healthvape®

HealthVape disposables consist of 2 components, each with different certifications.

## LITHIUM BATTERY

CE RoHS COMPLIANT

## E-LIQUID

SGS ISO PRODUCED IN GMP CERTIFIED LABORATORY

healthvape®

# BATTERY CERTIFICATIONS

The HealthVape inhaler uses a disposable battery to warm the e-liquid concentrate which produces the breathable vapor.

The e-liquid is made in an FDA registered facility that conforms to Good Manufacturing Standards.

- CE Certified
- ROHS Certified
- Battery MSDS Sheet Available
- EMC Test Report Available

POCE
POCE Technology

**Shenzhen POCE Technology Co.,Ltd.**

H Building, Hongfa Science and Technology Park,
Tangtou, Shiyan, Bao'an District, Shenzhen, China

# CERTIFICATE OF CONFORMITY

| | | |
|---|---|---|
| **Certificate No.** | : | POCE190927029CCE |
| **Applicant** | : | Big River Ventures, LLC DBA: HealthVape |
| **Address** | : | 8635 W Sahara Avenue, Suite 3001 Las Vegas, NV 89117-5858 |
| **Manufacturer** | : | >> REDACTED << |
| **Address** | : | >> REDACTED << |
| | | >> REDACTED << |
| **Product** | : | HealthVape |
| **Trade Name** | : | HealthVape |
| **Model(s)** | : | HV-0011 |
| **Test Report No.** | : | POCE190927028ERE |
| **Test Standards** | : | **EN 61000-6-3:2007+A1:2011+AC:2012** |
| | | **EN 61000-6-1:2007** |
| | | **EN 61000-3-2:2014** |
| | | **EN 61000-3-3:2013** |

The EUT described above has been tested by us with the listed standards and found in compliance with the council **EMC** Directive 2014/30/EU. It is possible to use CE marking to demonstrate the compliance with this EMC Directive.

CE

Jade Wang

For Chief Executive / Jade Wang
Date: Sep. 27, 2019

This certificate of conformity is based on a single evaluation of the submitted sample(s) of the above mentioned product. It does not imply an assessment of the whole production and other relevant directives have to be observed.

POCE
POCE Technology

**Shenzhen POCE Technology Co.,Ltd.**

H Building, Hongfa Science and Technology Park,
Tangtou, Shiyan, Bao'an District, Shenzhen, China

# CERTIFICATE OF CONFORMITY

| | | |
|---|---|---|
| **Certificate No.** | : | POCE190927036HCR |
| **Applicant** | : | Big River Ventures, LLC DBA: HealthVape |
| **Address** | : | 8635 W Sahara Avenue, Suite 3001 Las Vegas, NV 89117-5858 |
| **Manufacturer** | : | >> REDACTED << |
| **Address** | : | >> REDACTED <<<br>>> REDACTED << |
| **Product** | : | HealthVape |
| **Trade Name** | : | HealthVape |
| **Model(s)** | : | HV-0011 |
| **Test Report No.** | : | POCE190927035RRR |
| **Test Standards** | : | **IEC 62321-4:2013; IEC 62321-5:2013; IEC 62321-6:2015; IEC 62321-7-1:2015; IEC 62321-8:2017** |

The EUT described above has been tested by us with the listed standards and found in compliance with the council **RoHS Directive(EU) 2015/863 amending Annex II to Directive 2011/65/EU.** It is possible to use **CE** marking to demonstrate the compliance with this **RoHS** Directive.

RoHS
POCE
CE

For Chief Executive / Jerry Yang
Date: Sep. 27, 2019

This certificate of conformity is based on a single evaluation of the submitted sample(s) of the above mentioned product. It does not imply an assessment of the whole production and other relevant directives have to be observed.

Web: http://www.poce-cert.com Tel: +86-755-29113252 E-mail: service@poce-cert.com

healthvape®

# E-LIQUID CONCENTRATE CERTIFICATIONS

The HealthVape e-liquid contains a combination of safe, tested ingredients including vitamins and essential oil extracts. There is no nicotine in any HealthVape formulas.

