ECG, Respiration and SpO_2 Transmitter

LX-7230N
LX-7230N(G)

Operation Manual

* Before using the product, please read this manual thoroughly.
* Store this manual where it can be always referred to.

FUKUDA

This manual is for the LX-7230N Version 02 and 03.

This manual is for the LX-7230N (G) Version 01 and 02.

⚠ CAUTION	Federal Law restricts this device to sale by or on the order of a physician.

CAUTION

- Only physician or persons instructed by physicians are allowed to use the equipment.
- The information contained in this document is subject to change without notice due to improvement in the equipment.

Thank you for purchasing this product.
Before using this product, read this operation manual thoroughly for correct handling and operation.

In this manual, the operation procedure of LX-7230N/LX-7230N(G) is explained using the illustration and screen of the LX-7230N as examples.

Safety Precautions

- Read the “Safety Precautions” thoroughly before use to ensure correct and safe use of the product.
- Make sure to follow the precautions indicated below, as these are important messages related to safety.

⚠DANGER	Failure to follow this message may cause immediate threat of death or serious injury.

⚠WARNING	Failure to follow this message may result in death or serious injury.

⚠CAUTION	Failure to follow this message may cause injury or failure to the equipment.

NOTE	A note is not related to product safety, but provides information about the correct use and operating procedures to prevent incorrect operation and malfunction of the equipment.

Precaution from Fukuda Denshi

Fukuda Denshi is liable for the safety, reliability, and performance of its equipment only if;

- Maintenance, modifications, and repairs are carried out by authorized personnel.
- Components are used in accordance with Fukuda Denshi operating instructions.

If the equipment is used incorrectly and become unusable, Fukuda Denshi is not liable for the malfunction. Use the equipment only for the purpose specified in this manual.

Intended Use of this Equipment

This equipment is designed for the following <Intended Use>.

<Intended Use>
This equipment is intended for patient monitoring in surgery room, ICU, ward, emergency room in the medical facility by measuring ECG, Respiration and SpO_2 and transmitting the measured data by wireless network to the central monitor continuously.

This equipment is intended to be used by healthcare professionals. Users should have a thorough knowledge of the function and operation before using this equipment. The maintenance of this equipment should be performed by skilled personnel who received a training of possible hazards and measures to avoid those hazards. Also, your local regulation must be followed. If this equipment is used for the purpose other than intended, or if the user does not follow the safety instructions, the following hazard may result.

- Hazard to the Life and Health of the Patient or the User
- A Problem Related to Medical Practice
- Damage to the Equipment

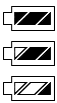
Graphic Symbols

The following symbols are used for this equipment.

Symbols indicated on the main unit

Symbol	*Description*
	Warning (indicated in yellow)
	Follow operating instructions (Warning); (indicated in blue) Indicates that the failure to follow operating instructions could place the patient or operator at risk.
	Type CF Applied Part with Defibrillation-Proof Indicates that the degree of protection against electric shock is Type CF Applied Part with defibrillation-proof.
	No Alarm Function Indicates no alarm function.
	Battery compartment lid lock Indicates locking and unlocking of the battery compartment lid.
	Battery Type and Direction Indicates the battery type and direction.
	Name and Address of Manufacturer Date of Manufacture Indicates the name and address of manufacturer, and date of manufacture.
	Non-ionizing electromagnetic radiation Indicates the radio transmitting device.

Symbols indicated on the LCD screen

Symbol	*Description*
	Heart Rate Synchronization Mark This mark flashes synchronizing to the heartbeat.
	Battery Mark Indicates the remaining battery level.

Precautions for Safe Operation of Medical Electrical Equipment

Cautions described here are regarding the general instructions for safety use to the patient and users. As for cautions about the LX-7230N/LX-7230N(G), please refer to the following pages.

⚠CAUTION

1. **Users should have a thorough knowledge of the operation before using this equipment.**
2. **Pay attention to the following when installing or storing the equipment.**
   - Do not install or store in an area where the equipment will be subject to splashing water.
   - Do not install or store in an area where the environmental conditions, such as atmospheric pressure, temperature, humidity, ventilation, sunlight, dust, sodium, sulfur, will adversely affect the system.
   - Place the equipment on a stable surface where there is no inclination, vibration, or shock (including during transportation).
   - Do not install or store in an area where chemicals are stored or gases are evolved.
3. **Before operating the equipment, verify the following items.**
   - Check the cable connection and polarity to ensure proper operation of the equipment.
   - Ensure that all cables are firmly and safely connected. Especially, recheck the attachment and connection condition of electrode and the probe (sensor).
   - Pay special attention when the equipment is used in conjunction with other equipment as it may cause erroneous judgment and danger.
   - Check the remaining battery level.
   - When replacing the battery, make sure that the battery polarity is correct. Do not charge the battery.
4. **During operation of the equipment, verify the following items.**
   - Do not operate the equipment beyond the time period required for diagnosis and medical care.
   - Do not pick up and/or swing the equipment pulling/grabbing the probe (sensor) or cable part. It may damage the equipment and lead to measurement error.
   - Always observe the equipment and patient to ensure safe operation of the equipment.
   - If any abnormality is found on the equipment or patient, take appropriate measures such as ceasing operation of the equipment and/or detaching the probe (sensor) and/or electrode, in the safest way for the patient.
   - Do not allow the patient to come in contact with other equipments.

⚠CAUTION

5. After using the equipment, verify the following items.

- Make sure to turn off the power of the equipment.
- When unplugging the cables, do not apply excessive force on the cable and pull from its connector.
- Clean the accessories and cables, and keep them together in one place.
- Keep the equipment clean to ensure proper operation for the next use.
- Make sure to remove the batteries if the equipment is not used for a long time. The leakage from the batteries may damage the equipment or an explosion from the batteries may occur.

6. If the equipment is damaged and in need of repair, ensure patient safety by immediately turning the equipment off and remove the electrodes and/or probe from the patient. User should not attempt service. Label the unit "OUT OF ORDER" and contact Fukuda Denshi representative.

7. Do not remodel the equipment.

8. Maintenance Check

- Make sure to periodically check the equipment, and accessories.
- Before reusing the equipment that has been left unused for a while, make sure that the equipment works normally and safely.

9. When using electrosurgical knives or defibrillator with this equipment, take care of the following.

- To prevent burn injury to the patient, verify proper attachment of patient ground plate, ECG electrode type for the electrosurgical knives, and the quantity of gel, output energy for the defibrillator. Also, verify that a proper ground is selected.
- Some types of equipment other than the above may cause accidental hazards to the patient and operator due to the conditions of the equipment. Read the operation manual attached to each equipment and understand the precautionary instructions prior to use.

Non-Explosion Proof

⚠DANGER

- Never operate the equipment in the presence of flammable anesthetics, high concentration of oxygen. It may cause an explosion or fire.
- Never operate the equipment inside a hyperbaric chamber. It may cause an explosion or fire.
- Never operate the equipment where flammable gas or fluid such as anesthetic, oxygen, and hydrogen are used. It may cause an explosion or fire.

Defibrillation Safety

⚠WARNING

- Use only the lead cable specified by Fukuda Denshi when defibrillating. If used by unspecified lead cable, the equipment may be damaged, resulting in a safety hazard.
- When using the defibrillator, keep away from the electrodes or medicament applied to the patient chest. If this is not possible, remove the electrodes or medicament before using it.
  If the defibrillator paddles are directly in contact with the electrodes or medicament, an electrical shock may result from the discharged energy.
- When using the defibrillator, do not touch the patient and the metal part of the equipment or cables. Electric shock may result from the discharged energy.
- This equipment will return to standard operating mode within 10 seconds after defibrillating. The stored data will not be affected. The measurement accuracy will temporarily decrease during defibirillation, but it will not compromise the safety of patient and the equipment.

Precautions about Magnetic Resonance Imaging (MRI)

⚠WARNING

- Do not operate this equipment in magnetic resonance imaging (MRI) environments.
- When conducting MRI test, remove the electrodes and sensors connected to the patient (test subject).
  The local heating caused by the induced electromotive force may cause burn injury to the patient (subject). For details, refer to the operation manual for the MRI testing device.

Electrosurgery Safety

⚠WARNING

The monitoring system contains protection against interference generated by electrosurgical instruments. However, depending on the operating conditions, surgery site with respect to the location of ECG electrodes, ground plate attachment condition, or the type of instrument used, it may cause burn injury at the electrode site or noise on the ECG. The noise is generated at the tip of the electrosurgical knife and is difficult to completely eliminate because of the frequency components of the ECG. To reduce electrosurgical interference, take the following precautions:

Location:

Locate the electrosurgical unit as far as possible from this equipment and the patient cable. This will help reduce interference on the ECG through the monitor or cables.

Electrode Placement:

The amount of noise interference is considerably different depending on the electrode position and surgery site. Place the ECG electrodes as far away as possible from the surgery site and the ground plate. Do not place electrodes in the path between the surgery site and the ground plate. If the electrodes are placed in this path, the amount of interference will be quite large. Position (+) and (–) electrodes as close as possible to each other.

Ground Plate:

When using electrosurgical instruments, make sure the contact between the patient and the ground plate is secure. If the connection is incomplete, the patient may suffer from burn at the electrode site.

Precautions about the Pacemaker

⚠WARNING

- Minute ventilation rate-adaptive implantable pacemakers can occasionally interact with certain cardiac monitoring and diagnostic equipment, causing the pacemakers to pace at their maximum programmed rate. The cardiac monitoring and diagnostic equipment may possibly send wrong information.
  If such event occurs, disconnect the cardiac monitoring and diagnostic equipment, or follow the procedures described in the operation manual of the pacemaker.
  (For more details, contact FUKUDA DENSHI personnel, your institution's professionals, or your pacemaker distributors.)
- Rate meters may continue to count the pacemaker rate during occurrences of cardiac arrest or some arrhythmias. Do not rely entirely upon rate meter alarms. Keep pacemaker patients under close surveillance.

  ➲ Reference
  "Minute Ventilation Rate-Adaptive Pacemakers"
  FDA alerts health professionals that minute ventilation rate-adaptive implantable pacemakers can occasionally interact with certain cardiac monitoring and diagnostic equipment, causing pacemakers to pace at their maximum programmed rate.
  [October 14, 1998 – FDA]

Precautions about the LX-7230N/LX-7230N(G)

⚠WARNING

- Do not connect cables not authorized by Fukuda Denshi to any I/O connector. If done so by mistake, the LX-7230N/LX-7230N(G) cannot deliver its maximum performance and may be damaged, resulting in a safety hazard.
- Do not use this equipment with multiple patients simultaneously.
- This equipment itself has no alarm function. Do not use it if an alarm function is necessary. The alarm function with the receiving monitor, refer to the operation manual of the receiving monitor.

⚠CAUTION

Do not pick up and/or swing the LX-7230N/LX-7230N(G) pulling/grabbing the probe (sensor) or cord part. The cable could break or get disconnected from the LX-7230N/LX-7230N(G). And it may hit people or damage other equipment around.

Precautions about Waterproof

⚠CAUTION

- Replace the “Battery Compartment Lid” of the LX-7230N/LX-7230N(G) regularly to keep the performance of waterproof. If not regularly replaced, the quality of the lid will deteriorate and cannot keep the waterproof performance. For details about the regular replacement, contact your local Fukuda Denshi service representative.
- The lid may be damaged from high impact. If the LX-7230N/LX-7230N(G) is dropped or is subjected to a high impact, make sure that the lid is not damaged.
- However, the SpO_2 probes (sensors) are not waterproof. Do not take a bath with them, and ensure to be away from liquid.
- Do not use the LX-7230N/LX-7230N(G) wet. Always wipe the LX-7230N/LX-7230N(G) with a soft cloth and dry it thoroughly before use.

Precautions about ECG

⚠CAUTION

- When removing electrodes from the patient, remove them carefully and slowly. Do not apply excessive force to remove them. Otherwise, it may damage the skin.
- If any electrodes get detached from the patient after being connected to the lead cable and the patient monitor, pay attention that the metal part of the electrode does not get in touch with any metal parts of the bed or any conductive parts. Also, the operator should not touch any conductive parts with bare hands. Otherwise, it may cause electric shock to the patient and/or operator due to excessive leakage current.
- The indication for continuous use of the electrode is about one day.
- Replace the electrode if the skin contact gets loosen due to perspiration, etc.
- When an electrode is attached to the same location for a long period, some patients may develop skin irritation. Check the patient's skin condition periodically and change the electrode site as required.
- For stable ECG monitoring, verify proper electrode placement, lead, and waveform size. If not properly selected, it may cause erroneous detection.
- There are some cases when the pacemaker pulse cannot be detected depending on the pacemaker type, pulse voltage, pulse width, electrode lead type (unipolar, bipolar), electrode placement, or lead method which causes the pacemaker pulse amplitude to decrease and disables pacemaker pulse detection.
- If signals similar to a pacemaker pulse are present, such as electric blanket noise or excessive AC frequency noise, these may be erroneously detected and displayed as a pacemaker pulse. In this case, check the condition of the electrodes and ECG lead cable to resolve the cause or turn off the pacemaker detection setting on the receiving monitor.

⚠CAUTION

- If a pacemaker pulse is continuously detected due to AC frequency interference, QRS detection will be suspended and the heart rate will be reduced.
- If an arrhythmia accompanied by an irregular rhythm occurs, there is a possibility of temporarily displaying an incorrect heart rate. When using this device in conjunction with an electrocautery scalpel, noise from an electric scalpel may be misrecognized as heartbeat or arrhythmia.

Precautions about SpO_2

⚠WARNING

- During SpO_2 monitoring, always use the probe (sensor) specified by Fukuda Denshi. Also, check the probe before usage to make sure that it is the specified probe. If unspecified probe is used, not only that the equipment cannot deliver its maximum performance, the equipment may be damaged and safety cannot be ensured.
- If irritation such as skin reddening appears with the sensor use, change the attachment site or stop using the sensor.
- Do not use a tape to attach the sensor.
- When the SpO_2 probe (sensor) is in a connector-off condition, the SpO_2 alarm will not be generated on the receiving monitor. Make sure that the SpO_2 probe (sensor) is securely connected. If the SpO_2 waveform/numeric data is not displayed, check the patient's condition and pay attention not to miss the connector-off condition.
- When not performing measurement, unplug the sensor from the SpO_2 connector. Otherwise, the measurement data may be erroneously displayed by the ambient light.
- Check the sensor attachment site constantly every 4 hours when probes or reusable sensor are used, and at least every 8 hours when single patient use sensors are used. Be especially careful of a patient with bad perfusion. If the sensor attachment position is not changed constantly, skin irritation or skin necrosis due to compression may be developed. For the patient with bad perfusion, check the sensor attachment position at least every 2 hours.
- As skin for neonate, premature infant is immature, change the sensor attachment site more frequently depending on the condition.
- When measuring the SpO_2 of a patient with high fever or peripheral circulatory insufficiency, check the probe (sensor) attachment periodically and change the attachment site. The temperature of the attachment site will rise 2°C to 3°C due to the sensor heat which may result in compression necrosis and burn injury.
- Even attachment for a short duration may inhibit the blood flow and generate compression necrosis or burn injury. Also, blood flow inhibition may prevent correct measurements.

⚠WARNING

- Direct sunlight to the sensor area can cause measurement error. Place a black or dark cloth over the sensor.
- The pulse wave is normalized for SpO_2 measurement, and does not indicate perfused blood volume. Check proper probe attachment by observing the pulse wave.
- Precautions for Reusable Sensors
  The light-emitting part of the sensor should be over the root of the fingernail or as instructed per the related sensor instruction manual.
  Do not insert the finger too far into the sensor as it may hurt the patient.
  For details, refer to the SpO_2 sensor instruction manual.
- Precautions for Single-Patient-Use Type Sensors
  The sensor can be reused on the same patient as long as the adhesive tape attaches without slippage. But do not reuse on other patients to avoid cross contamination. It is intended for single patient use only. For details, refer to the SpO_2 sensor instruction manual.
- Measuring on a limb with NIBP cuff, arterial catheter, or intracatheter may result in incorrect measurement.
- Venous congestion may cause under reading of actual oxygen saturation. Therefore, assure proper venous outflow from monitored site. Sensor should not be below heart level (e.g. sensor on hand of a patient in a bed with arm dangling to the floor).

⚠CAUTION

For the following case, accurate measurement may not be possible.

