



Operator's Manual

Models 8500 & 8500M
Handheld Pulse Oximeters

English

About the Manual

There are many contraindications, warnings, and cautions throughout this manual. Read them carefully; they are important to the use of the product.

The information in this manual has been carefully checked and is believed to be accurate. In the interest of continued product development, NONIN reserves the right to make changes and improvements to this manual and the products it describes at any time, without notice or obligation.

**CAUTION! Federal law (USA) restricts this device to sale by or on the order of a physician.
CAUTION! Read this manually carefully before using the Model 8500.**

Nonin Medical, Inc.
13700 1st Avenue North
Plymouth, MN 55441-5443 USA

(763) 553-9968
(800) 356-8874 (USA and Canada)
Fax (763) 553-7807
E-mail: info@nonin.com
www.nonin.com

Authorized EC Representative:
MPS, Medical Product Service GmbH
Borngasse 20
D-35619 Braunfels, Germany



References to “NONIN” in this manual shall imply Nonin Medical, Inc.

NONIN, nVISION, Flexi-Form, and FlexiWrap are registered trademarks or trademarks of Nonin Medical, Inc.

References to “8500” in this manual imply Models 8500 and 8500M.

© 2005 Nonin Medical, Inc.

Table of Contents

Precautions for Use 1
 Unpacking the Model 8500 4
 Introduction..... 4
Basic Operation..... 5
 Installing Batteries 5
 Connecting Sensors 6
Features 9
 Controls..... 9
 Displays 14
 Printer/Serial Output 15
 Memory Option (8500M Only) 16
Specifications..... 17
Service 18
Warranty..... 18
Accessories 19
Troubleshooting Guide 20

Guide to Symbols



Attention: See instructions for use.



Type BF Applied Part
(Patient isolation from electric shock)



Not for Continuous Monitoring
(No alarm for SpO₂)



UL Mark for Canada and the United States with respect to electric shock, fire, and mechanical hazards only in accordance with UL 60601-1 30EM and CAN/CSA C22.2 No. 601.1



CE Marking indicating conformance to EC directive No. 93/42/EEC concerning medical devices



Indicates separate collection for electrical and electronic equipment (WEEE).

Precautions for Use

Contraindications

- Do not use the 8500 in an MRI environment.
- Do not use the 8500 in a situation where alarms are required. The 8500 has no alarms.

Warnings

- Explosion Hazard. Do not use the 8500 in an explosive atmosphere.
- The 8500 is intended only as an adjunct in patient assessment. It must be used in conjunction with other methods of assessing clinical signs and symptoms.
- Use only NONIN-manufactured pulse oximeter sensors. These sensors are manufactured to meet the accuracy specifications for NONIN Pulse Oximeters. Using other manufacturers' sensors may cause improper pulse oximeter performance.
- As with all medical devices, carefully route patient cabling to reduce the possibility of patient entanglement or strangulation.
- Check the application site *frequently* to determine the positioning of the sensor and the circulation and skin sensitivity of the patient. Each patient's sensitivity to NONIN sensors might vary, depending upon medical status or skin condition.
- Discontinue use of adhesive tape strips if the patient exhibits an allergic reaction to the adhesive material.
- Do not stretch the adhesive tape while applying the sensor. This may cause inaccurate readings or skin blisters.
- General operation of the 8500 might be affected by the presence of an electrosurgical unit (ESU).
- Do not use a damaged sensor.
- This device should not be used adjacent to or stacked with other equipment. If adjacent or stacked use is necessary, the device should be observed carefully to verify normal operation.
- The use of accessories, sensors, and cables other than those listed in this manual may result in increased emission and/or decreased immunity of this device.

Cautions

- Federal law (USA) restricts this device to sale by or on the order of a physician.
- Read this manual carefully before using the 8500.
- Before use, carefully read the package insert provided with the sensor(s).
- The 8500 is intended for spot-checking or monitoring by a health care professional. Because the 8500 has no audible alarms, international labeling requirements dictate that it be labeled "Not for continuous monitoring."
- The 8500 pulse oximeter is not an apnea monitor.
- Verify that all visible indicators illuminate during the startup (initialization) sequence. If any indicator is not lit, do not use the 8500. Contact NONIN Customer Support for repair or replacement.
- The 8500 must be able to measure the pulse properly in order to obtain an accurate SpO₂ measurement. Verify that nothing is hindering the pulse measurement before relying on the SpO₂ measurement.
- The 8500 might not work on all patients. If you are unable to achieve stable readings, discontinue use.
- The 8500 might misinterpret motion as good pulse quality. Minimize patient motion as much as possible.
- The 8500 is designed to determine the percentage of arterial oxygen saturation of functional hemoglobin. Significant levels of dysfunctional hemoglobin may affect the accuracy of the measurement.
- Cardiogreen and other intravascular dyes, depending on the concentration, might affect the accuracy of the SpO₂ measurement.
- Ear Clip and Reflectance pulse oximeter sensors are not recommended for pediatric or neonatal use. The accuracy of these sensors has not been established for pediatric or neonatal use.
- Do not immerse the 8500 or NONIN sensors in liquid.
- Do not use caustic or abrasive cleaning agents on the 8500 or on the sensors.
- The 8500 must be repaired only by trained NONIN personnel.
- Do not use different types of batteries or mix fully- and partially-charged batteries at the same time.; battery leakage might result.
- Do not remove any covers (except the battery cover) when replacing batteries.
- Follow local ordinances and recycling instructions regarding disposal or recycling of the device and device components, including batteries. Batteries might leak or explode if used or disposed of improperly.
- Remove the batteries to avoid the risk of leakage if the 8500 will be stored for more than one month.
- This equipment complies with International Standard EN 60601-1-2:2001 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 healthcare 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.
- This device has not been tested for immunity to electromagnetic disturbances.
- Portable and mobile RF communications equipment can affect medical electrical equipment.
- In compliance with the European Directive on Waste Electrical and Electronic Equipment (WEEE) 2002/96/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.

