ePM 15/ePM 15A/ePM 15C ePM 12/ePM 12A/ePM 12C ePM 10/ePM 10A/ePM 10C

Patient Monitor

Service Manual

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Preface

Manual Purpose

This manual provides detailed information about the assembling, dissembling, testing and troubleshooting of the equipment to support effective troubleshooting and repair. It is not intended to be a comprehensive, in-depth explanation of the product architecture or technical implementation. Observance of the manual is a prerequisite for proper equipment maintenance and prevents equipment damage and personnel injury.

Intended Audience

This manual is for biomedical engineers, authorized technicians or service representatives responsible for troubleshooting, repairing and maintaining the monitors.

Passwords

A password may be required to access different modes. The passwords are listed below:

- User maintenance: 888888
- Manage Configuration: 315666
- Factory maintenance: 332888
- Demo mode: 2088

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FOR YOUR NOTES

1.1 Safety Information

\mathbb{N} warning

 Indicates a potential hazard or unsafe practice that, if not avoided, could result in death or serious injury.

 Indicates a potential hazard or unsafe practice that, if not avoided, could result in minor personal injury or product/property damage.

NOTE

• Provides application tips or other useful information to ensure that you get the most from your product.

1.1.1 Warnings

/ WARNING

- This equipment is used for single patient at a time.
- This equipment and its accessories are suitable for use within the patient environment.
- To avoid explosion hazard, do not use the equipment in the presence of oxygen-rich atmospheres, flammable anesthetics, or other flammable agents.
- Use and store the equipment in specified environmental condition. The monitor and accessories may not meet the performance specification due to aging, stored or used outside the specified temperature and humidity range.
- The equipment is not intended to be used within the Magnetic Resonance (MR) environment.
- Before connecting the equipment to the power line, check that the voltage and frequency ratings of the power line are the same as those indicated on the equipment's label or in this manual.
- Before putting the system into operation, the operator must verify that the equipment, connecting cables and accessories are in correct working order and operating condition.

- To avoid risk of electric shock, the equipment must only be connected to mains power with protective earth. If a protective earth conductor is not provided, operate it on battery power, if possible.
- Do not use the multiple portable socket outlets (MPSO) or AC mains extension cords. Insure that the sum of the individual ground leakage currents does not exceed the allowable limits.
- Do not touch the patient and live parts simultaneously. Otherwise patient injury may result.
- Do not come into contact with the patient during defibrillation. Otherwise serious injury or death could result.
- Do not open the equipment housings. All servicing and future upgrades must be carried out by trained and authorized personnel.
- Do not rely exclusively on the audible alarm system for patient monitoring. Turning the alarm volume to a low level or off may result in a hazard to the patient. Remember that alarm settings should be customized according to patient situations. Always keep the patient under close surveillance.
- The physiological data and alarm messages displayed on the equipment are for reference only and cannot be directly used for diagnostic interpretation.
- Route, wrap and secure the cables to avoid inadvertent disconnection, stumbling and entanglement.
- The software equipment copyright is solely owned by Mindray. No organization or individual shall resort to modifying, copying, or exchanging it or to any other infringement on it in any form or by any means without due permission.

1.1.2 Cautions

- Use only parts and accessories specified in this manual.
- Ensure that the equipment is supplied with continuous electric power during work. Sudden power failure may cause data loss.
- Magnetic and electrical fields are capable of interfering with the proper performance of the equipment. For this reason make sure that all external devices operated in the vicinity of the equipment comply with the relevant EMC requirements. Mobile phone, X-ray equipment or MRI devices are a possible source of interference as they may emit higher levels of electromagnetic radiation.
- Always install or carry the equipment properly to avoid damage caused by drop, impact, strong vibration or other mechanical force.
- Dry the equipment immediately in case of rain or water spray.
- Some settings are password protected and can only be changed by authorized personnel. Contact your department manager or biomedical engineering department for the passwords used at your facility.

- Dispose of the package material as per the applicable waste control regulations. Keep it out of children's reach.
- At the end of its service life, the equipment, as well as its accessories, must be disposed of in compliance with the guidelines regulating the disposal of such products. If you have any questions concerning disposal of the equipment, please contact us.

1.1.3 Notes

NOTE

- Put the equipment in a location where you can easily view and operate the equipment.
- The equipment use a mains plug as isolation means to the mains power. Do not locate the equipment in a place difficult to operate the mains plug.
- The typical operator's position is in front of the monitor.
- The software was developed in compliance with IEC62304. The possibility of hazards arising from software errors is minimized.
- This manual describes all features and options. Your equipment may not have all of them.
- Keep this manual in the vicinity of the equipment so that it can be obtained conveniently when needed.

1.2 Equipment Symbols

See the **ePM Series Patient Monitor Operator's Manual (P/N: 046-012606-00)** for information about the symbols used on this product and its packaging.

FOR YOUR NOTES

2.1 Overview

The ePM series multi-parameter monitors enable to provide complete patient management, adequate physiological parameter monitoring and physiological alarm, possessing powerful data review function, flexible wired or wireless network configuration and application ability. It provides a series of CAA applications to assist physicians to make a diagnosis. Meanwhile, the ePM series provide hospital management with superior monitor management applications to help hospitals to improve the efficiency and quality of monitor equipment management.

Based on the needs of clinical applications, ePM series could provide product models with displays of different sizes. Users can operate the monitor by touch screen or shortcut key. The ePM series have good human-computer interaction design, clinical applicability, complete hospital IT solution capabilities and lots of CDS applications.

2.2 Product System Structure

The ePM series monitors have only one main unit:

- ePM 10/ePM 10A/ePM 10C host uses 10.1" TFT WXGA display
- ePM 12/ePM 12A/ePM 12C host uses 12.1" TFT WXGA display
- ePM 15/ePM 15A/ePM 15C host uses 15.6" TFT WXGA display
- Both of them use a touch screen and shortcut buttons as input devices, with an optional remote control
- Optional WiFi module
- Optional built-in recorder



Fig. 1-1 System Block Diagram

2.2.1 Main Control Board/Parameter/Interface Board

The main control board includes the main control CPU, program memory, data memory, system configuration memory, WiFi module (optional), power management MCU, charging circuit for batteries and DC-DC circuit. A multi-parameter module circuit (ECG/Resp/SpO₂/NIBP/IBP) is also integrated on this board. In addition, there are internal and external interfaces. The internal interfaces include an interface for the recorder, an internal parameter module interface, an interface between the AC-DC and the battery. The external interfaces include a VGA display interface, a USB interface, an Ethernet interface and a multi-function interface. (There is an additional DC_IN interface for ePM 10/ePM 10A/ePM 10C).



Fig. 1-2 Main Control Board Block Diagram





Fig. 1-3 Power Supply Architecture Diagram

The AC/DC power module outputs 15V to the main control board. By the internal DC-DC_2 and DC_DC_1 conversion circuits of the main control board, it can generate 3.3V, 5V and 12V to supply power for other modules or boards in the host. The battery charging circuit is powered by 15V and can be switched between AC power supply and battery power supply by AC connection test. The ePM12/ePM 12A/ePM 12C can support dual batteries. The first battery is installed in the main unit and the second battery is installed in the external battery compartment. The external battery compartment is installed at the bottom of the main unit. For the ePM15/ePM 15A/ePM 15C, the internal battery compartment can support two batteries. The ePM Series monitors support 3 types of batteries, 2600mAh, 4500mAh and 5600mAh. 2600mAh and 4500mAh are interchangeable. 5600mAh cannot be used interchangeably with these two batteries. The ePM 10/ePM 10A/ePM 10C can be mounted to an ambulance via a charging dock. There is an isolated power module in the charging dock which converts the 28~12V input to a 15V output. The output of the charging dock is connected to the DC_in interface of the ePM 10/ePM 10A/ePM 10C. +12V is the power supply for the integral module rack, recorder and parameter acquisition circuit. The power module for parameter acquisition adopts the DC-DC isolation design. DC_DC_3 is used to power the main processor.

2.2.3 Alarm Indicator Light Board

There are LED alarm indicator lights and light sensors (optional) on the board. The light sensors conduct ambient light detection in order to adjust the brightness of LCD background light.

2.2.4 Power On/Off Button Board/Shortcut Button Board

For the power switches, indicator lights and shortcut buttons are integrated on one board.



2.3 Data Logic Flow

Fig.2- 3 Data Flow Diagram

The monitoring parameters are collected and analyzed by the module and then forwarded to the system software via an integral or external module rack. The system software displays waveforms, values and alarms, meanwhile the data, alarms and values will also be stored in the internal data memory. It can also be sent to a central station or other monitors via network (wired or wireless).

2.4 Power On/Off Signal Flow Diagram



Power ON/OFF chart



Power OFF process.

3.1 Introduction

This section describes how to install the Wireless LAN (WLAN) for Mindray patient monitor.

3.2 Network Deployment Process

If the hospital has established a WLAN, follow the installation process below:



Fig. 3-1 Network deployment flow chart

3.2.1 Output List

Operation	Output	Acceptance Criteria	Template
A0	Wireless network requirements	Determine the wireless network	Wireless network
	for Mindray patient monitor	deployment requirements for the	requirement table
		Mindray patient monitor.	
A3	Network inspection and	Confirm whether the customer	Wireless network
	acceptance report	network meets the requirements of	inspection and
		the Mindray patient monitor by	acceptance table
		questionnaires and measurements.	
A5	Installation confirmation	Confirm the actual operation of the	Installation
	report	Mindray patient monitor after	confirmation table for
		installation.	patient monitor

If the hospital plans to create a new WLAN for the Mindray patient monitor, make sure there is at least one WiFi channel that is not in use. Otherwise, after establishing a new WLAN, it is impossible to meet the requirements of the Mindray patient monitor in terms of co-channel interference. See the installation process below:



Fig. 3-2 New WLAN installation process

Operation	Output	Acceptance Criteria	Template
A0	Wireless network	Determine the wireless network	Wireless network
	requirements for	deployment requirements for the	requirement table
	Mindray patient	Mindray patient monitor.	
	monitor		
A2	Network design	/	/
	document, list of		
	materials		
A5	Installation	Confirm the actual operation of the	Installation confirmation
	confirmation report	Mindray patient monitor after installation.	table for patient monitor

Precautions

• Network design and deployment engineering are complex and need a professional IT engineer to help get the job done. This document does not contain these contents.

3.3 Network Requirements

The wireless network needs to meet the following requirements.

No.	ltem	Specific requirements		
Wireless	Wireless coverage requirements			
1	WiFi coverage signal	-65dBm		
	strength (RSSI)	RSSI is the value displayed on the patient monitor		
2	Co-channel interference	20dB (co-channel interference AP signal is at least 20dB lower		
		than the AP signal used by the monitor)		
3	Ping	The average latency of a PC or mobile phone is less than 100		
	latency	milliseconds and the packet loss rate should be lower than 1%.		
AP capab	AP capability requirements			
1	AP capability	The expected number of devices connected to an AP must be		
		less than 50% of the AP capacity. For example, within the		
		coverage of an AP, usually there are 16 devices connected to the		
		AP, so the entitled number of devices that are allowed to be		
		connected to the AP at the same time must be greater than 32.		
		An AP can create multiple SSIDs.		
2	Equipment density	The maximum number of devices that can be connected to an		
		AP at the same time is 16.		
		(including patient monitors and other equipment).		
WLAN fea	atures			

Table 3-1 Wireless network requirement table

No.	ltem	Specific requirements
1	AP channel width	Set the AP channel width to 20MHz. Do not use the HT40 or
		НТ80.
2	802.11 protocol	The WLAN cannot use protocols that are not supported by the
		Mindray patient monitor, such as 802.11ac
3	Security mode	The WLAN cannot use the security mode that is not supported by
		the Mindray patient monitor.
		WPA2-PSK is highly recommended. WPA2-Enterprise may
		increase the off-line probability while roaming, so it is not
		recommended.
4	Virtual local area	The patient monitor requires a special VLAN.
	network for special use	The use of VLAN can minimize the broadcast or multicast data
	(VLAN)	that may affect the stability of patient monitor.
Key settin	ngs	
1	DHCP	The DHCP server needs to keep a sufficient number of IP
		addresses to ensure that the patient monitor can obtain an IP
		address.
2	IGMP snooping	Enable IGMP snooping if the patient monitor adopts multicast
3	Multicast	If the patient monitor adopts multicast, the network multicast
		function should be enabled.
4	Beacon and DTIM	AP DTIM = 1, Beacon = 100 milliseconds
5	Service port	See the Mindray Patient Monitor Network White Paper; require
		network devices to turn on certain TCP/UDP ports for patient
		monitors

3.4 Network Inspection and Acceptance

3.4.1 Tools and Resources

- A laptop with Windows 7 (or a later version) and a wireless network card installed. It is recommended that the laptop be equipped with an Intel Centrino wireless adapter. If your laptop is configured with a different wireless adapter, make sure it has high precision.
- In terms of wireless network survey tools, it is recommended to use professional survey tools such as Tamograph, Wirelessmon or other professional network survey tools.
- Professional network engineer.