- Patient with excessive abnormal hemoglobin (COHb, MetHb)
- Patient with the pigment injected to the blood
- Patient receiving CPR treatment
- Placement of SpO_2 probe (sensor) on limb with a blood-pressure cuff, arterial catheter, or intravascular line
- When measuring at placement position with venous pulse
- Patient with body motion
- Patient with small pulse
- Excessive body motion (patient's motion)
- Excessive light (direct sunlight, fluorescent, light therapy equipment, surgical light, infrared heat ramp, etc.)
- External colorant such as nail polish
- Abnormally low or high hemoglobin concentration
- Electrosurgery
- Influence of electromagnetic waves from other electronics devices
- High-intensity radio waves from mobile phones

Precautions about Output Signal

⚠WARNING

Do not use the output signal of the monitor that receives radio wave signal from the LX-7230N/LX-7230N(G) as the trigger signal for IABP, MRI echocardiographic, or defibrillator for the following reasons.

- It may lead to a delay of operating timing due to the delay time of waveform transmission.
- A trigger signal unrelated to the heart rate may be generated due to the interfusion of spike noise at weak electric field.

Precautions about Accessories and Optional Accessories

⚠WARNING

Use only the accessories such as disposable electrodes, electrode codes and SpO_2 probe (sensor) specified by Fukuda Denshi. Otherwise, this equipment cannot deliver its maximum performance and may be damaged, resulting in a safety hazard.

⚠CAUTION

- Do not reuse disposable products.
- Store the disposable products properly as mentioned in their user manuals.

Precautions about Battery

⚠WARNING

- Use new "AA" size (“LR06” size) alkaline cell.
- Install the battery with the correct polarity.
- Do not charge the battery. Any attempt to charge the battery may cause it to leak or break.
- Do not short the (+) and (-) terminals. It may result in exothermic heat and fire.
- Do not throw the battery into fire. It may explode.

Precautions about Disposing of Equipment, Accessories, or Components

⚠CAUTION

- When disposing of the equipment, accessories, or components, use an industrial waste distributor. Do not dispose of as ordinary waste.
- Used disposal items (ECG electrodes, etc.) shall be discarded as medical waste.

Precautions about Disposing of Battery

⚠CAUTION

Obey the local municipal rule to dispose the used dry cell battery.

Precautions for Use of Medical Telemeter

⚠ WARNING

- The LX-7230N/LX-7230N(G) transmitter must not be co-located or operated in conjunction with any other antenna or transmitter.
- For the receiving monitor of the LX-7230N transmitter, make sure to use the Fukuda Denshi products with the receiving range of 608 MHz-614 MHz.
- For the receiving monitor of the LX-7230N(G) transmitter, make sure to use the Fukuda Denshi products with the receiving range of 1395 MHz-1400 MHz and 1427 MHz-1432 MHz.
- This equipment complies with FCC/IC radiation exposure limits set forth for an uncontrolled environment and meets the FCC radio frequency (RF) Exposure Guidelines and RSS-102 of the IC radio frequency (RF) Exposure rules. This equipment has very low levels of RF energy that are deemed to comply without testing of specific absorption rate (SAR).
- Operation of LX-7230N/LX-7230N(G) requires the prior coordination with a frequency coordinator designated by the FCC for the Wireless Medical Telemetry Service.
- This radio frequency device is susceptible to interference from outside sources. Interference may prevent the monitoring of patients connected to this equipment. If a problem exists, contact your local service representative.
- The LX-7230N/LX-7230N(G) transmits vital signs to the receiving monitor using radio wave signal. Under unstable radio wave signals, the receiving monitor will not generate any alarms. This situation may miss sudden change in the patient's condition and may cause a serious accident. Under unstable radio wave signals, check the patient status consistently under this situation. To get stable radio wave signals, make sure to have a proper telemetry installation.

⚠CAUTION

- For installation, make sure of the following.
  - The medical institution (hereinafter referred to as the “Institution”) must decide the telemetry installation plan for the medical department in order to prevent interference and interference between transmitters (telemetry based on destination country’s radio law). When telemetry has already been installed and been used, radio format, frequency, and antenna power are required to be examined to prevent interference.
  - When laying receiver antenna for each transmitter, the Institution has to examine the installation so that electronic interference does not occur.
  - Based on the above examination result, the Institution should install each receiver antenna as required.
- For management, make sure to follow the precautions below.
  - The Institution should appoint a person (hereinafter referred to as the “Coordinator”) to manage the wireless channels for the whole Institution.
  - The Coordinator must be selected from people who understand the characteristics and functionality of telemetry systems, and are skilled in operating telemetry.
  - When installing telemetry, the Coordinator has to understand the precautions for use of telemetry in advance.
  - The Coordinator is responsible for maintenance of wireless channels and storage and maintenance of telemeter in the overall medical facilities to give proper instructions to the telemetry users.
  - The Coordinator should create a management log (hereinafter referred to as the “log”), which contains a list of the management status of the wireless channels for the whole Institution. When changing a wireless channel, register it in the log and give proper instructions to the Coordinator or to the user.
  - The Coordinator assumes responsibility for managing the wireless channels, storing, and managing telemetry.
  - The Coordinator assigns the transmitter to the user, and provides enough education for use.
  - The telemetry user verifies operation of the transmitter/receiver before use.
  - When interference or breakdown occurs in telemetry communication, the user is required to inform the Coordinators of the problems. The Coordinators are to deal with the problem properly and/or contact their nearest Fukuda Denshi representative for service.

Electromagnetic Compatibility

The performance of this equipment under electromagnetic environment complies with IEC 60601-1-2 (2007).

Precautions for Safe Operation under Electromagnetic Influence

⚠CAUTION

If any sorts of electromagnetic wave, magnetic field, or static electricity exist around the equipment, noise interference or malfunction of the equipment may occur. If any unintended malfunction or noise occurs during monitoring, check the magnetic influence and take appropriate countermeasures.
The following are examples of the common cause and countermeasures.

●Mobile Phone

The radio wave may cause malfunction to the equipment.
Mobile phones and radio sets should be turned off in the room (building) where medical device is located.

●Static Electricity

In a dry environment (room), static electricity is likely to occur. Take the following countermeasures.

- Both operator and patient should remove any static electricity before entering the room
- Humidify the room

EMC Guidance

This equipment complies with IEC 60601-1-2 (2007). However, if portable transmitter or wireless LAN equipment is used extremely nearby, the electromagnetic influence may largely exceed the compliance level and may cause unexpected phenomenon such as noise interference on the waveform, etc.
Therefore, this equipment should be used in a location specified by each medical institution. If any unexpected noise interference on the waveform or failure to the peripheral device occurs, stop using the equipment and follow the instruction of the technician.

The following is the information relating to EMC (Electromagnetic Compatibility).
(When using this equipment, verify that it is used within the environment specified below.)

●Compliance to the Electromagnetic Emissions

The LX-7230N/LX-7230N(G) is intended for use in the electromagnetic environment specified below. The customer or the user of the LX-7230N/LX-7230N(G) should assure that it is used in such an environment.

Emission Test	***Compliance***	***Electromagnetic Environment – Guidance***
RF Emission CISPR 11	Group 1	The LX-7230N/LX-7230N(G) 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 A	The LX-7230N/LX-7230N(G) is suitable for use in all establishments other than domestic establishments.
Harmonic Emission IEC 61000-3-2	N/A	
Voltage Limit / Flicker Emission IEC 61000-3-3	N/A	

●Compliance to the Electromagnetic Immunity (1)

The LX-7230N/LX-7230N(G) is intended for use in the electromagnetic environment specified below. The customer or the user of the LX-7230N/LX-7230N(G) should assure that it is used in such an environment.

Immunity Test	***IEC 60601-1-2 Test Level***	***Compliance Level***	***Electromagnetic Environment Guidance***
Electrostatic Discharge (ESD) IEC 61000-4-2	±2,4,6kV: Contact ±2,4,8kV: Air	±2,4,6kV: Contact ±2,4,8kV: 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	±2kV: Power supply lines ±1kV: Input/output lines	N/A	
Surge IEC 61000-4-5	±1kV: differential mode ±2kV: common mode	N/A	
Voltage dips, short interruptions and voltage variations on power supply input lines. IEC61000-4-11	<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 5sec.	N/A	
Power Frequency (50/60Hz) Magnetic Field IEC 61000-4-8	3A/m	3A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.
Note: U_T is the AC mains voltage prior to application of the test level.			

●Compliance to the Electromagnetic Immunity (2)

The LX-7230N/LX-7230N(G) is intended for use in the electromagnetic environment specified below. The customer or the user of the LX-7230N/LX-7230N(G) should assure that it is used in such an environment.

Immunity Test	***IEC 60601-1-2 Test Level***	***Compliance Level***	***Electromagnetic Environment Guidance***
			Portable and mobile RF communications equipment should be used no closer to any part of the LX-7230N/LX-7230N(G), including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. **Recommended Separation Distance**
Conducted RF IEC 61000-4-6	3Vrms 150kHz to 80MHz	3Vrms	$d = 1.2\sqrt{P}$
Radiated RF IEC 61000-4-3	3V/m 80MHz to 2.5GHz	3V/m	$d = 1.2\sqrt{P}$ 80MHz to 800MHz $d = 2.3\sqrt{P}$ 800MHz to 2.5GHz Where *P* is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and *d* is the recommended separation distance in meters (m). Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey [a], should be less than the compliance level in each frequency range [b]. Interference may occur in the vicinity of equipment marked with the following symbol:

Note 1: At 80MHz and 800MHz, the higher frequency range applies.
Note 2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

[a] Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the LX-7230N/LX-7230N(G) is used exceeds the applicable RF compliance level above, the LX-7230N/LX-7230N(G) should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the LX-7230N/LX-7230N(G).

[b] Over the frequency range 150kHz to 80MHz, field strength should be less than 3V/m.

●Recommended Separation Distances between Portable and Mobile RF Communications Equipment and the LX-7230N/LX-7230N(G)

The LX-7230N/LX-7230N(G) is intended for use in an environment in which radiated RF disturbances are controlled. The customer or the user of the LX-7230N/LX-7230N(G) can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the LX-7230N/LX-7230N(G) as recommended below, according to the maximum output power of the communications equipment.

Rated Maximum Output Power of Transmitter (W)	***Separation Distance according to Frequency of Transmitter (m)***		
	150kHz to 80MHz $d = 1.2\sqrt{P}$	80MHz to 800MHz $d = 1.2\sqrt{P}$	800MHz to 2.5GHz $d = 2.3\sqrt{P}$
0.01	0.12	0.12	0.23
0.1	0.38	0.38	0.73
1	1.2	1.2	2.3
10	3.8	3.8	7.3
100	12	12	23

For transmitters rated at a maximum output power not listed above, the recommended separation distance *d* in meters (m) can be determined using the equation applicable to the frequency of the transmitter, where *P* is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer.

Note 1 : At 80MHz and 800MHz, the separation distance for the higher frequency range applies.

Note 2 : These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

Contact

If you need more information, please contact the following.

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CONTENTS

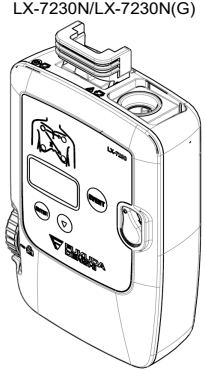
1. General Description

The LX-7230N/LX-7230N(G) is a radio telemetry transmitter designed to measure the ECG, respiration waveform, SpO_2 (functional oxygen saturation of arterial hemoglobin), pulse waveform with two “AA” size (“LR06” size) alkaline batteries.
Information such as ECG measurements, respiration waveform, SpO_2 measurements, pulse waveform, battery level, and the conditions of the ECG electrodes and SpO_2 probe (sensor) are displayed on the LCD of the front panel.
ECG lead selection is available using the two buttons (ENTER and ▽) on the front panel. (In case of using a 3-electrode lead cable or a 5-electrode chest lead cable)
The LX-7230N/LX-7230N(G) can also function as a transmitter to measure only the ECG/Respiration without SpO_2 or to measure only the SpO_2 without ECG/Respiration.

Before using the LX-7230N/LX-7230N(G), read also the operation manual of the patient monitor at the receiving side thoroughly.

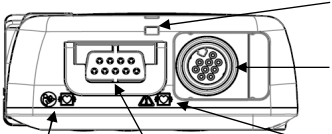
External Appearance

LX-7230N/LX-7230N(G)



1. General Description

2. Names of Parts and Their Functions



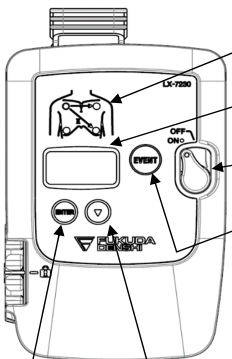
Neck Strap Hole
Attaches the neck strap.

ECG/RESP Input Connector
(Type CF Applied Part)
Connects the ECG lead cable to measure ECG and respiration waveform.

Refer to “Safety Precautions” in this manual’s preface.

Refer to “Safety Precautions” in this manual’s preface.

SpO_2 Input Connector (Type CF Applied Part)
Connects the SpO_2 probe (sensor).



Electrode Position Label
Indicates standard ECG electrode position.

LCD
Displays measurement waveform and transmitter information.

Power Switch
Turn ON/OFF the power.

EVENT Button
Activates the function assigned on the receiving monitor.

ENTER button
For setup.

▽ button
For setup.

2. Names of Parts and Their Functions

Battery Compartment Lid
This is the lid for the battery compartment.
Make sure to close it when the transmitter is operated.

NOTE
When using this equipment, use the supplied neck strap. Or use it by putting it in the pocket of the patient's clothing.

3. Preparation

1) Installing the Batteries

The LX-7230N/LX-7230N(G) functions with two "AA" size (“LR06” size) alkaline batteries.
With new batteries, the LX-7230N/LX-7230N(G) is capable of the following operation.

LX-7230N: approximately 2.5 days
LX-7230N(G): approximately 1.5 days

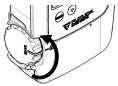
However, continuous operating time may be shorter than the above mentioned time depending on the application of the SpO_2 probe (sensor).

⚠WARNING

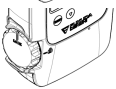
- Unplug the ECG lead cable when the battery compartment lid is opened. Otherwise, patient leakage current beyond the allowable value may occur.
- Use new "AA" size (“LR06” size) alkaline batteries.
- Do not short out the (+) and (-) terminals. It may result in exothermic heat and fire, the leakage from the batteries may damage the equipment, or an explosion from the batteries may occur.
- Install the batteries with the correct polarity.
- Do not charge alkaline batteries. Any attempt to charge the batteries may cause them to leak or break.
- Do not use a disassembled or a damaged battery due to drop or shock. The leakage from the batteries may damage the equipment, or an explosion from the batteries may occur.
- Do not use different types of batteries at the same time. The leakage from the batteries may damage the equipment, or an explosion from the batteries may occur.
- Remove the exhausted batteries immediately. The leakage from the batteries may damage the equipment, or an explosion from the batteries may occur.
- If the equipment is not in use for a long period of time, remove the batteries and store the equipment in an appropriate place. If the batteries are left in the equipment for a long period of time, the leakage from the batteries may damage the equipment or an explosion from the batteries may occur.
- Make sure to replace the two batteries simultaneously. If a new and used battery are mixed, a leakage from the batteries may damage the equipment or an explosion from the batteries may occur.

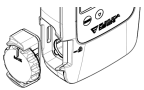
⚠CAUTION

- Use only alkaline battery (AA). Other battery will shorten the continuous operating time.
- Once the power switch is on the OFF position, then open the battery compartment lid.
- Do not replace the batteries with wet hands.
- In case of storing the used or unused batteries, make sure that the terminals are not touching other batteries or metal parts.

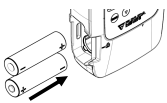


Unlock the battery compartment lid by turning the lock as shown on the left.

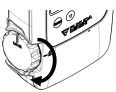


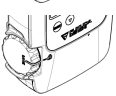


Remove the battery compartment lid.



Install new batteries according to the polarity.





Fix on the battery compartment lid by turning the lock.

⚠CAUTION

- Make sure that any foreign particles, such as hairs, are not held on the battery compartment lid and dust is not adhered to the edge of the lid to prevent water entering into the battery compartment area.
- Make sure to only turn ON the LX-7230N/LX-7230N(G) after closing the battery compartment lid.

2) Operating Power Switch

Turning the power switch to “ON”

Rotate the power switch to the left until it clicks.

LCD screen turns ON and measurement starts.
Regarding the LCD screen, refer to “7. Measurement”.
The screen automatically turns itself OFF after 180 seconds.

