LIFESMART

CE 0123

M70C

Operator's Manual Fingertip Pulse Oximeter

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General Description

The measurement of oxygen saturation of arterial blood (also known as pulse oxygen saturation, usually shortened as SpO_2) adopts the principles of light spectra and volume tracing. The LED emits lights with two specific wavelengths, which are selectively absorbed by oxygenated hemoglobin and deoxyhemoglobin. The optical receptor measures the changes in the light intensity after the light passes the capillary network and estimates the ratio of oxygenated hemoglobin and the total hemoglobin.

$$SpO_2\% = \frac{\text{oxygenated hemoglobin}}{\text{oxyhemoglobin} + \text{deoxyhemoglobin}} \times 100\%$$

Method for evaluating SpO2 accuracy: The SpO2 accuracy of pulse oximeter is measured by comparing SpO2 readings of the pulse oximeter to values of SaO2 determined with a CO-OXIMETER.The healthy volunteers who consent to induced hypoxia and arterial blood sampling as part of the experimental procedure (see ISO80601-2-61 Annex EE.2)

Caution

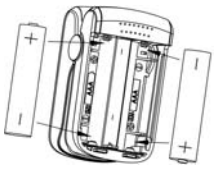
- Please read the user manual carefully prior to operating.

Intended Use

The Fingertip Pulse Oximeter is intended to measure functional arterial oxygen saturation (SpO2) and pulse rate of adult, pediatric and adolescent patients in hospital, hospital type facilities as well as in the home care environment.
The oximeter is not suitable to monitor patient continuously for long term

Battery Installations

1. Push the battery cover.
2. Install two AAA batteries into battery cabin in correct polarities, as shown on the right.
3. Close the battery cover.

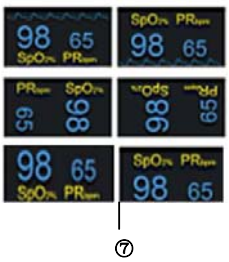


Notes:

- **For battery installation, see picture to the right. Put or remove batteries in right order, or it may damage the bracket.**
- **Battery polarities must be correctly installed. Otherwise, damage might be caused to device.**
- **Please remove the battery if the oximeter will not be used for long time.**

Operation Instructions

1. Install two AAA batteries into battery cassette before closing its cover.
2. Nip the oximeter, then insert one of fingers into the rubber hole of the oximeter before releasing the oximeter, and your nail surface must be upward.
3. Press the function button once on front panel.
4. Your finger and body should not tremble during measuring.
5. Read corresponding data on the display screen.
6. After turning the oximeter on, each time you press the power switch, the display screen will change to another direction. There are two display modes. If you long press the power switch, you can adjust the brightness of screen from 1 to 5.



① SpO_2 Plethysmogram
② SpO_2 reading
③ Pulse rate reading
④ Indication of battery capacity
⑤ Indication of pulse intensity
⑥ Indication of screen brightness
⑦ Display modes

Precautions for Use

- The patient is the operator when the device is used at home.
- Patients can maintain and use all functions of the device safely according to this user's manual.
- Keep this product out of reach of children to avoid injury to children.
- Explosion hazard. Do not use the oximeter in the presence of flammable anesthetics mixture with air, oxygen, or hydrogen.
- When the oximeter is in use, there should not be any great power appliances as high voltage cables, X-ray machine, ultrasound equipment and electrizer in use nearby.
- Keep the oximeter away from lint, dust, vibration, corrosive substances, explosive materials, high temperature and moisture.
- This oximeter does not have an alarm function; please do not use this product in the environment where an alarm is required.
- The oximeter should be handled with care so as to avoid shocks and falls.
- When the oximeter is in use, it must be ensured the batteries have sufficient capacity; otherwise, there might be such phenomena as starting-up abnormalities or inaccurate measurement data, etc.
- Please do not use pointed objects such as pen point or nails for pressing operation, otherwise it might cause permanent damage to the surface of the keyboard.
- Do not make any clinical judgments based solely on the oximeter. The oximeter is intended only as an adjunct in patient assessment. It must be used in conjunction with clinical signs and symptoms, as well as doctor's diagnoses.
- To ensure accurate performance and prevent device failure, do not expose the oximeter to extreme moisture, such as direct exposure to rain. Such exposure may cause inaccurate performance or device failure.
- Do not conduct SpO_2 measurement on the finger smeared with nail polish, otherwise this will lead to unreliable measurement results.
- Please do not open the enclosure. The enclosure shall only be opened by the authorized person.
- In order to have more accurate measurements of SpO_2 and PR, the oximeter should be used in quiet and comfortable environment.
- Follow local ordinances and recycling instructions regarding disposal or recycling of the device and device components, including batteries.
- Prolonged continuous monitoring may increase the risk of unexpected changes in skin characteristics, such as irritation, redness, blisters or burns. Inspect the sensor site every two hours and move the sensor if the skin quality changes.
- Pulse oximeter simulator can not be used to access the accuracy of the pulse oximeter.
- The expected service life of the device is five years.
- For assistance with installation, use or maintenance, contact the manufacturer or manufacturer's representative.
- To validate the PR accuracy, we refer to the electronic pulse simulator.
- When used at room temperature from the lowest or highest storage temperature environment, the product can be used directly without needing to be placed for a period of time to achieve its expected function.

