

IntelliVue MP5 Patient Monitor

Philips M8105A Technical Data Sheet

The MP5 portable patient monitor is compact in size, ergonomic, and modular in design. It shares a common user interface and technological platform with the Philips IntelliVue MP20-MP90 patient monitors.

The MP5 can be connected to one of the MP20 to MP90 monitors and used like a Multi-Measurement Module (MMS), providing monitoring continuity in transport situations.

The monitor is highly customizable. Dedicated configurations are available for the anesthesia, critical and cardiac, and neonatal care environments.

The IntelliVue series offers a complete monitoring solution that is flexible and modular, designed to suit a broad spectrum of monitoring needs.

Measurement Features

- Compact, rugged, lightweight monitor with built in measurements
- ECG monitoring using any combination of three to 10 electrodes.
- 12-lead ECG monitoring with five electrodes using the EASI method or with 10 electrodes using the conventional method.
- Multi-lead arrhythmia and ST segment analysis at the bedside on all available leads.
- Respiromics Mainstream or Sidestream CO₂
- FAST SpO₂ for accurate performance even with low perfusion.
- Invasive Pressure and Temperature measurement
- Choice of either auscultatory or intra-arterial measurements as a measurement reference for Non-Invasive Blood Pressure

PHILIPS

- Predictive temperature measurement, providing temperature readings within 6 to 15 seconds.
- Built in Recorder
- Telemetry devices can be directly connected to monitor telemetry data (ECG/SpO₂) on the MP5 screen (Telemetry as a parameter (TAAP)).
- The MP5 with IntelliVue Instrument Telemetry can be declared as a telemetry device at the Information Center and paired with a monitor.
- The monitor can operate using battery power for up to four hours with basic monitoring configuration and up to three hours with extended monitoring configuration, to let you safely and easily monitor patients during in-hospital transfer. See “Battery Specifications” on page 7.

Usability Features

- Touchscreen as input device.
- Intuitive user interface.
- Simple menu hierarchy gives fast access to all basic monitoring tasks.
- Patient data management with tabular and graphic trends.
- Settings “Profiles” for rapid case turnover.
- Patented automatic alarm limits help clinicians provide care more efficiently.
- Neonatal Event Review keeps a record of rapidly changing condition of neonatal patients.
- Bed-to-bed overview provides clinicians with an overview of all the patient beds in their care.
- 8.4" TFT flat panel display with SVGA (800 x 600) resolution, wide viewing angle, large numerics, permanently visible alarm limits, and up to four real-time waves.
- Capable of functioning in a wireless infrastructure (WLAN or IIT)
- Timer application allows you to set timers to notify you when a specific time period has expired.

Intended Use

The monitor is intended to be used for monitoring and recording of, and to generate alarms for, multiple physiological parameters of adults, pediatrics, and neonates in a hospital environment and during patient transport inside and outside of hospitals. The monitor is intended for use by health care professionals.

The monitor is only for use on one patient at a time. It is not intended for home use. Not a therapeutic device.

Rx only: U.S. Federal Law restricts this device to sale by or on the order of a physician.

ST segment monitoring is intended for use with adult patients only and is not clinically validated for use with neonatal and pediatric patients.

The Predictive Temperature unit is intended for use with adult and pediatric patients in a hospital environment.

The ECG measurement is intended to be used for diagnostic recording of rhythm and detailed morphology of complex cardiac complexes (according to AAMI EC 11).

The derived measurement Pulse Pressure Variation (PPV) is intended for use with sedated patients receiving controlled mechanical ventilation and mainly free from cardiac arrhythmia. The PPV measurement has been validated only for adult patients.

Hospital Environment:

The monitor is suitable for use in all medically used rooms which fulfill the requirements regarding electrical installation according to IEC60364-7-710 “Requirements for special installations or locations - Medical locations”, or corresponding local regulations.

EMC Environment:

The following measurements and system interfaces are, in addition, suitable for use in establishments directly connected to the public low-voltage supply network that supplies buildings used for domestic purposes:

- ECG/Respiration, NBP, SpO₂, Pressure, Temperature, CO₂ (only Mainstream Sensor M2501A)
- LAN, Video Out, Battery, Nurse Call, RS232, and recorder interfaces.

The monitor is available as a standalone or networked solution

Upgradability

The MP5 monitor allows new capabilities to be added in the future as your monitoring requirements evolve. This upgradability gives the security of knowing that the monitors can be enhanced and updated as practices and technologies advance, and it protects long-term investments.

Main Components

Monitor

The monitor has a color 8.4” LCD TFT display with a wide viewing angle, providing high resolution waveform and data presentation.

The display, processing unit, measurements and power supply are integrated into one device.

User Interface

The user interface is designed for fast and intuitive operation. The color graphical user interface ensures that clinicians quickly feel at ease using the monitor.

Configurable SmartKeys with intuitive icons allow monitoring tasks to be performed quickly and easily, directly on the monitor screen.

Waves and numerics are color-coded.

The monitor displays up to four measurement waves simultaneously. For 12-lead ECG monitoring it can display 12 real-time ECG waves, with a rhythm strip and all ST values.

The MP5 monitor is supplied with a resistive touchscreen.

Simulated Keyboard

If alpha or numeric data entry is required, for example to enter patient demographics, an on-screen keyboard will automatically appear on the screen.

Mounting

The mounting options available enable flexible, space saving placement of the monitors for an ergonomic work space. The monitor is shipped with a low cost mounting plate if not specified otherwise.

Application Features

Critical and Cardiac Care Features

- The monitor performs multi-lead **arrhythmia detection** analysis on the patient's ECG waveform at the bedside. It analyzes for ventricular arrhythmias, calculates heart rate, and generates alarms, including asystole, bradycardia, and ventricular fibrillation.
- Up to 12 leads of **ST segment analysis** can be performed on adult patients at the bedside, measuring ST segment elevation and depression and generating alarms and events. The user can trend ST changes, set high and low alarm limits, and set both ST and isoelectric measurement points. Using ST Snippets, one-second wave segments can be compared with a baseline segment for each measured ST lead.
- **QT/QTc interval monitoring** provides the measured QT interval, the calculated heart-rate corrected QTc value and a Δ QTc value, which tracks variation in the QT interval in relation to a baseline value.
- optional **ST Map** application shows ST changes over time in two multi-axis spider diagrams.
- optional **12-lead ECG** data can be measured, using either the EASI placement method with five standard electrodes or conventional electrode placement with 10 electrodes.¹
- 12 realtime ECG waveforms can be displayed simultaneously.
- **FAST-SpO₂**, using Fourier Artifact Suppression Technology, performs accurately even in cases with low perfusion.

1. EASI-derived 12-lead ECGs and their measurements are approximations to conventional 12-lead ECGs. As the 12-lead ECG derived with EASI is not exactly identical to the 12-lead conventional ECG obtained from an electrocardiograph, it should not be used for diagnostic purposes.

- Choice of sidestream or mainstream **CO₂ monitoring** for high quality measurements with intubated and non-intubated patients.
- **Telemetry devices** (TRx4841A/TRx4851A TRx/TRx+ IntelliVue Transceiver) can be connected via a cable to the MP5 to monitor telemetry data (ECG/SpO₂) on the MP5 screen.
- The optional **Drug Calculator** helps you to manage intravenous (IV) drug infusions by calculating drug dose, rate, amount, volume, concentration, and standardized rate.

Anesthesia Features

- The **IntelliVue G1** and **G5²** anesthetic gas modules measure up to 5 respiratory gases and one or two agents and produce waves and numerics for display on the MP5 monitor.
- **Screens** and **Profiles** provide flexible viewing of patient information during different procedures or phases of an anesthesia case.

Neonatal Monitoring Features

- The **OxyCRG** screen provides a simultaneous presentation of up to three trends:
 - beat-to-beat heart rate (btbHR)
 - an oxygenation measurement trend
 - compressed respiration wave.This customized display gives clinicians a convenient overview of the neonatal patient's most important vital signs, helping them to identify significant events. Continuous OxyCRG recordings can be made on the built-in recorder, and reports can be printed on locally or centrally-connected printers.
- Optional **Neonatal Event Review (NER)** is optimized for monitoring neonatal patients. For each event, an episode of four minutes of data sampled four times a second is stored, to help you keep a record of the rapidly-changing condition of neonatal patients. Combi-events correlate apnea events with bradycardia and/or desaturations.

Ease of Use

- **Screen layouts** are easily adjustable, allowing flexible display of measurement information.
- Temperature, height, and weight can be configured either in metric or imperial units. Pressure measurements can be displayed in kPa or mmHg. Gases can be displayed in kPa, mmHg.

Trends

- The **trend database** stores patient data from up to 16 measurement numerics. The measurement information can be sampled every 12 seconds, one minute, or five minutes, and stored for a period ranging from four to 48 hours.

ProtocolWatch

- ProtocolWatch allows clinicians to run clinical protocols that can monitor developments in the patient's condition. The SSC Sepsis
 - 2. IntelliVue G5 in combination with MP5 is only available in the USA

Protocol runs on the ProtocolWatch application and is used in screening for severe sepsis.

Transport Features

- The monitor's portable design means it can be used for in-hospital transport: a basic monitor weighs 4 kg.
- The monitors can operate using battery power for up to four hours, to let you safely and easily monitor patients during procedures or in-hospital transfer.
- The monitor's network capability means that it is ready for use as an integrated part of the hospital system.
- Specially-designed mounting solutions let you quickly disconnect the monitor for transport and reconnect to the mount after transport.
- The Universal Admit, Discharge and Transfer (ADT) feature means that all ADT information is shared between the networked monitor and the Information Center. Information need only be entered once.

