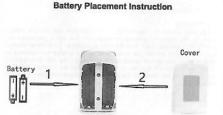
# PULSE OXIMETER



Battery case

Part 2 Operation Instructions ...... 6

Part 3 EMC Related Summary ...... 12

Part 4 Repair and Warranty ...... 12

.............

Content

# Pulse Oximeter user manual Part 1 General Information ......1

## Part 1 General Information

1. Preduct Information

Product name: Pulse Oximeter

2. Explanation of Symbols

The following symbols appear on the Pulse Oximeter and its packaging. Table 1 Pulse Oximeter and Packaging Symbols

| Symbol   | Description  | Symbol      | Description                                  |
|----------|--|-------------|--|
| 潦        | BF applied part:<br>including F applied<br>part (float/insulation) | $\triangle$ | Caution,consult<br>accompanying<br>documents |
| 11       | Keep upright   | M           | Date of<br>manufacture                       |
| *        | Keep dry   | I           | Fragile, handle<br>with care                 |
| 10%      | Humidity<br>limitations  | -30°C       | Temperature<br>limitations                   |
| 107.4LPs | Atmospheric<br>pressure<br>limitations                             | SN          | Serial Number                                |
|          |  | ***         | Manufacturer address                         |
| IPX1     | Protective grade   | Z           | Complicance to<br>WEEE standard              |

3. Safety Standards The following table describes the safety standards of the Pulse Oximeter.

Table 2 Safety Standards

| Parameter  | Specification   |
|--|---|
| Protection class   | Class I, anti-shock, externally and internally<br>powered equipment, per IEC 60601-1  |
| Degree of protection                                     | Type BF defibrillator-proof: per IEC 60601-1 Degree of noxious-liquid proof as IPXI Anti-shock degree as BF applied part According to the degree of safety of application in the presence of a flammable annesthetic mixture with air or with oxygen or introus oxide, the equipment is not suitable for use in the presence of a flammable anaesthetic mixture with air or with oxygen or nitrous oxide. |
| Sterilization and disinfection                           | As recommended by manufacturer  |
| Mode of operation  | Continuous  |
| I. Safety Conventions The manual uses the followarnings. | wing conventions for Notes, Cautions, and   |
| wanings.   |   |
|  | on to an important point in the text.   |

\* FUNCTIONAL TESTER cannot be used to assess ACCURACY.

\* The skin temperature is initially at 35 °C for each PULSE OXIMETER, the

APPLIED PART temperature can not exceed 41 °C. \* The material that the pulse oximeter contacted to body is Non-toxic silica gel

which meet the ISO10993 requirements, so can be safety used. But to some natural rubber latex sensitive user, there may be some allergic reactions. \* Only use accessories recommended by the manufacturer. Using other kinds

of accessories might cause damage or personal injury. Modification of the

Pulse Oximeter could be unsafe as applicable.

\* To avoid personal injury, only use accessories and parts produced or

lectromagnetic energy is extremely high. Ensure that any nearby instruments are also in compliance with EMC standards, Never turn on or use portable communication devices like mobile phones or portable dual-channel radios

near a Pulse Oximeter. \* When power is lost for less than or equal to 30 s, the ALARM SETTINGS

the Pulse Oximeter. \* Periodically check the Pulse Oximeter for damage. When necessary according

may cause choking.

\* Never use the Pulse Oximeter during MRIs or CT scans.

" Never use the Pulse Oximeter in an environment of anesthetic gases.

\* The Pulse Oximeter is not for diagnostic or therapeutic use.

"This device is intended for sport and recreational use.

5. Safety Requirements

recommended by distributor. Otherwise, damage to the Pulse Oximeter can occur

\* Inhalation or swallowing of small parts that be detached from Pulse Oximeter

\* The degrade sensor may degrade the performance.

4 The Pulse Oximeter must conform to the international standard IEC 60601-1-2

prior to the power loss shall be restored automatically.

\* Clean and sterilize the Pulse Oximeter and accessories according to local

6 Keep all Pulse Oximeter packing materials away from children, pet or pests,

and other applicable EMC standards. Interference takes place when

\* Ensure that qualified service representatives annually calibrate and maintain

to your local hospital waste disposal regulations.

requirements. Turn off the Pulse Oximeter before cleaning or sterilization.

or dispose of them in accordance with your local environmental regulations.

Always properly dispose of the Pulse Oximeter and all accessories at the end

of their service life. Dispose of batteries according to your local regulations.

Never incinerate batteries or expose them to high temperatures.