Manufacturer's Declaration

Refer to the following table for specific information regarding this device's compliance to IEC Standard 60601-1-2.

Table 1: Electromagnetic Emissions

Emissions Test	Compliance	Electromagnetic Environment— Guidance
<p><i>This device is intended for use in the electromagnetic environment specified below. The customer and/or user of this device should ensure that it is used in such an environment.</i></p>		
RF Emissions CISPR 11	Group 1	This device 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 Emissions CISPR 11	Class B	This device is suitable for use in all establishments, including domestic and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.
Harmonic Emissions IEC 61000-3-2	N/A	
Voltage Fluctuations/ Flicker Emissions IEC 61000-3-3	N/A	

Unpacking the Model 8500

Confirm that the items listed below are packed with the 8500 Handheld Pulse Oximeter. The 8500 shipment includes:

- 8500 or 8500M Handheld Pulse Oximeter
- Operator's Manual for Models 8500 and 8500M Pulse Oximeter
- Six AA batteries
- One sensor

If any item on this list is missing or damaged, contact your distributor. Contact the carrier immediately if the shipping carton is damaged.

Introduction

Indications for Use

The NONIN® 8500 Handheld Pulse Oximeter is indicated for measuring and displaying functional oxygen saturation of arterial hemoglobin (SpO₂) and pulse rate for adult, pediatric, and neonatal patients in hospital, ambulatory, home, and EMS (emergency medical service) environments. The 8500 is intended for spot-checking and/or continuous monitoring when attended by a trained healthcare professional.

General Description

The Model 8500 is a digital handheld pulse oximeter that displays numeric values for blood oxygen saturation and pulse rate. It typically operates for 100 hours continuously between alkaline battery replacements and requires no routine calibration or maintenance other than battery replacement.

The pulse oximeter determines functional oxygen saturation of arterial hemoglobin (SpO₂) by measuring the absorption of red and infrared light passing through perfused tissue. Changes in absorption caused by the pulsation of blood in the vascular bed are used to determine oxygen saturation and pulse rate.

Oxygen saturation and pulse rate values are displayed by light-emitting diode (LED) digital displays. On each detected pulse, the pulse quality LED blinks. Patient pulse quality signals are graded as good, marginal, or inadequate and are indicated as such by the pulse quality indicator blinking green, yellow, or red, respectively. This simple method gives the user a pulse-by-pulse visual indication of waveform signal quality without requiring complex waveform analyses.

The Model 8500 pulse oximeter may be used with a variety of NONIN pulse oximeter sensors.

Because the Model 8500 has no patient alarms, SpO₂ and pulse rate displays must be observed frequently.

A sensor disconnect or malfunction is indicated by a lack of good pulse quality blinking and/or a dash to the left of the SpO₂ value on the LED display. When adequate pulse signals are not received, the SpO₂ and/or pulse rate numeric values will be replaced by dashes. Low and critically low battery conditions will be indicated by the low battery LED.

Basic Operation

Installing Batteries

CAUTION! Do not use different types of batteries at the same time, and do not mix fully- and partially-charged batteries at the same time. Battery leakage might result.

Do not remove any covers (except the battery cover) when replacing batteries.

Follow local ordinances and instructions regarding disposal or recycling of the device and components, including batteries. Batteries may leak or explode if used or disposed of improperly.

Remove the batteries to avoid the risk of leakage if the Model 8500 will be stored for more than one month.

The 8500 Pulse Oximeter is powered by 6 AA Alkaline cells that will typically provide 100 hours of continuous operation. *The 8500 indicates when the batteries are low by flashing the digital displays once each second.* When the displays begin flashing, the batteries should be replaced as soon as possible. Replace the batteries by removing the battery door on the back of the 8500. Be sure to follow the polarity markings on the rear label of the pulse oximeter when installing new batteries. Refer to Figure I for an illustration of battery replacement.

Rechargeable Nickel Cadmium batteries may be used in the 8500 if desired. Since NiCad batteries have less than half the capacity of alkaline batteries, the batteries will have to be recharged more often than every 100 hours.

NOTE: Reducing the display brightness can extend battery life up to 100%.

