Root with 03° Regional Oximetry

Available for Adult, Pediatric, Infant, and Neonatal Applications





O3 Regional Oximetry

O3 Regional Oximetry may help clinicians monitor cerebral oxygenation in situations in which peripheral pulse oximetry alone may not be fully indicative of the oxygen in the brain.

O3 Regional Oximetry monitors the regional hemoglobin oxygen saturation of blood (rSO2) in the cerebral region for infant, neonatal, pediatric, and adult patients.

With their flexible design, O3 sensors easily conform to and allow for ergonomic application on foreheads of all sizes.



Infant and Neonatal Application

- > 3% A_{RMS}² trending accuracy specification
- > Patients less than 10kg



Pediatric Application

- > 5% A_{RMS} absolute accuracy and 3% A_{RMS} trending accuracy specifications
- > Patients between 5kg and 40kg



Adult Application

- 4% A_{RMS} absolute accuracy and 3%
 A_{RMS} trending accuracy specifications
- > Patients greater than 40kg

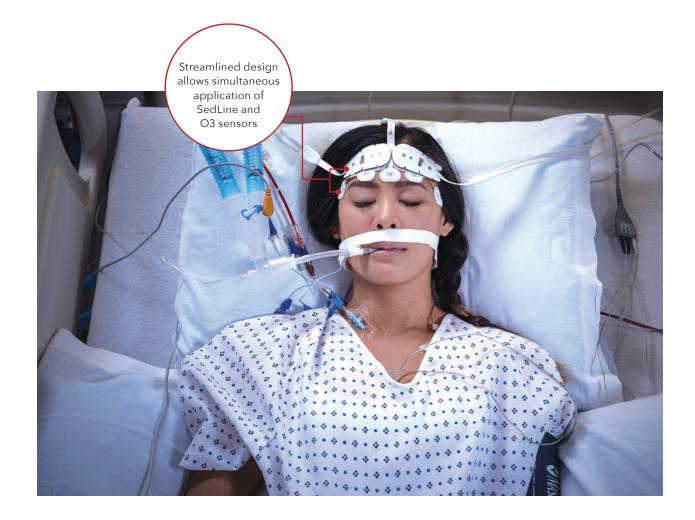
Expansion with Root

The expandable, versatile, and customizable Root patient monitoring and connectivity platform allows O3 Regional Oximetry to be combined with other monitoring modalities and automatically charts patient data in electronic medical records (EMRs).

Expanded Visibility of the Brain

Root with O3 Regional Oximetry and Next Generation SedLine® Brain Function Monitoring provides a more complete picture of the brain

Root with **Next Generation SedLine brain function monitoring** helps clinicians monitor the state of the brain under anesthesia with bilateral data acquisition and processing of four leads of electroencephalogram (EEG) signals, enabling continuous assessment of both sides of the brain.



When used together on Root, SedLine and O3 provide a more complete picture of the brain on an instantly interpretable, integrated display.

Patient State Index,

PSi, a processed EEG parameter related to the effect of anesthetic agents



rSO2 provides tissue oxygen saturation

Expanded Visibility of Oxygenation Status

Root with O3 Regional Oximetry and Masimo SET® Pulse Oximetry (SpO2)

O3 is displayed with Masimo SET® pulse oximetry on Root, providing clinicians with expanded visibility of a patient's oxygenation status.



Expanded Visibility of Patient Data

Iris Gateway® for Advanced Connectivity and Interoperability

Integrate data from Root and third-party devices using Iris® ports for automated charting in EMRs.



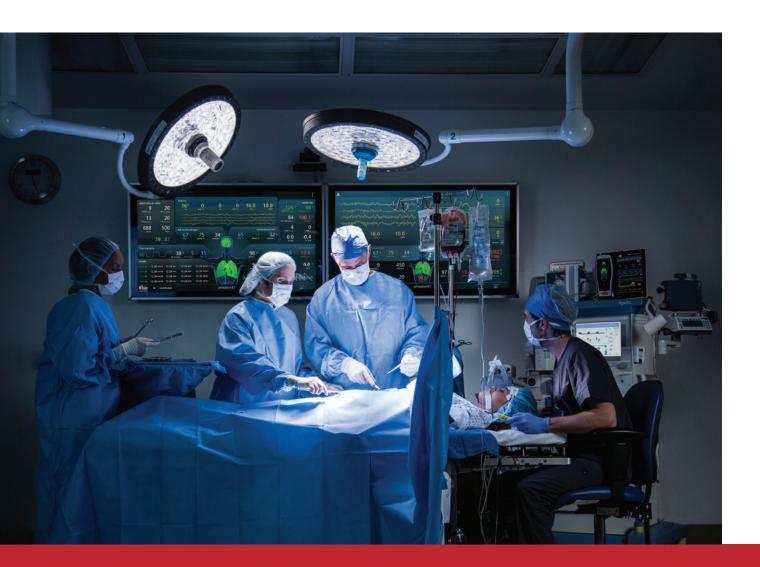
Data from Root and connected third-party devices

Device data and alarms are automatically charted in EMRs

Expanded Visibility Through Supplemental Display

UniView™ aggregates data and alarms from multiple Masimo and third-party devices – such as patient monitors, ventilators, anesthesia machines, IV pumps and others connected through Masimo systems – on a supplemental display.

