacturer's Name : SUZHOU BEING MEDICAL DEVICE CO., LTD

Manufacturer's Address : NO. 108 GONGXIANG RD QIANDENG TOWN, KUNSHAN CHINA

Tel: +86-21-56633709

Email: JILL.SHEN@BLUEPARD.COM

Authorised Representative :Giorgio Bormac S.r.l – Via della Meccanica, 25 41012 Carpi (MO)-ITALY

Object of Declaration: : FORCED AIR INCUBATORS

This declaration of conformity is issued under the sole responsibility of the manufacturer.

Product names:

Product description FORCED AIR INCUBATORS

Model: BI-120FL, BI-120F, BI-200FL, BI-200F, BI-400FL

Serial Number: from s/n 200100001 to 2600100001

Product options: This declaration covers all options of the above products

• The object of the declaration describe above complies with the essential requirements of the following applicable European Directives, and carries the CE marking accordingly:

EMC directive: 2014/30/UE	Directive 2014/30/EU of the European Parliament and of the Council of 26 February 2014 on the harmonization of the laws of the Member States relating to electromagnetic compatibility.
RoHS Directive 2011/65/EU	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.
LVD Directive: 2014/35/UE	Directive 2014/35/EU of the European Parliament and of the Council of 26 February 2014 on the harmonization of the laws of the Member States relating to the making available on the on the market of electrical equipment designed for use within certain voltage limits Text with EEA relevance.
Machinery Directive: 2006/42/EC	DIRECTIVE 2006/42/EC OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 17 May 2006 on machinery, and amending Directive 95/16/EC (recast)

and conforms with the following standards:

EN 61010-1:2010+A1:2019

EN 61326-1:2013

EN 61000-3-2:2014

EN 61000-3-3:2013

EN 60204:2018

EN ISO 12100:2010

NAME AND ADDRESS OF THE PERSON AUTHORISED TO COMPILE THE TECHNICAL FILE $\,$

Giorgio Bormac S.r.l. - Via della Meccanica, 25 41012 Carpi (MO) - ITALY

Signed for and on behalf of: XIE WEIMIN

Place 24/09/2020

SHANGHAI SIGNATURE_





Suzhou Being Medical Device Co., Ltd

It is stated that the models we produce correspond to the following names distributed by Giorgio Bormac ltd.

Model names

Giorgio Bormac	Being
TCN-30 Plus	BO-30NL
TCN-50 Plus	BO-50NL
TCN-50 Super	BO-50N
TCN-115 Plus	BO-115NL
TCN-115 Super	BO-115N
TCN-200 Plus	BO-200NL
TCN-200 Super	BO-200N
TCF-50 Plus	BO-50FL
TCF-50 Super	BO-50F
TCF-120 Plus	BO-120FL
TCF-120 <mark>Super</mark>	BO-120F
TCF-200 Plus	BO-200FL
TCF-200 Super	BO-200F
TCF-400 Plus	BO-400FL
TCF-400 Super	BO-400F
ICN-16 Plus	BIT-16
ICN-16 Super	BI-16T
ICN-35 Plus	BIT-35
ICN-35 Super	BI-35T
ICN-55 Plus	BIT-55
ICN-55 Super	BI-55T
ICN-120 Plus	BIT-120
ICN-120 Super	BI-120T
ICN-200 Plus	BIT-200
ICN-200 Super	BI-200T
ICF-120 Plus	BI-120FL
ICF-120 Super	BI-120F
ICF-200 Plus	BI-200FL
ICF-200 Super	BI-200
ICF-400 Plus	BI-400FL
ICF-400 Super	BI-400F
IC 150-R Plus	BPC-150F
SKI 4	BSI-3
SB-35	DKZ-1
WB 12	BWS-12
WB 22	BWS-27
WB 22 Pump	BWS-27G
WB 40 Pump	BWS-40G
CB 5-10	MP-10C
CB 5-20	MP-20C
CB 5-30	MP-30C
CH 150	LHH-150SD
CH 250	LHH-250SD
WB-5	BWB-05

Place, Date 23rd Nov 2020

Manufacturer signature_

lorgio Bormac S r L signature

