CARESCAPE V100

Vital signs monitor

The CARESCAPE* V100 monitor is designed for care areas where patients require vital signs measurements. It can go with you from one patient to the next, and because of its speed, accuracy and connectivity, the CARESCAPE V100 monitor collects the right information at the point of care to help you make fast, quality care decisions.

Features

- Can be used for spot-checking or for continuous monitoring, providing you the flexibility of two devices in one
- Designed for adult and pediatric use, as well as neonatal patients with very low perfusion rates
- Includes the same advanced parameters and algorithms as other higher acuity GE monitors, helping ensure measurement consistency across all care areas
- Non-invasive blood pressure measurement uses GE's exceptional DINAMAP* technology
- Three choices for pulse oximetry include GE Ohmeda TruSignal*; Nellcor OxiMax** or Masimo SET**
- Three options for temperature include Exergen** TemporalScanner**, Alaris** Turbo Temp** and Alaris Tri-Site
- Allows for inflation setpoints, so you can be sensitive to patients' special circumstances and ensure their comfort
- Large display makes it easy to read even from a distance
- Stores up to 40 measurements for up to 24 hours with the capability to print strips
- Designed for easy serviceability with a removable panel for easy access and simple field-replacement kits
- Typical battery life of up to 11 hours before requiring recharge. If the battery is discharged, it maintains the data.
- Connect up to three additional accessories simultaneously with the DINAMAP Serial Hub, via the monitor's HostComm (sold separately)







Technical specifications

Portability	Carried by recessed handle or on a roll stand
Printer	
Printer type	Thermal dot array
Resolution	384 dots/inch horizontal
Paper type	Must be compatible with
	GE PN 770137
Languages printed	English, German, French, Italian, Spanish, Portuguese, Hungarian, Polish, Czech, Finnish, Swedish, Danish, Dutch, Norwegian, and Slovak
Temperature options	

Exergen TemporalS*canner* temporal artery thermometer Alaris Turbo Temp** thermometer Alaris Tri-site thermometer

Performance specifications

GE TruSignal SpO₂ specifications

Measurement range	
SpO ₂	1 to 100%
Pulse rate	30 to 250 bpm
Accuracy	Saturation
Adult	70 to 100% ±2 digits (without motion)
Neonate ¹	70 to 100% ±3 digits (without motion)
Adult/Neonate ²	70 to 100% ±3 digits (during clinical motion)
Low perfusion	70 to 100% ±2 digits (during clinical low perfusion)
Pulse rate	
Adult/Neonate	30 to 250 bpm: ± 2 digits or ± 2%, whichever is greater, (without motion)
	30 to 250 bpm: ± 5 digits (during motion)
Low perfusion	30 to 250 bpm ±3 digits
	sed on deep hypoxia studies using TruSignal

1 SpO₂ measurement accuracy is based on deep hypoxia studies using TruSignal sensors on healthy adult volunteer subjects. Arterial blood samples were analyzed simultaneously on multiple CO-oximeters. This variation equals plus or minus one standard deviation. Plus or minus one standard deviation encompasses 68% of the population.

2 Applicability: TS-AF sensors.

NOTE: Accuracy may vary for some sensors; always check the instructions for the sensor.

GE sensor accuracy Sensor model SpO₂ range 70 to 100% TruSignal TS-F-D³ ±2 digits without motion TS-W-D³ ±2 digits without motion TS-E-D³ ±3 digits without motion TS-SE-3³ ±2 digits without motion TS-AF-10³ ±2 digits without motion TS-AF-25³ ±2 digits without motion TS-F2-GE ±2 digits without motion TS-F4-GE ±2 digits without motion TS-E2-GE ±3 digits without motion TS-E4-GE ±3 digits without motion

For TS-SA4-GE and TS-SA-D sensors the accuracy range is as following

70 to 100%	90 to 100%	80 to 90%	70 to 80%	below 70%
± 2 digits	± 1 digits	± 2 digits	± 3 digits	unspecified

Sensor light source

TS-SA4-GE

TS-SA-D³

Wavelength ⁴	Infrared: Red:	930 to 950 (nominal) 650 to 670 (nominal)
Maximum output power for each LED	< 15mV	