\* Ensure that no water condenses into or on the Pulse Oximeter. Condensation can occur from changes in temperature or exposure to humidity.

### 6. Product Operation Scope

This Pulse Oximeter is a kind of innovated detection device with non-invasive and continuous features for artery SpO2 and PR detection. It is portable and easy to measure the SpO2 and PR value quickly and precisely. This can be through the finger Pulse Oximeter to measure human blood oxygem saturation and heart rate. This product is untable for family, climic, oxygen bar, sports health (use before and after exercise is not recommended for use during exercise), community health and other ranges. It's for ages from 15 to 60 years old users. This product is not suitable for Pulse Oximetering the users's proloned use.

### 7. General Description

Haemoglobin Saturation is percentage of Oxyhemoglobin (HbO2) capacity, compounded with oxygen, by all combinativable haemoglobin (HbO obin (HbO2) capacity in blood. In other words, it is consistency

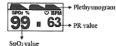
of Oxyhemoglobin in blood. It is a very important ecological parameter for Respiratory circulation System. Many respiratory diseases can result in haemoglobin saturation being lowered in human blood. Moreover, the following factors can also lead to problems in oxygen supply, so that human haemoglobin saturation might be reduced: Automatic Organic Regulation Malfunction caused by Anesthesia, Intensive Postoperative Trauma, hurts resulted in by some medical examination and etc. In the situation, illnesses, such as light head, asthenia, vomitory and etc., might happen to users and even endanger the users' s life. Therefore, it is very important to know Hemoglobin saturation of users timely in clinical medical aspects. So that doctors can find problems in time. The fingertip pulse oximeter features in small volume, low power consumption convenient operation and being portable. It is only necessary for users to put one of his fingers into a fingertip photoelectric sensor for diagnosis. and a display screen will directly show measured value of hemoglobin Saturation. It has been proved in clinical experiments that it features in rather high precise and repeatability.

### 8. Measurement principle

Principle of the oximeter is as follows: An experience formula of data process is established taking use of Lambert beer Law according to Spectrum Absorption Characteristics of reductive hemoglobin (R Hb) and Oxyhemoglobin (O2 Hb) in glow and near-infrared zones. Operation principle of the instrument is photoelectric Oxyhemoglobin Inspection Technology, is adopted in accordance with capacity pulse scanning and recording Technology, so that two beams of different wavelength of lights (660am glow and 940nm near infrared light) can be focused onto human anil tip through perspective clamp finger-type sensor. Then measured signal can be obtained by a photosensitive element, information acquired through which will be shown on two groups of LED through process in electronic circuits and microprocessor.

### 9. Appearance introduction





### 10. Features

- Two colors OLED display, figure and Oxygen volume chart display together on interface:
- Adjust the display interface direction manually, according to the users observation data needs;
- 3.audible alarm function;
- 4 Low Power consumption. 20 hours continuous to work
- 5.Low Perfusion ≤0.6%.
- Low voltage indicator.
   Automatic power off when no signal in 8s.
- 8.Small and light weight, convenient to carry.
- A menu function can set up oxygen alarm limit, the upper and lower limits of PR, alarm switch, sound switch

### Part 2 Operation Instructions

- 11. Operation Instructions
- 11.1 The product Operation Instructions
- Installing two AAA batteries into battery cassette in correct polarities and cover it.
- Plug one of fingers into rubber hole of the Oximeter (it is best to plug the finger thoroughly) nail surface upward, then releasing the clamp.





- 3.Press the switch button once on front panel.
- Your finger do not tremble during the Oximeter is working. Your body is not recommended in moving status.
- Read correspondent date from display screen.

Note — Please use the medical alcohol to clean the subber touching the finger inside of Oximeter, and clean the test finger using medical alcohol before and after each test. (The subber inside of the Oximeter belongs medical subber, which has no toxin, and no harmful to the skin of human

being).
When your finger is plugged into the Oximeter, you nail surface must

be upward.

Disposable batteries should be unloaded when used for a long interval.

### Warning

- \* The maximum application time for a PULSE OXIMETER PROBE at a single site is 4 hours.
- The misapplication of a PULSE OXIMETER PROBE with excessive pressure for prolonged periods can induce pressure injury.
- \* Ambient light (including photodynamic therapy), physical movement (USERS and imposed motion), diagnostic testing; low perfusion; electromagnetic interference; HF SURGICAL EQUIPMENT; dysfunctional haemoglobin; there will be wundage for SpO2value.

