Pulse Oximeter

User Manual

CX130

Charmcare Co., Ltd.

Contents

Part 1 Before You Begin

1.1 Overview

This manual contains information for collecting patient oxygen saturation data while operating the CX130

1.2 Intended Audience

This manual provides information to health-care professionals acting as caregivers for operation and user of the monitoring system. Before use, carefully read this manual, accessory Directions for Use, and all precautionary information and specifications.

1.3 Safety Information, Warning, Caution and Note

This is contains safety information requiring users to exercise appropriate caution while using the monitoring system.

1.3.1 Safety Symbol Definitions

Warning

"Warning" is used to refer to factors, which, when ignored, may result in severe and/or fatal injuries and property damage.

Caution

"Caution" is used to refer to factors, which, when ignored, may result in moderate, but non-life-threatening, injuries.

Note

"Note" is used to highlight factors that are not dangerous, but should be paid close attention to during installation, use, and maintenance.

1.3.2 Warning

- Do not use the battery with other manufacturer's batteries.
- Do not use any monitoring system or Pulse Oximeter cables, sensors, or connectors that appear damaged.
- Do not use any monitoring system, sensor, cable, or connector that appears damaged. Remove any damaged equipment from service for inspection by a qualified service technician.
- No modification of this equipment is allowed.

  Do not lift by the sensor or interface cable. The cable may disconnect, potentially dropping the monitoring system on a patient or damaging surface.
- When installing the AC power cord, ensure the cord is carefully positioned to prevent tripping and entanglement.
- Do not spray, pour, or spill any liquid on the monitoring system, its accessories, connectors, switches, or openings in the chassis, since this may cause damage to the monitoring system.
- To ensure accurate performance and prevent device failure, do not subject to extreme moisture, such as direct exposure to rain. Such exposure may cause inaccurate performance or device failure.
- Use only when connected to a grounded outlet to avoid electric shock.
- Do not Pause or disable audible alarms or decrease the volume of the audible alarm if patient safety could be compromised. Do not dim or disable visual alarms if patient safety could be compromised..
- Use of portable phones or other radio frequency (RF) emitting equipment near the system may cause unexpected or adverse operation.
- Operator shall not touch the battery compartment and the patient simultaneously.
- The measured values of the monitoring system can be affected by patient conditions, excessive patient movement, sensors, environmental conditions, and nearby electromagnetic external conditions.
- Ensure the monitoring system is clear of any obstructions that prevent awareness of visual or audible alarms. Failure to do so may result in inadvertently missing a visual alarm or an inaudible alarm tone.
- The use of accessories, transducers and cable other than those specified may result in increased emissions or decreased immunity performance of the equipment.
- The equipment or system should not be used adjacent to, or stacked with, other equipment. If adjacent or stacked use is necessary, the equipment or system should be tested to verify normal operation in the configuration in which it is being used.

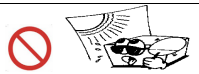
1.3.3 ⚠ Caution

- Accessory equipment connected to the monitoring system's data interface must be certified according to IEC Standard 60950-1 for data-processing equipment. All combinations of equipment must be in compliance with IEC Standard 60601-1:2012 Requirements for Medical Electrical Systems. Anyone who connects additional equipment to the signal input or signal output port configures a medical system and is therefore responsible for ensuring the system complies with the requirements of IEC Standard 60601-1:2012 and IEC Standard 60601-1-2:2007.
- When connecting the monitoring system to any instrument, verify proper operation before clinical use.
- Both the monitoring system and the instrument connected to it must utilize a grounded outlet.
- Any equipment connected to the data interface must be certified according to the latest IEC/EN 60950-1 standard for data-processing equipment, the latest IEC/EN 60601-1 standard for electro-medical equipment, or the latest IEC/EN safety standards relevant to that equipment.
- All combinations of equipment must be in compliance with Requirements for Medical Electrical Systems IEC Standard 60601-1:2007.
- Anyone who connects equipment to the data interface is configuring a medical system and, therefore, is responsible for ensuring the system complies with the Requirements for Medical Electrical Systems IEC/EN Standard 60601-1 and the electromagnetic compatibility IEC/EN Standard 60601-1-2:2007.
- Accuracy may degrade if it is connected to secondary I/O devices when the equipment is not connected to earth reference.
- Use only approved sensors and interface cables when connecting to the sensor port. Connecting any other cable or sensor influences the accuracy of sensor data, which may lead to adverse results.
- The top of the monitoring system screen indicates the sensor type when connecting a recommended sensor to the monitoring system or when the monitoring system completes POST with an attached sensor.
- If the pulse beep tone does not sound with each pulse, the pulse beep volume is set to zero, the speaker is malfunctioning, or the signal is corrupt. Reset the device.
- Perform the following checks every 12 months.
- Inspect the monitoring system for mechanical and functional damage or deterioration.
- Inspect the internal fuses for proper value and rating.

- Ensure all user interface items, cables, and accessories function normally.
- Have a qualified service technician replace the battery at least every 6 months.
- The battery is recyclable. Do not dispose of the battery by placing it in the regular trash.
- Dispose of the battery in accordance with local guidelines and regulations or contact Charmcare to arrange for disposal.
- Inspect the safety relevant labels for legibility. Contact Charmcare or a local representative, if labels are damaged or illegible.
- Instructions provided on preventive inspection, calibration, maintenance and its frequency
- Information provided for safe performance of routine maintenance necessary to ensure continued safe use of ME EQUIPMENT
- Instructions provided to ensure adequate maintenance of ME EQUIPMENT containing rechargeable batteries to be maintained by anyone other than SERVICE PERSONNEL
- Magnetic and electrical fields are capable of interfering with the proper performance of the device.
- X-ray equipment or MRI devices are possible sources of interference as they may emit higher levels of electromagnetic radiation.

1.3.4 Proper Environments for the Product

- Do not use or store the product in the following conditions:

 **Exposure to Humidity and/or Moisture** Do not use the product with wet hands.	 **Do not store or place the product in direct sunlight.**
 **Do not store or place the product in areas where temperature changes are drastic.** Use the product between temperatures of 5°C to 35°C and at a humidity between 5% to 95%.	 **Do not place the product near heat sources.**
 **Do not store or place the product in very humid areas or areas where air circulation is a problem.**	 **Do not subject product to severe shock and or vibrations.**
 **Do not store or place the product in places where product is exposed to chemicals or flammable gas.**	 **Keep product free of dust and debris, particularly metallic objects.**
 **Do not attempt to disassemble the product yourself. Charmcare is not liable for any problems that may occur should you attempt to do so.**	 **Do not connect the power during installation. This may damage the product.**
 **Grip the plug when unplugging the product from a power source.**	

1.3.5 Warranty Period

- To obtain information about a warranty, if any, for this product, contact your local Charmcare representatives.
- This product has been manufactured and inspected following the strict quality assurance guidelines of Charmcare.
- Refer to the Economic Planning Board's "Regulations Regarding Consumer Compensation" for more information on conditions for product repairs and exchanges.
- Product malfunctions occurring from regular use shall be repaired for free at the Charmcare service center during the term of the warranty period.
- During the term of the warranty period, report all problems with the product to Charmcare by including the model no., the device no., date of purchase and a detailed description of the error.
- Manufacturer and/or the store where the product was purchased do not assume any responsibility for any and/or all problems resulting from improper use or improper storage of the product.

1.3.6 Battery Replacement

- If the monitor has not been used for a long period of time, the batter will need charging. To charge the battery, contact your local Charmcare representatives.
- Charmcare recommends that the Lithium Ion Rechargeable battery be replaced at 6 months intervals. Refer to the monitor service manual for battery replacement and general service instruction. Follow local governing ordinances and recycling plans regarding disposal or recycling of their battery and other device components.
- Remove the battery if the equipment is not likely to be used for some time.
- When incorrect battery replacement by inadequately trained personnel would result in an unacceptable risk(e.g. excessive temperatures, fire or explosion)

1.3.7 Contact Us

- Please contact us for better service and products.
- Charmcare service is always open.

Need to purchase products or parts?	**Charmcare Co., Ltd.**
Need service or repairs?	(Gasan-Dong , Woolim Lions 2-cha), 714, 2, Gasandigital1-ro, Geumcheon-Gu, Seoul, Korea
Need technical advice?	Tel : +82-2-862-5052 Fax : +82-2-862-5065
Website	http://www.charmcare.com

1.3.8 Electrical Safety

Please check the following conditions before attempting to use the product.

- Are you using the proper power source line? (100-240VAC)
- Are you using the power source supplied by the manufacturer? (DC 9V, 3.34A)
- Are all parts (power cord and optional parts) connected properly to the product?

Warning

The AC power plug is a means to isolate its circuits electrically from the supply mains on all poles simultaneously. Do not place the device in an area when there is difficult to disconnect from the supply mains.

Warning

Read and check the conditions under “Electrical Safety” before using the product. Failure to do so may result in severe injuries and damage to the product.

Warning

The CX130 Pulse Oximeter is a prescription device and is to be operated by qualified personnel only.

Caution

To prevent noise, install the product away from generators, X-ray machines, speakers and power cords. Proximity of the product to such equipment may result in improper functioning of the product and lead to undesirable results. A separate power circuit and secure grounding of the product are very important. Sharing of a power source with other equipment(s) may lead to undesirable results.

1.3.9 Maintenance and Cleaning

- Using various methods can clean Pulse Oximeter and its accessories. Please follow the methods mentioned below to avoid unnecessary damage or contamination to the Equipment.
- In the event that harmful (unauthorized) materials are used for cleaning, the damaged or contaminated Equipment shall not be serviced without charges regardless of warranty period.