After the power is turned ON, make sure to check the remaining battery level on the LCD screen.
Refer to the following symbol about the remaining battery level.

Battery Symbol	Remaining Battery Level
	Full
	Getting low but still available
	Nearly empty; Replace the battery

The battery level estimation is in case of using alkaline batteries.

Turning the power switch to “OFF”

Rotate the power switch to the right until it clicks.

4. ECG Monitoring

When the transmitter is used without the SpO_2 probe (sensor), it will measure only ECG and respiration.

⚠CAUTION

When using the transmitter with only the ECG lead cable, SpO_2 measurements on the receiving monitor shall be turned off to prevent an erroneous alarm.

■ Connecting the ECG Lead Cable and Electrodes

The optional ECG lead cables for LX-7230N/LX-7230N(G) are as follows.

ECG Lead Cables

Item No.	Applicable Lead	Remark
CMT-01CTH-0.8DA	Limb Lead (1CH)	3-electrode Clip Type (White, Black, Red)
CMT-02CTH-0.8DA	Limb Lead (2CH)	4-electrode Clip Type (White, Black, Red, Green)
CMT-03CTH-0.8DA	Limb Lead (1CH) +Chest (1CH)	5-electrode Clip Type (White, Black, Red, Green, Brown)
CMT-01HTH-0.8DA	Limb Lead (1CH)	3-electrode Hook Type (White, Black, Red)
CMT-02HTH-0.8DA	Limb Lead (2CH)	4-electrode Hook Type (White, Black, Red, Green)
CMT-03HTH-0.8DA	Limb Lead (1CH) +Chest (1CH)	5-electrode Hook Type (White, Black, Red, Green, Brown)

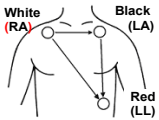
⚠WARNING

Use only the specified lead cable from Fukuda Denshi. Otherwise, proper monitoring may not be performed, and also defibrillation may fail or cause a malfunction of the equipment when the equipment is used with a defibrillator.

The relations between the attached electrode positions and lead method are as follows. Attach the electrodes to monitor proper waveform.

For 3-electrode lead cable

For AHA color code electrode position



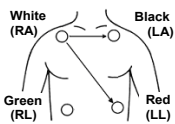
Standard Limb leads

Standard Limb leads can be selected from lead I, lead II, or lead III under the setting of the equipment.

Refer to “8. Operation ■Changing Setup ●ECG Display Screen (1) <<Switching Lead>>”.

For 4-electrode lead cable

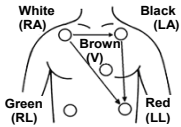
For AHA color code electrode position



Standard Limb leads

Two leads measurements, lead I and II are fixed. Lead III, aVR, aVL, and aVF can be also displayed from the setting on the receiving monitor. For details, refer to the operation manual of the receiving monitor.

For 5-electrode (Chest) lead cable
For AHA color code code electrode position



Standard Limb lead
and Chest lead

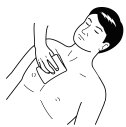
One limb lead and one chest lead (Brown) measurements are available.
Standard Limb leads can be selected from lead I, lead II, or lead III under the setting of the equipment.
The chest lead waveform is measured from the chest lead (Brown) positioned on the chest.

Refer to “8. Operation ■Changing Setup ●ECG Display Screen (1) <<Switching Lead>>”.

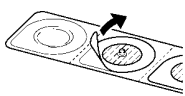
■ Attaching the Electrodes

⚠CAUTION

- Always use the same type of electrodes. If different types of electrodes are used at the same time, the difference between the polarization potential from each electrode may interfere with monitoring.
- Do not reuse the disposable electrodes. It is intended for single patient use only.



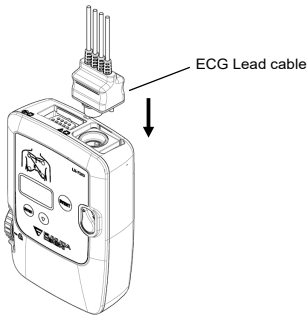
Clean the electrode sites with alcohol wipes or other skin preparation. If necessary, shave the electrode sites to remove excessive hair.



Peel off the disposable electrode.
Pay attention not to touch the electrode gel.
Attach the lead cable end to the electrode (convex part).
Turn right and left to verify that it is securely attached.

■ Connecting the ECG Lead Cable to the LX-7230N/LX-7230N(G)

Insert the ECG lead cable firmly into the ECG/RESP input connector matching the transmitter's connector guide and the direction of the notched part on the connector.



⚠CAUTION

- The threshold level for HR detection of this equipment and the receiving monitor changes with ECG waveform size. Set a proper waveform size for monitoring.
- There are some cases when pacemaker pulse cannot be detected depending on the pacemaker type, pulse voltage, pulse width, electrode lead type (unipolar, bipolar), electrode placement, or lead method which causes the pacemaker pulse amplitude to decrease and disables pacemaker pulse detection.
- If signals similar to a pacemaker pulse are present, such as electric blanket noise or excessive AC frequency noise, these may be erroneously detected and displayed as a pacemaker pulse. In this case, check the condition of the electrodes and ECG lead cable to resolve the cause or turn off the pacemaker detection setting on the receiving monitor.
- Time constant of this equipment is shorter than Fukuda Denshi monitors (direct ECG connection). Therefore, there is a difference in the ST measurement value between them. Pay attention to the difference when monitoring a patient from a transmitter or a monitor.
- When an electrode is attached on the same location for a long time, some patients may develop skin irritation. Check the patient's skin condition periodically and change the electrode position as required.
- The indication for continuous use of an electrode is about one day.
- Replace the electrode if the skin contact gets loosen due to perspiring, etc.
- Make sure to use new disposable electrodes. Otherwise, the waveform quality may become poor and it may fail to perform correct monitoring.
- When "Check Electrode" message is displayed on the screen of the receiving monitor or the LCD of this equipment, check the condition of the electrodes and ECG lead cable to resolve the cause.
- When removing electrodes from the patient, remove them carefully and slowly. Do not apply excessive force to remove them. Otherwise, it may damage the skin.
- A correct measurement may not be performed depending on the attached position of the electrodes. Attach the electrodes on the patient referring to "■ Connecting the ECG Lead Cable and Electrodes" and make sure that the correct waveform is displayed on the LCD.

5. Respiration Monitoring

Follow the preparation of “4. ECG Monitoring” to allow the respiration monitoring.

This respiration monitoring is performed with impedance method.
The ECG electrodes are also used for detecting the respiration. Each lead cable specifies the electrodes to detect the respiration. For 3-electrode and 5-electrode (chest) lead cable, the electrodes to detect the respiration are fixed as follows. Even if lead method is switched, they are no changes.

Lead Cable	Color of Electrode
3-electrode	White (RA) and Red (LL)
4-electrode	White (RA) and Red (LL)
5-electrode (Chest)	White (RA) and Red (LL)

⚠WARNING

Minute ventilation rate-adaptive implantable pacemakers can occasionally interact with certain cardiac monitoring and diagnostic equipment, causing the pacemakers to pace at their maximum programmed rate. The cardiac monitoring and diagnostic equipment may possibly send wrong information.
If such event occurs, please disconnect the cardiac monitoring and diagnostic equipment, or follow the procedures described in the operation manual of the pacemaker.
(For more details, contact FUKUDA DENSHI personnel, your institution’s professionals, or your pacemaker distributors.)

➲Reference
“Minute Ventilation Rate-Adaptive Pacemakers”
FDA alerts health professionals that minute ventilation rate-adaptive implantable pacemakers can occasionally interact with certain cardiac monitoring and diagnostic equipment, causing pacemakers to pace at their maximum programmed rate.
[October 14, 1998 – FDA]

⚠CAUTION

- Even if the electrodes are attached on the proper positions for ECG monitoring, it may not be always the proper ones for respiration monitoring as well.
- When a defibrillator is used during respiration monitoring, a large offset voltage will be placed on the ECG electrodes, which may cause interruption of monitoring for a few seconds.

5. Respiration Monitoring

6. SpO_2 Monitoring

When the transmitter is used without the ECG lead cable, it will measure only SpO_2.

⚠WARNING

When the SpO_2 probe (sensor) is in a connector-off condition, the SpO_2 alarm will not be generated on the receiving monitor. Make sure that the SpO_2 probe (sensor) is securely connected. If the SpO_2 waveform/numeric data is not displayed, check the patient's condition and pay attention not to miss the connector-off condition.

⚠CAUTION

- When using the transmitter with only the SpO_2 sensor cable, ECG and respiration measurements on the receiving monitor shall be turned off to prevent an erroneous alarm.
- The pulse wave and level meter are normalized for SpO_2 measurement. It does not indicate perfused blood volume. Check proper probe attachment by observing the pulse wave.

SpO_2 Monitoring

The LX-7230N/LX-7230N(G) measures SpO_2 with a built-in Nellcor® SpO_2 module.

< SpO_2 Sensors for LX-7230N/LX-7230N(G) **>**

The optional SpO_2 sensors available for LX-7230N/LX-7230N(G) are as follows. The following table shows applicable patient and proper site for each SpO_2 sensor. Select the proper one depending on the purpose and intended use.

Sensor Types	Applicable Patient	Applied Site
OxiMax® MAX-I	Infant (weight of 3 kg to 20 kg)	Toe
OxiMax® MAX-P	Pediatric (weight of 10 kg to 50 kg)	Finger
OxiMax® MAX-A/AL	Adult (weight of 30 kg and over)	Finger
Durasensor® DS-100A	Adult (weight of 40 kg and over)	Finger
OxiMax® MAX-R	Adult (weight of 50 kg and over)	Nose
OxiMax® MAX-FAST	Adult/Pediatric (weight of 10 kg and over)	Forehead
OxiMax® MAX-N	Adult (weight of 40 kg and over)	Finger
	Neonate (weight of less 3 kg)	Foot

⚠WARNING

- For SpO_2 monitoring, always use the sensor specified by Fukuda Denshi. Also, check the probe before usage to make sure that it is the specified probe. If unspecified probe is used, not only that the equipment cannot deliver its maximum performance, the equipment may be damaged and safety cannot be ensured.
- As with all medical equipment, carefully route cables to reduce the possibility of patient entanglement and strangulation.

⚠CAUTION

- SpO_2 sensors are not waterproof. Keep away from liquids.
- Do not pick up the equipment pulling the sensor or cable part. It may get disconnected from the equipment and the equipment may be dropped.
- A message is displayed when the SpO_2 sensor is disconnected from the equipment.
- A message is displayed when the equipment detects that the SpO_2 sensor is disconnected from the patient. Properly attach the SpO_2 sensor to the patient.
- Do not reuse the single-use SpO_2 sensor. It may cause incorrect measurements.
- Read through the instruction of the SpO_2 sensor as well.

⚠CAUTION

The accuracy of SpO_2 measurement may be influenced by abnormal hemoglobin, such as carbon monoxide hemoglobin (COHb) and methemoglobin (MetHb). It may be also affected by cardiogreen or intravascular dyes.

In addition, the following case may affect the accuracy of SpO_2 and pulse rate measurement.

- Outside light (direct sunlight, fluorescent, light therapy equipment, surgical light, infrared heat ramp, etc.)
- Hypoperfusion
- Excessive body motion (patient's motion)
- Pigment injected to the blood for testing
- In case of measurement during receiving CPR treatment
- Placement of SpO_2 sensor on limb with a blood-pressure cuff, arterial catheter
- External colorant such as nail polish
- Abnormally low or high hemoglobin concentration
- Venous pulse
- Electrosurgery
- Influence of electromagnetic waves from other electronics
- High-intensity radio waves from mobile phones

■ Applying the OxiMax® MAX-I Sensor

This Nellcor® adhesive sensor, model MAX-I, is indicated for continuous noninvasive arterial oxygen saturation and pulse rate monitoring and can be reused on the same patient as long as the adhesive tape attaches without slippage.

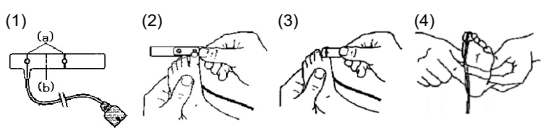
1. Remove the plastic backing from the MAX-I and locate the two transparent windows on the adhesive side. Windows cover optical components. Note the corresponding alignment marks (a) on the non-adhesive side and the dashed line (b) midway between the marks (Figure (1)).

   The big toe is the preferred MAX-I location. Alternatively, apply the sensor to another digit of similar size, for example, the thumb.

NOTE

When selecting the sensor site, priority should be given to an extremity free of an arterial catheter, blood pressure cuff, or intravascular infusion line.

2. Orient the MAX-I so that the window next to the cable is aligned on the bottom of the big toe as shown. The cable should extend towards the heel (Figure (2)).
3. Wrap the MAX-I firmly, but not too tightly around the toe. Windows must oppose each other for correct measurement (Figure (3)).
4. Wrap any excess tape loosely around the toe. Use additional tape provided to secure the cable across the bottom of the foot, loosely enough to maintain good circulation (Figure (4)).
5. Connect the MAX-I into the LX-7230N/LX-7230N(G). Verify proper operation as described in the operation manual.



NOTE

If the sensor does not track the pulse reliably, it may be incorrectly positioned – or the sensor site may be too thick, thin, or deeply pigmented, or otherwise deeply colored (for example, as a result of externally applied coloring such as nail polish, dye, or pigmented cream) to permit appropriate light transmission. If any of these situations occurs, reposition the sensor or choose an alternate Nellcor sensor to use on a different site.

Reapplication

1. The MAX-I can be reused on the same patient as long as the adhesive tape attaches without slippage.
2. Enclosed adhesive “dots” are provided for reapplication. Place a transparent dot over each window as shown, and then remove the protective paper that covers each dot (Figure (5)). The sensor is now ready to be reapplied to the same patient. For the reapplication, do not remove the previous adhesive dot, but place the enclosed adhesive dot over it.

(5)

⚠CAUTION

- Precautions for Use of Adhesive Sensor, MAX-I
  - Do not reuse the sensor on other patients. This is a sterilized product and it is intended for single patient use only.
  - Circulation distal on the sensor site should be checked routinely. The site must be inspected every 8 hours to ensure adhesion, skin integrity, and correct optical alignment. If skin integrity changes, move the sensor to another site
  - Do not use the sensor on patients who exhibit allergic reactions to the adhesive tape.
  - Failure to apply the sensor properly may cause incorrect measurements.
  - While the sensor is designed to reduce the effects of ambient light, excessive light may cause inaccurate measurements. In such cases, cover the sensor with an opaque material.
  - If the sensor is wrapped too tightly or supplemental tape is applied, venous pulsations may lead to inaccurate saturation measurements.
  - Excessive motion may compromise performance. In such cases, try to keep the patient still, or change the sensor site to one with less motion.
  - Intravascular dyes or externally applied coloring such as nail polish, dye, or pigmented cream may lead to inaccurate measurements.
  - Do not pull the sensor cable to remove the sensor from the equipment.
  - In the event of damage to the sterile packaging, do NOT use. Make sure to check whether the packaging and product is cracked or damaged before use. If there is any damage, do not use the product.
  - Do not immerse in water or cleaning solutions. Do not resterilize.

For additional warnings, cautions or contraindications when using sensors with the LX-7230N/LX-7230N(G), refer to each Nellcor SpO_2 sensor instruction manual.

■ Applying the OxiMax® MAX-P/ MAX-A/ MAX-AL Sensor

This Nellcor® adhesive sensor, model MAX-P/ MAX-A/ MAX-AL, is indicated for continuous noninvasive arterial oxygen saturation and pulse rate monitoring and can be reused on the same patient as long as the adhesive tape attaches without slippage.

1. Remove the plastic backing from the MAX-P/MAX-A/MAX-AL and locate the transparent windows (a) on the adhesive side. Windows cover optical components (Figure (1)).

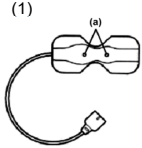
   The index finger is the preferred MAX-P/MAX-A/MAX-AL location. Alternatively, apply the sensor to the small thumb, smaller finger, or big toe.

NOTE

When selecting the sensor site, priority should be given to an extremity free of an arterial catheter, blood pressure cuff, or intravascular infusion line.

2. Orient the MAX-P/MAX-A/MAX-AL so that the dashed line in the middle of the sensor is centered on the tip of the finger/toe (Figure (2)). Wrap the adhesive flaps around the digit. Note that the cable must be positioned on the top of the hand or foot.
3. Fold the cable end over the top of the finger/toe so that the windows are directly opposite to each other. Wrap the adhesive securely around both sides of the digit (Figure (3)).
4. Connect the MAX-P/MAX-A/MAX-AL into the LX-7230N/LX-7230N(G). Verify proper operation as described in the operation manual.

