

# Sentec Digital Monitor with OxiVenT™ Sensor



## tcPCO<sub>2</sub> | tcPO<sub>2</sub> | SpO<sub>2</sub> | PR | HP

Continuous or V-Check™ Mode

Neonates, Pediatrics & Adults

Noninvasive & Easy to Use

Accurate & Fast

Gentle & Safe

## Overall System Performance

### Transcutaneous Carbon Dioxide Partial Pressure (tcPCO<sub>2</sub>)<sup>1</sup>

**Measurement range:** 0–200 mmHg (0–26.7 kPa)

**Resolution:** 0.1 mmHg (0.01 kPa) below 100 mmHg (10 kPa) / 1 mmHg (0.1 kPa) above 100 mmHg (10 kPa)

**Drift:** Typically < 0.5%/hour

**Response time (T90):** Typically < 80 sec

**Linearity:** Typically < 1 mmHg (0.13 kPa)

**Interferences by anesthetic gases:** Negligible

**Stabilization/ artifact detection:** After sensor application or occurrence of a tcPCO<sub>2</sub> artifact, tcPCO<sub>2</sub> is displayed in grey until it (re)stabilizes.

### Oxygen Saturation (SpO<sub>2</sub>)

**Measurement range:** 1–100%

**Resolution:** 1%

**Accuracy:** ±2.25% (A<sub>rms</sub> over 70% to 100%)<sup>3</sup>

**Averaging mode:** 2, 4, 6, 8, 12, 16, and 32 sec

**Approved sites for SpO<sub>2</sub>/PR monitoring:** Earlobe, low on forehead, cheek, upper arm, on scapula (shoulder blade)

### Sensor Temperature

**Measurement range:** 0.0–70.0 °C

**Resolution:** 0.1 °C

**Accuracy:** ±0.2 °C (over 37.0 to 45.0 °C)

### Transcutaneous Oxygen Partial Pressure (tcPO<sub>2</sub>)<sup>4</sup>

**Measurement range:** 0–800 mmHg (0–106.7 kPa)

**Resolution:** 1 mmHg (0.1 kPa)

**Drift:** Typically < 0.1%/hour

**Response time (T90):** Typically < 150 sec

**Linearity:** Typically < 1 mmHg (0.13 kPa)

**Interferences by anesthetic gases:** Negligible

**Stabilization/ artifact detection:** After sensor application or occurrence of a tcPO<sub>2</sub> artifact, tcPO<sub>2</sub> is displayed in grey until it (re)stabilizes.

### Pulse Rate (PR)

**Measurement range:** 30–250 bpm

**Resolution:** 1 bpm

**Accuracy:** ± 3 bpm

### Pulsation Index (PI)

**Measurement range:** 0.1–10.0%

**Resolution:** 0.1%

### Sensor Heating Power (HP)

**Measurement range:**

Absolute Heating Power (AHP): 0–999 mW

Relative Heating Power (RHP): –999–999 mW

**Resolution:** 1 mW

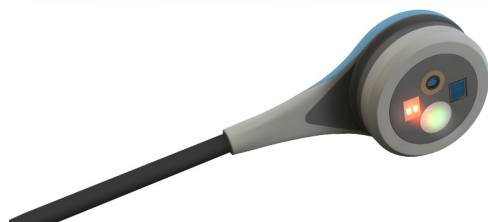
<sup>1</sup> An algorithm developed by J.W. Severinghaus is used to calculate tcPCO<sub>2</sub> from the measured cutaneous PCO<sub>2</sub>. This algorithm accounts for temperature and metabolic correction factors. The tcPCO<sub>2</sub> values displayed by the SDM are corrected/normalized to 37 °C and provide an estimate of arterial PCO<sub>2</sub> (PaCO<sub>2</sub>) at 37 °C. Correction factors can be customized by institution. Additionally, and subject to institution's permission, In-vivo Correction (IC) of tcPCO<sub>2</sub> values is possible at the bedside.

<sup>2</sup> Respective specifications based on in vitro tests performed as per IEC 60601-2-23:2011 at a sensor temperature of 43 °C.

<sup>3</sup> SpO<sub>2</sub> accuracy specification is based on controlled hypoxia studies on healthy, adult volunteers over the specified saturation range by applying a Sentec TC Sensor to each of the specified measurement sites.