NOTE: Replacing batteries erases the memory and clock settings of the 8500.

NOTE: Batteries should be removed if the 8500 is going to be stored for more than 30 days. Batteries may leak if left in the device for a long period of time.

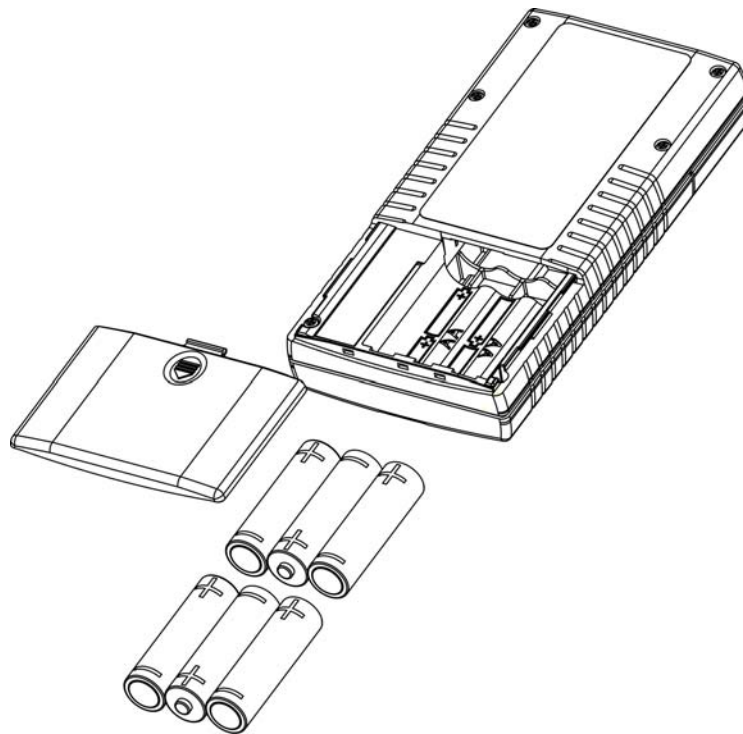


Figure I: Replacing Batteries in the 8500

Connecting Sensors

Connect the sensor to its 9-pin mating jack on the top of the 8500 as shown in Figure II. If additional cable length is necessary, connect the Model 8500I Patient Cable between the sensor and the 8500 Pulse Oximeter. Position the appropriate sensor on the patient.

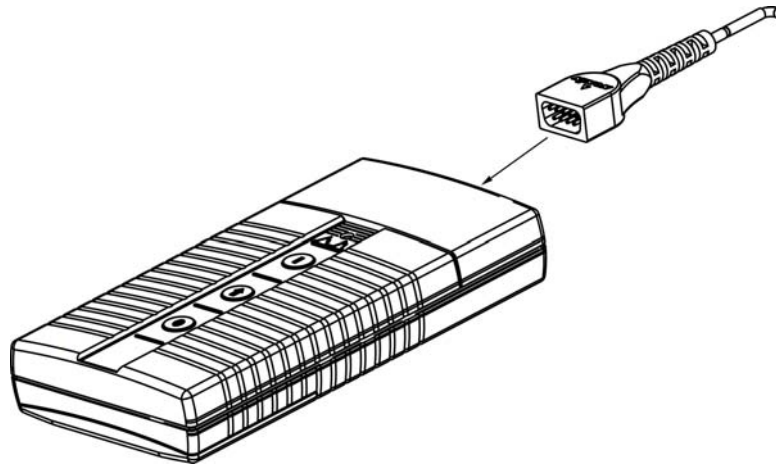


Figure II: Connecting Sensors to the 8500

Turning on the Pulse Oximeter

Turn on the Model 8500 by pressing the " | " button on the front of the pulse oximeter. Refer to Figure III.

When the 8500 is powered on, the SpO₂ and ♥ displays will cycle through the following sequence before displaying valid data values:

- "888 888"
- time saved in memory in hours and minutes
- software revision number
- " _ _ "

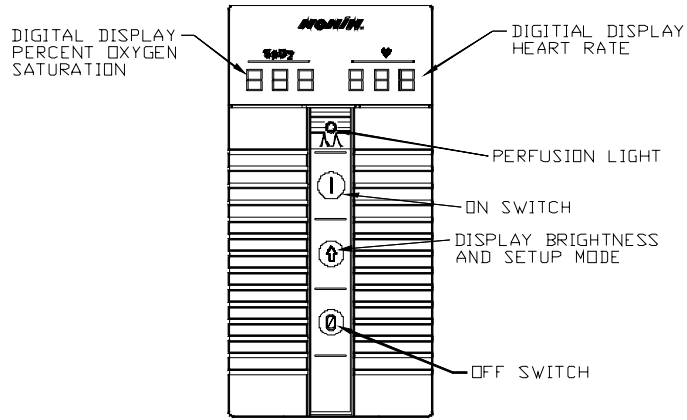


Figure III: Front View of the 8500

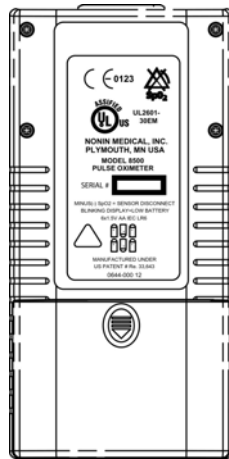



Figure IV: Rear View of the 8500

Verifying Operation

CAUTION! The 8500 must be able to measure the pulse properly to obtain accurate SpO₂ measurement. Verify that nothing is hindering the pulse measurement before relying on the SpO₂ measurement.