(1)



(2)

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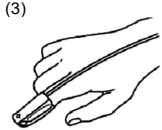


NOTE

If the sensor does not track the pulse reliably, it may be incorrectly positioned – or the sensor site may be too thick, thin, or deeply pigmented, or otherwise deeply colored (for example, as a result of externally applied coloring such as nail polish, dye, or pigmented cream) to permit appropriate light transmission. If any of these situations occurs, reposition the sensor or choose an alternate Nellcor sensor to use on a different site.

⚠CAUTION

- Precautions for Use of Adhesive Sensors, MAX-P/MAX-A/MAX-AL
  - Do not reuse the sensor on other patients. This is a sterilized product and it is intended for single patient use only.
  - Circulation distal on the sensor site should be checked routinely. The site must be inspected every 8 hours to ensure adhesion, skin integrity, and correct optical alignment. If skin integrity changes, move the sensor to another site
  - Do not use the sensor on patients who exhibit allergic reactions to the adhesive tape.
  - Failure to apply the sensor properly may cause incorrect measurements.
  - While the sensor is designed to reduce the effects of ambient light, excessive light may cause inaccurate measurements. In such cases, cover the sensor with an opaque material.
  - If the sensor is wrapped too tightly or supplemental tape is applied, venous pulsations may lead to inaccurate saturation measurements.
  - Excessive motion may compromise performance. In such cases, try to keep the patient still, or change the sensor site to one with less motion.
  - Intravascular dyes or externally applied coloring such as nail polish, dye, or pigmented cream may lead to inaccurate measurements.
  - Do not pull the sensor cable to remove the sensor from the equipment.
  - In the event of damage to the sterile packaging, do NOT use. Make sure to check whether the packaging and product is cracked or damaged before use. If there is any damage, do not use the product.
  - Do not immerse in water or cleaning solutions. Do not resterilize.

For additional warnings, cautions or contraindications when using sensors with the LX-7230N/LX-7230N(G), refer to each Nellcor SpO_2 sensor instruction manual.

■ Applying DURASENSOR® DS-100A

This Nellcor® reusable sensor, model DS-100A, is indicated for continuous noninvasive arterial oxygen saturation and pulse rate monitoring for patients weighing greater than 40 kg. The DS-100A is contraindicated for use on active patients or for prolonged use.

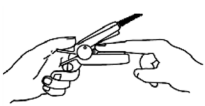
1. Place the index finger over the sensor window of the DS-100A with the finger tip against the stop (Figure (1)).
2. If the fingernail is long, the nail tip will extend over the finger stop (Figure (2)).
3. Spread open the rear tabs of the sensor to provide even force over the length of the pads (Figure (3)). If the index finger cannot be positioned correctly, or is not available, a smaller finger can be used, or use other OxiMax® sensor. Do not use the DS-100A on a thumb or toe or across a child's hand or foot.

NOTE

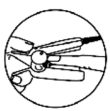
When selecting the sensor site, priority should be given to an extremity free of an arterial catheter, blood pressure cuff, or intravascular infusion line.

4. The sensor should be oriented in such a way that the cable is positioned along the top of the hand (Figure (4)).
5. Connect the DS-100A into the LX-7230N/LX-7230N(G). Verify proper operation as described in the operation manual.

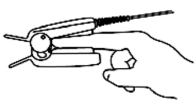
(1)



(2)



(3)



(4)

NOTE

If the sensor does not track the pulse reliably, it may be incorrectly positioned – or the sensor site may be too thick, thin, or deeply pigmented, or otherwise deeply colored (for example, as a result of externally applied coloring such as nail polish, dye, or pigmented cream) to permit appropriate light transmission. If any of these situations occurs, reposition the sensor or choose an alternate Nellcor sensor to use on a different site.

⚠CAUTION

- Precautions for Use of Reusable Sensors, DS-100A
  - Do not apply the sensor on the thumb or toe. It may cause incorrect measurements.
  - Do not use the sensor for long-term monitoring.
  - Circulation distal on the sensor site should be checked routinely. Reusable sensors must be moved to a new site at least every 4 hours. Because individual skin condition affects the ability of the skin to tolerate sensor placement, it may be necessary to change the sensor site more frequently with some patients. If skin integrity changes, move the sensor to another site. If long-term monitoring is required, use an OxiMax® sensor (MAX-A, MAX-AL, or MAX-N).
  - Failure to apply the sensor properly may cause incorrect measurements.
  - While the sensor is designed to reduce the effects of ambient light, excessive light may cause inaccurate measurements. In such cases, cover the sensor with an opaque material.
  - Do not apply tape to secure the sensor in place or to tape it shut; venous pulsations may lead to inaccurate saturation measurements.
  - Excessive motion may compromise performance. In such cases, try to keep the patient still, or change the sensor site to one with less motion.
  - Intravascular dyes or externally applied coloring such as nail polish, dye, or pigmented cream may lead to inaccurate measurements.
  - Do not pull the sensor cable to remove the sensor from the equipment.
  - In the event of damage to the sterile packaging, do NOT use. Make sure to check whether the packaging and product is cracked or damaged before use. If there is any damage, do not use the product.
  - Do not immerse in water or cleaning solutions. Do not resterilize.

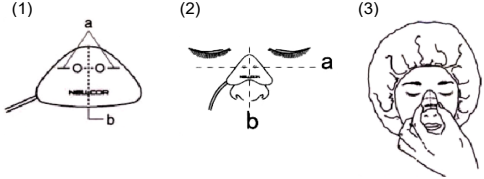
For additional warnings, cautions or contraindications when using sensors with the LX-7230N/LX-7230N(G), refer to each Nellcor SpO_2 sensor instruction manual.

■ Applying the OxiMax® MAX-R Sensor

This Nellcor® adhesive sensor, model MAX-R, is indicated for continuous noninvasive arterial oxygen saturation and pulse rate monitoring. The MAX-R is designed for use only on the nose. Use this sensor when finger pulsatile flow is inadequate, or monitoring a finger/toe is not possible.

1. Clean the bridge of the patient's nose with the contents of the enclosed acetone/alcohol ampule to remove skin oils. Do not allow the acetone/alcohol solution to get in the patient's eyes.
2. Remove the plastic backing from the MAX-R and locate the transparent windows on the adhesive side. Windows cover optical components. Note the corresponding alignment marks on the non-adhesive side (a) and the dashed center line (b) midway between the marks (Figure (1)).
3. Orient the MAX-R so that the dashed line is centered on the nose (a) and the alignment marks are at the bone-cartilage junction (b). The cable should extend toward the patient's right side (Figure (2)).
4. Press the MAX-R firmly onto the nose and hold in place for 10 seconds to ensure adhesion (Figure (3)). The MAX-R must be secured firmly for proper operation.
5. As with all medical equipment, carefully route cables to reduce the possibility of patient entanglement or strangulation.
6. Connect the MAX-R into the LX-7230N/LX-7230N(G). Verify proper operation as described in the operation manual.

(1) (2) (3)



NOTE
If the sensor does not track the pulse reliably, it may be incorrectly positioned – or the sensor site may be too thick, thin, or deeply pigmented to permit appropriate light transmission. If any of these situations occurs, try another MAX-R or choose an alternate Nellcor Puritan Bennett sensor.

⚠CAUTION

- Precautions for Use of Adhesive Sensor, MAX-R
  - Do not reuse the sensor on other patients. This is a sterilized product and it is intended for single patient use only.
  - Circulation distal on the sensor site should be checked routinely. The site must be inspected every 8 hours to ensure adhesion, skin integrity, and correct sensor site. If skin integrity changes, move the sensor to another site
  - Do not use the sensor on patients who exhibit allergic reactions to the adhesive tape.
  - Do not get the acetone/alcohol cleaning solution in the patient's eyes.
  - Failure to apply the sensor properly may cause incorrect measurements.
  - While the sensor is designed to reduce the effects of ambient light, excessive light may cause inaccurate measurements. In such cases, cover the sensor with an opaque material.
  - Intravascular dyes or externally applied coloring such as dye or pigmented cream may lead to inaccurate measurements.
  - Take care when removing the MAX-R so that the adhesive does not damage delicate facial tissue.
  - The MAX-R is not recommended for patients wearing oxygen or anesthesia masks.
  - Excessive motion may compromise performance.
  - Do not pull the sensor cable to remove the sensor from the equipment.
  - In the event of damage to the sterile packaging, do NOT use. Make sure to check whether the packaging and product is cracked or damaged before use. If there is any damage, do not use the product.
  - Do not immerse in water or cleaning solutions. Do not resterilize.

For additional warnings, cautions or contraindications when using sensors with the LX-7230N/LX-7230N(G), refer to each Nellcor SpO_2 sensor instruction manual.

■ Applying the OxiMax® MAX-N Sensor

This Nellcor® adhesive sensor, model MAX-N, is indicated for continuous noninvasive arterial oxygen saturation and pulse rate monitoring and can be reused on the same patient as long as the adhesive tape attaches without slippage.

1. Remove the plastic backing from the MAX-N and locate the two transparent windows on the adhesive side. Windows cover optical components. Note the corresponding alignment marks (a) on the non-adhesive side and the dashed line (b) midway between the marks (Figure (1)).
2. Orient the MAX-N so that the dashed line is on the lateral edge of the site (a):

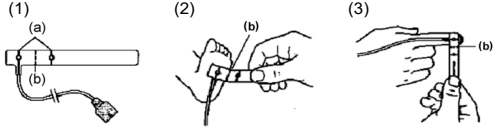
   Neonates: The preferred site is the foot. Alternatively, use the hand. The window next to the cable goes on the sole of the foot as shown (Figure (2)).

   Adults: The preferred site is the index finger. Alternatively, other fingers may be used. The window next to the cable goes on the nail side, distal to the first joint. Do not place on a joint. Note that the cable must be positioned on the top of the hand (Figure (3)).

NOTE
When selecting a sensor site, priority should be given to an extremity free of an arterial catheter, blood pressure cuff, or intravascular infusion line.

3. Wrap the MAX-N firmly, but not too tightly around the foot or finger. Windows must oppose each other.
4. Connect the MAX-N into the LX-7230N/LX-7230N(G). Verify proper operation as described in the operation manual.

(1) (2) (3)

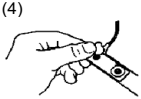


NOTE
If the sensor does not track the pulse reliably, it may be incorrectly positioned - or the sensor site may be excessively wrinkled, or too deeply pigmented or otherwise deeply colored (for example, as a result of externally applied coloring such as dye or pigmented cream) to permit appropriate light transmission. If any of these situations occurs, reposition the sensor in a different location or choose an alternate Nellcor sensor to use on a different site.

Reapplication

1. The MAX-A can be reused on the same patient as long as the adhesive tape attaches without slippage.
2. Enclosed adhesive “dots” are provided for reapplication. Place the transparent dot over each window as shown, and then remove the protective paper that covers each dot (Figure (4)). The sensor is now ready to be reapplied to the same patient. For the reapplication, do not remove the previous adhesive dot, but place the enclosed adhesive dot over it.

(4)



⚠CAUTION

- Precautions for Use of adhesive sensor, MAX-N
  - Do not reuse the sensor on other patients. This is a sterilized product and it is intended for single patient use only.
  - Circulation distal on the sensor site should be checked routinely. The site must be inspected every 8 hours to ensure adhesion, skin integrity, and correct optical alignment. If skin integrity changes, move the sensor to another site.
  - Do not use the sensor on patients who exhibit allergic reactions to the adhesive tape.
  - Failure to apply the sensor properly may cause incorrect measurements.
  - While the sensor is designed to reduce the effects of ambient light, excessive light may cause inaccurate measurements. In such cases, cover the sensor with an opaque material.
  - If the sensor is wrapped too tightly or supplemental tape is applied, venous pulsations may lead to inaccurate saturation measurements.
  - Excessive motion may compromise performance. In such cases, try to keep the patient still, or change the sensor site to one with less motion.
  - Intravascular dyes or externally applied coloring such as nail polish, dye, or pigmented cream may lead to inaccurate measurements.
  - Do not pull the sensor cable to remove the sensor from the equipment.
  - In the event of damage to the sterile packaging, do NOT use. Make sure to check whether the packaging and product is cracked or damaged before use. If there is any damage, do not use the product.
  - Do not immerse in water or cleaning solutions. Do not resterilize.

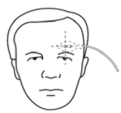
For additional warnings, cautions or contraindications when using sensors with the LX-7230N/LX-7230N(G), refer to each Nellcor SpO_2 sensor instruction manual.

■ Applying the OXIMAX® MAX-FAST Sensor

This is an adhesive sensor, model MAX-FAST, for continuous noninvasive arterial oxygen saturation and pulse rate monitoring and can be reused on the same patient as long as the adhesive tape attaches without slippage.

1. Clean the sensor site with an alcohol wipe to remove skin oils. See illustration for the recommended site. (Figure (1))

(1)



2. Remove the white paper backing to expose the first of three adhesive pads (Figure (2)). The OXIMAX® MAX-FAST sensor is now ready to be applied on the patient.

(2)

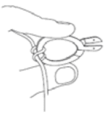
NOTE
There are three adhesive pads attached to the sensor, each with a pull-tab for removal. When repositioning the sensor on the same patient, first expose the new adhesive pad by grasping the tab and peeling off the old adhesive pad. The sensor is now ready to be reapplied to the patient.

3. Place the sensor onto a flat, hairless portion of the patient's forehead just above the left or right eyebrow. If the patient is lying on their side, place the sensor above the eye on the side of the patient's head not in contact with the bed. Press the MAX-FAST sensor firmly in place for 10 seconds, ensuring that the entire surface area of the adhesive pad makes contact with the skin (Figure (3)).

(3)

4. If desired, the sensor cable can be secured to the patient's clothing or other material by using the clip located on the cable. To open, pinch the sides of the clip; release to close (Figure (4)).

(4)



5. Connect the OXIMAX® MAX-FAST oximetry sensor into the LX-7230N/ LX-7230N(G). Verify proper operation as described in the operation manual.

NOTE

If the sensor does not track the pulse reliably, it may be incorrectly positioned - or the sensor site may be excessively wrinkled, or too deeply pigmented or otherwise deeply colored (for example, as a result of externally applied coloring such as dye or pigmented cream) to permit appropriate light transmission. If any of these situations occurs, reposition the sensor in a different location or choose an alternate Nellcor sensor to use on a different site.

⚠WARNING

- Precautions for Use of headband
  - Do not use headband on children age 24 months and younger.
  - Do not use headband on children with open fontanelles.

For details, refer to the instruction manual of OXIMAX® MAX-FAST oximetry sensor.

⚠CAUTION

- Precautions for Use of headband
  - Applying the headband too loose or too tight can cause inaccurate readings. Make sure the headband applies equal pressure to the entire sensor. The sensor must be completely covered by the headband.

For details, refer to the instruction manual of OXIMAX® MAX-FAST oximetry sensor.

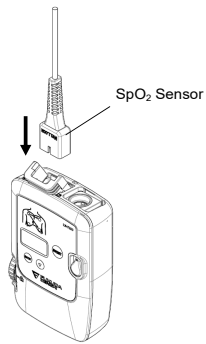
⚠CAUTION

- Precautions for Use of adhesive sensors, MAX-FAST
  - Do not reuse the sensor on other patients. This is a sterilized product and it is intended for single patient use only.
  - Circulation distal on the sensor site should be checked routinely. The site must be inspected every 12 hours to ensure adhesion, skin integrity, and correct position. Because individual skin condition affects the ability of the skin to tolerate sensor placement, it may be necessary to change the sensor site more frequently with some patients.
  - Do not use the OXIMAX® MAX-FAST sensor on patients who exhibit allergic reactions to the adhesive pad; for patients who perspire profusely; or under conditions where the patient is in the Trendelenburg position (head lower than the heart).
  - Failure to apply the sensor properly may cause incorrect measurements.
  - While the sensor is designed to reduce the effects of ambient light, excessive light may cause inaccurate measurements. In such cases, cover the sensor with an opaque material.
  - Do not use tape with the sensor. Use of additional tape or other types of adhesives may cause skin damage.
  - Applying the headband too tightly can lead to inaccurate saturation measurements, or possibly to temporary pressure marks from sensor.
  - Excessive motion may compromise performance. In such cases, try to keep the patient still, or change the sensor site to one with less motion.
  - For patients in a prone position, venous pooling and/or pulsation may cause inaccurate SpO_2 readings. Use of the headband is advised.
  - Do not pull the sensor cable to remove the sensor from the equipment.
  - In the event of damage to the sterile packaging, do NOT use. Make sure to check whether the packaging and product is cracked or damaged before use. If there is any damage, do not use the product.
  - Do not immerse in water or cleaning solutions. Do not resterilize.

