

Reprocessed by



Sustainability Solutions







Instructions for Use
Reprocessed Pulse Oximeter Sensor

Reprocessed Device for Single Use

Caution: Federal (U.S.A.) law restricts this device to sale by or on the order of a physician.

- STERILE
- LATEX-FREE

Explanation of Symbols

	Sterilized by Ethylene Oxide Gas
	Date of Reprocessing
	Use by Date
	Product Code
	Do Not Reuse
	See Instructions For Use

Stryker Sustainability Solutions, Inc. © 2006, 2011
 1810 W Drake Dr.
 Tempe AZ, 85283
 sustainability.stryker.com
 888.888.3433

Reprocessed Pulse Oximeter Sensor Description

This Reprocessed Oxygen Transducer is a previously used Nellcor Puritan Bennett™ (NPB) Oxisensor II® or OxiMax™ transducer which has been reworked, inspected, tested, packaged and sterilized by Stryker Sustainability Solutions.

This insert is intended for the following oxygen transducers:

Model	Description
D-25	Adult Sensor for use on finger of patients > 30 kg, 18" cable
D-25L	Same as D-25, with 36" instrument cable
D-20	Pediatric Sensor for use on finger of patients 10-50 kg, 18" cable
N-25	Neonatal Sensor for use on patient foot if < 3 kg or finger if > 40 kg, 36" cable
I-20	Infant O2 Transducer for use on toe of patient 3-20 kg., 36" cable
Max-A	Adult Sensor for use on finger of patients > 30 kg, 18" cable
Max-AL	Same as Max-A, with 36" instrument cable
Max-P	Pediatric Sensor for use on finger of patients 10-50 kg, 18" cable
Max-N	Neonatal/Adult Sensor for use on patient foot if < 3 kg or finger if > 40 kg, 36" cable
Max-I	Infant O2 Transducer for use on toe of patient 3-20 kg., 36" cable

Indications for Use

This sensor is indicated for use in continuous noninvasive arterial oxygen saturation and pulse rate monitoring.

Contraindications for Use

This device should not be used in patients who exhibit allergic reactions to the adhesive tape.

Warnings

- Prior to use, read and follow these instructions as well as those of the Operator's Manual for your pulse oximetry system.
- This oxygen transducer package is provided sterile by method of ethylene oxide gas. Do not use if there is any evidence of damage to the sterile package.
- Inspect the sensor site periodically to ensure correct sensor alignment and adhesion. Skin integrity and circulation distal to the site should be checked routinely and the sensor relocated to another site if found to be compromised.
- Incorrect application or duration of use of a sensor can cause tissue damage.
- Do not use oximetry sensors during magnetic resonance imaging (MRI), as the conducted current may cause burns. Cross-interference between the two devices can also cause inaccuracies in the measurements of either system.
- Do not attempt to repair, modify or clean the sensor. Immersion in water will compromise the device performance.
- Reprocessed OxiMax™ sensors have the event history recording feature disabled.
- When uncertain about any measurement accuracy, check the patient's vital signs by alternate means, then make sure the pulse oximeter is working properly.
- In conjunction with clinical signs and symptoms, pulse oximeter sensors are exclusively designed to be used as an adjunct in patient assessment.
- Do not use a sensor or pulse oximeter cable if it is damaged and/or if optical components are exposed.
- Do not attach any cable intended for computer use into the sensor's port connector.

- Sensor application errors, certain patient and ambient environmental conditions, can affect pulse oximeter's readings and signal.
- Do not lift the sensor by the power cord or cable; this may cause the sensor to disconnect and drop on the patient.

Any of the following conditions can cause inaccurate oxygen measurements

- Failure to properly apply the sensor to the patient or to align the optical transducers.
- Application of sensor to an extremity with an arterial catheter, blood pressure cuff or intravascular infusion line in place.
- Application of sensor to a site that is too thick, thin or deeply pigmented.
- Venous pulsations if the sensor or supplemental tape is wrapped too tightly.
- Transducer exposure to excessive light. Cover the sensor with opaque material if it is suspected that the transducer is exposed to excessive ambient light.
- Intravascular dyes.
- Excessive motion. Locate sensor at a stationary site and try to keep patient still.