Patient Data Documentation

- An extensive range of **Patient Reports** can be printed:
 - Event Review and Episode Reports
 - OxyCRG Reports
 - 12-lead ECG Reports
 - Alarm Limit Reports
 - Vital Signs
 - Graphic Trends
 - Drug Calculator Reports
 - Realtime Wave ReportsReport templates can be defined in advance, enabling print-outs tailored to each hospital's specific requirements to be started quickly. Reports can be printed on locally or centrally-connected printers, and they can be initiated manually or automatically at user-defined intervals.

Alarms

The alarm system can be configured to present either the traditional HP/Agilent/Philips alarm sounds or sounds compliant with the draft ISO/IEC 9703-2 Standard.

Alarm limits are permanently visible on the main screen. The Alarm Limits page provides a graphic depiction of alarm limits in relation to the currently monitored measurement values and lets you adjust alarm limits. It also lets you preview wide and narrow automatic alarm limits before you apply them.

When an alarm limit is exceeded, it is signalled by the monitor in the following ways:

- an alarm tone sounds, graded according to severity
- an alarm message is shown on the screen, color-coded according to severity
- the numeric of the alarming measurement flashes on the screen
- alarm lamps flash for red and yellow alarms and are illuminated for technical INOPs

If the monitor is connected via a network to a central monitoring station, alarming is simultaneous at the monitor and at the Information Center.

The nurse call relay has active closed contacts and a user-definable delay time.

Alarms are graded and prioritized according to severity:

- **Red Alarms***** identify a potentially life threatening situation for a patient .
- **Yellow Alarms**** indicate conditions violating preset vital signs limits.
- **Technical Alarms (INOPS)** are triggered by signal quality problems, equipment malfunction or equipment disconnect. The Silence/Pause Alarms function (equivalent to Silence/Suspend with previous monitor generations) allows the user to switch off alarm tones with one touch. All alarms can be paused for a period of one, two, three, five, or 10 minutes or turned off indefinitely. Alarm strip recordings are available on the optional built-in recorder or on a centrally-connected recorder. Patented automatic alarm limits automatically adapt the alarm limits to the patient's currently measured vital signs within a safe margin defined individually for each patient. Visual and/or audible latching and non-latching alarm handling is available.

Profiles

Profiles are predefined configuration settings for Screens, measurement settings, and monitor settings. Each Profile can be designed for a specific application area and patient category, for example OR adult, or ICU neonatal. Profiles enable a quick reaction to patient and care location changes: activating a Profile with a particular patient category (Adult, Pediatric or Neonatal) automatically applies suitable alarm and safety limits and saves time usually spent carrying out a complete set-up procedure.

Profiles can be created directly on the monitor or remotely on a personal computer and transferred to the monitor using the IntelliVue Support Tool. A selection of Profiles for common monitoring situations is provided with the monitor. These profiles can be changed, added to, renamed, or deleted.

Other Bed Overview Capability

The alarm status of beds in the same Care Group on the hospital network can be permanently displayed on the screen of each monitor in the Care Group. The user can also view measurement data from all other monitors connected to the hospital network.

Service Features

- The Support Tool helps technical personnel to
 - carry out configuration, upgrades and troubleshooting via the network, or on an individual monitor
 - share configuration settings between monitors
 - back up the monitor settings.
- A password-protected Service Mode ensures that only trained staff can access service tests and tasks.
- The Configuration Mode is password-protected and allows trained users to customize the monitor configuration.

Device Connections

The monitor can be connected to:

- an Information Center (for example M3150B)
- a PC
- IntelliVue G1/G5 gas module
- TRx4841A/TRx4851A TRx/TRx+ IntelliVue Transceiver

Network Interface

The network interface provides the system with networking capability via a wired or wireless network connection.

Wireless Network

The monitor can function within a wireless infrastructure based on an IEEE 802.11 a/b/g network in the 2.4 GHz / 5 GHz bands (ISM). Additionally, the monitor can function within a telemetry infrastructure compatible with the Philips Cellular Telemetry System (CTS) in the WMTS and ISM bands. Additional components are required to complete the system. Please refer to the M3185A IntelliVue Clinical Network Technical Data Sheet for further information.

MIB/RS-232 Interface

The monitor has an optional serial MIB/RS-232 interface board with a fully-isolated port. The port can be configured to be used for:

- input for connection to a touchscreen
- data export using a computer interface, to an automated anesthesia record keeper or a personal computer (not available in all geographies)
- connection to an IntelliVue G1/G5 anesthetic gas module

Companion Mode Interface

The Companion Mode interface is used to connect the MP5 to a host monitor of the IntelliVue MP20-90 family.

Monitor Specifications

Safety Specifications

The monitor complies with the Medical Device Directive 93/42/EEC (CE₀₃₆₆) and with IEC 60601-1:1988 + A1:1991 + A2:1995; EN60601-1:1990 + A1:1993 + A2:1995; UL 60601-1:2003; CAN/CSA C22.2#601.1-M90; JIS T 0601-1:1999; IEC 60601-1-1:2000; EN 60601-1-1:2001.

All applied parts are Type CF unless otherwise specified. They are protected against damage from defibrillation and electrosurgery.

The possibility of hazards arising from software errors was minimized in compliance with ISO 14971:2000, EN60601-1-4:1996 + A1:1999 and IEC 60601-1-4:1996 + A1:1999.

The monitor complies with the EMC standards IEC 60601-1-2:2001; EN 60601-1-2:2001

This ISM device complies with Canadian ICES-001. Cet appareil ISM est conforme a la norme NMB-001 du Canada.

The MP5 patient monitor, with the following measurements and interfaces:

- ECG/Respiration, NBP, SpO₂, Pressure, Temperature, CO₂ (only Mainstream Sensor M2501A)
- LAN, Video Out, Battery, Nurse Call, RS232, and recorder interfaces

can be used in a transport environment such as a road ambulance, airplane or helicopter. For this purpose the monitor fulfills the following additional mechanical, EMC and environmental requirements:

- **Shock Tests** according to IEC TR 60721-4-7, Class 7M3. Test procedure according to IEC/EN 60068-2-27 (peak acceleration up to 100g).
- **Random Vibration** according to IEC TR 60721-4-7, Class 7M3. Test procedure according to IEC/EN 60068-2-64 (RMS acceleration 5g).
- **Sinusoidal Vibration** according to IEC TR 60721-4-7, Class 7M3. Test procedure according to IEC/EN 60068-2-6 (acceleration up to amplitude 2g).
- **Bump Test** according to IEC/EN60068-2-29 (peak acceleration 15g, 1000 bumps).
- **Free Fall Test** according to EN1789 (covers also IEC TR 60721-4-7 and Class 7M3). Test procedure according to EN 60068-2-32 (height 0.75 m).
- Specification for degrees of protection provided by enclosures according to **IEC/EN 60529: IP 32**
- **EN 1789 +A1:2003** Medical vehicles and their equipment - Road ambulances (chapter 6 - Medical Devices).
- **Radiated susceptibility 20 V/m** according to EN ISO 9919 (SpO₂)

and EN ISO 21647 (CO2).

- **Altitude Range** from -500 to 3000 m operating and -500 to 4600 m storage and transportation.
- Extended radiated susceptibility tests

The MP5 patient monitor with its out-of-hospital parameter set provides a general immunity level of 20 V/m with only few restrictions. Details are as listed below:

- GSM 900: Immunity at 900 MHz (uplink mobile phone), 20V/m (ECG:10V/m), duty cycle 1:8
- GSM 1800: Immunity at 1800 MHz (uplink mobile phone), 20V/m, duty cycle 1:8.
- DECT: Immunity at 1800 MHz (digital cordless phone), 20V/m, duty cycle 1:24
- AM: 1 kHz Immunity from 80 MHz to 1.0 GHz (any radio communication unit, broadcasting and TV transmitter), 20V/m, modulation factor 80%. (ECG: 20 V/m except 600-950 MHz where it is 10V/m and Temperature which holds 3V/m over the full range)

Temperature measurement accuracy may be compromised in the presence of strong electromagnetic fields (>3V/m) in certain small frequency bands.

- **Magnetic Field emission** according to MIL STD 461E, Chapter RE101: Radiated emissions, magnetic field, 30 Hz to 100 kHz. Limit class: Army.
- **Magnetic Field susceptibility:** Radiated susceptibility, magnetic field, 50, 60 and 400 Hz, 18 μT(15 A/m)
- **Operating ambient temperature** testing over the range from 0 to 40 °C (32 to 100 °F).
- **Operating ambient humidity** testing up to 95% RH at 40 °C (100 °F), non condensing.

