AS/3[™] Compact Anesthesia Monitor upgraded with U-LIFE2

User's Guide Part I: For Monitor Setup and Reference



Datex-Ohmeda AS/3 Compact Anesthesia Monitor upgraded with U-LIFE2

User's Guide

Part I: For Monitor Setup and Reference

Related to software licenses L-ANE03 and L-ANE03A

Conformity according to the Council Directive 93/42/EEC concerning Medical Devices

CAUTION: U.S. Federal law restricts this device to sale by or on the order of a licensed medical practitioner. Outside the USA, check local laws for any restriction that may apply.

All specifications subject to change without notice.

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About this guide

This User's Guide describes the most common features and functions offered by the Datex-Ohmeda AS/3 Compact Anesthesia Monitor. Descriptions refer to S/5 software licenses L-ANE03 and L-ANE03A.

If you are a new user of the monitor, we suggest you begin with sections "Safety precautions" "System introduction" and "Monitoring basics." Also read Part II of this User's Guide.

The following conventions are used:

- Names of the hard keys on the Command Board, Remote Controller and modules are written in the following way: ECG.
- Menu items are written in bold italic typeface: ECG Setup.
- Menu access is described from top to bottom. For example, the selection of the Screen Setup menu item and the Waveform Fields menu item would be shown as Screen Setup - Waveform Fields.
- Messages (alarm messages, informative messages) displayed on the screen are written inside single quotes: 'Learning'.
- When referring to different sections in this manual, section names are enclosed in double quotes: "Cleaning and care."
- In this manual, the word "select" means choosing and confirming.

Related documentation

Clinical aspects, basic methods of measurement and technical background: S/5 Compact Anesthesia Monitor, User's Reference Manual

Instructions for daily use: AS/3 Compact Anesthesia Monitor upgraded with U-LIFE2, User's Guide Part II

Installation, technical solutions and servicing: AS/3 and CS/3 Compact Monitors upgraded with U-LIFE2 and S/5 Modules, Technical Reference Manuals

Options and selections of the software: Default Configuration Worksheet

Other devices closely related to the AS/3 Compact Anesthesia Monitor:

S/5 Central and S/5 Arrhythmia Workstation User's Reference Manuals

Intended purpose (Indications for use)

The Datex-Ohmeda AS/3 Compact Anesthesia Monitor upgraded with L-ANE03 or L-ANE03A software is intended for multiparameter patient monitoring with optional patient care documentation.

The AS/3 Compact Anesthesia Monitor upgraded with L-ANE03 or L-ANE03A software is indicated for monitoring of hemodynamic (including arrhythmia and ST-segment analysis), respiratory, ventilatory, gastrointestinal/regional perfusion, Bispectral index (BIS), Entropy (State Entropy and Response Entropy) and neurophysiological status of all hospital patients.

The AS/3 Compact Anesthesia Monitor with L-ANE03 and L-ANE03A software when using BIS is for monitoring the state of the brain by data acquisition and processing of electroencephalograph signals and may be used as an aid in monitoring the effects of certain anesthetic agents.

The AS/3 Compact Anesthesia Monitor with L-ANE03 or L-ANE03A software is also indicated for documenting patient care related information.

The AS/3 Compact Anesthesia Monitor with L-ANE03 or L-ANE03A software is indicated for use by qualified medical personnel only.

Classifications

In accordance with IEC 60601-1:

- Class I and internal powered equipment the type of protection against electric shock.
- Type BF or CF equipment. The degree of protection against electric shock is indicated by a symbol on each parameter module.
- Equipment is not suitable for use in the presence of a flammable anesthetic mixture with air or with oxygen or nitrous oxide.
- Continuous operation according to the mode of operation.

In accordance with IEC 60529:

IPX1 - degree of protection against harmful ingress of water

In accordance with EU Medical Device Directive: IIb

Responsibility of the manufacturer

Datex-Ohmeda Division, Instrumentarium Corp. is responsible for the safety, reliability and performance of the equipment only if:

- assembly, extensions, readjustments, modifications, service and repairs are carried out by personnel authorized by Datex-Ohmeda.
- electrical installation complies with appropriate requirements.
- the equipment is used in accordance with this User's Guide.

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A portion of the entropy software is derived from the RSA Data Security, Inc. MD5 Message-Digest Algorithm.

Safety precautions

These precautions refer to the entire system. Warnings and cautions specific to parts of the system can be found in the relevant section.

Warnings

A WARNING indicates a situation in which the user or the patient may be in danger of injury or death.

- Connect only one patient to the monitor at a time.
- Use only hospital-grade grounded power outlets and power cord.
- Some equipment malfunctions may not generate a monitor alarm. Always keep the patient under close surveillance.
- To avoid explosion hazard, do not use the monitor in presence of flammable anesthetics. The monitor measures only non-flammable anesthetics.
- Do not use the monitor in high electromagnetic fields (for example, during MRI).
- Do not touch battery operated monitor during defibrillation procedure.
- If the integrity of the external protective earth conductor arrangement is in doubt, use the monitor with battery operation.
- Do not connect any external devices to the system other than those specified by Datex-Ohmeda.
- Keep the monitor horizontal when the Compact Airway Module is used. Tilting the monitor may cause erroneous results in the Compact Airway Module's readings and damage the module.
- Do not touch the patient, table, instruments or the monitor during defibrillation.
- If the integrity of the external protective earth conductor arrangement is in doubt, use the monitor with battery operation.

- Use only accessories and defibrillator-proof cables and invasive pressure transducers approved by Datex-Ohmeda. Other cables, transducers and accessories may cause a safety hazard, damage the system, result in increased emissions or decreased immunity of the system or interfere with the measurement. Protection against cardiac defibrillator discharge is due in part to the accessories for pulse oximetry (SpO₂), temperature (T) and invasive pressure (P) measurement.
 Single-use accessories are not designed to be re-used. Re-use may cause a risk of contamination and affect the measurement
- The monitor or its components should not be used adjacent to or stacked with other equipment. If adjacent or stacked use is necessary, the monitor and its components should be observed to verify normal operation in the configuration in which it will be used.
- Pins of connectors identified with the ESD warning symbol should not be touched. Connections should not be made to these connectors unless ESD precautionary procedures are used. See "Safety precautions: ESD precautionary procedures" in the "User's Reference Manual" for details.
- If liquid has accidentally entered the equipment, disconnect the power cord from the power supply and have the equipment serviced by authorized service personnel.
- If the unit fails to respond as described, do not use the monitor until tested and repaired by authorized service personnel.

accuracy.

Cautions

A CAUTION indicates a situation in which the unit or devices connected to it may be damaged.

- Before connecting the power cord to the power supply, check that the local voltage and frequency correspond with the rating stated on the device plate on the rear panel of the monitor.
- Turn off the power before making any rear panel connections.
- Vibrations during transport may disturb SpO₂, ECG, impedance respiration and NIBP measurements.
- Leave space for circulation of air to prevent the monitor from overheating.
- Do not store or use the monitor outside the temperature and humidity ranges specified in the "Performance" section of this manual.
- Refresh the batteries completely once a month (see section "Cleaning and care").
- Do not subject memory cards to excessive heat, bending or magnetic fields.

Disposal

Dispose of the whole device or parts of it in accordance with local environmental and waste disposal regulations.

Points to note

- Medical electrical equipment needs special precautions regarding EMC and needs to be installed and put into service according to the EMC information provided in the "Technical Reference Manual" by qualified Datex-Ohmeda trained personnel.
- Portable and mobile RF communications equipment can affect the medical electrical equipment.
- The allowed Datex-Ohmeda cables, transducers and accessories for the system are listed in the "Supplies and accessories" section of this manual.
- The equipment is suitable for use in the presence of electrosurgery. Please notice the possible limitations in the parameter sections and in the "Performance" section.
- Service and reparations are allowed for authorized service personnel only.

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Symbols



Attention, consult accompanying documents.

- When displayed next to the O₂ value, indicates that the FiO_2 low alarm limit is set below 21 %.
- When displayed next to the HR value, indicates that the pacer is set on R or a wide ORS is selected.
- BIS: On the Aspect DSC indicates that the converter must not be opened for any reason, or autoclaved.
- On the interface module, M-INT, indicates that it is for connecting external devices. Do not connect patient cables to the module.
- On the M-PRESTN, M-PRETN and M-RESTN module indicates that protection against cardiac defibrillator discharge is due in part to the accessories for pulse oximetry (SpO₂), temperature (T) and invasive pressure (P) measurement.
- On the M-TONO module indicates that the module should be used only with Tonometrics catheters.
- On the rear panel this symbol indicates the following warnings and cautions:

- Electric shock hazard. Do not open the cover or the back. Refer servicing to qualified personnel.

- For continued protection against fire hazard, replace the fuse only with one of the same type and rating.

- Disconnect from the power supply before servicing. - Do not touch battery operated monitor during

defibrillation.

- Lithium battery on CPU board: follow the local regulations for disposal.



Type BF (IEC 60601-1) protection against electric shock



Type BF (IEC 60601-1) defibrillator-proof protection against electric shock



Type CF (IEC 60601-1) protection against electric shock



Type CF (IEC 60601-1) defibrillator-proof protection against electric shock



When displayed in the upper left corner of the screen, indicates that the alarms are silenced. When displayed in the menu or digit fields, indicates that the alarm source has been turned off or alarm does not meet the alarm-specific activation criteria.



Equipotentiality. Monitor can be connected to potential equalization conductor.

Alternating current



Fuse. Replace the fuse only with one of the same type and rating.



Battery operation and remaining capacity (green bar)



Battery charging (white bar)



The monitor is connected to the Datex-Ohmeda Network.



Data Card (green) and/or the Menu Card (white) is inserted.



ESD warning symbol for electrostatic sensitive devices. Pins of connectors identified with the ESD warning symbol should not be touched. Connections should not be made to these connectors unless ESD precautionary procedures are used. See "Safety precautions: ESD precautionary procedures" in the "User's Reference Manual" for details.



Symbol for non-ionizing electromagnetic radiation. Interference may occur in the vicinity of equipment marked with this symbol.



Serial number



V



Submenu. Selecting a menu item with this symbol opens a new menu.

A blinking heart next to the heart rate or pulse rate value indicates the beats detected.

A lung next to the respiration rate value indicates that respiration rate is calculated from the impedance respiration measurement.

System introduction

- (1) AS/3 Compact Anesthesia Monitor with modules inside
- (2) Remote Controller, K-REMCO
- (3) Printer
- (4) Datex-Ohmeda Network



NOTE: Your system may not include all these components. NOTE: Connect only one patient to the Compact Anesthesia Monitor at a time.