Caution

Please check carefully both frame and sensor, after cleaning the Equipment, Do not use the Equipment that is worn out or damaged.

- At least once a month, clean and wipe off the frame by using the soft cloth after wetting it with lukewarm water and alcohol. Do not use lacquer, thinner, ethylene, or oxides, which could be harmful to the Equipment.
- For surface cleaning and disinfection of the monitoring system, follow institutional procedures or the recommended actions below.
- Surface cleaning – Use a soft cloth dampened with either a commercial, nonabrasive cleaner or a solution of 70% alcohol in water, lightly wiping the surfaces of the monitoring system.
- Disinfection – Use a soft cloth saturated with a solution of 10% chlorine bleach in tap water, lightly wiping the surface of the monitoring system.
- Before attempting to clean a Charmcare sensor, read the Instructions for Use enclosed with the sensor. Each sensor model has cleaning instructions specific to that sensor. Follow the sensor cleaning and disinfecting procedures in the particular sensor’s Instructions for Use.

- Make sure both cables and accessories are free of dust or contaminants, and wipe them off with soft cloth wetted with lukewarm water (40℃ / 104℉), and at least once a week, clean them by using the clinical alcohol.
- Do not submerge the accessories under any liquid or detergent. Also, make sure any liquid not to penetrate into the Equipment or probe.

Caution
Do not dispose single use probe to any hazard place, Always think about environmental contamination

Perform the following checks every 12 months.

- Inspect the monitoring system for mechanical and functional damage or deterioration.
- Inspect the internal fuses for proper value and rating.
- Ensure all user interface items, cables, and accessories function normally.
- At least once a month, clean and wipe off the frame by using the soft cloth after wetting it with lukewarm water and alcohol. Do not use lacquer, thinner, ethylene, or oxides, which could be harmful to the Equipment.

Warning
A functional tester cannot be used to assess the accuracy of a Pulse Oximeter probe or a Pulse Oximeter monitor

1.3.10 Classifications

The CX130 patient monitor is classified, according to IEC 60601-1 as:

Equipment Classification	EN ISO 80601-2-61 : 2012 EN IEC 60601-1:2012
Type of protection	Class II
Degree of protection	BF-Type applied part
Electromagnetic Compatibility	IEC 60601-1-2:2007
Liquid Ingress	IPX1
Degree of Safety	Not suitable for use in the presence of flammable anesthetics

1.3.11 Description of product & Label symbols

Icon	Comments	Icon	Comments
	SYSTEM POWER On/Off		Alarm On/Off
	DC Input Indicator		Connect DC power
	Battery Input Indicator		Use for Program Upgrade
	Type BF applied part complying with IEC 60601-1	**%SPO2**	Oxygen Saturation
	Manufacturer	**IPX 1**	Conforms to IEC60601-1 sub-clause 44.6 and IEC60529 standard.
REF	Reference number		Date of manufacture
SN	Serial number		Crossed-out wheeled bin
CE 0120	CE Mark	EC REP	EU representative
	Do not use hand hooks		Keep away from water
	Fragile, handle with care		This side up
	Caution		Warning
	Refer to instruction manual / booklet		Prohibition
	Note		Class II equipment
	Touch Key Control Button		Menu button

Part 2 Product Summary

2.1 Intended use

The intended use of CX130 is detecting and measurement as below parameters to provide and help doctors for making figure out the patient's vital condition.

- Exact and stable Oxygen saturation of arterial blood
- Accurate Pulse Rate

Especially, CX130 has AGC (Auto Gain Controller) function for weak pulsatile patients.
The device is to detect and control its gain automatically in accordance with patient's pulse rate strength.
Designed for hospital , transport and home use .

Note

Hospital use typically covers such areas as general care floors (GCFs), operating rooms, special procedure areas, intensive and critical care areas within the hospital and in hospital-type facilities. Hospital-type facilities include physician office-based facilities, sleep labs, skilled nursing facilities, surgicenters, and sub-acute centers.

Note

Intra-hospital transport includes transport of a patient within the hospital or hospital-type facility.

▣ Frequently used functions
- Connect the SpO2 probe
- Patient mode setting
- Alarm system (according to IEC 60601-1-8:2012)
- Measure SpO2, Pulse rate
- Battery (Internally powered)

▣ Essential performance

- SpO2 accuracy
- Heart rate accuracy
- Alarm limit
- Battery condition alarm

▣ Patient population

a) Age: newborn, infant, adult

b) Weight: not relevant

c) Health: not relevant

d) Nationality: multiple

e) Patient state: patient is not user – not relevant, unless patient is agitated

▣ Part of the body or type of tissue applied to

a) Measurement site: finger, the back of hand, arm, toe, top of the foot.

b) Condition: the area which wish to place the probe is wiped down with alcohol.

▣ User profile

Intended conditions of use

Considerations	Requirement description
Education	• Nurse who had gotten a specialized education • Physician who have the national license and can treat a patient
Knowledge	• Read and understand 'westernized Arabic' numerals when written in Arial font • Can perceive alarm conditions and distinguish alarm priority • Understand means of the parameter used on the equipment • Can distinguish SpO2 and Pulse rate • Understands hygiene
Language understanding	• Understand how the equipment functions in the IFU • Understand abbreviations used in the IFU
Experience	• A training related to how the equipment functions • Be enough familiar with the IFU • Can use SpO2 probe correctly • Can diagnose patient states through measured parameters

▣ Intended conditions of use

Considerations		Requirement description
Environment	General	• Professional use, not intended for home use • Indoor use only • When it is functioning it shall keep its calibration / precision • Use environment away from X-ray equipment, any mobile device • Battery charging and replace for maintaining normal operation
	Conditions of visibility	• Operation distance: within 1m
	Physical	• Operating environment temperature: 5℃ to 35℃ humidity: 5% to 95% • Storage environment temperature: -20℃ to 70℃ humidity: 10% ~ 100% • Background sound pressure level: <55dB in the range of 100Hz – 8kHz
Frequency of use	• Measure patient state continuously	
Mobility	• Portable medical device to be used on a resting patient	

2.2 List of Parts

▣ Basic Parts

Type	Quantity
CX130 Unit	1

▣ Basic Accessories

Type	Quantity
SpO2 Sensor	1
AA Alkaline battery or 3.7V Lithium Ion Rechargeable battery, DC Power Unit, Power Cord	4 or each 1
User Manual	1

▣ Optional Parts

Type	
1	SpO2 Sensor(Pediatric, Neonate)
2	SpO2 Extension cable
3	DC Power Unit
4	Power Cord
5.	IV Pole Bracket
6	Battery

▣ Sensor and Cable Length

Type	Length
SpO2 Adult Sensor	1m, 3m
SpO2 Pediatric Sensor	1m, 3m
SpO2 Neonate Sensor	1.5m, 3m
SpO2 Neonate Sensor	1m, 3m
SpO2 Extension cable	2.5m
Power Cord	1.8m
DC Power Unit	2m

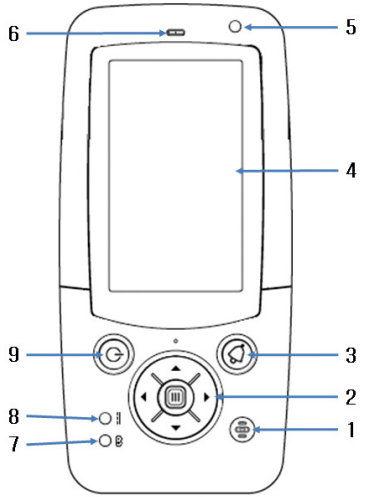
Warning

The use of accessories, pulse oximetry sensors, and cables other than those specified may result in inaccurate readings of the monitoring system and increased emission of the monitoring system.

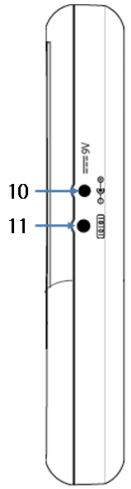
2.3 Nomenclature of the Parts

This section identifies the symbols, controls, displays, and indicators on the CX130

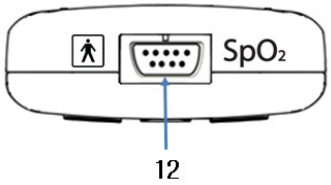
▣ Buttons & Front



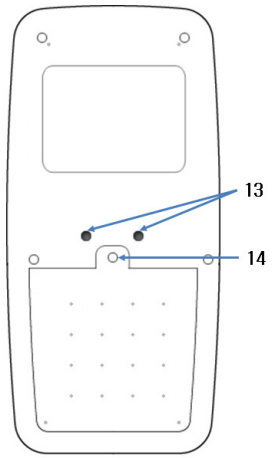
▣ Right side



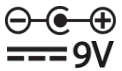
▣ Upper side

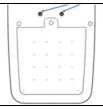


▣ Rear side



No.	Indicator	Name	Description
1		Speaker	The speaker indicates audio alarms. Care should be taken not to cover the speaker and muffle the audible alarm volume.
2		Touch Key Control Button	Press a Touch Key Control Button to select the corresponding touch key icon. See Part 4, Touch Key Control Buttons and Icons for more details.
3		Alarm Paused button	Press the Alarm Paused button to temporarily Pause patient and low battery alarms. Press the Alarm Paused button when the SENSOR OFF or FINGER OFF messages are flashing (i.e. the sensor is removed from the patient) to acknowledge the end of monitoring. In these states, all further alarms are suspended until the Pulse Oximeter starts measuring SpO2 and pulse rate again. Note: System failure alarms can be Paused by pressing the Power/Standby or Alarm Paused button. If the Power/Standby Button does not Pause the system fault alarm, press the Alarm Paused button.
4		SpO2 Measurement Display	The functional arterial hemoglobin oxygen saturation is displayed in units of percentage SpO2. The upper and lower SpO2 alarm limits are also displayed next to the SpO2 measurement. When a sensor is not connected to a patient the display will show dashed lines. When the measured value is outside of the alarm limits, the SpO2 Measurement Display flashes and an alarm will sound. The oxygen saturation is calculated and the display is updated at a frequency of once per second.

