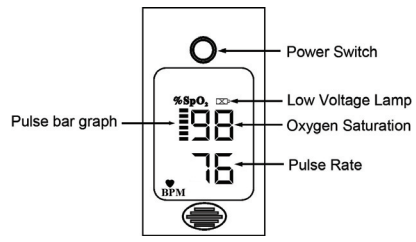
OPERATOR' S MANUAL

Fingertip Pulse Oximeter

CONCORD HEALTH SUPPLY, INC.

Concord Oximeter

Concord Emerald	CCI-300-OG
Concord Sapphire	CCI-300C3-BLUE
Concord Topaz	CCI-300C
Concord Pink	CCI-300-PINK



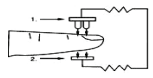
General Description

Thank you very much for purchasing a Concord Pulse Oximeter.
This Manual describes the Pulse Oximeter' s features and requirements, functions, specifications, correct methods for transportation, installation, usage, operation, repair, maintenance and storage, as well as the safety procedures to protect both the user and equipment. Please read and follow the User Manual carefully before using this product. Failure to follow the User Manual may cause measuring abnormality, equipment damage and human injury. The manufacturer is NOT responsible for the safety, reliability and performance issues and any monitoring abnormality, human injury and equipment damage due to improper use. The specific products you received may not be exactly as described in this User Manual. This product can be used repeatedly for sports and aviation use. If you have any questions regarding to the use of this product, please call us at 888-970-2999 Monday- Friday from 8:00 AM to 5:00 PM Central Time.

Measurement Principle

Principle of the oximeter is as follows: A mathematical formula is established making use of Lambert Beer Law according to Spectrum Absorption Characteristics of Reductive hemoglobin (RHb) and Oxyhemoglobin (Hb_2O) in red and near-infrared zones. Operation principle of the instrument: Photoelectric Oxyhemoglobin Inspection Technology is adopted in accordance with Capacity Pulse Scanning and Recording Technology, so that two beams of different wavelength of lights (660nm red and 905nm near infrared light) can be focused onto a human nail tip through a clamping finger-type sensor. A measured signal obtained by a photosensitive element, will be shown on the oximeter' s display through process in electronic circuits and microprocessor.

Diagram of Operation Principle
1. Red and Infrared-ray Emission Tube
2. Red and Infrared-ray Receipt Tube



Precautions For Use

1. Before use, carefully read the manual.
2. Check the main unit and all accessories periodically to make sure that there is no visible damage that may affect user's safety and monitoring performance. It is recommended that the device should be inspected once a week.
3. The fingertip pulse oximeter must be able to measure the pulse properly to obtain an accurate SpO_2 measurement. Verify that nothing is hindering the pulse measurement before relying on the SpO_2 measurement.
4. Keep the oximeter away from dust, vibration, corrosive substances, explosive materials, high temperature and moisture.
5. Do not use the fingertip pulse oximeter in an explosive atmosphere.
6. Not intended for Continuous monitoring
7. The fingertip pulse oximeter is intended only for sports and aviation monitoring
8. Do not sterilize the device using autoclaving, ethylene oxide sterilizing, or immersing the device in liquid. The device is not intended for sterilization.
9. Follow local ordinances and recycling instructions regarding disposal or recycling of the device and device components, including batteries.
10. This equipment complies with IEC 60601-1-2:2014 for electromagnetic compatibility for electrical equipment and/or systems. However, because of the proliferation of radio-frequency transmitting equipment and other sources of electrical noise the environment, it is possible that high levels of such interference due to close proximity or strength of a source might disrupt the performance of this device.
11. Portable and mobile RF communications equipment can affect electrical equipment.
12. This equipment should not be used adjacent to or stacked with other equipment.
13. It may be unsafe to:
    —use accessories 、 detachable parts and materials not described in the instructions for use
    —interconnect this equipment with other equipment not described in the instructions for use
    —disassemble, repair or modify the equipment.
14. These materials that contact with the users skin contain medical silicone and ABS plastic enclosure all pass the ISO10993-5 Tests for invitro cytotoxicity and ISO10993-10 Tests for irritation and delayed-type hypersensitivity.

Contraindication

It is not for continuous monitoring.

Inaccurate measurements may be caused by

1. Significant levels of dysfunctional hemoglobin (such as carbonyl - hemoglobin or methemoglobin).
2. High ambient light. Shield the sensor area if necessary.
3. Excessive patient movement.
4. High-frequency interference.
5. Poor blood perfusion.
6. Placement of a sensor on an extremity with a blood pressure cuff.
7. Fingernail polish or false fingernails.
8. Weak pulse quality.
9. Low hemoglobin.

Product Features

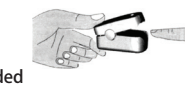
1 Brightness LED display SpO_2 , PR, and Pulse bar.
2 Two display modes.
3 2 pcs AAA-size alkaline batteries; battery-low indicator.
4 When no operation or low signal is detected, the pulse oximeter will power off automatically in 8 seconds.