Physical Specifications

| Product | Max Weight | W x H x D |
|--|-------------------|--|
| M8105A IntelliVue MP5 (including battery, without options) | 4 kg ±5% (8.8 lb) | < 259 x 248 x 186 mm (10.2 x 9.76 x 7.32 in) |

Environmental Specifications

| Item | Condition | Range |
|---|---|--|
| Temperature Range | Operating | 0 to 40°C (32 to 104 °F) |
| | Storage (without battery) and Transport | -20 to 60°C (-4 to 140 °F) |
| Temperature Range with IntelliVue 802.11 Bedside Adapter or IntelliVue Instrument Telemetry Wireless Network or when charging the battery | Operating | 0 to 35°C (32 to 95°F) |
| Temperature Range with Predictive Temperature Unit | Operating | 10 to 40°C (50 to 104 °F) |
| | Storage | -20 to 50°C (-4 to 120 °F) |
| Humidity Range | Operating | 15% to 95% Relative Humidity (RH) (non condensing) |
| | Storage and Transport | 5% to 90% Relative Humidity (RH) |
| Altitude Range | Operating | -500 m to 3000 m (10000 ft) |
| | Storage and Transport | -500 m to 4600 m (15000 ft) |
| Ingress Protection | Monitor without the Predictive Temperature unit | IP32 |
| | Monitor with the Predictive Temperature unit | IPX1 |

Performance Specifications

| Monitor Performance Specifications | | |
|---|--|--|
| Power Specifications | Power consumption | < 40W average, <65W peak |
| | Line Voltage | 100 to 240 V ~ |
| | Current | 1.3 to 0.7A |
| | Frequency | 50/60 Hz |
| SVGA Display 8.4 inch | Resolution | 800 x 600 |
| | Refresh rate | 60 Hz |
| | Useful screen | 170.4 x 127.8 mm |
| | Pixel size | 0.213 x 0.213 mm |
| Sweep Speeds | 6.25, 12.5, 25 and 50 mm/s with ±5% accuracy | |
| Indicators | Alarms Off | red LED |
| | Alarms | red/yellow/cyan LED |
| | On/Standby/Error | green/red LED |
| | AC Power | green LED |
| | Battery | red-yellow-green LED |
| Sounds | Audible feedback for user input. Prompt tone. Two different QRS tones, SpO ₂ modulation tone. Four different alarm sounds. Remote tone for alarms on other beds in network. Tone for Timer expired. | |
| <p>Trends: 12, 16, 24 or 32 numerics @ 12 sec, 1 minute, 5 minute resolution. Multiple choices of number of numerics, resolution and duration depending on application area.</p> | | |
| High-Res Trend Waves | Measurements OxyCRG | HR, SpO ₂ , Resp |
| | Resolution | Measurement samples are taken at a resolution of four samples per second |
| | Update Speed | waves are drawn at a speed of 3 cm/minute |

| Monitor Performance Specifications | | |
|--|---|---|
| Events | Information | trigger condition and time, event classification and associated detailed view of episode data |
| | Episode data | 4 minutes of high resolution trend |
| Alarm Signal | System delay | less than 3 seconds |
| | Pause duration | 1,2,3 minutes or infinite, depending on configuration |
| | Extended alarm pause | 5 or 10 minutes |
| Review Alarms Window | Information: all alarms / inops, main alarms on/off, alarms acknowledged and time of occurrence | |
| | capacity | 100 items |
| Real Time Clock | Range: from: January 1, 1997, 00:00 to: December 31, 2080, 23:59 | |
| | Accuracy: < 2 seconds per day (typically) | |
| | Hold Time: infinite if powered by AC; otherwise at least 48 hours (typical: > 72 hours) | |
| Buffered Memory | Contents: Active settings, trends, snapshots, events, review alarms | |
| | Hold Time: infinite if powered by AC; otherwise at least 48 hours (typical: > 72 hours) | |
| Restart time: After power interruption, an ECG wave will be shown on the display after 30 seconds maximum. | | |

Battery Specifications

A battery can be used to operate the monitor.

Special Philips high-power batteries M4605A 10.8 V 6000mAh Lithium Ion Battery

- PN 989 8031 31111 (removable)
- Weight: 480g per battery
- Status LEDs indicate charge status of battery

Battery Operating Time:

A new and fully charged battery :

- with basic monitoring configuration (automatic brightness reduction, ECG/RESP and SpO₂ measurements in use, NBP every 15 minutes):4 hours
- with extended monitoring configuration (maximum brightness, ECG/RESP, SpO₂, PRESS/TEMP and CO₂ measurements in use, NBP every 15 minutes, recorder every 15 minutes): 3 hours

Battery Charge Time:

- when monitor is switched off: about 4 hours when monitor is switched on and fully functional: 5 hours and above, depending on configuration (in some configurations the battery may not completely recharge in the monitor, in this case the M8043A Smart Battery Charger should be used).

Interface Specifications

| Monitor Interface Specifications | | |
|----------------------------------|----------------|---|
| Network | Standard | IEEE 802.3 10-Base-T |
| | Connector | RJ45 (8 pin) |
| | Isolation | 1.5 kV |
| Companion Mode Interface | Connectors | Female ODU (Proprietary) |
| | Power Sync. | RS-422 compliant input 78.125 kHz (typical) |
| | LAN signals | IEEE 802.3 10-Base-T compliant |
| | Serial signals | RS-422 compliant |
| | Local signals | Internal use only |
| MIB/RS232 | Standard | IEEE 1073-3.2-2000 |
| | Connectors | RJ45 (8 pin) |
| | Mode | BCC (Rx/D/TxD cross over) |
| | Power | 5V +/- 5%, 100mA (max.) |
| | Isolation | 1.5kV |

| Monitor Interface Specifications | | |
|---|--|---|
| ECG Sync Pulse Mode | Pulse Width | 100 +/- 10ms (high) |
| | Delay from R-wave peak to start of pulse | 20 ms maximum per AAMI EC13 |
| | Output voltage swing | +/- 5V minimum |
| 802.11 Bedside Adapter | Wireless Technology | IEEE 802.11 a/b/g |
| | Frequency Band | 2.4 GHz and 5 GHz ISM |
| Internal WMTS Adapter (US only) | Technology | compatible with Philips Cellular Telemetry System (CTS) cellular infrastructure |
| | Frequency Band | WMTS, 1395-1400 MHz and 1427-1432 MHz |
| Internal ISM Adapter | Technology | compatible with Philips Cellular Telemetry System (CTS) cellular infrastructure |
| | Frequency Band | 2.4 GHz ISM |
| Basic Nurse Call Relay | Connector | 3.5 mm phone jack, active closed contact only |
| | Contact | <= 100 mA, <= 24 V DC |
| | Isolation | 1.5 kV |
| | Delay | < (Configured Latency + 0.5 sec) |
| ECG Output/Marker Input (1/4" stereo phone jack with tip, ring, sleeve) | | |
| General | Connector | 1/4" phone each with tip, ring, sleeve |
| | Isolation | 500 V |

| Monitor Interface Specifications | | |
|----------------------------------|------------------------|--|
| ECG Output (ring, tip) | Signal Gain | 320 to 3200 in 19 steps |
| | Full Scale on Display | Signal Gain * measured ECG Voltage |
| | Gain Error | <20% |
| | Baseline Offset | <150mV |
| | Bandwidth | 1 to 80Hz |
| | Output Impedance | ECG Output (ring): <2.2kΩ±20% ECG Output/Marker Input (tip) <2.5kΩ ±20% |
| | Signal delay | ≤30ms |
| Marker Input Requirements (tip) | Signal Type | 0 to -12V, negative edge pulse |
| | Pulse Source Impedance | <7kΩ |
| | Pulse Fall Time | <100µs |
| | Pulse Duration | >4ms |

Measurement Specifications

ECG/Arrhythmia/ST

Complies with IEC 60601-2-25:1993 + A1:1999 /EN60601-2-25:1995 + A1:1999, IEC 60601-2-27/EN60601-2-27:1994, IEC 60601-2-51:2003 /EN 60601-2-51:2003 and AAMI EC11/EC13:1991/2002.

| ECG/Arrhythmia/ST Performance Specifications | | |
|--|-------------|---|
| Cardiotach | Range | Adult/pedi: 15 to 300 bpm Neo range: 15 to 350 bpm |
| | Accuracy | ±1% of range |
| | Resolution | 1 bpm |
| | Sensitivity | ≥200 µV _{peak} |
| PVC Rate | Range | 0 to 300 bpm |
| | Resolution | 1 bpm |

| ECG/Arrhythmia/ST Performance Specifications | | |
|--|--------------------------|---|
| ST Numeric | Range | -20 to +20 mm |
| | Accuracy | ±0.5 mm or 15%, whichever is greater |
| | Resolution | 0.1 mm |
| Sinus and SV Rhythm Ranges | Brady | Adult: 15 to 60 bpm Pedi: 15 to 80 bpm Neo: 15 to 90 bpm |
| | Normal | Adult: 60 to 100 bpm Pedi: 80 to 160 bpm Neo: 90 to 180 bpm |
| | Tachy | Adult: > 100 bpm Pedi: >160 bpm Neo: >180 bpm |
| Bandwidth | Diagnostic Mode | Adult/neo/pedi: 0.05 to 150Hz |
| | Extended Monitoring Mode | Neo/pedi: 0.5 to 150Hz |
| | Monitoring Mode | Adult: 0.5 to 40Hz Neo/pedi: 0.5 to 55Hz |
| | Filter Mode | Adult/neo/pedi: 0.5 to 20Hz |
| Differential Input Impedance | | >2MΩ RA-LL leads (Resp) >5MΩ at all other leads (at 10Hz including patient cable) |
| Common Mode Rejection Ratio | | Diagnostic mode: >86 dB (with a 51 kΩ/47 nF imbalance). Filter mode: >106 dB (with a 51 kΩ/47 nF imbalance). |
| Electrode Offset Potential Tolerance | | ±500mV |
| Auxiliary Current (Leads off Detection) | | Active electrode: <100 nA Reference electrode: <900 nA |
| Input Signal Range | | ±5 mV |