Precautions

• Those who implement WiFi network surveys should have received good training on WiFi. If you do not have a professional network engineer, please ask a third party for help.

3.4.2 WiFi Signal Calibration

Before testing network coverage by the use of a wireless network survey tool that runs on a laptop, follow the steps below to use the patient monitor to calibrate the RSSI of the wireless network survey tool.

- 1. Keep the patient monitor close to the wireless network survey tool. The distance between the patient monitor and the wireless network survey tool should not exceed 30cm and the distance from a human body should exceed 50cm.
- 2. Simultaneously move the patient monitor and wireless network survey tool (keep the same distance as before).
- 3. When the patient monitor displays the following RSSI values: -50dBm, -60dBm, -70dBm, and -80dBm, record the RSSI value read by the wireless network survey tool.
- 4. When conducting a site survey, calibrate the RSSI of the wireless network survey tool relative to the patient monitor (the RSSI of the patient monitor is the judge criterion for wireless coverage).

3.4.3 Network Inspection and Acceptance Process

This is done in two ways: First, complete the project that requires the hospital IT department to conduct self-test, as shown in the network inspection and acceptance table. Then, the customer service personnel or authorized party will conduct site test and confirm the remaining contents, and finally fill in the network inspection and acceptance table. If any items are found to be non-conformed during the network inspection and acceptance, adjust them before installing the patient monitor.

During the testing, need to enable the WiFi network SSID broadcast to ensure that the SSID of the WiFi can be scanned.

No.	ltem	Specific requirements	Inspection and acceptance	Inspection results
			method	
Wirele	ess coverage requ	irements		
1	Received	≥ -65dBm	Service personnel use the	
	signal	RSSI is the value	network survey tool to	
	strength(RSSI)	displayed on the patient	perform tests.	
		monitor	Be sure to test all expected	
			coverage areas (such as	
			wards, corridors, toilets, stairs	
			and elevators).	
2	Co-channel	-20dB	Service personnel use the	
	interference		network survey tool to	
			perform tests.	
			Be sure to test all expected	
			coverage areas (such as	
			wards, corridors, toilets, stairs	
			and elevators).	

Table 3-2 Wireless network inspection and acceptance table

No.	ltem	Specific requirements	Inspection and acceptance method	Inspection results
3	Ping latency	The average latency of a PC or mobile phone using a normal WiFi module is less than 100 milliseconds and the packet loss rate should be lower than 1%.	Steps for the service personnel to perform the test: 1. Connect your PC or mobile phone to the AP. 2. Connect another PC to the LAN port to which the central monitoring system is connected. 3. Run the "ping –t –l 32 –w 1000 IPaddress-of -cellphone" command for 10 minutes. 4. Run "ctrl+c".	
AP cap	bacity requirement	nts		
1	AP capability	The expected number of devices connected to an AP must be less than 50% of the AP capacity. For example, within the coverage of an AP, usually there are 16 devices connected to the AP, so the entitled number of devices that are allowed to be connected to the AP at the same time must be greater than 32. An AP can create multiple SSIDs.	Service personnel obtain the AP model number from hospital personnel or by direct observation. Obtain an AP data manual as per this model number to confirm related AP capabilities.	
2	Equipment density	The maximum number of devices that can be connected to an AP at the same time is 12 (including patient monitors and other devices).	Check together with hospital IT personnel to confirm whether this requirement is met.	
WLAN	features			
1	AP channel width	Set the channel width to 20MHz. Do not use the HT40 or HT80.	Check together with hospital IT personnel to confirm whether this requirement is met.	

No.	ltem	Specific requirements	Inspection and acceptance method	Inspection results
2	802.11	The WLAN cannot use	Check together with hospital	
	protocol	protocols that are not	IT personnel to confirm	
		' supported by the	whether this requirement is	
		Mindray patient	met.	
		monitor, such as		
		802.11ac		
3	Security mode	The WLAN cannot use	Check together with hospital	
		the security mode that	IT personnel to confirm	
		is not supported by the	whether this requirement is	
		Mindray patient	met.	
		monitor.		
		WPA2-PSK is highly		
		recommended.		
		WPA2-Enterprise may		
		increase the off-line		
		probability while		
		roaming, so it is not		
		recommended.		
4	Virtual local	The patient monitor	Check together with hospital	
	area network	requires a special VLAN.	IT personnel to confirm	
	for special use	The use of VLAN can	whether this requirement is	
	(VLAN)	minimize the broadcast	met.	
		or multicast data that		
		may affect the stability		
		of patient monitor.		
Key se	ttings			
1	DHCP	The DHCP server needs	Check together with hospital	
		to keep a sufficient	IT personnel to confirm	
		number of IP addresses	whether this requirement is	
		to ensure that the	met.	
		patient monitor can		
		obtain an IP address.		
2	IGMP snooping	Enable IGMP	Check together with hospital	
		snooping if the	IT personnel to confirm	
		patient monitor	whether this requirement is	
		adopts multicast	met.	
3	Multicast	If the patient monitor	Check together with hospital	
		adopts multicast, the	IT personnel to confirm	
		network multicast	whether this requirement is	
		function should be	met.	
		enabled.		
4	Beacon and	AP DTIM = 1, Beacon =	Check together with hospital	

No.	ltem	Specific requirements	Inspection and acceptance	Inspection results
			method	
	DTIM	100 milliseconds	IT personnel to confirm	
			whether this requirement is	
			met.	
5	Service port	See the Mindray Patient	Check together with hospital	
		Monitor Network White	IT personnel to confirm	
		Paper; need to keep	whether this requirement is	
		certain TCP/UDP ports	met.	
		open for patient		
		monitors		

3.5 Use Patient Monitors to Assess Network Coverage

To confirm the coverage, perform a coverage test in the areas where patients often go around.

Confirm whether the coverage meets the requirements by observing the signal strength (RSSI) displayed on the patient monitor and whether an off-line event occurs.

If necessary, adjust the AP position or increase APs to ensure adequate coverage.

Please follow the steps below:

- 1. Set the patient monitor to access the central monitoring system.
- 2. Perform a Ping command on the patient monitor via the central monitoring system (enter "ping t l 32 w 1500 IP address" in the CLI window) (continue to run the Ping command on the patient monitor. The data packet is 32 bytes and the reply timeout is 1500 milliseconds). Ten minutes later, enter "ctrl + c" (complete Ping) to ensure that the average latency is less than 250 milliseconds and the packet loss rate should be lower than 1%.
- 3. Hold the patient monitor by hand and avoid being held up by other people. Walk around the expected coverage areas (such as wards, toilets, smoking areas, corridors, and all corners of the elevator).
- 4. The number of disconnections from the central station should be less than 10% of the roaming times of the patient monitor, and the RSSI value displayed on the patient monitor should not be lower than -65dBm.
- 5. If the signal strength is below -65dBm while walking around, stop walking at that location and observe for 30 seconds. If the RSSI value is not lower than -65dBm for more than 66% of the time, then it meets the coverage requirements.

Table 3-3 Installation confirmation table for patient monitor

Testing or observing items	Results (pass, fail or not applicable)
Perform a Ping command on the patient monitor via the central	
monitoring system and ensure that the average latency is less than	
250 milliseconds and the packet loss rate should be lower than 1%.	
Hold the patient monitor by hand and walk around different AP	
coverage areas. After walking around all the expected coverage areas,	
observe the continuous waveforms on the central monitoring system.	
The off-line time should be less than 10% of the patient monitor's	
roaming time.	
In the position of worst coverage, the signal strength displayed on	
the screen is higher than -65dBm.	

Precautions

• If the monitor is assessed only for use in a fixed position instead of roaming among various APs, then it's unnecessary to walk around coverage area during testing. It is only necessary to place the monitor at the possible installation position of the worst signal, and then confirm the signal strength and Ping effect.

3.6 Recommended Network Equipment

The Cisco devices listed in the table below are recommended.

Device	Parts No.	
2500 wireless controller	AIR-CT2504-x-K9	
2600 wireless APs	AIR-CAP2602I-x-K9	

3.7 Seting the Wireless Parameters of the Patient Monitor

Configure the patient monitor's WLAN parameters as per the table below:

Parameters	Settings recommended	Remarks	
[Main Menu] → [Maintena	ance] \rightarrow [User Maintenance] -	→ [Network Setup] → [WLAN]	
SSID	Set the actual network name to be used	1	
Security mode	WPA2-PSK	The security mode should be the same as that of the WLAN deployed for the patient monitor. If you are using EAP, choose the security mode based on your WLAN deployment.	
Password	Set the actual network password to be used.	/	
[Main Menu] → [Maintenance] → [User Maintenance] → [Network Setup] → [WLAN]→ [WLAN Setup]			

Parameters	Settings recommended	Remarks	
WLAN frequency band	5G	The options include: 2.4G, 5G and Auto.	
		2.4G = use only 2.4GHz band	
		5G = use only 5GHz band	
		Auto = use 2.4GHz and 5G Hz bands (5GHz shall	
		prioritize)	
ID verification server	ACS	Options include: ACS and SBR.	
type		ACS refers to the Cisco access control server.	
		SBR refers to other servers except ACS.	
		It applies only when the security type is Enterprise.	
BG channel	Designate	The options include: All, Designate and None.	
		Stability and roaming performance can be improved	
		by limiting the number of channels that the monitor	
		can be connected to. For example, on a 2.4GHz	
		network, set the channels to 1, 6, and 11, then the	
		network card will not scan or connect to other	
		channels.	
		The BG channel settings on the patient monitor must	
		match the AP channel settings.	
Channel A	Designate	The options include: All, Designate and None.	
		Stability and roaming performance can be improved	
		by limiting the number of channels that the monitor	
		can be connected to.	
		The 5GHz channel settings on the patient monitor	
		must match the WLAN AP channel settings.	
[Main Menu] → [Maintena	ance] \rightarrow [User Maintenance] -	→ [Network Setup] → [WLAN] → [Certificate	
Management]			
Local	/	Display the existing EAP certificate in patient monitor	
USB driver	/	Display the existing EAP certificate in the USB driver	
[Main Menu] → [Maintena	ance] → [Factory Maintenance	e] → [Setup] → [WLAN Setup]	
Adjustment area	International area	South Korea, Turkey, Russia and Brazil need to be	
		configured separately. Other countries/regions only	
		need to choose international area.	
		The patient monitor needs to be restarted for the	
		patient monitor settings to take effect.	
CCX features	Support	This means it supports CCX 4.0 and fast roaming	
PMK cache	Criteria	The options include: Standard and OPMK.	
		The option of Standard refers to PMK cache.	
		OPMK refers to random key cache.	
Trigger	-70	When the RSSI is below the roam trigger value, the	
		network card will attempt to roam.	
Scan cycle	5	When the RSSI is below the roam trigger value, the	
		probe request cycle is 5 seconds.	

The security modes supported by the monitor include:

menu	Basic	Authentication	Encryption	Whether support CCKM
	algorithm	mode	mode	
WPA PSK	WPA	PSK	TKIP/RC4	No
WPA2 PSK	WPA2	PSK	CCMP/AES	No
WPA PSK AES	WPA	PSK	CCMP/AES	No
WPA TKIP	WPA	EAP	TKIP/RC4	No
WPA2 AES	WPA2	EAP	CCMP/AES	No
WPA AES	WPA	EAP	CCMP/AES	No
ССКМ ТКІР	ССКМ	EAP	TKIP/RC4	Yes
CCKM AES	ССКМ	EAP	CCMP/AES	Yes

After the EAP authentication mode is selected, the system will display corresponding configuration items. The table below lists the configuration items for different EAP methods.

	Identity	Anonymous	Password	CA	User	PAC	PAC
				certificate	certificate	certificate	Password
PEAP-MSCHAPV2	Y	0	Υ	Y	Ν	Ν	Ν
PEAP-GTC	Y	0	Υ	Y	Ν	Ν	Ν
PEAP-TLS	Y	0	Υ	Y	Y	Ν	Ν
TTLS	Y	0	Υ	Y	Ν	Ν	Ν
TLS	Y	Ν	Y	Y	Y	Ν	Ν
FAST	Y	0	Y	N	N	Y	Y
LEAP	Y	Ν	Υ	Ν	Ν	Ν	Ν

Remarks: Y means "yes", N means "no", and O means "optional".

