by anandic

CARESCAPE Patient Data Module

High-acuity mobile patient monitoring

The CARESCAPE* Patient Data Module helps you to transport patients to the right place at the right time so you can deliver a consistent level of care virtually anywhere.

Features

- Helps eliminate the traditional tangle of cables connected to multiple individual monitors by uniting common parameters in one convenient, compact, ergonomically designed unit, allowing better access to the patient in emergency situations and quickly prepare for transport
- With either the CARESCAPE Monitor B450 or the Transport Pro* monitoring device, provides uninterrupted flow of clinical intelligence before, during and after intra-hospital transport
- Powers the Transport Pro device in case of a depleted or missing battery to ensure monitoring continuity
- Refreshes the patient's record with the data collected before and during transport, when reconnected to the network in the new location, eliminating time-consuming ECG template resets and critical data gaps
- Supports patient monitoring in—and between—the highest-acuity clinical environments
- Exceptional parameter set including GE's clinical algorithms, including 12SL* 12-lead ECG, 12RL* derived 12-lead ECG, GE EK-Pro four-lead arrhythmia analysis, GE DINAMAP* SuperSTAT non-invasive blood pressure, and Masimo® SET® or Nellcor® OxiMax® SpO₂





Performance specifications

ECG

Standard leads available I, II, III, V1 to V6, aVR, aVL, and aVF

Leads analyzed Twelve (I, II, III, V1 to V6, aVR,

simultaneous aVL, and aVF)

Lead fail Identifies failed electrodes and

switches to those intact

Lead fail sensing current Active electrodes: < 30 nA each,

referenced electrode < 270 nA

Gain selections 0.5x = 5 mm/mV

1x = 10 mm/mV

2x = 20 mm/mV

4x = 40 mm/mV

Display bandwidth

Diagnostic 0.05 to 100 Hz

Monitoring 0.05 to 32 Hz (with 50 Hz

powerline frequency)

0.05 to 40 Hz (with 60 Hz

powerline frequency)

0.05 to 150 Hz

Moderate 0.05 to 22 Hz

Maximum 5 to 25 Hz

ECG diagnostic (12SL)

analysis signal bandwidth

Differential offset voltage ±1 V

Input impedance

Common mode > 10 M Ω at 50/60 Hz

Differential > 2.5 M Ω from dc to 60 Hz

Maximum tall t-wave

rejection capability

Overall system error Less than ±5%; using the method described in AAMI EC11 3.2.7.1

Leadwire supported 3-, 5-, 6-, and 10-leadwire

Leddwire supported 5,5,6, drid 10-leddwir

Input voltage range for ± 2 mV to ± 700 mV

pace detection and rejection

Pacemaker marker 5 V, 2 ms pulse; summed with

the ECG analog output

For a 1 mV QRS test signal is 1.5 mV

Defibrillator sync delay < 35 ms

Defibrillation protection 5000 V, 360 J

Analog output

ECG signal output 1 V/1 mV

ECG signal bandwidth 0.05 to 100 Hz

ECG analog output delay < 35 ms

Input specification

QRS detection range ± 0.5 mV to ± 5 mV

Signal width 40 ms to 120 ms (Q to S)

Heart rate range 30 to 300 beats per minute

Common mode rejection 90 dB minimum at 60 Hz

Gain accuracy ±5% (diagnostic mode)

Linearity deviation ±5%

Noise $< 30 \,\mu\text{V}$ (referred to input)

Sampling rate

Monitoring mode 240 samples/second

Diagnostic mode 500 samples/second

Heart rate

The ECG heart rate indicates a new heart rate for a simulated step increase of 80 to 120 bpm in less than 8.6s (range 7.9 to 9.5s), and a step decrease of 80 to 40 bpm in less than 9.8s

(range .2 to 11.2s).

Heart rate calculation operates with irregular rhythms of

ANSI/AAMI EC13 4.1.2.1e as follows:

a): 80 bpm

b): 60 bpm

c): 120 bpm

d): 87 bpm

Heart rate is computed by converting the total time duration of the most recent 4 or 8 RR intervals into

an equivalent heart rate.