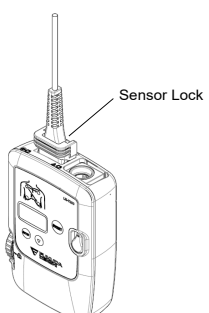
For additional warnings, cautions or contraindications when using sensors with the LX-7230N/LX-7230N(G), refer to each Nellcor SpO_2 sensor and headband instruction manual.

■ Connecting the Nellcor® SpO_2 Sensor to the LX-7230N/LX-7230N(G)

1 Insert the SpO_2 sensor into the SpO_2 input connector on the LX-7230N/LX-7230N(G).



2 Attach the sensor lock as shown in the following illustration to prevent the SpO_2 sensor to be disconnected.

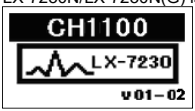


7. Measurement

Turn ON the power and the measurement starts.
*The screen examples of the LX-7230N are used for explanation.

■ Starting Screen

When the power is turned ON, the channel number configured on the LX-7230N/LX-7230N(G) is displayed at the top of the LCD.



Make sure whether the channel number on the LCD matches the channel number indicated on the label of the LX-7230N/LX-7230N(G) and the channel number configured on the receiving monitor.
This screen automatically moves onto the next waveform display screen.

■ Waveform Display Screen

ECG waveform (1CH when using 3-electrode lead cable, 2CH when using other lead cable), heart rate, pacemaker marker, respirogram, respiration rate, pulse wave, SpO_2 measurement value, remaining battery level, and various messages are displayed.

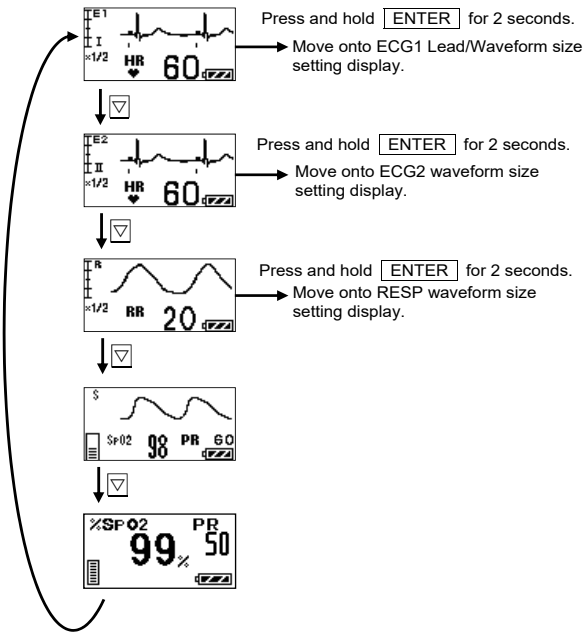
⚠CAUTION

- The LX-7230N/LX-7230N(G) does not have a diagnostic function. Check the diagnostic function on the receiving monitor.
- The LX-7230N/LX-7230N(G) does not have an alarm function. Check the alarm function on the receiving monitor.
- The ECG waveform size and sweep speed settings displayed on the LCD of the LX-7230N/LX-7230N(G) do not interface with the ones displayed on the screen of the receiving monitor.
- The heart rate and respiration rate displayed on the LCD of the LX-7230N/LX-7230N(G) may be different from the ones displayed on the receiving monitor. This is because the algorithm of the ECG and respiration rate is different.

●Display Switch

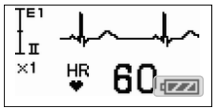
The screen (e.g. ECG) can be switched to other selected screens (respiration, pulse, or SpO_2, etc.)
The LCD display will automatically turn itself OFF after 180 seconds if no operation is done. To restart the LCD display, refer to “8. Operation ■ Restarting the LCD ”.

When the LCD display is active, press the ▽ button to move onto the next screen. The screen will be switched in the following order.



■ Battery Level Check

Check the battery level on the waveform display screen.



Battery Symbol	Remaining Battery Level
	Full
	Getting low but still available
	Nearly empty Replace the battery. A message that prompts the battery check appears on the screen of the receiving monitor.

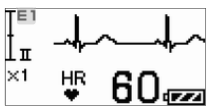
■ Waveform Display

●ECG Display Screen (1)

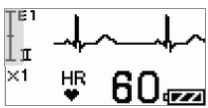
ECG1 waveform, heart rate, pacemaker marker, remaining battery level, and electrode check message are displayed.
The LCD display will automatically turn itself OFF after 180 seconds if no operation is done. To restart the LCD display, refer to “8. Operation ■ Restarting the LCD ”.

When the LCD display is active, press the ▽ button to move onto the next waveform display screen.

【Descriptions of the Screen】
The descriptions of contents displayed on the LCD are as follows.

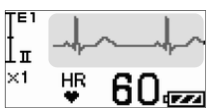


Indicates ECG 1.



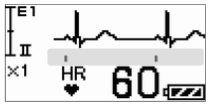
Indicates the scale of the displayed ECG.

One scale corresponds to 1 mV.
In the left illustration, it can display ECG waveform between -1 mV and +1 mV.

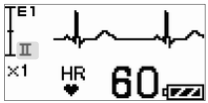


Displays the ECG waveform.

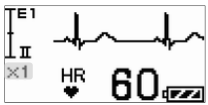
Displays the heart rate.
♥ is displayed in synchronization with the heart rate.



Displays the detection marker when the pacemaker pulse is detected.



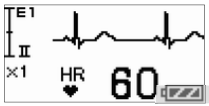
Indicates the measuring lead.



Indicates the ECG waveform size displayed on the LCD.

⚠CAUTION

- The displayed ECG waveform size setting does not interact with the one displayed on the receiving monitor, as the LX-7230N/LX-7230N(G) cannot transmit the setting information of the waveform size to the receiving monitor. If the ECG waveform size displayed on the receiving monitor is changed, follow the instruction in the operation manual of the receiving monitor.
- When “Wide” is selected in the “QRS Detection” and if HR is outside the display range (12 bpm to 300 bpm), 0 bpm will be displayed if 11 bpm and below is measured and 300 bpm will be displayed if 300 bpm and above is measured.
- When “Narrow” is selected in the “QRS Detection”, and if HR is outside the display range (30 bpm to 300 bpm), 0 bpm will be displayed if 29 bpm and below is measured and 300 bpm will be displayed if 300 bpm and above is measured.



Indicates the remaining battery level.

For details of the battery level, refer to “3. Preparation”.

“ELECTRODE?” message appears when the ECG electrode is detached.

●ECG Display Screen (2)

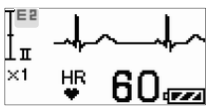
ECG2 waveform, heart rate, pacemaker marker, remaining battery level, and electrode check message are displayed.

NOTE
If a 3-electrode lead cable is used, this screen will not appear.

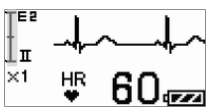
The LCD display will automatically turn itself OFF after 180 seconds if no operation is done. To restart the LCD display, refer to “8. Operation ■ Restarting the LCD ”.

When the LCD display is active, press the ▽ button to move onto the next waveform display screen.

【Descriptions of the Screen】
The descriptions of contents displayed on the LCD are as follows.

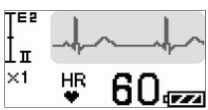


Indicates ECG 2.

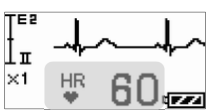


Indicates the scale of the displayed ECG.

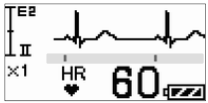
One scale corresponds to 1 mV.
In the left illustration, it can display ECG waveform between -1 mV and +1 mV.



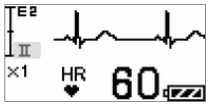
Displays the ECG waveform.



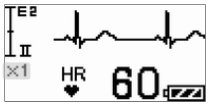
Displays the heart rate.
♥ is displayed in synchronization with the heart rate.



Displays the detection marker when the pacemaker pulse is detected.



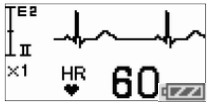
Indicates the measuring lead.



Indicates the ECG waveform size displayed on the LCD.

⚠CAUTION

- The displayed ECG waveform size setting does not interact with the one displayed on the receiving monitor, as the LX-7230N/LX-7230N(G) cannot transmit the setting information of the waveform size to the receiving monitor. If the ECG waveform size displayed on the receiving monitor is changed, follow the instruction in the operation manual of the receiving monitor.
- When “Wide” is selected in the “QRS Detection” and if HR is outside the display range (12 bpm to 300 bpm), 0 bpm will be displayed if 11 bpm and below is measured and 300 bpm will be displayed if 300 bpm and above is measured.
- When “Narrow” is selected in the “QRS Detection”, and if HR is outside the display range (30 bpm to 300 bpm), 0 bpm will be displayed if 29 bpm and below is measured and 300 bpm will be displayed if 300 bpm and above is measured.



Indicates the remaining battery level.

For details of the battery level, refer to “3. Preparation”.

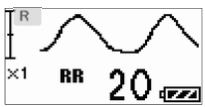
Displays the electrode check “ELECTRODE?” message appears when the ECG electrode is detached.

●Respiration Display Screen

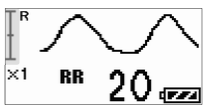
Respiration waveform, respiration rate, remaining battery level, and electrode check message are displayed.
The LCD display will automatically turn itself OFF after 180 seconds if no operation is done. To restart the LCD display, refer to “8. Operation ■ Restarting the LCD ”.

When the LCD display is active, press the ▽ button to move onto the next waveform display screen.

【Descriptions of the Screen】
The descriptions of contents displayed on the LCD are as follows.

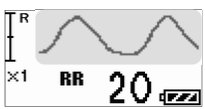


Indicates the respiration waveform display screen.



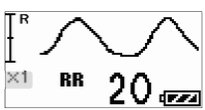
Indicates the scale of the displayed respiration waveform.

One scale corresponds to 1Ω.
In the left illustration, it can display the change of respiration waveform between -1Ω and +1Ω.



Displays the respiration waveform.

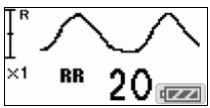
Displays the respiration rate.



Indicates the respiration waveform size displayed on the LCD.

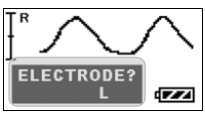
⚠CAUTION

- The displayed respiration waveform size setting does not interact with the one displayed on the receiving monitor, as the LX-7230N/LX-7230N(G) cannot transmit the setting information of the waveform size to the receiving monitor. If the respiration waveform size of the receiving monitor is changed, follow the instruction in the operation manual of the receiving monitor.
- If RR is outside the display range (9 Bpm to 150 Bpm), 0 Bpm will be displayed if 8 Bpm and below is measured and 150 Bpm will be displayed if 150 Bpm and above is measured.



Indicates the remaining battery level.

For details of the battery level, refer to “3. Preparation”.



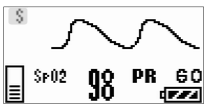
Displays the electrode check “ELECTRODE?” message appears when the ECG electrode is detached.

●SpO_2 Display Screen

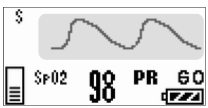
Pulse wave, pulse rate, SpO_2 measurement value, remaining battery level, and probe condition are displayed.
The LCD display will automatically turn itself OFF after 180 seconds if no operation is done. To restart the LCD display, refer to “8. Operation ■ Restarting the LCD ”.

When the LCD display is active, press the ▽ button to move onto the next waveform display screen.

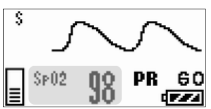
【Descriptions of the Screen】
The descriptions of contents displayed on the LCD are as follows.



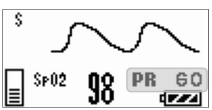
Indicates the SpO_2 display screen.



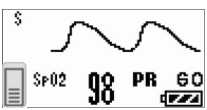
Displays the pulse wave.
The waveform size displayed on the LCD is adjusted automatically.



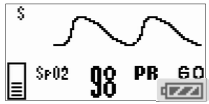
Displays the SpO_2 measurement value.



Displays the pulse rate.

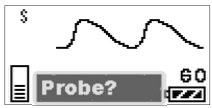


Displays the level meter of the pulse wave.
Indicates the amplitude level of the pulse wave in 8 steps.



Indicates the remaining battery level.

For details of the battery level, refer to “3. Preparation”.



Displays messages such as probe off.

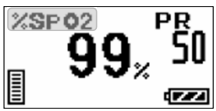
For details of messages, refer to “11. Troubleshooting”.

●SpO_2 Enlarged Display Screen

Pulse rate, SpO_2 measurement value, remaining battery level, and probe condition are displayed.
The LCD display will automatically turn itself OFF after 180 seconds if no operation is done. To restart the LCD display, refer to “8. Operation ■ Restarting the LCD ”.

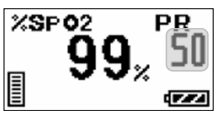
When the SpO_2 Enlarged Display Screen is displayed, press the ▽ button to move onto the next waveform display screen.

【Descriptions of the Screen】
The contents displayed on the LCD are as follows.

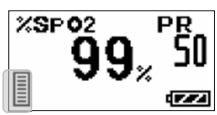


Indicates the SpO_2 display screen.

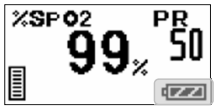
Displays the SpO_2 measurement value.



Displays the pulse rate.

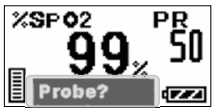


Displays the level meter of the pulse wave.
Indicate the amplitude level of the pulse wave in 8 steps.



Indicates the remaining battery level.

For details of the battery level, refer to “3. Preparation”.



Displays messages such as probe off.

For details of messages, refer to “11. Troubleshooting”.

8. Operation

* The screen examples of the LX-7230N are used for explanation.

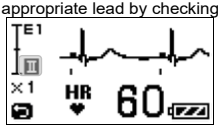
■ Changing Setup

●ECG Display Screen (1)

In the ECG display screen (1), the ECG waveform size and lead displayed on the LCD of the LX-7230N/LX-7230N(G) can be changed.

【Setting Method】
How to enter the setup mode:
Press and hold the [ENTER] button for 2 seconds in the ECG display screen (1).

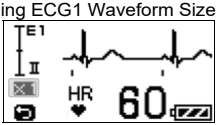
<<Switching Lead>>
Lead of ECG 1 can be switched when 3-electrode lead cable or 5-electrode (Chest) lead cable is used.
Select an appropriate lead by checking the ECG waveform on the LCD.



The lead indication of ECG 1 is highlighted.

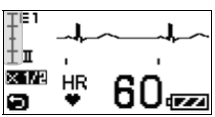
Pressing the [ENTER] button will sequentially change the lead of ECG 1.
Lead I → II → III → I

<<Changing ECG1 Waveform Size on LCD>>



Press the [▽] button to highlight the size indication of ECG 1.

Pressing the [ENTER] button will sequentially change the size of ECG 1.
Size ×1 → ×1/2 → ×1

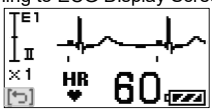


When changing the size of the ECG waveform on the LCD, the ECG scale will also change.
In the left illustration, it can display the ECG waveform between -2 mV and +2 mV.

⚠CAUTION

The displayed ECG waveform size setting does not interact with the one displayed on the receiving monitor, as the LX-7230N/LX-7230N(G) cannot transmit the setting information of the waveform size to the receiving monitor. If the ECG waveform size of the receiving monitor is changed, follow the instruction in the operation manual of the receiving monitor.

<< Returning to ECG Display Screen (1) >>



Press the ▽ button to highlight [↩].
Press the ENTER button to return to the ECG display screen (1).

⚠CAUTION

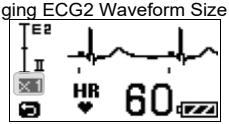
Do not operate the LX-7230N/LX-7230N(G) with the setup screen open to prevent the settings to be changed due to an unintended operation. Make sure to highlight [↩] and press ENTER to terminate the setup screen. Otherwise, the LCD display will automatically turn itself OFF after 180 seconds.