Verify that the sensor is properly positioned, and ensure that the system is sensing adequate perfusion by observing that the  (perfusion) indicator is blinking green, and the blinking is correlated to the pulse rate for 10 seconds. Should the perfusion light be red or yellow or flashing erratically, reposition the sensor or try a different sensor.

Cleaning the Pulse Oximeter

CAUTION! Do not immerse the 8500 in liquid, and do not use caustic or abrasive cleaning agents.

The 8500 Pulse Oximeter may be cleaned with a mild detergent and a soft cloth or with an isopropyl alcohol wipe. Allow enough time for the 8500 to dry thoroughly before reusing.

Features

Controls

All functions of the 8500 are controlled by switches found on the front of the unit. Refer to Figure III for an illustration of these switches.

Power

Pressing the ON switch ("|") causes power to be applied to all internal circuitry. Pressing the OFF switch ("Ø") causes power to be removed from the displays and puts the pulse oximetry circuitry into a low power standby mode.

In order to conserve battery life, the 8500 automatically powers off after 10 minutes of inactivity. Inactivity is indicated by dashes on the displays and is caused by:

- no sensor connected to the pulse oximeter
- patient pulse too low
- sensor not attached to a patient

Each time a reading is displayed, the 10 minute timer is restarted.

The "|" switch has additional clock and printer mode setting functions when used in conjunction with the "↑" switch.

Display Brightness

The arrow switch ("↑") causes the brightness of the digital displays to change. When powered up, the digital display defaults to the maximum brightness. Pressing the "↑" will advance the brightness to the lowest setting, and each subsequent press will advance the brightness through 8 different settings. The function is circular, which means it will cycle through the entire brightness range and then start at the beginning again. Lower brightness may be used to preserve battery life; higher brightness may be used to view the displays from a distance.

NOTE: *Reducing the LED display brightness can extend battery life up to 100%.*

The "↑" switch has additional clock mode setting functions when used with the "|" switch.

Setup Mode

Setup mode is used to control the internal time-of-day clock and the external real-time printer (purchased separately). The setup mode is initiated by holding the "↑" switch when the unit is turned on by pressing the "|" switch. In setup mode, the "|" switch and the "↑" switch are used to make the selections.

Advance to the next sequential mode by pressing the "|" switch. Each time the "↑" switch is pressed, the number on the ♥ display will increment. It starts with the current value stored in memory for the parameter designated in the SpO₂ display. When the correct value appears in the ♥ display, pressing the "|" switch will advance the SpO₂ display to the next sequential parameter as listed in Table I. This process is continued until all parameters are set. The settings can be checked easily, since the first value displayed for each parameter represents the current setting. When the setting sequence has been completed, the 8500 exits setup mode and begins normal operation.

Sequence	Appears in SpO ₂ Display	Range of Values	
		From	To
Printer	P r n	00	15
Year	y	00	99
Month	m m	00	12
Day	d	01	31
Hours	h h	00	23
Minutes	m m	00	59

Table I: Printer, Clock, and Calendar Mode Parameters

WARNING: The use of the 8500P may result in increased emission and/or decreased immunity of this device.

Printer Settings

NOTE: The 8500P prints real-time data only. Data stored in the 8500M cannot be downloaded to the 8500P.

"P r n" will appear in the SpO₂ display, indicating print setup mode. There are 16 options for the printer mode: 00 through 15.

The modes determine how often and in which format data is written to the printer. Refer to Figure V for a flowchart of setting the printer mode. When the printer setting sequence is completed, the 8500 continues to the calendar settings.

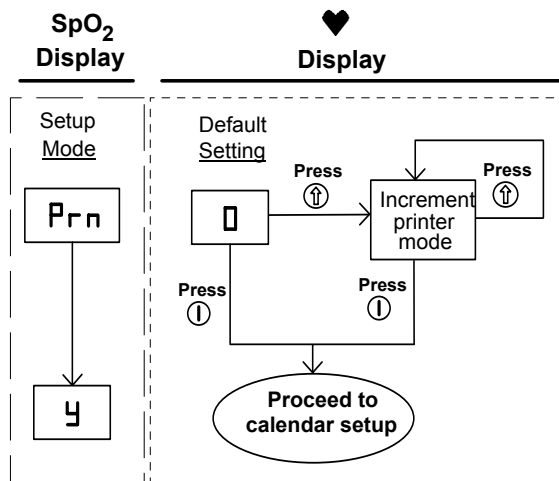


Figure V: Flow Chart for Setting Printer Mode

Printer Mode	Seconds per data point	Minimum SpO₂ printed?*	Touch print capability?***
00	10	No	Yes
01	30	No	Yes
02	120	No	Yes
03	10	Yes	Yes
04	30	Yes	Yes
05	120	Yes	Yes
15	---	No	Yes

* For modes where the minimum SpO₂ data is written, there are two lines of data written for each data output. The first line contains the minimum value for SpO₂ since the last printout, and the second line contains the current data values.