The meaning of each configuration item is shown below:

- Identity Verification Protocol (Phase 2 Identity Verification): When PEAP in the EAP method is selected, the user can configure the following PEAP internal methods: EAP-MSCHAPV2, EAP-GTC and EAP-TLS.
- Identity: user identity, i.e. the user name in AD, LDAP, or local user management of the RADIUS server.
- Anonymous: This item does not affect the identity verification process. The function of this item is to hide the real name (identity).
- Password: the password for the identity.
- CA certificate: Select the CA certificate from the imported certificates.
- User certificate: Select the user certificate from the imported certificates.
- PAC certificate: When EAP-FAST is selected, also select the PAC certificate from the imported certificates. If the RADIUS server supports intra-band PAC deployment and deploy PAC for customers, there is no need to set a PAC certificate or password.
- PAC password: Enter the PAC password for the PAC certificate when EAP-FAST is selected. If the RADIUS server supports intra-band PAC deployment and deploy PAC for customers, there is no need to set a PAC certificate or password.

3.8 Troubleshooting

Sign	Possible causes	Recommended measures
The patient monitor	The nearby AP is not turned on.	Make sure the AP is turned on and it belongs to the
cannot be		VLAN for the patient monitor.
connected to the	The patient monitor is not turned	Go to the AP coverage area and turn on the patient
AP and an X is	on within the AP coverage area.	monitor. Ensure that the signal strength displayed
displayed on the		on the patient monitor is greater than –65dBm.
patient monitor's		Ensure the co-channel interference meets the
WiFi signal icon.		requirements.
	The SSID and IP address	Refer to this manual to reconfigure this
	acquisition mode and security	information.
	mode are not properly	
	configured on the patient	
	monitor.	
	Patient monitor fault.	Check if another patient monitor can get online. If
		possible, restart the patient monitor and make sure
		the two patient monitors have the same
		configuration. If the patient monitor is still unable
		to get online then return it to Mindray for repair.
The patient monitor	The patient monitor is not	Patient monitor is allowed to access the central
can access AP but is	licensed to access the central	monitoring system.
unable to be	monitoring system.	
connected to the	The patient monitor is unable to	Use other network devices to connect to the
central monitoring	obtain any IP addresses, and the	central monitoring system and check if the IP
system.	IP addresses in the IP address	address can be obtained.
	pool have all been taken.	If the problem still exists, contact IT department.
	A static IP address conflict has	Observe if a prompt indicating an IP address
	occurred.	conflict is displayed on the patient monitor.
		If this prompt is displayed, ensure that all network
		devices have unique IP addresses.
	Network link trouble.	Confirm if the PC or mobile phone can ping the
		central monitoring system after being connected
		to the AP.
		If the problem still exists, contact IT department.
	The service port required for the	Check if the service port required for the patient
	patient monitor is not enabled	monitor is enabled on the hospital network. If not,
	on the hospital network.	enable related services (such as some UDP ports
		and multicast).
		If the problem still exists, contact IT department.
Intermittent	Move the patient monitor to the	Check if the WiFi signal strength is greater than
disconnection	blind coverage area.	-65dBm in the location where disconnection
occurs to a single		occurs.

Sign	Possible causes	Recommended measures
patient monitor	Patient monitor fault.	Check if disconnection occurs easily to the patient
		monitor at the same location. If the problem
		cannot be resolved after restarting the patient
		monitor, return the patient monitor to Mindray for
		repair.
	A static IP address conflict has	Observe if a prompt indicating an IP address
	occurred.	conflict is displayed on the patient monitor.
		Check if an IP address has been assigned to
		multiple devices.
Intermittent	APs in some areas are destroyed.	Make sure the AP has been turned on and works
disconnection		properly.
occurs to multiple	There is strong interference in	Use the network survey tool to see if the
patient monitors	some areas.	interference is strong and remove significant
		sources of interference or adjust the WLAN
		deployment to meet Mindray requirements.
	Insufficient signal coverage in	Use the network survey tool to check signal
	some areas.	coverage. If the signal coverage is insufficient in a
		certain area, adjust the AP position or increase APs.
Intermittent	Improper wired network	Use a wired patient monitor to check wired
disconnection	configuration	network configuration Ensure that the WLAN
occurs to all patient		bandwidth configured on the switch is sufficient
monitors		with a 50% surplus
	Radio interference exists	Use the network survey tool to check if there is
		radio interference and remove obvious sources of
		interference or adjust the WLAN deployment to
		meet Mindray requirements.

FOR YOUR NOTES

4.1 Introduction

Service personnel need to inspect, maintain and test the monitor regularly to ensure that it can keep working stably for a long time. This chapter provides basic test methods for the monitor as well as recommended appropriate testing frequency and testing tools. Please maintain and test the monitor with proper testing tools and according to actual needs.

The testing and inspection methods provided in this chapter are mainly used to confirm if the performance of the monitor can meet the specifications. During the testing, if test results do not meet the requirements, it indicates that the monitor or a certain function module of the monitor goes wrong and needs to be repaired or replaced immediately. Any other questions, please contact our after-sales service department.

Attention

- All tests can only be performed by qualified professional service personnel.
- Be careful when setting up and changing the contents in the maintenance and configuration menus, otherwise a data loss may be caused.
- Before the testing, service personnel need to ensure proper test tools and connection lines are used. Service personnel should be able to use these test tools proficiently.

4.1.1 Test Device

See the following testing sections.

4.1.2 Test Report

After our service personnel have done the test, please record the details according to the maintenance and testing report at the last page of this chapter and return it to the after-sales service department of the company.

4.1.3 Preventive Maintenance

A list of items requiring preventive maintenance for this monitor is provided below. Regular maintenance is recommended to be conducted at least once every two years (once a year for the CO₂ module). (See the following sections for detailed testing procedures and contents) Visual inspection NIBP test CO₂ testing and calibration

Inspection/maintena	ance items	Recommended frequency
Preventive maintena	ance	
Visual inspection		Installation for the first time, or after each re-installation.
	Pressure Test	
	Leakage Test	1. When the user suspects that the measured value is not
NIBP test	Overpressure	accurate.
	protection circuit	2. After repairing or replacing the relevant module.
	test	3. At least once every 2 years for the NIBP module, and once
Sidestream and	Leakage Test	a year for the CO ₂ module.
microstream CO ₂	Performance test	
test	Module calibration	
Performance test		
FCG test	Performance test	
	Module calibration	
Resp performance tes	t	
Spo ₂ test		
	Pressure Test	1 When the user suspects that the measured value is not
NIBP test	Leakage Test	1. when the user suspects that the measured value is not
Temp test		2. After repairing or replacing the relevant module.
	Performance test	3. At least once every two years At least once a year for CO_2
IBP test	Pressure calibration	module.
C.O. test	•	1
Mainstream CO ₂ test		
Sidestream,	Leakage Test	
microstream CO ₂	Performance test	
test	Module calibration	
Nurse call test		When the user suspects that the nurse call or analog output
Analog output test		function is not normal.
Electric safety test		
	Housing leakage	
	current test	
	Earth leakage	1 After repairing or replacing the power module
Electric safety test	current	2 Or after the monitor falls off
Licethe safety test	Patient leakage	3 At least once every two years or as needed
	current	
	Patient auxiliary	
	current	
Other tests		
Power On test		1. Installation for the first time, or after each re-installation.
		2. After repairing or replacing the parts of the main unit.

4.1.4 Recommended Frequency
Recorder check		After repairing or replacing the recorder.	
Network printing check		1. Installation for the first time.	
		2. After repairing or replacing the printer.	
Battery check	1. Installation for the first time.		
	Function check	2. After replacing the battery.	
	Performance check	Every two months or when the battery running hours are	
		significantly shortened.	

4.2 Preventive Maintenance

4.2.1 Visual Inspection

Visual check mainly refers to a comprehensive visual check on the monitor appearance. If the monitor has no obvious physical damage, then it passes the visual check. The specific check items are as follows:

- Check if there's physical damage to the monitor housing, display and buttons.
- Whether the module suffers physical damage.
- Whether the power cord, bracket and module accessories suffer physical damage.
- Whether the external cable wears out, whether the connector pin is loose or twisted.
- Whether the monitor's peripheral interface is loose, whether the pins are twisted.
- Whether the safety label and nameplate are clearly legible.

4.2.2 NIBP Test

Pressure check

Test tools:

- T connector
- Airway tube
- Spherical air pump
- Rigid container: 500 ± 25 ml
- Standard pressure meter: has been calibrated with a precision of at least 1 mmHg

See the below for check steps:

1. Connect the monitor, standard pressure meter, spherical air pump and rigid container as shown below.



- 2. The standard pressure meter's reading should be zero before air inflation. If it is not zero, open the spherical air pump valve so that the airway leads to the atmosphere until the standard pressure meter reads zero, then close the valve.
- Select [Main Menu] → [Maintenance] → enter password → [Module] → [NIBP] → [NIBP Pressure Test].
- 4. Check the readings of the standard pressure meter and the monitor, both of which should show a pressure value of 0mmHg.
- 5. Inflate the rigid container with a spherical air pump until the internal pressure reaches 50mmHg, then stop inflating and wait for 10s in order that the measurement value keeps stable.
- 6. Check the readings of the standard pressure meter and the monitor. The difference between the two should be within 3mmHg. If the difference is greater than 3mmHg, please contact service personnel.
- 7. Inflate the rigid container with a spherical air pump until the internal pressure reaches 200mmHg, then stop inflating and wait for 10s in order that the measurement value keeps stable. Repeat step 6.

Precautions

- You can also use a blood pressure simulator instead of a spherical air pump and a standard pressure meter to form a test system.
- You can also replace the rigid container with cylinders and cuffs of a right size.

Leakage Test

Testing tools:

- Adult cuff
- Airway tube
- Cylinder

See the below for testing steps:

- 1. Set [Patient Category] to [Adult].
- 2. Connect well the cuff with the NIBP cuff connector of the monitor.
- 3. Wrap the cuff around a cylinder of a right size, as shown in the figure.



- Select [Main Menu] → [Maintenance] → enter password → [Module] → [NIBP] → [NIBP Leakage Test], the NIBP parameter area will display [Leakage Test...].
- 5. After approximately 20 seconds, the system will automatically deflate which indicates the leakage test is completed.
- 6. If there is no prompt message in the NIBP parameter area, it means there is no air leakage in the system. If [**NIBP Air Leakage**] is displayed, it indicates that there may be an air leakage. At this time, the operator should check whether there is loose connection, whether the cuff and the inflation tube are damaged or leaked. When the connection is confirmed to be proper and there's no leakage in the cuff or the airway tube, then conduct a leakage test once again.

You can also perform a leakage test manually:

- 1. Perform steps 1 to 4 in the Pressure Calibration section.
- 2. Inflate the metal container with a spherical air pump until the internal pressure reaches 250mmHg, then stop inflating and wait for 5s in order that the measurement value keeps stable.
- 3. Record the current pressure and start timing using a timer. Record the pressure value after 60s.
- 4. The displayed pressure value after 60s minus the pressure value displayed at the beginning of the leakage test should not exceed 6mmHg.

Overpressure protection circuit test

Testing tools:

- T connector
- Airway tube
- Spherical air pump
- Rigid container: 500 ± 25 ml
- Standard pressure meter: has been calibrated with a precision of at least 1 mmHg
- 1. Perform steps 1 through 4 in the NIBP Pressure Check section.
- Select [Main Menu] → [Maintenance] → enter password → [Factory Maintenance] → [NIBP] → [Test].
- 3. In [Overpressure Protection Circuit Test], select [Adult/Child] for [Patient Category], adjust the output pressure of the air pump to 320-330mmHg. After it keeps stable, select the [Test] button to

start **calibration**. When the **test** succeeds, the NIBP menu will display the prompt of [**Test** Successful]. If the pressure exceeds 320-330mmHg, [**Test** Failed] will be displayed.

4. In [Overpressure Protection Circuit Test], select [Neo] for [Patient Category] and adjust the air pump output pressure to 160-165mmHg. When the value keeps stable, select the [Test] button on the right side of the menu to start Test. When the test succeeds, the NIBP menu will display the prompt of [Test Successful]. If the pressure exceeds 160-165mmHg, [Test Failed] will be displayed.

4.2.3 Sidestream and Microstream CO₂ Testing and Calibration

Leakage Test

- 1. After the CO₂ preheating startup is completed, block the module or the water tank inlet hole completely by hand or other objects, the sidestream and microstream CO₂ modules will have different actions respectively:
 - Sidestream: Plug the sidestream CO₂ module into the module rack of the main unit. Wait for 1min., after the module completes preheating startup, block the module air intake with your fingers or other objects. The monitor will display the alarm message of [CO₂ Airway Occluded]. Keep blocking it for about 60s, check [Maintenance] → enter password → [module] → [CO₂] → [CO₂ Module Calibration], check that the current flow rate of the module is <10ml/min, and the alarm message continues, then it proves that the module has no air leakage. If the alarm message of [CO₂ Airway Occluded] disappears or the current flow rate is ≥10ml/min, it indicates that there is air leakage.
 - Microstream: Keep blocking the module air intake for 3s, the screen displays the alarm message of [CO₂ Purging]; continue to keep blocking it for about 30s, the alarm message of [CO₂ Airway Occluded] is displayed, then it proves that the module has no air leakage.

