

Infinity[®] M300 Patient-worn Monitor

Infinity[®] M300 provides the performance of a full-size patient monitor, packaged in a patient-worn telemetry device for adult and pediatric patients. Built-in ACE[®] (Arrhythmia Classification Expert) and pacemaker detection algorithms enhance ECG processing and help to reduce false alarms.



Infinity M300 provides continuous standalone monitoring – even if the patient inadvertently moves out of the network coverage area. Two-way communication between Infinity M300 and the Infinity CentralStation facilitates wireless data exchange and signal integrity. Working together, the Infinity CentralStation and Infinity M300 enhance patient care management by providing fast data access, rapid assessment, decision support and clinical reporting.

FEATURES

- 3- to 6-wire ECG monitoring with TruST™ 12-lead
- Vital information access in color
- Alarms alerts and controls to support the telemetry workflow
- SpO₂ – ready in every device
- True battery management solution
- Wireless networking using commercial WiFi components

TECHNICAL DATA

SUPPORTED PARAMETERS

ECG

Available leads	3-wire: I, II, III
Adult/Pediatric	5-wire: I, II, III, aVR, aVL, aVF, V 6-wire: I, II, III, aVR, aVL, aVF, V, V+ 6-wire with Infinity TruST 12-lead: I, II, III, aVR, aVL, aVF, dV1, V2, dV2, dV3, dV4, V5, dV6
Leads analyzed	Any two user-selected ECG leads (simultaneously), or any one user-selected ECG lead
Detected events/rhythms	Asystole, Ventricular Fibrillation, Ventricular Tachycardia, Bradycardia, Ventricular Run, Accelerated Idioventricular Rhythm, Supraventricular Tachycardia, Ventricular Couplet, Ventricular Bigeminy, Tachycardia, Pause, Artifact, PVC/min
HR level alarm adjustment range	20 to 300 bpm
Measurement range	15 to 300 bpm
Accuracy	± 2 bpm or ± 1%, whichever is greater
Degree of protection against electrical shock	Type CF



Infinity M300
Patient-worn telemetry device

CONTINUING TECHNICAL DATA

Defibrillation protection	In accordance per IEC 60601-2-27
Bandwidth or resolution	Filter Monitoring: 0.5 – 40 Hz
Sweep speed	25 mm/sec \pm 10%
QRS detection	Amplitude: 0.5 – 5.0 mV Duration: 40 – 120 ms
Electrosurgery and cauterly	Not intended for use during ESU

Pacemaker

Detection leads	I and (II or III)
Detection amplitude	\pm 2 to \pm 900 mV
Detection width	0.1 to 2.0 ms
Precautions	Contains a tiny magnet which generates an extremely low static magnetic field of approximately 2 gauss at 12.7 mm (0.5 in) distance. Please refer to the manufacturer's Instructions for Use of any third party medical devices in the patient vicinity for compatibility.

ST Segment Analysis

Leads analyzed	3-wire: I, II, or III 5-wire: I, II, III, aVR, aVL, aVF, V 6-wire: I, II, III, aVR, aVL, aVF, V, V+ 6-wire with Infinity TruST 12-lead: I, II, III, aVR, aVL, aVF, dV1, V2, dV2, dV3, dV4, V5, dV6
ISO point	Default: - 28 msec
ST measurement point	Default: +80 msec
ST complex	Length: 900 msec (250 samples) Frequency response: 0.05 to 40 Hz
Update interval	15 seconds
ST level alarm adjustment range	-15.0 to 15.0 mm, -1.5 to 1.5 mV
ST accuracy	\pm 0.1 mm (\pm 0.01 mV)
ST measurement range	-15.0 to 15.0 mm, -1.5 to 1.5 mV
ST resolution	0.1 mm, 0.01 mV

Pulse Oximetry (optional)

Parameter display	Percentage of functional (oxygen-saturated) hemoglobin (%SpO ₂); pulse rate
Measuring method	Absorption-spectrophotometry
Measurement and display range	SpO ₂ : 1 – 100% Pulse rate: 30 – 250 bpm
Calibration range	70 – 100%
Display update period	2 seconds nominal
Maximum hold from previous update	30 seconds (in the event of artifact or other error)
SpO ₂ Alarm Adjustment Range	20 to 100%
Pulse Rate Alarm Adjustment Range	30 to 240 bpm

