Instructions for Use - English

NoninConnect™ Model 3230 Bluetooth® Smart Pulse Oximeter

 R_{XOnly} (ϵ 0123

Installing AAA Batteries

WARNING: Before changing off and is not applied to a digit.

1. Hold the 3230 so you see the back of the device and the arrows on the battery door point away from you.





3. Slide the battery door awa from you and off the 3230



- 4. If applicable, remove the old batteries from the 3230. Properly dispose of the batteries.
- 5. Insert two new 1.5 volt AAA size batteries. Carefully match the polarity markings (+ and -). The 3230 will not work if the batteries are inserted the wrong way.





6. Carefully slide the battery door



Turning On the NoninConnect

. Insert a digit into the Model 3230 until it touches the built-in stop.

Model 3230





NOTE: Make sure the finger is centered within the finger guide and flat (not on its side). For best results. keep the device at heart or chest

- 2. If the CorrectCheck screen (see Display Symbols table) displays, slide finger further into device. Correct positioning of the finger is critical for accurate measurements.
- 3. The 3230 begins sensing the pulse and displaying readings.



4. View about 4 seconds of readings before relying on the displayed values. It is common for the displayed values to vary slightly over a period of several seconds

NOTE: While on the finger, do not press the device against any surface and do not squeeze or hold it together. The internal spring provides the correct pressure: additional pressure may cause inaccurate readings.

Indications for Use

The NoninConnect Model 3230 Finger Pulse Oximeter is a small, lightweight, portable device indicated for use in measuring and displaying functional oxygen saturation of arterial hemoglobin (% SOO_2) and pulse rate of patients who are well or poorly perfused. It is intended for spot-checking of adult and pediatric patients with digits between 0.8 - 2.5 cm (0.3 - 1.0 inch) thick.

NOTE: Use Environment—Home healthcare environments under the supervision of qualified medical professionals. Users include current/potential users of pulse oximetry in the home and caregivers/ potential caregivers of such a user.

Warnings

Do not use the device in an MR environment, in an explosive atmosphere, or on neonatal patients. This device is not defibrillation proof per IEC 60601-1.

- Use the Model 3230 within its designated range (approximately 10 m/32 ft, spherical radius, line of sight
 when connected to a Bhotooth Smart Ready device). Moving outside this range may cause missing, lost, and/or inaccurate data.
- Inspect the sensor application site at least every 4 hours to ensure correct sensor alignment and skin integrity. Patient sensitivity to sensor may vary due to medical status or skin condition.
- oid excessive pressure to the sensor application site as this may cause damage to the skin beneath the sensor This device is intended only as an adjunct in patient assessment. It must be used in conjunction with
- other methods of assessing clinical signs and symptoms.

 The device must be able to measure the pulse properly to obtain an accurate SpO₂ measurement. Verify
- The cucked miss is bindered in the pulse measurement before relying on the SpO₂ measurement. Vernly that nothing is bindered by the pulse measurement before relying on the SpO₂ measurement.

 Operation of the SpO₂ measurement.

 Operation of the device below the minimum amplitude of 0.3% on the spO₂ measurement.

 Solvent of the device minimum amplitude of 0.3% on the spO₂ measurement.
- Keep the oximeter away from young children. Small items such as the battery door and battery are
- Before changing batteries, make sure the device is off and is not applied to a digit.

- This device has no audible alarms and is intended only for spot-checking.
 This device is designed to determine the percentage of arterial oxygen saturation of functional hemoglobin. Factors that may degrade pulse oximeter performance or affect the accuracy of the measurement include the following:
- measurement include the tollowing:

 applying the pulse someter on improperly applied device the same arm as a blood finger is outside recommended size range size range anemia or low hemogo concentrations carboxyhemoglobin
- or infusion line(s) (IVs)

 excessive light, such as sunlight
 venous pulsations
 or direct home lighting
 cardiogreen and other
- excessive motion intravascular dves · moisture in the device
- methemoglobin
 - dysfunctional hemoglobin
 artificial nails or fingernail polish
 residue (e.g., dried blood, dirt,
 - grease, oil) in the light path
- · The device may not work when circulation is reduced. Warm or rub the finger, or re-position the device. The device is designed to be attached only to a digit.
 This device's display will shut off after 30 seconds of no readings or poor readings.
- In some circumstances, the device will interpret motion as good pulse quality. Minimize patient motion
- as much as possible.