3 Requires compatible interconnect cable TS-G3

4 Information about wavelength range can be especially useful to clinicians.

Masimo SET specifications⁵ Measurement range		Masimo sensor accuracy ¹⁰	
		Sensor model	SpO ₂ range 70% to 100%
SpO ₂	1 to 100%	LNOP	2
Pulse rate	25 to 240 bpm	LNOP ADT	± 2 digits without motion
Perfusion range	0.02 to 20%	LNOP NEO	± 3 digits without motion
Accuracy and motion tolerance	Saturation	LNOP NEO-L	Foot \pm 3 digits without motion Finger \pm 2 digits without motion
Without motion adult/pediatric ⁶	70 to 100% ±2 digits	LNOP NEO PT-L	± 3 digits without motion
Without motion neonate ⁶	70 to 100% ± 2 digits 70 to 100% ± 3 digits	LNOP Adtx	± 2 digits without motion
With motion	70 to 10070 10 digits	LNOP Pdtx	± 2 digits without motion
adult/ped/neonate ^{7,8}	70 to 100% ±3 digits	LNOP DCI	± 2 digits without motion
Low perfusion ⁹	70 to 100% ±2 digits	LNOP DCIP	± 2 digits without motion
Pulse Rate	0 to 69% unspecified	LNOP Hi Fi-Neo/adult	Foot \pm 3 digits without motion Finger \pm 2 digits without motion
Without motion	25 to 240 bpm ±3 digits	LNOP Hi Fi-Infant/Ped	± 2 digits
With motion	Normal physiologic range 25 to 240 bpm ±5 digits	LNOP Blue Infant	Thumb/Toe ¹¹ \pm 3 digits (for 80-100) without
Low perfusion performance			motion
0.02% Pulse amplitude	Saturation (% SpO ₂)		± 4 digits (for 60-80) without motion
% transmission >5%	± 2 digits Pulse rate ± 3 digits		± 3.3 digits (for 70-100) without motion
Interfering substances: Carboxyhemoglobin may erroneously increase readings. The level of increase is approximately equal to the amount of carboxyhemoglobin		LNOP YI Multi-Site	Foot/hand ± 3 digits without motion Finger/toe ± 2 digits without

approximately equal to the amount of carboxyhemoglobin present. Dyes, or any substance containing dyes, that change usual arterial pigmentation may cause erroneous readings.

5 Masimo CSD-1201 (MS-2011 specifications cleared by the FDA).

- 6 The Masimo SET** SpO, parameter with LNOP-Adt sensors has been validated for no-motion accuracy in human blood studies on healthy adult volunteers in induced hypoxia studies in the range of 70-100% SpO₂ against a laboratory CO-oximeter and ECG monitor. This variation equals plus or minus one standard deviation. Plus or minus one standard deviation encompasses 68% of the population.
- 7 The Masimo SET SpO, parameter with LNOP-Adt sensors has been validated for motion accuracy in human blood studies on healthy adult volunteers in induced hypoxia studies while performing rubbing and tapping motions at 2 to 4 Hz at an amplitude of 1 to 2 cm and a nonrepetitive motion between 1 to 5 HZ at an amplitude of 2 to 3 cm in induced hypoxia studies in the range of 70-100% ${\rm SpO}_{\rm _2}$ against a laboratory COoximeter and ECG monitor. This variation equals plus or minus one standard deviation. Plus or minus one standard deviation encompasses 68% of the population.
- 8 The Masimo SET SpO₂ parameter with LNOP-Neo Pt sensors has been validated for neonatal motion accuracy in human blood studies on neonates while moving the neonate's foot at 2 to 4 cm against a laboratory CO-oximeter and ECG monitor. This variation equals plus or minus one standard deviation. Plus or minus one standard deviation encompasses 68% of the population.
- 9 The Masimo SET SpO, parameter has been validated for low-perfusion accuracy in bench-top testing against a Bio-Tek Index 2 simulator and Masimo's simulator with signal strengths of greater than 0.02% and a % transmission of greater than 5% for saturations ranging from 70 to 100%. This variation equals plus or minus one standard deviation. Plus or minus one standard deviation encompasses 68% of the population.

