

## DECLARATION OF CONFORMITY

We

Name and address of manufacturer:

**Shenzhen Brav Electronic Technologies Co.,Ltd**

4/F, Block11, Tongfuyu Industry District, Lezhujiao, Jiuwei, Xixiang, Baoan, Shenzhen, P.R.China

declare on our own responsibility that

the medical device

**Infrared Thermometer**

IT-901, IT-121, IT-121S, T-905, IT-906, IT-907, IT-122, IT-123, IT-124, IT-125,

IT-126, IT-127, IT-128, IT-129

**Digital Clinical Thermometer**

DT-121, DT-122

meets all applicable requirements of the Directive 93/42/EEC Annex I, as amended by Directive 2007/47/EC.  
The manufacturer is exclusively responsible for the declaration of conformity.

Applied standards :

EN ISO 14971:2012, EN 60601-1:2006+A1:2013, EN 60601-1-2:2015, EN ISO 10993-5:2009,  
EN ISO 10993-10:2010, EN ISO 15223-1:2016, EN 60601-1-11:2010, EN 60601-1-6:2010,  
EN 62366:2008, EN ISO 80601-2-56:2012, EN 60601-1-8:2007, EN 62304:2006, EN ISO 13485:2016

Classification:

*Classified as class **Ila** according to Annex IX,*

*Rule 4. 9.10 of the Directive 93/42/EEC.*

Conformity assessment procedure:

*Conformity assessment was performed according to  
Annex V of the Directive 93/42/EEC.*

The Authorized Representative within the  
EU who has been empowered to enter into  
commitments on our behalf is:

*MedNet GmbH*

*Borkstrasse 10*

*48163 Münster, Germany*

Notified Body:



*DNV GL Nemko Presafe AS*

*Veritasveien 3, N-1363 Høvik Postbox 116,*

*N-1300 Sandvika*

2018.11.05-2023.09.29

Validity period

Shenzhen, 2018.12.24

place, date



legally binding signature

Kay Wang (General Manager)

name and function