 Clean the device before applying it to a new patient.
- Do not sterilize, autoclave, or immerse this device in liquid. Do not pour or spray any liquids into the device
- Do not use caustic or abrasive cleaning agents, or any cleaning products containing ammonium chloride or isopropyl alcohol.
- Do not use cleaning solutions other than those recommended here, as permanent damage could result. This device is a precision electronic instrument and must be repaired by Nonin Technical Service. Field
- Insucvice is a precision receitable institutent and must be reparted by whom te cause covered reactive repair of the device is not possible. Do not attempt to open the case or repair the electronics. Opening the case may damage the device and void the warranty.

 This equipment complies with IEC (6001-1-2 for electromagnetic compatibility for medical electrical equipment and/or systems. This standard is designed to provide reasonable protection against harmful interference in a typical medical installation. However, because of the proliferation of radio-frequency transmitting equipment and other sources of electrical noise in health care and other environments, it is possible that high levels of such interference due to close proximity or strength of a source might disrupt the performance of this device. Medical electrical equipment needs special precautions regarding EMC, and all
- equipment must be installed and put into service according to the EMC information specified in this manual

 Portable and mobile RF communications equipment including CT, diathermy, RFID, and electronic article security systems can affect medical electrical equipment.

 Batteries may leak or explode if used or disposed of improperly. Remove batteries if the device will be stored
- for more than 30 days. Do not use different types of batteries at the same time. Do not mix fully charged and
- partially charged batteries at the same time. These actions may cause the batteries to leak.
 Follow local, state, and national governing ordinances and recycling instructions regarding disposal or recycling of the device and device components, including batteries.

 In compliance with the European Directive on Waste Electrical and Electronic Equipment (WEEE)
- 2002/6/EC, do not dispose of this product as unsorted municipal waste. This device contains WEEE materials; please contact your distributor regarding take-back or recycling of the device. If you are unsure how to reach your distributor, please call Nonin for your distributor's contact information.

Symbols Symbol

,			.,
Δ	Caution!		4
③	Follow Instructions for Use.		+
(8)	MR unsafe		
*	Type BF Applied Part (patient isolation from electrical shock)		IP
	UL Mark for Canada and the United		
, The second	States with respect to electric shock, fire, and mechanical hazards only in		S
1(1)11	accordance with IEC 60601-1, UL 60601-1 and CAN/CSA-C22.2 No. 601.1.		В
(€0123	CE Marking indicating conformance to EC Directive No. 93/42/EEC concerning medical devices.		_/
	Non-ionizing electromagnetic		
((<u>@</u>))	radiation. Equipment includes RF transmitters. Interference may occur in		- 7
	the vicinity of equipment marked with		
	this symbol.		1
{	Date of Manufacture		R _x
	Continua Certified TM signifies this device meets Continua certification		22.
4	testing requirements, which support		œ
•	interoperability in personal health devices (continuaalliance.org).		-
Ħ	Indicates separate collection for electrical and electronic equipment		EC
-	(WEEE)		Ri

2	Not for continuous monitoring (no alarm for SpO ₂)		
(+ 1)	Battery orientation		
IP32	Protected against vertically falling wat drops when enclosure is tilted up to 15 degrees and ingress of solid foreigr objects greater than or equal to 2.5 m (0.1 in.) in diameter per IEC 60529.		
SN	Serial Number		
BDA	Bluetooth Device Address		
1	Storage/shipping temperature range		
Ī	Handle with care		
*	Keep dry		
	Indoor use (France only)		
$R_{\!\!X\!\text{Only}}$	Federal law (USA) restricts this device to sale by or on the order of a licensed practitioner.		
COMP ¹	Country of Manufacture		
ш	Manufacturer		
EC REP	Authorized Representative in the European Community		
REF	Catalogue number		
QTY	Quantity		

Definition

NOTE: Where applicable, an additional label bearing your country radio communications license information will appear on the side of your device. This is not a serial number or device identifier.

