EC Certificate

Directive 93/42/EEC Annex II, excluding Section 4
Full Quality Assurance System
Medical Devices

Registration No.: HD 60137930 0001

Report No.: 17062851 006

Manufacturer:

FEELLIFE HEALTH INC.
Room 1903, Building A
No. 9 Furong Road, Tantou Community
Songgang Subdistrict, Bao'an District
Shenzhen
518104 Guangdong
China

Products: Ultrasonic Nebulizers

Replaces Approval, Registration No.: HD 60122817 0001

Expiry Date: 2022-09-29

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

Effective Date: 2019-08-07

Date: 2019-08-07

Notified Body

Fuxiu Sheng

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.