Heart rate averaging 8/4 beats

Display update interval < 2 seconds

Response time < 6 seconds

Heart rate alarm range 0 to 300 beats/minute, high limit

> low limit

PVC range Solar	1 to 100 PVCs/minute	Pace detection/rejection Input voltage range	±2 mV to ±700 mV	
		, ,		
CARESCAPE modular monitor	0 to 300 PVCs/minute	Input pulse width:	0.1 ms to 2 ms	
Method	QRS morphology classification and timing based on single or multiple-lead analysis	Rise time	0 μs to 100 μs	
rietriod		Over/under shoot	Overshoot measured using Method A of AAMI EC13 4.1.4.2	
Arrhythmia calls	Full, lethal only, or no arrhythmia	Detection/rejection mode	Pacemaker artifact rejection "On" or 'Off'	
PVC rate resolution	1 PVC/minute	Standard leads available	I, II, RL, LL	
ST segment analysis Measurement description	ST segment deviation is measured and displayed for all acquired leads	Accuracy	±1% or ±1 bpm, whichever is greater	
ST display	Lead label, ST deviation, current complex superimposed over a reference complex, J-point indicator and 15-minute mini-trends are shown for all acquired leads	Resolution	1 bpm	
ST display		Sensitivity	≥ 0.5 mV peak	
		ST numeric accuracy	±0.3 mm or 20%, whichever is greater	
Measurement point	Measured at user-selectable measurement points (0, 30, 40, 50, 60, and 80 ms) following the J-point	QT numeric range	100 to 900 ms	
		QT numeric accuracy	±30 ms	
		QT numeric resolution	1 ms	
Measurement range	-12.0 mm to 12.0 mm	QTc numeric range	100 to 900 ms	
Display resolution	0.1 mm	QTc numeric resolution	1 ms	
ST measurement	16 beats averaging	Descination		
ST alarm limits	±12mm, high limit > low limit, for any event within a lead group (inferior, lateral, or anterior) that exceeds the alarm limit for that group	Respiration Respiration range limit	1 to 200 breaths/minute	
		Input impedance range Dynamic	0.4 to 10 Ω	
		Static	100 to 1500 Ω @52.7 kHz	
		Respiration rate alarm range	1 to 200 breaths/minute	
		No Breath alarm range	3 to 30 seconds	
		Impedance respiration me	easurement	
		Accuracy	±1 breath/minute over the range of 0 to 120 breaths per minute	
			±3 breaths/minute over the range of 121 to 200 breaths per minute	
		Impedance respiration update interval	1s	

Temperature Number of channels	up to 2 (with Y-adapter cable)	Transducer requirements Excitation voltage	+2.5 V DC ±0.1%
Input specifications		Transducer output	5μV/V/mmHg
Probe type	Series 400 or 700 (determined by input cable)	Input specifications range Solar	-25 mmHg to 349 mmHg
Temperature range	0°C to 45°C (32°F to 113°F)	CARESCAPE	-98 mmHg to 349 mmHg
Resolution	±0.1°C (±0.1°F)	modular monitors	
Output specifications Parameters displayed	T1, T2	Output specifications Displayed frequency response	0 to 12 Hz or 0 to 40 Hz (-3dB) user-selectable
Accuracy	±0.1°C (±0.2°F) for series 400 probes	Zero balance range	±150 mmHg (±20.0 kPa)
(independent of source)	±0.3°C (±0.5°F) for series	Zero balance accuracy	±1 mmHg (±0.1 kPa)
Alarms	700 probes User-selectable upper and lower limits	Accuracy	±2% or ±1 mmHg, whichever is greater (exclusive of transducer)
Test measurement cycle	Every minute		±2% or ±2 bpm, whichever is greater
Invasive pressures		Sweep speed options	6.25, 12.5, 25, and 50 mm/s
Number of channels	Up to 4 (with appropriate cables)	Pulse rate	1 bpm
Transducer sites, site name, and displayed values		display resolution	
Arterial (ART)	Systolic, diastolic, mean and rate	Pulse rate range	30 to 300 bpm
Femoral (FEM)	Systolic, diastolic, mean and rate	Display scale selections	0-30, 0-40, 0-60, 0-100, 0-160, 0-200, 0-300 mmHg (0.0-2.0, to 0.0-40.0 kPa, with a step size of 2.0 kPa)
Pulmonary artery (PA)	Systolic, diastolic, mean		
Central venous pressure (CVP)	Mean		
Left atrial (LA)	Mean	Alarms	User selectable upper and lower limits for systolic, diastolic, and mean pressures
Right atrial (RA)	Mean		
Intracranial pressure (ICP)	Mean	Alarm range	-99 to 350 mmHg
Umbilical artery (UAC)	Systolic, diastolic, mean, and rate	Analog output Invasive pressure output	
Umbilical vein (UVC)	Mean		1V/100 mmHg
Special pressure (SP)	Mean	Invasive pressure analog	< 35 ms