SpO₂ accuracy^{1,2,3,4}

0 to 69% not specified
70 to 100% sensor-specific as follows:

Masimo® LNOP® Sensors

LNOP adt, LNOP Pdt, LNOP neo, LNOP DCI, LNOP TC-I, LNOP DCIP, LNOP YI: \pm 3%
Pulse Rate Accuracy: \pm 3 bpm or \pm 3% (whichever is greater)
Low Perfusion Accuracy, SpO₂: \pm 2%
Low Perfusion Accuracy, Pulse Rate: \pm 3 bpm or \pm 3% (whichever is greater)

Masimo® LNCS® Sensors

LNCS DCI®, LNCS DCIP, LNCS Adtx, LNCS Pdtx, LNCS Inf: \pm 2%
Pulse Rate Accuracy: \pm 3 bpm or \pm 3% (whichever is greater)
Low Perfusion Accuracy, SpO₂: \pm 2%
Low Perfusion Accuracy, Pulse Rate: \pm 3 bpm or \pm 3% (whichever is greater)

Nellcor® Sensors

OxiMAX® MAX-A, OxiMAX MAX-AL, OxiMAX MAX-P, DS100A: \pm 3%

Dräger Sensors

MS16444 Disposable Foam Pedi,
MS16445 Disposable Foam Adt,
MS16449 Disposable Vinyl Adt,
MS16448 Disposable Vinyl Pedi,
MS13235 Reusable Sensor: \pm 2%

Notes:

¹ Since pulse oximeter measurements are statistically distributed, only about two-thirds of those measurements can be expected to fall within \pm 1 Arms of the value measured by a co-oximeter.

² These accuracies have been validated using blood samples obtained from healthy adult volunteers during induced hypoxia studies in the range of 70-100% SpO₂ against a laboratory co-oximeter and ECG monitor.

³ SpO₂ accuracies are expressed as \pm "X" digits between indicated saturation levels. Accuracy of the SpO₂ measurement is specified within 1 Arms of the value measured by a co-oximeter.

⁴ The pulse rate accuracy has been validated on healthy adult volunteers during induced hypoxia studies in the range of 70-100% SpO₂ against a laboratory co-oximeter and ECG monitor. This variation equals \pm 1 Arms of the pulse rate value measured by the ECG monitor.

User Interface

Controls	6 function keys: alarm pause, view screen, staff alert, record/mark event, up/down scroll
Alarms	Audible & visible alarm indication (user controlled) 3 severity levels: Life threatening, Serious, Advisory

Display

Size/viewing area	5.08 x 5.08 cm (2 x 2 in) diagonal LCD
Resolution	220 x 176 pixels

Communications

Network	IEEE 802.11b/g
Wireless encryption	WEP, WPA2 - Personal Mode
Radio power output	30 mW maximum

Physical Specifications

Size (H x W x D)	142.2 x 76.2 x 30.5 mm (5.6 x 3 x 1.2 in)
Weight	276.4 g (9.75 oz) with battery
Cooling	Convection
Connections	ECG, Communication port for SpO ₂ or Programming Cable, Bedside Charger, Central Charger

Electrical Specifications

Power source	Rechargeable 3.75 V lithium ion battery, available from Dräger
Battery operating time	ECG only: 17 to 19 hours ECG + continuous SpO ₂ : 14 to 16 hours Operation time varies according to use of display, alarm alerts, and wireless environment (roaming)
Battery recharging time	Using Bedside Charger to 100%, approximately: 0 to 25% = 2 hours 0 to 50% = 4 hours 0 to 75% = 6 hours 0 to 100% = 8 hours Using Central Charger to 100%, approximately: 0 to 25% = 40 minutes 0 to 50% = 1.5 hours 0 to 75% = 2 hours 0 to 100% = 3 hours

Environmental Requirements**Temperature**

Operating	0° C to 40° C (32° F to 104° F)
Storage	-20° C to 60° C (- 4° F to 140° F)

Humidity (non condensing)

Operating	10% to 85%
Storage	10% to 85%

Atmospheric pressure

Operating	64.7 to 106 kPa
Storage	50 to 106 kPa
Free fall	IEC 60068-2-32, Procedure 1 Height of fall: 1 m Number of falls: 1 on each of six surfaces
Protection against water ingress	IPX7, temporary immersion

