

EC Declaration of Conformity

Manufacturer:

whose single Authorized Representative:

ANDON HEALTH CO., LTD.

No.3 Jinping Street, Ya An Road, Nankai
District, Tianjin, China

iHealthLabs Europe SARL

3 rue Tronchet, 75008, Paris, France

We, the manufacturer, herewith declare that the product

iHealth iGluco App

which is used together with the following devices

Name of the product	Types
Blood Glucose Monitor	BG1
	BG3
	BG5

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



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

Following the procedure relating to the EC Declaration of Conformity set out in Annex VII 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 2014/12/15
Place, date

[Signature]
Legally binding signature, Function