Precision test

- A steel cylinder containing $6 \pm 0.05\%$ CO₂ and a balance gas of N₂.
- Steel cylinder with O₂ gas (its concentration >40%) and a balance gas of N₂ (for sidestream CO₂ module with oxygen module)
- T connector
- Airway tube
- Flowmeter
- 1. Insert the module into the module rack.
- 2. After the CO₂ preheating startup is completed, check the airway and conduct leakage test to ensure that there is leakage.
- 3. Go to [Maintenance] \rightarrow [Module] \rightarrow [CO₂] \rightarrow [CO₂ Module Calibration].
- 4. Connect the test system as shown below.



- Open and adjust the pressure reducing valve switch until the flow rate indicated by the flowmeter is 10~50mL/min and it keeps stable.
- 6. Check and ensure that the real-time concentration of CO_2 displayed in the calibration menu is $6 \pm 0.2\%$ (45 ± 2 mmHg for microstream CO_2).

Module calibration

- A steel cylinder containing $6 \pm 0.05\%$ CO₂ and a balance gas of N₂
- T connector
- Airway tube
- Flowmeter
- 1. Ensure that the sidestream CO₂ module or the microstream CO₂ module has completed preheating or startup.
- 2. Check the airway and conduct leakage test to ensure there is no leakage.
- Open the [CO₂ Calibration] menu: select [Main Menu] → [Maintenance] → enter the user maintenance password → [Module] → [CO₂].
- 4. Select [**Zero**] in the [**CO**₂] menu.
- 5. After it is successfully set to zero, connect it as shown below.



- 6. Open and adjust the pressure reducing valve switch until the flow rate indicated by the flowmeter is 10~50mL/min and it keeps stable.
- 7. Enter 6% (CO₂ concentration value) in the [**CO**₂%] text box of the [**CO**₂ **Calibration**] menu.
- 8. The current measured CO₂ concentration will be displayed in the [**CO**₂ **Calibration**] menu. Wait until the measured CO₂ concentration keeps stable, select [**Calibration**] to calibrate the CO₂ module.

After the calibration is successful, the [**Calibration Completed**] message will be displayed on the [**CO**₂ **Calibration**] menu; if the calibration fails, the [**Calibration Failed**] message will be displayed. In this case, please check if the calibration operation is correct and re-calibrate. If multiple calibrations fail, please return to the factory for repair.

4.3 Power On Test

The power-on test is conducted to confirm whether the monitor can be turned on and work properly. If the monitor can be started as follows, then it passes the power-on test. Specific steps:

- 1. Connect the monitor to an AC power. The AC power indicator lights up and the battery indicator light is on.
- 2. Press the power switch to turn on the monitor, the system will give a "beep" sound (indicating that the alarm sound passes the self-test); the red, yellow and green alarm lights light up respectively, and finally go out (indicating that the alarm lights pass the self-test)).
- 3. The startup screen disappears and enters the main interface of the system, and the normal startup is completed.

4.4 Module Performance Test

4.4.1 ECG Test

ECG performance test

Test tools:

- Patient simulator Medsim300B
- 1. Connect the patient simulator to the ECG parameter module with the ECG lead wire.
- 2. Set the simulators as follows: ECG sinus rhythm, HR = 60 bpm, amplitude is 1 mV.
- 3. Check and ensure that the ECG waveform display is normal and with no noise, and the HR value displayed on the monitor should not exceed 60 ± 1 bpm.
- 4. Disconnect each lead in turn, and observe that the screen displays the corresponding lead-off information.
- 5. Set the output pace signal of the simulator, set the [**Pace**] to [**Yes**] for the monitor, and confirm that the screen displays the pacing sign.

ECG calibration

Test tools:

- Vernier caliper
- 1. Set the [Filter Mode] of ECG parameters to be [Diagnose].
- 2. Select [Main Menu] \rightarrow [Maintenance] \rightarrow enter password \rightarrow [Module].
- 3. Select [**Calibration**], square wave signal will appear on the screen, and the technical alarm area will display [**ECG is Calibrating**].
- 4. Compare the amplitude of the square wave with the scale, and the error range should be no more than 5%.
- 5. After completing the calibration, select [**Stop Calibration**].

When needed, you can also output the above wave and scale by the recorder and measure the accurate error.

4.4.2 Resp Test

- Patient simulator Medsim300B
- 1. Connect the simulator and monitor with a non anti-ESU cable and set the monitor's resp lead to II.

- 2. Simulator settings: set the resp lead to II; basic damping to 500 Ω ; variable damping to 1 Ω ; and resp rate to 20 rpm.
- 3. Check if the monitor displays the RESP waveform with no distortion and that the displayed Resp value must not exceed 20 ± 1 rpm.

4.4.3 SpO₂ Test

Test tools:

- None
- Connect the adult SpO₂ sensor to the monitor SpO₂ interface, set the [Patient Category] of the monitor to [Adult], and set [PR Source] to SpO₂.
- 2. Measure the SpO_2 of your finger (assuming you are in a healthy state).
- 3. Check that the monitor displays the pleth waveform and PR value of blood oxygen, and the displayed saturation value range of blood oxygen should be between 95%-100%.
- 4. Remove the SpO₂ sensor from your finger and confirm that an alarm is given to indicate the SpO₂ sensor is off.

Confirmation of measurement accuracy:

The accuracy of the MPM SpO₂ module has been confirmed in human experiments by comparison with the reference values of arterial blood sample measured by a CO-oximeter. The measurements of pulse oximeter are statistically consistent, and it is expected that only about two-thirds of the measurements will be within the specified accuracy compared to the measurements of CO-oximeter.

Precautions

• The simulator cannot be used to verify the accuracy of the oxygen saturation monitor or the SpO₂ sensor, and can only prove that the working state of the monitor is normal. The accuracy of the oxygen saturation monitor or SpO₂ sensor needs to be verified by clinical data.

4.4.4 NIBP Test

See 4.2.2 NIBP test.

4.4.5 Temp Test

- Resistance box (accuracy not less than 0.1Ω)
- 1. Use a wire to connect both ends of any individual temperature interface of the parameter module to both ends of the variable resistor box.
- 2. The resistance box is set to 1354.9 Ω (corresponding to a temperature value of 37°C),

3. Verify each temperature channel of the monitor, and ensure that the monitor's display value does not exceed $37 \pm 0.1^{\circ}$ C.

4.4.6 IBP Test

Performance test

Test tools:

- Patient simulator Medsim300B, MPS450 or other equivalent devices
- Adapter cable for IBP test (300B, P/N: 00-002199-00) (MPS450, P/N: 00-002198-00)
- 1. Connect the simulator to the monitor's IBP parameter module.
- 2. Set that the simulator outputs 0mmHg to each IBP channel.
- 3. Press the "Zero" button of IBP module to conduct zero calibration for the parameter module.
- 4. Set the static pressure of the monitor = 200mmHg.
- 5. The display value on the monitor should not exceed $200 \pm 2mmHg$
- 6. If the error exceeds ± 2mmHg, perform a pressure calibration for the IBP parameter module. If the IBP parameter module has been calibrated for pressure with a reusable IBP sensor connected, then connect the IBP sensor and check the pressure calibration results.
- 7. Set the simulator to output 120/80mmHg ART signal and 120/0mmHg LV signal to each IBP channel respectively, and check if the IBP waveform is displayed correctly.

Pressure calibration

Method 1:

Test tools:

- Patient simulator Medsim300B, MPS450 or other equivalent devices
- Adapter cable for IBP test (300B, P/N: 00-002199-00) (MPS450, P/N: 00-002198-00)
- 1. Connect the simulator to the monitor's IBP parameter module.
- 2. Set the simulator to zero pressure.
- 3. Press the "Zero" button of IBP module to conduct zero calibration for the parameter module.
- 4. Set the static pressure of the monitor = 200mmHg.
- Select [Main Menu] → [Maintenance] → enter the user maintenance password → [Module] →
 [IBP] → [Calibration].

In the [IBP] menu, set the calibration value of the target pressure to 200mmHg.

- 6. Select the [**Calibration**] button on the right side of the target channel and the monitor will begin calibrating.
- 7. When calibration completes, it will display [**Calibration Completed**]. If it cannot be calibrated, it will display corresponding prompt message.

Method 2:

Calibration tools:

- Pressure meter
- Spherical air pump
- Airway tube
- T connector
- 1. Connect the three-way switch to the sphygmomanometer and the spherical air pump using a T-connector, as shown below.
- 2. First perform zero calibration. After the calibration is completed, the three-way switch leads to the sphygmomanometer.



Sphygmomanometer

- Select [Main Menu] → [Maintenance] → enter the user maintenance password → [Module] → [IBP]
 → [Calibration]. In the open interface, set the target pressure calibration value of the target channel. Input the range: 80 ~ 300mmHg.
- 4. Inflate with a spherical air pump to allow the pressure reading of the sphygmomanometer to get close to the pressure set in the menu.
- 5. Repeat the adjustment of the calibration pressure value in the menu and the pressure value of the sphygmomanometer until they are the same.
- 6. Select the [**Calibration**] button of the target channel and the defibrillation monitor will begin calibrating.
- 7. When calibration completes, it will display [**Calibration Completed**]. If it cannot be calibrated, it will display corresponding prompt message.

4.4.7 C.O. Test

- Patient simulator Medsim300B
- C.O. adapter box (for 300B)

- 1. Connect the simulator to the monitor C.O. parameter module using the main cable of C.O. accessory.
- 2. Set the simulator's blood temperature (TB) to 37 °C and ensure that the display value on the monitor does not exceed 37 ± 0.1 °C.
- 3. Set the [Injection Temp Source] to [Manual], adjust the [Injection Temp Value] (TI) until it displays 24 °C, select [C.O. Measure], open the C.O. measure window, and set [Comp Const] to 0.595.
- Set the injection temperature of the C.O. simulator to 24 °C, and the cardiac output to 5 L/min. Press
 [Start] in the C.O. measure window to start C.O. measure, then within 3 to 10s, press the RUN button of the simulator.
- 5. The displayed C.O. measure result is 5 ± 0.25 L/min.

4.4.8 Mainstream CO₂ Test

Precautions

• Before performing the mainstream CO₂ test, check and ensure that the atmospheric pressure setting in the [CO₂ Module Maintenance] of [Maintenance] is consistent with the local atmospheric pressure value.

- A steel cylinder containing $6 \pm 0.05\%$ CO₂ and a balance gas of N₂.
- Steel cylinder with 100% N₂ gas
- T connector
- Airway tube
- Flowmeter
- 1. Insert the module into module rack and connect a sensor.
- 2. When the CO₂ preheating startup completes, go to the [**CO**₂ **Setup**] menu, click [**Zero**] to conduct zero calibration for the CO₂ module. If the zero calibration fails, the screen prompts [**Zero Failed**]; if the zero calibration succeeds, the baseline of the CO₂ waveform on the screen returns to the zero position.
- 3. Go to [**CO**₂ **Setup**] menu and set [**Apnea Delay**] to 10s.
- Place the sensor in front of the mouth and breathe, so that the CO₂ waveform is generated on the screen, then place the sensor in the air and ensure that the monitor generates an alarm message [CO₂ Apnea].
- 5. Connect the test system as shown below.



- 6. Open the pressure reducing valve of the N2 cylinder and the CO₂ cylinder respectively, and ensure that only one cylinder is connected to the T-connector at the same time.
- Adjust the pressure reducing valve switch respectively so that the flow rate indicated by the flow meter is 2~5L/min and it keeps stable.
- 8. Switch the cylinder connected to the T-connector at an interval of $6s\sim10s$ and ensure that the CO_2 display value should be $45 \pm 2mmHg$.

4.4.9 Sidestream and Microstream CO₂ Testing and Calibration

See 4.2.3 Sidestream and microstream CO₂ testing and calibration.

4.5 Nurse Call Test

- Multimeter
- 1. Connect the nurse call line to the nurse call output port of the monitor.
- Set the monitor to [Demo] status, select [Main Menu] → [Maintenance] → enter the user maintenance password → [Alarm] → [Nurse Call].
- 3. In the [Nurse Call] menu, select [Alarm Priority] and [Alarm Category], and set [Trigger Mode] to [Normally Open].
- 4. In the [Nurse Call] menu, set [Signal Type] to [Pulse]. Set the monitor to generate an alarm, check the pulse with a pulse width of 1s when there is an alarm, and the multimeter measures the conducting state.

5. In the [Nurse Call] menu, set [Signal Type] to [Continuous]. Set the monitor to generate an alarm, check the continuous high level outputted when there is an alarm, and the multimeter can measure the conducting state.

4.6 Analog Output Test

Test tools:

- Patient simulator
- Oscilloscope
- 1. Connect the patient simulator to the monitor via an ECG line or IBP measurement line, and connect the oscilloscope to the analog output port.
- 2. Verify that the waveform displayed on the oscilloscope matches that displayed on the monitor screen.