No.	Indicator	Name	Description
5		Visual Alarm Indicator	The Visual Alarm Indicator is illuminated when an alarm condition is active and the Alarm Status Indicator is shown.
6		Buzzer	A buzzer is an audio signaling device.
7		Battery Status Indicator	The Battery Status Indicators show the capacity of the CX130 Handheld Station batteries. The indicator flashes when less than 15 minutes of battery life is left and the battery needs to be recharged.
8		DC Power Usage Indicator	Indicates connection to alternating current power source.
9		Power Button	Press the Power/On/Off Button to turn the instrument on. Press, hold the button for more than 2 seconds and then release the button to turn the instrument off.
10		DC Power Input port	The power entry module contains the input connector for DC power. The DC input provides power to the system from the DC line. Always connect the Pulse Oximeter to the mains power for continuous operation and/ or battery recharging. Note : Use the power cord as the means to disconnect the instrument from the mains power supply.
11		RS-232 Port	Software upgrade port
12		SpO2 Probe Port	Connect the patient cable to the Handheld CX130 by PATIENT CABLE plugging the cable into the Patient Cable Connector. Use only CONNECTOR CHARMCARE compatible sensors and cables with this

			oximeter. See Part4, Sensors and Patient Cables for more details.
13		IV Pole Clamp Hole	Attaching clamp holes,
14		Battery Container Cover Hole	If device use AA batteries, can exchange AA batteries as open cover .

Warning

To prevent electric shock, keep the top cover closed and do not attempt to disassemble the product on your own. Product should only be disabled by authorized service personnel.

Part 3 Product Installation

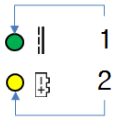
First-time users: Read the installation instructions thoroughly and install the product in a safe place to ensure product longevity.

Caution When Installing

- Use the product in a place where the temperature is between 5~35℃ and the humidity is 5~95%.
- Make sure that the power cord is plugged in properly.
- Do not plug in more than one device into one outlet.
- Do not use a power cord that causes noise.
- The product is very sensitive to shock. Caution is advised.
- Keep the product dust-free and install away from flammable materials.

Connecting the Power

- Connecting the Power – Connect the power to the power port in the side of the product.
- When connecting the enclosed authentic DC power adaptor to the DC power adaptor connection jack located on the rear of the product, a green light appears in the DC Power Usage Indicator. Product usage is possible without DC power adaptor connection via the embedded battery. In such cases, a red light appears in the Battery Usage Indicator.



① DC Power Usage Indicator

② Battery Usage Indicator

Caution

When the charge level of the embedded battery is running low, the battery usage indicator blinks and an information alarm is emitted. Depending on usage parameters, power may be cut instantly, and thus the DC power adaptor should be connected for use.

Caution

Both the monitoring system and the instrument connected to it must utilize a grounded outlet.

Warning

The battery usage warranty period is six months. Following this period, battery life can diminish significantly.

Warning

Use the power supply device supplied by the manufacturer. Failure to do so may result in electric shock and damage to the product. Edge or points of the Product can hurt to patient and user.

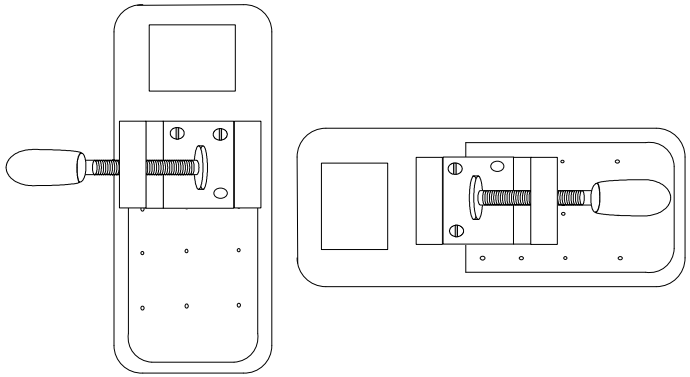
Warning

Accuracy may degrade if it is connected to secondary I/O devices when the equipment is not connected to earth reference.

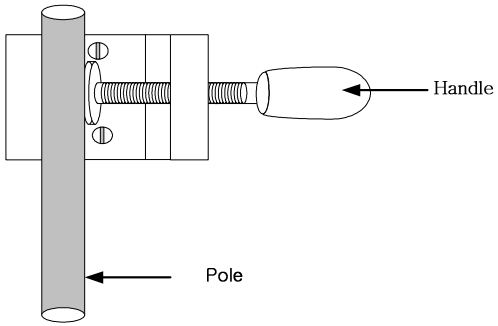
Installing the IV Pole

The product may also be mounted on a stand, which can then be attached to an IV Pole. This makes the product more portable. (*IV Pole is optional.)

Step 1) The product may be used horizontally or vertically. First, decide how you want the product to be mounted. Then, insert two bolts into the holes on the stand and tighten.



Step 2) Mount the stand to the pole by sliding the stand onto the pole and then, turn the handle to tighten



Caution

The pole on which the stand is mounted should be less than 30mm in diameter. Do not use poles that are any thicker than this.

Part 4 Using The Product

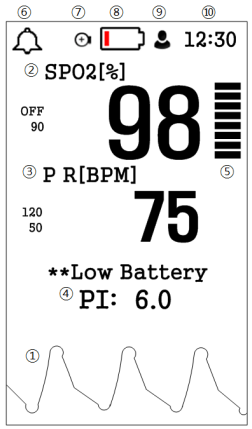
This section contains information on the basic nomenclature and directions for using the product.

- Check for defects on the exterior of the SpO2 sensor(Oxygen sensor).
- Check for defects on the exterior of the AC power cord, oximetry use cable.
- Connect the power, oximeter use cable to the SpO2 sensor and product. (If the oximetry use cable is not utilized, SpO2 sensor is directly connected to the product.)
- Check the connection status of the cable and status of SpO2 sensor before turning the product on and connecting the product to the patient.
- Since this product uses the principle of a spectrophotometer, check if a surgery light is located where the product is placed, and if the ray blocks the usability cover the SpO2 sensor with a opaque cloth.
- Check the attachment region of the SpO2 sensor and the compatibility, and avoid certain locations according to the status and use a SpO2 sensor that can be attached on to the bridge of the nose or forehead.
- When attaching the SpO2 sensor on to the finger, no manicure should be on the finger nails.
- For sterilization or antiseptic. cleaning is necessary for the SpO2, select the methods shown below according to the environment conditions of the medical institution equipment, and use the usage and application of the professional prefers.

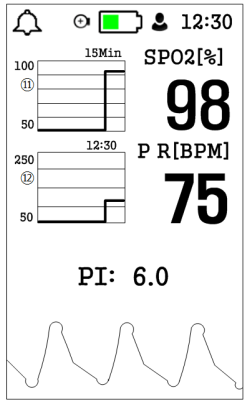
*How to use antiseptic .cleaning solution

Chemical Product		Method
Alcohol	Isopropyl alcohol	Use soaked gauze.
Liquid Soap	Benzalkonium Chloride	0.05 W/V%(x200 diluted) Use soaked gauze.
Iodine	Povidone-iodine	0.02 W/V%(x50 diluted) Use soaked gauze.
Glutaral	C5H8O2	2 W/V %(test solution) Use soaked gauze.

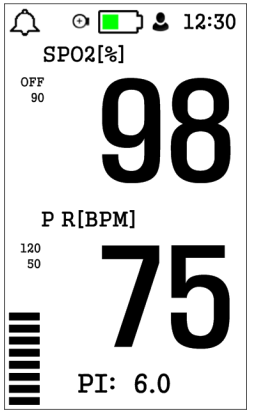
Screen



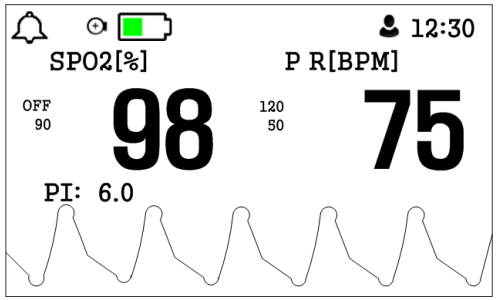
Screen1



Screen2



Screen3



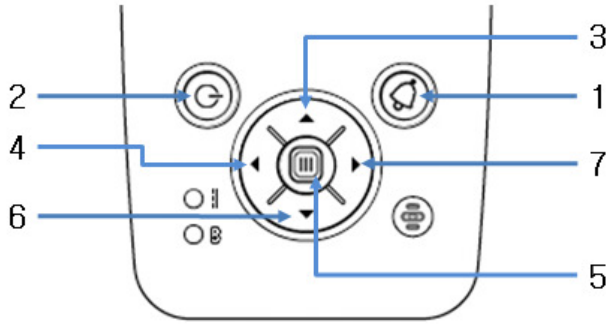
Screen4

No.	Name	Description
1	Pulse Waveform Display	The Pulse Waveform Display shown the acquired plethysmographic waveform. The plethysmographic waveform is scaled with signal strength. Signal strength is defined as the relation of arterial pulsatile signal to the non-pulsatile
2	Saturation SpO2 Alarm Display	The Saturation Alarm Limits Display shows the upper and lower saturation alarm limits. When an alarm limit is exceeded, the SpO2 value and the violated limit flashes.
3	Pulse Rate Alarm Limits Displays	The Pulse Rate Alarm Limits Displays Shows the upper and lower pulse rate value and the violated limit flashes.
4	Perfusion Index Display(PI)	The Perfusion Index indicates numerically the percentage of pulsatile signal to non-pulsatile signal (Pulse strength).
5	Level Bar	Indicates pulse beat and the relative (non-normalized) pulse amplitude in numbers only view. As the detected pulse becomes stronger, more bars light with each pulse.
6	Menu Bar	Shows the current time, alarm, battery status and other main menu items.
7	SpO2 Short Trend	Continuously displays the latest SpO2 measurements of the last 15 minutes on the screen.
8	PR Short Trend	Continuously displays the latest PR measurements of the last 15 minutes on the screen.

