EU Certificate

Quality Management System
REGULATION (EU) 2017/745 on Medical Devices
Annex IX Chapter I, Section 2 and 3 and Chapter III

Registration No.:	HZ 2139746-1
Manufacturer:	FEELLIFE HEALTH INC. Room 2501, 1903, 2002, Building A, No.9 Furong Road, Tantou Community, Songgang Subdistrict, Bao'an District, Shenzhen 518104 Guangdong P.R. China
EUDAMED Single Registration No.:	CN-MF-000000782
Products:	Products of class IIa: R060101 - COLD NEBULISATION SYSTEMS
Authorized representative(s):	Lotus NL B.V. Koningin Julianaplein 10,1e Verd,2595AA,The Hague,Netherlands.

Certificate history		
Revision:	Description:	Issue date:
0	Initial certification	2024-11-15

The Notified Body hereby declares that the requirements of Annex IX, Chapter I, Section 2 and 3 of the REGULATION (EU) 2017/745 have been met for the listed products. The above named manufacturer has established and applies a quality management system, which is subject to periodic surveillance, defined by Annex IX, Chapter I, Section 3 of the aforementioned regulation. The requirements of Annex IX, Chapter III are fulfilled.
If class III devices or class IIb implantable devices referred to in the second subparagraph of Article 52(4) are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 4.9 is required before placing them on the market.

Report No.:	10922763-120
Effective date:	2024-11-15
Expiry date:	2029-11-14
Issue date:	2024-11-15

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This certificate can be validated on https://www.certipedia.com

TÜV Rheinland LGA Products GmbH is a Notified Body according to REGULATION (EU) 2017/745 concerning medical devices with the identification number 0197.

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