4.7 Electric Safety Test

Warning

- Electrical safety test is used to test the electrical safety of the monitor under test. It's used to detect abnormal electrical hazards. If these hazards are not detected in time, it may cause harm to patients or operators.
- Electrical safety tests can be carried out using test equipment such as commercially available safety analyzers. Service personnel are required to ensure the applicability, functional integrity and safety of the test equipment, and familiarize with the use of it.
- Electrical safety test should comply with the following standards: EN 60601-1 and UL60601.
- If it's stipulated otherwise by local laws and regulations, please carry out relevant electrical safety tests in accordance with applicable regulations.
- In the patient area, all the equipment that uses AC power and is connected to medical equipment must comply with IEC 60601-1 and must receive electrical safety test at the intervals as required by the monitor.

Electrical safety test is used to detect electrical hazards that may cause harm to patients, operators and service personnel. Perform electrical safety tests under normal conditions (including temperature, humidity and atmospheric pressure).

The electrical safety test given in this chapter uses the 601 safety analyzer as an example. The safety analyzers used may vary by different regions. Please ensure the applicability of your electrical safety test plan.

See the figure below for equipment connection.



A: AC power (programmable power supply, regulate frequency)

B: Isolation figure for leakage current test on tooling

C: Safety tester

Test tools:

- Safety analyzer
- Isolation transformer

4.7.1 Housing Leakage Current Test

- 1. Connect the 601 safety analyzer to a 264 VAC, 60 Hz power supply.
- 2. Connect the equipment under test to the auxiliary power output jack of the 601 safety analyzer via a power line.
- 3. Connect one end of the red test lead to the "Red input terminal" of the safety analyzer and press the other end against the metal foil that is in close contact with the housing surface of the equipment under test.
- 4. Power on the 601 safety analyzer and press "5-Enclosure leakage" on the panel of the 601 safety analyzer to enter the interface for housing leakage current test.
- 5. Normally the leakage current of the housing is not more than 100 μ A, and not more than 300 μ A in a single fault condition.

4.7.2 Earth Leakage Current Test

- 1. Connect the 601 safety analyzer to a 264 VAC, 60 Hz power supply.
- 2. Connect the application part of the equipment under test to the RA side of the safety analyzer.
- 3. Connect the equipment under test to the auxiliary power output jack of the 601 safety analyzer via a power line.

- 4. Power on the 601 safety analyzer and press "4-Earth leakage" on the panel of the 601 safety analyzer to enter the interface for earth leakage current test.
- 5. Normally the earth leakage current is not more than 300 μ A, and not more than 1000 μ A in a single fault condition.

4.7.3 Patient Leakage Current Test

- 1. Connect the 601 safety analyzer to a 264 VAC, 60 Hz power supply.
- 2. Connect the application part of the equipment under test to the RA side of the safety analyzer.
- 3. Connect the equipment under test to the auxiliary power output jack of the 601 safety analyzer via a power line.
- 4. Power on the 601 safety analyzer and press "6-Patient leakage" on the panel of the 601 safety analyzer.
- 5. Press the "APPLIED PART" button continuously to select AC and DC measurements. When it's DC, "DC" is displayed behind the limit value.
- 6. Normally the patient leakage current is not more than 10 μ A, and not more than 50 μ A in a single fault condition.

4.7.4 Patient Auxiliary Current Test

- 1. Connect the 601 safety analyzer to a 264 VAC, 60 Hz power supply.
- 2. Connect the equipment under test to the auxiliary power output jack of the 601 safety analyzer via a power line.
- 3. Connect the ECG line of the equipment under test to the RA end of the safety analyzer.
- 4. Power on the 601 safety analyzer and press the "8-Patient Auxiliary Current Test" on the panel of the 601 safety analyzer to enter the interface for patient auxiliary current test.
- 5. Press the "APPLIED PART" button continuously to select AC and DC measurements. When it's DC, "DC" is displayed behind the limit value.
- 6. Normally the patient auxiliary current is not more than 10 μ A, and not more than 50 μ A in a single fault condition.

4.8 Recorder Check

- None
- 1. Print the ECG waveform, the recorder should be able to print normally with sound clarity and consistency.
- 2. Setup should ensure that relevant prompts should be displayed if it is out of paper or the buckle cambers or other malfunctions, and it should work normally after recovery.

3. Perform alarm printing for each parameter, turn on the alarm recording switch of each parameter, set different alarm limits, and ensure to conduct printing for each parameter alarm.

4.9 Network Printing Check

Test tools:

Hub and network cable

4.9.1 Equipment Connection and Setup

1 Equipment connection: Connect the monitor and network printer to the hub via a common network cable, as shown in the following figure:



2 IP setup: Go to the "Main Menu" interface, select [Main Menu] → [Maintenance] → enter the user maintenance password, select [Network Setup] in the maintenance interface, and set the IP address of the monitor to be in the same network segment with the IP address of the network printer (See the instructions for the printer).

4.9.2 Printing Function Test

- 1 Enter demo mode of the monitor.
- 2 Go to the "Main Menu" interface, select [Interface Setup] → [Interface Layout], and select "ECG Full-Screen 7 Lead Interface".
- 3 Go to the "Main Menu" interface, select [**Normal Report**] → [**ECG Report**], select normal report, click "Print", and the network printer should print out the ECG report.

4.10 Battery Check

Test tools:

None

Function check

- 1. Check that the monitor works properly when powered by AC.
- 2. Remove the AC power line and check that the monitor continues to operate properly.

Performance check

Please refer to the relevant contents in the **battery** section of the operator's manual to check that the battery power supply time meets the specifications.

4.11 Charging Dock Check

Test tools:

Multimeter

Function check

- 1. Disconnect the charging dock from the monitor and connect the charging dock to the DC power supply (12V-28) on the ambulance.
- 2. When the charging dock is powered on, the green power indicator lights are lit.
- 3. Use a multimeter to measure the output port voltage of the transmission dock. The voltage should be (15V +5%).
- 4. Connect the monitor and check if it can be turned on properly.

4.12 Factory Maintenance

4.12.1 Enter Factory Maintenance

Select [**Main Menu**] hot key \rightarrow turn to the page 3 \rightarrow select [**Maintenance**] from the [**System**] column \rightarrow enter the factory maintenance password \rightarrow select [**Ok**], then select factory maintenance.

4.12.2 Monitor linformation (Exporting Log)

Before inserting the USB flash drive, confirm that its format is FAT32, and then insert it into the USB port of the monitor's main unit (Note: Not the USB port of the iView's main unit!),

Open the [Monitor Information] menu, you can see the information about the monitor's main unit, such as: CPU temperature, Wifi signal strength, hard disk capacity and so on.

At this point, you can export all the monitor's log information by clicking the "Export Log" button in the bottom left corner of the window.

4.12.3 Production Test

Open the [**Production Test**] menu and you can perform basic function tests related to the monitor's hardware interface.

Production test includes auto test and single test.

- Auto test: After selecting [**Start Auto Test**], the system will automatically completes all tests in order.
- Single test: Select one for a single test.

4.12.4 Setup

Open the [Setup] menu and you can do ECG alarm settings and other configurations.

4.12.5 Debugging

Open the [**Debug**] menu and you can make settings related to debugging tests.

4.12.6 Power Information

Open the [Power Info] menu and you can check the power status of the monitor.

4.12.7 Clinical Data

Open the [**Clinical Data**] menu and you can do settings related to clinical data collection. When the Clinical Data Location is selected as "Local", the clinical data is stored in the monitor. You can export the data to the USB flash drive by the same way as exporting the log. When the Clinical Data Location is selected as "Udisk", the clinical data is directly stored in the U disk.

4.12.8 Sending Clinical Data

Open the Send Clinical Data menu and you can select the clinical data you need to send.

4.12.9 Software Version

Open the maintenance menu and select the [Version] menu, you can view all the software versions in the system that can be viewed.

Maintenance test report

(For detailed test steps and contents, see the above section)

Client name		
Client address		
Maintenance people		
Maintenance company name		
Name of device under test		
Model of device under test		
Serial No. of device under test		
Hardware version		
Software Version		
Test device	Model/No.	Validity of calibration

Test Item		Test result	Test result (Pass/Fail)		
Visual inspection					
There should be no physical dar	nage to the monitor housing,				
display, buttons, SMR, modules,	power cords, brackets and module				
accessories					
The external connection cable s	hould be free of wear and the				
connector pins should have no	looseness or distortion				
The monitor's peripheral interfa	ce should have no looseness or				
distorted pins					
Safety labels and nameplates sh	ould be clearly legible				
Power On test					
It passes power-on test, the pov	ver indication and alarm system work				
normally, and the monitor starts	s normally.				
Performance test					
ECG performance test					
The ECG waveform is normal an	d noise-free, and the HR value				
displayed on the monitor must					
The "ECG Lead Off" alarm works					
When the pacing is turned on, t	he pacing signal can be detected				
and the pacing mark is displaye	d.				
The difference between ECG cal	ibrated waveform amplitude and				
scale amplitude should not exce	eed 5%				
RESP test					
The monitor shows that the RES	P waveform is not distorted, and the				
displayed Resp value must not e	exceed 20 \pm 1 rpm.				
SpO ₂ test					
Connect healthy people's finger	to test, the monitor displays the				
blood oxygen pleth waveform a	nd the PR value, and the displayed				
SpO₂ value range should be 95% to 100%.					
SpO ₂ sensor off alarm function is normal					
NIBP test					
For NIBP pressure test, 0, 50 and 200mmHg test difference does not					
exceed ± 3mmHg					
There is no gas leakage in NIBP, or the manual leakage test result					
does not exceed 6mmHg/min.					
Temp test					
The display value of each tempe					

not exceed $37 \pm 0.1^{\circ}C$	
IBP test	·
Static pressure display value of each IBP channel does not exceed	
200 ± 2mmHg	
The ART and LV waveforms of each IBP channel are displayed	
correctly	
C.O. test	·
Monitor's TB display value does not exceed $37 \pm 0.1^{\circ}$ C	
The displayed C.O. measurement result is 5 ± 0.25 L/min.	
Mainstream CO ₂ test	
The mainstream CO_2 zero calibration is successful, and the baseline	
of the CO ₂ waveform returns to the zero position on the screen.	
The CO ₂ Apnea alarm is normal.	
The CO $_2$ display value should be 45 \pm 2mmHg	
Sidestream CO ₂ test	·
After blocking the air inlet of the water tank, the flow rate of the	
sidestream CO ₂ is <10ml/min, and it alarms CO ₂ airway is blocked,	
and there is no air leakage	
CO_2 display value should be 6 \pm 0.2%	
Microstream CO ₂ test	·
After the microstream CO_2 is blocked for about 30s, it alarms	
CO2sampling airway is blocked, and the module has no air leakage	
The CO ₂ display value should be 45 \pm 2mmHg	
Nurse call test	·
Nurse call cable conducts when the monitor alarms	
Analog output test	·
The waveform displayed by the oscilloscope should be consistent	
with that displayed on the monitor's screen.	
Electric safety test	
Normally the housing leakage current is not more than 100 μ A, and	
not more than 300 μ A in a single fault condition.	
Normally the earth leakage current is not more than 300 μ A, and not	
more than 1000 μ A in a single fault condition.	
Normally the patient leakage current is not more than 10 μ A, and	
not more than 50 μ A in a single fault condition.	
Normally the patient auxiliary current is not more than 10 μA , and	
not more than 50 μ A in a single fault condition.	
Recorder check	
Can print ECG waveforms normally with sound consistency and	
clarity	
Setup should ensure that relevant prompts should be displayed if it	
is out of paper or the buckle cambers or other malfunctions, and it	
should work normally after recovery	

The alarm recording function of each parameter is normal.	
Network printing check	
4 The network printer can print an ECG report properly	
Battery check	
Batteries support monitor to continue to operate normally when AC	
power is accidentally disconnected	
Battery power supply time meets product's specifications	

Test conclusion:

Pass or not: (Yes No)

Sign by the tester:

Test Date:

FOR YOUR NOTES

5.1 Introduction

This chapter lists the problems that may occur during the use of the monitor and recommended measures. Check the monitor as per the table given in this chapter to identify and resolve problems. For more information on troubleshooting, please contact Mindray after-sales service department.

5.2 Component Replacement

The PCB, main parts and components of the monitor can be replaced. Once PCB failure is identified, replace the PCB as per the instructions given in the Repair and Disassembly section. Then confirm that the monitor can work properly and passes all performance tests. For information on replaceable parts, please refer to *8 Parts*.

5.3 Checkup before Starting the Monitor

In addition, it is necessary to check whether the appearance is damaged before starting the machine. In particular, if the touch screen of the screen assembly is damaged, do not use it.

