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

iHealth MyVitals App

Which is used together with the following devices:

Name of the product	Types
Electronic Sphygmomanometer	BP5S, BP7S, KN-550BT
Pulse Oximeter	PO3M
Pedometer	AM4

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

The medical device has been assigned to class I according to Rule 12 in Annex IX of the Directive 93/42/EEC. It bears the mark

CE

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

Following the procedure relating to the EC Declaration of Conformity set out in Annex VII of the 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 2016/10/19
Place, date

Legally binding signature, Function