### 11.2 Operation Instructions

Interface 3

Interface 4

Interface 5

Interface 6

11.2.1 Display Description (6 interface diagrams) Alm Settings V1.02 Setting interface Restore Pulse Oximeter Interfaces Without detection signal interface A detection signal interface diagram SP02 % Interface I 10111011101110 Interface : 6 001

6 SPO2% PREEDIN SPO:% PRbpm

~~~ 79 "66 M48 O % <sup>20</sup>dS M48 O % 40dS

1111111 100 80

SPO2% PRbpm

11.2.2 Button operating instructions

I put into two AAA batteries according to the instructions, the device will turn on automatically and display interface 1; then put into finger for measuring, if there is finger for detection and without operation, it power off automatically in 8s

2. When there is battery, but the Pulse Oximeter power off, press the button, it

3 During the measurement (there is a measurement signal and figure), press the button shortly, the interface can turns from interface 1 to interface 6 circularly. Press the button longly during the measurement; it can turn to settings menu interface.

4. In the Settings menu interface, press button longly, it can choose 8 options circularly, they are Alarm switch (Alarm Sound on/off). PR Sound switch (BeeP) and the withdraw from the menu (Exit). The selected

options foreground color will become Blue Square 5. All the setting value will restore to default value: alarm off, oxygen lower limit value 90. PR upper limit value: 110. PR lower limit value: 50. PR

sound off when we replaced the battery. 6.In the Settings menu interface, choose to Exit option and press button

shortly, it will quit the setting menu interface.

12. Alarm

The Pulse Oxieter uses the following alarm indicators: audible alarm and visual alarms

Note - To correctly identify visual alarms, always observe the Pulse Oximeter within 1 meter of its position.

12.1 Low power alarm

when battery power is at lowest level, the battery capacity indicates empty in OLED, remind users of replacement of battery.

Note - When power is lost for less than or equal to 30 s, the ALARM

SETTINGS prior to the power loss shall be restored automatically.

45dB-70dB.

12.2 Audible alarm The audible alarm is buzzer alarm; the auditory sound pressure range is

12.3 SpO2 and PR alarm

" Default alarming Value of SpO2: 90%.

" Default alarming Value of PR: High Limit: 110BPM: Lower Limit: 50BPM

"When it is under operation conditions, if the measurement of the PR and SpO2 value exceed the setting alarm limit and the alarm switch is on. It

has a sound alarm and its long buzzer alarm.

\* When it is under operation conditions, if the PR sound switch is on, the

measurement PR will go with a PR sound.

To test the alarm according the following a)Connect the Pulse Oximeter to simulator:

b)the simulator setting : SpO2 90%, PR 200bpm.;

c)the Pulse Oximeter alarm setting: SpO2 95%, PR High Limit 110 bpm; d)to verify the Pulse Oximeter alarm.

13. SpO2 Specifications

The update rate for the SpO2 value and pulse rate is typically 1 second Data averaging and other signal processing on the displayed and transmitted data values of SpO2 and pulse rate is not more than 20 seconds. Depending on the magnitude of difference between the alarm limit and the displayed value, the alarm signal generation delay may be from 1 second to 20 seconds. The maximum alarm condition delay is 4 seconds the maximum alarm signal delay is 20 seconds, the average alarm condition is 2 seconds, the average alarm signal is 10 seconds.

Because pulse oximeter measurements are statistically distributed, only approximately two-thirds of pulse oximeter equipment measurements can be expected to fall within the Arms value measured by a CO-oximeter.

The signal adequacy is indicated by waveform, and it is NORMALIZED.

SpO2 calibration when the Pulse Oximeter is in use.

Note - The PULSE OXIMETER EQUIPMENT is calibrated to display FUNCTIONAL OXYGEN SATURATION. It is not necessary to have a The following table describes the SpO2 specifications. Table 3 SpO2 Specifications

Parameter

Specification

SpO2 measurement range 35% - 99% ±1% in the range of 90%-99%. SpO2 accuracy1 ±2% in the range of 70%-89%. Other scope is not defined Consistent with the display range, the SpO2 Alarm range set step length is 1% Pulse rate range 30 bpm -240 bpm ±1 bpm Pulse rate accuracy Consistent with the display range, the Pulse rate Alarm range set step length is 5bpm Data undate cycle 0.25 5 - 2 5 SpO: PR average onise rate sound PR tone Peak Wavelength range 2 500 nm - 1000nm Maximum optical output power 150 mW Pulse rate display Numeric Display Specifications two color OLFD 0.96" in normal measurement, less than power dissipation 40 mA in power off state, less than

(1). Sensor accuracy was obtained by performing controlled hypoxia studies on healthy, non-smoking adult volunteers (according to ISO 80601-2-61). The SpO2 readings have been compared to CO-oximeter measurements on arterial

blood samples. To repre-sent the general population, data from at least