5.4 Software Version Check

In some troubleshooting, it may involve the compatibility of software version. In this case, you need to know the monitor configuration and software version. For details about version compatibility, please contact our after-sales service department. Please follow the steps below to check software version:

Select [Main Menu] \rightarrow turn to the third page \rightarrow select [Maintenance >>] from the [System >>] column \rightarrow enter the factory maintenance password \rightarrow [Version >>]. In the pop-up menu, you can view the information about the system software version.

5.5 Technical Alarm Messages

This section lists technical alarms, their default priority, indication on alarm reset, and the actions that can be taken when an alarm occurs.

Technical alarms give different alarm indicators when the alarm system is reset. In this section we classify the technical alarms into three categories for easy clarification:

• A: technical alarms are cleared. The monitor gives no alarm indications.

- B: technical alarms are changed to the prompt messages.
- C: the alarm is silenced and a √ appears before the alarm message, indicating that the alarm is acknowledged.

In the following tables we will use A, B, and C to refer to the indications on alarm reset.

5.5.1 General Technical Alarm Messages

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
XX Module Error	High	С	XX module does not work properly. Replug
			the module, if the alarm persists, contact your
			service personnel.

Note: XX represents a measurement or parameter label, such as HR, RR, SpO₂, EtCO₂, and so on.

5.5.2 ECG Technical Alarm Messages

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
ECG Noisy	Low/Prompt	А	The ECG signal is noisy. Check for any possible
			sources of signal noise around the cable and
			electrode, and check the patient for excessive
			motion.
ECG Amplitude Too	Low	с	The ECG amplitude does not reach the
Small			detected threshold. Check for any possible
			source of interference around the cable and
			electrode.
ECG Lead Off	High, Med, or	В	The electrode has become detached from the
	Low,		patient or the lead wire has become
	configurable		disconnected from the adapter cable. Check
			the connections of the electrodes and
			leadwires.
ECG XX Lead Off	High, Med, or	В	The electrode has become detached from the
	Low,		patient or the lead wire has become
	configurable		disconnected from the adapter cable. Check
			the connections of the electrodes and
			leadwires.
ECG Signal Invalid	Low	А	Patient skin impedance is too high. Check
			ECG electrode application.
ECG Learning	Prompt	/	ECG learning is manually or automatically
			triggered.
Cannot Analyze QT	Prompt	/	/
D12L not available	Prompt	с	The current Va and Vb combination does not
			support D12L. Choose an available Va and Vb
			combination. For more information, see 9.5
			Using 6-lead Placement to Derive 12-lead ECG

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
			(D12L).

Note: XX represents ECG lead name, for example RL, LL, V, Va, Vb, and so on.

5.5.3 Resp Technical Alarm Messages

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
Resp Interference	Prompt	/	The respiration circuit is disturbed. Check for
			any possible sources of signal noise.
Electrode Poor Contact	Prompt	/	Check the electrode application. Reposition or
			replace the electrodes if necessary.

5.5.4 SpO₂ Technical Alarm Messages

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
SpO2 Sensor Off	Adjustable	В	The SpO ₂ sensor has become detached from
			the patient or the module. Check the sensor
			connection. If the alarm persists, replace the
			sensor.
SpO2 No Sensor	Low	A	The SpO ₂ extension cable is detached from
			the SpO ₂ module, or the SpO ₂ sensor is
			detached from the SpO ₂ extension cable.
			Check the SpO $_2$ cable and the sensor
			connection. If the alarm
			persists, replace the sensor.
SpO2 Excess Light	Low	с	Ambient light is too strong. Move the sensor
			to a place with lower level of ambient light or
			cover the sensor to minimize the ambient
			light.
SpO ₂ No Pulse	Low	с	The SpO $_2$ sensor failed to obtain pulse signal.
			Check the patient's condition and replace the
			sensor application site. If the alarm persists,
			replace the sensor.
SpO ₂ Sensor	Low	с	Incompatible or an unspecified SpO ₂ sensor is
Incompatible			used. Use specified sensors.
SpO ₂ Low Signal Quality	Low	с	1. Check the sensor and sensor position.
			2. Make sure the patient is not shivering or
			moving.
			3. The patient's pulse may be too low to be
			measured.

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
SpO ₂ Interference	Low	с	The SpO_2 signal has been interfered. Check
			for any possible sources of signal noise and
			check the patient for excessive motion.
SpO ₂ Sensor Error	Low	С	Replace the sensor and measure again.
SpO ₂ Searching Pulse	Prompt	/	SpO ₂ is searching for pulse.
SpO ₂ Low Perfusion	Prompt	/	The SpO ₂ sensor is not properly placed or the
			patient's perfusion index is too low.
			1. Check the sensor and sensor position.
			2. Reposition the sensor if necessary.

5.5.5 Temp Technical Alarm Messages

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
T XX Sensor Off	Low	А	Check the sensor connection and reconnect
			the sensor.

Note: XX represents a temperature site, for example skin, core, axil, T1, and so on.

5.5.6 NIBP Technical Alarm Messages

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
NIBP Cuff Loose	Low	А	There is a leak in the cuff or air tubing. Use a
			cuff of correct type based on the patient size.
			Apply the cuff and connect the air tubing as
			instructed in the manual.
NIBP Cuff or Airway Leak	Low	А	Check the NIBP cuff and pump for leakages.
NIBP Airway Error	Low	A	The air tubing may be occluded. Check the air
			tubing for an occlusion or kinking. If the
			alarm persists, contact your service
			personnel.
NIBP Weak Signal	Low	A	The patient's pulse is weak or the cuff is loose.
			Check the patient's condition and replace the
			cuff application site.
NIBP Overrange	Low	A	The measured NIBP value exceeds the
			module measurement range. Check the
			patient's condition.
NIBP Excessive Motion	Low	A	Check the patient's condition and reduce
			patient motion.
NIBP Cuff Overpressure	Low	A	The NIBP airway may be occluded. Check the
			airway and measure again. If the alarm
			persists, contact your service personnel.

NIBP Timeout	Low	A	The measurement time exceeds 120 seconds
			in the adult or pediatric mode, or exceeds 90
			seconds in the neonatal mode, and the BP
			value cannot be obtained. Check the patient's
			condition and NIBP connections, or replace
			the cuff and measure again.
NIBP Cuff and Patient	Low	A	The cuff type mismatches the patient
Mismatch			category. Verify the patient category or
			replace the cuff if necessary. If patient
			catergory is correct, check that the tubing is
			not bent and the airway is not occluded.
NIBP Airway Leak	Low	A	Airway leakage is found during the NIBP
			leakage test. Check the NIBP cuff and pump
			for leakages.

5.5.7 IBP Technical Alarm Messages

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
XX Sensor Error	Med	С	The IBP sensor fails. Replace the sensor.
XX No Sensor	High, Med, or	A	The IBP patient cable and/or corresponding
	Low, configurable		IBP sensor is not connected or detached.
			Check the cable and sensor connection.
XX No Pulse	Low	A	The catheter may be occluded. Please flush
			the catheter.
XX Disconnected	High	С	The liquid way is disconnected from the
			patient, or the three-way valve is open to the
			air. Check the connection of the liquid way, or
			check the valve is open to the patient. If the
			alarm persists, contact your service
			personnel.

Note: XX represents an IBP label, for example PA, CVP, FAP, P1, and so on.

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
TB Sensor Off	Low	A	Check the sensor connection and reconnect
			the sensor.
T1 Sensor Off	Low	A	Check the sensor connection and reconnect
			the sensor.

5.5.8 C.O. Technical Alarm Messages

5.5.9 CO₂ Technical Alarm Messages

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
CO ₂ Module High Temp	Low	С	Ambient temperature is too high or there is a
			module failure.
			1. Lower the operating temperature.
			2. Replug the module.
			3. If the alarm persists, the CO_2 module may
			fail, contact your service personnel.
CO ₂ Module Low Temp	Low	С	Ambient temperature is too low or there is a
			module failure.
			1. Raise the operating temperature.
			2. Replug the module.
			3. If the alarm persists, the CO_2 module may
			fail, contact your service personnel.
CO ₂ Zero Failed	Low	С	For mainstream CO ₂ module, check the
			connections between the adapter and CO_2
			transducer. Wait till the sensor's temperature
			becomes stabilized, and then perform a zero
			calibration again.
			For sidestream CO ₂ module, replug the
			module. If the alarm persists, contact your
			service personnel.
CO ₂ No Watertrap	Low	В	Check the watertrap connections.
CO ₂ High Airway	Low	С	1. Check the airway pressure settings of the
Pressure			ventilator/anesthesia machine.
			2. Disconnect the module from the
			ventilator/ anesthesia machine.
			3. Replug the module.
			4. If the alarm persists, contact your service
			personnel.

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
CO ₂ Low Airway Pressure	Low	С	1. Check the airway pressure settings of the
			ventilator/anesthesia machine.
			2. Disconnect the module from the
			ventilator/ anesthesia machine.
			3. Replug the module.
			4. If the alarm persists, contact your service
			personnel.
CO ₂ High Barometric	Low	с	The ambient pressure exceeds the operating
			pressure range or CO ₂ module fails.
			1. Make sure that the ambient pressure
			meets the specifications, and check for
			sources that affect the ambient pressure.
			2. Replug the module.
			3. If the alarm persists, contact your service
			personnel.
CO ₂ Low Barometric	Low	С	The ambient pressure exceeds the operating
			pressure range or CO_2 module fails.
			1. Make sure that the ambient pressure
			meets the specifications, and check for
			sources that affect the ambient pressure.
			2. Replug the module.
			3. If the alarm persists, contact your service
			personnel.
CO ₂ Airway Occluded	Low	с	1. Check if the sample line is kinked or
			occluded.
			2. Replace the sample line.
			3. Replug the module.
			4. If the alarm persists, contact your service
			personnel.
CO ₂ No Filterline	Low	А	Make sure that the filterline is connected.
CO ₂ Calibration Required	Low	С	Perform a calibration.
CO ₂ Airway Error	Low	С	1. Check if the sample line is kinked or
			occluded.
			2. Replace the sample line.
			3. Replug the module.
			4. If the alarm persists, contact your service
			personnel.
CO ₂ Adapter Error	Low	A	Check, clean or replace the airway adapter.
			Perform a zero calibration.
CO ₂ No Sensor	Low	A	Make sure that the CO ₂ transducer is
			connected.

Alarm message	Default priority	Indication on alarm reset	Cause and solution
CO ₂ : Change Watertrap	Low	С	Replace the watertrap.
CO ₂ Watertrap and	Low	С	Check the patient category and use a correct
Patient Mismatch			watertrap.
CO ₂ : Change O ₂ Cell	Low	с	The oxygen sesnor is depleted or fails.
			Replace the oxygen sensor.

5.5.10 AG Technical Alarm Messages

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
AG No Watertrap	Low	В	Check the connections of the watertrap and
			re- connect it.
AG: Change Watertrap	Low	с	Replace the watertrap.
AG Watertrap and	Low	с	Check the patient category and use a correct
Patient Mismatch			watertrap.
AG Zero Failed	Low	с	There is external electromagnetic
			interference, airway occlusion or module
			failure.
			1. Check for external inference sources.
			2. Check for "AG Airway Occluded" alarm
			message. Remove the occlusion.
			3. If the alarm persists, contact your service
			personnel.
Anesthetic Mixture	Low	С	Anesthetic mixture is detected.
AG Airway Occluded	Low	с	1. Check if the sample line is occluded.
			2. Check the sample line.
			3. Replug the module.
			4. If the alarm persists, contact your service
			personnel.

5.5.11 BIS Technical Alarm Messages

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
BIS Sensor Off	Low	A	Check and reconnect the BIS sensor. If the
			alarm persists, replace the sensor.
BIS Electrode XX Off	Low	A	Check the electrode connection, and
			re-attach the electrodes if necessary.
BIS Electrode XX Poor	Low	A	Enter the Sensor Check menu, and check the
Contact			connections of the sensor and electrodes.
BISx Error	High	с	Replug the module. If the alarm persists,
			contact your service personnel.

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
BIS No Sensor	Low	А	BIS sensor is not properly connected, BIS
			cable fails, BISx or BISx4 fails.
			1. Check the BIS sensor connection.
			2. Replug the BIS module.
			3. Replace the BIS cable.
			4. Replace BISx or BISx4.
BIS Sensor Too Many	Low	А	Replace the sensor.
Uses			
BIS Sensor Old	Low	А	Replace the sensor.
BIS XX Signal Quality Too	Low	A	SQI < 15
Low			1. Check the patient's condition.
			2. Check the sensor position, and its contact
			with the patient's skin.
			3. Check that BISx or BISx4 is away from the
			electrically radiating equipment.
BIS XX Low Signal	Low	A	SQI < 15
Quality			1. Check the patient's condition.
			2. Check the sensor position, and its contact
			with the patient's skin.
			3. Check that BISx or BISx4 is away from the
			electrically radiating equipment.
BIS Wrong Sensor Type	Low	А	Check or replace the sensor.
BIS Sensor Error	Low	С	Replace the sensor.
Disconnect/Reconnect	Low	А	Replug the BIS Module.
BIS			

Note: XX represents BIS label, for example G, C, LE, LT, RL-RA, L-R, F-R, 1, 2, and so on.

