EC Declaration of Conformity

Name and address of the manufacturer: **FEELLIFE HEALTH INC.**
Room 1903,Building A, No.9 Furong Road, Tantou Community, Songgang Subdistrict, Bao'an District, Shenzhen, 518104, Guangdong, China

Name and address of the authorized European Representative: EC REP **Prolinx GmbH Brehmstr. 56, 40239 Duesseldorf Germany**

We declare under our sole responsibility that

the medical device: **Ultrasonic Nebulizers (Portable Mesh Nebulizer)**

model: **Air 360+, mini Air 360+, A5, Air Pro, Air Angel, Air Force, Air Mask, AeroCentre, Air Q+, Air Bee, Air Garden, Air Pro Ⅱ, Air Pro Ⅲ, Aerogo , Air Plus , AeroCentre+ , Air Kids , AirICU , Air Pro Ⅷ , Air Pro Ⅸ , Air Mask Ⅱ,Air Plus 2**

of class: **Ⅱa, Rule 11**

according to annex IX of directive 93/42/EEC

meets the provisions of the directive 93/42/EEC and its transpositions in national laws which apply to it.

Conformity assessment procedure: **Directive 93/42/EEC Annex II, excluding Section 4**

Registration No.: HD 60147222 0001
Report No.: 17062851 010
Effective Date: 2020-07-30
Expiry Date: 2024-05-26

Notified Body: **TÜV Rheinland LGA Products GmbH**
Tillystraße 2,90431 Nürnberg,Germany

This Declaration of Conformity covers all medical devices as specified in the product list belonging to this declaration and is only valid in connection with a batch specific Certificate of Compliance for all products concerned bearing the CE mark.

CE 0197

The above mentioned declaration of conformity is exclusively under the responsibility of

Company: **FEELLIFE HEALTH INC.**

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

Shenzhen / 2020-08-06
Place, date

HUA JIAN / General Manager
Name and function