NOTE: After transferring or reinstalling the monitor, always check that it is properly connected and securely attached.

NOTE: The monitor display is fragile. Ensure that it is not placed near a heat source or exposed to mechanical shocks, pressure, moisture, or direct sunlight.

WARNING: Before starting to use the system, ensure that the entire combination complies with the international standard IEC 60601-1-1 and with the requirements of the local authorities. Do not connect any external devices to the system other than those specified by Datex-Ohmeda.

WARNING: A printer must be supplied from an additional transformer providing at least basic isolation (isolating or separating transformer).

WARNING: Never install the monitor so that it is above the patient.

AS/3 Compact Anesthesia Monitor connections



- (1) Air filter
- (2) Built-in recorder (optional)
- (3) Connector for anesthesia record keeping keyboard and Remote Controller
- (4) Potential equalization
- (5) Receptacle for power cord
- (6) Place for network connector B-NET
- (7) Device plate
- (8) Place for network connector B-UPINET or B-UPI
- (9) Insertion slots for memory cards (optional)

System possibilities

The Datex-Ohmeda AS/3 Compact Anesthesia Monitor provides places for up to four single-width or two double-width modules. You can use the same modules and patient accessories in the Compact Anesthesia Monitor, Compact Critical Care Monitor, Anesthesia Monitor, and Critical Care Monitor.

Optional components for the AS/3 Compact Anesthesia Monitor are:

- built-in recorder in the F-CMREC frame or the recorder module, M-REC
- network functionality with B-UPINET board or B-CMNET board
- Data card and MemCard functionality with memory board, B-CMMEM, or the memory module, M-MEM

Communication between monitors

You can use the AS/3 Compact Anesthesia Monitor as a stand-alone monitor or for

- viewing and receiving data (alarms, vital signs) from other patient monitors
- gathering and storing data during transportation.

To view other patient monitors, the monitor needs to be connected to the Datex-Ohmeda Network. To gather, store, and transfer data between different Datex-Ohmeda monitors, use memory cards.

The green Data Card is for storage and transfer of patient data. The white Menu Card is used for storing the anesthesia record keeping menus and configurations, and for loading patient data and user modes from the Data Card.

Monitoring basics

You can control monitoring through the keys on the monitor's Command Board and side panels, modules, anesthesia record keeping keyboard and Remote Controller. The commands introduced in this User's Guide mainly focus on the Command Board hard keys and module keys.

Command Board - functions of some hard keys



Side panel keys

There are keys on the monitor's side panel for ON/Standby, NIBP, Invasive Pressures and Recorder functions. With these keys, you can start or end a function immediately.

NOTE: Recorder functions are available with the optional internal recorder or recorder module only.

Remote Controller

The Remote Controller allows access to the same menus as the Command Board. It also has some direct function keys that start or end a function immediately. To enter functions that do not have their own key, press the **Menu** key.



Navigating in menus

A menu is a list of functions or commands. To display a menu, press one of the hard keys. Selections in the menus are made with the ComWheel. For example, to change the ECG display:

- 1. Press the key to open the function menu.
- 2. Turn the ComWheel to select the desired function in the menu.
- 3. Push the ComWheel to open a submenu or an adjustment window.
- 4. Push the ComWheel to confirm the selection.

Common parts for modules

- Insertion guide slot (1)
- Module keys (2)
- (3) Patient cable connectors
- Module release latch (4)

Inserting a module

- 1. Align the module with the insertion guide.
- 2. Push the module into the Compact Monitor frame until it clicks.

Removing a module

- 1. Press the release latch.
- 2. Pull out the module.

You can connect and disconnect modules during monitoring.

Identical modules

Some modules are identical and cannot be used in the same system. See the following table when choosing modules for different parameters.

To monitor:	Select only one of these identical modules	
ECG (E), NIBP (N), SpO ₂ (S), Temp (T), Pressure (P) or Resp (R)	M-PRESTN, M-PRETN, M-RESTN, M-NE(12)STPR, M-NE(12)TPR, M-NE(12)STR, M-ESTPR, M-ETPR, M- ESTR, M-ESTP, M-EST or M-ETP.	
	NOTE: M-ESTP rev. 01, M-EST rev. 00 and M-ETP rev. 00 work only with S-STD93, S-STD94, S- ARK94, S-STD95, S-ARK95, S-STD96 and S- ARK96 software.	
	NOTE: M-PRESTN, M-PRETN and M-RESTN work only with L-ANEO1(A) and with rev. 10.9 of S-00A01, S-00A02, L-00A03 and L-00A04 software, or later versions.	
NIBP (N)	M-NIBP, M-PRESTN, M-PRETN, M-RESTN, M-NE(12)STPR, M-NE(12)TPR, M-NE(12)STR	
Pressure (P)	M-P, M-PP or M-PT	
C.O. (CO), Pressure (P) or SvO ₂ (Sv)	M-COP or M-COPSv	
CO ₂ (C), O ₂ (O), Spirometry (V), anesthetic agents(A), agent identification (i), gas exchange (X)	M-CO, M-COV, M-CAiOV, M-CAiO, M-CAiOVX, M-miniC	
Separate SpO ₂ measurement	M-NSAT or M-OSAT	
	NOTE: M-NSAT and M-OSAT can be used with M-PRESTN, M-PRETN, M-RESTN, M-NE(12)STPR, M-NE(12)STR, M-ESTPR, M-ESTR, M-ESTP and M-EST. The SpO ₂ measurement in M-NSAT and M-OSAT automatically overrides the SpO ₂ measurement in these modules.	
The following parameters have no identical modules:		
NMT	M-NMT	
EEG and AEP	M-EEG	
Tonometry (PgCO2)	M-TONO	
BIS	M-BIS	
Entropy	M-ENTROPY	



Setting up the monitor before use

Before starting to use the monitor, check the monitor installation settings and what is configured in different user modes, and make the necessary changes. The user modes are hospital-specific. The default modes are CPB Mode, General, Invasive, Regional, Neuro, Pediatric and Recovery. If you wish to make permanent changes, we recommend that you contact the person responsible for the configuration of the system. For more information about the default user modes, see the "Default Configuration Worksheet" delivered with the monitor. For more information about the installation settings and user modes, see the "User's Reference Manual."

Passwords

The default password for entering the *Install/Service* menu is 16 4 34. The default password for entering the *Save Modes* menu is 13 20 31.

Interfacing

- 1. Press Monitor Setup and select Interfacing.
- 2. Select desired interfaced internal modules or external monitors.

Setting time and date

NOTE: If the monitor is connected to the S/5 Central, it follows the Central's time settings and the *Time and Date* menu is not available.

NOTE: You cannot change the monitor's time settings after the case has been started.

- 1. Press Monitor Setup and select Time and Date.
- 2. Turn and push the ComWheel to set the time and date.

Changing the monitor installation settings

The monitor installation settings are the same in all user modes. The changes are preserved until changed again. Press **Monitor Setup** and select *Install/Service*, enter the password and select *Installation:*

Printer

ECG Printout Type: Select 25 mm, 50 mm or USA. Snapshot Printout: Select 12.5 or 25 mm/s. Printer Connection: Select printer connection (default: None). Paper Size: Select A4 or Letter (default A4).

Alarm options

Show limits: Select YES to show alarm limits in digit fields. Audio OFF: Select NO to disable alarm silencing. This hides the silencing options in the Audio ON/OFF menu in Alarms Setup. Latching Alarms: Select YES to keep alarm messages on screen until Silence Alarms key is pressed. Perminder Volume: Adjust volume of audible alarm reminder topo

Reminder Volume: Adjust volume of audible alarm reminder tone. **Alarm Tones**: Select alarm tones.

Display setup

Change the number of colors on the display (the result depends on the LCD display type.)

Monitor settings

Monitor Type: Monitor type defines the text on the start screen. *ARRWS Network*: Select **YES** if Arrhythmia Workstation is in the network.

ARK Settings: Set **Record Keeping ON/OFF** and change settings. **Parameter Settings**: Set tidal volume calculation conditions, CO₂ humidity compensation and inspired flow drawing direction.

Units

- Change units for height, weight, parameters, energy expenditure, laboratory values and calculations.
- You can change temperature units in the *Temp Setup* menu (Others - *Resp&Temp Setup* - *Temp Setup*) and CO₂ units in the *CO2 Setup* menu (Airway Gas - *CO2 Setup*). The changes are permanent.

Changing the user modes

NOTE: If you want to make changes in user modes, we recommend you contact the person responsible for the configuration. When new settings are saved, they should be marked in the "Default Configuration Worksheets." See below for instructions on how to change the modes permanently. Select the user mode you wish to change by pressing **Monitor Setup** and selecting **Select Mode**.

- 1. To make changes in:
- sweep speed, parameter colors, screen setup, Normal Screen layout, pages layout, trends, snapshots, press Monitor Setup. If necessary, select Install/Service. For instructions, see below.
- parameter setup, press a parameter key and go to the setup menu. For instructions, see parameter sections in User's Guide Part II.
- alarm limits and volume, press Alarms Setup. For instructions, see section "Alarms" in User's Guide Part II.
- 3. Confirm changes through Monitor Setup Install/Service Save Modes Save. You can save the changes also in other modes. If you do not save the changes in the modes, they are temporary and valid only until you reset the case or change the mode or until more than 15 minutes has elapsed from the turn-off of the monitor. Entering Save Modes requires a password, see "Passwords."

Changing the startup mode

- 1. Select Monitor Setup Install/Service Save Modes.
- 2. Select Startup Mode 1, 2, 3, 4, 5 or 6.

Renaming a mode

- 1. Select Monitor Setup- Install/Service Save Modes.
- 2. Select the mode, select *Name* and give a new name.

Loading modes

- 1. Select Monitor Setup Install/Service Save Modes.
- 2. Select *Load Modes* and load from/to card or from/to network.

NOTE: To load modes from the Data Card, you need the Menu Card in the other slot.

Changing the waveform sweep speeds

- 1. Select Monitor Setup Sweep Speeds.
- 2. Select the parameters and adjust the values. Slow waveforms show the amplitude changes better.

Changing the parameter colors

To change the colors for parameter waveforms, digits and trends,

select Monitor Setup - Install/Service -Colors.

Changing the recorder settings

- 1. Press Record/Print.
- 2. Select **Record Waveforms** and select the recorded waveforms, delay, paper speed and length, and select if you wish to record waveforms on alarms.
- Select *Record Trends* and set the numerical trend resolution and trend type, default trend type, and select the graphical trend recorded in upper and lower field.