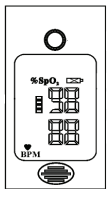
Intended Use

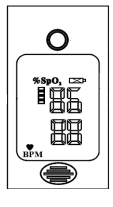
Fingertip pulse oximeter is a portable non-invasive device intended for spot-checking of oxygen saturation of arterial hemoglob in (SpO_2) and pulse rate of adult and pediatric users in sports and aviation.

Operation Instructions

1. Install two AAA batteries according to the Battery Installation instructions.
2. Place one of your fingers into the rubber opening of the pulse oximeter.
3. Press the switch button one time on front panel to turn the pulse oximeter on.
4. Keep your hands still for the reading. Do not shake your finger during the test. It is recommended that you do not move your body while taking a reading.
5. Read the data from the display screen. There are two display modes. After turning on the pulse oximeter, each time you press the power switch, the pulse oximeter will switch to another display mode.







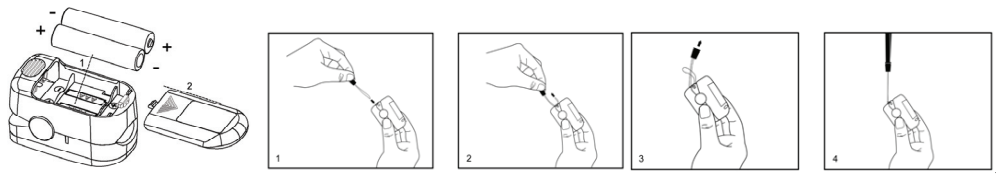
Front Panel

The pulse bar less than 30% indicates signal inadequacy and the displayed SpO_2 and pulse rate value is potentially incorrect.

Battery Installation

1. Install two AAA batteries into the battery compartment. Match the plus (+) and minus (-) signs in the compartment. If the polarities are not matched, damage may be caused to the oximeter.
2. Slide the battery door cover horizontally along the arrow shown as the picture.

Note:
✧ Please remove the batteries if the pulse oximeter will not be used for long periods of time.
✧ Please replace the battery when the power indicator start flickering.



Using the Lanyard

1. Thread thinner end of the lanyard through the loop.
2. Thread thicker end of the lanyard through the threaded end before pulling it tightly.

Warnings!
✧ Please notice that the lanyard which is tied to the oximeter may cause strangulation due to excessive length.

Maintenance and Storage

1. Replace the batteries in a timely manner when low voltage lamp is lighted.
2. Clean surface of the fingertip oximeter before it is used in diagnosis for patients.
3. Remove the batteries if the oximeter is not operated for a long time.
4. It is best to store the product in-4℉~+131℉ and ≤93% humidity
5. Keep in a dry place. Extreme moisture may affect oximeter lifetime and may cause damage.
6. Dispose of battery properly; follow any applicable local battery disposal laws.

Cleaning the fingertip pulse oximeter
Please use medical alcohol to clean the silicone touching the finger inside of oximeter with a soft cloth dampened with 70 % isopropyl alcohol. Also clean the finger being tested using alcohol before and after each test.
Do not pour or spray liquids onto the oximeter, and do not allow any liquid to enter any openings in the device. Allow the oximeter to dry thoroughly before reuse.
The fingertip pulse oximeter requires no routine calibration or maintenance other than replacement of batteries.
The use life of the device is five years when it is used for 15 measurements every day and 10 minutes per one measurement. Stop using and contact local service center if one of the following cases occurs:

- An error in the Possible Problems and solutions is displayed on screen.
- The oximeter cannot be powered on in any case and not the reasons of battery.
- There is a crack on the oximeter or damage on the display resulting readings cannot be identified; the spring is invalid; or the key is unresponsive or unavailable.

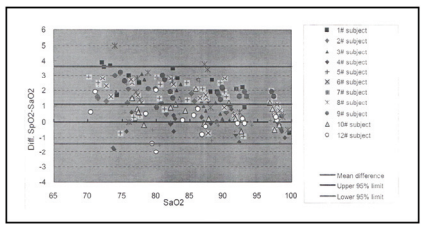
Specifications

1. Display Type
LED/LCD display

2. SpO_2
Display range: 0%~100%
Measurement range: 70%~100%
Accuracy: 70%~100%±2digits; 0%~69% no definition
Resolution: 1%
A_{RMS} Value Analysis

Item	70--100	90--100	80--<90	70--<80
#pts	231	82	89	60
Bias	1.10	0.49	1.35	1.62
A_{RMS}	1.68	1.09	1.77	2.14

Bland-Altman plot analysis of sampled data points on all subjects as below



Note: A functional tester cannot be used to assess the accuracy of a pulse oximeter monitor or sensor. Clinical testing is used to establish the SpO_2 accuracy. The measured arterial hemoglobin saturation value (SpO_2) of the sensors is compared to arterial hemoglobin oxygen (SaO_2) value, determined from blood samples with a laboratory CO-oximeter. The accuracy of the sensors in comparison to the CO-oximeter samples measured over the SpO_2 range of 70%~100%. Accuracy data is calculated using the root-mean-squared (Arms value) for all subjects, per ISO 9919:2005, Medical Electrical Equipment- Particular requirements for the basic safety and essential performance of pulse oximeter.
A functional tester is used to measure how accurately Fingertip Pulse Oximeter is reproducing the specified calibration curve and the PR accuracy.
The model of functional tester is Index2 FLUKE simulator and the version is 2.1.3.