Sensor Specifications**Oxisensor®**

- Accuracy: SpO_2 : $\pm 3\%$ ($\pm 4\%$ in Neonates) over the range of 70% to 100%*
Pulse Rate: ± 3 beats/min over the range of 30-240 BPM
- Operating Environment: Temperature 5o to 50o C. 10% to 75% Relative Humidity

OxiMax™

- Accuracy: SpO_2 : $\pm 3\%$ ($\pm 4\%$ in Neonates) over the range of 70% to 100%*
Pulse Rate: ± 3 beats/min over the range of 30-180 BPM
- Operating Environment: Temperature 5o to 50o C. 10% to 75% Relative Humidity

*Oxisensors were validated with a Nellcor™ N-395 oximeter against co-oximetry measurements of arterial saturation during a controlled hypoxia "breathe-down" study. OxiMax™ sensors were validated with Nellcor™ N-595 oximeters against co-oximetry measurements of arterial saturation during a controlled hypoxia "breathe-down" study. Specified accuracy range is increased by $\pm 1\%$ for neonates in order to account for potential influences of fetal hemoglobin upon oximetry measurements.

Directions for Use

1. The package label is detachable and may be affixed to the medical record of the patient.
2. When selecting a sensor, consider patient's weight and activity level, need for sterility, perfusion adequacy, sensor site availability, and expected monitoring duration.
3. Locate a suitable site for monitoring oxygen saturation and pulse rate.
For pediatric and adult patients, an index finger is the preferred sensor site, or alternatively another finger or a great toe. For newborns and infants, the preferred site is the great toe or sole of the foot (alternatively the hand).
4. Peel the pouch open and remove the transducer from its package. Remove the plastic backing from sensor. On the non-adhesive side of the device are two solid lines with a dashed line at their midpoint. The solid lines are aligned with the transparent windows on the reverse (adhesive) side of the sensor that cover the optical transducers.
5. Orient the sensor so that the dashed line is at the tip of the finger/toe or on the lateral side of the foot/hand. The solid line closest to the cable should be positioned on the nail side of the finger/toe, the sole of the foot or palm of the hand. Wrap the sensor firmly (but not too tightly) around the finger, toe, foot or hand so that the solid lines oppose each other. Plug the sensor into the pulse oximeter module and verify proper operations as described in the system operator's manual. If the sensor does not indicate reliable tracking of the pulse rate, relocate sensor to an alternative site or choose an alternative sensor. The oxygen transducer can be reused on the same patient for as long as the adhesive provides an adequate attachment.

Returning the sensor to Stryker for Reprocessing

- Only sensors that functioned properly during clinical use should be placed in the collections container for reprocessing.
- Gently coil the sensor and place in the Stryker provided collection container.
- Once the container is full, place it in the pre-addressed carton provided by Stryker, seal the carton and deliver it to the hospital shipping department.

Warranty

Reprocessed Products

Stryker warrants all reprocessed products, subject to the exceptions provided herein, to be free from defects in reprocessing and to substantially conform to the product specifications contained in the documentation provided by Stryker with the products for one use in accordance with the instructions for use of such product.

STRYKER SHALL NOT BE LIABLE FOR ANY DAMAGES TO THE EXTENT CAUSED BY ANY DEFECT IN MATERIAL, WORKMANSHIP OR DESIGN BY THE ORIGINAL MANUFACTURER OF THE PRODUCT OR ANY ACT OR OMISSION OF THE ORIGINAL MANUFACTURER OF THE PRODUCT.

Products for which Stryker is the Original Manufacturer

Stryker warrants all products for which it is the original manufacturer, subject to the exceptions provided herein, to be free from defects in design, materials and workmanship and to substantially conform to the product specifications contained in the documentation provided by Stryker with the products for a period of one year from the date of purchase.