●ECG Display Screen (2)

In the ECG display screen (2), the ECG waveform size displayed on the LCD of the LX-7230N/LX-7230N(G) can be changed.

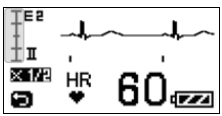
【Setting Method】
How to enter the setup mode:
Press and hold the ENTER button for 2 seconds in the ECG display screen (2).

<< Changing ECG2 Waveform Size on LCD >>



The size indication of ECG 2 is highlighted.

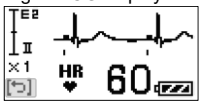
Pressing the ENTER button will sequentially change the size of ECG 2.
Size ×1 → ×1/2 → ×1



When changing the size of the ECG waveform on the LCD, the ECG scale will also change.
In the left illustration, it can display ECG waveform between -2mV and +2mV.

⚠CAUTION
The displayed ECG waveform size setting does not interact with the one displayed on the receiving monitor, as the LX-7230N/LX-7230N(G) cannot transmit the setting information of the waveform size to the receiving monitor. If the ECG waveform size of the receiving monitor is changed, follow the instruction in the operation manual of the receiving monitor.

<< Returning to ECG display screen (2) >>



Press the ▽ button to highlight [↩].
Press the ENTER button to return to the ECG display screen (2).

⚠CAUTION

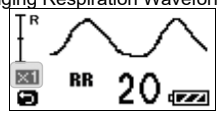
Do not operate the LX-7230N/LX-7230N(G) with the setup screen open to prevent the settings to be changed due to an unintended operation. Make sure to highlight [↩] and press ENTER to terminate the setup screen. Otherwise, the LCD display will automatically turn itself OFF after 180 seconds.

●Respiration Display Screen

In the respiration display screen, the respiration waveform size displayed on the LCD of the LX-7230N/LX-7230N(G) can be changed.

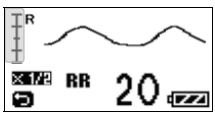
【Setting Method】
How to enter the setup mode:
Press and hold the ENTER button for 2 seconds in the respiration display screen.

<< Changing Respiration Waveform Size on LCD>>>



The size indication of the respiration is highlighted.

Pressing the ENTER button will sequentially change the size of respiration.
Size ×1 → ×1/2 → ×1

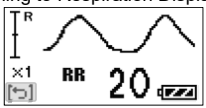


When changing the size of the respiration waveform on the LCD, the respiration scale will also change.
In the left illustration, it can display the respiration waveform until 4Ω of change.

⚠CAUTION

The displayed respiration waveform size setting does not interact with the one displayed on the receiving monitor, as the LX-7230N/LX-7230N(G) cannot transmit the setting information of the waveform size to the receiving monitor. If the respiration waveform size displayed on the receiving monitor is changed, follow the instruction in the operation manual of the receiving monitor.

<< Returning to Respiration Display Screen>>



Press the ▽ button to highlight [↩].
Press the ENTER button to return to the respiration display screen.

⚠CAUTION

Do not operate the LX-7230N/LX-7230N(G) with the setup screen open to prevent the settings to be changed due to an unintended operation. Make sure to highlight [↩] and press ENTER to terminate the setup screen. Otherwise, the LCD display will automatically turn itself OFF after 180 seconds.

●SpO_2 Display Screen

LX-7230N/LX-7230N(G) has no setup item in the SpO_2 display screen.

●SpO_2 Enlarged Display Screen

LX-7230N/LX-7230N(G) has no setup item in the SpO_2 display screen

■ Restarting the LCD Display

The LCD display will automatically turn itself OFF after 180 seconds if no operation is done.
Press the [ENTER] button or press and hold the [▽] button to restart the LCD display.
The starting screen with telemetry channel number appears, and then the waveform display screen appears.

■ Pressing the EVENT Button

Press and hold the [EVENT] button for 2 seconds to activate the function assigned on the receiving monitor. The following message appears on the LCD while the “EVENT” is transmitted.

After the transmission is completed, the starting screen with the telemetry channel number appears, and then the waveform display screen appears.
“EVENT” operation is available as a remote recording.
For details of the receiving monitor operation and settings related to the “EVENT” function, refer to the operation manual of the receiving monitor.

9. Other Setting Items

The following settings are available for the LX-7230N/LX-7230N(G) depending on the usage and condition of the patient. For details of the settings, contact our service representative.

Items		Selection	Default	Backup
ECG				
	Time Constant	0.4 s, 0.1 s	0.4 s	Yes
	Pace Detect.	L, M, H	M	Yes
	QRS Width	Narrow, Wide	Wide	Yes
RESP				
	Respiration Detection Signal (Display)	ON, OFF	ON	Yes
Display				
	Contrast	8 steps	8	Yes
Channel				
	LX-7230N Channel	One from the following channels. 0801 to 0879 0900 to 0979 1000 to 1079 1100 to 1179 1200 to 1279 1300 to 1379	1100	Yes
	LX-7230N(G) Channel	One from the following channels. 9501 to 9539 9600 to 9639 9700 to 9739 9800 to 9839 9900 to 9938 2701 to 2739 2800 to 2839 2900 to 2918 2921 to 2939 3000 to 3039 3100 to 3118	9501	Yes
	Group	One from 00 to 63	00	Yes

■ Changing the Time Constant

The default setting of the time constant is “0.4 second”.
If a stable monitoring is difficult with excessive change in the baseline due to excessive body motion of the patient or an interference noise, such as AC frequency, by changing the time constant to “0.1 second” the monitoring may become relatively stable.

For details of the setting change, contact your local Fukuda Denshi service representative.

⚠CAUTION

- The threshold level for HR detection of this equipment and the receiving monitor changes with ECG waveform size. Set a proper waveform size for monitoring.
- When changing the time constant to “0.1 second”, the lower frequency characteristic becomes 1.6Hz ±25%. This setup does not meet EN 60601-2-27 standard. It may lead to a change in the ECG waveform and ST measurement value may be especially affected. Fukuda Denshi recommends “0.4 second” setting in normal use.
- The LCD screen in normal use does not indicate the selection of time constant. Make sure to take measures, such as marking on the LX-7230N/LX-7230N(G), to distinguish whether the selection of time constant is changed.

■ Changing the Detection Sensitivity of the Pacemaker Pulse

The default setting of pacemaker pulse detection sensitivity is “Mid”.
The “Mid” setting can detect and reject the following pacemaker pulse specified in IEC 60601-2-27 standard.

Detection/ Rejection of Pacemaker Pulse:

a) Pacemaker Pulse without Over/Undershoot:
Capable to reject pulses of pulse width 0.1 to 2ms, amplitude ±2 to ±700mV

b) Pacemaker Pulse with Over/Undershoot:
Rejection is not possible.

Fukuda Denshi recommends the “Mid” setting in normal use.

There may be some cases when the pacemaker pulse cannot be detected depending on the pacemaker type, pulse voltage, pulse width, electrode lead type (unipolar, bipolar). In this case, change the lead or the position of the electrodes to be able to detect the pacemaker pulse.
Nonetheless, if the detection is still undetectable, change the setting to “High” in order to increase the detection sensitivity. So that smaller pacemaker pulse can be detected. However, the “High” setting may lead to erroneous detection due to interference noise, such as AC frequency.

If erroneous detections occur due to interference noise, such as AC frequency, turn OFF the setting of the pacemaker pulse detection in the receiving monitor.
If erroneous detections occur due to interference noise, such as AC frequency, while monitoring a patient with a pacemaker, and the setting of the pacemaker pulse detection cannot be turned OFF, replace the electrodes or change the lead to remove the interference noise, such as AC frequency.

Nonetheless, if erroneous detections still occur, change the setting to “Low” in order to decrease the detection sensitivity. It makes the LX-7230N/LX-7230N(G) less likely to be interfered by the noise, such as AC frequency.
The “Low” setting decreases the detection sensitivity. Therefore, it cannot detect the pacemaker pulse specified in IEC 60601-2-27 standard.

For details of the setting change, contact your local Fukuda Denshi service representative.

⚠CAUTION
The LCD screen in normal use does not indicate the setting status of the pacemaker pulse detection. Make sure to take measures, such as marking on the LX-7230N/LX-7230N(G), to distinguish whether the setting of the pacemaker pulse detection is changed.

■ Changing the Respiration Detection Signal ON/OFF

The default setting of the respiration detection signal is “ON”.
The respiration waveform can be detected when the setting of the respiration detection signal is turned “ON”.

⚠WARNING

If the LX-7230N/LX-7230N(G) is used with minute ventilation rate-adaptive implantable pacemaker, the respiration detection signal may cause the pacemaker to pace at its maximum programmed rate. If such event occurs, change the setting to “OFF” to prevent an occurrence of erroneous pacing rate.

For details of the setting change, contact your local Fukuda Denshi service representative.

⚠CAUTION

- The respiration waveform cannot be measured if the setting of the respiration detection signal is turned “OFF”.
- Make sure to turn OFF the respiration measurement function on the receiving monitor to prevent an erroneous detection of the respiration alarm (on the receiving monitor side).
- The LCD screen in normal use does not indicate the setting status of the respiration detection signal ON/OFF. Make sure to take measures, such as marking on the LX-7230N/LX-7230N(G), to distinguish whether the setting of the respiration detection signal ON/OFF is changed.

■ Changing the LCD Contrast

The LCD display contrast of the LX-7230N/LX-7230N(G) can be changed in 8 steps.

For details of the setting change, contact your local Fukuda Denshi service representative.

10. Changing the Transmitter Channel and Group ID

■ Changing the Transmitter Channel

The LX-7230N/LX-7230N(G) is a transmitter of PLL synthesizer type, and its transmitter channel can be programmed. It can be set up with an arbitrary channel among the channels assigned by the Telemetry Laws (according to each country).

For details of the setting change, contact your local Fukuda Denshi service representative.

⚠WARNING

- If the transmitter channel is changed, follow the instruction of the person in charge of the radio telemetry channel in your facility. Mismanagement may result in a serious accident, such as interference and mixing up patients.
- Replace promptly the new channel label if the transmitter channel has been changed.

■ Changing the Group ID

The LX-7230N/LX-7230N(G) transmits its group ID, which it belongs to, to prevent interference with neighboring hospital's transmitter.
The receiving monitor checks whether the incoming group ID is the same as that of the receiving monitor. There are 64 group codes available. The default setting is “00”.
The transmitter group ID can be changed if there is interference with a neighboring hospital's transmitter.

For details of the setting change, contact your local Fukuda Denshi service representative.

⚠CAUTION

Possible causes of interference other than radio telemetry from neighboring hospital's transmitter, are the proximity of mobile phone, amateur radio station, radio taxi, and illegal citizens band, which may be a cause of interference. In such case, the situation should be carefully observed to find the cause of interference.

10. Changing the Transmitter Channel and Group ID

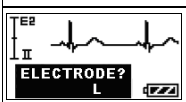
11. Troubleshooting

■ List of Displayed Messages

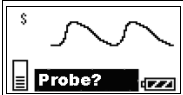
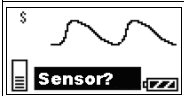
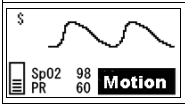
Transmitter (Main Unit)

Message	Cause	Solution
	Failed to transmit waveform and value.	Contact your local Fukuda Denshi service representative.
	Faulty EEPROM.	
	Failed to initialize CPU.	

ECG

Message	Cause	Solution
 Character string displayed, such as L, depends on the detached electrode position.	Electrode is off.	Check the electrode condition.

SpO_2

Message	Cause	Solution
	Probe is off.	Check the attached condition of the probe.
	Faulty Probe.	Replace the probe with a new one.
	SpO_2 is not measured correctly.	Check the attached condition of the probe.
		Cover the probe with an opaque material to cut off the outside light.
	Due to excessive body motion, SpO_2 is not measured correctly	Keep the patient still.

■ Troubleshooting

Make sure of the following. However, if there is no improvement in the phenomenon, contact your local Fukuda Denshi service representative.

Transmitter (Main Unit)

Phenomenon	Cause	Solution
Nothing is displayed on the LCD when the power switch is turned ON.	No battery or wrong polarity	Install the battery correctly.
	Battery level is empty.	Replace the battery with a new one.
Nothing is displayed on the receiving monitor screen.	The channel number between the transmitter and the receiving monitor do not match up.	Set the same channel number for the transmitter and the receiving monitor.
Transmission problem.	Same channel number is already used.	Make sure to not duplicate channel numbers. Follow the instruction by the person in charge of radio telemetry channel in your facility and use the LX-7230N/LX-7230N(G) with the correct channel setting.
	Channel interference	Follow the instruction by the person in charge of radio telemetry channel in your facility and use the LX-7230N/LX-7230N(G) with the correct channel setting.
	Transmitter failure	Contact your local Fukuda Denshi service representative.

ECG

Phenomenon	Cause	Solution
“ELECTRODE?” message is displayed.	Lead cable is off.	Check the connection between the lead cable and the LX-7230N/LX-7230N(G).
		Check the connection between the lead cable and the electrode.
	Faulty Lead cable.	Replace the ECG cable with a new one.
	Electrode is peeling off.	Replace the electrode with a new one.
	Polarization potential of the electrode is too high.	Replace the electrode with a new one.
ECG waveform contains noise	Electrode gel is dry.	Replace the electrode with a new one.
	Electrode is peeling off.	
	Electric blanket is used.	Cover the electric blanket with a shield cover.
	AC filter setting of the receiving monitor is OFF.	Set the AC filter up as ON.
Respiration waveform cannot be measured.	Electrode gel is dry.	Replace the electrode with a new one.
	Electrode is peeling off.	
	The positions of the electrodes are improper.	Attach the electrodes where the respiration waveform can be measured appropriately.

SpO_2

Phenomenon	Cause	Solution
SpO_2 value is unstable.	The probe size is improper.	Use a probe, which fit properly.
	The probe is peeling off or is affected by the outside light due to the poor condition	Attach the probe properly following the instruction.
	Transmitting and measuring LEDs sensor are dirty.	Clean both LED sensors from dirt.

■ In Case of Dropping the LX-7230N/LX-7230N(G) into Water

In case of dropping the LX-7230N/LX-7230N(G) into water containing disinfectant, pick up the LX-7230N/LX-7230N(G) as soon as possible.
In case of dropping the equipment into dirty water, clean it without disconnecting the ECG lead cable and SpO_2 probe (sensor), and make sure that the battery compartment lid is locked. After cleaning, wipe off any moisture thoroughly before removing the ECG lead cable, SpO_2 probe (sensor), or batteries.

⚠CAUTION

- Do not use a dryer. The LX-7230N/LX-7230N(G) shape may change or be broken.
- When rinsing the LX-7230N/LX-7230N(G) with running water, make sure to close the battery compartment lid.
- In case of dropping the equipment into dirty water, it is recommended to contact Fukuda Denshi or your nearest service representative.
- If it is difficult to clean the connector part, or if an inadequate contact occurs, contact Fukuda Denshi or your nearest service representative.

12. Cleaning and Disinfection

The cleaning and disinfection of the LX-7230N/LX-7230N(G), ECG lead cable, and SpO_2 probe (sensor) shall be performed as follows.

⚠CAUTION

Do not sterilize the LX-7230N/LX-7230N(G), ECG lead cable, and SpO_2 probe (sensor) in any manners, such as radioactive rays, steam, or ethylene oxide.

■ Cleaning and Disinfecting the LX-7230N/LX-7230N(G)

●Cleaning

【About Cleaning the Housing】

1. Clean the equipment using squeezed gauze or an absorbent cotton cloth dampened with alcohol or a neutral cleanser. When cleaning, do not allow any solution to enter the equipment or connectors. Also, do not use organic solvents, thinner, toluene and benzene to avoid damaging the resin case.
2. In case of dropping the equipment into dirty water, clean it without disconnecting the ECG lead cable and SpO_2 probe (sensor), and make sure that the battery compartment lid is locked. After cleaning, wipe off any moisture thoroughly before removing the ECG lead cable, SpO_2 probe (sensor), or batteries.

⚠CAUTION

- Clean the equipment frequently so stains can be removed easily.
- To prevent injury, it is recommended to wear gloves when cleaning the equipment.
- Do not allow any chemical solution to enter the LX-7230N/LX-7230N(G) or connectors.
- The LX-7230N/LX-7230N(G) cannot be sterilized.
- Do not use organic solvents, thinner, toluene and benzene to avoid damaging the resin case.
- Do not polish the housing with abrasive or chemical cleaner.
- Use only neutral detergent to clean the housing. Do not use chemical cloth, scrub brush, abrasive, polishing powder, hot water, volatile solvent and chemicals (cleanser, thinner, benzine, benzol, and synthetic detergent for house and furniture), or sharp-edged tools. The surface resin coating may be damaged, resulting in discoloration, scratches, and other problems.