Symbol	Description
	Displays the alarm enabled or disabled status.
	 Uses DC adaptor power only but battery is fully charged  Uses DC adaptor power and battery is being charged  Uses a battery and capacity is FULL  Uses a battery and capacity is 1/2  Uses a battery and capacity is 1/4  Uses a battery and capacity is low(about10min)
12 : 30	The Time Indicator displays the current time. The time is displayed in 12 or 24 hour format. Select the time display formats in the time menu.
	The coin battery capacity is not enough.
	Adult Patient Icon
	Pediatric Patient Icon
	Neonate Patient Icon

Button

Single buttons can be used for speedy and convenient use of functions from the product menu.



No.	Name	Description
1	Alarm Button	On the main screen, in the cases of high priority and mid priority alarms, the alarm will not ring for 2 minutes when you press the alarm button. At the menu screen, this action cancels the current menu and returns you to the previous menu.
2	Power Button	Used to turn on/off the power to the product
3	▲ Up Button	Used to change the product screen; Press once per instance to switch screens following a sequence of Screen 1→ Screen 2→ Screen 3→ Screen 4
4	◀ Left Button	Press ◀Button
5	MENU Button	Used to enter the SETUP screen
6	▼ Down Button	Press ▼Button
7	▶ Right Button	Press ▶Button

Post-Use Storage and Management

- After use, store the product in an area that does not exceed 85% humidity, and maintain the temperature in the range of 5°C ~ 35°C.
- Do not store the product together with chemical products and/or in an area subject to gas exposure.
- Avoid exposing the product directly or indirectly to heat.
- Make sure to review the product usage instructions prior to use.
- Store the product in an area where it will not come in contact with water.
- Store the product in an area free of dust and/or other foreign substances.
- Store the product away from direct sunlight.
- Maintain the product in a safe and stable condition away from vibration and/or other sources of shock.
- If the product and/or sensor area becomes dirty, wipe it clean with rubbing alcohol and cotton swabs, and then allow it to dry at room temperature for 30 minutes.

Power Off

- Press power button during 3seconds to power off. When display is off, separate the DC Power Unit, Power Cord by pulling off the cords connector from device side .

Part 5 Measuring The SpO2

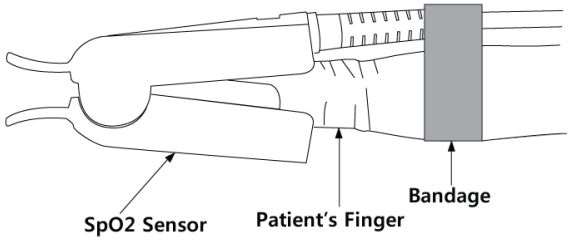
Attaching the SpO2 Probe

Step 1) Wipe down the area where you wish to place the probe with alcohol.

Step 2) Attach the probe to the patient's finger.

Step 3) To get an accurate reading, make sure the patient minimizes all movement and Please attach the probe wire to patient's finger firmly. Attach the bandage loosely so as to not cut of circulation to the finger.

Step 4) Check the patient's finger and the probe every two to three hours to make sure that the sensor is properly placed over the finger. If there is a change in the appearance of the patient's finger due to the prolonged exposure to the probe, switch the probe to another finger.



Warning

Measuring SpO2 on a patient undergoing an MRI may result in severe burns for the patient. To minimize risk for burns, use a non-inductive wire. In the event that this does occur, immediately remove the probe from the patient.

The area around the SpO2 sensor shall not exceed 37℃. The sensor will not work in temperatures above 37℃.

Do not attach the probe near arterial or venous catheters.

Make sure that the sensor emits a light and that the sensor is properly placed over the patient's finger.

Excessive pressure for prolonged periods from the sensor may cause necrosis of the skin.

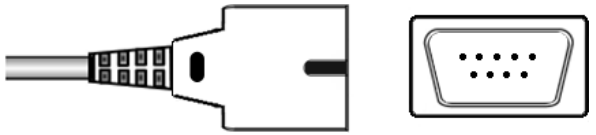
Sensor sites should be changed at least every 8 hours

SpO2 Sensor Port

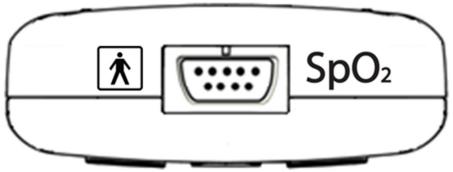
Use only approved sensors and interface cables when connecting to the sensor port. Connecting any other cable or sensor influences the accuracy of sensor data, which may lead to adverse results.

If the pulse beep tone does not sound with each pulse, the pulse beep volume is set to zero, the speaker is malfunctioning, or the signal is corrupt. Reset the device.

- Wire connection



- SpO2 sensor Port



Warning

Before use the probe sensor, operator needs to verify that the SpO2 sensor type match

Caution

Handle the probe sensor and wire with caution. Careless handling may damage the sensitive sensor. Protect the wire from sharp objects.

The skin of patients who have high fevers or have problems with distal circulation will be 2-3 degrees higher than normal.

Patients with abnormally high oxyhemoglobin or methemoglobin levels will not give a proper SpO2 reading.

Note

Taking the NIBP can affect the SpO2 reading. When taking NIBP, place the SpO2 probe on the other arm.

Avoid using the probe with other medical equipment that affects blood flow. Avoid placing the probe near an area that requires medical attention.

Measuring the SpO2

Step 1) Insert 2 x AA battery or connect adapter to adapter jack on the side.

Step 2) Connect SpO2 sensor to upper side.

Step 3) Press the power button.

Step 4) After all LCD indications are lightened up, the measurements are ready.

Step 5) Attach the SpO2 sensor to finger.

Step 6) When the display reads “Learning,” this means that SpO2 measurement has begun. Minimize patient movement until the screen no longer reads “Learning” to ensure a proper SpO2 reading.

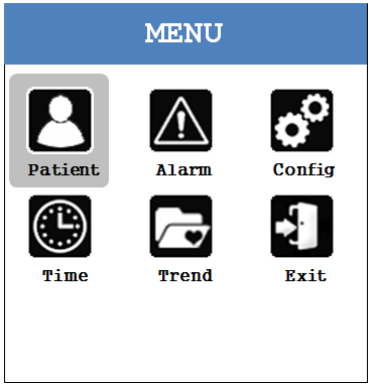
Step 7) After receiving data for 3secs, SpO2 results will be indicated.

Part 6 How To Use Menu

The SETUP menu allows for the adjustment of various settings including Current Time, Patient Mode, alarm Limits, Alarm Volume, Pulse Volume, Backlight, Trend, etc.

Entering the SETUP menu

Step1) Press the menu button to enter the MENU screen.



Setting the Current Time

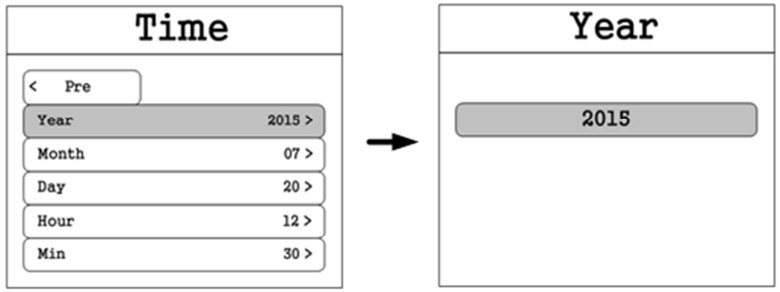
Step 1) Use the directional buttons of ▲ (up), ▼ (down), ▶ (right) and ◀ (left) to choose a menu and designate a “Time”.

Step 2) Click the menu button once to enter the selected menu.

Step 3) Press the ▲ (up) or ▼ (down) button to move between menus and click the menu button once you have moved to the menu you wish to select.

Step 4) Use the ▲ (up) or ▼ (down) button to adjust the value you wish to set. Press the ▼ (down) button to reduce the value and the ▲ (up) button to increase the value. If you press and hold the button, the value increases rapidly.

Step 5) Pressing the menu button allows you to return to the previous screen and selecting the "Pre" menu and then pressing the menu button brings you back to the initial menu screen. Pressing the ALARM button cancels the current settings and takes you back to the previous menu.



