DINAMAP® PRO 100-400
PATIENT MONITOR

It's What's Inside That Counts!

With the DINAMAP PRO Series, you can take BP measurements that are fast, accurate and comfortable. Get reliable temperature readings in seconds and get oxygen saturation readings with your choice of two superior $Sp0_2$ technologies. Which means you can spend less time checking vitals and more time on actual patient care.

Market-Leading Technologies

Designed for quality driven healthcare professionals who demand proven world-class technology in every parameter. The DINAMAP PRO 100-400 Series brings these technologies together for the first time in one complete line of flexible monitors.

Built to Last

Above all, you need a monitor you can depend on. As we strive to continuously improve our products, our latest generation includes enhancements in performance, reliability and serviceability. In addition to incorporating the highest-quality materials for "hospital grade" durability, the new PRO Series is also designed to require little maintenance and be easy to repair. Furthermore, our warranty is the best in the business and fully backed by GE - one of the most respected companies in the world. That's peace of mind that delivers day in and day out.

Mechanical

Dimensions	Height: 9.8 in (25.0 cm) Width: 9.8 in (24.8 cm) Depth: 6.9 in (17.5 cm)
Weight, Including Battery	7.8 lb (3.5 kg)
Mountings	Self-supporting on rubber feet or pole mountable
Portability	Carried by recessed handle or pole mounted
Classification Information	Mode of operation: continuous Degree of protection against harmful ingress of water: Drip-proof IPX1
Power Requirements	AC input voltage: 100-240 VAC, 50 / 60 Hz (nominal) 90 ~ 253 VAC, 47 ~ 63 Hz (range), 50VA. Protection against electrical shock: Class 1 DC input voltage: 24 VDC (nominal), 12-30 VDC, 36VA, supplied from a source conforming to IEC 601-1. AC input is protected by two internal fuses, replaceable by qualified service personnel only. DC input line is protected by an internal auto-resetting fuse. Battery: 12 volt, 2.3 amp-hours protected by internal auto-resetting fuse. Minimum operation time: 2 hrs (5 min cycle with adult cuff at 25 ∞C with power save mode enabled) from full charge. Time for full recharge: 1 hr 50 min from full discharge when the Monitor is switched off and 8 hrs when the Monitor is switched on.
Environmental Operating Temperature	+ 5 ∞C to + 40 ∞C (+ 41 ∞F to + 104 ∞F)
Operating Atmospheric Pressure	700 hPa to 1060 hPa
Storage Temperature	ñ 20 ∞C to + 50 ∞C (ñ 4 ∞F to + 122 ∞F)
Storage/Transportation Atmospheric Pressure	500 hPa to 1060 hPa
Humidity Range	0% to 95% noncondensing
Radio Frequency	Complies with IEC Publication 601-1-2 (April 1993) Medical Electrical Equipment,
Electromagnetic	Compatibility Requirements and Tests and CISPR 11 (Group 1, Class A) for radiated and conducted emissions.

IPX1

The DINAMAP PRO Monitor is protected against vertically falling drops of water and conforms to IEC-529 standard at level of IPX1. Vertically falling drops of water shall have no harmful effects to the Monitor.

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NIBP

Cuff Pressure Range	0 to 290 mmHg (adult)
(Normal operating range)	0 to 140 mmHg (neonate)
Default Target: Cuff Inflation	160 ± 15 mmHg (adult)
	110 ± 15 mmHg (neonate)
Target Cuff Inflation:	100 to 250 mmHg (adult)
Adjustment Range	100 to 140 mmHg (neonate)
(in 5 mmHg increments)	
Blood Pressure Measurement	

	Range (mmHg)	Systolic	MAP	Diastolic
	Adult 30 - 245	15 - 215	10 -195	
	Neonate	40 - 140	30 - 115	20 -100

Blood Pressure Accuracy	Meets or exceeds ANSI/AAMI standard SP-10 (mean error £5 mmHg, standard deviation £8 mmHg)
Maximum Determination Time	120 s (adult) 85 s (neonate)
Overpressure Cutoff	300 to 330 mmHg (adult) 150 to 165 mmHg (neonate)
Pulse Rate Range	30 to 200 beats/min (adult) 30 to 200 beats/min (neonate)
Pulse Rate Accuracy	± 3.5%

Turbo•Temp Temperature

Scale	∞Fahrenheit (F) ∞Celsius (C)
Range	
Monitor mode	Predictive mode Max: 41.1∞ C; 106.0∞ F Min: 33.6∞ C; 96.0∞ F Max: 41.1∞ C; 106.0∞ F Min: 26.7∞ C; 80.0∞ F
Monitor mode accuracy	± 0.1∞ C ± 0.2∞ F (when tested in a calibrated liquid bath; meets ASTM E1112, Table 1, in range specified)
Predictive mode accuracy	± 1.0∞ F ± 0.6∞ C
Determination time	less than 60 seconds

Use only IVAC probes and probe covers. The size, shape, and thermal characteristics of the probe covers can affect the performance of the instrument. Inaccurate readings or retention problems may occur unless IVAC probes and probe covers are used.