Symbol	Description			
	Nonin's CorrectCheck TM senses that the finger has not been correctly inserted. If yo see this symbol, slide finger further into device.			
% Sp0 ₂	The number next to this symbol is the amount of oxygen in your blood (functional oxygen saturation of arterial hemoglobin).			
	The number next to this animated symbol is your pulse rate. Pulse rate is the number times your heart beats per minute.			
	Dashes replace the readings when the 3230 is unable to detect a usable signal.			
*	White symbol – Radio is on. Green symbol – 3230 is connected. Flashing symbol – Connection error. The radio will reset. No symbol – Radio is off.			
0	Poor signal. Steady your hand, reposition finger, warm finger by rubbing, or select a different finger.			
	Low battery. Replace batteries.			
	Critical battery. Flashing indicator on full screen. The device will not work until the batteries are replaced.			
Ø	†Spot-check complete. While Spot-check is in progress, a clockwise spinning circulation displays.			
\subseteq	† Measurement complete (full screen).			

Using the NoninConnect Model 3230

Installing AAA Batteries

Use only alkaline batteries. When batteries are low, alkaline batteries as soon as possible. See the "Installing AAA Batteries" instructions and figures at left.

Turning On the NoninConnect Model 3230

See the "Turning on the NoninConnect Model 3230" instructions and figures at left.

Connection via Bluetooth Wireless Technology

When the Model 3230 is placed on the finger and turns on, it is ready for a Bluete The 3230 stays in this mode until it is shut off or the Bluetooth radio turns off. The 3 symbol is white when the Bluetooth radio is on, green when the 3230 is connected, and flashes when communication error

The Bluetooth symbol is useful for the product installer.

Due to the wide variety of wireless environments, the Bluetooth connection between the 3230 and the host device must be tested before using the 3230's Bluetooth capabilities.

Turning Off the NoninConnect Model 3230

The Model 3230 will automatically turn off approximately 10 seconds after the digit is removed, or after a 2-minute period of poor signals

Cleaning the NoninConnect Model 3230

CAUTIONS:

- Clean the device before applying it to a new patient.
 Do not sterilize, autoclave, or immerse this device in liquid. Do not pour or spray any liquids into the device.
- . Do not use caustic or abrasive cleaning agents, or any cleaning products containing ammonium chloride or isopropyl alcohol.
- Do not use cleaning solutions other than those recommended here, as permanent damage could result.
- 1. To clean, wipe the device's surfaces with a soft cloth dampened with one of the following:
- A 10% bleach solution (household bleach [5.25% sodium hypochlorite]) Warm, soapy water (hand dishwashing detergent – see note below), and then rinse the cleaned surfaces with a soft cloth dampened with water (home use only).
- 2. Dry with a soft cloth, or allow to air dry. Ensure that all surfaces are completely dry

NOTE: The hand dishwashing detergent that was tested includes these ingredients: Sodium Lauryl Sulfate, Sodium Laureth Sulfate, Lauramine Oxide, Sodium Chloride, PPG-26, PEG-8 Propylhepty Ether, and Phenoxyethanol.

NONIN MEDICAL, INCORPORATED, (Nonin) warrants to the purchaser, for a period of 2 years from the date of purchase, each Model 3230 exclusive of the batteries and spring. The device's expected

Nonin shall repair or replace any 3230 found to be defective in accordance with this warranty, free of charge, for which Nonin has been notified by the purchaser by serial number that there is a defect, provided notification occurs within the applicable warranty period. If unable to repair, Nonin shall replace with a 3230 or a comparable device. This warranty shall be the sole and exclusive remedy by the purchaser hereunder for any 3230 delivered to the purchaser which is found to be defective in any manner whether such remedies be in contract, tort or by law

This warranty excludes cost of delivery to and from Nonin. All repaired units shall be received by the purchaser at Nonin's place of business. Nonin reserves the right to charge a fee for a warranty repair request on any 3230 found to be within specifications.

Model 3230 is a precision electronic instrument and must be repaired by trained Nonin personnel only. Any sign or evidence of opening the 3230, field service by non-Nonin personnel, tampening, or any kind of misuse of the 3230, shall void the warranty. All non-warranty work shall be done at Nonin's standard rates and charges in effect at the time of delivery to Nonin.