** The touch print mode enables the user to print out data at any time. This is activated by pressing the " | " switch.

Table II: 8500P Printer Modes

Calendar Settings

After the printer setting has been determined in the setup mode, "y" will appear in the SpO₂ display, indicating calendar setup mode for the year. The year may be set to "00" through "99". After selecting the year, the display will show "nn", indicating the setup mode for the month. The month may be set to "00" through "12". After selecting the month, the display will show "d", indicating the setup mode for the day of the month. The day may be set to "01" through "31". Refer to Figure VI for a flowchart of setting the calendar. When the calendar setting sequence is completed, the 8500 continues to the clock settings.

NOTE: Setting the month to "00" disables the clock function and helps conserve battery life.

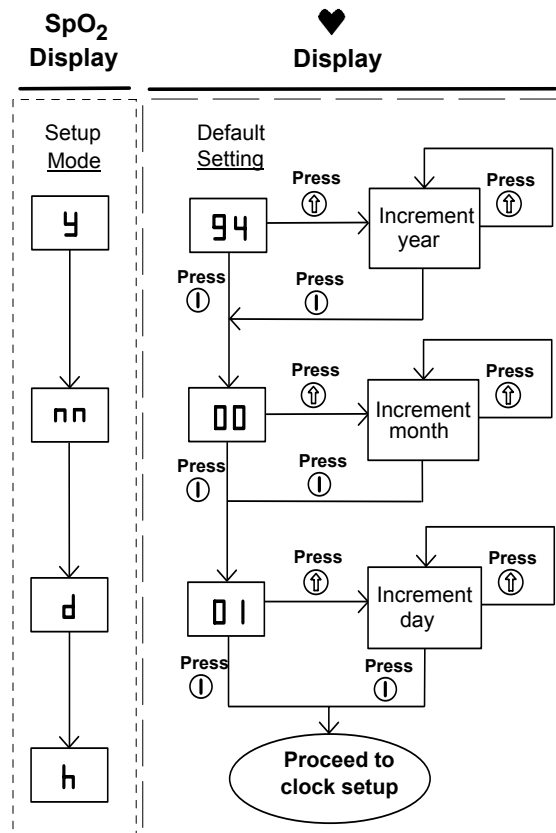


Figure VI: Flow Chart for Setting Calendar

Clock Settings

After the calendar settings have been determined in the setup mode, "h" will appear in the SpO₂ display, indicating clock setup mode for the hour. The hour may be set to "00" through "23". After selecting the hour, the display will show "n n," indicating the setup mode for the minutes. The minutes may be set to "00" through "59". After selecting the minutes, the display will return to normal operation. Refer to Figure VII for a flowchart of setting the clock.

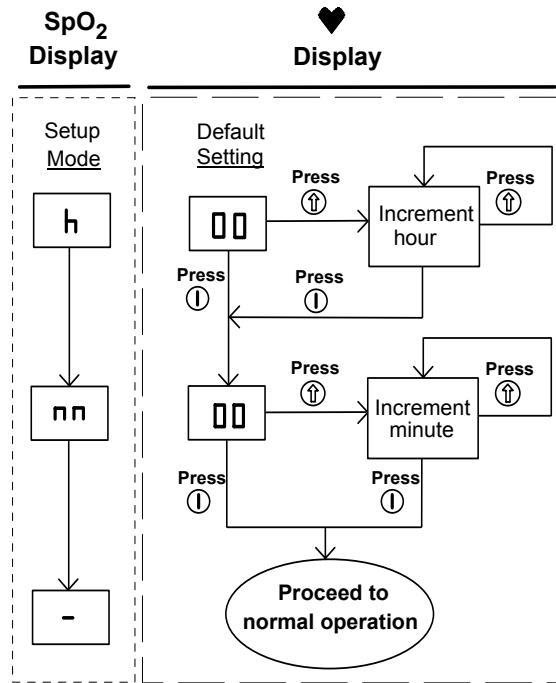


Figure VII: Flow Chart for Setting Clock

Printer modes determine how often and in which format data is written to the printer. For modes where the minimum SpO₂ data is written, two lines of data are written for each data output. The first line contains the minimum value for SpO₂ since the last printout, and the second line contains the current data values. Touch print mode enables users to print data at any time by pressing the "1" switch.

Displays


SpO₂ Display

The left digital display is a 3-digit light emitting diode (LED) digital display that indicates oxygen saturation percentage. Refer to Figure III.

♥ (Pulse Rate) Display

The right digital display is a 3-digit LED digital display that indicates pulse rate in pulses per minute. Refer to Figure III.