| ECG/ Arrhythmia/ST Alarm Specifications | Range | Adjustment |
|--|---|---|
| HR | 15 to 300 bpm maximum delay: 10 seconds according to AAMI EC 13- 1992 standard | Adult:1 bpm steps (15 to 40 bpm) 5 bpm steps (40 to 300 bpm) Pedi/Neo:1 bpm steps (15 to 50 bpm) 5 bpm steps (50 to 300 bpm) |
| Extreme Tachy | Difference to high limit 0 to 50 bpm | 5 bpm steps |
| | Clamping at 150 to 300 bpm | 5 bpm steps |
| Extreme Brady | Difference to low limit 0 to 50 bpm | 5 bpm steps |
| | Clamping at 15 to 100 bpm | 5 bpm steps |
| Run PVCs | 2 PVCs | Not adjustable by user |
| PVCs Rate | 1 to 99 PVCs/ minute | 1 PVC |
| Vent Tach HR | 20 to 300 bpm | 5 bpm |
| Vent Tach Run | 3 to 99 PVCs/ minute | 1 PVC |
| Vent Rhythm Run | 2 to 99 PVCs/ minute | 1 PVC |
| SVT HR | 120 to 300 bpm | 5 bpm |
| SVT Run | 3 to 99 SV beats | 1 SV beat |
| ST High | -19.8 to +20 mm | 0.2 mm |
| ST Low | -20 to +19.8 mm | 0.2 mm |

| ECG/Arrhythmia/ST Supplemental Information as required by AAMI EC11/13 | | |
|---|---|--|
| Respiration Excitation Waveform | | Sinusoidal signal, 260 μ A, 39 kHz |
| Noise Suppression | | RL drive gain 44 dB max., max. voltage 1.8 Vrms |
| Time to Alarm for Tachycardia | Vent Tachycardia 1mV _{pp} , 206 bpm | Gain 0.5, Range 6.5 to 8.4 seconds, Average 7.2 seconds |
| | | Gain 1.0 Range 6.1 to 6.9 seconds, Average 6.5 seconds |
| | | Gain 2.0, Range 5.9 to 6.7 seconds, Average 6.3 seconds |
| | Vent Tachycardia 2 mV _{pp} , 195bpm | Gain 0.5, Range 5.4 to 6.2 seconds, Average 5.8 seconds |
| | | Gain 1.0, Range 5.7 to 6.5 seconds, Average 6.1 seconds |
| | | Gain 2.0, Range 5.3 to 6.1 seconds, Average 5.7 seconds |
| Tall T-Wave Rejection Capability | | Exceeds ANSI/AAMI EC 13 Sect. 3.1.2.1(c) minimum recommended 1.2 mV T-Wave amplitude |
| Heart Rate Averaging Method | | Three different methods are used: Normally, heart rate is computed by averaging the 12 most recent RR intervals. For runs of PVCs, up to 8 RR intervals are averaged to compute the HR. If each of 3 consecutive RR intervals is greater than 1200 ms (that is, rate less than 50 bpm), then the 4 most recent RR intervals are averaged to compute the HR. |
| Response Time of Heart Rate Meter to Change in Heart Rate | | HR change from 80 to 120 bpm: Range: [6.4 to 7.2 seconds] Average: 6.8 seconds HR change from 80 to 40 bpm: Range: [5.6 to 6.4 sec] Average: 6.0 seconds |

| ECG/Arrhythmia/ST Supplemental Information as required by AAMI EC11/13 | |
|--|--|
| Heart Rate Meter Accuracy and Response to Irregular Rhythm | Ventricular bigeminy: 80 bpm Slow alternating ventricular bigeminy: 60 bpm Rapid alternating ventricular bigeminy: 120 bpm Bidirectional systoles: 90 bpm |
| Accuracy of Input Signal Reproduction | Methods A and D were used to establish overall system error and frequency response. |

| Respiration Alarm Specifications | Range | Adjustment | Delay |
|----------------------------------|------------------|----------------|-------|
| Apnea Alarm | 10 to 40 seconds | 5 second steps | |

Respiration

| Respiration Performance Specifications | | |
|--|------------|--|
| Respiration Rate | Range | Adult/pedi: 0 to 120 rpm Neo: 0 to 170 rpm |
| | Accuracy | at 0 to 120 rpm ± 1 rpm at 120 to 170 rpm ± 2 rpm |
| | Resolution | 1 rpm |
| Bandwidth | | 0.3 to 2.5Hz (-6dB) |
| Noise | | Less than 25m Ω (rms) referred to the input |

SpO₂

Complies with EN ISO 9919:2005 (except alarm system; alarm system complies with IEC 60601-2-49:2001).

Measurement Validation: The SpO₂ accuracy has been validated in human studies against arterial blood sample reference measured with a CO-oximeter. Pulse oximeter measurements are statistically distributed, only about two-thirds of the measurements can be expected to fall within the specified accuracy compared to CO-oximeter measurements. Display Update Period: Typical: 2 seconds, Maximum: 30 seconds. Max. with NBP INOP suppression on: 60 seconds.

| Respiration Alarm Specifications | Range | Adjustment | Delay |
|----------------------------------|---|---|---|
| High | Adult/pedi: 10 to 100 rpm Neo: 30 to 150 rpm | under 20 rpm: 1 rpm steps over 20 rpm: 5 rpm steps | max. 14 seconds |
| Low | Adult/pedi: 0 to 95 rpm Neo: 0 to 145 rpm | under 20 rpm: 1 rpm steps over 20 rpm: 5 rpm steps | for limits from 0 to 20 rpm: max. 4 seconds for limits above 20 rpm: max. 14 seconds |

| SpO ₂ Performance Specifications | |
|---|--|
| Sensors | Wavelength range: 500 to 1000 nm Emitted Light Energy: ≤ 15mW Information about the wavelength range can be especially useful to clinicians (for instance, when photodynamic therapy is performed) |
| Pulse Oximeter Calibration Range | 70 - 100% |

a. The specified accuracy is the root-mean-square (RMS) difference between the measured values and the reference values

| SpO ₂ Alarm Specifications | Range | Adjustment | Delay |
|---------------------------------------|---|---|---------------------------------|
| SpO ₂ | Adult: 50 to 100% Pedi/Neo: 30 to 100% | 1% steps | (0, 1, 2, 3,... 30) + 4 seconds |
| Desat | Adult: 50 to Low alarm limit Pedi/Neo: 30 to Low alarm limit | 1% steps | |
| Pulse | 30 to 300 bpm | Adult: 1 bpm steps (30 to 40 bpm) 5 bpm steps (40 to 300 bpm) Pedi/Neo: 1 bpm steps (30 to 50 bpm) 5 bpm steps (50 to 300 bpm) | max. 14 seconds |
| Tachycardia | Difference to high limit 0 to 50 bpm | 5 bpm steps | max. 14 seconds |
| | Clamping at 150 to 300 bpm | 5 bpm steps | |

| SpO ₂ Performance Specifications | | |
|---|-----------------------|--|
| SpO ₂ | Range | 0 to 100% |
| | Accuracy ^a | <p>Philips Reusable Sensors: M1191A, M1191AL, M1191ANL, M1191B, M1191BL, M1192A, M1192AN = 2% (70% to 100%) M1193A, M1193AN, M1194A, M1194AN, M1195A, M1195AN, M1196A = 3% (70% to 100%)</p> <p>Philips Reusable Sensors with M1943A(L): M1191T, M1192T, M1193T (Adult), M1196T = 3% (70% to 100%) M1193T (Neonate) = 4% (70% to 100%)</p> <p>Philips Disposable Sensors with M1943A(L): M1132A, M1133A (adult/infant) = 2% M1131A, M1133A (neonate), M1901B, M1902B, M1903B, M1904B = 3% (70% to 100%)</p> <p>NellcorPB[®] Sensors with M1943A(L): MAX-A, MAX-AL, MAX-P, MAX-I, MAX-N, D-25, D-20, I-20, N-25, OxiCliq A, P, I, N = 3% (70% to 100%)</p> <p>Masimo Reusable Sensors[®] with LNOP MP12 or LNC MP10: LNOP DC-I, LNOP DC-IP, LNOP YI, LNCS DC-I, LNCS DC-IP, LNCS TF-I: 2% (70% to 100%) LNOP TC-I, LNCS TC-I: 3.5% (70% to 100%)</p> <p>Masimo Disposable Sensors[®] with LNOP MP12 or LNC MP10: LNOP Adt, LNOP Adtx, LNOP Pdt, LNOP Pdtx, LNOP Inf-L, LNCS Adtx, LNCS Pdtx, LNCS Inf-L: 2% (70% to 100%) LNOP Neo-L, LNOP NeoPt-L, LNCS Neo-L, LNCS NeoPt-L: 3% (70% to 100%)</p> |
| | Resolution | 1% |
| Pulse | Range | 30 to 300 bpm |
| | Accuracy | ±2% or 1 bpm, whichever is greater |
| | Resolution | 1 bpm |

| SpO ₂ Alarm Specifications | Range | Adjustment | Delay |
|---------------------------------------|-------------------------------------|-------------|-----------------|
| Bradycardia | Difference to low limit 0 to 50 bpm | 5 bpm steps | max. 14 seconds |
| | Clamping at 30 to 100 bpm | 5 bpm steps | |

NBP

Complies with IEC 60601-2-30:1999/EN60601-2-30:2000.