<sup>4</sup> tcPO<sub>2</sub> designates an estimate of PaO<sub>2</sub> and corresponds to the measured PcO<sub>2</sub>. In newborns, PO<sub>2</sub> measured at the skin surface (PcO<sub>2</sub>) correlates with arterial PO<sub>2</sub> (PaO<sub>2</sub>) almost in a one-to-one relationship at a sensor temperature of 43 to 44 °C. The accuracy of PcO<sub>2</sub> compared to PaO<sub>2</sub> is best up to a PaO<sub>2</sub> of 80 mmHg (10.67 kPa), above which it increasingly tends to read lower than PaO<sub>2</sub>. Refer to J. W. Severinghaus, The Current Status of Transcutaneous Blood Gas Analysis and Monitoring, Blood Gas News 1998, 7(2): 4–9 and references contained therein.

## OxiVenT™ Sensor (OV-A/P/N) (Digital, combined tcPCO<sub>2</sub>/tcPO<sub>2</sub>/SpO<sub>2</sub>/PR sensor)



### General Characteristics

Suitable for neonatal, pediatric, and adult patients  
Reusable, waterproof

### Measurement Principle

Severinghaus-type PCO<sub>2</sub> sensor combined with reflectance 2-wavelength pulse oximetry and an optical fluorescence quenching PO<sub>2</sub> sensor.

### Digital Microtechnology

Highly integrated opto-electronic sensor head comprising micro pH-electrode, optical oximetry unit, temperature sensors, heating unit, optical fluorescence excitation/ sensing unit all combined in a fully digital design. High definition digitizer and pre-processing in the sensor head provides robust and low noise signals that are digitally transmitted to the Sentec Digital Monitor (SDM).

### Sensor Memory

Sensor-specific data are stored in the sensor's memory after manufacturing (serial number, factory PCO<sub>2</sub> sensitivity/ calibration, factory PO<sub>2</sub> sensitivity, calibration etc.) and during operation (sensor calibration, membrane change, etc.).

### Sensor Membrane Change

Up to 31 days (default 28 days). Patented '4 Press-and-Turn steps' membrane tool for simple and highly reproducible membrane change.

### Sensor Calibration

**Calibration duration:** Typically 3 minutes (ex. factory)

**Calibration interval:** Up to 12 hours. Once the calibration interval has elapsed, sensor

calibration is **recommended**, monitoring is possible for another 4 to 6 hours with tcPCO<sub>2</sub> marked as 'questionable'. Thereafter, sensor calibration is **mandatory** and tcPCO<sub>2</sub>/tcPO<sub>2</sub> are marked as 'invalid' (tcPCO<sub>2</sub>/tcPO<sub>2</sub> values replaced by '---'). tcPO<sub>2</sub> is calibrated during each mandatory calibration and subsequently approximately once every 24 hours during one of the anyways ongoing PCO<sub>2</sub> calibrations.

### SMART CALMEM

Supports the disconnection of the sensor for up to 30 minutes without losing the calibration status. Furthermore, the sensor can be removed from the Docking Station for up to 10 minutes without initiating a calibration upon reinsertion of the sensor into the Docking Station. Overall, SMART CALMEM significantly reduces the number of required calibrations and the calibration gas consumption.

### Sensor Internal Temperature Control

Sensor Temperature is reliably supervised/ controlled by two independent circuits. In case of errors the sensor's power consuming parts are switched-off.

### Sensor Dimensions/ Sensor Cable

**Diameter x height of sensor head:** 14 mm x 9 mm (0.55" x 0.35")

**Weight of sensor head:** < 2.9 g (0.1 oz)

**Length of sensor cable:** Approx. 80 cm (31") [plus Digital Sensor Adapter Cable of 150 cm (59"), 250 cm (98"), or 750 cm (295") length]

**Sensor cable:** Highly flexible, shielded cable with coating withstanding cleaning agents and irradiation commonly used in busy hospital environments.

### Transport/Storage of Sensor

**Transport temperature:** 0–50 °C (32–122 °F)

**Long term storage temperature:** 15–26 °C (59–78 °F)

Transport/ store sensor with membrane and protected from light/ radiation.

### Sensor Life Time

The sensor is granted in the sense of a license to the customer and its use is only granted and permitted during the life time (license period). With expiration of the life time, its operation automatically terminates. For details, please refer to 'Warranty and License Periods' on <https://www.sentec.com/info/legal-disclaimer/>.

## Sentec Digital Monitor (SDM) (Software version SMB SW-V08.00 and higher; with activated PO<sub>2</sub> measurement)

### Physical Characteristics

**Weight:** 2.3 kg (5.1 lbs) – including gas cylinder

**Size:** 10.2 cm x 27.0 cm x 23.0 cm (4.00" x 10.63" x 9.06")

**Flip feet:** Flip feet serving as carrying handle or to adjust angle for improved table-top viewing.

**Mountable:** Mountable on roll/ infusion stands, wall mounts/ railings, transport incubators, etc.

### Sensor Calibration

Built-in sensor calibration chamber for 1-point calibration. Automatic calibration ensures that system is 'Ready for use' if sensor is stored in calibration chamber. Comprehensive controls guarantee reliable calibrations.