5.5.12 Power Supply Technical Alarm Messages

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
Low Battery	Med	с	Connect the monitor to the external power
			supply and allow the batteries to charge.
Critically Low Battery	High	с	Connect the monitor to the external power
			supply and allow the batteries to charge.
Power Board Comm	High	с	Restart the monitor. If the alarm persists,
Error			contact your service personnel.
Battery Error	High	с	The battery may fail. Contact your service
			personnel.
RT Clock Need Reset	High	С	Contact your service personnel.
RT Clock Not Exist	High	С	Contact your service personnel.

Alarm message	Default priority	Indication on alarm reset	Cause and solution
XX V Too High	High	С	There is a problem with the system power
XX V Too Low	High	С	supply. Restart the monitor.

Note: XX represents 2.5 V, 3.3 V,5 V, or 12 V.

5.5.13 Recorder Technical Alarm Messages

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
Recorder Init Error	Low	A	An error occurred during the recorder
			initialization. If the alarm persists, contact
			your service personnel.
Recorder Comm Error	Low	A	Restart the monitor if not solved. If the alarm
			persists, contact your service personnel.
Recorder Head Hot	Low	С	The recorder has been working for too long
			time. Stop the recording and resume the
			recording till the recorder's print head cools
			down.
Recorder Initializing	Prompt	/	Wait until the recorder initialization is
			completed.
Recorder out of Paper	Prompt	/	The recorder paper is not loaded or the recorder
			door is not closed. Check the recorder, load the
			recorder paper or close the recorder door.
Recorder Busy	Prompt	/	The buffer queue for recording is full.

5.5.14 Printer Technical Alarm Messages

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
Printer Buffer Full	Prompt	/	The printer buffer is full. Wait till the printer
			finishes the printing task.
Fail	Prompt	/	The printer runs out of paper or cannot be
			connected. Check the printer.
Printing Stopped	Prompt	/	Printing is manually stopped.
Printer Unavailable	Prompt	/	The printer may fail. Check the printer.
PDF storage space is	Prompt	/	Delete the files saved under the PDF file path
nearly full			to release storage space. Otherwise you
			cannot save new PDF files.
Error storing PDF file	Prompt	/	The PDF file path settings on the printer
			server and the PDFCreator are not consistent
			or the PDF storage space is full. Check the
			PDF file path settings for consistency, or
			delete the files saved under the PDF file path
			to release storage space.

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
Change the print server	Prompt	/	Verify that the language settings of the
language to be			printer server and the monitor are consistent,
consistent with this			Otherwise you cannot perform printing.
monitor			
Print Server	Prompt	/	Check that the monitor is properly connected
Disconnected			with the printer server.

g
9

Alarm message	Default	Indication on	Cause and solution
	priority	alarm reset	
No CMS	Low	В	The monitor is disconnected from the CMS.
			Check the network connection.
View Bed XX YY-ZZ,	Low	A	The network is interrupted when the monitor
Network Disconnected.			is viewing the remote device. Check the
			network connection.
Viewed by Bed XX YY-ZZ,	Low	A	The network is interrupted when the monitor
Network Disconnected.			is viewed by another remote device. Check the
			network connection.
WLAN IP Address Conflict	Low	с	Wireless network IP network conflicts. Check
			the network settings.
LAN1 IP Address Conflict	Low	с	Wired network LAN1 IP network conflicts.
			Check the network settings.
Fail To Get WLAN IP	Low	с	Unable to automatically obtain the wireless
Address			network IP address. Check the network
			settings.
Fail To Get LAN1 IP	Low	с	Unable to automatically obtain the wired
Address			network LAN1 IP address. Check the network
			settings.

Note: XX refers to the department name, YY refers to the room number, and ZZ refers to the bed number.

5.5.	16	Other	System	Technical	Alarm	Messages
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Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
Storage Card Error	High	с	The storage card fails or files are damaged.
			Restart the monitor to format the storage
			card. If the alarm persists, contact your service
			personnel.
Loading Default Config	Low	А	The default configuration is not correctly
Failed			loaded. The monitor will restore to the factory
			default configuration for the current patient
			category.

Alarm message	Default priority	Indication on	Cause and solution
		alarm reset	
XX Conflicts	Prompt	/	The same type of corresponding module
(XX refers to the module			being used exceeds the supported number.
label)			Remove the conflict module.
XXX Measurement has	Prompt	/	The parameter module is disabled. Switch on
been closed			the module if you want to use it. For more
(XX refers to the module			information, see 3.11.1 Switching On or Off a
label)			Parameter.
The display setup for	Prompt	/	The parameter of the newly inserted module
XXX is disabled.			is not displayed on the screen. Select a
(XX refers to the			desired area to display the parameter
parameter label)			numerics and waveforms. For more
			information, see 26.11 The Other Settings.
The patient data storage	Configurable	В	Delete unnecessary earlier discharged
space is nearly full.			patient.
Please delete some			
discharged patients.			

5.6 Troubleshooting Guide

5.6.1 Problems with Turning on/off

Troubles	Possible causes	Troubleshooting	
Unable to turn the monitor on	AC power is not connected, or a low battery or a damaged battery	 Please confirm if the AC power is connected properly. Check if the battery is low or if the battery is damaged. 	
	Connection line failure	 Check if the line between the ON/OFF button board and the main control board is connected properly. Check if the plug of the connection line and the corresponding socket are damaged. 	
	ON/OFF button board is damaged	Replace the ON/OFF button board	
	Power module is damaged	Replace the power module	
	The main control board is damaged	Replace the main control board	
	Power protection	If there are other devices connected to the main unit, such as external parameter modules, USB flash drives and scanners, disconnect them from the main unit first. If you can turn it on after the disconnection, it's possible that abnormal external devices trigger power protection.	

Troubles	Possible causes	Troubleshooting
		If there are no other devices connected to the main
		unit, check the backboard of integral module rack
		or the main control board to see if there are short
		circuits which result in power protection.



Troubleshooting flowchart for the inability to turn it on due to battery failure



Troubleshooting flowchart for the inability to turn it on due to AC power failure
5.6.2 Display Failures

Troubles	Possible causes	Troubleshooting
Unable to	Connection line	 Check if the display is properly connected to the main
display, or	failure	control board (screen and backlight line).
abnormal		Check if the line plug and corresponding socket are
display, but the		damaged
main unit can	LCD display is	Replace the front housing assembly
work	damaged	
	Main control board	Update the main control board software
	software goes wrong	
	The backlight drive is	Change the main control board
	damaged	
	The display drive is	Change the main control board
	damaged	
The touch screen	Connection line	 Check if the touch screen is properly connected to the main
does not	failure	control board
respond.		 Check if the line plug and corresponding socket are
		damaged
	The touch screen is	Replace the front housing and touch screen repair kit
	damaged	
	Main control board	Update the main control board software
	software goes wrong	
	The main control	Change the main control board
	board is damaged	



Display troubleshooting flowchart

5.6.3 Alarm Failures

Troubles	Possible causes	Troubleshooting
The alarm indicator light is always off or	Connection line failure	 Check if the alarm light board is properly connected to the main control board. Check if the connection line or plug is damaged.
always on, but the alarm sounds	Alarm indicator light board is damaged	Replace the alarm indicator light board
generated properly.	The main control board is damaged	Change the main control board
No alarm sound is generated, but the alarm	Alarm sound is turned off	Check whether the alarm sound is set to be silent; select [Main Menu] \rightarrow [System >>] \rightarrow [Maintenance >>] \rightarrow enter the user maintenance password \rightarrow [Alarm >>], and adjust the [Minimum Alarm Volume] to an appropriate value in the pop-up menu. Select [Alarm] \rightarrow [Setup] on the main menu to adjust the alarm volume.
indicator light	Speaker damage	Replace the speaker
works properly.	Connection line failure	Check if the speaker is properly connected to the main control board.
	The main control board is damaged	Change the main control board

5.6.4 Recorder Failures

Troubles	Possible causes	Troubleshooting	
	Recorder module is not	Check if the recorder's work indicator light is on.	
	allowed	If it is on, restore its function in the factory maintenance.	
	There's a paper jam	Reinstall printing paper	
The recorder is		Check if the recorder is properly connected to the main	
unable to print	Connection line failure	control board.	
unable to print		Check if the connection line or plug is damaged.	
	The recorder is damaged	Replace the recorder	
	The main control board is	Change the main control board	
	damaged		
	Thermal-sensitive		
	coating of the printing	Replace the printing paper	
Poor printing effect	paper fails		
	The thermal head is dirty	Clean the thermal head	
	The recorder is damaged	Replace the recorder	

5.6.5 Output Interface Failures

Troubles	Possible causes	Troubleshooting
Nurse call signal is not outputted	The main control board is damaged	Change the main control board.
USB Device Unusable.	The main control board is damaged	Change the main control board.
The network interface cannot be used	The main control board is damaged	Change the main control board.
VGA interface cannot be used	The display does not match with the timing sequence of VGA interface	Please confirm whether the display supports the resolution of 1280*800 (10-inch/12-inch model) or 1376 * 768 (15-inch).
	The main control board is damaged	Change the main control board.

5.6.6 Battery Failures

Troubles	Possible causes	Troubleshooting
	Battery is damaged	Replace the battery
Battery cannot supply power	Connection line failure	 Check if the main control board is properly connected to the battery interface board. Check if the connection line or plug is damaged.
	Battery is damaged	Replace the battery
The battery cannot be fully charged or cannot be	Connection line failure	 Check if the main control board is properly connected to the battery interface board. Check if the connection line or plug is damaged.
charged	The main control board is damaged	Change the main control board

5.6.7 Parameter Module Failures

Troubles	Possible causes	Troubleshooting
	Incorrect software version	Check the MPM and system software versions and update the software
Troubles ECG/Resp/SPO2 (Mindray)/NIBP/Temp/IBP do not work SPO2 module (nellcor) cannot work CO2 (built-in) module cannot work	Parameter circuit is damaged	Change the main control board
	Connection line failure	 Confirm whether the parameter board and the main board are properly connected. Confirm if these connection lines and connectors are intact.
	Accessories may be damaged	Replace the accessories.
	Check whether the parameter configuration is correct	According to the user manual, confirm whether the corresponding parameter configuration has been opened
CPO, modulo (nollcor) connet work	Accessories may be damaged	Replace accessories
SPO ₂ module (nelicor) cannot work	SPO2 module failure	Replace the module.
CO ₂ (built-in) module cannot work	Internal air pipe connection failure	 Check if the internal air pipe is properly connected Check if the internal air pipe is damaged

Troubles	Possible causes	Troubleshooting
	System software configuration error	Check if the modules in the system software are configured correctly.
	The module is damaged	Replace the module
	Connection line failure	 Check if the CO₂ module is properly connected to the main control board. Check if the line is damaged
	The main control board is damaged	Change the main control board
C.O. (built-in) module cannot work	System software configuration error	Check if the modules in the system software are configured correctly.
	The module is damaged	Replace the module
	Line connection failure	 Check if the parameter board is properly connected to the C.O. module and the C.O. module is properly connected to the main control board. Check if the connection line is damaged
	The main control board is damaged	Change the main control board

5.6.8 Network Failures

Troubles	Possible causes	Troubleshooting
Frequent disconnection of network	The network cable is not properly connected.	Check if the network cable is connected properly, or whether the network cable is too close to the power cable of a high-power device, or whether the network cable is too long (cannot exceed 50m).
	Network settings error	Check for IP conflicts in the network. If yes, reset the network.
The network has been connected, the Viewbed	The network cable is not properly connected	Check if the network cable is connected properly, or whether the network cable is too close to the power cable of a high-power device, or whether the network cable is too long (cannot exceed 50m).
function cannot be realized	Have got too many monitors under observation	Confirm the number of monitors that can be connected at the same time according to the user manual.

Troubles	Possible causes	Troubleshooting		
	Network settings error	Check for IP conflicts in the network. If yes, reset the network.		
WLAN cannot be connected	Network settings error	Check whether WLAN settings are correct.		
	The antenna is not properly installed	Please check if the antenna of the wireless network card is properly connected to the wireless module.		
	Wireless module is damaged	Please replace the wireless module.		
	The main control board is damaged	Replace the main control board.		

5.6.9 Software Upgrade Failures

Troubles	Possible causes	Troubleshooting	
Program upgrade fails	Connection error	 Check if the network cable is connected to the monitor's network interface instead of iView's network interface. Make sure the hub or switch can work properly and check if the hub is properly connected. 	
	Program upgrade package error	Please select the correct upgrade package.	
	Error in the IP address setting of PC	Set a fixed IP address for the upgraded monitor within the specified Class C addresses. It is not recommended to upgrade the program in a network with multiple PCs connected.	