| NBP Performance Specifications | | |
|--------------------------------|------------|--|
| Measurement Ranges | Systolic | Adult: 30 to 270 mmHg (4 to 36 kPa) Pedi: 30 to 180 mmHg (4 to 24 kPa) Neo: 30 to 130 mmHg (4 to 17 kPa) |
| | Diastolic | Adult: 10 to 245 mmHg (1.5 to 32 kPa) Pedi: 10 to 150 mmHg (1.5 to 20 kPa) Neo: 10 to 100 mmHg (1.5 to 13 kPa) |
| | Mean | Adult: 20 to 255 mmHg (2.5 to 34 kPa) Pedi: 20 to 160 mmHg (2.5 to 21 kPa) Neo: 20 to 120 mmHg (2.5 to 16 kPa) |
| | Pulse Rate | Adult: 40 to 300 Pedi: 40 to 300 Neo: 40 to 300 |
| Accuracy | | Max. Std. Deviation: 8 mmHg (1.1 kPa) Max. Mean Error: ±5 mmHg (±0.7 kPa) |

| NBP Performance Specifications | | |
|---------------------------------|---|------------------------------|
| Pulse Rate Measurement Accuracy | 40 to 100 bpm: ± 5 bpm 101 to 200 bpm: ± 5% of reading 201 to 300 bpm: ± 10% of reading (average over NBP measurement cycle) | |
| Heart Rate Range | 40 to 300 bpm | |
| Measurement Time | Typical at HR > 60bpm Auto/manual: 30 seconds (adult) 25 seconds (neonatal) Stat: 20 seconds Maximum time: 180 seconds (adult/pediatric) 90 seconds (neonates) | |
| Cuff Inflation Time | Typical for normal adult cuff: Less than 10 seconds Typical for neonatal cuff: Less than 2 seconds | |
| Initial Cuff Inflation Pressure | Adult: 165 ±15 mmHg Pedi: 130 ±15 mmHg Neo: 100 ±15 mmHg | |
| Auto Mode Repetition Times | 1, 2, 2.5, 3, 5, 10, 15, 20, 30, 45, 60 or 120 minutes | |
| STAT Mode Cycle Time | 5 minutes | |
| Venipuncture Mode Inflation | | |
| Inflation Pressure | Adult | 20 to 120 mmHg (3 to 16 kPa) |
| | Pediatric | 20 to 80 mmHg (3 to 11 kPa) |
| | Neonatal | 20 to 50 mmHg (3 to 7 kPa) |
| Automatic deflation after | Adult/pediatric | 170 seconds |
| | Neonatal | 85 seconds |

Measurement Validation: In adult and pediatric mode, the blood pressure measurements determined with this device comply with the American National Standard for Electronic or Automated Sphygmomanometers (ANSI/AAMI SP10 - 1992) in relation to mean error and standard deviation, when compared to intra-arterial or auscultatory measurements (depending on the configuration) in a representative patient population. For the auscultatory reference the 5th Korotkoff sound was used to determine the diastolic pressure.

In neonatal mode, the blood pressure measurements determined with this device comply with the American National Standard for Electronic or Automated Sphygmomanometers (ANSI/AAMI SP10 - 1992 and AAMI/ANSI SP10A -1996) in relation to mean error and standard deviation, when compared to intra-arterial measurements in a representative patient population.

| NBP Alarm Specifications | Range | Adjustment |
|--------------------------|---------------------------------------|---|
| Systolic | Adult: 30 to 270 mmHg (4 to 36 kPa) | 10 to 30 mmHg: 2 mmHg (0.5 kPa) > 30 mmHg: 5 mmHg (1kPa) |
| | Pedi: 30 to 180 mmHg (4 to 24 kPa) | |
| | Neo: 30 to 130 mmHg (4 to 17 kPa) | |
| Diastolic | Adult: 10 to 245 mmHg (1.5 to 32 kPa) | |
| | Pedi: 10 to 150 mmHg (1.5 to 20 kPa) | |
| | Neo: 10 to 100 mmHg (1.5 to 13 kPa) | |
| Mean | Adult: 20 to 255 mmHg (2.5 to 34 kPa) | |
| | Pedi: 20 to 160 mmHg (2.5 to 21 kPa) | |
| | Neo: 20 to 120 mmHg (2.5 to 16 kPa) | |

| NBP Overpressure Settings | | |
|---------------------------|-----------------------------|---------------------|
| Adult | > 300 mmHg (40 kPa) > 2 sec | not user adjustable |
| Pedi | > 300 mmHg (40 kPa) > 2 sec | |
| Neo | > 150 mmHg (20 kPa) > 2 sec | |

Invasive Pressure and Pulse

Complies with IEC 60601-2-34:2000/EN60601-2-34:2000.

| Invasive Pressure Performance Specifications | | |
|--|------------------------------|--|
| Measurement Range | | -40 to 360 mmHg |
| Pulse Rate | Range | 25 to 350 bpm |
| | Accuracy | ±1% Full Range |
| | Resolution | 1 bpm |
| Input Sensitivity | | Sensitivity:5µV/V/mmHg (37.5µV/V/kPa) Adjustment range:±10% |
| Transducer | | Load Impedance:200 to 2000 Ω (resistive) Output Impedance:≤3000 Ω (resistive) |
| Frequency Response | | dc to 12.5 Hz or 40 Hz |
| Zero Adjustment | Range: | ±200 mmHg (±26 kPa) |
| | Accuracy | ±1 mmHg (±0.1 kPa) |
| | Drift | Less than 0.1mmHg/°C (0.013 kPa/°C) |
| Gain Accuracy | Accuracy | ±1% |
| | Drift | Less than 0.05%/°C |
| | Non linearity and Hysteresis | Error of ≤ 0.4% FS (@CAL 200 mmHg) |
| Overall Accuracy | (including transducer) | ± 4% of reading or ± 4 mmHg (± 0.5 kPa), whichever is greater |
| Volume displacement of CPJ840J6 | | 0.1 mm ³ /100 mmHg |

| Invasive Pressure Alarm Specifications | Range | Adjustment | Delay |
|--|---------------------------------------|---|-----------------|
| Pressure | -40 to 360 mmHg (-5.0 to 48 kPa) | -40 to 30 mmHg 2 mmHg (0.5 kPa) > 30 mmHg 5 mmHg (1 kPa) | max. 12 seconds |
| Extreme High | Difference to high limit 0 to 25 mmHg | 5 mmHg steps (0.5 kPa) | |
| | Clamping at -40 to 360 mmHg | 5 mmHg steps (1.0 kPa) | |
| Extreme Low | Difference to low limit 0 to 25 mmHg | 5 mmHg steps (0.5 kPa) | |
| | Clamping at -40 to 360 mmHg | 5 mmHg steps (1.0 kPa) | |
| Pulse | 25 to 300 bpm | Adult: 1 bpm steps (25 to 40 bpm) 5 bpm steps (40 to 300 bpm) Pedi/Neo: 1 bpm steps (25 to 50 bpm) 5 bpm steps (50 to 300 bpm) | |
| Tachycardia | Difference to high limit 0 to 50 bpm | 5 bpm steps | max. 14 seconds |
| | Clamping at 150 to 300 bpm | 5 bpm steps | |
| Bradycardia | Difference to low limit 0 to 50 bpm | 5 bpm steps | max. 14 seconds |
| | Clamping at 25 to 100 bpm | 5 bpm steps | |

Temp

Complies with EN 12470-4:2000

| Temp Performance Specifications | | |
|---------------------------------|------------|---|
| Temp | Range | -1 to 45 °C (30 to 113 °F) |
| | Resolution | 0.1 °C (0.2 °F) |
| | Accuracy | ±0.1 °C (±0.2 °F) |
| Average Time Constant | | Less than 10 seconds |
| Alarms | Range | -1 to 45 °C (30 to 113 °F) |
| | Adjustment | -1 to 35 °C (30 to 95 °F): 0.5 °C (1.0 °F) steps 35 to 45 °C (95 to 113 °F): 0.1 °C (0.2 °F) steps |

| Temp Alarm Specifications | Range | Adjustment |
|---------------------------|----------------------------|---|
| Temp High/Low Alarms | -1 to 45 °C (30 to 113 °F) | -1 to 35 °C (30 to 95 °F), 0.5 °C (1.0 °F) steps 35 to 45 °C (95 to 113 °F), 0.1 °C (0.2 °F) steps |

Predictive Temperature

| Performance Specifications | |
|---|--|
| Technology | Welch Allyn® SureTemp Plus® |
| Probe Types | oral/axillary, rectal |
| Temperature Measurement Range | 26.7 to 43.3 °C (80 to 110 °F) |
| Resolution | ±0.1 °C (±0.2 °F) |
| Accuracy (Oral, axillary and rectal measurements) | ±0.1 °C (±0.2 °F) (in continuous mode, complies with ASTM 1112-00) |

CO₂

The CO₂ measurement complies with EN ISO 21647:2004 + Cor.1:2005 (except alarm system; alarm system complies with IEC 60601-2-49:2001).

| Mainstream CO ₂ Performance Specifications | | |
|---|---|---|
| CO ₂ | Range | 0 to 150 mmHg (0 to 20.0 kPa) |
| | Accuracy | after 2 minutes warmup: For values between 0 and 40 mmHg: ±2.0 mmHg (±0.29 kPa) For values from 41 to 70 mmHg: ±5% of reading For values from 71 to 100 mmHg: ±8% of reading The specifications are valid for standard gas mixtures, balance air, fully hydrated at 35°C, P _{abs} = 760 mmHg, flow rate = 2 l/min. |
| | Resolution | Numeric: 1.0 mmHg (0.1 kPa) Wave: 0.1 mmHg (0.01 kPa) |
| | Stability: Short term drift Long term drift | ±0.8 mmHg over four hours Accuracy specification will be maintained over a 120 hour period |
| awRR | Range | 2 to 150 rpm |
| | Accuracy | ±1 rpm |
| Warm-up Time | | 2 minutes with CO ₂ transducer attached for full accuracy specification |
| Response Time | | Less than 60 ms (with adult or infant reusable or disposable adapter) |

Ordering Information

Ordering information for the M8105A patient monitor is given here.

| Basic Functionality | MP5 (M8105A) |
|--|--------------|
| Order one Hxx option | |
| General/ICU Configuration | H10 |
| Neonatal Configuration | H20 |
| OR/Anesthesia Configuration ^a | H30 |
| Cardiac Configuration | H40 |
| Order one Axx option | |
| 3 Realtime Wave Segments | A03 |
| 4 Realtime Wave Segments | A04 |

a. If IntelliVue G1/G5 is required, H30 must be ordered.