### Sensor Temperature

**Selectable sensor temperature range:** Configurable by institution between 37.0 and 44.5 °C (in steps of 0.5 °C; default range=40.0–44.0 °C). Safety controls of the SDM may restrict the selectable range depending on the type of the connected sensor, the selected patient mode, or the enabled parameters.

**Selectable range:** 37.0–44.5 °C with OxiVen™ Sensor and 37.0–43.5 °C with V-Sign™ Sensor 2 (or as restricted by institution and/ or safety controls of the SDM). In steps of 0.5 °C. PO<sub>2</sub> only available with 41 °C or higher.

**Default sensor temperature:** If tcPO<sub>2</sub> is enabled, 43.0 °C in Neonatal Mode and 44.0 °C in Adult Mode. Otherwise, 41.0 °C in Neonatal Mode and 42.0 °C in Adult Mode (or closest setting of selectable range if default Sensor Temperature is outside selectable range).

### Initial Heating

Temporarily increases sensor temperature after sensor application for faster perfusion and results (in Adult Mode only; +2 °C max. 44.5 °C; can only be switched on if enabled by institution).

### Redundant Sensor Temperature Control on SDM

To guarantee safe operation should the sensor's temperature control fail, the SDM firmware redundantly controls the temperature of the connected sensor. Restarts or switches off sensor in case of errors.

### Site Time

**Maximal selectable 'Site Time':** Configurable by institution between 0.5 and 12.0 hours (in steps of 0.5 hours; max. 4.0 hours in Neonatal Mode at 43 °C or in Adult Mode at 44 °C). Depending on the selected patient mode and with increasing sensor temperature safety controls of the SDM may enforce a safer setting.

**Selectable range:** 0.5–12.0 hours (or as restricted by institution and/ or safety controls of the SDM). In steps of 0.5 hours.

**Default 'Site Time':** 2.0 hours in Neonatal Mode at 43.0 °C or in Adult Mode at 44.0 °C (or as restricted by institution and/ or safety controls of the SDM).

### Site Timer

Timer indicating remaining 'Site Time' during monitoring. Triggers an alarm once 'Site Time' has elapsed.

### Site Protection

Safety feature which reduces sensor temperature (to 39 °C if SpO<sub>2</sub> disabled and to 41 °C if SpO<sub>2</sub> enabled) once 'Site Time' has elapsed (can only be switched off if enabled by institution).

### Alarm System

**Alarm signals:** Visual/auditory alarm signals for high/low tcPCO<sub>2</sub>, tcPO<sub>2</sub>, SpO<sub>2</sub>, PR, technical alarms. 'Alarm Melodies' are institution-selectable.

**Alarm inhibition:** Auditory alarm signals can be PAUSED (1 or 2 minutes) or switched off permanently (if enabled by institution).

**Alarm system status indicators:** Alarm Status Icon, AUDIO Status Icon, AUDIO PAUSED/OFF Indicator (LED indicator), AUDIO OFF Reminder (can only be switched off if enabled by institution).

### Display/Indicators

**LED indicators:** ON/OFF; AUDIO PAUSED/OFF; AC Power/Battery; Battery Charging

**Display size:** 16 cm (6.3") diagonal TFT Color Display (LED backlight)

**Data update rate:** 1 sec for tcPCO<sub>2</sub>, tcPO<sub>2</sub>, SpO<sub>2</sub>, PR, RHP; between 1.5 and 30 mm/sec for Pleth Wave

**Data validity:** Clear representation of data validity/quality for tcPCO<sub>2</sub>, tcPO<sub>2</sub>, SpO<sub>2</sub>, PR, PI, RHP (valid, questionable, unstable, invalid)

**Measurement screens:** Various preconfigured, user-selectable measurement screens displaying values/ online trends for tcPCO<sub>2</sub>, tcPO<sub>2</sub>, SpO<sub>2</sub>, PR, RHP; alarm limits for tcPCO<sub>2</sub>, tcPO<sub>2</sub>, SpO<sub>2</sub>, PR; Baseline, Baseline values and Delta-x values for tcPCO<sub>2</sub>, tcPO<sub>2</sub>, SpO<sub>2</sub>, RHP; values for pulsation index, AHP, IC indicator; wiper bar Pleth Wave or blip bar reflecting relative pulse amplitude; visual alarm signals; status icons (e.g. remaining monitoring time) and status messages; 'Patient info' during remote monitoring with V-CareNet™

**'Calibration / Ready for use' screens:** 'Calibration' and 'Ready for use' screens displaying important system information (patient mode, sensor temperature and 'Site Time' related

settings, name of profile, 'Patient info' during remote monitoring with V-CareNet™, etc.)