Changing the printer settings

Select **Record/Print** - *Print Graphical* and select the pages to print and how many hours to print on one page.

Setting the Normal Screen format

Press Monitor Setup and select Screen Setup.

- Waveform Fields: Select the displayed waveforms.
- Digit Fields: Change the contents of a field or turn it off.
- **Split Screen**: Select what you wish to display with the waveforms (minitrends, spirometry, EEG, EP, ST or **None**).
- Minitrend Length: Select the length of the minitrend.

Changing the layout of other pages

You can check the contents of the pages by pushing the ComWheel in Normal Screen. To change the layout of the pages:

- 1. Press Monitor Setup and select Install/Service -Pages Setup.
- 2. Select the page and make the changes.

Setting the default trend

You can select graphical or numerical trends to be displayed by default.

- 1. Press Monitor Setup and select Install/Service Trends & Snapshot.
- 2. Select Default Trends and Graph or Num.

Configuring trends

To set the parameters displayed on the graphical trend pages:

- 1. Press Monitor Setup and select Install/Service Trends & Snapshots Graphical Trends.
- 2. Select the graphical trend page you want to change.
- 3. Select parameters for fields.

Setting trend length and time scale

Press Trends.

- Select *Time Scale* and the value.
- Select Trend Scales and adjust the scales.

Configuring snapshots

To change the snapshot setting, press **Monitor Setup** and select **Install/Service - Trends & Snapshots - Snapshot**:

- **Field x:** Select to display waveform, graphical trend or numerical trend.
- **Create on Alarms**: Select **Yes** (default) to create automatic snapshots for Tachy, Brady, Art high, Art low alarms. You can select other arrhythmia alarms to create snapshots through the **Arrhythmia Alarms** menu.
- Automatic Print: Select All to print all the snapshots immediately after creation, Alarms to print snapshots created on alarms or No to print only on request.
- **Print Loops**: Select **Yes** to print Patient Spirometry loops when snapshots are taken.

Cleaning and care

Daily and between patients	Once a month	Every six months
 Wipe the monitor surface. Wipe the ECG trunk cable, NIBP cuff and cables and SpO₂ sensors. Change or sterilize all airway and invasive patient accessories. Clean, disinfect or sterilize reusable temperature probes. Change the Tonometrics catheter between patients. Empty the D-fend water trap, see below. Change the BIS sensor between patients. Change the entropy sensor between patients. Check that the accessories, cables and monitor parts are clean and intact. 	 Perform gas calibration for gas exchange monitoring, see below. Refresh the batteries, see below. Check the fan filter on the monitor's side panel, on the gas module's front panel and under the display unit. Clean if necessary: Pull out the filter. Wash it in detergent solution and allow to dry before reinserting. Do not use pressurized air. Replace the filter if it is damaged. Change the D-fend water trap every two months and when 'Replace D-fend' appears. 	 Perform gas calibration for tonometry and airway gas monitoring, see below.
Permitted detergents	Permitted disinfectants	DO NOT !
 Datex-Ohmeda Cleaning Fluid Other mild detergents 	 Ethanol Isopropyl alcohol Chlorite compounds Glutaraldehyde 	 Do not use hypochlorite, acetone-, phenol- or ammonia based cleaners. Do not autoclave the device or its parts. Do not immerse any part of the device in liquids, or allow liquid to enter the interior. Do not apply pressurized air to any outlet or tubing connected to the monitor.

Before cleaning

1. Turn off the monitor from the power switch.

2. Disconnect the power cord.

After cleaning

- 1. Let dry completely.
- $2. \ \ \, {\rm Reconnect\ the\ power\ cord\ and\ turn\ on\ the\ monitor.}$

More comprehensive checking

See the "Technical Reference Manual."

WARNING: After cleaning or if liquid has accidentally entered the monitor, ensure that every part of the monitor is dry before reconnecting it to the power supply.

D-fend water trap

- Empty the container whenever half full.
- Change the D-fend or Mini D-fend water trap every two months and when 'Replace D-fend' appears.
- The water trap cartridge is disposable. Do not wash or reuse the cartridge.
- Change the green D-fend+ water trap every 24 hours and when 'Replace D-fend' appears.

Reusable D-lite sensor

The reusable D-lite sensor can be washed in a washing machine and steam autoclaved. Make sure that the sensor is dry and connectors are not damaged. A tight connection is essential for correct measurement.

Other accessories

For information on how to clean and check reusable accessories, see the accessory package. Do not reuse single-use disposable accessories.

Refreshing the batteries

- 1. Turn on the monitor.
- 2. Connect the monitor power cord to power supply for at least 10 hours.
- 3. Remove the power cord and wait until the monitor turns off. Then wait for another 15 minutes.
- 4. Reconnect the power cord to power supply and charge the battery for at least 10 hours.

Changing fuses

- 1. Remove the power cord.
- 2. Remove the fuse holder by pushing the locking pin and pulling the holder gently out.
- 3. If a fuse is blown, ensure that you replace it with a fuse of the correct type and rating.

Calibrating

- 1. Turn on the power. Let the monitor warm up for 30 minutes.
- 2. Attach a regulator to the calibration gas container.

NOTE: % is used for CO_2 regardless of selected units.

NOTE: See section "Supplies and accessories" for correct regulator and gas.

NOTE: Ensure that the calibration gas and regulator are functioning properly before calibration. Perform annual maintenance on the regulator as required.

Calibrating airway gases

Follow the recommended calibration intervals (every six months in normal use and every two months in continuous use) to ensure that the measurement accuracy remains within specifications.

- 1. Attach a new sampling line to the water trap. Connect the other end of the sampling line to the regulator on the gas container.
- 2. Press the **Airway Gas** key.
- 3. Select Gas Calibration.
- 4. Wait until the texts 'Zero OK' and then 'Feed Gas' appear on the screen, open the regulator and start feeding gas. Push the ComWheel and continue feeding gas until the text 'Adjust' is displayed.
- 5. Check that the displayed gas values match the values on the calibration gas container. Adjust with the ComWheel, if necessary.

NOTE: Calibrate M-miniC module with calibration gas 755580 only and set the ${\rm O}_2$ concentration to 20%.

Calibrating Patient Spirometry

Perform flow calibration once a year or when the difference between the inspiratory and expiratory volumes is permanent. See the "Technical Reference Manual" for more information.

Calibrating gas exchange

To ensure gas exchange accuracy, perform the airway gas calibration once a month and patient spirometry calibration once a year. Use 2-m (7-ft) airway gas sampling line.

Calibrating tonometry

- 1. Connect the calibration gas sampling line to the D-gate regulator and to the module's catheter connector.
- 2. Press the Others key and select Tonometry PgCO2 Calibration.
- 3. Wait until the text 'Feed gas' appears. Open the regulator and start feeding gas until the text 'Adjust' appears.
- 4. Close the regulator.
- 5. Check that the displayed values match the values on the calibration gas container. Adjust with the ComWheel, if necessary, and confirm.
- 6. If airway gases are monitored, calibrate the gas module at the same time.

NOTE: Use only Datex-Ohmeda Calibration gas sampling line; wrong line length or diameter can cause incorrect calibration.

NOTE: Do not wash or disinfect calibration gas sampling lines.

Calibration check of temperature, NIBP and invasive blood pressures

Calibration check of temperature, NIBP and invasive blood pressures should be performed at least once a year by qualified service personnel as a part of the Planned Maintenance, see the "Technical Reference Manual."

Alarm basics

Enabling the alarms

To enable the alarms, connect the patient cables. If the alarm source is selected, the alarms are operative also when the measurement is not displayed (except the respiration measurement alarms).

Alarm indications

When an alarm becomes active, messages appear in the order of priority. The alarming measurement value flashes and its background color indicates the alarm category (see the table below). In some cases, there may be a message on the display giving more detailed information. An audible alarm is also triggered.

For details about alarms, see section "Alarms" in User's Guide Part II.

NOTE: If the monitor is connected to the network, the alarms can be heard and seen on the S/5 Central as well. Please consult the "Datex-Ohmeda S/5 Central User's Reference Manual: Alarms" for details.

Alarm categories

The priority depends primarily on the cause and alarm duration.

Visual	Meaning	Tone pattern (selected when the system is configured)	Front panel LED
Red	For life threatening situations	Triple + double beep every 5 seconds or continuous beep	red LED lit
Yellow	For serious but not life threatening problems	Triple beep every 19 seconds or double beep every 5 seconds 19 / 5 5	yellow LED blinking
White	Advisory	Single beep -	yellow LED lit

Performance

WARNING: Operation of the monitor outside the specified values may cause inaccurate results.

Datex-Ohmeda Compact Anesthesia Monitor

Power supply

100 to 240 V 50/60 Hz
±10 %
140 VA

Battery operation

Batteries:	NiCd
Charging time:	5 hours
Operation time:	up to 1 hour

Environmental conditions

Operating	
temperature:	+10 to+35°C (50 to 95°F)
Storage and transpo	ort
temperature:	-10 to +50°C (14 to 122°F)
Relative humidity:	10 to 90 % noncondensing,
in airway	0 to 100 % condensing
Atmospheric	
pressure:	660 to 1060 mbar
	(500 to 800 mmHg)

Alarm behavior

The maximum alarm delay of the alarm at the monitor signal output to network: 1.1 seconds If the alarm mode is latched, the technical alarms are latched as well. This does not comply with the NIBP (IEC 60601-2-30) and invasive pressure (IEC 60601-2-34) standard requirements. Silencing alarms for 5 minutes does not comply with the SpO₂ (ISO 9919) standard requirements.

Hemodynamic Modules M-NE(12)STPR, M-NE(12)TPR, M-NE(12)STR, M-ESTPR, M-ETPR, M-ESTR; M-P, M-PP, M-PT; M-NIBP

Letters in the module name stand for: N= NIBP, E= ECG, 12 = up to 12 ECG leads, S= Pulse oximetry, T= Temperature, P= Invasive blood pressure, R= Impedance respiration

ECG

Filter modes: monitoring filter 0.5 to 30 Hz ST filter 0.05 to 30 Hz 0.05 to 100 Hz diagnostic filter - 12-lead ECG: 0.05 to 150 Hz With 60 Hz power supply frequency: monitoring filter 0.5 to 40 Hz ST filter 0.05 to 40 Hz Defibrillation protection: 5000V, 360J Recovery time: 2 seconds Heart rate: Measurement range: 30 to 250 bpm Measurement accuracy: ±5% or ±5 bpm Display averaging time: 5 seconds Display update time: 5 seconds Pacemaker pulse detection: detection level: 2 to 500 mV pulse duration: 0.5 to 2 ms

Impedance respiration

Temperature

Invasive blood pressure

Measurement range: -40 to 320 mmHg Measurement accuracy: ±5% or ±2 mmHg

Pulse rate:

NIRP

Measurement range: adult 25 to 260 mmHg. child 25 to 195 mmHg. infant 15 to 145 mmHg Pulse rate range accepted: 30 to 250 bpm Typical measuring time: adult 23 s. infant 20 s

NOTE: NIBP measurement is intended for patients weighing over 5 kg (11 lb.)