Nellcor SpO2

Measurement Range	
SpO2	1 to 100%
Pulse Rate	20 to 250 beats/min
Perfusion Range	0.03 to 20%
Accuracy and Motion Tolerance	
Saturation	
Without Motion - Adults*	70 to 100% ±2 digits
Without Motion - Neonate*	70 to 100% ±3 digits
With Motion - Adults/Neo**	70 to 100% ±3 digits
Low Perfusion	70 to 100% ±2 digits
Pulse Rate	
Without Motion	20 to 250 beats/min ±3 digits
With Motion	normal physiologic range 55 to 125 beats/min ±5 digits
Low Perfusion	20 to 250 beats/min ±3 digits

*Adult specifications are shown for OXIMAX MAX-A and MAX-N sensors. Neonate specifications are shown for OXIMAX MAX-N. Saturation accuracy will vary by sensor type.
**Applicability: OXIMAX MAX-A, MAX-AL, MAX-P, MAX-I, and MAX-N sensors.

Default Settings	
SpO2 (%)	HIGH: 100
SpO2 (%)	LOW: 90
Response Mode	2 (for Mode 2: Fast Response)
Sat Seconds	0
Audible Indicator	Pitch changes continuously with saturation; volume from 0 (off) to 9
Waveforms	Pulse plethysmograph waveform on LCD gain compensated
Sensor Connect/ Disconnect From Patient	Monitor detect attachment or disconnection of sensor from patient within 15 s
Pulse Detection attached to patient	Monitor will detect pulse or enter no signal state within 15 s of being
Loss of Pulse within 10 s	Monitor will detect loss of pulse from patient and enter no signal state
Sensor Light Source	
Wavelength	Infrared: 890 nm (nominal) Red: 660 nm (nominal)
Power Dissipation	52.5 mW (max)

Masimo SET SpO2

Measurement Range	
SpO2	1 to 100%
Pulse Rate	25 to 240 beats/min
Perfusion Range	0.02 to 20%
Accuracy and Motion Tolerance	
Saturation	
Without Motion - Adult/Ped*	70 to 100% ±2 digits
Without Motion - Neonate*	70 to 100% ±3 digits
With Motion - Adult/Ped/Neo**Ü	70 to 100% ±3 digits
Low Perfusioná	70 to 100% ±2 digits 0 to 69% unspecified
Pulse Rate	
Without Motion	25 to 240 beats/min ±3 digits
With Motion	normal physiologic range 25 to 240 beats/min ±5 digits

*The Masimo SETÆ SpO2 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.
**The Masimo SETÆ SpO2 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 non repetitive motion before 1 to 5 Hz at an amplitude of 2 to 3 cm 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.
ÜThe 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 neonateís foot at 2 to 4 cm against a laboratory co-oximeter and ECG monitor. This validation equals plus or minus, one standard deviation. Plus or minus one standard deviation encompasses 68% of the population.
áThe Masimo SET SpO2 parameter has been validated for low perfusion accuracy in bench top testing against a Biotek Index 2 simulator and Masimoís 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.

Masimo Sensor Accuracy		
Sensor Model		SpO2 Range 70% - 100%
LNOP		
LNOP-ADT		±2 digits
LNOP-ADT	Long	±2 digits
LNOP-PDT		±2 digits
LNOP-NEO		±3 digits
LNOP-NEO PT		±3 digits
LNOP-DCI (reusable)		±2 digits
LNOP-DCSC (reusable)		±2 digits
LNOP-DCIP (reusable)		±2 digits
NRI25 (reusable)		±2 digits
Resolution		
Saturation (% SpO2)	1%	
Pulse Rate (bpm)	1	
Low Perfusion Performance		
>0.02% Pulse Amplitude and % Transmission >5%	Saturation (% SpO2) ±2 digits Pulse Rate ±3 digits	

Interfering Substances	
Carboxyhemoglobin may erroneously increase readings. The level of increase is approximately equal to the amount of carboxyhemoglobin present. Dyes, or any substance containing dyes, that change usual arterial pigmentation may cause erroneous readings.	
Sensor Light Source	
Wavenlength	Infrared: 905 nm (nominal) Red: 660 nm (nominal)
Power Dissipation	Infrared: 22.5 mW (max) Red: 27.5 mW (max)

Default Settings	
SpO2 (%)	HIGH: 100
SpO2 (%)	LOW: 90
Sensitivity Mode	2 (for low perfusion-Default)
Averaging Time	12 seconds
FastSAT Mode	0 (for Off)

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GE Medical Systems
Information Technologies

We bring good things to life.

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