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nonin.com

Specifications

regulatory@nonin.com for accuracy data.

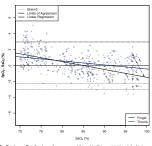
Oxygen Saturation Display Range: 0% to 100% SpO₂

Pulse Rate Display Range: 18 to 321 beats per minute (BPM) The table below shows A_{rms} values measured using the Model 3230 in a Declared Accuracy*

NOTE: If your national regulatory authority recognizes accuracy in motion, please contact

Accuracy Summary - Finger and Thumb

Range	Specified Oxygen Saturation (A _{rms})	Finger Oxygen Saturation (A _{rms})	Thumb Oxygen Saturation (A _{rms})
70 – 100%	± 2	± 1.31	± 1.56
70 – 80%	± 2	± 1.65	± 1.91
80 - 90%	± 2	± 1.05	± 1.21
90 - 100%	± 2	± 1.18	± 1.49



This graph shows plots of the error (SpO₂ – SaO₂) by SaO₂ using the 3230 with a linear regression fit and upper 95% and lower 95% limits of agreement. Each sample data point is identified by subject from a clinical study in non-motion conditions

SpO₂ Low Perfusion Accuracy (A_{rms})*: 70 to 100% ± 2 digits Pulse Rate Declared Accuracy Range (A_{rms})*: 20 to 250 BPM ± 3 digits Low Perfusion Pulse Rate Declared Accuracy Range (A_{rms})*: 40 to 240 BPM ± 3 digits

 Measurement Wavelengths and Output Power**:

 Red:
 660 nanometers @ 0.8 mW max. average Infrared:

 910 nanometers @ 1.2 mW max. average

Temperature: -5 °C to 40 °C / 23 °F to 104 °F Operating: Storage/Transportation -40 °C to 70 °C / -40 °E to 158 °E

Time (from storage) for device to be ready for its intended use:

3 minutes to warm from -40 °C to -5 °C 8 minutes to cool from 70 °C to 40 °C Humidity

Operating: Storage/Transportation: 10% to 95% non-condensing 10% to 95% non-condensing Altitude: Up to 4,000 meters / 13,123 feet

Operating: Hyperbaric Pressure Battery Life:

Approximately 2,200 spot checks (25 sec. per spot-check), within 10 meters/32 feet of collector with streaming data 1 month, with batteries installed. CAUTION: Remove batteries if the device will be stored for more than 30 days.

* ±1 A.... represents approximately 68% of measurements at zero bias.

** This information is especially useful for clinicians performing photodynamic therapy.

Bluetooth Wireless Technology Information

Bluetooth Compliance: Version 4.0 single mode low energy 2.4 to 2.4835 GHz Operating Frequency: Output Power TX: +3 dBM 10 meter radius (line of sight) Operating Range:

Network Topol Star - bus Operation: Model 3230

Antenna Type Integrated chip type antenna Modulation Type: Frequency Hopping Spread Spectrum 1 Mbit/second

Data Rate Data Latency 6 ms Data Integrity Adaptive Frequency Hopping 24-bit CRC (cyclic redundancy check)

32-bit message integrity check Nonin Proprietary: Sends data packets once per second. Includes a second counter that allows the host to detect if packets are missing and the device Data Format:

Bluetoath SIG Standard: Compliant with Bluetoath SIG Pulse Oximeter

Profile specifications adopted by Continua. This device uses Bluetooth Smart technology for wireless communications, which allows for reliable communications in electrically noisy

environments, and transmits physiological data. If the connection is lost, the device will become available for a connection in a few second Bluetooth Profiles Supported: GATT-based Nonin Proprietary Oximeter Profile; GATT-based Bluetooth

SIG Pulse Oximeter Profile Authentication and Encryption: Supported
Encryption Key Size: 128 bits AES (advanced encryption standard)

Bluetooth Security

Quality of Service:

The Bluetooth radio contained in the 3230 is a Bluetooth Smart single-mode, low-energy radio. The 3230 supports an encryption key size of 128 bits. While the 3230 is in a Bluetooth connection, it will be unavailable for other connections. Apart from the standard Bluetooth security measures, Nonin has two non-standard security measures that are available

For additional technical information, please see the insert, "NoninConnect Model 3230 Technical