- > Integrated real-time data visualization reduces cognitive overload and promotes data sharing among multiple clinicians, helping them to spot trends and coordinate care
- > Visual alarm indicators, relayed from connected devices, help care teams recognize patient distress and target areas for clinical focus
- > Tailored use-case-specific screen layouts optimize the presentation of advanced and integrated parameters, trend data, and waveforms in critical care areas
- > Adaptable layout automatically reconfigures based on connected devices





Kite® expands visibility by providing a supplemental display of patient data from Root, with the ability to customize the layout differently from Root.

By allowing customization of what can be displayed, Kite allows clinicians to focus on the most pertinent data for each stage of a patient's journey, empowering them to make more informed decisions.

With Kite, all clinicians in the OR can view brain monitoring information instantly, simultaneously.

O3 Module Specifications

PHYSICAL CHARACTERISTICS	ENVIRONMENTAL
Length (including cable) 12.1 ft (3.7 m) Width 1.8 in (4.6 cm) Thickness 0.6 in (1.5 cm) Weight 7.1 oz max (200 g max)	Operational Temperature .32 to 104° F (0 to 40° C) Storage Temperature 40 to 158° F (-40 to 70° C) Operating and Storage Humidity .10 to 95%, non-condensing Altitude Up to 12,000 ft (3700 m)

O3 Sensor Specifications

Application Site	4
Adult rSO ₂ Sensor Accuracy (A _{RMS}) ²	≥40 kg
Absolute Regional Oxygen Saturation (rSO2)	4%
Trending Regional Oxygen Saturation (rSO2)	3%
Pediatric rSO ₂ Sensor Accuracy (A _{RMS}) ²	. ≥5 kg and $<$ 40 kg
Absolute Regional Oxygen Saturation (rSO2)	5%
Trending Regional Oxygen Saturation (rSO2)	3%
Neonatal rSO ₂ Sensor Accuracy (A _{RMS}) ²	<10 kg
Trending Regional Oxygen Saturation (rSO2)	3%

ENVIRONMENTAL

Operating Temperature at Ambient Humidity	41 to 104° F (5 to 40° C
Storage Temperature at Ambient Humidity	40 to 140° F (-40 to 60° C
Storage Humidity	15% to 90%, 86 to 140° F (30 to 60° C

SedLine Module Specifications

PHYSICAL CHARACTERISTICS	ENVIRONMENTAL
Module Physical Dimensions 1.3 in (3.3 cm) Width 4.0 in (10.2 cm) Thickness 0.8 in (2.0 cm)	Module Operating Conditions 41-104°F (5-40°C) Operating Temperature. 41-104°F (5-40°C) Operational Humidity. 15-95%, non-condensing Module Storage Conditions .40-158°F (-40-70°C) Storage Temperature. .40-158°F (-40-70°C) Storage Humidity .15-95%, non-condensing
	Exposure to Pressure

SedLine Sensor Specifications

Application Site	Forehead
Active Channels	4
Active Electrodes L1,	L2, R1, and R2

Ground Electrode	CB
Reference Electrode	СТ
Duration of Use	Maximum of 24 hours
Latex Content	Does not contain natural rubber latex
Adult SedLine EEG Sensor	>18 years

Root Specifications

ELECTRICAL	PHYSICAL CHARACTERISTICS
Root AC Power Requirements	Weight <8 lbs (3.63 kg)
Operating Temperature. 32°F to 122°F (0°C to 50°C) Transport/Storage Temperature40°F to 158°F (-40°C to 70°C) Operating Humidity 10% to 95%, Non-Condensing Storage Humidity 10% to 95%, Non-Condensing Operating Altitude 500 mbar to 1060 mbar -1,000 ft to 18,000 ft (-304 m to 5,486 m)	ConnectorType (Number of Ports)Nurse Call1/4-in Round Female (1)MOC-9Masimo Connector (3)USBUSB 2.0 (2)

¹ ARMs accuracy is a statistical calculation of the difference between device measurements and reference measurements. Approximately two-thirds of the device measurements fell within ± ARMs of the reference measurements in a controlled study. ² This represents approximate run time at the lowest indicator brightness, using a fully charged battery.

Caution: Federal (USA) law restricts this device to sale by or on the order of a physician. See instructions for use for full prescribing information, including indications, contraindications, warnings, and precautions.