output delay

Mean

Special pressure (SP)

Non-invasive blood pressure		Pressure accuracy	
Measurement technique	Oscillometric	Static	±2% or ±3 mmHg (0.4 kPa),
Displayed parameters	Systolic, diastolic, and mean pressures, pulse rate, time of last measurement		whichever is greater
		Clinical	±5 mmHg (0.7 kPa) average error, 8 mmHg (1.1 kPa) standard deviation
Modes	Manual, Auto and Stat	Auto zoro	
Heart rate detection		Auto zero	Zero pressure reference prior to each cuff inflation
Adult and Pediatric	30 to 240 beats/min	Tubing length	Variable
Neonate	30 to 240 beats/min	Automatic cuff deflation	Cycle time exceeding 2 minutes
Total cycle time	20 to 40 seconds typical (Dependent on heart rate		(85 seconds neonatal), power off, or cuff pressure exceeds
	and motion artifact)	Adult	290 ±6 mmHg (38.7 ±0.8 kPa)
Systolic pressure range Adult	30 to 290 mmHg (4.0 to 38.7 kPa)	Pediatric	250 ±5 mmHg (33.3 ±0.7 kPa)
Pediatric	30 to 240 mmHg (4.0 to 32.0 kPa)	Neonatal	145 ±5 mmHg (19.3 ±0.7 kPa)
Neonatal	30 to 140 mmHg (4.0 to 18.7 kPa)	Cuff sizes	
Diastolic pressure range	00 to 110g (no to 10m u,	Disposable	Large adult, adult, small adult,
Adult	10 to 220 mmHg (1.3 to 29.3 kPa)	Reusable	pediatric, child, and neonatal
Pediatric	10 to 200 mmHg (1.3 to 26.7 kPa)		Adult thigh, large adult, adult, small adult, small adult, small adult/child,
Neonatal	10 to 110 mmHg (1.3 to 14.7 kPa)		child, and infant
Mean pressure range		Alarms	User selectable upper and lower limits for systolic, diastolic, and mean pressures
Adult	20 to 260 mmHg (2.7 to 34.7 kPa)		
Pediatric	20 to 215 mmHg (2.7 to 28.7 kPa)	Maximum inflation pressures	
Neonatal	20 to 125 mmHg (2.7 to 16.7 kPa)	Adult/pediatric	315 ±5 mmHg (42.0 ±0.7 kPa)
Cuff pressure range		Infant	157 ±5 mmHg (20.9 ±0.7 kPa)
Adult	0 to 290 mmHg	Automatic cycle times 1 min, 2 min, 2.5 min, 3 min, 4 min, 5 min,	
Pediatric	0 to 250 mmHg		
Neonatal	0 to 145 mmHg	10 min, 15 min, 20 min, 30 min, 1 h, 2 h and 4 h	
		Default NIBP measuremer Adult	nt initial inflation pressures 135 mmHg (18.0 kPa)
		Pediatric	125 mmHg (16.7 kPa)
		Infant	100 mmHg (13.3 kPa)

Pulse oximetry

Display resolution 1 digit (% of SpO₂)