*Current Time Setting Ranges

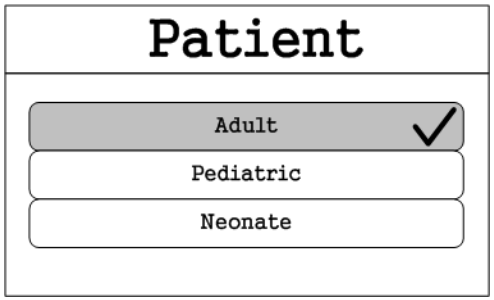
Setting	Year	Month	Day	Hour	Minute
Range	2000~2099	1~12	1~31	0~23	0~59

Patient Mode Setting

Step 1) Use the directional buttons of ▲ (up), ▼ (down), ▶ (right) and ◀ (left) to choose a menu and designate a “Patient”.

Step 2) Click the menu button once to enter the selected menu.

Step 3) Press the ▲ (up) or ▼ (down) button to move between menus and press the menu button at the desired value to save the value and return to the previous menu. Pressing the ALARM button cancels the current settings and takes you back to the previous menu. The current value is displayed with a ✓.



Alarm Limit Setting

The factory default setting provides both visual and audible alarms for alarm conditions.

When the monitoring system detects an error condition, it displays the alarm message, suggests corrective action, and sounds an alarm.

The monitoring system automatically uses alarm limit settings at power-on. Any temporary changes to the Alarm menu remain active until power off or power loss greater than 30 seconds.

Step 1) Use the directional buttons of ▲ (up), ▼ (down), ▶ (right) and ◀ (left) to choose a menu in MENU, and designate a “Alarm”.

Step 2) Click the menu button once to enter the selected menu.

Step 3) Press the ▲ (up) or ▼ (down) button to move between menus and click the menu button once you have moved to the menu you wish to select.

Step 4) Use the ▲ (up) or ▼ (down) button to adjust the value you wish to set. Press the ▼ (down) button to reduce the value and the ▲ (up) button to increase the value. If you press and hold the button, the value increases or decrease rapidly.

Step 5) Pressing the menu button allows you to return to the previous screen and selecting the "Pre" menu and then pressing the menu button brings you back to the initial menu screen. Pressing the ALARM button cancels the current settings and takes you back to the previous menu.

Managing Alarms and Alarm Limits

When the monitoring system detects certain conditions that require user attention, the monitoring system enters an alarm state. The monitoring system uses both visual and audible indicators to identify high-priority, low-priority alarms. High priority alarms take precedence over low-priority alarms.

- Patient Alarm : This alarm occurs from the patient data. Occurs when the measured physiological signal value initiate the limit alarm or alerts through the patient status.
- Technical Alarm : This alarm occurs from the system. Occurs for product related actions or system errors.
- Information alarm : Information alarm notifies user on necessary information or when something is wrong at measuring. For example, it alarms when the product is running out of battery.

  The alarm does not occur and shows a related message to the user. This is information such as “LEARNING” and “ MOVING” that starts the measurement after sensing a signal.

Alarm Levels

- High priority alarm : The high priority alarm is activated only when measurement values are highly abnormal. For example, when the heart rate in adult mode is higher than 180 or lower than 40, the high priority alarm is activated regardless of alarm limit settings.
- Medium Priority Alarm: The medium priority alarm is activated only when measurement values fall outside the alarm limit settings.
- Visual Alarm Indicators : Visual alarms appear on the screen in order of highest priority, regardless of any audible alarm status

Basic Alarm Settings and Ranges

1) Adult

	Basic Settings		Alarm Setting Range	
	HIGH	LOW	HIGH	LOW
HR	120	50	15 ~ 300, OFF	OFF, 15 ~ 300
SpO2	OFF	90	50 ~ 100, OFF	OFF, 50 ~ 100

2) Pediatric

	Basic Settings		Alarm Setting Range	
	HIGH	LOW	HIGH	LOW
HR	160	75	15 ~ 300, OFF	OFF, 15 ~ 300
SpO2	OFF	90	50 ~ 100, OFF	OFF, 50 ~ 100

3) Neonatal

	Basic Settings		Alarm Setting Range	
	HIGH	LOW	HIGH	LOW
HR	200	100	15 ~ 300, OFF	OFF, 15 ~ 300
SpO2	OFF	90	50 ~ 100, OFF	OFF, 50 ~ 100

Characteristics of alarm indicator lights

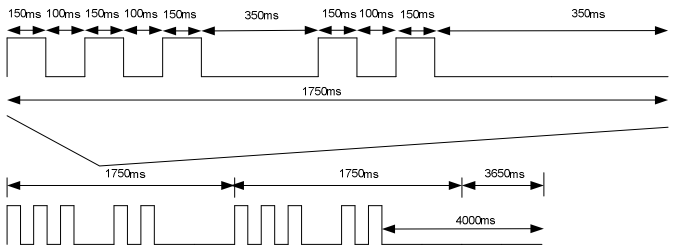
Alarm category	Indicator color	Flashing frequency	Duty cycle
High	Red	2Hz	60% on
Medium	Yellow	1Hz	60% on
Low	Yellow	Constant	100% on

Alarm Tone Definitions

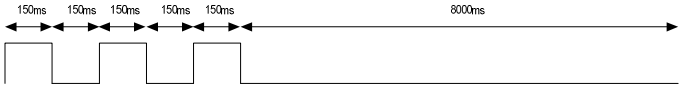
Tone Category	Description
High priority alarm signal	
Pulse Frequency	1000Hz
Pulse width	150ms(IEC60601-1-8)
Number of pulse in burst	10, interburst interval of 4 seconds (IEC60601-1-8)
Repetitions	continually
Sound level range	70~76dB
Medium priority alarm signal	
Pulse Frequency	666Hz
Pulse width	150ms(IEC60601-1-8)
Number of pulse in burst	3, interburst interval of 8 seconds (IEC60601-1-8)
Repetitions	continually
Sound level range	60~67dB
Low priority alarm signal	
Pulse Frequency	500Hz
Pulse width	250ms(IEC60601-1-8)
Number of pulse in burst	2, interburst interval of 16 seconds (IEC60601-1-8)
Repetitions	continually
Sound level range	57~62dB

Priority Signal Timing

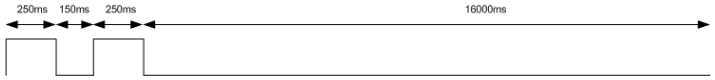
1) High priority



2) Medium priority



3) Low priority



Alarm Conditions

Alarm condition	Auditory alarm	Alarm signal	Messages
High heart rate limits violated	Medium	Visual alarm Auditory alarm	High PR
Low heart rate limits violated	Medium	Visual alarm Auditory alarm	Low PR
High SpO2 limits violated	Medium	Visual alarm Auditory alarm	High SpO2
Low SpO2 limits violated	Medium	Visual alarm Auditory alarm	Low SpO2
Unable to determine pulse rate or SpO2.	Low	Visual alarm Auditory alarm	Check Probe

Communication Error	High	Visual alarm Auditory alarm	SpO2 fault
SpO2 cable/sensor disconnect	Low	Visual alarm Auditory alarm	Sensor off
Finger off form SpO2 sensor	Low	Visual alarm Auditory alarm	Finger off
Low battery	Medium	Visual alarm Auditory alarm	Low Battery
Low Internal Coin Battery	Low	Visual alarm Auditory alarm	BATT ERROR
Clock settings lost (Invalid date and time value detected during start up)	Low	Visual alarm Auditory alarm	CLOCK ERROR

Alarm Condition Delay

*Average Time

Average time 2

- Responds to changes in blood oxygen saturation in 2 to 4 seconds.

Average time 4

- Responds to changes in blood oxygen saturation in 4 to 8 seconds.

Average time 8

- Responds to changes in blood oxygen saturation in 8 to 16 seconds.

Average time 12

- Responds to changes in blood oxygen saturation in 12 to 24 seconds.

Audible Alarm Indicators

- Alarm Status Symbol

Alarm Icon	Status
	Alarm monitoring
	Alarm sound paused
	Alarm sound OFF , Remind signal (interval of 3 min.)

- Alarm sound paused

  Alarm occur condition is exceed SpO2 high or low, PR high or low set value. It can set to the MENU > Alarm

  Press the  button when the alarm occurs and the alarm sound is paused. The alarm does not sound when it is paused. During the audible alarm pause period, you can press Alarm button again to re-enable the audible alarm tone. Alarm pause time is set to MENU > Config > Alarm Duration.

  For any alarm condition still active for more than two (2) minutes, the monitoring system will increase the urgency level of the audible alarm signal by increasing its frequency.

- Remind signal

  Remind signal is implemented to give an audible signal at 3 min interval.

- There are operations according to the alarm when alarm sound is paused.
  - Patient Alarm : Stops the alarm sound during ALARM DURATION
  - Technical Alarm : When SENSOR OFF, FINGER OFF, LOW BATTERY occurs,

  Alarm sound stops and is maintained before each state is canceled.

Warning

Do not pause button the audible alarm or decrease its volume if patient safety could be compromised.

Warning

Each time the monitor is used, check alarm limits to make sure that they are appropriate for the patient being monitored.

- Alarm sound off

  This turns the alarm souund off. It operates the alarm message and alarm lamp when an alarm occurs.

  Available alarm volume to "0" by after set to the MENU > Config > Maintenance > enter password > Alarm 0 use to "ON", then set MENU > CONFIG > Alarm or Pulse Volume to "OFF".

  The alarm sound stops, and the alarm watch status of SpO2 and PR switches to  0.

  When the alarm sound is off, an alarm sounds every 120 seconds to alert the current alarm status. This can be set when the alarm occurs and when MENU > Config > Alarm Duration is checked.

*When initially powered on the device, finger off and sensor off alarm does not work. It work after measuring SpO2 by patient finger.