General Warranty Terms Applicable to All Products

TO THE FULLEST EXTENT PERMITTED BY LAW, THE EXPRESS WARRANTY SET FORTH HEREIN IS THE ONLY WARRANTY APPLICABLE TO THE PRODUCTS AND IS EXPRESSLY IN LIEU OF ANY OTHER WARRANTY BY STRYKER, EXPRESSED OR IMPLIED, INCLUDING, BUT NOT LIMITED TO, ANY IMPLIED WARRANTY OR MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE. IN NO EVENT WILL STRYKER'S LIABILITY ARISING IN CONNECTION WITH THE SALE OF THE PRODUCT (WHETHER UNDER THE THEORIES OF BREACH OF CONTRACT, TORT, MISREPRESENTATION, FRAUD, WARRANTY, NEGLIGENCE, STRICT LIABILITY OR ANY OTHER THEORY OF LAW) EXCEED THE PURCHASE PRICE, CURRENT MARKET VALUE OR RESIDUAL VALUE OF THE PRODUCTS, WHICHEVER IS LESS. STRYKER SHALL NOT BE LIABLE FOR INDIRECT, SPECIAL, INCIDENTAL, PUNITIVE OR CONSEQUENTIAL DAMAGES RESULTING FROM ANY BREACH OF WARRANTY OR UNDER ANY OTHER LEGAL THEORY.

This warranty shall apply only to the original end-user purchaser of products directly from Stryker or a Stryker authorized distributor. This warranty may not be transferred or assigned without the express written consent of Stryker.

This warranty does not apply to: (1) products that have been misused, neglected, modified, altered, adjusted, tampered with, improperly installed or refurbished; (2) products that have been repaired by any person other than Stryker personnel without the prior written consent of Stryker; (3) products that have been subjected to unusual stress or have not been maintained in accordance with the instructions in the user manual or as demonstrated by a Stryker representative; (4) products on which any original serial numbers or other identification marks have been removed or destroyed; or (5) products that have been repaired with any unauthorized or non-Stryker components.

If a valid warranty claim is received within thirty (30) days of the expiration of the applicable warranty period, Stryker will, in its sole discretion: (1) replace the product at no charge with a product that is at least functionally equivalent to the original product or (2) refund the purchase price of the product. If a refund is provided by Stryker, the product for which the refund is provided must be returned to Stryker and will become Stryker's property. In any event, Stryker's liability for breach of warranty shall be limited to the replacement value of the defective or non-conforming part or component.

If Stryker determines in its reasonable discretion that the claimed defect or non-conformance in the product is excluded from warranty coverage as described hereunder, it will notify the customer of such determination and will provide an estimate of the cost of repair of the product. In such an event, any repair would be performed at Stryker's standard rates.

Products and product components repaired or replaced under this warranty continue to be warranted as described herein during the initial applicable warranty period or, if the initial warranty period has expired by the time the product is repaired or replaced, for thirty (30) days after delivery of the repaired or replaced product. When a product or component is replaced, the item provided in replacement will be the customer's property and the replaced item will be Stryker's property. If a refund is provided by Stryker, the product for which the refund is provided must be returned to Stryker and will become Stryker's property.

The OEM information listed on the label is provided as device ID prior to reprocessing and may contain the trademarks of unrelated third parties that do not sponsor this device.

Sterilization: This product and its packaging have been sterilized with ethylene oxide gas (EtO). Even though the product then is processed in compliance with all applicable laws and regulations relating to EtO exposure, Proposition 65, a State of California voter initiative, requires the following notice:

Warning: This product and its packaging have been sterilized with ethylene oxide. The packaging may expose you to ethylene oxide, a chemical known to the State of California to cause cancer or birth defects or other reproductive harm.

Nellcor™ and OxiMax™ are trademarks of Nellcor Puritan Bennett LLC.
Oxisensor II® is a registered trademark of Nellcor Puritan Bennett LLC.

POX Rev. H 09/2017 RM702018