Peripheral pulse

1 bpm

rate resolution

Display update period

Less than 30s

Sweep speed options

6.25, 12.5, 25, and 50 mm/s

Waveform scale options

1x, 2x, 4x, and 8x

Wavelength of SpO₂ probe LEDs using Masimo

LNOP and LNCS sensors

Infrared LED 905 nm

Red 660 nm

LNOP and LNCS tip clips

Infrared LED 880 nm

Red 653 nm

LNOP and LNCS TF-I

Infrared LED 880 nm

Red 660 nm

Wavelength of SpO₂ probe LEDs using Nellcor

Infrared LED 900 nm

Red 660 nm

Parameters monitored Arterial oxygen saturation (SpO₂)

and pulse rate

Probe types Masimo (reusable/single use)

Nellcor (reusable/single use)

Masimo range SpO₃: 1 to 100%

> Pulse Rate: 25 to 240 beats per minute

Nellcor range SpO₂: 1 to 100%

> Pulse rate: 20 to 300 beats per minute

Masimo accuracy¹

Without motion SpO₂ (70% to 100%):

±2 Adult, ±3 Neonatal

SpO₂ (< 70%): Unspecified

With motion SpO₂ (70% to 100%):

> ±3 Adult. ±3 Neonatal SpO₂ (< 70%): Unspecified

Low perfusion SpO₂ (70% to 100%):

±2 Adult, ±3 Neonatal SpO₂ (< 70%): Unspecified

Nellcor accuracy¹

With/without motion SpO₂ (70% to 100%):

> ±2 Adult, ±2 Neonatal SpO₂ (60% to 80%): ±3 Adult, ±3 Neonatal SpO₂ (< 60%): Unspecified

SpO₂ (70% to 100%): Low perfusion

> ±2 Adult, ±2 Neonatal SpO₂ (< 70%): Unspecified

Messages No Sensor, Defective Sensor,

> Sensor Off, Unrecognized Sensor, Low Perfusion, Pulse Search, Interference Detected,

Ambient Light, Low Signal IQ

Nellcor Probe off patient, low quality,

pulse search

⁽¹⁾ Refer to Probe Manufacturer's specifications for probe accuracy statement.

Cardiac output

Method Thermodilution

Cardiac output range 0.2 to 15 liters per minute

Blood temperature range 17°C to 42°C (62.6°F to 107.6°F)

Blood

±0.5°C (0.9°F):

temperature accuracy BT 17°C to 30°C (62.6°F to 86.0°F)

±0.2°C (0.4°F):

BT30°C to 42°C (86.0°F to 107.6°F)

Injectate

0°C to 30°C (32°F to 86°F)

temperature range

Injectate ± 0.3 °C (± 0.6 °F)

temperature accuracy

Blood temperature 0.1°C (0.1°F)

display resolution

Output parameters Cardiac output, blood temperature,

injectate temperature, real-time cardiac output washout curve,

last average CO

Cardiac output review accept/reject individual measurements

and store average

Catheter sizes 5, 6, 7, 7.5, or 8 French

Injectate 3, 5, or 10

volume selections

Environmental specifications

Operating conditions

Heat dissipation 15.36 BTU/hour

Temperature 10°C to 35°C (50°F to 95°F)

Relative humidity 15% to 95% (non-condensing)

Storage conditions

Temperature -40°C to 60°C (-40°F to 140°F)

Relative humidity 15% to 95% (non-condensing)

Power specifications

Cooling Natural convection

Batteries

Type Removable lithium ion

Quantity One

Voltage 11.1 Volt (nominal)

Capacity 1.8 Amp hour (nominal)

Charge time Approximately 2.5 hours

Run time Approximately 1.5 hours

(new, fully charged)

Battery Life 300 cycles to 60% capacity

Physical specifications

Dimensions (H x W x D) $7.0 \times 14.6 \times 21.6 \text{ cm}$

 $(2.75 \times 5.75 \times 8.5 \text{ in})$

Weight 1.1 kg (2.4 lb) without battery

1.3 kg (2.9 lb) with battery

Warranty

One year



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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.

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