Pulse oximetry

Proportional scaling of plethysmographic waveform. Sp02

Measurement and display range: 40 to 100% Calibration range: 50 to 100% Calibrated against functional oxygen saturation. Measurement accuracy (% SpO₂ \pm 1SD) **1**):

100 to 80%, ±2 digits; 80 to 50%. ±3 digits: 50 to 40%, unspecified Display update time: 5 seconds Display averaging time: adjustable: 10 sec. 20 sec or beat-to-beat Display resolution: $1 \text{ digit} = 1\% \text{ of } \text{SpO}_2$ Pulse rate: Measurement and display range: 30 to 250 bpm Measurement accuracy: ±5% or ±5 bpm

2)

Default alarm limits

SpO₂ high Off. low 90%: PR high 160. low 40

NOTE: Pulse oximetry measurement is intended for patients weighing over 3 kg (6.6 lb).

Hemodynamic modules M-PRESTN, **M-PRETN and M-RESTN**

Letters in the module name stand for: P= Invasive blood pressure: R= Impedance respiration; E= ECG; S= Pulse oximetry; T= Temperature: N= NIBP

ECG³⁾

Filter modes: 0.5 to 30 Hz monitoring filter ST filter 0.05 to 30Hz diagnostic filter 0.05 to 150 Hz With 60 Hz power supply frequency: monitoring filter 0.5 to 40 Hz ST filter 0.05 to 40 Hz Defibrillation protection: 5000 V. 360 J Recovery time: 5 seconds Heart rate:

```
Measurement range: 30 to 250 bpm
Measurement accuracy: ±5 % or ±5 bpm
Display averaging time: 5 seconds
Display update time: 5 seconds
```

Average heart rate response time and time range of response time:

Response time 80 to 120 bpm: 7.9s (6.4 to 9.1 s) Response time 80 to 40bpm; 9.9s (8.3 to 11.4 s) Maximum Tall T wave amplitude that does not disturb the heart rate calculation: 2.2 mV The heart rate calculation operates with irregular rhythms of ANSI/AAMI EC13 3.1.2.1 as follows: a): 75bpm b): 61bpm

c): 115bpm

d): 97bpm

Pacemaker pulse detection: 2 to 700 mV detection level: pulse duration: 0.5 to 2 ms The monitor is specified for both of the methods A and B required in EC13 section 4.1.4.2. Offset voltage range: +800 mV CMRR: >95 dB Pacer pulse rejection of fast ECG signals: 1.29 V/s according to the test defined in FC13 section 4.1.4.3.

Pacemaker detector may not operate correctly during the use of high-frequency (HF) surgical equipment. The disturbances of HF surgical equipment typically cause false positive pacer detection.

1)

Accuracy is based on deep hypoxia studies using Datex-Ohmeda FingerSat Sensors on volunteered subjects. Arterial blood samples have been analyzed by a Radiometer OSM Co-oximeter, 1 standard deviation = 68 % of all readings in the specified range in stable conditions. 2)

Limits are adjustable: OFF to 51% for SpO₂ high 50 to 100% for SpO_2 low 250 to 35 bpm for PR high 30 to 245 bpm for PR low

³⁾ The isolation barrier capacitance in the module has been minimized to reduce the hazard of burns in the event of a defect in the ESU return electrode connection.

Direct current for leads-off detection through an active patient electrode : 25 nA

Direct current for leads-off detection through a reference electrode: 225 nA

The normalized respiration sensing current between RA (R) and LL (F) or RA (R) and LA (L) or LA (L) and LL (L): $3.2 \ \mu$ A

Frequency of respiration sensing current: 31.25 kHz Minimizing the effects of the line isolation monitor transients:

Crystal controlled oscillator used as the operating frequency source of the patient isolation power supply.

The average time and time range () to alarm for tachycardia are as follows (ANSI/AAMI EC13 3.1.2.1.g):

Figure 4a halved amplitude: $6.5 \pm (6.1 \pm 7.1 \pm 1.5)$ Figure 4a normal amplitude: $5.3 \pm (4.9 \pm 5.7 \pm 5.7 \pm 5.5)$ Figure 4a doubled amplitude: $5.8 \pm (5.5 \pm 5.6 \pm 6.2 \pm 5.5)$ Figure 4b halved amplitude: $5.0 \pm (4.5 \pm 5.6 \pm 5.6)$ Figure 4b normal amplitude: $5.4 \pm (4.6 \pm 5.6)$ Figure 4b doubled amplitude: $5.3 \pm (4.6 \pm 5.8 \pm 5.6)$

The ECG measurement fulfils the requirements of the ANSI/AAMI EC11 3.2.7.2/4.2.7.2 by using the test methods a, b, c and e.

Direct cardiac application:

The display area reserved for the ECG measurement in the monitoring system screen may not be adequate for displaying the complete ECG amplitude when measuring ECG direct from the surface of the heart. Clipping of the signal can be reduced by adjusting the size of the signal on the display (for example, from the default 1.0 to 0.2) in the ECG menu.

Auxiliary output, ECG:

Bandwidth of auxiliary output: 0.5 to 30Hz Gain: 1mV ECG signal is 1V at the auxiliary output.

Propagation delay: < 15ms.

The pacemaker pulses have been replaced with 2ms fixed digital pulses at the ECG output for IABP or defibrillator equipment.

An auxiliary device that fulfils the requirements of the IEC 60601-1 standard can be connected to the auxiliary output. There are no other limitations, because the auxiliary output of the monitor is galvanically isolated from patient applied part of the ECG measurement.

Impedance respiration

Respiration range:4 to 120 resp/minAccuracy: \pm 5% or \pm 5 resp/min

The EMC immunity of the respiration measurement has been tested with 1 Vrms and 1 V/m. This level has been used for optimizing the immunity of the respiration measurement to damp the operating frequency of the electrosurgery equipment.

NOTE: Impedance respiration measurement is intended for patients over three years old.

Invasive blood pressure⁴⁾

Measurement range:-40 to 320 mmHg Measurement accuracy: ± 5 % or ± 2 mmHg

Pulse rate:

Temperature ⁴⁾

Measurement range: 10 to 45°C (50 to 113°F) Measurement accuracy:

±0.1 °C (25 to 45.0 °C) ±0.2°C (10 to 24.9°C) Measurement accuracy with single-use sensors: ±0.2 °C (25 to 45 °C) ±0.3 °C (10 to 24.9 °C) Probe type: Use only Datex-Ohmeda temperature probes or defibrillator-proof YSI 400 series probes. Time constant of temperature probes: Reusable skin temperature probe: 3 s Reusable adult central temperature probe: 6 s Reusable pediatric central temperature probe: 4 s Disposable skin temperature probe: 3 to 6s Disposable central temperature probe, 12F: 5 to 8 s Disposable central temperature probe, 9F: 5 to 8 s Esophageal stethoscope with temperature probe, 9F: 15 s Esophageal stethoscope with temperature probe, 12F: 16 s Esophageal stethoscope with temperature probe, 18F⁵⁾: 23 s Esophageal stethoscope with temperature probe, 24F⁵): 32 s

4) The isolation barrier capacitance in the module has been minimized to reduce the hazard of burns in the event of a defect in the ESU return electrode connection.

5) Response time of the probe exceeds 150 s.

NIRP

Measurement range: adult 25 to 260 mmHg child 25 to 195 mmHg infant 15 to 145 mmHg Pulse rate range accepted: 20 to 250 bpm Typical measuring time: adult 23 s. infant 20 s Overall system accuracy:

Meets or exceeds SP10-2002 AAMI standards

The ESU does not cause a burn hazard through the NIBP cuff, because there is no electrical connection between the cuff and the NIBP measuring electronics.

NOTE: NIBP measurement is intended for patients weighing over 5 kg (11 lb.)

$Sp0_{2}^{6}$

Measurement and display range: 40 to 100 % Calibration range: 70 to 100 % Calibrated against functional oxygen saturation. Measurement accuracy $^{7)}$ (% SpO₂ ±1SD): 100 to 70 %, ±2 digits, ±3 digits during clinical patient motion: 69 to 40 %, unspecified Display update time: 5 seconds continuous, defined by the main software of the monitor Display resolution: $1 \text{ digit} (1\% \text{ of } \text{SpO}_2)$

Wavelength of SpO₂ probe LEDs:

Infrared I FD 900 nm Red I FD 660 nm Maximum energy of SpO₂ probe LEDs: Infrared I FD 42 µJ/pulse Red I FD 62 µJ/pulse

Pulse rate:

Measurement and display range: 30 to 250 bpm Measurement accuracy: ±5 % or ±5 bpm

Default alarm limits⁸⁾.

SpO₂ high Off. low 90% PR high 160. low 40

Cardiac Output Modules, M-COP and M-COPSv

Pressure performance as above.

Cardiac output

Measurement range: 0.1 to 20 l/min Repeatability: ±2 % or 0.02 l/min Catheters: Edwards Lifesciences Corp. compatible

SvO₂

Catheters:

Measurement range: 1 to 98% Measurement accuracy:

±2% SvO₂ equals 1 standard deviation for range of 30% to 95% SvO2 and 6.7 to 16.7 g/dl Hb when using in vivo calibration. Edwards Lifesciences Corp. SvO₂ catheter

RFF

Measurement range: 1 to 85% Repeatability: ±2% as measured by

electronically generated pulsatile curves for range 10 to 60%. For other ranges accuracy is unspecified. Edwards Lifesciences Corp. RFF catheter

Catheters:

6) The isolation barrier capacitance in the module has been minimized to reduce the hazard of burns. in the event of a defect in the ESU return electrode connection.

7) Accuracy is based on deep hypoxia studies with volunteered subjects during motion and nonmotion conditions over a wide range of arterial blood oxygen saturations as compared to arterial blood CO-Oximetry.