± 2 digits without motion LNCS Adult Adtx ± 2 digits without motion LNCS Ped Pdtx ± 2 digits without motion LNCS Infant-L ± 2 digits without motion ± 3 digits without motion

motion

± 2 digits without motion

± 3.5 digits without motion

± 3.5 digits without motion

± 2 digits without motion

LNCS Neo PT-L Resolution

LNOP DC-195

LNOP TC-I

LNCS TCI

LNCS DC-I

LNCS DC-IP

LNCS

Saturation (% SpO ₂)	1%
Pulse rate (bpm)	1

10 Masimo CSD-1109 (sensor specification)

¹¹ Masimo SET Technology with LNOP Blue sensors have been validated for no-motion accuracy in human blood studies on neonatal, infant and pediatric patients with congenital, cyanotic cardiac lesions in the range of 60% to 100% SpO₂ against a laboratory CO-oximeter. This variation equals plus or minus one standard deviation, which encompasses 68% of the population.

Sensor light source

Power dissipation

Wavelength¹²

Infrared: 905 nm (nominal) Red: 660 nm (nominal)
Infrared: 22.5 mW (max)
Red: 27.5 mW (max)

Nellcor OxiMax specifications¹³

Measurement range

1 to 100%
20 to 250 bpm
0.03 to 20%
Saturation
70 to 100% ±2 digits
70 to 100% ±3 digits
70 to 100% ±2 digits
20 to 250 bpm ±3 digits
20 to 250 bpm ±3 digits

13 Nellcor N600x Operator's Manual

14 Adult specifications are shown for OxiMax** MAX-A and MAX-N sensors with the N-600. Saturation accuracy will vary by sensor type. This variation equals plus or minus one standard deviation. Plus or minus one standard deviation encompasses 68% of the population. Accuracy is based on deep hypoxia studies on healthy adult volunteer subjects. Arterial blood samples were analyzed simultaneously on multiple CO-oximeters

15 Applicability: OxiMax MAX-A, MAX-AL, MAX-P, MAX-I, and MAX-N sensors.

Oxi-Max sensor accuracy¹⁶

NOTE: All Nellcor** OxiMax sensors must be used with the Nellcor cable; the SCP-10 cable. RS-10 and Oxisensor** II sensors are not compatible with the CARESCAPE V100.

Sensor model	SpO, range 70 to 100%
OxiMax	
MAX-A, MAX-AL	±2 digits
MAX-N (adult)	±2 digits
MAX-N ¹⁷ (neonate)	±3 digits
MAX-P	±2 digits
MAX-I	±2 digits
MAX-FAST	±2 digits
SC-A (adult)	±2 digits
SC-PR (neonate)	±3 digits
SC-NEO	±3 digits
MAX-R ¹⁸	±3.5 digits
OxiCliq**	-
OxiCliq A	±2.5 digits
OxiCliq P	±2.5 digits
OxiCliq N (adult)	±2.5 digits
OxiCliq N ¹⁷ (neonate)	±3.5 digits
OxiCliq I	±2.5 digits
Reusable sensor models	
D-YS (infant to adult)	±3 digits
D-YS (neonate)	±4 digits
D-YS & D-YSE	±3.5 digits
D-YS & D-YSPD	±3.5 digits
DS-100A	±3 digits
OXI-A/N (adult)	±3 digits
OXI-A/N (neonate)	±4 digits
OXI-P/I	±3 digits
Sensor light source	
Wavelength ¹⁹	Infrared: 890 nm (nominal) Red 660 nm (nominal)
Power dissipation	Infrared: 22.5mW (max) Red: 30 mW (max)
16 Nellcor oxygen saturation accuracy	
17 The MAX-N, D-YS, OXI-A/N, and OxiC	ling in were tested on patients >40 kg.

18 The accuracy specification has been determined between saturations of 80% to 100%

19 Information about wavelength range can be especially useful to clinicians.

Note: Neonatal Sensor Accuracy: When sensors are used on neonatal subjects as recommended, the specified accuracy range is increased by ± 1 digit, as compared to adult usage, to account for the theoretical effect on oximeter measurements of fetal hemoglobin in neonatal blood. For example, MAX-N accuracy on neonates is ± 3 digits, rather than ± 2 digits.