**'Quick Access Menus':** To set new Baseline, new RHP reference, or 'Operator Events' during monitoring (and other functions).

**Languages:** Català, čeština, dansk, deutsch, english, español, français, italiano, japanese (katakana), polski, nederlands, norsk, português, русский (russian), svenska, suomi, türkçe

**Highly configurable:** Patient Mode, Enabled Parameters, Severinghaus Correction Mode, Heating Power Mode, V-Check™ Mode, Parameter Display Color, PCO<sub>2</sub>/PO<sub>2</sub> Unit, (time) ranges for online trends, sweep speed of Pleth wave, Sleep Mode, Brightness, Audio Settings, Menu Access, Profile Mode ('Basic' or 'Institutional')

### SDM Profiles

'SDM Profiles' help to ensure that all your SDMs can be configured the way you want them to. Within V-STATS™ preconfigured 'SDM Profiles' tailor-made to meet the specific needs of varying clinical settings are available. With V-STATS™, 'SDM Profiles' can be customized and up to 4 'SDM Profiles' can be stored on the SDM. During use of the SDM, the operator at any time can restore the active 'SDM Profile' (if modified) or select a different profile in the menu of the SDM. If at power-up the settings differ from those of the active 'SDM Profile' the modified settings can be maintained, the active 'SDM Profile' can be restored, or a different 'SDM Profile' can be selected.

### Special (safety-relevant) SDM Parameters

Within a password-protected area of V-STATS™ the institution can configure all menu accessible parameters as well as special parameters not being accessible in the menu of the SDM. Several of these special parameters permit to disable or to restrict operator-access to menu accessible parameters. The maximal 'Sensor Temperature' or the maximal 'Site Time' selectable at the bedside, for example, can be adapted to settings being safe for your typical patients.

### Patient Data Management

Data Recording Interval institution-selectable between 1 and 8 seconds; non-volatile memory providing between 35/ 227 hours monitoring data (at 1-/8-seconds resolution); automatic determination of measurement start/end enables convenient selection of measurement(s) for on-screen viewing/ printing of graphical trends and statistical summary. V-STATS™ provides fast data download to PC (approx. 3 min. for 8 hours data at 4-seconds resolution) for subsequent display, analysis, and reporting within V-STATS™. With V-CareNet™ simultaneous download is possible from multiple SDMs.

### Interfaces (isolated from sensor port)

**Serial output (RS-/EIA-232):** Supported protocols: SenTecLink, Philips VueLink/ IntelliBridge, Spacelabs Flexport, Serial Printer, TCB

**LAN port (Ethernet 10 BaseT):** For Remote Monitoring with V-CareNet™

**Analog output (0-1V):** tcPCO<sub>2</sub>, SpO<sub>2</sub>, PR, pleth wave (selectable ranges)

**Nurse-call:** Open and close type relays

### Electrical

**Instrument:** AC Power: 100 – 240 V (50/60 Hz), max. 900 mA/ Electrical Safety (IEC 60601-1): Class I, Type BF, Applied Part – Defibrillation Proof, IPX1.

**Internal battery:** Type: rechargeable, sealed Lilon Battery/ Capacity (new fully charged battery): up to 10 hours (if Sleep Mode=OFF, AUTO) and up to 12 hours (if Sleep Mode=ON)/ Charging Time: approx. 7 hours

### Environmental

**Transport/storage temperature:** 0 – 50 °C (32 – 122 °F)

**Transport/storage humidity:** 10 – 95% non-condensing

**Operating temperature:** 10 – 40 °C (50 – 104 °F)

**Operating humidity:** 15 – 95% non-condensing

**Operating altitude:** -400 – 4000 m (-1300 – 13120 ft) if connected to mains; -400 – 6000 m (-1300 – 19600 ft) if operated on battery.

**Built-in barometer:** Range: 350 – 820 mmHg (47 – 109 kPa)/ Accuracy: ± 3 mmHg (0.4 kPa)

### Compliance

IEC 60601-1, ANSI/ AAMI ES60601-1, CAN/CSA-C22.2 No. 60601-1, IEC 60601-1-6, IEC 60601-1-8, IEC 60601-2-23, ISO 80601-2-61, 60601-1-11, 60601-1-2

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Specifications are subject  
to change without notice

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