⁸⁾ Limits are adjustable: OFF to 51% for SpO_2 high

50 to 100% for SpO₂ low 250 to 35 bpm for PR high 30 to 245 bpm for PR low

Datex-Ohmeda Compatible Saturation Module, M-OSAT

Automatic scaling of plethysmographic waveform.

SpO₂

Calibrated against functional saturation.

Pulse rate

Measurement and display range: 30 to 250 bpm Measurement accuracy:

±2 % or ±2 bpm (whichever is greater) Resolution: 1 bpm Display averaging time: 12 seconds Display update time: 5 seconds

9)

Default alarm limits SpO₂ high Off, low 90% PR high 160, low 40

Nellcor Compatible Saturation Module, M-NSAT

Automatic scaling of plethysmographic waveform.

$\pmb{\text{Sp0}}_2$

Measurement and display range: 20 to 100 % Calibration range: 70 to 100 %

Calibrated against functional oxygen saturation.

Pulse rate

Measurement and display range: 30 to 250 bpm Measurement accuracy: ±3 bpm

9)

Default alarm limits

SpO₂ high Off, low 90% PR high 160, low 40

Tonometry Module, M-TONO Gastrointestinal PCO₂ (PgCO₂)

9)

10)

The accuracy value depends on the sensor used. Accuracy is based on Nellcor protocol #081400-N, Non-invasive Controlled Hypoxia Study Rev. B. **11**

Typical value.

Compact Airway Modules, M-CO, M-COV, M-CAiO, M-CAiOV, M-CAiOVX

Letters in the name stand for: $C = CO_2$ and N_2O , O = Patient O_2 , A = Anesthetic agents, i = Agent identification, V = Patient Spirometry, X = Gas exchange

 12)

 Sampling rate

 12)

 Sampling rate

 12)

 Sampling delay

 2.5 s typical with a 3-m sampling line

Total system response time:

2.9 seconds typical with a 3-m sampling line, including sampling delay and rise time

12) Warm-up time :

2 to 5 min, 30 min for full spec.

13) Default alarm limits

 $\label{eq:starsest} \begin{array}{l} \mbox{EtCO}_2 \mbox{ high 8\%, low 3\%} \\ \mbox{FiEnf high 5.1\%, low 0ff} \\ \mbox{FiCO}_2 \mbox{high 3\%, low 0ff} \\ \mbox{EtEnf high 3.4\%, low 0ff} \\ \mbox{EtO}_2 \mbox{high 0ff, low 10\%} \\ \mbox{Filso high 3.4\%, low 0ff} \\ \mbox{FiO}_2 \mbox{high 0ff, low 18\%} \\ \mbox{EtIso high 2.3\%, low 0ff} \\ \mbox{FiDes high 18\%, low 0ff} \\ \mbox{FiDes high 12\%, low 0ff} \\ \mbox{FiDes high 1.5\%, low 0ff} \\ \mbox{EtIso high 5.1\%, low 0ff} \\ \mbox{FiSev high 5.1\%, low 0ff} \\ \mbox{FiSev high 3.4\%, lo$

Non-disturbing gases, maximum effect on readings $CO_2 < 0.2$ vol%, N₂O, O₂ < 2 vol%, anesth, agents: < 0.15 vol%: Ethanol C₂H₅OH <0.3% Acetone < 0.1% < 0.2% Methane CH₄ Nitrogen N₂ Carbon monoxide CO <200 ppm Nitric oxide NO water vapor Effect of Helium: decreases CO₂ readings <0.6 vol% typically

Carbon dioxide (CO₂)

Oxygen (O₂)

Nitrous oxide (N₂0)

Respiration rate (RR)

Anesthetic agent (AA)

Measurement rise time: < 400 ms typical Gas cross effects: $< 0.15 \text{ vol}\% \text{ N}_2 \text{O}$ Halothane, Isoflurane, Enflurane Measurement range: 0 to 6 % 12) Accuracy : ± 0.2 vol % Sevoflurane Measurement range: 0 to 8 % 12) ± 0.2 vol % Accuracy : Desflurane Measurement range: 0 to 20 % 12) Accuracy : $0 \text{ to } 5\% \pm 0.2 \text{ vol }\%$. 5 to 10 % ± 0.5 vol % $10 \text{ to } 20 \% \pm 1.0 \text{ vol } \%$

Agent identification

12) Identification threshold : 0.15 vol %

12)

Typical value.

13) Alarm limits and their adjustment range may vary depending on the mode used.

Patient Spirometry

Using D-lite (+) or Pedi-lite(+) flow sensor and gas sampler:

D-lite(+)	Pedi-lite(+)
Tidal volume:	
Measurement range	

weasurer	nent lange.	
	150 to 2000 ml	15 to 300 ml
	14)	
Accuracy	:	
	$\pm 6\%$ or 30 ml	$\pm6\%$ or 4 ml
Minute v	olume:	
Measurer	nent range:	

Measurement range: 2 + 201/min

2 to 20	0 l/min 0	.5 to 5 l/min
14)		
Accuracy : ±6	% ±	6%

Airway pressure:

Measurement range: -20 to +100 cmH₂0 - 20 to +100 cmH₂0 I:E: 1:4.52:1 14) Accuracy $\pm 1 \text{ cmH}_2\text{O}$ $\pm 1 \text{ cmH}_2\text{O}$

Flow:

Measurement range:

1.5 to 100 l/min 0.25 to 25 l/min

Compliance:

Measurement range:

4 to 100 ml/cmH₂O 1 to 100 ml/cmH₂O

Airway resistance:

Measurement range:

0 to 40 cm H₂O/I/s 0 to 40 cm H₂O/I/s

Sensor specifications:

	D-lite(+)	Pedi-lite(+)
Dead space:	9.5 ml	2.5 ml
Resistance:		
at 30 l/min	0.5 cmH ₂ 0	
at 10 l/min		$1.0 \text{ cmH}_2\text{O}$

Gas Exchange

VO₂ and VCO₂:

Measurement range: 20 to 1000 ml/min 14) Accuracy (valid for respiration rates 4 to 35 breaths/min): ±10% or 10 ml FiO₂ < 65% 65%<Fi02<85% ±15% or 15 ml

RQ, Respiratory Quotient $(=VCO_2/VO_2)$

Measurement range: 0.6 to 1.2

Detection through D-lite flow sensor and gas sampler (see the measurement ranges and sensor specifications above.)

Measurement not valid with $O_2 + N_2O$ mixtures.

Specifications apply only without condensation at measurement point.

Single-width Airway Module, M-miniC

Sampling rate:	150 ±25 ml/min (sampling line 2 to 3 m, normal conditions)
Sampling delay:	2.1 s typical with a 3-m sampling line
Total system respon	nse time: 2.4 seconds typical with a 3-m sampling line, including sampling delay and rise time (typically 3.7 seconds with a 6-m sampling line)
Warm-up time:	1 min for operation with CO_2 30 min for full specification

Non-disturbing gases are those with a maximum effect on the CO₂ reading < 0.2 vol%. The effect is valid for specific concentrations shown in parentheses of the non-disturbing gas: Ethanol C_2H_5OH (<0.3%) Acetone (<0.1%) Methane CH_4 (<0.2%) Nitrogen N₂ water vapor Trichloromonofluoromethane (<1%) Dichlorotetrafluoroethane (<1%) Dichlorofluoromethane (<1%)

14)

Typical value

Single-width Airway Module, M-miniC (cont.)

Disturbing gases and their effect on the CO_2 reading at 5.0 vol-% CO_2 are shown below. Errors listed reflect the effect of specific concentrations (shown in parentheses) of an individual disturbing gas and should be combined when estimating the effect of gas mixtures:

 $\label{eq:second} \begin{array}{l} \mbox{Halothane (4\%) increases < 0.3 vol\%} \\ \mbox{Isoflurane (5\%) increases < 0.4 vol\%} \\ \mbox{Enflurane (5\%) increases < 0.4 vol\%} \\ \mbox{Desflurane (24\%) increases < 1.2 vol\%} \\ \mbox{Sevoflurane (6\%) increases < 0.4 vol\%} \\ \mbox{Helium (50\%) decreases < 0.3 vol\%} \\ \mbox{If } O_2 \ compensation is not activated:} \\ \mbox{O}_2 \ (40 \ to \ 95\%) \ decreases < 0.3 vol\% \\ \mbox{If } O_2 \ compensation is activated:} \\ \mbox{O}_2 \ (40 \ to \ 95\%) \ error < 0.15 \ vol\% \\ \mbox{If } N_20 \ compensation is not activated:} \\ \mbox{N}_20 \ (40 \ to \ 80\%) \ increases < 0.4 \ vol\% \\ \mbox{N}_20 \ (40 \ to \ 80\%) \ increases < 0.3 \ vol\% \\ \mbox{If } N_20 \ compensation is activated:} \\ \mbox{N}_20 \ (40 \ to \ 80\%) \ increases < 0.3 \ vol\% \\ \mbox{If } N_20 \ compensation is activated:} \\ \mbox{N}_20 \ (40 \ to \ 80\%) \ increases < 0.3 \ vol\% \\ \mbox{If } N_20 \ compensation is activated:} \\ \mbox{N}_20 \ (40 \ to \ 80\%) \ increases < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \\ \mbox{If } N_20 \ (40 \ to \ 80\%) \ error < 0.3 \ vol\% \ error < 0$

15)

Default alarm limits

 $EtCO_2$ high 8%, low 3% $FiCO_2$ high 3%, low Off

Carbon dioxide (CO₂)

Valid for respiration rate < 40 1/min at I:E ratio of 1:1. (Relative error is typically 10% for respiration rate 80 1/min at I:E ratio of 1:1.) The accuracy is specified in simulated ventilation. With higher respiration rates and with varying ventilation methods the specifications may not be met.

Respiration rate

Breath detection: 1% change in CO₂ level Measurement range: 4 to 80 breaths/min Accuracy:

 $\begin{array}{ll} \pm 1/\text{min} \text{ in the range 4 to 20 I}/\text{min} \\ \pm 5\% \text{ in the range 20 to 80 I}/\text{min} \\ \text{Resolution:} & 1/\text{min} \end{array}$

NOTE: M-miniC is intended for patients weighing over 5 kg (11 lb).

Neuromuscular Transmission Module, M-NMT

NMT

Stimulation modes: Train of four, TOF Double burst (3.3), DBS; Single twitch, ST 50 Hz tetanic & post tetanic count, PTC Stimulus current range: supramax 10 to 70mA manual 10 to 70 mA with 5 mA steps Stimulus current accuracy: 10% or ±3 mA

Regional block mode (plexus)

Stimulation mode: Single twitch Stimulus current range: 0 to 5.0 mA with 0.1 mA steps Stimulus current accuracy: 20% or ±0.3 mA

15)

Alarm limits and their adjustment range may vary depending on the mode used.