NIBP specifications

Cuff pressure range	0 to 290 mmHg (adult/ped)
(Normal operating range)	0 to 145 mmHg (neonate)
Blood pressure accuracy	
SuperSTAT	Mean error ≤5 mmHg,
NIBP algorithm	Standard deviation ≤8 mmHg (Meets ANSI/AAMI Standard SP10:1992)
Classic and auscultatory	Mean error ≤5 mmHg, standard deviation ≤8 mmHg (Meets ANSI/AAMI Standard SP10:2002)
Maximum determination time	120 s (adult/ped) 85 s (neonate)
Overpressure cutoff	300 to 330 mmHg (adult/ped) 150 to 165 mmHg (neonate)
Blood pressure range	
SuperSTAT NIBP Algorithm	
Systolic	30 to 290 mmHg (adult/ped) 30 to 140 mmHg (neonate)
MAP	20 to 260 mmHg (adult/ped) 20 to 125 mmHg (neonate)
Diastolic	10 to 220 mmHg (adult/ped) 10 to 110 mmHg (neonate)
Classic and auscultatory	
Systolic	30 to 245 mmHg (adult/ped) 40 to 140 mmHg (neonate)
MAP	15 to 215 mmHg (adult/ped) 30 to 115 mmHg (neonate)
Diastolic	10 to 195 mmHg (adult/ped) 20 to 100 mmHg (neonate)
Pulse rate range	
SuperSTAT	30 to 240 beats/min (adult/ped)
NIBP algorithm	30 to 240 beats/min (neonate)
Classic and auscultatory	30 to 200 beats/min (adult/ped) 30 to 220 beats/min (neonate)
Pulse rate accuracy	± 3.5% or 3 bpm, whichever is

 $\label{eq:greater} greater$ NOTE: To ensure accurate measurements, use only recommended blood pressure cuffs available from GE.

Exergen TemporalScanner specifications

Accuracy	± 0.1°C or 0.2°F
Temperature range	16° to 43°C (61° to 110°F)
Operating environment	16° to 40°C (60° to 104°F) (ambient)
Arterial heat balance range	
for body temperature ²⁰	34.5° to 43°C (94° to 110°F)
Resolution	0.1 °C or 0.1°F
Response time	0.04 seconds (approx.)

Alaris Turbo Temp specifications

Accuracy ²¹	± 0.1°C or 0.2°F
Temperature range	
Predictive mode	35.6° to 41.1°C (96° to 106°F)
Monitor mode	26.7° to 42.1°C (80° to 107.9°F)
Response time	As fast as 7 seconds

Alaris Tri-Site specifications

Accuracy ²¹	± 0.1°C or 0.2°F	
Temperature range		
Predictive mode	35° to 41.1°C (95° to 106°F)	
Monitor mode	26.7° to 42.1°C (80° to 107.9°F)	
Response time	As fast as 11 seconds	

20 Automatically applied when temperature is within normal body temperature range, otherwise reads surface temperature.

21 When tested in a calibrated liquid bath; meets ASTM E1112, Table 1, in range specified. Accuracy measured in continuous (monitor) mode.

NOTE: To ensure accurate measurements, use only recommended blood pressure cuffs available from $\ensuremath{\mathsf{GE}}$.