【About Cleaning the Connector】

Do not wipe the ECG connector and SpO_2 connector with a swab, gauze, or absorbent cotton.
Use an air duster to clean the dust and dirt in the connector.

⚠CAUTION
If a swab, gauze, or cotton is used to clean the connector, dust or cotton fibers may enter the connector causing inadequate contact. Also, chemical solution may enter the connector causing inadequate contact. If it occurs, correct measurement cannot be performed and the alarm may not be generated. If cleaning the connector is needed, contact Fukuda Denshi or your nearest service representative.

●Disinfection

If there is a possibility of being infected, clean the LX-7230N/LX-7230N(G) using a squeezed gauze or an absorbent cotton cloth dampened with alcohol or invert soap.

⚠CAUTION
● Do not immerse the connector parts of the LX-7230N/LX-7230N(G) in any chemical solution to prevent connection failure. ● When disinfecting the entire room using a spray solution, pay close attention not to have liquids get into the LX-7230N/LX-7230N(G) or connectors.

■ Cleaning the ECG lead cable

After using the cable, clean it with neutral detergent or 70% isopropyl alcohol.

⚠CAUTION
● Do not use thinner, toluene, or other organic solvents to clean the cables. ● Do not pull the cable and do not hold the connector part when cleaning. (It may degrade the cable coating and result in damage. Particularly organic solvents and antiseptic solution such as cresol soap solution will degrade the cable coating.) ● After cleaning, dry it completely before usage. ● Do not use high temperature sterilization such as steam or EOG method.

■ Cleaning and Disinfecting the SpO_2 Probe (Sensor)

●NELLCOR® SpO_2 sensors

After using the the Durasensor® (DS-100A) on each patient, clean it with 70% isopropyl alcohol. If low-level disinfection is required, use a 1:10 bleach solution. Any unproved cleaning solutions may case early failure of the sensors.

The OxiMax® can be reused on the same patient as long as the adhesive tape attaches without slippage. Do not resterilize and reuse it on other patients. It is intended for single patient use only.

13. Maintenance and Inspection

This section explains the daily checks and periodic checks of the LX-7230N /LX-7230N(G).
To ensure safety, reliability, and high performance, a “Daily Check” and “Periodic Check” must be performed. We are not liable for any accident arising from lack of maintenance.

⚠CAUTION
● Do not open the housing or attempt service. The service should be done by Fukuda Denshi or Fukuda Denshi’s representative. ● Do not allow excessive moisture or cleaning agents into the connectors or inside the equipment.

■ Daily Check

Perform daily checks using the “Daily Check List”.

■ Periodic Check

Periodic check of medical electronic equipment is mandatory to prevent failures and accidents and to ensure safety and reliability.
Periodic maintenance may be performed by the medical institution or by a third party by concluding a “Maintenance Contract”.
For more details, contact your local Fukuda Denshi service representative.

Perform Periodic check using the “Periodic Check List”.The periodic check should be performed once a year.
If there is an item with “Fail” judgement, the overall judgement will be “Fail”.
Make sure to take countermeasures for the “Fail” item.
Use the equipment only if the judgements of all the items are “Pass”.

■ Periodic Replacement Parts

The “**Battery Compartment Lid (Waterproof)**” is the only periodic replacement part.
To ensure the reliability of waterproof (IPX8) performance of the LX-7230N/LX-7230N(G), replace it once a year.
It may be possible to keep using the LX-7230N/LX-7230N(G) without periodic replacement of the lid. However, as it gets older, the reliability of water resistance (IPX8) performance will not be ensured.
When replacing the lid, contact your local Fukuda Denshi service representative.

⚠CAUTION
The periodic replacement parts must be replaced at specified period.

Daily Check List

No.

Inspected Date Inspected by Location

Device Type S/No. Date of Purchase

Items	*Details*	*Criteria*	*Judgment*
Appearance	Visually check for any damage, cracks, chip, peeled label, and loosen screw on the housing.	No abnormality should be found.	☐OK/ ☐NG
Battery Compartment	Visually check for the ring condition of the battery compartment lid.	No damage, kink, floating, and adhesion of dust should be found.	☐OK/ ☐NG
	Visually check for the contact springs, inside the LX-7230N/LX-7230N(G), to the battery and the lock lever of the battery compartment lid.	No deformation, cracks, and rust should be found.	☐OK/ ☐NG
Power Supply	Turn the power ON/OFF to verify proper switch operation.	With batteries installed, the LCD should turn ON.	☐OK/ ☐NG
ECG Connectors	Visually check the connectors of the cable and the LX-7230N/LX-7230N(G).	No damage, chip, and adhesion of dust should be found.	☐OK/ ☐NG
ECG Lead cable	Visually check each lead for damages.	No crack and damage should be found.	☐OK/ ☐NG
SpO_2 Sensor (Probe)	Visually check the cable, optical receiver, LED, and connector for damages.	No crack, chip, damage, and adhesion of dust should be found.	☐OK/ ☐NG
Wireless Channel	Verify whether the transmitting channel and group ID are the same with the receiving monitor.	Must match the wireless channel check list.	☐OK/ ☐NG
Transmission Function	Turn the power ON and make sure the information is displayed on the receiving monitor.	Waveforms and values should be received without any problem.	☐OK/ ☐NG
Display Function	Turn the power ON and verify each display condition, such as SpO_2 value and bar graph.	All data should be properly displayed.	☐OK/ ☐NG
Periodic Check	Check the date of the previous periodic check.	Should be within one year.	☐OK/ ☐NG

Comment

■Periodic Check

The periodic maintenance check is intended to check the medical equipment used daily in a medical institution to prevent failures and accidents and to ensure safety and reliability.
The check procedures are described for daily and periodic checks. Each check item must be performed according to the described check procedure.
The consignee can select the check items according to the product quality, frequency of usage, and maintenance check period. However, electrical safety items must also be performed.
For details of the electrical safety check procedure, refer to IEC 60601-1.

●Periodic Check Items

The periodic check items are as follows.

No.	Check Item
1	External Appearance
2	Power Supply Switch
3	Display / Operation / Record
4	ECG
5	Respiration
6	Arterial Oxygen Saturation (SpO_2)
7	Electrical Safety

No.	Check Item	Check Procedure	Criteria
1. External Appearance, Accessories			
01	Appearance	Visually check the exterior for scratches, cracks, deformation, and rust.	No abnormality should be found.
02	Cables	Visually check all cables for any damage or being disconnected.	No damage should be found.
03	Operation Manual	Check if the operation manual and other accompanying documents are stored in the specified places.	Should be stored in the specified place.

No	Check Item	Check Procedure	Criteria
2. Power Supply Switch			
01	Power Supply Switch	Turn ON/OFF the power switch.	Should turn ON/OFF the power switch properly.

No	Check Item	Check Procedure	Criteria
3. Display, Operation, Record			
01	Labels	Visually check the labels, caution labels, etc.	Should be clean, clear and firmly attached.
02	Operation, Switches and keys	Check by operating the switches and keys.	Should operate properly.
03	LCD	Check that the characters and waveforms appear on the LCD.	The characters and waveforms should be clearly displayed. The brightness should be sufficient.

No	Check Item	Check Procedure	Criteria
4. ECG			
01	Input Impedance*	Measure input impedance with safety tester.	Should be 5MΩ or above.
02	Suppression Characteristic of Common-Mode Signal*	Input 50Hz or 60Hz 20Vr.m.s. signal, and measure fluctuation of each lead.	Should be 10 mmp-p or below for standard sensitivity (sensitivity 1).
03	Transient Characteristic*	With comprehensive tester, apply standard voltage of 1 mV, and check the time the amplitude natural logarithmically drops and becomes 37% of the waveform of 0.04 sec. after the application of standard voltage.	Should be 0.4 ± 0.1 seconds. (When the time constant is set to 0.4 seconds.)
04	Frequency Characteristic*	With comprehensive tester, apply sinusoidal voltage. Measure the frequency characteristic at test voltage of 40 Hz.	Should be 40 Hz (-3 dB) or above.
05	Heart Rhythm Detection*	With comprehensive tester, input both positive and negative polarity of 0.3 mV and 3 mV with sensitivity 1.	The heart beat rhythm should be detected with sensitivity 1 according to the peak-to-peak signal of 0.3 mV and 3 mV.
06	ECG Sensitivity	With comprehensive tester, apply 1mV voltage and measure the displayed amplitude.	Wave form size on the receiving monitor should be within 0mm ±10% at sensitivity 1.

07	Heart Rate Accuracy*	With ECG simulator, test heartbeat 60, 180 beats/min., and check the displayed HR value.	For reference heartbeat signal of 60, 180 beats/min., error of the displayed HR value should be within ±3 beats/min.
08	ECG Lead Switch	With ECG simulator, check that each lead is displayed properly. (Check for 3-electrode, and 4-electrode, 5-electrode.)	For each lead cable, lead should be correctly switched, and waveform should be correctly displayed.
09	Lead-Off Indication	Remove each electrode, and check that lead-off message is displayed.	Lead-Off message for the corresponded lead should be displayed.
10	ECG Lead Cable Recognition	Switch the ECG lead cable or switch the setup of lead cable.	Should correctly recognize the connected lead cable.

* As these functions are dependent on the design or software, these items are not mandatory for periodic checks. Perform the test as necessary.

No	Check Item	Check Procedure	Criteria
5. Respiration			
01	Respiration Waveform Sensitivity	With comprehensive tester or reference respiration signal generator, input sinusoidal waveform of 0.5 Hz with base resistance of 1.5 kΩ / 1 Ω change.	The amplitude displayed on the receiving monitor screen should be within 10 mm ± 2 mm.
02	Respiration Rate Accuracy*	Input reference respiration signal to comprehensive tester or respiration simulator, and check the respiration rate display. Test with the respiration reference load signal of 60, 120/min.	Error should be within ±5 Bpm.
03	Frequency Characteristic*	With comprehensive tester or reference respiration signal generator, input 0.5 Hz and 1.5 Hz or 2.5 Hz, and measure the frequency characteristic.	The crest value at 1.5 Hz should be more than 70% of the crest value at 0.5 Hz input.

* As these functions are dependent on the design or software, these items are not mandatory for periodic checks. Perform the test as necessary.

No	Check Item	Check Procedure	Criteria
6. Arterial Oxygen Saturation (SpO_2)			
01	SpO_2 Accuracy	Measure the error at 75%, 90% using a SpO_2 simulator.	Error should be within ±2% for SpO_2 of 70−100%.
		Prepare other reference device, perform measurement on healthy subject, and compare the value.	Error between the 2 devices should be within ±4%.
02	Pulse Rate Accuracy	Input 60, 200bpm using the SpO_2 simulator, and measure the error.	Error should be within ±3bpm (20-250bpm)
		Prepare other reference device, perform measurement on healthy subject, and compare the value.	Error between the 2 devices should be within ±6%.
03	SpO_2 Probe-Off Detection	Check the display by disconnecting the probe.	Waveform and numeric data should disappear from the display.

No	Check Item	Check Procedure	Criteria
7. Electrical Safety			
01	Contact current	Measure the leakage current that runs through the ground from the enclosure of the device under normal condition using a leak measurement safety tester. According to test procedure of IEC 60601-1 8.7.4	From the enclosure to the ground (NC) ≤0.1mA.
02	Patient leakage current that runs through the ground from the patient connecting part (NC)	Measure the patient leakage current that runs through the ground from the patient connecting part using a leak measurement safety tester. According to test procedure of IEC 60601-1 8.7.4	【AC/DC】 From the patient connecting part to the ground (NC) ≤0.01mA.
03	Leakage current when external voltage is applied to the patient connection in the Type F attaching part (SFC)	Measure the leakage current when external voltage is applied to the patient connection in the Type F attaching part using a leak measurement safety tester. According to test procedure of IEC 60601-1 8.7.4	Leakage current when external voltage is applied to the patient connection in the Type F attaching part. (SFC) ≤0.05mA.

No	Check Item	Check Procedure	Criteria
04	Total patient leakage current that runs through the ground from the total patient connecting part. (NC)	Measure the total patient leakage current that runs through the ground from the patient. According to test procedure of IEC 60601-1 8.7.4	【AC/DC】 From the patient connecting part to the ground (NC) ≤0.05mA.
05	Leakage current when external voltage is applied to the patient connection in the Type F attaching part (SFC)	Measure the leakage current when external voltage is applied to the patient connection in the Type F attaching part using a leak measurement safety tester According to test procedure of IEC 60601-1 8.7.4	Leakage current when external voltage is applied to the patient connection in the Type F attaching part. (SFC) ≤0.01mA.
06	Patient auxiliary current (NC)	Measure the patient auxiliary current (NC) using a leak measurement safety tester According to test procedure of IEC 60601-1 8.7.4	Patient auxiliary current (NC) ≤0.01mA.

No	Check Item	Check Procedure	Criteria
7. Electrical Safety (*) Perform the following check item as appropriate. Check these items when you have disassembled the equipment to check/ replace the boards or units.			
07	Withstand Voltage Test (the enclosure – isolated connecting part)	Apply AC 1500V for 1 minute between the enclosure and a connecting part. Note: The voltage differs depending on the internal protective circuit composition of the equipment. According to test procedure of IEC 60601-1 8.8.3	Should withstand applied voltage.

Periodic Check List

Telemetry Transmitter Periodic Check Report

Check Date

Location		Delivery Date		Periodic Check Contract ☐Yes ☐No
		Customer Code		Month Check
Model Name	Serial No.:	Product Code		Next Check Date
Requested Item			Acceptance Date	

No.	Check Item	Judge	Check
1	**Exterior, Accessories**		
01	Exterior	OK NG	
02	Cables	OK NG	
03	Operation Manuals	OK NG	
2	**Power Supply Switch**		
01	Power Supply Switch	OK NG	
3	**Display, Operation, Record**		
01	Labels	OK NG	
02	Operation Switch/Key	OK NG	
03	LCD	OK NG	
4	**ECG**		
01	Input Impedance	OK NG	
02	Suppression Characteristic of Common-Mode Signal	OK NG	
03	Transient Characteristic	OK NG	
04	Frequency Characteristic	OK NG	
05	Heart Rhythm Detection	OK NG	
06	ECG Sensitivity	OK NG	
07	Heart Rate Accuracy	OK NG	
08	ECG Lead Switch	OK NG	
09	Lead-Off indication	OK NG	
10	ECG Lead Cable Recognition	OK NG	

No.	Check Item	Judge	Check
5	**Respiration**		
01	Respiration Waveform Sensitivity	OK NG	
02	RR Accuracy	OK NG	
03	Frequency Characteristic	OK NG	
6	**SpO_2**		
01	SpO_2 Accuracy	OK NG	
02	PR Accuracy	OK NG	
03	SpO_2 Probe-Off Detection	OK NG	

No.	Check Item	Judge	Check
7	**Electrical Safety**		
01	From the patient connecting part to the ground NC ( ) mA	OK NG	
02	Patient leakage current that runs through the ground from the patient connecting part NC ( ) mA	OK NG	
03	Leakage current when external voltage is applied to the patient connection in the Type F attaching part. SFC ( ) mA	OK NG	
04	Total patient leakage current that runs through the ground from the total patient connecting part. NC ( ) mA	OK NG	
05	Leakage current when external voltage is applied to the patient connection in the Type F attaching part. SFC ( ) mA	OK NG	
06	Patient auxiliary current NC ( ) mA	OK NG	
07	Withstand Voltage Test	OK NG	

Description

Lv	Check	A	Adjustment
×	Replacement part	C	Cleaning
／	Not covered	R	Repair

The check result is as follows:
☐Normal Operation ☐Malfunctioning. ☐Needs to be repaired.
(Details of malfunction and repair)

Company

Inspector

Person in charge

Replacement parts

Classification
☐On-site ☐Taking-over
☐Holiday ☐Night

■Repairing the Equipment

This equipment is basically repaired at Fukuda Denshi factory.
If detailed information about the repair is needed, contact Fukuda Denshi.

On-site repair is possible for the following parts.

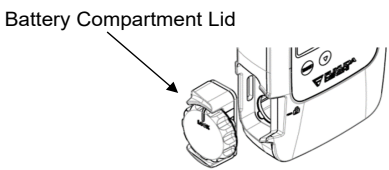
- Replacing the battery compartment lid

⚠CAUTION
Make sure to replace the parts correctly. Otherwise, it may cause damage and heat generation of the equipment.