EEG Module, M-EEG

EEG

Sampling frequency: 100 Hz per channel Range: +400 uV Frequency range: 0.5 to 30 Hz 60 nV Resolution: Input impedance: $>8 M\Omega$ at 10 Hz <0.5 µV rms from Noise level: 0.5 Hz to 30 Hz CMRR: >100 dB at 50 Hz Parameters from power spectrum: SEF, MF, relative power in frequency bands Burst suppression ratio (BSR)

AEP

Stimulation

Click (condensating): duration 100 µs Frequency: 1.1 to 9.1 Hz (1 Hz steps at 10 ms meas.) Intensity: 10 to 90 dB nHL, 10 dB steps

Measurement

Sampling frequency: 2400 Hz for MLAEP/ 4800 for BAEP

Frequency range: 0.5 to 1000 Hz Highpass filter: off/10/30/50/75/100/150 Hz

Single average:

Averaged responses: 100 to 2000 stimuli

Moving average:

Gross average: 100 to 2000 stimuli Update interval: after every 100 stimuli (200, when gross average is 2000)

EMG

Frequency range: 60 to 300 Hz Parameter displayed: Amplitude (RMS)

BIS Module, M-BIS

EEG

Epoch duration: 2 seconds Artifact rejection: automatic 25 to 400 uV EEG scales: EEG sweep speeds: 12.5/25/50 mm/sec **Bispectral index:** 0 to 100 Signal quality index: 0 to 100 30 to 80 dB (70 to 110 Hz) FMG: Suppression Ratio: 0 to 100% Update rate: 1 second for BIS index ON (2 to 70 Hz with notch), Filters: OFF (0.25 to 100 Hz) Smoothing rate: 15 seconds (default) or 30 seconds Mode: sensor automatically selects mode

DSC (Digital signal converter)

Analog to digital converter:

 noise-shaped sigma-delta

 Sampling rate:
 16 384 samples/second

 Resolution:
 16 bits at 256

 samples/second
 samples/second

 Input impedance:
 50 Mohms minimum

 Noise:
 <0.3 μV RMS</td>

 (2.0 μV peak-to-peak)
 0.25 Hz to 50 Hz

Common mode rejection (Isolation mode): 110 dB at 50/60 Hz to earth ground Bandwidth: 0.16 to 450 Hz

Entropy Module, M-ENTROPY

Entropy parameters:

Response Entropy (RE): range 0 to 100 State Entropy (SE): range 0 to 91 Burst Suppression Ratio (BSR): range 0 to 100 % Display resolution: 1 digit

Entropy EEG

Scales:+Sweep speed:1Resolution:6

 $\pm 25/50/100/250/400~\mu\text{V}$ 12.5/25/50 mm/s 60 nV

Amplifier and A/D conversion

Amplification:10000Sampling frequency:1600 HzFrequency range:0.5 to 118 HzResolution:60 nV

CAUTION: The entropy measurement is to be used as an adjunct to other physiological parameters in assessing the effects of certain anesthetic agents.

27

Abbreviations

/min	beats per minute, breaths per minute
°C	
°F	Celsius degree
-	Fahrenheit degree
μg	microgram
A	arm (describing location)
A	alveolar
a	arterial
	arterio-alveolar PO ₂ ratio
AaDO ₂	alveolo-arterial oxygen difference
AA	anesthetic agent
AAMI	Association for the Advancement
	of Medical Instrumentation
ABG	arterial blood gases
ABP	arterial pressure
ADU	Anesthesia Delivery Unit
AEP	auditory evoked potential
AirW	airway temperature
Alpha, Al	alpha frequency band
AM	Anesthesia Monitor
Amp	amplitude
Ant	anterior
APN	apnea
Arrh.	arrhythmia
Art	arterial pressure
ASY	asystole
ATMP	atmospheric pressure
ATPD	atmospheric/ambient
	temperature and pressure, dry gas

ATPS AV aVF avg aVL aVR aw Axil	ambient temperature and pressure, saturated gas atrioventricular left foot augmented lead average left arm augmented lead right arm augmented lead airway axillatory temperature
BAEP	brainstem auditory evoked potential
Bal	balance gas
bar	1 atmosphere
Beta, Be	beta frequency band
Bigem.	bigeminy
BIS	bispectral index
Blad	bladder temperature
Blood	blood temperature (C.O.
	measurement)
Body	body temperature
BP	blood pressure
Brady	bradycardia
BSA	body surface area
BSR	burst suppression ratio
B-to-B	beat-to-beat
BTPS	body temperature and pressure, saturated gas

С	chest
C(a-v)02	arteriovenous oxygen content
	difference
C.C.O.	continuous cardiac output
C.I.	cardiac index
C.O.	cardiac output
cal.	calibration
Calc	calculated/derived value
Calcs	calculations
CAM	Compact Anesthesia Monitor
	arterial oxygen content
Casc.	cascaded (ECG)
CC	cubic centimeter
CCCM	Compact Critical Care Monitor
CCM	Critical Care Monitor
	capillary oxygen content
CCU	cardiac (coronary) care unit
CEL	Celsius degree
CISPR	International Special Committee
	on Radio Interference
cmH ₂ O	centimeter of water
CMRR	common mode rejection ratio
CO	carbon monoxide
CO_2	carbon dioxide
COHb	carboxyhemoglobin
Compl	compliance
Cont.	continuous
Contrl	controlled ventilation

calculated/derived value

С

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Core	core temperature	ECG2	second ECG waveform	Exp	expiratory
Count	count of responses	ECG3	third ECG waveform	F	foot (describing location)
CPB	cardiopulmonary bypass	ED	emergency department	FAH	Fahrenheit degree
CPP	cerebral perfusion pressure	EDV	end-diastolic volume	FEMG	frontal electromyogram
CSA	compressed spectral array	EDVI	end-diastolic volume index	FFT	fast Fourier transform
СТ	computer tomography	EE	energy expenditure (kcal/24h)	FI, Fi	fraction of inspired gas
CvO ₂	(mixed) venous oxygen content	EEG	electroencephalogram	FiAA	fraction of inspired anesthetic
CVP	central venous pressure	EEG1	first EEG waveform		agent
		EEG2	second EEG waveform	Fib	fibrillation
d	day	EEG3	third EEG waveform	FiBal	fraction of inspired balance gas
dB	decibel	EEG4	fourth EEG waveform	FiCO ₂	fraction of inspired carbon dioxide
DBS	double burst stimulation (NMT)	EEMG	evoked electromyogram	FiN ₂	fraction of inspired N ₂
DEL	delete	EEtot	total energy expenditure	FiN ₂ O	fraction of inspired nitrous oxide
Delta, De	delta frequency band	elect	electrode	FiO ₂	fraction of inspired oxygen
depr.	depression	elev.	elevation	Flow	airway gas flow
Des	desflurane	EMC	electromagnetic compatibility	Freq.	frequent
Dia	diastolic pressure	EMG	electromyogram	ft	foot, feet
Diagn	diagnostic (ECG filter)	Enf	enflurane	FVloop	flow volume loop
DIFF	difference	Entr	entropy		
DIS	S/5 Device Interfacing Solution	EP	evoked potential	g	gram
DO_2	oxygen delivery	ESD	electrostatic discharge	Graph.	graphical
DO_2I	oxygen delivery index	Eso	esophageal temperature		
DSC	digital signal converter	ESV	end-systolic volume	h	hour
DSSS	Direct Sequence Spread	ESVI	end-systolic volume index	Н	hand (describing location)
	Spectrum	ET, Et	end-tidal concentration	Hal	halothane
dyn	dynamic	EtAA	end-tidal anesthetic agent	Hb	hemoglobin
		EtBal	end-tidal balance gas	Hbtot	total hemoglobin
е	estimated	EtCO ₂	end-tidal carbon dioxide	HCO3-	bicarbonate
ECG	electrocardiogram	EtN_2O	end-tidal nitrous oxide	Hemo	hemodynamic
ECG1	first ECG waveform (top)	EtO ₂	end-tidal oxygen	Hemo Calc	s hemodynamic calculations
ECG1/r	real-time ECG	ET-tube, E	Π endotracheal tube	HHb	reduced hemoglobin

HME	heat and moisture exchanger
HMEF	heat and moisture exchanger with
	filter
hPa	hectopascal
HR	heart rate
HRdiff	heart rate difference
ht	height
HW	hardware
Hz	hertz
	liona
150	
IEC	International Electrotechnical
	Comission
I:E	inspiratory-expiratory ratio
IABP	intra-aortic balloon pump
IC	inspiratory capacity
ICP	intracranial pressure
ICU	intensive care unit
ID	identification
Imped.	impedance; impedance
•	respiration
in	inch
Inf	inferior
Infl.	inflation (limit)
Insp	inspiratory
Inv.	invasive
Inv. BP	invasive blood pressure
Irreg.	irregular
lso	isoflurane
ISO	International Standards
	Organisation
11/10	i di a constati a col a contra sta con

- idioventricular rhythm IVR joule
- J

K kelvin kcal kilocalorie kJ kilojoule kPa kilopascal	
L leg (describing location)	
L left (describing location)	
L, I liter	
l/min liters/minute	
Lab laboratory	
LAN local area network	
LAP left atrial pressure	
Lat lateral	
lb pound	
LCD liquid crystal display	
LCW left cardiac work	
LED light emitting diode	
LVEDP left ventricular end diastolic pressure)
LVEDV left ventricular end diastolic volume	2
LVSW left ventricular stroke work	
LVSWI left ventricular stroke work i	ndex

MAC	minimum alveolar concentration
Max	maximum
mbar	millibar
mcg	microgram
Mean	mean blood pressure
mEq	milliequivalent
MetHb	methemoglobin
MF	median frequency
mg	milligram
min	minute
Min	minimum
ml	milliliter
MLAEP	middle-latency auditory evoked
	potential
mmHg	millimeters of mercury
mol	mole
Monit	monitoring (ECG filter)
MRI	magnetic resonance imaging
Mult.	multiple
Multif. PVC	S
	multifocal PVCs
MV	minute volume
MVexp	expired minute volume (l/min)
MVexp(BTF	PS) expired minute volume in BTPS
	conditions
MVexp(STF	PD) expired minute volume in STPD conditions
MVinsp	inspired minute volume (I/min)
MVspont	spontaneous minute volume
Муо	myocardiac temperature
myo	

