EC Declaration of Conformity

Manufacturer:

Shenzhen Jiacom technology Co., ltd

No.3/F,A7 Building and 2-3/F, A6 Building, Silicon Valley Power, Qinghu Park, Longhua Street, Bao ' an District 518109 Shenzhen, China

whose single Authorized Representative:

Shanghai International Holding Corp. GmbH (Europe)

Eiffestraße 80,
D-20537 Hamburg, Germany

We, the manufacturer, herewith declare that the products

Compressor Nebulizer

UMDNS-Code: **12712**;

meet the provisions of Directive 93/42/EEC amended by 2007/42/EC 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 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.: SX 60127618 0001
Issue date: 2018-05-08
Expiry date: 2021-04-29

following the procedure relating to the EC Declaration of Conformity set out in Annex II 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

Company: Shenzhen Jiacom technology Co., ltd
Address: 3/F,A7 Building and 2-3/F, A6 Building, Silicon Valley Power, Qinghu Park, Longhua Street, Bao ' an District, Shenzhen

Shenzhen 2020/3/18
Place, date

Legally binding signature, Function.