Application Options

| Application Options | M8105A |
|---|--------|
| Cardiac Applications | |
| Basic Arrhythmia | Incl. |
| Full Arrhythmia | C01 |
| Neonatal Applications | |
| Neonatal Event Surveillance (includes OxyCRG) | C04 |
| Clinical Applications | |
| Drug Calculator | C05 |
| 12-Lead ECG Application (conventional) | C12 |
| ST Map | C13 |

ProtocolWatch

| Application Options | M8105A |
|--|--------|
| Protocol Watch - Sepsis Screening Tool | P01 |

Measurement Options

| Parameter Add-ons | M8105A |
|---|--------|
| ECG, Resp, NBP, SpO2 | B20 |
| ECG, Resp, NBP, SpO2, Predictive Temp. | B21 |
| ECG, Resp, NBP, SpO2, Press & Temp (1x) | B22 |
| ECG, Resp, NBP, SpO2 + TAAP + Predictive Temp | B31 |
| ECG, Resp, NBP, SpO2 + TAAP + Press & Temp (1x) | B32 |
| ECG, Resp, NBP, SpO2, Press & Temp, Predictive Temp | B41 |
| ECG, Resp, NBP, SpO2, Press & Temp (2x) | B42 |
| ECG, Resp, NBP, SpO2, Press & Temp, CO2- ready | B43 |

Hardware Options

| Hardware Add-Ons | M8105A |
|-----------------------------------|--------|
| Built-in recorder | E05 |
| Bed hanger Mount | E21 |
| Quick Release Mount | E22 |
| 1x High Power Lithium-Ion battery | E24 |

Basic Mandatory Interface Options

One of the following three options must be ordered

| Interfaces | M8105A |
|---------------------------|--------|
| LAN & Video Output | J01 |
| LAN & Battery Operation | J02 |
| Advanced System Interface | J40 |

Interface Options

| Interfaces | M8105A |
|-----------------------------------|--------|
| Full Networking software | C15 |
| Companion Mode Interface | J21 |
| IntelliVue 802.11 Bedside Adapter | J35 |
| Instrument Telemetry 1.4 GHz | J45 |
| Instrument Telemetry 2.4 GHz | J47 |

Upgrade Options

| Upgrade | M8105AU |
|--|---------|
| Waves | |
| 4 waves | A04 |
| Interfaces | |
| Advanced System Interface | J40 |
| Parameter Add-ons | |
| Add Predictive Temperature to B20 | B21 |
| Add Predictive Temperature to B22 | B41 |
| Add Press & Temp to B22 | B42 |
| Add CO ₂ -ready to B22 | B43 |
| Clinical Applications | |
| Full Arrhythmia Capability | C01 |
| Neonatal Event Review | C04 |
| Drug Calculator | C05 |
| ST-Map | C13 |
| Full Networking Software | C15 |
| Protocol Watch | |
| Sepsis Screening | P01 |
| Rev SW + Sepsis Screening | P41 |
| Hardware Add-Ons | |
| Built-in Recorder | E05 |
| Bed Hanger Mount | E21 |
| Interfaces | |
| Companion Mode Interface for System Software F.0 or higher | J21 |
| IntelliVue 802.11 bedside adapter | J35 |
| Instrument Telemetry 1.4 Ghz | J45 |
| Instrument Telemetry 2.4 Ghz | J47 |
| Companion Mode Interface for System Software E.0 - incl. upgrade to Rev. F.0 | J99 |
| Upgrade to the latest software release | SU0 |

Sensors and Disposables

| Accessory | M8105A |
|---|--------|
| 3-lead Accessories Bundle ICU-AAMI one piece ECG cable (Trunk/Lead set) | G06 |
| 3-lead Accessories Bundle ICU-IEC one piece ECG cable (Trunk/Lead set) | G07 |
| 5-lead Accessories Bundle ICU-AAMI one piece ECG cable (Trunk/Lead set) | G08 |
| 5-lead Accessories Bundle ICU-IEC one piece ECG cable (Trunk/Lead set) | G09 |
| 5-lead Accessories Bundle ICU-AAMI | H06 |
| 5-lead Accessories Bundle ICU-IEC | H07 |
| 5-lead Accessories Bundle OR-AAMI | H08 |
| 5-lead Accessories Bundle OR-IEC | H09 |
| Accessories Bundle Neonatal-AAMI | H14 |
| Accessories Bundle Neonatal-IEC | H15 |
| 3-lead Accessories Bundle ICU-AAMI | H16 |
| 3-lead Accessories Bundle ICU-IEC | H17 |
| 3-lead Accessories Bundle OR-AAMI | H18 |
| 3-lead Accessories Bundle OR-IEC | H19 |
| CO ₂ Mainstream Sensor | N01 |
| Reusable Adult Airway Adapter (msCO ₂) | N02 |
| Reusable Infant Airway Adapter (msCO ₂) | N03 |
| Single Use Adult Airway Adapter (msCO ₂) | N04 |
| Single Use Infant Airway Adapter (msCO ₂) | N05 |
| CO ₂ Sidestream Sensor | N11 |
| Non-intubated Adult Airway Adapter (ssCO ₂) | N12 |

| Accessory | M8105A |
|---|--------|
| Non-intubated pediatric Airway Adapter (ssCO ₂) | N13 |
| Intubated Adult Airway Adapter (ssCO ₂) | N14 |
| Intubated Pediatric Airway Adapter (ssCO ₂) | N15 |
| Predictive Temp Oral with 25 probe covers | T01 |
| Predictive Temp Rectal with 25 probe covers | T02 |
| Calibration Key (453564033691) | n/a |

Related Products

M3086A Support Tool

Mounting Information

The Intellivue MP5 Roll Stand Mounting Kit (Order No. 989803153021) is compatible with the table top mount and the standard mounting plate. For information on other mounting hardware, contact your local Philips sales representative. For GCX mounting hardware information, see www.gcx.com/philips.

Documentation

All documentation is available in .pdf format on documentation CD-ROM. Additionally, a printed copy of the Instructions for Use and Quick Guide ships with each monitor.

- Instructions for Use (printed)
- Quick Guide (printed)
- Installation and Service Guide
- Configuration Guide
- Documentation CD-ROM
- Training Guide (printed)

Cables M8022A

| Length | Description ^a | Product/Option |
|--------------|--------------------------|----------------|
| Analog Video | | |

| Length | Description ^a | Product/Option |
|--|--|----------------|
| 1.5 m | Monitor to Display | M8022A #VA2 |
| 3.0 m | Monitor to Display | M8022A #VA3 |
| 10.0 m | Monitor to Display | M8022A #VA6 |
| 15.0 m | Monitor to Display | M8022A #VA7 |
| 25.0 m | Monitor to Display | M8022A #VA9 |
| MIB RS/232 Cables | | |
| 1.5 m | Serial cable | M8022A #SR2 |
| 3.0 m | Serial cable | M8022A #SR3 |
| 10.0 m | Serial cable | M8022A #SR6 |
| 15.0 m | Serial cable | M8022A #SR7 |
| 25.0 m | Serial cable | M8022A #SR9 |
| Touch Cables | | |
| 1.5 m | Touch cable | M8022A #TC2 |
| 3.0 m | Touch cable | M8022A #TC3 |
| 10.0 m | Touch cable | M8022A #TC6 |
| 15.0 m | Touch cable | M8022A #TC7 |
| 25.0 m | Touch cable | M8022A # TC9 |
| Nurse Call Relay Cable | | |
| 3.0 m | standard (backward compatible) nurse paging relay cable ^b | M8022A #NC3 |
| 10.0 m | cable | M8022A #NC6 |
| ECG Out Cable | | |
| 3.0 m | standard ECG out cable ^c | M8022A #SY3 |
| PWD (Patient Worn Device) Interface Cables | | |
| 0.5 m | PWD Interface Cable (PWD side) | 989803143481 |
| 2.0m | PWD Interface Cable (monitor side) | 989803146911 |

a. Both ends terminated with HDSUB15 (VGA) connectors.

b. One end terminated with phone plug; other end w/o connector.

c. Both ends terminated with 1/4" phone plug.

ECG Accessories



This symbol indicates that the cables and accessories are designed to have special protection against electric shocks (particularly regarding allowable leakage currents), and are defibrillator proof.