To Pause an Audible Alarm

- Press the **Alarm Paused button**.
- To cancel, press the **Alarm Paused button** again.
- To re-enable the audio tones during the **Alarm Audio Paused** period, press the Alarm Paused button again. If another alarm occurs during the **Alarm Audio Paused** period, the monitoring system re-enables all audio tones.

- If the **Alarm Audio Paused** period is enabled, the audible alarm is not active for the specified time interval(Alarm Duration) and the **Alarm Audio Paused** icon appears above the appropriate alarm limit icon.

Warning

Ensure the speaker is clear of any obstruction. Failure to do so could result in an inaudible alarm tone.

Warning

A potential hazard can exist if different alarm presets are used for the same or similar equipment in any single area

Warning

Auditory alarm signal sound pressure levels that are less than ambient levels can impede operator recognition of alarm conditions

Note

As the alarm sounds emitted are difficult to describe in words, it is recommended that the user listen directly to the sounds of the high priority and medium priority alarms prior to product use.

Caution

When turning alarm sound OFF, you should take extra caution because there is a chance that you could not recognize the status of patient. If you have to turn it OFF, check on patient frequently.

Caution

If alarm limits set to extreme values, that can render the alarm system useless.

Verifying Visual and Audible Alarm Indication

If the device fails to perform as specified in this test, contact qualified service personnel or your local supplier for assistance.

You can verify the alarm operation for all parameters SpO2 by following the below procedures.

(1) Connect the device to an DC Power Unit or AA size Alkaline battery.

(2) Press Power button to turn on the device

(3) Connect the simulator to sensor input cable and connect cable to device

(4) Set the simulator to smaller value than the lower alarm limit on the device.

(5) Verify following the device reaction:

a. The device begins to track the physiological signal from the simulator.

b. After about 10 to 20 seconds, the device displays the value measured as specified by simulator. Verify values are within the tolerances specified in Specification section for each parameter.

c. Audible alarm sounds.

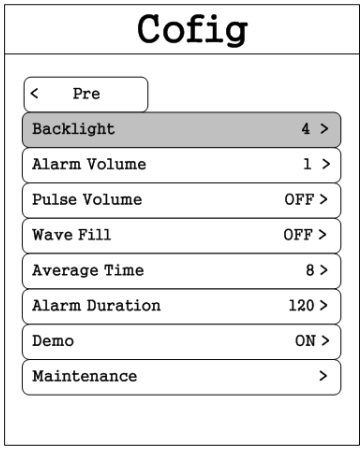
d. The numerical area flashes, indicating the parameter has violated default alarm limits.

Note

It is recommended that you listen to the high and medium alarms beforehand in order to be cognizant of the sounds when the alarms do go off.

CONFIG Setting

CONFIG Setting is used to adjust the settings for Backlight, Alarm Volume, Pulse Volume, Wave Fill, Average Time, Alarm Duration, Demo, Maintenance.



Step 1) Use the directional buttons of ▲ (up), ▼ (down), ▶ (right) and ◀ (left) to choose a menu and designate a “CONFIG” by press menu button.

Step 2) Click the menu button once to enter the selected menu.

Step 3) Press the ▲ (up) or ▼ (down) button to move between menus and click the menu button once you have moved to the menu you wish to select.

Step 4) Use the ▲ (up) or ▼ (down) button to adjust the value you wish to set. Press the ▼ (down) button to reduce the value and the ▲ (up) button to increase the value. If you press and hold the button, the value increases rapidly.

Step 5) Pressing the menu button allows you to return to the previous screen and selecting the "Pre" menu and then pressing the menu button brings you back to the initial menu screen. Pressing the ALARM button cancels the current settings and takes you back to the previous menu.

*Range for Each Setting

Setting	Description	Range
Backlight	Sets the LCD backlight to ON/OFF	1~10
Alarm Volume	Adjusts the alarm volume	OFF, 1~7
Pulse Volume	Adjusts the pulse volume	OFF, 1~7
Wave Fill	Sets the drawing of wave fill	ON, OFF
Average Time	Sets the measurement speed (average time)	2, 4, 8, 12
Alarm Duration	Set the duration (second) when in alarm sound pause and alarm sound off.	30, 60, 90,120
Demo	Runs Demo mode	On, OFF
Maintenance	Factory Default, User Config Save, Alarm 0 Use, Alarm 0 Remind	-

⚠ Caution

Take care in setting alarm volume, as the user will not be able to ascertain the occurrence of any alarm if the alarm volume is turned off. Thus, if the alarm volume is turned off, make sure to check on the patient status frequently.

Exiting the SETUP Menu

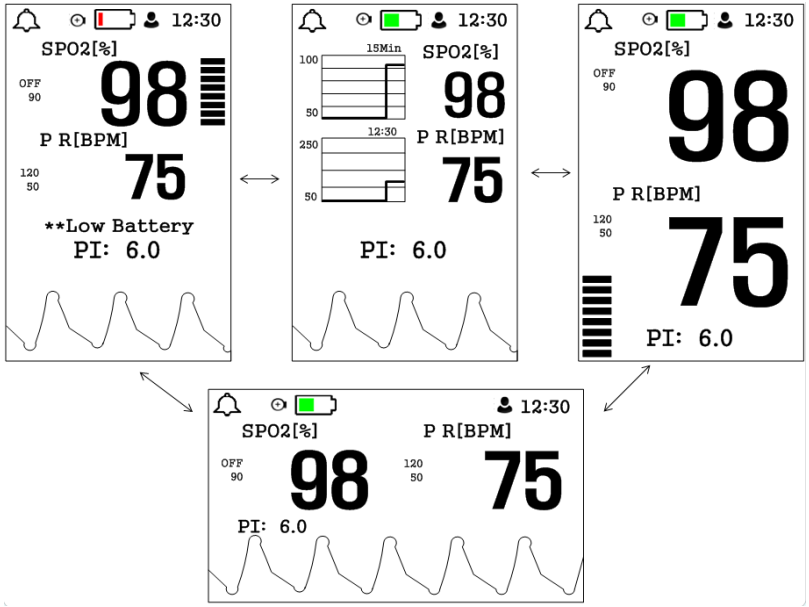
Press the menu button once on "Pre" or press the ALARM button to return to the main menu.

Part 7 Using DISPLAY Menu

Changing the Measurement Screen

The CX130 has 4 screen modes, allowing the user use the menu of their choice.

To change screen, use the directional buttons of ▲ (up) on initial screen.



The SpO2 parameter is calibrated to read functional arterial oxygen saturation. Significant levels of dysfunctional hemoglobins such as carboxyhemoglobin or methemoglobin may affect the accuracy of the SpO2 measurement.

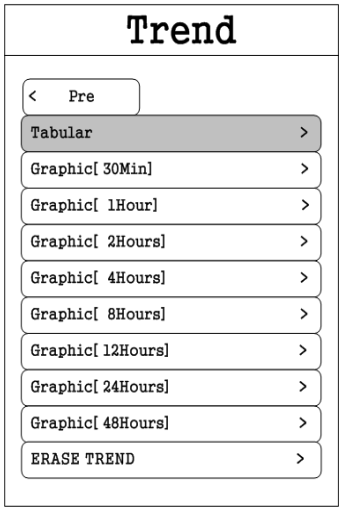
The movement of the plethysmographic waveform are visual indicators of real-time data. The pulse beep tone is an audible indicator of the real-time patient data.

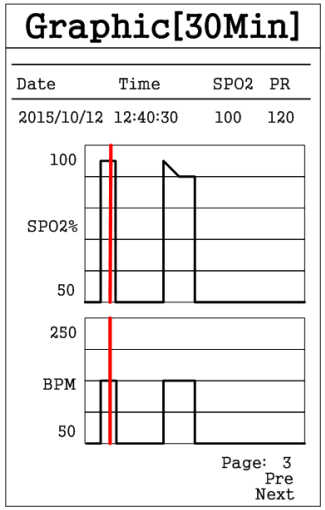
Part 8 Using TREND Menu

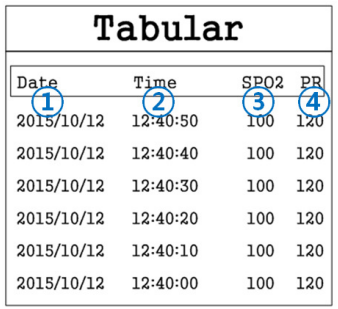
Entering the TREND Screen

Step 1) Press the menu button to enter the MENU screen.

Step 2) Press the ▲ (up) or ▼ (down) button to choose a Trend and press the menu button.







No.	Description
1	Displays measurement date
2	Displays measurement time.
3	Displays SpO2 value of measurement
4	Displays PR value of measurement.

* Data update period

During normal measurement conditions in the Normal response mode, the averaging time is 6 to 7 second and the response time is 2 to 4 seconds in Fast mode. The sum of alarm condition delay and alarm signal generation delay less than 1 second.

*ERASE TREND

Erase functional is erase all data for patient trend in CX130 device.

