

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 SAS
36 rue de Ponthieu, 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



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 WangYang Management Representative
Place name function

WangYang 2019.12.24
Legally binding signature, date