■Replacing the Battery Compartment Lid

●Life of the Battery Compartment Lid

Life of the waterproof battery compartment lid is one year. If this unit is used for more than a year, the waterproof (IPX8) performance cannot be guaranteed. Replace the battery compartment lid to maintain its waterproof performance.

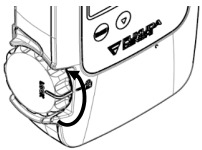


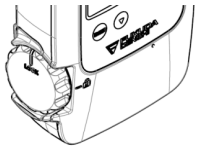
⚠CAUTION
The battery compartment lid must be replaced at specified period. Even if the LX-7230N/LX-7230N(G) is used less than one (1) year, the unit may be damaged from high impact. If the LX-7230N/LX-7230N(G) is dropped or is subjected to a high impact, make sure that the unit is not damaged.

●Assembly and Disassembly

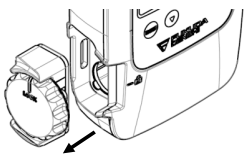
Follow the procedure below to remove the battery compartment lid.

1. Unlock the battery compartment lid by turning the lock as shown below.



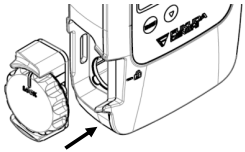


2. Remove the old battery compartment lid.

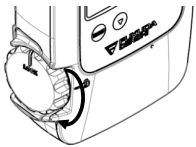


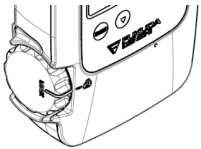
Follow the procedure below to attach the battery compartment lid.

1. Set the new battery compartment lid adjusting to the unit.



2. Lock the battery compartment lid by turning the lock as shown below.





14. Standard and Optional Accessories

This section lists the accessories for the LX-7230N/LX-7230N(G).

⚠WARNING

Use only the accessories, such as ECG Lead cable and SpO_2 probe (sensor) specified by Fukuda Denshi for the LX-7230N/LX-7230N(G). Otherwise, the LX-7230N/LX-7230N(G) cannot deliver its maximum performance and may be damaged, resulting in a safety hazard.

⚠CAUTION

For quality improvement, specifications are subject to change without prior notice.

■ Standard Accessories

No.	*Item*	*Q'ty*	*Note*
1	ECG Lead Cable	1	AHA color code
2	SpO_2 Protective Cap	1	
3	Operation Manual	1	

■ Optional Accessories

The following optional accessories are available for the LX-7230N/LX-7230N(G).
Purchase them as required.

●ECG, Impedance Respiration Measurement

Item	*Model Type*	*Note*
ECG Clip Type Lead Cable	CMT-01CTH-0.8DA	3-electrode (White, Black, Red), Limb Lead (1CH)
ECG Clip Type Lead Cable	CMT-02CTH-0.8DA	4-electrode (White, Black, Red, Green), Limb Lead (2CH)
ECG Clip Type Lead Cable	CMT-03CTH-0.8DA	5-electrode (White, Black, Red, Green, Brown), Limb Lead (1CH)+Chest (1CH)
ECG Hook Type Lead Cable	CMT-01HTH-0.8DA	3-electrode (White, Black, Red), Limb Lead (1CH)
ECG Hook Type Lead Cable	CMT-02HTH-0.8DA	4-electrode (White, Black, Red, Green), Limb Lead (2CH)
ECG Hook Type Lead Cable	CMT-03HTH-0.8DA	5-electrode (White, Black, Red, Green, Brown), Limb Lead (1CH)+Chest (1CH)

●SpO_2 Measurement

Item	*Model Type*	*Note*
Durasensor®	DS-100A	Reusable Adult (weight of 40 kg and over) Finger
OxiMax®	MAX-P	Single-Patient-Use Pediatric (weight of 10 kg to 50 kg) Finger
OxiMax®	MAX-A/AL	Single-Patient-Use Adult (weight of 30 kg and over) Finger
OxiMax®	MAX-I	Single-Patient-Use Infant (weight of 3 kg to 20 kg) Toe
OxiMax®	MAX-R	Single-Patient-Use Adult (weight of 50 kg and over) Nose
OxiMax®	MAX-N	Single-Patient-Use Adult (weight of 40 kg and over) Finger
		Neonate (weight of less than 3 kg) Foot
OxiMax®	MAX-FAST	Single-Patient-Use Adult /Pediatric (weight of 10 kg and over) Forehead

●Others

Item	*Model Type*	*Note*
Battery case lid	OAT-7230A	
SpO_2 Protective cap	-	Ten pieces

15. Specification

■ Specification

⚠CAUTION

For quality improvement, specifications are subject to change without prior notice.

Standard Specification

Size:	76.0 (W) mm x 115.0 (H) mm x 31.8 (D) mm (not including the protrusion)
Weight:	Approximately 200 g (with batteries)
Transmitting Waveform:	ECG 1CH or 2CH (selectable from the ECG lead cable), Respiration waveform, pulse waveform (with SpO_2 value)
ECG Lead cable Type:	3-electrode, 4-electrode, or 5-electrode (Limb+Chest) lead cable Automatically detects the type after inserting the lead cable
Transmitting Status Data:	Electrode Off, Low Battery, Event Switch, Pacemaker Detection, Channel ID, 64 group codes, SpO_2 Sensor Off
LCD:	Built-in
Waterproof:	IPX8 (If periodic replacements are performed) / IPX5 IPX5: Protection from water IPX8: Protection from submerge
Power Supply:	DC: Two 1.5 V “AA” size (“LR06” size) alkaline batteries
LX-7230N Continuous Operating Time:	Approximately 2.5 days
LX-7230N(G) Continuous Operating Time:	Approximately 1.5 days

*Continuous operating time is assumed when using new “AA” size (“LR06” size) alkaline batteries specified by Fukuda Denshi.

ECG

Numbers of Lead Electrode:	3-electrode, 4-electrode, or 5-electrode (Limb+Chest)
Numbers of Input Channel:	1CH (3-electrode) or 2CH
Accuracy of Sensitivity:	10 mm / 1 mV ± 20% (Display sensitivity on the receiving monitor)
ECG Input Impedance:	5 MΩ and above
Input dynamic range and Offset voltage:	Input dynamic range: ±5 mV Offset voltage: ±300 mV Change of amplitude caused by offset voltage: Within ±10%
Common Mode Rejection Ratio:	Less than 10 mVp-p (95 dB and above)
Accuracy of Heart Rate Measurement:	±10% or ±5 bpm, whichever is greater
HR Display Range:	0, 12 bpm to 300 bpm (1 bpm step)
Frequency Characteristic:	0.5 Hz to 40 Hz (within -3dB)
Time Constant	0.4 sec ± 25% Can be switched to 0.1 sec ± 25%.
Rejection of Pacemaker Pulse:	a) Pacemaker Pulse without Over/Undershoot Capable to reject pulses of pulse width 0.1 ms to 2 ms, amplitude ±2 mV to ±700 mV b) Pacemaker Pulse with Over/Undershoot Rejection is not possible.
Protection to Defibrillation:	Complies with IEC 60601-2-27
Lead-off Detection Current:	100 nA and below

Respiration (Impedance Method)

Accuracy of Sensitivity:	10 mm / 1Ω ± 2 mm (When standard Impedance is 480Ω.)
Resp. Display Range:	0, 9 Bpm to 150 Bpm
Display Error of Respiration Rate:	±3 Bpm
Measured Current of Respiration:	Below 100μA (at 42 kHz)

SpO_2

SpO_2 Measurement Range: 1%SpO_2 to 100%SpO_2
Resolution: 1%SpO_2
Measurement Accuracy: Accuracy of measurement with SpO_2 probe is as follows.

SpO_2 Probe	Measurement Accuracy (±1SD)
OxiMax® MAX-I	±2%SpO_2
OxiMax® MAX-P	±2%SpO_2
OxiMax® MAX-A/AL	±2%SpO_2
Durasensor® DS-100A	±3%SpO_2
OxiMax® MAX-R	±3.5%SpO_2
OxiMax® MAX-FAST	±2%SpO_2
OxiMax® MAX-N	±2%SpO_2

(When SpO_2 is 70% to 100%. Less than 70% is not specified.)
The measurement accuracy is determined by measuring on 11 healthy subjects with 5 types of target oxygen saturation controlling the inspired oxygen to maintain steady state, and comparing with the CO-oximeter measurement value.

Wavelength: Approx. 660 nm (Red light)
Approx. 900 nm (Infrared light)
Measurement Response Time: 6 sec. to 7 sec. (Averaging time)
Measurement Value Update Rate: 1 second
Optical Output Power: Under 15 mW

NOTE

The SpO_2 measurement accuracy is determined based on the values of the root-mean-square (rms) difference between SpO_2 readings of the pulse oximeter equipment and values of SaO_2 determined with a CO-oximeter, by healthy adult volunteers. The pulse oximeter equipment measurements are statistically distributed; ±2% measurement accuracy means that only about two-thirds of pulse oximeter equipment measurements can be expected to fall within ±2% of the value measured by a CO-oximeter.

US Patents

Covered by one or more of the following U.S. patents and foreign equivalents:

5,485,847; 5,676,141; 5,743,263; 6,035,223; 6,226,539; 6,411,833; 6,463,310; 6,591,123; 6,708,049; 7,016,715; 7,039,538; 7,120,479; 7,120,480; 7,142,142; 7,162,288; 7,190,985; 7,194,293, 7,209,774; 7,212,847; 7,400,919.

Pulse Wave

Pulse Rate Measurement Range:	20 bpm to 250 bpm
Measurement Accuracy:	±3 bpm
Measurement Response Time:	6 sec. to 7 sec. (Averaging time)
Measurement Value Update Rate:	1 second

Transmission Method of LX-7230N

Modulation Mode:	Digital, Frequency shift keying
Frequency:	608 MHz to 614 MHz
Oscillation Method:	PLL synthesizer method by crystal control
Channel Spacing:	12.5 kHz
Occupied Frequency Bandwidth:	8.5 kHz
RF output power:	1 mW ± 2 dB
Transmitting Antenna:	ECG lead cable and/or SpO_2 Probe

Transmission Method of LX-7230N(G)

Modulation Mode:	Digital, Frequency shift keying
Frequency:	1395 MHz to 1400 MHz, 1427 MHz to 1432 MHz
Oscillation Method:	PLL Synthesizer method by crystal control
Channel Spacing:	25.0 kHz
Occupied Frequency Bandwidth:	16 kHz
RF Output Power:	5 mW ± 2 dB
Transmitting Antenna:	Dielectric Antenna

Safety

General Standard:	ANSI / AAMI ES 60601-1: 2005(R)2012 and A1:2012, C1:2009(r)2012 and A2: 2010(r)2012 (Medical electrical equipment - Part 1: General requirements for basic safety and essential performance)
EMC Standard:	IEC 60601-1-2: 2007 (Medical electrical equipment - Part 1-2: General requirements for basic safety and essential performance - Collateral standard: Electromagnetic compatibility - Requirements and tests)
Operation Mode	Continuous Operating Equipment
The class of protection against electric shock:	Internally Powered Equipment
The type of protection against electric shock:	ECG/RESP: Type CF Applied Part SpO_2: Type CF Applied Part

Operating Environment

Temperature:	10°C to 40°C / 50°F to 104°F
Humidity:	30%RH to 85%RH (non-condensing)
Atmospheric Pressure:	70 kPa to 106 kPa
Vibration/Shock:	Complies with EN ISO 80601-2-61:2011 201.15.3.5.101.1 b), EN ISO 80601-2-61:2011 201.15.3.5.101.1 a)1)

Transport / Storage Environment

Temperature:	-10°C to 60°C / 14°F to 140°F
Humidity:	10%RH to 95%RH (40°C / 104°F, non-condensing)
Atmospheric Pressure:	70 kPa to 106 kPa

■ Functional Testers and Patient Simulator for SpO_2 measurement

Some models of commercially available bench top functional testers and patient simulators can be used to verify the proper functionality of pulse oximeter sensors, cables and pulse oximeters. See the operator's manual of the individual testing device for the procedure which is specific to the model of tester being used.

While such devices may be useful for verifying that the pulse oximeter sensor, cabling, and pulse oximeter are functional, they are incapable of providing the data required to properly evaluate the accuracy of a system's SpO_2 measurements.

Fully evaluating the accuracy of the SpO_2 measurements requires, at a minimum, accommodating the wavelength characteristics of the sensor and reproducing the complex optical interaction of the sensor and the patient's tissue. These capabilities are beyond the scope of known bench top testers.

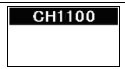
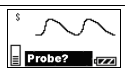
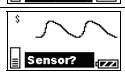
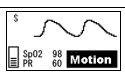
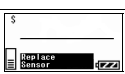
SpO_2 measurement accuracy can only be evaluated in vivo by comparing pulse oximeter readings with values traceable to SaO_2 measurements obtained from simultaneously sampled arterial blood using a laboratory Co-oximeter.

Many functional testers and patient simulators have been designed to interface with the pulse oximeter's expected calibration curves and may be suitable for use with pulse oximeters and/or sensors. Not all such devices, however, are adapted for use with the calibration system. While this will not affect use of the simulator for verifying system functionality, displayed SpO_2 measurement values may differ from the setting of the test device.

For a properly functioning pulse oximeter, this difference will be reproducible over time and from pulse oximeter to pulse oximeter within the performance specifications of the test device.

■ Displays

The following displays are shown on the LCD of the LX-7230N/LX-7230N(G).

	Display	Description
Starting Screen		Displayed after turning on the power. Automatically moves onto the channel display screen.
Channel Display Screen		Displays the transmitter channel after turning on the power and also when refreshing the screen. Automatically moves onto the waveform display screen.
EVENT		Displayed when the EVENT button is pressed. Automatically moves onto the channel display screen.
ELECTRODE?		Displayed when the ECG electrode is disconnected or the ECG/respiration waveform cannot be measured normally. For details about electrode check message, refer to “15. Specification ■ Details of the “ELECTRODE?” Message”.
SpO_2 Measurement Status		
Probe?		Displayed when the SpO_2 probe is disconnected from the equipment.
Sensor?		Displayed when the SpO_2 probe is off from the measuring position or SpO_2 cannot be measured normally due to outside light, etc.
Motion		Displayed when the measurement cannot be executed due to an artifact such as body motion.
Replace Sensor		Displayed when it is necessary to replace the sensor.

		Display	Description
Error Message			
	Telemeter Error		Displayed when the transmitter is faulty
	EEPROM Error		Displayed when the EERPROM is faulty
	CPU Error		Displayed when the CPU is faulty

■ Details of the “ELECTRODE?” Message

The following “ELECTRODE?” messages are displayed on the LCD depending on the selected lead cable and lead.

Check Position	3-electrode lead cable Lead I display	3-electrode lead cable Lead II display	3-electrode lead cable Lead III display
LL			
RA			
LA			
Several Position Simultaneously			

Check Position	4-electrode lead cable	For 5-electrode (Chest) lead cable
LL		
RA		
LA		
RL		
V		
Several Position Simultaneously		

■ List of Setup Items

This section lists the available selection, default setting, and backup status for each setup item, which is available for the LX-7230N/LX-7230N(G).

Items	Selection	Default	Backup
ECG Lead	I, II, III	II	Yes
Display Size of ECG (1)	×1, ×1/2	×1	Yes
Display Size of ECG (2)	×1, ×1/2	×1	Yes
Display Size of Respiration Waveform	×1, ×1/2	×1	Yes

- For details of the following settings, contact our service representative.

Items		Selection	Default	Backup
ECG				
	Time Constant	0.4 s, 0.1 s	0.4 s	Yes
	Pace Detect.	L, M, H	M	Yes
	QRS Width	Narrow, Wide	Wide	Yes
RESP				
	Respiration Detection Signal (Display)	ON, OFF	ON	Yes
Display				
	Contrast	8 steps	8	Yes
Channel				
	LX-7230N Channel	One from the following channels. 0801 to 0879 0900 to 0979 1000 to 1079 1100 to 1179 1200 to 1279 1300 to 1379	1100	Yes
	LX-7230N(G) Channel	One from the following channels. 9501 to 9539 9600 to 9639 9700 to 9739 9800 to 9839 9900 to 9938 2701 to 2739 2800 to 2839 2900 to 2918 2921 to 2939 3000 to 3039 3100 to 3118	9501	Yes
	Group	One from 00 to 63	00	Yes

15. Specification

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