MENU > Trend > ERASE TREND

Using the Buttons

* In TABULAR TREND Screen

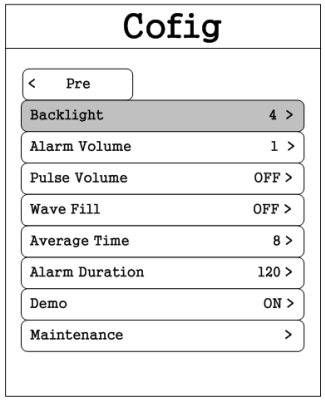
Button	Description
▶ Right	Displays the TREND based on current time
▲ Up	Displays the TREND with an increase in time based on the currently viewed time
▼ Down	Displays the TREND with a decrease in time based on the currently viewed time
◀ Left	Prints the patient measurement value of the currently viewed TREND based on the most recent time

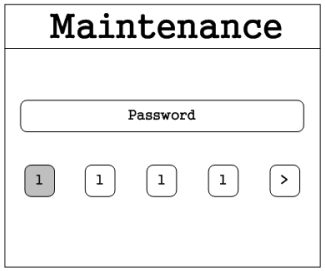
* In GRAPHIC TREND Screen

Button	Description
◀ Left	Moves the value bar once to the left and displays the TREND of the location indicated by the value bar.
▶ Right	Moves the value bar once to the right and displays the TREND of the location indicated by the value bar.
▲ Up	Moves to the TREND before the currently displayed TREND. You can check whether a change has occurred through the Page numbers.
▼ Down	Moves to the TREND after the currently displayed TREND. You can check whether a change has occurred through the Page numbers.

Part 9 Using The Maintenance

Entering Maintenance Menu





Step1) Press the menu button to enter the MENU screen.

Step2) Press the button to choose a "Config" and press the menu button.

Step3) Press the button to choose a Maintenance and press the menu button.

Step4) You can press the ◀(Left) button, ▶(Right) button to go to each place and change the number by ▲ (Up) button, ▼ (Down) button. Press the Menu button on ">" and can into Maintenance menu.

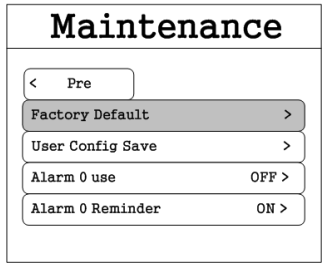
(3) If enter Incorrect PASSWORD, display again the Config menu.

(4) Press the menu button on the "Pre" menu, will leave the Maintenance.

⚠ Caution

MAINTENANCE setting can be manipulated by authorized personnel only. Be aware that the contents of MAINTENANCE are of nature that is not to be altered by normal users.

Setup Maintenance



(1) Press the ▲ (up) button or ▼ (down) button to move to the set.

(2) You can adjust the settings in the item set press by menu button.

(3) Press the menu button on the "User Config Save" to save the changed setting value.

(4) Press the menu button on the "Pre" will leave the SETUP menu.

The contents of each item are as shown as below.

Item	Description	Range
Factory default	Press the Menu button to change the factory default settings, all settings.	LOAD
User config save	The user-defined value is stored in memory, brings to the default settings when you reboot.	SAVE
Alarm 0 use	From SOUND menu it enables to turn ALARM Volume 0.	ON, OFF
Alarm 0 remind	When the alarm volume is 0 and the alarm occurs, sets if the alarm should beep in 3 minutes duration.	ON, OFF

Part 10 Basic Troubleshooting

Here are some basic troubleshooting hints. There may be times when the product does not work as it is supposed to and you may not know how to proceed. Try the following steps to work through the problems.

▣ General

Situation	Cause
The power will not come on.	-Make sure the product is plugged in. -Make sure the battery is fully charged.
The screen is dim and hard to read.	-Turn on LCD backlight.
I can't hear the alarm or the pulse sounds.	-Make sure the alarm is not turned off.
Coin battery image (⊕) is shown in the lower portion of the screen.	-Make sure the coin battery is not lacking capacity.

▣ Measuring the SpO2

Situation	Cause
The red sensor light does not come on.	-The probe is not connected properly. -The probe is broken.
I can't see the waves.	- The probe is not connected properly.

▣ Error Message

Message	Cause
High PR	SpO2 is above the alarm limit. Check patient immediately.
Low PR	SpO2 is above the alarm limit. Check patient immediately.
High SpO2	SpO2 is above the alarm limit. Check patient immediately.
Low SpO2	SpO2 is above the alarm limit. Check patient immediately.
Check Probe	Sensor not attached to patient. Reposition or replace sensor.
SpO2 fault	Analog board resulted in unstable contact with the IC Communication. Return to a qualified service technician.
Sensor off	Sensor not attached to patient. Reposition or replace sensor.
Finger off	Sensor not attached to patient finger. Reposition or replace sensor.
Low Battery	Connect to DC adaptor power. Continue charging on DC adaptor
BATT ERROR	Low Internal Coin Battery. Return to a qualified service technician.
CLOCK ERROR	Date and time setting lost. Set the date and time. Return to a qualified service technician.

Part 11 Product Specifications

10.1 Measurement Range

- Pulse Rate : 30 to 250bpm
- % SpO2 : 0 to 100%

10.2 Saturation Accuracy

- Saturation : 70% to 100%
- Adults, Pediatrics : ±2 digits
- Neonates : ±3 digits

10.3 Pulse Rate Accuracy

- Pulse Rate : 30 to 250bpm
- Adults, Pediatrics, Neonates: ±3 digits

10.4 Resolution

- Saturation (%SpO2) : 1%
- Pulse Rate : 1bpm

10.5 Perfusion index(PI)

- Range: 0.05 ~ 20%

10.6 Environment

- Operating Temperature : 5℃ to 35℃
- Storage Temperature : -20℃ to 70℃
- Operating Humidity : 5% to 95%
- Operation Altitude : 0~2.000m
- Air Pressure: 80kPa ~ 106kPa

10.7 Alarms

- Low SpO2 and pulse rate, system failure, low battery, Finger off.
- The alarm will sound and a message will appear on the LCD.

- Alarm limit

  1) Adult, Pediatric, Neonatal

  HR: 30 ~ 250bpm

  SpO2: 0 ~ 100%

10.8 Physical Characteristics

- Weight : Main : 273g

  Power module : 100g
- Dimensions : 72(H) x 154(W) x 26(D) mm

10.9 Display/Indicators

- Type : TFT LCD (3")
- Pixels : 240 x 400 dots
- Dot Pitch : 0.53mm
- Displayed Data : SpO2, pulse rate, SpO2 limits, pulse rate limits, waveform, alarm status, trend, battery status, current time
- Indicators : AC power or battery operated indicators

10.10 Trend Memory(option)

- SpO2 and pulse rate (20 days trend memory 10 sec)

10.11 Power

- Main Machine
  - Input : +9Vdc, 1.7A
  - Power Consumption : 9VA (max)
  - Fuse Rating : 4A, 250V
- Alkaline Battery
  - Type : 4AA Alkaline Batteries(4 * 1.5V)
  - Capacity : Up to 10 hours
- Rechargeable Battery
  - Type : Lithium Ion Rechargeable Battery(2360mA)
  - Capacity : Up to 8 hours
  - Charging Time : 6 hours
- Adaptor(option)
  - Input : 100-240Vac, 50-60Hz, 1.0A
  - Output : +9Vdc, 3.34A

10.12 SpO2 Sensor light source

- Wavelength
  - Infrared : 905 nm (nominal)
  - Red : 660 nm (nominal)
- Power dissipation
  - Infrared : 100mW (max)
  - Red : 100mW (max)

10.13 Manufacturer's Declaration

- Electromagnetic Compatibility (EMC)

⚠ Caution

This monitoring system is intended for use by healthcare professionals only. This monitoring system may cause radio interference or may disrupt the operation of nearby equipment, regardless of whether it is CISPR compliant or not. It may be necessary to take mitigation measures, such as re-orienting or relocating the monitoring system or shielding the location.

⚠ Caution

The use of accessories, Sensors, and cables other than those specified may result in inaccurate reading s of the monitoring system and increased emission and/ or decreased electromagnetic immunity of the monitoring system.

- Electromagnetic Emissions

Emissions Test	Compliance	Electromagnetic Environment Guidance
RF emissions CISPR 11	Group 1 Class A	Not intended for use in a residential environment. If used in a domestic environment, may not offer adequate protection to radio-frequency communication services. The user may be required to take mitigation measures, such as relocating or re-orienting the equipment..
Harmonic emissions	Class A	.

IEC 61000-3-2		
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Complies	

■ Electromagnetic Immunity

Immunity test	EN/IEC 60601-1-2 Test level	Compliance level	Electromagnetic Environment-Guidance
Electrostatic discharge (ESD) IEC 61000-4-2	±6 kV contact ±8 kV air	±6 kV contact ±8 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30 %.
Electrical fast transient/burst IEC 61000-4-4	±2 kV for power supply lines ±1 kV for input/output lines	±2 kV for power supply lines ±1 kV for input/output lines	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	±1 kV line(s) to line(s) ±2 kV line(s) to earth	±1 kV line(s) to line(s) ±2 kV line(s) to earth	Mains power quality should be that of a typical commercial or hospital environment.
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	<5 % U_T (>95 % dip in U_T) for 0,5 cycle 40 % U_T (60 % dip in U_T) for 5 cycles 70 % U_T (30 % dip in U_T) for 25 cycles <5 % U_T (>95 % dip in U_T) for 5 s	<5 % U_T (>95 % dip in U_T) for 0,5 cycle 40 % U_T (60 % dip in U_T) for 5 cycles 70 % U_T (30 % dip in U_T) for 25 cycles <5 % U_T (>95 % dip in U_T) for 5 s	Mains power quality should be that of a typical commercial or hospital environment. If the user of the [ME EQUIPMENT or ME SYSTEM] requires continued operation during power mains interruptions, it is recommended that the [ME EQUIPMENT or ME SYSTEM] be powered from an uninterruptible power supply or a battery.
Power frequency	3 .0A/m	3 .0A/m	Power frequency magnetic fields

(50/60 Hz) magnetic field IEC 61000-4-8			should be at levels characteristic of a typical location in a typical commercial or hospital environment.