Download and Install APP Software

1. Software download methods:
   - IOS device users, login to your ***App Store*** account and search for LifeSmart Health.
   - Android device users, login to your ***Google Play*** account and search LifeSmart.

QR Code for download:

Android system IOS system

2. Download the **Free LifeSmart App**

Note: App software only applied for the oximeter with Bluetooth.

Maintenance

1. Using a soft cloth dampened with either a commercial nonabrasive cleaner, or a solution of 70% alcohol in water, lightly wipe the surfaces of the oximeter.
2. The most commonly used hospital cleaning agent and non-corrosive detergents can be used for cleaning the oximeter, but please be careful that most detergents must be diluted before use; Please use them according to the directions of the manufacturers of the detergents.
3. Avoid using alcohol-based, amido or acetone-based detergents
4. The casing of the oximeter should be kept from the contamination of filth and dirt, and it can be wiped with non-velvet soft cloth. When cleaning, do not spill the liquid onto the instrument. Ensure no liquid is allowed to enter the inside of the oximeter.
5. It is forbidden to use such grinding materials as wire brush or metal polishing agent, because these materials may cause damage to the panels of the oximeter.
6. Please do not soak the oximeter in liquid.
7. Under normal circumstances, it is unnecessary for the oximeter to have special maintenance, and cautions must be exercised on the following points during the use of the oximeter:

- Please use the oximeter in the environment according to the requirements of the performance criteria.
- Avoid exposure to direct sunlight.
- Avoid excessive radioactive infrared rays or ultraviolet rays.
- Avoid contacts with organic solutions, dusts or corrosive gases.

Product Specifications

◆ **Measurement specifications**

SpO_2	
Measuring Range	0~100%
Resolution	1%
Accuracy	At 70~100%, ±2%; At 0~69%, unspecified
Data update period	<13 s

EC REP

PR	
Measuring Range	25~250 bpm
Resolution	1 bpm
Accuracy	±1% or ± 1 bpm, whichever is greater
Data update period	< 13 s

◆ **Battery specifications**

Type	Voltage
Two AAA alkaline battery	1.5 Volts DC (per battery)
The oximeter uses two 1.5 V AAA type batteries. A set of new batteries can be used after 18 hours, depending on battery types.	

◆ **Environmental specifications**

Operation

Temperature	+5℃～+40℃
Atmospheric Pressure	700hPa～1060hPa
Relative Humidity	15%～85% （non condensing）

Transport and Storage

Temperature	-20℃～+55℃
Atmospheric Pressure	500hPa～1060hPa
Relative Humidity	10%～93%（non condensing）

◆ **Physical specifications**

Weight	about 21g (exclude battery) about 54g (include battery)
Dimensions	60mm(length) × 35mm(width) ×35mm(height)

◆ **Sensors specifications**

Wavelength	Pulse oximetry sensors contain LEDs that emit red light at a wavelength of approximately 660 nm and infrared light at a wavelength of approximately 905 nm. The total optical output power of the sensor LEDs is less than 15 mW. This information may be useful to clinicians, such as those performing photodynamic therapy. **Note:** Sensor LED light emissions fall within Class 1 level, according to IEC 60825-1. No special safety precautions are required.