Standards

Compliances	IEC 60601-1:1988 + A1:1991 + A2:1995, IEC 60601-1-2:2001, IEC 60601-2-27:2006, IEC 60601-2-49:2001, ANSI/AAMI EC13:2002(R)2007.
-------------	--

**Central Charger****Bedside Charger**

CONTINUING TECHNICAL DATA

INFINITY M300 CENTRAL CHARGER

Physical Specification

Size (H x W x D)	520.7 x 215.9 x 190.5 mm (20.5 x 8.5 x 7.5 in)
Weight	6.5 kg (14.4 lb)
Cooling	Convection
Connections	Up to ten (10) Infinity M300 devices

Electrical Specifications

Input voltage	92 – 264 VAC
Input frequency (Hz)	50/60 Hz
Protection class	Class 1
Mode of operation	Continuous

Environmental Requirements

Temperature

Operating	10° C to 45° C (50° F to 113° F)
Storage	- 40° C to 70° C (- 40° F to 158° F)

Humidity (non condensing)

Operating	10% to 95%
Storage	10% to 95%

Atmospheric pressure

Operating	70 kPa to 106 kPa
Storage	50 kPa to 106 kPa

Protection against water ingress	IPX1, dripping water
----------------------------------	----------------------

Standards

Compliances	IEC 60601-1, IEC 60601-1-2
-------------	----------------------------

INFINITY M300 BEDSIDE CHARGER

Physical Specifications

Size (H x W x D)	45.72 x 162.56 x 99.06 mm (1.8 x 6.4 x 3.9 in)
Weight	224 g (7.9 oz)
Cooling	Convection
Connections	One (1) Infinity M300

Electrical Specifications

Input voltage	92 – 264 VAC
Input frequency (Hz)	50/60 Hz ± 5%
Protection class	Class 2
Mode of operation	Continuous

Environmental Requirements

Temperature

Operating	0° C to 40° C (32° F to 104° F)
Storage	- 20° C to 60° C (- 4° F to 140° F)

Humidity (non condensing)

Operating	10% to 85%
Storage	10% to 85%

Atmospheric pressure

Operating	64.7 kPa to 106 kPa
Storage	50 kPa to 106 kPa

Protection against water ingress	IPX4, splashing water
----------------------------------	-----------------------

Free fall	IEC 60068-2-32, Procedure 1
-----------	-----------------------------

Standards

Compliances	IEC 60601-1, IEC 60601-1-2
-------------	----------------------------

ORDERING INFORMATION

Infinity M300	MS18501
Contact Dräger for ordering details.	

HEADQUARTERS

Drägerwerk AG & Co. KGaA
Moislinger Allee 53–55
23558 Lübeck, Germany

www.draeger.com

REGION EUROPE CENTRAL
AND EUROPE NORTH

Dräger Medical GmbH
Moislinger Allee 53–55
23558 Lübeck, Germany
Tel +49 451 882 0
Fax +49 451 882 2080
info@draeger.com

REGION EUROPE SOUTH

Dräger Médical S.A.S.
Parc de Haute
Technologie d'Antony 2
25, rue Georges Besse
92182 Antony Cedex, France
Tel +33 1 46 11 56 00
Fax +33 1 40 96 97 20
dlmfr-contact@draeger.com

REGION MIDDLE EAST, AFRICA,
CENTRAL AND SOUTH AMERICA

Dräger Medical GmbH
Branch Office Dubai
Dubai Healthcare City, P.O. Box 505108
Dubai, United Arab Emirates
Tel + 971 436 24 762
Fax + 971 436 24 761
contactuae@draeger.com

REGION ASIA / PACIFIC

Dräger Medical
South East Asia Pte Ltd
25 International Business Park
#04-27/29 German Centre
Singapore 609916, Singapore
Tel +65 6572 4388
Fax +65 6572 4399
asia.pacific@draeger.com

REGION NORTH AMERICA

Draeger Medical, Inc.
3135 Quarry Road
Telford, PA 18969-1042, USA
Tel +1 215 721 5400
Toll-free +1 800 437 2437
Fax +1 215 723 5935
info.usa@draeger.com

Manufacturer:

Dräger Medical Systems, Inc.
Telford, PA 18969, USA
The quality management system at
Dräger Medical Systems, Inc. is
certified according to ISO 13485,
ISO 9001 and Annex II.3 of Directive
93/42/EEC (Medical devices).