Trunk Cables

| | 3-Electrode Cable Set | 5-Electrode Cable Set | 6-Electrode Cable Set | 10-Electrode Cable set (5+5) | 10-Electrode Cable set (6+4) |
|----------|-----------------------|-----------------------|-----------------------|------------------------------|------------------------------|
| Part No. | M1669A | M1668A | M1667A | M1663A | M1665A |
| Length | 2.7m | 2.7m | 2.7m | 2.0m | 2.7m |

3-Electrode Cable Sets

| Description | Length | AAMI Part No. | IEC Part No. |
|-----------------------|--------|---------------|--------------|
| OR Grabber shielded | 1.0m | M1675A | M1678A |
| ICU Grabber shielded | 1.0m | M1671A | M1672A |
| ICU snap shielded | 1.0m | M1673A | M1674A |
| ICU Clip non-shielded | 0.45m | M1622A | -- |
| ICU Clip non-shielded | 0.7m | M1624A | M1626A |

5-Electrode Cable Sets

| Description | Length | AAMI Part No. | IEC Part No. |
|---------------------|-----------|---------------|--------------|
| OR Grabber shielded | 1.0m/1.6m | M1973A | M1974A |

| Description | Length | AAMI Part No. | IEC Part No. |
|---------------------------|-----------|---------------|--------------|
| ICU Grabber shielded | 1.0m/1.6m | M1968A | M1971A |
| ICU Snap shielded | 1.0m/1.6m | M1644A | M1645A |
| ICU Miniclip non-shielded | 0.7m/1.3m | M1647A | M1648A |

6-Electrode Cable Sets

| Description | Length | AAMI Part No. | IEC Part No. |
|-------------|-----------|---------------|--------------|
| OR Grabber | 1.0m/1.6m | M1684A | M1685A |
| ICU Grabber | 1.0m/1.6m | M1680A | M1681A |
| ICU Snap | 1.0m/1.6m | M1682A | M1683A |

10-Electrode (5+5) Cable Sets

| Description | Length | AAMI Part No. | IEC Part No. |
|---|--------|---------------|--------------|
| ICU Grabber, chest, shielded | 1.0m | M1976A | M1978A |
| ICU Snap, chest, shielded | 1.0m | M1602A | M1604A |
| OR Grabber, chest, shielded | 1.0m | M1979A | M1984A |
| For Limb Leads see 5-electrode cable sets | | | |

10-Electrode (6+4) Cable Sets

| Description | Length | AAMI Part No. | IEC Part No. |
|------------------------------|--------|---------------|--------------|
| ICU Grabber, chest, shielded | 1.0m | M1532A | M1533A |
| ICU Snap, chest, shielded | 1.0m | M1537A | M1538A |
| OR Grabber, chest, shielded | 1.0m | M1557A | M1558A |

| Description | Length | AAMI Part No. | IEC Part No. |
|---|--------|---------------|--------------|
| For Limb Leads see 6-electrode cable sets | | | |

One-piece Cables

| Description | Length | AAMI Part No. | IEC Part No. |
|---------------------|--------|---------------|--------------|
| 3-lead Grabber, ICU | 1.0m | 989803143181 | 989803143171 |
| 5-lead Grabber, ICU | 1.0m | 989803143201 | 989803143191 |

Radio-translucent Cables

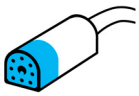
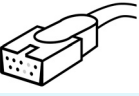

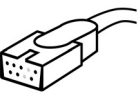

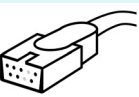
Pack of five single wires, radio-translucent, 0.9m, M1649A




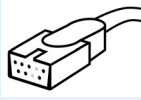
Set Combiners and Organizers

| Set combiners and organizers | | Part No. |
|--|-------------|--------------|
| Set combiner | 3-electrode | M1501A |
| | 5-electrode | M1502A |
| Set organizer for shielded leadsets - grabber and snap | 3-electrode | M1503A |
| | 4-electrode | M1664A |
| | 5-electrode | M1504A |
| | 6-electrode | M1679A |
| Set organizer for non-shielded lead sets - miniclip | 3-electrode | M1636A |
| | 5-electrode | M1638A |
| Bedsheet clip | | M1509A |
| Replacement red cover for trunk cable (for 5-electrode cable sets) | | 989808148861 |

Philips FAST SpO₂ Accessories

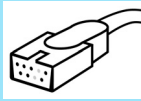
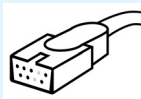
Philips Reusable Sensors

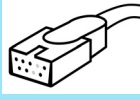
| Part Number | Description | Connector Type |
|-------------------|---|--|
| M1191A/B | Adult Sensor (2m cable) | Philips 8-pin  |
| M1191AL/BL | Adult Sensor (3m cable) | |
| M1191ANL | Adult Sensor (3m cable) Nellcor OxiMax-compatible ^a | |
| M1191T | Adult Sensor (requires M1943A (1.1m) or M1943AL (3m) adapter cable) | Generic D-Sub  |
| M1192A | Small Adult/Pediatric sensor (1.5m cable) | Philips 8-pin  |
| M1192AN | Small Adult/Pediatric sensor (1.5m cable) Nellcor OxiMax-compatible ^a | |
| M1192T | Small Adult Pediatric sensor (requires M1943A (1.1m) or M1943AL (3m) adapter cable) | Generic D-Sub  |
| M1193A | Neonatal Hand/Foot Sensor (1.5m cable) | Philips 8-pin  |
| M1193AN | Neonatal Hand/Foot Sensor (1.5m cable) Nellcor OxiMax-compatible ^a | |
| M1193T | Neonatal Sensor (requires M1943A (1.1m) or M1943AL (3m) adapter cable) | Generic D-Sub  |

| Part Number | Description | Connector Type |
|----------------|--|--|
| M1194A | Adult/Pediatric Clip Sensor (ear) (1.5m cable) | Philips 8-pin  |
| M1194AN | Adult/Pediatric Clip Sensor (ear) (1.5m cable) Nellcor OxiMax-compatible ^a | |
| M1195A | Infant Sensor (1.5m cable) | Philips 8-pin  |
| M1195AN | Infant Sensor (1.5m cable) Nellcor OxiMax-compatible ^a | |
| M1196A | Adult Clip Sensor (3m cable) | Philips 8-pin  |
| M1196T | Adult Clip Sensor (requires M1943A (1.1m) or M1943AL (3m) adapter cable) | Generic D-Sub  |

a. only in combination with Philips FAST-SpO₂ and Philips OxiMax-compatible patient monitors.

Philips Disposable Sensors

| Part Number | Description | Connector Type |
|---------------|---|--|
| M1131A | Adult/Pediatric Sensor (requires M1943A (1.1m) or M1943AL (3m) adapter cable) | Generic D-Sub  |
| M1132A | Infant Sensor (requires M1943A (1.1m) or M1943AL (3m) adapter cable) | Generic D-Sub  |

| Part Number | Description | Connector Type |
|---------------|--|--|
| M1133A | Adult/Infant/ Neonatal Sensor (requires M1943A (1.1m) or M1943AL (3m) adapter cable) | Generic D-Sub  |

NELLCOR® Disposable Sensors¹:

Purchase Nellcor OxiCliq sensors and adapter cables directly from Tyco Healthcare.

| Product Number | Description | Philips Part Number |
|--------------------------------|---------------------------|---------------------------|
| OxiMax MAX-A ^a | Adult Sensor | M1904B^b |
| OxiMax MAX-AL ^a | Adult Sensor (long cable) | n/a |
| OxiMax MAX-P ^a | Pediatric Sensor | M1903B^b |
| OxiMax MAX-I ^a | Infant Sensor | M1902B^b |
| OxiMax MAX-N ^a | Neonatal Sensor | M1901B^b |
| Oxisensor II D-25 ^a | Adult Sensor | n/a |
| Oxisensor II D-20 ^a | Pediatric Sensor | n/a |
| Oxisensor II I-20 ^a | Infant Sensor | n/a |
| Oxisensor II N-25 ^a | Neonatal Sensor | n/a |
| OxiCliq A ^c | Adult Sensor | n/a |
| OxiCliq P ^c | Pediatric Sensor | n/a |
| OxiCliq I ^c | Infant Sensor | n/a |
| OxiCliq N ^c | Neonatal Sensor | n/a |

a. Requires M1943 A(L) adapter cable

b. not available from Philips in the U.S.A.

c. Requires M1943 A(L) and OC3 adapter cables

1. Nellcor, OxiMax and OxiCliq are trademarks of Nellcor Puritan Bennett Inc., a part of Tyco Healthcare.

MASIMO LNOP®¹ Reusable Sensors:

| Product Number | Description | Philips Part Number |
|----------------|----------------------------|---------------------|
| LNOP DC-I | Adult Sensor | 989803140321 |
| LNOP DC-IP | Pediatric Sensor | 989803140331 |
| LNOP-YI | Reusable Multi-Site Sensor | n/a |
| LNOP TC-I | Tip Clip reusable Sensor | 989803140341 |

MASIMO LNCS®¹ Reusable Sensors:

| Product Number | Description | Philips Part Number |
|----------------|--------------------------|---------------------|
| LNCS DC-I | Adult Sensor | 989803148281 |
| LNCS DC-IP | Pediatric Sensor | 989803148291 |
| LNCS-TC-I | Reusable Ear Sensor | 989803148301 |
| LNCS TF-I | Reusable Forehead Sensor | 989803148311 |

MASIMO LNOP® Disposable Adhesive Sensors:

| Product Number | Description | Philips Part Number |
|----------------|--|---------------------|
| LNOP Adt | Adult Sensor | 989803140231 |
| LNOP Adtx | Adult Sensor | n/a |
| LNOP Pdt | Pediatric Adhesive Sensor | 989803140261 |
| LNOP Pdtx | Pediatric Sensor | n/a |
| LNOP INF-L | Neo/Infant Adhesive Sensor | 989803140311 |
| LNOP NEO-L | Neo Adhesive Sensor | 989803140291 |
| LNOP NEOPT-L | Neo Pre-Term Sensitive Skin Adhesive Sensors | 989803140301 |

MASIMO LNCS® Disposable Adhesive Sensors:

| Product Number | Description | Philips Part Number |
|----------------|--|---------------------|
| LNCS Adtx | Adult Finger Sensor | 989803148231 |
| LNCS Pdtx | Pediatric Finger Sensor | 989803148241 |
| LNCS INF-L | Infant Toe Sensor | 989803148251 |
| LNCS NEO-L | Neo Foot Sensor or Adult Finger Sensor | 989803148271 |
| LNCS NEOPT-L | Neo Pre-Term Sensitive Skin Adhesive Sensors | 989803148261 |



The Philips M8105A uses Masimo certified pulse oximetry for reduced noise and low perfusion performance with Masimo Sensors under the Masimo NR&LP protocol available from Masimo.