Possible Problems and Resolutions

Problems	Possible causes	Solution
There is no response to the function button.	The button cannot be pressed to its position	Ensure that the button is fully depressed.
	Battery capacities are low	The batteries may be missing, discharged, or oriented incorrectly. Replace them with new ones, or insert them correctly
The Pulse search time is too long	Perfusion may be too low	Check the patient. Change the measuring site.
	Patient movement	Interference due to patient activity may be preventing the oximeter from tracking the pulse. Keep the patient still, if possible.
	Electromagnetic interference may be preventing the oximeter from tracking the pulse.	Remove the source of interference.
	There may be interference due to ambient light, or the oximeter may be on an extremity with a blood pressure cuff, arterial catheter, or intravascular line.	Reposition oximeter, as necessary.
Display is dark-or-bright	Battery capacities are low.	Replace the batteries.

Symbols Definitions

Symbol	Definition
	Type BF equipment (Refer to IEC 60601-1)
%SpO₂	Oxygen saturation of arterial blood
♡/ Min	Pulse rate
	Non-Alarm indication (The device does not have alarm function)
IP22	Enclosure degree of ingress protection.
SN	Serial number
	Refer to this user's manual.
	Symbol for the marking of electrical and electronics devices according to Waste Electrical and Electronic Equipment Directive. The device, accessories and the packaging have to be disposed of waste correctly at the end of the usage. Please follow Local Ordinances or Regulations for disposal. **Note: The Oximeter is applied to this regulation.**

Guidance and manufacturer's declaration–electromagnetic emissions -for all EQUIPMENT and SYSTEMS

Guidance and manufacturer's declaration – electromagnetic emission		
The Fingertip Pulse Oximeter is intended for use in the electromagnetic environment specified below. The user of the Fingertip Pulse Oximeter should assure that it is used in such an environment.		
Emission test	**Compliance**	**Electromagnetic environment – guidance**
RF emissions CISPR 11	Group 1	The Fingertip Pulse Oximeter uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emission CISPR 11	Class B	The Fingertip Pulse Oximeter is suitable for use in all establishments, including domestic establishments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.

Instructions on Environmental Aspects

■ **Instructions for minimizing environmental impact during normal use.**

1. Instructions on how to install the fingertip pulse oximeter in order to minimize the ENVIRONMENTAL IMPACT during its EXPECTED SERVICE LIFE;

Try to keep the integrity of the non-disposable packing material and put away the packing materials for future use or put into the specified location where complying with the rules and regulations of the local council or the hospital. Avoid overusing the cleaning reagents and other substances.

2. Instructions on how to use and maintain the fingertip pulse oximeter in order to minimize the ENVIRONMENTAL IMPACT during its EXPECTED SERVICE LIFE;

Do not mix disinfecting solutions (such as bleach and ammonia) as this may result in hazardous or poisonous gases or liquids. When there is a need to maintain, please follow the instructions for use or follow the rules and regulations of the hospital.

3. Consumption during NORMAL USE (e.g. energy, consumable materials/parts, disposables, water, gasses, chemicals/reagents etc.);

During normal use of this device, it will consume electricity (battery). The batteries shall be disposed following the rules. For cleaning or disinfection for the machine, the water and ethanol will be used and the waste liquid shall be thrown following the rules.

4. Emissions during NORMAL USE (e.g. WASTE water, WASTE consumable materials, acoustic, energy, heat, gasses, vapours, particulates, HAZARDOUS SUBSTANCES and other WASTE);

Consumption of the battery during use.

5. Information on the location within the device of HAZARDOUS SUBSTANCES, radioactive sources and induced radioactive materials.

This product has no hazardous substances, such as radioactive sources or induced radioactive materials.

■ **Information for end of life management.**

1. The location of components and parts within the device that contain stored energy or pose other hazards that can result in an unacceptable risk to disassemblers or others and methods for controlling such risks.

The device uses an alkaline battery. May heat, explode or leak if shorted, recharged, disposed of in fire or dissected.

2. The identity and location of hazardous substances requiring special handling and treatments.

The battery is installed in the battery case.

3. Disassembly instructions sufficient for the safe removal of these hazardous substances including radioactive sources and induced radioactive materials within the monitor.

For other hazards that may result in unacceptable risk, the main concern is handling the battery. Do not store the battery in a high temperature environment and store the battery in a cool, ventilated environment.

As for disposing or recycling of the device and device components at end of life, follow local ordinances and recycling instructions regarding

Applicable Models

M70C

Packing List

NO.	Item	Quantity	
1	Oximeter	1	☐
2	AAA battery	2	☐
3	Cord	1	☐
4	User's manual	1	☐

Sponsor: Genesis Biotech Pty Ltd.
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