Power specifications

Universal power converter	P/N 2018859-001	Operating conditions	
AC input voltage	100 to 250VAC, 12VA	Temperature	5° to 40°C (41° to 104°F)
DC output voltage	12VDC at 1A	Atmospheric pressure	500 hPa to 1060 hPa
	The AC mains power adapter contains a non-resettable and non-replaceable fuse.	Storage conditions	
		Storage temperature	–20° to 50°C (– 4° to 122°F)
Protection against electrical shock	Internally powered or Class II when powered from specified external power supply	Atmospheric pressure	500 to 1060 hPa
		Humidity range	5 to 95% noncondensing
		Radio frequency	Complies with IEC 60601-1-2.
DC input voltage	12 VDC, supplied from a source conforming to IEC 60601-1.		Medical Electrical Equipment, Electromagnetic Compatibility Requirements and Tests and CISPR 11 (Class B, Group 1) for radiated and conducted emissions
Fuses	Monitor contains three fuses, mounted within. The fuses protect the low voltage DC input, the battery, and the remote alarm output. The +5 V output on the host port connector is regulated by internal supply.		
		Physical specifications	
		Dimensions (H \times W \times D)	19.5 x 21.9 x 13.5 cm (7.7 x 8.6 x 5.3 in)
Battery			19.5 x 25.4 x 13.5 cm (7.7 x 10 x 5.3 in)
Туре	Sealed lead acid, 6V, 3.3 Ahr		with Alaris temperature option
Battery life	5 hours with NIBP every 5 minutes and SpO ₂ , temperature and printer active	Weight	2.4 kg (5.4 lb) including battery
		Mountings	Self-supporting on rubber feet, pole mounted, or wall mount bracket
	11.5 hours non-SpO ₂ versions with a usage scenario of: NIBP determinations every 15 minutes without temperature active.		
		Roll stand (optional)	
		Height to mounting	
Charge time	Approximately 5 hours from full discharge when the monitor is off.	platform	100 cm (39 in) from floor to lowest position 125 cm (49 in) from floor to highest position
	Approximately 8 hours when the monitor is on.		
		Base Diameter	48 cm (19 in) 5 - 7.6 cm (3 in) casters—all non locking
			1/ C 0/ 7 1/ 0

Basket ($H \times W \times D$)

Certifications

Weight

14.6 x 26.3 x 16.8 cm (5.75 x 10.375 x 6.625 in)

11 kg (24 lb)

IEC 60601-1-2, IEC 60601-1-4, IEC 60601-1-8, IEC 60601-2-30,

UL 60601-1, CAN/CSA C22.2 No. 601.1, IEC 60601-1,

CE marked to the Medical Devices Directive - 93/42/EEC

IEC 60601-2-49, EN 1060-1, EN 1060-3, ISO 9919

Environmental specifications

© 2012 General Electric Company – All rights reserved.

* GE, GE Monogram, CARESCAPE, CARESCAPE iPanel and CARESCAPE Mobile Viewers, CRITIKON, DINAMAP, Ohmeda and TruSignal are trademarks of General Electric Company.

** Exergen and TemporalScanner are trademarks of Exergen Corporation. Alaris, Turbo-Temp and IBAC are trademarks of CareFusion Corp. Oxicliq, OxiMAX, Oxisensor, SatSeconds and Nellcor are trademarks of Nellcor Puritan Bennett, Inc. Masimo SET is a trademark of Masimo Corporation.

GE Healthcare reserves the right to make changes in specifications and features shown herein, or discontinue the product described at any time without notice or obligation. Contact your GE Healthcare representative for the most current information.

GE Medical Systems *Information Technologies*, Inc. a General Electric Company, doing business as GE Healthcare.

GE Healthcare Finland Oy, a General Electric company, doing business as GE Healthcare.

GE Healthcare, a division of General Electric Company.

About GE Healthcare

GE Healthcare provides transformational medical technologies and services that are shaping a new age of patient care. Our broad expertise in medical imaging and information technologies, medical diagnostics, patient monitoring systems, drug discovery, biopharmaceutical manufacturing technologies, performance improvement and performance solutions services help our customers to deliver better care to more people around the world at a lower cost. In addition, we partner with healthcare leaders, striving to leverage the global policy change necessary to implement a successful shift to sustainable healthcare systems.

Our "healthymagination" vision for the future invites the world to join us on our journey as we continuously develop innovations focused on reducing costs, increasing access and improving quality around the world. Headquartered in the United Kingdom, GE Healthcare is a unit of General Electric Company (NYSE: GE). Worldwide, GE Healthcare employees are committed to serving healthcare professionals and their patients in more than 100 countries. For more information about GE Healthcare, visit our website at www.gehealthcare.com.

GE Healthcare P.O. Box 900, FIN-00031 GE, Finland GE Direct United Kingdom: +44 (0)800 0329201

www.gehealthcare.com