■ . Electromagnetic Immunity Compliance

Immunity test	IEC 60601-1-2 Test level	Compliance level	Electromagnetic Environment-Guidance
	Frequency of Transmitter		Equation for Separation Distance (d)
Conducted RF IEC 61000-4-6	3 Vrms 150 kHz to 80 MHz	3 Vrms 150 kHz to 80 MHz	$d = 1.2\sqrt{P}$ 150 kHz to 80 MHz
Radiated RF IEC 61000-4-3	3 V/m 80 MHz to 800 MHz	3 V/m 80 MHz to 800 MHz	$d = 1.2\sqrt{P}$ 80 MHz to 800 MHz
	3 V/m 800 MHz to 2,5 GHz	3 V/m 800 MHz to 2,5 GHz	$d = 2.3\sqrt{P}$ 800 MHz to 2,5 GHz

■ Recommended Separation Distance

Rated maximum output power (P) of transmitter in watts	Separation Distance in Meters		
	$d = 1.2\sqrt{P}$ 150 kHz to 80 MHz	$d = 1.2\sqrt{P}$ 80 MHz to 800 MHz	$d = 2.3\sqrt{P}$ 800 MHz to 2,5 GHz
0.01	0.12	0.12	0.23
0.10	0.38	0.38	0.73
1.00	1.20	1.20	2.30
10.00	3.80	3.80	7.30
100.00	12.00	12.00	23.00

For transmitters rated at a maximum output power not listed above, estimate the separation distance (*d*) using the equation in the corresponding column, where *P* is the maximum output power rating of the transmitter in watts (*W*) according to the transmitter manufacturer.

Note

These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects, and people.

Note

Portable and mobile RF communications equipment can affect medical electrical equipment. Such RF equipment should be used no closer to any part of the monitoring system, including cables, than the recommended separation distance calculated from the equation appropriate for the frequency of the transmitter.

- Clinical Studies

Overview

This contains data from clinical studies conducted for Neonate Sensor Adult Sensor, Disposable Sensor and Pediatric Sensor used with the ACCURO oximeter

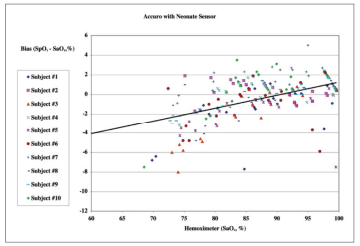
Methods

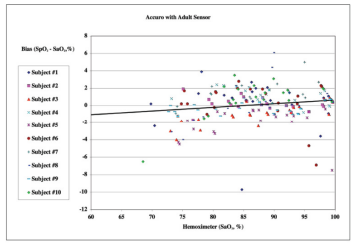
The current study for Charmcare, Inc. performed 3/18/10 included 10subjects- 7 women and 3 men. The ACCURO oximeter was studied with three different sensors: adult, disposable and neonate. A radial arterial cannula was placed in either the left or right wrist of each subject. Blood gas analysis to determine oxyhemoglobin saturation was performed on an OSM 3® multi- wavelength oximeter (Hemoximeter, Radiometer, Copenhagen, serial 89R0243 N010). This instrument underwent a full factory technician calibration and reference standards check on June 8, 2009. The instrument was also verified to read within specifications with the use of Radiometer Qualicheck standards within 48 hours of the study dates. No subject was anemic (Hemoglobin ≤ 10 gm•dl-1) and only healthy non-smoking individuals of age 21-49 were included in the study.

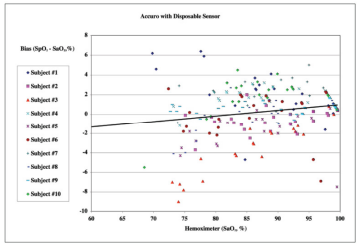
The study was reviewed and approved by the University of California at San Francisco Committee on Human Research. The approval date was 3/21/09, expiration April 21, 2010. The Approval number is H6301-01706-24. The approval letter is on file at UCSF. Each subject had control data taken at the beginning of each experiment, with two control blood samples drawn while breathing room air. Hypoxia was induced to different levels of oxyhemoglobin saturation (between 70-100%) by having subjects breathe mixtures of nitrogen, room air, and carbon dioxide. Each plateau level of oxyhemoglobin saturation was maintained for at least 30 seconds and until Pulse Oximeters readings had stabilized. Two arterial blood samples were then obtained, approximately 30 seconds apart. Each stable plateau therefore was maintained for at least 60 seconds with SpO2 fluctuating by less than 3%. The plateaus were nominally at 100%, room air saturation, 93%, 90%, 87%, 85%, 82%, 80%, 77%, 75%

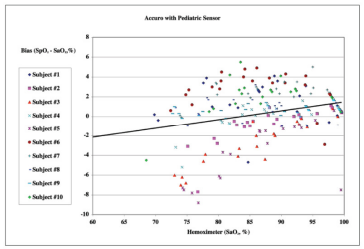
and 70%. A total of 22-24 samples were obtained that the plateaus across this span. Data were recorded by computer. At least 200 data points were collected for each type of oximeter and probe combination studied.

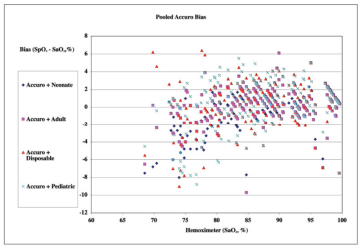
Study Results











Conclusion

The pooled results indicate that for a saturation range of 60-80% for SpO2, the acceptance criterion was met for the monitoring system when tested with Neonate Sensor Adult Sensor, Disposable Sensor and Pediatric Sensor. The pooled results indicate that for a saturation range of 70-100% for SpO2, the acceptance criterion was met.

Attachment 1 Factory Set

The default value of the product is as shown below.

Factory Default

* Timing to save

- **When altering parameter values** : When a user changes each parameter value at users discretion it will be saved in memory that after a quick restarting the last saved values will be applied.

- **USER CONFIG**: Even if the user changes parameter values at users discretion validity of this change will be preserved for only 30 seconds after powering off the device before restarting it. Authorized maintenance personnel can only change USER CONFIG setting(MAINT>USER CONFIG SAVE) that will be always retrieved if restarted more than 30 seconds after any other normal users manipulations.

Below shows the parameters that users can changes.

Menu	Item	Factory set	* Timing to save
CONFIG	BACKLIGHT	4	When changing parameter value.
	ALARM VOLUME	7	When changing parameter value.
	PULSE VOLUME	7	When changing parameter value.
	WAVE FILL	OFF	When changing parameter value.
	AVERAGE TIME	8	USER CONFIG
	ALARM DURATION	120	When changing parameter value.
	DEMO	OFF	USER CONFIG
ALARM	SpO2 High limit	OFF	USER CONFIG
	SpO2 Low Limit	90(neonate:85)	USER CONFIG

	PR High Limit	Adult : 120 Pediatric :160 Neonate : 200	USER CONFIG
	PR Low Limit	Adult : 50 Pediatric : 75 Neonate : 100	USER CONFIG
PATIENT	PATIENT	ADULT	When changing parameter value.

Maintenance Set

Below shows the parameters that authorized personnel with password can change.

Category	Name	Factory Default	* Timing to save
MAINTENANCE	FACTORY DEFAULT	-	When changing parameter value. (Authorized maintenance personnel)
	USER CONFIG SAVE	-	When changing parameter value. (Authorized maintenance personnel)
	ALARM 0 USE	OFF	When changing parameter value. (Authorized maintenance personnel)
	ALARM REMIND	OFF	When changing parameter value. (Authorized maintenance personnel)

Factory Set By Patient Type

* When transferred to Patient mode, the alarm limit value and SPO2 Average value automatically changes to the USER CONFIG set.

Item	Factory Default	
SPO2 HIGH (%)	ADULT	: OFF
	PEDIATRIC	: OFF
	NEONATE	: OFF
SPO2 LOW (%)	ADULT	: 90
	PEDIATRIC:	: 90
	NEONATE	: 85
PR HIGH (bpm)	ADULT	: 120
	PEDIATRIC:	: 160
	NEONATE	: 200
PR LOW (bpm)	ADULT	: 50
	PEDIATRIC	: 75
	NEONATE	:100
SpO2 Average (s)	ADULT	: 8
	PEDIATRIC	: 8
	NONATE	:12

Attachment 2 Alarm Message

The level and cause of a single alarm is presented. '*' mark can be changed according to the value, and when the value does not need to be changed it is set to a fixed value.

- **Patient Alarm Message**

Alarm Message	Level	Cause
HIGH SPO2	Medium	Occurs when the measured value is greater than the high limit or greater than the low limit. Check the status of the patient. Also, check if the limit set value is valid.
LOW SPO2	Medium	
HIGH PR	Medium	
LOW PR	Medium	

- **Technical Alarm Message**

Alarm Message	Level	Cause
SPO2 FAULT	High	Not connected with the oxygen saturation measuring part.
SENSOR OFF	Low	The sensor is not connected to the product. Occurs when the sensor is not correctly connected or a problem is detected in the sensor.
FINGER OFF	Low	The sensor is not attached to the patient.
CHECK PROBE	Low	Checking the sensor status is required. The light of the sensor may be exposed to an external light.
LOW BATTERY	Medium	Battery is low. Connect DC power and charging is necessary.
BATT. ERROR	Low	Coin battery is low.
CLOCK ERROR	Low	An error for time value occurs. reset is necessary.

CE 0120

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