andon

EC Declaration of Conformity

Manufacturer:

ANDON HEALTH CO., LTD.
No.3 JinPing Street, Ya An Road,Nankai District, Tianjin, China

whose single Authorized Representative:

iHealth Labs Europe
3 rue Tronchet, 75008, Paris, France

We, the manufacturer, herewith declare that the products

Electronic Sphygmomanometers

UMDNS-Code: **16-157**;

Model: KD-931,KD-936, KD-972, BP5(ABI)

Apps of the model listed above: iHealth MyVitals, Version 1.3.5
NOTE: *KD-931,KD-936 and KD-972 are the model used in the manufacture, the trade name of KD-931 is BP3, KD-936 is BP5, KD-972 is BP7.*

meet the provisions of Directive 93/42/EEC which apply to them.

The medical device has been assigned to class IIa according to Annex IX of the Directive 93/42/EEC. It bears the mark

CE **0197**

The product concerned has been designed and manufactured under a quality management system according to Annex II of Directive 93/42/EEC.

Compliance of the designated product with the Directive 93/42/EEC has been assessed and certified by the Notified Body

TÜV Rheinland LGA Products GmbH
Tillystraße 2, 90431, Nürnberg, Germany
Certificate No.: HD 60097963 0001
Issue date: 2014-11-25
Expiry date: 2019-11-05

Following the procedure relating to the EC Declaration of Conformity set out in Annex II of Directive 93/42/EEC.

This Declaration of conformity is valid in connection with the release document for the respective batch of produced devices.

The above mentioned declaration of conformity is exclusively under the responsibility of

ANDON HEALTH CO., LTD.
No.3 JinPing Street, Ya An Road,Nankai District, Tianjin, China

Tianjin 2020/12/15
Place, date

Legally binding signature, Function

V1.0:2014-03-20

CE-03

Digitally signed by Busuioc Anatol
Date: 2022.01.31 20:23:33 EET
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