1. LNOP and LNCS are federally registered trademarks of Masimo Corporation

Extension/Adapter Cables:

| Part Number | Description |
|---------------------------------|--|
| M1941A | Extension Cable (2m) (8-pin to 8-pin) |
| M1943A | Adapter Cable (1.1m) for Philips and Nellcor disposable sensors (8-pin to 9-pin D-Sub) |
| M1943AL | Adapter Cable (3m) for Philips and Nellcor disposable sensors (8-pin to 9-pin D-Sub) |
| OC3 | Adapter cable for OxiCliq Sensors (available from Nellcor only) |
| LNOP MP12 (451261000761) | LNOP MP Series Patient Cable (3.6 m) Adapter Cable for Masimo LNOP Sensors |
| LNC MP10 (989803148221) | LNCS MP Series Patient CABLE (3.0 m) Adapter Cable for Masimo LNCS Sensors |

Non Invasive Blood Pressure Accessories



These cuffs and tubings are designed to have special protection against electric shocks (particularly regarding allowable leakage currents), and are defibrillator proof.

| Multi-Patient Comfort Cuffs and Disposable Cuffs | | |
|--|-----------------|---------------|
| Patient Category | Disposable cuff | Reusable cuff |
| Adult (Thigh) | M1879A | M1576A |
| Large Adult | M1878A | M1575A |
| Adult | M1877A | M1574A |
| Small Adult | M1876A | M1573A |
| Pediatric | M1875A | M1572A |
| Infant | M1874A | M1571A |

Tubing: Use M1598B or M1599B

| Reusable Cuff Kits | Part No. |
|---|----------|
| Infant, pediatric, small adult, adult | M1577A |
| Small adult, adult, large adult, thigh | M1578A |
| Infant, pediatric, small adult, adult, large adult, thigh | M1579A |

| Adult/Pediatric Antimicrobial Coated Reusable cuffs | | | |
|---|--------------------|-----------------------|----------------------|
| Cuff Size (color) | Circumference (cm) | Bladder Width | Single-Hose Part No. |
| Infant (orange) | 9.0 - 14.8 | 5.4 cm 2.1 inches | M4552A |
| Pediatric (green) | 13.8 - 21.5 | 8.0 cm 3.1 inches | M4553A |
| Small Adult (royal blue) | 20.5 - 28.5 | 10.6 cm 4.2 inches | M4554A |
| Adult (navy blue) | 27.5 - 36.5 | 13.5 cm 5.3 inches | M4555A |
| Adult X-long (navy blue) | 27.5 - 36.5 | 13.5 cm 5.3 inches | M4556A |
| Large Adult (burgundy) | 35.5 - 46.0 | 17.0 cm 6.7 inches | M4557A |
| Large Adult X-long (burgundy) | 35.5 - 46.0 | 17.0 cm 6.7 inches | M4558A |
| Thigh (grey) | 45 - 56.5 | 21.0 cm 8.3 inches | M4559A |

Tubing: Use M1598B or M1599B

| Adult/Pediatric Soft Single Patient Single-Hose Disposable Cuffs | | | |
|--|--------------------|---------------|--------------------------|
| Patient Category | Limb Circumference | Bladder Width | Disposable cuff Part No. |
| Adult (Thigh) | 45.0-56.5 cm | 20.4 cm | M4579A |
| Large Adult X-long | 35.5-46.0 cm | 16.4 cm | M4578A |
| Large Adult | 35.5-46.0 cm | 16.4 cm | M4577A |
| Adult X-long | 27.5 to 36.5 cm | 16.4 cm | M4576A |
| Adult | 27.5-36.5 cm | 13.1 cm | M4575A |
| Small Adult | 20.5-28.5 cm | 10.4 cm | M4574A |
| Pediatric | 15.0-21.5 cm | 8.0 cm | M4573A |
| Infant | 9.0-15.0 cm | 5.6 cm | M4572A |

Tubing: Use M1598B or M1599B

| Neonatal/Infant Cuffs (Disposable, non-sterile) | | | |
|---|--------------------|---------------|----------|
| Cuffs | Limb Circumference | Bladder Width | Part No. |
| Size 1 | 3.1 to 5.7 cm | 2.2 cm | M1866A |
| Size 2 | 4.3 to 8.0 cm | 2.8 cm | M1868A |
| Size 3 | 5.8 to 10.9 cm | 3.9 cm | M1870A |
| Size 4 | 7.1 to 13.1 cm | 4.7 cm | M1872A |


Tubing: Use M1596B or M1597B

| Cuff Tubing | | |
|-------------|-------------|--------|
| Adult | 1.5 m /4.9' | M1598B |
| | 3.0 m/9.8' | M1599B |
| Neonatal | 1.5 m /4.9' | M1596B |
| | 3.0 m/9.8' | M1597B |

Temperature Accessories

| Temperature Probes | Part No. |
|---|----------|
| Reusable | |
| General purpose probe | 21075A |
| Small flexible vinyl probe (Infant/Pediatric) | 21076A |
| Attachable surface probe | 21078A |
| Disposable | |
| General purpose probe | M1837A |
| Skin probe | 21091A |
| Esophageal/Stethoscope Probe (12 French) | 21093A |
| Esophageal/Stethoscope Probe (18 French) | 21094A |
| Esophageal/Stethoscope Probe (24 French) | 21095A |
| Foley Catheter Probe (12 French) | M2255A |
| Foley Catheter Probe (16 French) | 21096A |
| Foley Catheter Probe (18 French) | 21097A |
| Adapter cable 1.5m/4.9' | 21082B |
| Adapter cable 3.0m/9.8' | 21082A |

PRESS Accessories

 These transducers and accessories are designed to have special protection against electric shocks (particularly regarding allowable leakage currents), and are defibrillator proof.

| Pressure Transducers and Accessories | Part No. |
|--|----------|
| Reusable | |
| Reusable pressure transducer 5 μ V/V/mmHg sensitivity | CPJ840J6 |
| Sterile disposable pressure domes for CPJ840J6 (pack of 50) | CPJ84022 |

| Pressure Transducers and Accessories | Part No. |
|---|----------|
| Transducer holder for CPJ840J6 (pack of 4) | CPJ84046 |
| IV pole mount for CPJ840J6 | CPJ84447 |
| Disposable (EU/EFTA only. Not available in USA) | |
| Single channel disposable sensor kit (20) | M1567A |
| Dual channel disposable sensor kit (20) | M1568A |
| Transducer holder for M1567/8A | M2271A |
| IV pole mount for M1567/8A | M2272C |
| Adapter cable for disposable sensor kit, 3.0m, for M1567/8A | M1634A |

Mainstream CO₂ Accessories

| Description | Part No. |
|--|----------|
| CO ₂ Sensor | M2501A |
| Adult/Pediatric Airway Adapter (reusable) | M2513A |
| Infant Airway Adapter (reusable) | M2516A |
| Adult Airway Adapter (single patient use) | M2533A |
| Infant Airway Adapter (single patient use) | M2536A |

Sidestream CO₂ Accessories

| Description | Part No. |
|--|----------|
| CO ₂ Sensor | M2741A |
| Nasal and Oral-Nasal Cannulas | |
| CO ₂ Nasal Cannula, Adult | M2744A |
| CO ₂ Nasal Cannula, Pediatric | M2745A |
| CO ₂ Nasal Cannula, Infant | M2746A |
| CO ₂ / O ₂ Nasal Cannula, Adult | M2750A |
| CO ₂ / O ₂ Nasal Cannula, Pediatric | M2751A |
| CO ₂ Oral-Nasal Cannula, Adult | M2756A |
| CO ₂ Oral-Nasal Cannula, Pediatric | M2757A |
| CO ₂ / O ₂ Oral-Nasal Cannula, Adult | M2760A |
| CO ₂ / O ₂ Oral-Nasal Cannula, Pediatric | M2761A |
| Airway Adapters | |
| Airway Adapter Set, ET > 4.0 mm | M2768A |
| Airway Adapter Set H, ET > 4.0 mm | M2772A |
| Airway Adapter Set H, ET =< 4.0 mm | M2773A |
| Straight Sample Lines | |
| Straight Sample Line | M2776A |
| Straight Sample Line H | M2777A |

Predictive Temperature Accessories

| Temperature Probes and Disposable Covers | Part No. | Welch Allyn Part No. |
|--|--------------|----------------------|
| Oral probe with probe well (holder), 2.7m cable (9ft) | 989803143381 | 02895-000 |
| Rectal probe with probe well (holder), 2.7 m cable (9ft) | 989803143391 | 02895-100 |
| Disposable probe covers: 1,000 (40 boxes, 25 per box) | M4823A | 05031-101 |

**Philips Healthcare is part of
Royal Philips Electronics**

Interested?

Would you like to know more about our imaginative products? Please do not hesitate to contact us. We would be glad to hear from you.

On the web

www.philips.com/healthcare

Via e-mail

healthcare@philips.com

By fax

+31 40 27 64 887

By mail

Philips Healthcare
Global Information Center
P.O Box 1286
5602 BG Eindhoven
The Netherlands

By phone

Asia

Tel: +852 2821 5888

Europe, Middle East, Africa

Tel: +49 7031 463 2254

Latin America

Tel: +55 11 2125 0764

North America

Tel: +1 425 481 7000

800 285 5585 (toll free, US only)



M8105A complies with the requirements of the Council Directive 93/42/EEC of 14 June 1993 (Medical Device Directive).



© 2008 Koninklijke Philips Electronics N.V.
All rights are reserved.

Philips Medical Systems Nederland B.V. reserves the right to make changes in specifications and/or to discontinue any product at any time without notice or obligation and will not be liable for any consequences resulting from the use of this publication.

Printed in The Netherlands.
4522 962 32371 MAR 2008