(Perfusion) Indicator

The perfusion indicator (identified by the waveform symbol ) will flash once for each pulse while measuring oxygen saturation. The perfusion indicator changes color to indicate changes in the pulse waveform signal that may affect SpO₂ data.

The perfusion indicator may blink one of three colors: green, yellow, or red. These colors are similar to the colors of a stoplight such that:

- **Red** indicates the pulse amplitude is too small. During red perfusion, SpO₂ and pulse rate values are not updated. After ten seconds, values are replaced with dashes, indicating SpO₂ measurement is not possible.
- **Yellow** indicates the pulse waveform amplitude is marginal or the pulse oximeter has detected artifact. Although SpO₂ data is acceptable, corrective measures should be considered to improve sensor placement, change sensor type, or reduce patient movement.
- **Green** indicates a good pulse waveform signal and accurate SpO₂ data.

CAUTION! The 8500 might interpret motion as good perfusion.

Flashing Display

The numerical displays will flash once each second if the 8500 determines that a battery low condition exists. Replace all six batteries immediately.

NOTE: *Inaccurate SpO₂ and/or pulse rate measurement might result if the 8500 is operated in a low battery condition.*

Dash in SpO₂ Display

If the 8500 determines that a sensor fault exists (sensor disconnect, sensor failure, or sensor dislodgment), a dash (-) appears in the left-most digit of the SpO₂ display. The readings displayed will remain unchanged while the sensor fault exists. If the sensor fault is not corrected, dashes will be displayed 10 seconds after the minus sign appears.

Printer/Serial Output

Both the 8500 and 8500M Hand Held Pulse Oximeters provide output capability to a custom printer via the 9-pin Sub-D connector. This connector serves as a sensor input connector as well as a printer interconnect device. The 9-pin Sub-D connector pin assignments are listed in Table III.

<u>Pin Number</u>	<u>Assignment</u>
1	Battery Voltage
2	Infrared Anode, Red Cathode
3	Infrared Cathode, Red Anode
4	Serial Data, TTL Levels
5	Detector Anode
6	Logic Level
7	Cable Shield
8	Coaxial Shield
9	Detector Cathode, +5 V

Table III: Printer/Sensor Interface Assignments

The information from the 8500 in the real-time mode is sent in an ASCII serial format at 9600 baud with 9 data bits, 1 start bit, and 1 stop bit. The data is output at a rate of once per second.

NOTE: *The 9th data bit is used for parity in memory playback mode. In real-time mode, it is always set to the mark condition. Therefore the real-time data may be read as 8 data bits, no parity.*

The data printed by the 8500P printer is in the following format:

HH:MM:SS SPO2=XXX HR=YYY

where "HH" represents the hour, "MM" represents the minutes, "SS" represents the seconds, "XXX" represents the SpO₂ value, and "YYY" represents the heart rate. SpO₂ and heart rate will be displayed as "---" if there is no data available for the data reading.

WARNING: The use of the 8500P may result in increased emission and/or decreased immunity of this device.

Memory Option (8500M Only)

The memory option is identified by the "M" in the model number (i.e. 8500M as opposed to the 8500). This model number is located just above the serial number on the back of the unit. The 8500M Pulse Oximeter can collect and store up to eighteen hours of SpO₂ and pulse rate information.

The memory in the 8500M functions as an "endless loop." When memory fills up, the unit begins overwriting the oldest data with the newest.

Each time the 8500M is powered up, the current time/date information (if the clock is set properly) is stored in memory to allow quick differentiation of recording sessions. Patient SpO₂ and pulse rate are sampled and stored every four seconds. The stored resolution of the oxygen saturation is in 1% increments in the range of 0 to 100 %. The stored pulse rate ranges from 18 to 300 BPM. The stored values have a resolution of 1 BPM from 18 to 200 and a resolution of 2 BPM from 201 to 300.

During the printing of the data, the last data recorded is the first data printed. For example, the last four minutes of data recorded would be the first four minutes of printout.

Recording Sessions

Each time the 8500M is turned on (except while setting the clock) data is automatically collected.

NOTE: *Only recording sessions greater than one minute in length are kept in memory for later printing.*

Memory Output Mode

To output the data stored in the memory of the 8500M, start with the unit OFF and then:

- 1) Hold the "↑" switch while pressing the "↓" switch;
- 2) Release the "↑" and "↓" switches when "000 000" is displayed in the SpO₂ and pulse rate LEDs;
- 3) Observe "Prrn XX" will be displayed in the SpO₂ and ♥ LEDs;
- 4) Data is automatically transferred from the memory in approximately 8 to 10 minutes.

Data is transferred at a rate of 20 minutes of collected data per second. An 18 hour recording session (the maximum memory saved) is transferred in approximately 1 minute. After all the data is transferred, the 8500M should be shut off before collecting new patient data. Patient information is held in memory as long as the batteries are good, so if the memory has to be cleared, remove the battery door for 60 seconds or longer. Outputting the memory does not clear any data from the memory.