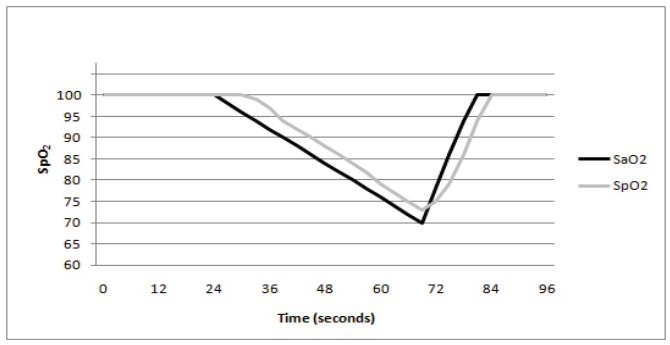
3. Pulse Rate
Display range: 0bpm~250bpm
Measure range: 30bpm~250bpm
Accuracy: 30bpm~99bpm, ±2bpm; 100bpm~250bpm, ±2%
Resolution: 1bpm

4. Probe LED Specifications

	Wavelength	Radiant Power
RED	660 ±2nm	3.2mW
IR	905 ±10nm	2.4mW

NOTE: The information about wavelength range can be especially useful to clinicians.
5. Power Requirements
Two AAA alkaline Batteries
Power consumption: Less than 25mA
Battery Life: Two AAA 1.5V, 1200mAh alkaline batteries could be continuously operated as long as 16 hours.
6. Environment Requirements
Operation Temperature: 5℃～40℃

Storage/ Transport Temperature: -20℃～+55℃
Ambient Humidity:≤80% no condensation in operation;≤93% no condensation in storage/transport
Atmosphere pressure: 86kPa～106kPa
7. Equipment data update period
As shown in the following figure. Data update period of slower average is 8s.



8. Classification
According to the type of protection against electric shock: INTERNALLY POWERED EQUIPMENT;
According to the degree of protection against electric shock: TYPE BF APPLIED PART, (applied part: the rubber hole of the device);
According to the degree of protection against ingress of water: IPX1
According to the mode of operation: CONTINUOUS OPERATION

Possible Problems and Solutions

Problems	Possible reason	Solution
SpO_2 or PR can not be shown normally	1. Finger is not inserted correctly 2. Patient's Oxyhemoglobin value is too low to be measured	1.Retry by inserting the finger 2. Try some more times. If you can make sure no problem exist in the product, please go to a hospital timely for exact diagnosis.
SpO_2 or PR is shown unstably	1. Finger might not be inserted deep enough. 2. Finger is trembling or patient's body is in movement status.	1. Retry by inserting the finger 2. Try not to move
The oximeter can not be powered on	1. Power of batteries might be inadequate or not be there at all. 2. Batteries might be installed incorrectly. 3. The oximeter might be damaged.	1. Please replace batteries 2. Please reinstall the batteries 3. Please contact with local customer service centre
Indication lamps are suddenly off	1. The product is automatically powered off when no signal is detected longer than 8 seconds 2. Power quantity of the batteries is started being inadequate	1. Normal 2. Replace the batteries
"Error3" or "Error4" is displayed on screen	1. Low power 2. Receiving tube being shielded or damaged together with broken connector. 3. Mechanical Misplace for receive-emission tube. 4. Amp circuit malfunctions.	1. Change batteries 2. Please contact local customer service center 3. Please contact local customer service center 4. Please contact local customer service center
Error 6	Err 6 means the screen is failure	Please contact local customer service center
"Error7" is displayed on screen	1. Low power 2. Emission tube damaged. 3. Current control circuit malfunctions.	1 Please change battery 2 Please contact local customer service center 3 Please contact local customer service center

Symbol Definitions

Symbol	Definition	Symbol	Definition
	Type BF applied part.		Attention.
IPX1	Protected against dripping water.	% SpO_2	Oxygen saturation
BPM	Pulse rate (BPM)		Low power indication
SpO_2	No SpO_2 Alarm		Power switch
+55℃ max -20℃ min RH≤93% non-condensing	Storage temperature and relative humidity		Follow instruction for use
	Date of Manufacture	SN	Serial No.
	Waste electrical and electronic equipment		Manufacturer's information

Box Content

1. Fingertip pulse oximeter
2. One lanyard
3. Two AAA batteries
4. One instruction manual

Notes:
1. The illustrations used in this manual may differ slightly from the appearance of the actual product.
2. The specifications are subject to change without prior notice.

Concord Health Supply, Inc.
Off: 888-970-2999
Fax: 888-970-8999
www.ConcordHealthSupply.com

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