N2 nitrogen gastrointestinal carbon dioxide Pbaro barometric pressure N2O nitrous oxide and arterial blood carbon dioxide PCWP pulmonary capillary wedge Na sodium concentration pressure Naso nasopharyngeal temperature P(g-ET)CO2 difference between PE polyethylene neo neonate gastrointestinal carbon dioxide pedi pediatric Net network and end tidal carbon dioxide PEP positive end-expiratory press	
N20nitrous oxideand arterial blood carbon dioxidePCWPpulmonary capillary wedgeNasodiumconcentrationpressureNasonasopharyngeal temperatureP(g-ET)CO2difference betweenPEpolyethyleneneoneonategastrointestinal carbon dioxidepedipediatric	
Nasonasopharyngeal temperatureP(g-ET)CO2difference betweenPEpolyethyleneneoneonategastrointestinal carbon dioxidepedipediatric	
neo neonate gastrointestinal carbon dioxide pedi pediatric	
Net network and end tidal carbon dioxide PEEP positive end-expiratory press	
	ure
NIBP non-invasive blood pressure concentration PEEPe extrinsic positive end expirat	ory
NMT neuromuscular transmission P(STPD) pressure in STPD conditions pressure	
NO nitric oxide P16 invasive pressure channel PEEPe+i total positive end expiratory	
NTPD normal temperature and pressure, identification on module pressure (ICU)	
dry gas PA pulmonary artery PEEPe+PEEPi	
Num.numericalPaPascal (unit of pressure)total positive end expiratory	
Paced paced beats pressure (ICU)	
O ₂ oxygen PaCO ₂ partial pressure of carbon dioxide PEEPi intrinsic positive end expirate	ory
O ₂ ER oxygen extraction ratio in the arteries pressure	
O ₂ Hb oxygenated hemoglobin PAO ₂ partial pressure of oxygen in the PEEPtot total positive end expiratory	
OR operation room alveoli pressure (anesthesia)	
Oxy oxygenation PaO ₂ partial pressure of oxygen in the PgCO ₂ gastrointestinal carbon dioxi	de
Oxy Calcs oxygenation calculations arteries concentration	
PAOP pulmonary artery occlusion pH pH	
P partial pressure pressure pHa arterial pH	
P pressure PA pulmonary arterial pressure pHi intramucosal pH	
P(BTPS) pressure in BTPS conditions pHv (mixed) venous pH	

PIC	patient interface cable	R
Pleth	plethysmographic pulse waveform	RA
PM	pacemaker	Ra
PM non-	capt.	RC
	, pacemaker non-capturing	RC
PM non-		RE
	pacemaker non-functioning	Re
Pmax	maximum pressure	RE
Pmean	mean pressure	ref
Pmin	minimum pressure	Re
Ppeak	peak pressure	Re
Pplat	plateau (pause) pressure	RF
PR	pulse rate	RN
Prev.	previous	Ro
psi	pounds per square per inch	RÇ
pt	patient	RF
PTC	post tetanic count (NMT)	rtn
pts	patients	RV
PVC	polyvinylchloride	RV
PVC	premature ventricular contraction	
PVloop	pressure volume loop	RV
PvO ₂	partial pressure of oxygen in	
	(mixed) venous blood	RV
PVR	pulmonary vascular resistance	RV
PVRI	pulmonary vascular resistance index	RV
Px	standard pressure label, x being 1,	S
	2, 3, 4, 5, or 6	SA
	· · · ·	Sa
QRS	QRS complex	SD
Q̃s∕Qt	venous admixture	SE
		SE

R	right (describing location)
RAP	right atrial pressure
Raw	airway resistance
RCW	right cardiac work
RCWI	right cardiac work index
RE	Response Entropy
Rect	rectal temperature
REF	right ventricular ejection fraction
ref.	reference
Resp	respiration rate (total) (set)
Resp Rate	respiration rate (total) (measured)
RF	radio frequency
RMS	average (root mean square) power
Room	room temperature
RQ	respiratory quotient
RR	respiration rate (total) (measured)
rtm	rhythm
RV	residual volume
RVEDV	right ventricular end-diastolic
	volume
RVESV	right ventricular end-systolic
	volume
RVP	right ventricular pressure
RVSW	right ventricular stroke work
RVSWI	right ventricular stroke work index
S	second
SA	sinoatrial
SaO ₂	arterial oxygen saturation
SD	standard deviation
SE	State Entropy
SEF	spectral edge frequency
SEMG	spontaneous electromyogram

Sev SI Skin SN, S/N Spiro SpO ₂ Spont	sevoflurane stroke index skin temperature serial number patient spirometry oxygen saturation spontaneous breathing
SQI SR	signal quality index suppression ratio
SR	sinus rhythm
SSEP ST	somatosensory evoked potentials single twitch (NMT)
ST	ST segment of electrocardiograph
stat	static
STAT	continuous NIBP cuff inflation for five minutes
STBY	standby
Stfilt	ST filter (ECG)
STPD	standard temperature and pressure, dry gas
Surf	surface temperature
SV	stroke volume
SVC	supraventricular contraction
SVI	stroke volume index
SvO ₂	(mixed) venous oxygen saturation
SVR SVRI	systemic vascular resistance
SVRI SW	systemic vascular resistance index software
Sys	systolic pressure

t	time (min)	V	venous
Т	temperature	V	ventricular
T(BTPS)	temperature in BTPS conditions	V	volume
T1%	first stimulus as % of the reference	V/Q	ventilation/perfusion ratio
	value (NMT)	V0.5	volume expired during the first 0.5
T14	temperature channel		seconds
	identification on module	V1.0	volume expired during the first
Tab.	tabular		second
Tachy	tachycardia	VA	alveolar ventilation
Tbl, Tblood	blood temperature	VC	vital capacity
Tcorr	temperature correction	VCO ₂	carbon dioxide production
Temp	temperature	Vd	dead space
Theta, Th	theta frequency band	Vd/Vt	dead space ventilation
Tinj	injectate temperature	Vent Calcs	ventilation calculations
TOF	train of four (NMT)	VFib	ventricular fibrillation
TOF%	ratio of the 4th to the 1st response	VO ₂	oxygen consumption
	(NMT)	VO ₂ Calc	calculated oxygen consumption*
Trigem.	trigeminy	VO ₂ Calcl	calculated oxygen consumption
TV	tidal volume		index*
TVexp	expired tidal volume (ml)	VO ₂ I	oxygen consumption index
TVinsp	inspired tidal volume (ml)	Vol	volume
Tx	temperature label, x being 1, 2, 3,	V Run	ventricular run
	or 4 or one of the other label	V Tachy	ventricular tachycardia
	choices		
Tymn	tympanic temperature		

WEP	Wired Equivalent Privacy	
WLAN	wireless local area network	
wt	weight	
Х	extreme	
yr	year	
yrs	years	
* with Fick equation		

Tymp tympanic temperature

Supplies and accessories

The accessories below are approved and specified for the Datex-Ohmeda AS/3 Compact Anesthesia Monitor. For more information, see the corresponding Datex-Ohmeda catalogs. Patient accessories designed for use with this device are made of biocompatible materials conforming to requirements of the standard EN 30993 Biological Evaluation of Medical Devices and therefore do not contain toxic ingredients or primary skin irritants. The conformity is based either on laboratory testing or material knowledge and the long history of the materials used. Please note that some products are not available worldwide. You can check the availability with your local Datex-Ohmeda office or distributor. Please, also refer to our catalog: http://supplies.datex-ohmeda.com/DO

ECG

Trunk cables, IEC color coding

545300	3 leadwire trunk cable, 3 m/10 ft		
545301	5 leadwire trunk cable, 3 m/10 ft		
545200	Multiparameter cable		
	(3- and 5-wire sets)		
545323	10 leadwire trunk cable, 3 m/10 ft		
FOR M-PRE	FOR M-PRESTN, M-PRETN and M-RESTN only:		
8003600	3 leadwire ECG trunk cable,		
	3 m/10 ft		
8003602	5/10 leadwire ECG trunk cable,		
	3 m/10 ft		
8003606	3 leadwire ECG DIN trunk cable,		
	3 m/10 ft		
8003610	5 leadwire ECG DIN trunk cable,		
	3 m/10 ft		

Leadwire sets, IEC color coding

545315	3 leadwire set, clip, 0.75 m/2.5 ft		
545316	5 leadwire set, clip, 1.25 m/4.1 ft		
545325	C2-C6 leadwire set, clip,		
	1.25 m/4.1 ft		
8001960	3 leadwire set, clip, 1.5 m/4.9 ft		
8001961	5 leadwire set, clip, 1.5 m/4.9 ft		
FOR M-PRE	FOR M-PRESTN, M-PRETN and M-RESTN only:		
8003610	3 leadwire set, clip, 0.75 m/2.5 ft		
8003613	3 leadwire set, clip, 1.5 m/4.9 ft		
8003620	5 leadwire set, clip, 0.75 m/2.5 ft or		
	1.25 m/4.1 ft		
8003623	10 leadwire set, clip, 1.5 m/4.9 ft		
8003630	10 leadwire set, clip, 0.75 m/2.5 ft or		
	1.25 m/4.1 ft		

One-piece ECG cables, IEC FOR M-PRESTN, M-PRETN and M-RESTN only:

adwire one-piece ECG cable, clip,
m/11.5 ft
adwire one-piece ECG cable, clip,
m/11.5 ft

Telemetry ECG leadwires, IEC

8003100	5 leadwire, snap
8003102	10 leadwire, snap

Trunk cables, AAMI color coding

545302	3 leadwire trunk cable, 3 m/10 ft	
545303	5 leadwire trunk cable, 3 m/10 ft	
545201	Multiparameter cable	
	(3 and 5 leadwire sets)	
545324	10 leadwire trunk cable, 3 m/10 ft	
FOR M-PRESTN, M-PRETN and M-RESTN only:		
8003601	3 leadwire ECG trunk cable,	
	3 m/10 ft	
8003603	5/10 leadwire ECG trunk cable,	
	3 m/10 ft	
8003606	3 leadwire ECG DIN trunk cable,	
	3 m/10 ft	
8003610	5 leadwire ECG DIN trunk cable,	
	3 m/10 ft	

