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:

iHealthLabs Europe SARL
3 rue Tronchet, 75008, Paris, France

We, the manufacturer, herewith declare that the products

Lancing Device

UMDNS-Code: **16-380**

Model: ALD-602

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

The medical device has been assigned to class Ⅰ according to 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 AnnexⅦ of Directive 93/42/EEC.

Following the procedure relating to the EC Declaration of Conformity set out in AnnexⅦ 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 2010/2/12
Place, date

Siyu MAO
Legally binding signature, Function