Lifemax

LIFEMAX LTD.
Unit 1 Pipers Court, Thatcham
Berkshire, RG19 4ER, England

T: 01635 874323
F: 01635 863622
info@lifemaxuk.co.uk
www.lifemaxuk.co.uk

# Declaration of Conformity

| Lifemax Model | Factory Model |
|---|---|
| 332 - Patient Alert | TS9643 |

**Electromagnetic Compatibility Directive:**
2004/108/EC. The equipment has been designed to comply with the relevant standards, EN 50130-4:2011+A1:2014, EN 61000-6-3:2007+A1:2011, EN 61000-6-1:2007, as listed in the attached certification.

**RoHS Declaration:**
We confirm that all products that we place on the market by importing after 2 January 2013 will be in compliance to the European RoHS 2 Directive 2011/65/EU and that none of the materials or components used in the production of our products will be either prohibited or exceed the legal limits as specified therein.

**Battery Directive Declaration:**
We confirm that all products that we place on the market by importing after 26 September 2008 are in compliance to the European Battery Directive 2006/66/EC, as addressed by The Batteries and Accumulators Regulations 2008 (issued by the Department for Business enterprise & Regulatory Reform). All products include clear guidance on the type of battery used and their safe removal. Any battery supplied with the product is within the material prohibition limits and subject to the appropriate markings outlined by the directive.

**WEEE Declaration:**
All Lifemax products comply with the Directive 2012/19/EU and are marked/labelled accordingly. Our WEEE number is WEE/EK0103WV.

Place of Issue: As Above
Name: Andrew Sivyer

Date of Issue: 13/12/2017
Position: Managing Director

CERTIFICATE

Anbotek
Product Safety

# CERTIFICATE

## Of Conformity
## EU Council Directive 2014/30/EU
## Electromagnetic Compatibility

**Registration No.: AT0917110059E**

**Report No.: R0917110059E**

| | | |
|---|---|---|
| **Applicant** | : | **TFE HONG KONG LIMITED**<br>7TH FLOOR, GEMMY FACTORY BUILDING, 12 HUNG TO ROAD, KWUN TONG, KOWLOON, HONG KONG |
| **Product** | : | **Patient Alert** |
| **Identification** | : | **Model No. : TS9643**<br>**Trade Mark : TFE**<br>**Rating : DC 3V** |
| **Test Standards** | : | **EN 61000-6-3: 2007+A1: 2011**<br>**EN 61000-6-1: 2007**<br>**EN 50130-4: 2011+A1: 2014** |

**The certificate of conformity is based on an evaluation of a sample of the above-mentioned product. Technical report and documentation are at the applicant's disposal. This is to certify that the tested sample is in conformity with all provisions of Annex II of Council Directive 2014/30/EU, in its latest amended version, referred to EMC Directive. The certificate does not imply assessment of the production and does not permit the use of Lab's logo. The applicant of the certificate is authorized to use this certificate in connection with EU declaration of conformity to Article 15 of the Directive.**

**Certified by**

ANBOTEK Anbotek CERTIFICATION

**Nov. 29, 2017**
**Date**

**Tom Chen**
**Manager**

CE **The CE Marking may only be used if all relevant and effective EU Directives are complied with** CE

**Shenzhen Anbotek Compliance Laboratory Limited**
1/F., Building 1, SEC Industrial Park, No.0409 Qianhai Road, Nanshan District, Shenzhen, Guangdong, China
Http://www.anbotek.com
Tel: (86)755-26066544
Fax: (86)755-26014772
Email: service@anbotek.com

CERTIFICATE

Anbotek
Product Safety

# CERTIFICATE

## Of Conformity
## EC Council Directive 2011/65/EU
## Restriction of the Use of Certain Hazardous Substances in Electrical and Electronic Equipment

**Registration No.: ATSZR171125008002**

**Report No.: SZR171125008002**

**Applicant** : **TFE HONG KONG LIMITED**
7TH FLOOR，GEMMY FACTORY BUILDING，12 HUNG TO ROAD，KWUN TONG，KOWLOON，HONG KONG

**Product** : **Patient Alert**

**Identification** : **Model No.** : **TS9643**

**Trade Mark** : **N.A.**

**Test method** : **IEC 62321-3-1:2013 Ed.1.0, IEC 62321-5:2013 Ed.1.0, IEC 62321-4:2013 Ed.1.0, IEC 62321-7-1:2015 Ed.1.0, IEC 62321-7-2:2017 Ed.1.0, IEC 62321-6:2015 Ed.1.0**

**This is to certify that, the certificate is based on Anbotek's test results and other related substance information provided by applicant. The submitted sample fulfills the requirement of the Directive 2011/65/EU ( RoHS ).**



**Certified by**

CE Product RoHS

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**Dec. 01, 2017**
**Date**

**Shenzhen Anbotek Compliance Laboratory Limited**
East of 4/F., Building A, Hourui No.3 Industrial Zone, Xixiang Street, Bao'an District, Shenzhen, Guangdong, China
Tel:(86)755-26066061
Fax: (86)755-26066021
Http://www.anbotek.com
Email:service@anbotek.com