The e-liquid concentrate is made in an FDA registered facility that conforms to Good Manufacturing Practice Standards.

- GMP / FDA Registered Lab
- SGS / ISO Certified - ISO2200:2005
- Proprietary formula ingredient lists available with NDA
- E-Liquid SDS contains the CAS numbers for each ingredient

SGS ISO PRODUCED IN GMP CERTIFIED LABORATORY

intertek
Total Quality. Assured.

# CERTIFICATE OF REGISTRATION

**This is to certify that the quality management system of:**

## FEEL LIFE CO., LIMITED

Area B, 2/F, Building 1, Phase 1, Yangbei Industrial Zone, Huangtian Village, Xixiang Subdistrict, Baoan District, Shenzhen City, Guangdong Province, China

**has been assessed by Intertek as conforming to the requirements of:**

**CURRENT GOOD MANUFACTURING PRACTICE BASED ON 21 CFR PART 110 (2020) PUBLISHED BY U.S. FOOD AND DRUG ADMINISTRATION**

**The scope of activities:**

Manufacturing of E-liquid

GMP
Intertek

**Certificate Number:**
SZ2107C5

**Initial Audit Date:**
01-02 Jun, 2015

**Certificate Issue Date:**
14 Jul, 2021

**Certificate Expiry Date:**
26 Jul, 2024

Certification Administration Centre
Intertek Testing Services

Jacob Lin
Vice President
Life & Environment Science



SH308E0 | 1st year | 2nd year

This certificate is valid as long as it bears a proper and authentic Intertek's Laser Logo dedicated for the year of initial certification and after satisfactory annual surveillance.

Certificate CN13/89396

The management system of

# FEELLIFE HEALTH INC

Floor2, Building 1, Huangtian Yangbei Industrial Zone Phase 1, Xixiang Street, Bao'an District, Shenzhen City, Guangdong Province, P.R. China

has been assessed and certified as meeting the requirements of

# ISO 22000:2018

For the following activities

**Processing of liquid flavours used as food ingredients**

**Food Category K - Production of (Bio) Chemicals**

This certificate is valid from 18 February 2021 until 30 March 2022
and remains valid subject to satisfactory surveillance audits.
Re certification audit due 60 days prior to expiry date.
Issue 4. Certified since 30 March 2010

Authorised by

SGS United Kingdom Ltd
Rossmore Business Park Ellesmere Port Cheshire CH65 3 EN UK
t +44 (0)151 350-6666 f +44 (0)151 350-6600 www.sgs.com
The certification information can be verified on the web site of Certification and Accreditation Administration of the People's Republic of China www.cnca.gov.cn

UKAS MANAGEMENT SYSTEMS
0005

HC SGS 22000 2018 0819

# Certificate Of FDA Registration

## 2020.08-2022.07

This is certified that:

At The Address Stated Below Has Completed U.S. FOOD And DRUG

ADMINISTRATION Food Facility Registration Through MANTONG.

# FEELLIFE HEALTH INC.

Floor 2, Building 1, Huangtian Yangbei Industrial Zone, Xixiang Street, Bao'an District, Shenzhen, Guangdong, China

Food Facility Registration Number: 17629110634

MTG FDA REGISTER CENTER CORPORATE SEAL CALIFORNIA

Jacky M. Chuang

Executive Director
Date: 08-12-2020

**MTG STANDARDS TESTING & CERTIFICATION CENTER** www.fdacn.org

*This certification affirm that the above device and company was registered with U.S. Food and Drug Administration pursuant to section 305 of the United States Public Health and Bioterrorism Preparedness and response Act of 2002. P.L. 107-188.on the date stated above, and makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder. for whole sole benefit it is issued MTG CO., Inc. assumes no liability to any person or entity in connection with the foregoing. MTG is a private registration agent not affiliated with the U.S. Food and Drug Administration*