



EC Declaration of Conformity

## 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 SAS**  
36 rue de Ponthieu, 75008, Paris, France

We, the manufacturer, herewith declare that the products

### Lancets

UMDNS-Code: **10-440**

**Model:** 28G, 30G, SL30G

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.: DD 60141763 0001  
Issue date: 2019-08-08  
Expiry date: 2024-05-27

Following the procedure relating to the EC Declaration of Conformity set out in Annex V 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 WangYang 2019/9/10  
Place name function Legally binding signature, date