NOTE: *The memory is cleared each time the batteries are changed.*

Specifications

1. Oxygen Saturation Range (SpO ₂)	0 to 100%
2. Pulse Rate Range	18 to 300 Pulses Per Minute
3. Displays	
Patient Indicator	Perfusion LED
Digital Displays	3-digit 7-segment LEDs
4. Measurement Wavelengths	
Red	660 nanometers
Infrared	910 nanometers
5. Accuracy	
SpO ₂ (±1 Standard Deviation)♦	<ul style="list-style-type: none">• 70 - 95% ±3 digits for neonates using infant or neonatal sensors• 70 - 100% ±2 digits for adults using the Finger Clip sensor• 70 - 100% ±3 digits for adults using Flex or Reflectance sensors• 70 - 100% ±4 digits using Ear Clip sensor• Below 70% is not specified for all sensors
Pulse Rate	±3% ±1 digit
6. Temperature	
Operating	-20 to +50°C
Non-operating	-30 to +50°C
7. Humidity	
Operating	10 to 90% non-condensing
Non-operating	10 to 95% non-condensing
8. Power Requirements	6 AA alkaline batteries 100 hours typical operation (maximum display brightness) 200 hours typical operation (minimum display brightness)
9. Dimensions	3" wide x 6" high x 1" deep 8 cm x 15 cm x 2 cm
10. Weight	10 oz. (280 g) with batteries

♦ Standard Deviation is a statistical measure: up to 32% of the readings may fall outside these limits.

Service

CAUTION! The 8500 Handheld Pulse Oximeters must be repaired by trained NONIN personnel only. Any evidence of opening the system, field service by non-NONIN personnel, tampering, or any kind of system misuse or abuse shall void the warranty.

The solid state circuitry within the 8500 requires no periodic maintenance or calibration other than battery replacement.

NONIN does not recommend field repair of the 8500 Handheld Pulse Oximeter. The circuit board in the 8500 series is a multi-layer board using traces 0.01" wide. Due to the very small trace size, extreme care must be used when replacing components to prevent permanent non-repairable damage to the circuit board. Most components are surface-mounted and require special hot air jet soldering and desoldering equipment. After any repairs are made, the pulse oximeter must be tested to ensure correct operation.

NOTE: *All repair work on the 8500 should be done by trained Nonin personnel. For Nonin Customer Support, contact:*

Nonin Medical, Inc.
13700 1st Avenue North
Plymouth, Minnesota
55441-5443 USA

1-800-356-8874
www.nonin.com

All non-warranty work shall be done according to NONIN standard rates and charges in effect at the time of delivery to NONIN. All repairs include a complete retest of the pulse oximeter using factory test fixtures.

Warranty

NONIN MEDICAL, INCORPORATED, (NONIN) warrants to the purchaser, for a period of three years from the date of purchase, each system exclusive of sensors, cables, and batteries. (Refer to the individual package inserts for specific warranty information for sensors, cables, and other accessories.) NONIN shall repair or replace any systems 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 said notification occurs within the applicable warranty period. This warranty shall be the sole and exclusive remedy by the purchaser hereunder for any systems or accessories delivered to the purchaser which are 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 device that is found to be within specifications.

These are precision electronic instruments and must be repaired by knowledgeable and specially trained NONIN personnel only. Accordingly, any evidence of opening the system, field service by non-NONIN personnel, tampering, or any kind of system misuse or abuse shall void the warranty in its entirety.

All non-warranty work shall be done according to NONIN standard rates and charges in effect at the time of delivery to NONIN.

DISCLAIMER/EXCLUSIVITY OF WARRANTY

THE EXPRESS WARRANTIES SET FORTH IN THIS MANUAL ARE EXCLUSIVE, AND NO OTHER WARRANTIES OF ANY KIND, WHETHER STATUTORY, WRITTEN, ORAL, OR IMPLIED, INCLUDING WARRANTIES OF FITNESS FOR A PARTICULAR PURPOSE OR MERCHANTABILITY, SHALL APPLY.

Accessories

The following NONIN accessories function with the 8500 and 8500M Handheld Pulse Oximeter:

Model	Description
<i>Pulse Oximeter Reusable Sensors</i>	
8000AA-1	Adult Articulated Finger Clip Sensor (1 meter)
8000AA-2	Adult Articulated Finger Clip Sensor (2 meters)
8000AA-3	Adult Articulated Finger Clip Sensor (3 meters)
8000AP	Pediatric Finger Clip Sensor (1 meter)
8000AP-3	Pediatric Finger Clip Sensor (3 meters)
8000J	Adult Flex Sensor
8000J-3	Adult Flex Sensor (3 meters)
8008J	Infant Flex Sensor
8001J	Neonatal Flex Sensor
8000Q	Ear Clip Sensor
8000R	Reflectance Sensor
<i>Pulse Oximeter Disposable Sensors</i>	
7000A	Adult Finger Flexi-Form® II Sensor, 10 per box
7000P	Pediatric Finger Flexi-Form® II Sensor, 10 per box
7000I	Infant Toe Flexi-Form® II Sensor, 10 per box
7000N	Neonatal Foot Flexi-Form® II Sensor, 10 per box
7000D	Flexi-Form® Sensor Assortment Pack, 10 per box
<i>Accessories</i>	
8000JFW	Adult FlexiWrap, 8000JFW, bag of 25
8001JFW	Neonatal FlexiWrap, 8001JFW, bag of 25
8008JFW	Infant FlexiWrap, 8008JFW, bag of 25
8000H	Reflectance Sensor Holder System
8000S	Patient Simulator
UNI-EXT-1	Extension Cable (3 feet/1 meter)
UNI-EXT-3	Extension Cable (10 feet/3 meters)
UNI-EXT-6	Extension Cable (20 feet/6 meters)
UNI-EXT-9	Extension Cable (30 feet/9 meters)
UNI-RA-0	Right Angle Connector (6 inches/0.15 meters)
UNI-RA-3	Right Angle Connector (10 feet/3 meters)
NiMHBC-NA	Nickel metal hydride battery charger with 12 AA NiMH batteries
NiMHBC-UNIV	Nickel metal hydride battery charger with 12 AA NiMH batteries
NiMHB	1800 mAh NiMH, 6 AA batteries for use with NiMHBC-NA and -UNIV
HHCC	Handheld carrying case, 8500/9840, black, holds 8500/9840 monitor, sensor, and airway adapter tubes
8500P	Attachable printer—8500/9840-series (Real-time pulse oximetry data)
8500PAP	Thermal Printer Paper (package of 20 rolls)
8500PCC	Carrying case for 8500, 8500P, and accessories (blue)
8500TS	Tabletop stand
8500CC-B	Carrying Case – Black
8500MB	Mounting Bracket (Wall or Pole Mount System)
1000MC	Memory Transfer Cable (for use with PC)
8500RB	Rubber Bumper
NVISION	nVISION software for Microsoft Windows 95/98/ 2000/NT 4.0 operating systems

For more information about NONIN parts and accessories, contact your distributor, or contact NONIN at (800) 356-8874 (USA and Canada) or (763) 553-9968.

Troubleshooting Guide

Symptom	Possible Cause	Possible Solution
8500 will not power up	Batteries are depleted	Replace all six batteries.
	Incorrect battery installation	Verify battery sequence.
Numeric displays are blinking at once per second	Battery voltage is low	Replace all six batteries of the 8500
	Incorrect battery installation	Verify battery sequence.
Minus sign appears in SpO ₂ display	Sensor fault exists. Sensor may have become dislodged from 8500 or from the patient	Verify sensor is connected to the 8500 and the patient correctly; Try a new sensor if condition persists
Segments of SpO ₂ or ♥ display are missing	Defective LED displays	Displayed values may not be reliable; discontinue use of 8500
Displayed pulse rate does not correlate to pulse rate displayed on ECG monitor	Excessive motion at sensor site may be prohibiting the 8500 from acquiring a consistent pulse signal	Eliminate or reduce cause of motion artifact or reposition sensor to new sensor site where motion is not present
	Patient may have an arrhythmia resulting in some heart beats that do not yield a perfusion signal at sensor sight	<i>Examine the patient:</i> Condition may persist even though both monitors are functioning properly if patient's arrhythmia persists
	Non-NONIN sensor is being used.	Replace sensor with a NONIN sensor
	ECG monitor may not be functioning properly	<i>Examine the patient:</i> Replace ECG monitor or refer to operator's manual for ECG monitor
Erratic ♥ display and/or yellow perfusion LED during use of electro-surgical equipment (ESU)	ESU may be interfering with oximeter performance	<i>Examine the patient:</i> move oximeter, cables, and sensor as far away from ESU as possible or refer to the ESU operator's manual
Perfusion is blinking yellow with each pulse	Perfusion signal at sensor site is marginal	<i>Examine the patient:</i> reposition sensor or select alternate sensor site
Unable to obtain green perfusion	Low patient pulse strength	Reposition sensor on patient
	Sensor site poorly perfused	
	Sensor not correctly positioned	
	Sensor attached too tightly or tape or other items are restricting perfusion at sensor site	Reapply sensor, select alternate sensor site, or remove restrictive material from sensor site
	Circulation reduced due to excess pressure between the sensor and a hard surface	Allow sensor and finger to rest comfortably on surface
	Excessive ambient light	Reduce ambient light
	Excessive patient motion	Reduce patient motion
	Sensor applied to a polished fingernail	Remove fingernail polish
Perfusion LED is blinking red and SpO ₂ and ♥ displays show dashes	Inadequate perfusion signal at sensor site	<i>Examine the patient:</i> reposition sensor or select alternate sensor site
	Excessive motion at sensor site may be prohibiting 8500 from acquiring a consistent pulse signal	Eliminate or reduce cause of motion artifact or reposition sensor to sensor site where motion is not present
Printer not printing out after changing Prn mode	Printer mode did not get updated internally	Reset pulse oximeter by turning it off and then on

If any of these solutions do not correct the problem, please contact NONIN Customer Support at (800) 356-8874 (USA and Canada) or (763) 553-9968.