Leadwire sets, AAMI color coding

545317	3 leadwire set, clip, 0.75 m/2.5 ft	
545318	5 leadwire set, clip, 1.25 m/4.1 ft	
545327	3 leadwire set, snap, 0.75 m/2.5 ft	
545328	5 leadwire set, snap, 1.25 m/4.1 ft	
545326	V2-V6 leadwire set, clip,	
	1.25 m/4.1 ft	
8001958	3 leadwire set, clip, 1.5 m/4.9 ft	
8001959	5 leadwire set, clip, 1.5 m/4.9 ft	
FOR M-PRESTN, M-PRETN and M-RESTN only:		
8003611	3 leadwire set, clip,	
	0.75 m/2.5 ft	
8003612	3 leadwire set, snap,	
	0.75 m/2.5 ft	
8003614	3 leadwire set, clip,	
	1.5 m/4.9 ft	
8003621	5 leadwire set, clip, 0.75 m/2.5 ft or	
	1.25 m/4.1 ft	
8003622	5 leadwire set, snap, 0.75 m/2.5 ft or	
	1.25 m/4.1 ft	
8003624	5 leadwire set, clip,	
	1.5 m/4.9 ft	
8003631	10 leadwire set, clip, 0.75 m/	
	2.5 ft or 1.25 m/4.1 ft	

One-piece ECG cables, AAMI FOR M-PRESTN, M-PRETN and M-RESTN only:

8003635	3 leadwire one-piece ECG cable, clip,
	3.5 m/11.5 ft
8003637	5 leadwire one-piece ECG cable, clip,
	3.5 m/11.5 ft

Telemetry ECG leadwires, AAMI

8003101	5 leadwire, snap
8003103	10 leadwire, snap

Electrodes

572683	Solid gel, Ag/AgCl, pkg of 50 pcs
572684	For infants, safety pin 60 cm,
	15 pcs

Pulse oximetry

OxyTip+ Reusable Finger Sensors

OXY-F4-NIntegrated finger sensor, 4 m/13 ftOXY-F-UN*Interconnect finger sensor, 1 m/3.3 ftOXY-F-DBInterconnect sensor, 2 m/7 ft

OxyTip+ Adhesive Sensors

OXY-AP-25* Adult and pediatric, pkg of 25 pcs OXY-AP-10* Adult and pediatric, pkg of 10 pcs OXY-AF-10* AllFit, pkg of 10 pcs OXY-DSP* Adhesive sensor sample kit *Requires the use of an OxyTip+ Interconnect Cable (OXY-OL3)

OxyTip+ Cables

OXY-OL3Interconnect cable, 3 m/10 ftOXY-SL3Interconnect cable, 3 m/10 ftOXY-SL4Adapter cable, 0.5 m/1.5 ftOXY-SLCAdapter cable, 2 m/7 ftOXY-C1Interconnect cable, 1.5 m/4.9 ftOXY-C3Interconnect cable, 3 m/10 ftOXY-C7Interconnect cable, 7 m/23 ft

Temperature

Reusable probes

16560	Skin temp probe, 3.5 m/11.5 ft
165602	Skin temp probe, 1.5 m/4.9ft
16561	Central temp probe, adult,
	2.8 m/9 ft
165622	Central temp probe, adult,
	1.5 m/4.9 ft
165611	Central temp probe, pedi,
	2.8 m/9 ft
165612	Central temp probe, pedi,
	1.5 m/4.9 ft

Disposable probes

8001642Skin temperature probe8001643Central temperature probe 12F8001644Central temperature probe 9F

Extension cables for disposable probes

165640	Extension cable 1.3 m/4.3 ft, used
	with multiparameter cables
165641	Extension cable 2.8 m/9.2 ft, used
	with Datex-Ohmeda monitor or module

Multiparameter cables (ECG, Sp02, Temp)

545200	Multiparameter cable, IEC
545201	Multiparameter cable, AAMI

Esophageal stethoscopes

8002910	Esophageal stethoscope with temperature probe, 9F
8002911	Esophageal stethoscope with temperature probe, 12F
8002908	Esophageal stethoscope with temperature probe, 18F
8002909	Esophageal stethoscope with temperature probe, 24F

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Invasive blood pressure Reusable transducers and cables

78000	SensoNor 844, 3 m/10 ft
165700	Spectramed P10EZ-1,
	0.45 m/1.5 ft
54586	Adapter cable for DTX disposable
	pressure transducers, 3.8 m/12 ft
875408	Cable for HP 1290C-type pressure
	transducers, 0.3 m/1 ft

Disposable flushing kits and domes

16577	Flushing kit for SensoNor 840, sterile,
	pkg of 10 kits
16578	Dome for SensoNor 840, sterile, pkg
	of 50 pcs
78001	Flushing kit for SensoNor 844
78002	Dome for SensoNor 844

NIBP

Reusable color coded latex-free cuffs

572429	Large adult cuff, red
572428	Standard adult cuff, blue
572427	Small adult cuff, gray
572426	Child cuff, green
572425	Infant cuff, tan
8002248	Long large adult cuff, red
8001997	Thigh NIBP cuff (for M-PRESTN,
	M-PRETN and M-RESTN only)

Disposable cuffs

8001991	Long large adult cuff
8001992	Large adult cuff
8002562	Long standard adult cuff
8001993	Standard adult cuff
8001994	Small adult cuff
8001995	Child cuff
572403	Infant cuff #3, pkg of 10 pcs
572404	Infant cuff #4, pkg of 10 pcs
572405	Infant cuff #5, pkg of 10 pcs
572404	Infant cuff #4, pkg of 10 pcs

Cuff hoses

Adult hose, black, 3 m/10 ft
Adult hose, black, 1.8 m/6 ft
Adult hose, black, 6 m/20 ft
Infant hose, white, 3 m/10 ft
Infant hose, white, 6 m/20 ft

Cardiac output

16573	Thermodil. cath. Edwards Lifesciences
	Corp. 93A-131-7F
16590	Catheter connecting cable
16591	Injectate bath temp probe
16592	Flow-through injectate temp probe
16593	Spectramed CO-set temp probe
16574	CO-set for room temp inject., 10 pcs

Patient Spirometry

Reusable sensors

733910	D-lite sensor
73393	Pedi-lite sensor

Single use sensors

733950	D-lite sensor, pkg of 50 pcs
896952	D-lite+ sensor
8002718	Pedi-lite+ sensor, pkg of 50 pcs

Disposable spirometry tubes

890031	2 m/7 ft, yellow, pkg of 5 pcs
884101	3 m/10 ft, yellow, pkg of 5 pcs

Disposable spirometry accessory kit

889560	pkg of 50 kits
8002718	pediatric spirometry kit, pkg of 50 pcs

Airway gases

Anesthesia gas sampling lines

73318Disposable, 2 m/7 ft, pkg of 10 pcs73319Disposable, 3 m/10 ft, pkg of 10 pcs

Disposable airway adapters

73385Straight T-adapter, pkg of 10 pcs73386Elbow adapter, pkg of 10 pcsThe following adapters are for low dead spacepediatric endotracheal tubes:

877583	ID 2.5 mm, pkg of 5 pcs
877584	ID 3.0 mm, pkg of 5 pcs
877585	ID 3.5 mm, pkg of 5 pcs
877586	ID 4.0 mm, pkg of 5 pcs

Reusable airway adapters

84995 Steel adapter, 15F-15M

D-fend water traps

D-fend, black, pkg of 10 pcs
D-fend+, green, pkg of 10 pcs
Mini D-fend, pkg of 10 pcs,
for M-miniC
Container, pkg of 5 pcs

Filtration

Machine side filter

557021200 Uni-Filter, pkg of 45 pcs Patient machine side filter 557022500 Uni-Filter/S, pkg of 60 pcs

Humidification/Filtration Heat and moisture exchangers with integrated bacterial/viral filters (HMEF) 557070100 HMEF 1000, pkg of 50 pcs 557070500 HMEF 500, pkg of 75 pcs

Dust filters

886236 For all M-Cxx modules

Calibration gases

755534	Regulator for calibration gases
	755580, 755581, 755583
755580	Quick Cal calibration gas for
	M-miniC, applicable for M-TONO
755581	Quick Cal calibration gas for M-CO,
	M-COV, M-COVX
755583	Quick Cal calibration gas for
	M-CAIO, M-CAIOV, M-CAIOVX

Tonometry

Catheters

TONO-8F	pkg of 5 pcs	
TONO-14F	pkg of 5 pcs	
TONO-16F	pkg of 5 pcs	
TONO-18F	pkg of 5 pcs	
Calibration gas for tonometry		
755580	Ouick Cal calibration gas	

755580	Quick Cal calibration gas
	(balance air)
755534	Regulator
733251	Calibration gas sampling line

NMT M-NMT sensors

888418 MechanoSensor, 0.3 m/1 ft 888416 ElectroSensor, 0.3 m/1 ft 888419 Pediatric MechanoSensor, 0.3 m/1 ft

M-NMT sensor cables

888414 3.3 m/11 ft 888415 1.5 m/5 ft

Miscellaneous for Datex-Ohmeda M-NMT

57268	NMT electrodes, solid gel, Ag/AgCl, pkg of 30 pcs
888417	Regional Block Adapter, cable 0.5 m/1.5 ft

EEG and AEP

Lead sets

898050	Basic EEG lead set, clip
898051	General EEG lead set, clip
898052	AEP lead set, clip

Electrodes

572685	EEG stick-on electrodes, 15 bags
	containing 5 electrodes each
572686	EEG cup electrodes, pkg of
	500 pcs
75349	Conductive paste

BIS

900507	Converter set: DSC and PIC Plus cable
545780	PIC Plus cable
545781	BIS Sensor Plus
545782	BIS Sensor Pediatric
545783	BIS Sensor Quatro

Entropy

8002858 Entropy sensor (pkg of 25 pcs) 8002964 Entropy sensor cable, 3.5 m/11.5 ft

Interface cables

881167	UPI-PC serial, 3 m/ 10 ft
889352	M-PT Universal ECG/P3 output, 6m/20
	ft
884988	M-PT Kontron IABP cable, 6 m/20 ft
884989	M-PT Datascope IABP cable 6 m/20 f
892385	INT Baxter Vigilance cable, 2 m/7 ft

Mounting elements

572238	Portable monitor wall mount
887053	CM universal mounting plate
886172	CM bed mount
572235	Portable monitor roll stand
800065	Remote Controller holder

Other monitor supplies

74205 85969	Thermal recorder paper, 20 rolls Cleaning fluid
	5
883387	Dust filter for Compact Monitor
886236	Dust filter for Compact Airway
	Modules

Fuses

51119	250 V, T3.15 A, 5*20 mm
511382	125 V,5 A slow,5*20 mm (UL/CSA)