10 subjects (male and female) with a wide range of skin color was taken to validate SpO2accuracy. Accuracy data for Masimo sensors is included in the sensor package insert. (2). Information about wavelength ranges can be useful for clinicians performing

photo dynamic therapy.

14. Power Specifications

The following table describes the power specifications.

Table 4 Power Specifications

Parameter Specification DC 3V AAA(X2) battery The Pulse Oximeter powers off if the battery power is almost depleted

15. Physical Specifications

The following table describes the physical specifications. Table 5 Physical Specifications

Item Parameter 35\*65\*40mm Size

16. Environmental Specifications The following table describes the environmental specifications.

Table 6 Environmental Specifications Specification Parameter Operating: 0°C -40°C Temperature Storage: -20 "C+55"C for the device Operating: 15% - 80% (noncondensing) Relative humidity Storage: 10% - 93% (noncondensing) Operating: 59 kPa~107.4 kPa Barometric pressure Storage: 22 kPa~107.4 kPa

SPO2% PRbpm

# Part 3 EMC Related Summary

# 17. EMC Related Summary

compatibility information which is attached with it. 3.Portable and mobile RF communication devices may influence pulse

1 Pulse Oximeter meets the requirement of electromagnetic compatibility in 2.The user needs to install and use according to electromagnetism

Oximeter performance, so pulse Oximeter should be kept away from them during using

Guidance and manufacturer's declaration stated in the appendix.

## 17.2 WARNING

17.1 NOTICE

IEC60601-1-2.

1.The user needs to install and use pulse Oximeter according to electromagnetism compatibility information which is attached with it.

2 Portable and mobile frequency communication devices may influence its performance, so it should be kept off these devices.

3. Pulse Oximeter should not be used adjacent to or stacked with other equipment and that if adjacent or stacked use is necessary, the Pulse Oximeter should be observed to verify normal operation in the configuration in which it will be used.

4. Speaking, pulse Oximeter could not be used for transport using, like ambulances (land and air ambulances).

# Part 4 Repair and Warranty

# 18. Repair and Maintenance

1. Regular inspection to make sure that no obvious damage existed to affect the safety and performance of device.

2. No flammable substance, overtop or lower temperature and humidity existed in operation conditions

3. When the device is dabbled or there is hydraulic set existed, stop operating, 4. When lower power capacity light in red pls replace the battery right away.

Pls clean the surface before applying for detection.

6.Pls take out the battery when device is not used for a period of time. 7.Pls disposure the battery according to the local statute.

8.If there is dust or dut on surface, 75% density of medical alcohol can be used to clean the surface. Pls use dry fabric with little alcohol to avoid alcohol permeates into the device.

9. The transportation and storage conditions are

Temperature: -20'C-55'C: Humidity: 10%-93%.

# 10 Troublechooting

| Trouble                                        | Possible Reason                                                                                                        | Solution                                                                                |
|------------------------------------------------|------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------|
| The SpO2 and<br>Pulse Rate<br>display instable | The finger is not places usade enough.     The finger is shaking of the users is moving.                               | 1 Place the finger properly and try<br>again<br>2 Let the users keep calm.              |
| The device can<br>not turn on                  | The batteries are drained or almost drained.     The batteries are not morried properly.     The device's malfunction. | Change batteries.     Reinstall batteries.     Please contact the local service center. |
| The indicator<br>light is off<br>suddenly      | The device will power off automatically when it gets no signal for 8 seconds.     The batteries are almost dramed.     | Normal.     Change batteries.                                                           |
| The SpO2 and<br>Pulse Rate alarm               | Thevalue has exceeded the high alarm limit or below the low alarm limit.                                               | Check the users physiological status, and alarm limit setup.                            |

# 20. Accessory

| AAA battery- | 2 | pc |
|--------------|---|----|
| Hang String- |   | pc |
| User Manual  | 1 | pc |

## 21. Warranty

The unit can not be repaired by users themselves. All services must be done by the engineers approved by distributor. The unit is guaranteed for a period of 12 months, valid from the date of purchase. Distributor warrants that each product we sell you is free from defects in labor and materials and shall conform to its product specifications as defined in the user documentation. If the product doesn't function as warranted during the warranty period, we will repair or replace it without charge. Misuse, improper maintenance may void the warranty.