6.1 Introduction

This monitor supports upgrade of the monitoring parameter function module, function component, and software.

Precautions

- Before upgrading the function of a monitor to be disassembled, eliminate static electricity. When disassembling components with electrostatic sensitive mark, wear ESD wristband or gloves to avoid damage to the components.
- During reinstallation, connect and place cables properly to avoid crushing the cables, which may cause a short circuit.
- During reinstallation, select screws with proper model. If you forcibly tighten improper screws, the device may be damaged. And the screws or components may become loose during use, which causes unpredictable product damage or personal injury.
- Disassemble the device based on the disassembly sequence. Otherwise, an irreversible damage may be caused to the device.
- Before disassembling components, ensure that all connected cables are removed. During disassembly, avoid pulling off the cables or damaging the connectors.
- Loosened screws and other parts should be categorized and placed properly so that they can be used for reinstallation and will not be dropped, polluted or lost.

6.2 Parameter Function Module Upgrade

This monitor supports upgrade of the following parameter function modules:

Embedded Parameter	DN	Name and Engrification	Bomarke
Module	FIN	Name and Specification	Rellidiks
IBP module	115-059953-00	Integrated IBP material package FRU	
C.O. module	115-059954-00	Integrated C.O. material package FRU	
Sidestream CO ₂ module		Integrated 10/12-inch sidestream CO ₂	
	115-059974-00	material package FRU	
Microstream CO ₂ module	115-050056-00	Integrated microstream CO ₂ material	
	115-059950-00	package FRU	
Mainstream CO ₂ module	115-059957-00	Integrated mainstream CO ₂ material	
	115-059937-00	package FRU	

For details about the use method of different parameter modules, see ePM series user manual. The following content describes the upgrade methods of the parameter modules:

6.3 Upgrading Parameter C.O. Function Module

When upgrading the C.O. function, install the upgrade material packag. The operation procedure is as follows:

- 1. Disassemble the main control board by referring to section "Disassembling Main Control Board" in the Disassembly and Repair part.
- 2. Install the C.O. board in the material package to the main control board by referring to "Disassembling C.O. Board".



- 3. Reassemble the machine.
- Start the monitor, and choose [Main Menu] > [Maintenance] > [Factory Maintenance] > [Factory Default], enable [C.O.], and restart the machine to validate the configuration.
- 5. Test the upgraded machine by referring to "C.O. Test" in the Module Performance Test part.

6.4 Upgrading the Gas Module

When upgrading the gas function, use the material in the upgrade material package to replace the gas module, cables connecting the gas module and main control board, and cables connecting the gas module and parameter face head.

The operation procedure is as follows:

- 1. Separate the front and rear housings of the machine by referring to "Disassembling Front and Rear Housings" in the Disassembly and Repair part.
- 2. Connect the mainstream CO₂ module, microstream CO₂ module, and sidestream CO₂ module in the material package using sheet metals and cables by referring to "Disassembling Gas Module" in the Disassembly and Repair part. The machine supports only single gas module upgrade.



Mainstream CO₂ module composition diagram



An microstream CO₂ module is used.



Sidestream CO2 module composition diagram

3. Fix the gas module to the bracket of the machine rear housing, and connect it with the main control board.



- 4. Reassemble the machine.
- Start the monitor, and choose [Main Menu] > [Maintenance] > [Factory Maintenance] > [Factory Default], select [Support CO₂ Class] to correspond with the upgraded gas module, and restart the machine to validate the configuration.
- 6. Test the upgraded machine by referring to "Gas Module Test" in the Module Performance Test part.

6.5 Function Component Upgrade

Precautions

• When upgrading the WiFi and analog output functions of a standard configuration host, use the board in the material package to replace the board in the monitor, in addition to installing corresponding function components into the monitor.

This monitor supports upgraded WiFi components and recorder components.

Function	PN	Name and Specification	Remarks
Component			
Recorder	115-059807-00	Recorder material package FRU	
WiFi	115-059923-00	Integrated WiFi material package FRU	

This monitor is configured with the WiFi function by building a WiFi through a wireless AP. Only the engineers or personnel specified by Mindray are qualified for the WiFi building and setting as well as performance test.

6.5.1 Upgrading WiFi Function

When upgrading the WiFi function, use the material in the upgrade material package to replace the WiFi module and connected dual-band antennas. The operation procedure is as follows:

- 1. Disassemble the main control board by referring to section "Disassembling Main Control Board" in the Disassembly and Repair part.
- 2. Install the WiFi module to corresponding socket of the main control board by referring to the "Disassembling WiFi Module", and attach the antennas to corresponding rear housing of the machine.



ePM 10/ePM 10A/ePM 10C Host



ePM 12/ePM 12A/ePM 12C Machine



ePM 15/ePM 15A/ePM 15C Machine

- 3. Sort the antennas, install the main control board, and reinstall the machine.
- 4. Start the machine, and test whether the WiFi function is normal.

5. To use the WiFi function, refer to its user manual.

Precautions

• If the antennas are not installed in the right position, the WiFi signal quality will be affected.

6.5.2 Upgrading Recorder Function

When upgrading the recorder function, use the material in the upgrade material package to replace the recorder and cables connecting the recorder and main control board.

The operation procedure is as follows:

- 1. Separate the front and rear housings of the machine by referring to "Disassembling Front and Rear Housings" in the Disassembly and Repair part.
- 2. Install the recorder to the bracket of the rear housing recorder by referring to "Disassembling Recorder".



3. Fix the recorder module to the bracket of the machine rear housing, and connect it with the main control board.



- 4. Reassemble the machine.
- Start the monitor, choose [Main Menu[> [Maintenance] > [Factory Maintenance] > [Factory Default], enable [Recorder], and restart the machine to validate the configuration.
- 6. Start the machine, and test whether the recorder function is normal.

6.6 Software upgrade

This monitor supports upgrade through a PC, network or USB disk. Using one of the methods, you can complete update of the monitor machine and peripheral related firmware.

- This monitor can be upgraded using the network upgrade software (PN: 110-006403-00 PC upgrade tool) of Mindray monitor. The software can directly run on a mobile or desktop PC. You can upgrade the following programs of the monitor by network connection or using crossover cables to connect the monitor with a PC.
- The monitor can also be upgraded using USB disk with specified authorization (containing the USB disk upgrade Bootstrap PN: 110-004854-00). You can upgrade the following programs of the monitor.

Software	PN	Name and Specification	Remarks				
System software	/	ePM system package	1				
package	,	er misystem package	7				
Module rack	/	EPGA chip-write software	1				
software	/		7				
		M51C Mindray monitoring algorithm	ECG				
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			software				
	/	DSP (BF70x) Bootstrap of ePM	DSP Bootstrap				
	/	multi-parameter module					
		IBP function program of ePM	IBP function				
	/	multi-parameter module	upgrade				

Software	PN	Name and Specification	Remarks
			program
			Mindray SpO ₂
	/	Mindray SpO ₂ function program of ePM	function
	7	multi-parameter module	upgrade
			program
			DSP function
	/	DSP (BF70x) function program of ePM	upgrade
		multi-parameter module	program
IBP module	/	Chip-write software of M03B module	/
C.O. module	/	Chip-write software of M03B module	/
M02D module	/	Program software of M02D function	
Power firmware	/	Chip-write software of power firmware	/
Receiver box module	/	Firmware software of receiver box module	/

6.6.1 Network Upgrade Tool 6.6.1.1 Installing Tool Software

1. Click the execution program SystemUpdateTool.exe of the tool software. A page is displayed. Click

to enter the page requiring you to input the Serial Number.

	🌍 MindraySystemUpdateTool 1.1.0 Setup		
	The following information must be entered before installat	tion.	
	Serial Number		
	Nullsoft Install System v3.0a1	Next > Cancel	
2.	Input the Serial Number, and click Next >	to enter the ins	tallation position page of the

Next >

to

program. Select corresponding installation folder, and follow the wizard to click complete the installation.

Appendix: Multilingual Reference Tab	le
Appendix. Multilingual herefellee tab	i C

Language (English)	Language (Chinese)	Remarks
ENGLISH	英语	/
SIM.CHINESE	简体中文	1
FRENCH	法语	/
GERMAN	德语	/
ITALIAN	意大利语	/
POLISH	波兰语	/
SPANISH	西班牙语	/
PORTUGUESE	葡萄牙语	/
RUSSIAN	俄语	1
CZECH	捷克语	1
TURKISH	土耳其语	1
HUNGARIAN	匈牙利语	1
Danish	丹麦语	1
Dutch	荷兰语	1
Finnish	芬兰语	1
Norwegian	挪威语	1
Swedish	瑞典语	/
Romanian	罗马尼亚	1
Serbian	塞尔维亚	1
GREEK	希腊语	1
TRA.CHINESE	繁体中文	1
Japanese	日语	/

6.6.1.2 Connecting the Monitor with a PC

Ensure that at least one NIC is installed on a PC correctly. The monitor is connected with the PC through this NIC.

- 1. Connect the monitor with the PC using a hub.
 - Connect the PC with the hub using a network cable. Insert one end of the network cable to the NIC slot of the PC, and insert the other end of the network cable to the slot of the hub.
 - Connect the monitor with the hub using a network cable. Refer to the PC and hub connection method. The hub has multiple slots, so it can be connected with at least 5 monitors to upgrade them concurrently.
- 2. Modify the IP address of the PC NIC.

Precautions

• Before running the upgrade program, set the IP address to 77.77.1.xx to ensure normal upgrade. The gateway and DNS can be set as necessary. For example, the IP address is set to 77.77.1.13, and the subnet mask is set to 255.255.255.0.

Methods of entering the upgrade mode:

- Connect the monitor with a USB keyboard, and press F4+F5 or * repeatedly during startup to enter the upgrade mode.
- Slide on the touch screen using two or more fingers to enter the upgrade mode during startup.

6.6.1.3 Setting Software Upgrade

Based on the previous configuration requirements, the software upgrade packages of different products need to be set. The setting is performed and managed by the administrator only. Set system software upgrade:

 Download the ePM system software package (stored in the model package directory), run the installed system (network) upgrade tool software, select [Select a New Model Package] >[Precise. Tool], and click [Open] and [Confirm], as shown in the following figure:

1	🎽 Mindray Patien	t Monitor Software Upgrade Tool v 1.0.0	×
	Model Information	1	
	Version	06.16.00	
	Model Description	Support product: Precise	
	Model Package In	formation	
	Model Path		
	Packaging Time	Select A New Model Package	
	Checksum		
		OK Cancel	

🚔 Open							X
60-	🛛 🍑 👻 Computer 👻 Local Disk (E:) 👻 UniformU	pgradeTools 🔻 Releas	2	\sim	👻 🔯 🛛 Search Release		2
Organize	 New folder 					= -	•
<u> </u>	Name 🔺	Date modified	Туре	Size			
	Merak.tool	11/19/2016 12:01	TOOL File	0 KB			
۲ ال							
	File <u>n</u> ame:				▼ files (*.tool,*.mp	ikg)	•
					Upen	Cancel	

2. In the [Product Type Selection] dialog box, select [Precise] in the [Select Product Type].

罩 Product Type Selection		\times
Select Product Type:	Precise	~
OK	Cancel	

The following page is displayed on the PC:

Mindray Patient Monitor Software Upgrade Tool(Merak)								
Operation(O) Setup(S)	View(V) Help(H)				1			
	•	<u>a</u>	4	6	6	J.		
Start	Stop	Create Package	Select Package	Create License	Create Mu	ulti-package	About	
Start Time	MAC Addr	Package Type			Percent (%)	State		
Time	MAC Addr	Package Type	State					

6.6.2 Software Upgrade

6.6.2.1 Upgrading System Software and Internal Plugin Cable

- 1. Enter the system upgrade and download page, and click Select Package
- 2. Browse the prepared system software upgrade package files, and verify whether the download content, including item, checksum, version, and note is correct, and click [**Confirm**].

影

The [Start] hot key is lit up in the main menu.

Select Package		×
Select Package	C:\Users\cms\Desktop\system(svn106461)+kernal(020700	004)+ Browse
Creation Time	2018-10-09 09:00:28	
Checksum	E8 68 D7 6D	
Item	Lhecksum Version	Note
LINUX KERNEL	56 10 7D 73 02.07.00.04	
Touchscreen binar	ry 36 69 48 89	
SPI Bridge	CB EB 20 01	
System program	B7 CB FB 47 2.9.0.1	Precise
Module software	DE A5 9E AF Powe	erBoard-function
	Ok	Cancel

3. Confirm that the network cable is connected correctly and the monitor is stopped, and click the [**Start**] hot key to download the software.

6.6.2.2 Upgrading Module Software

Upgrade the module program files by referring to the System Software Upgrade Method. After they are upgraded, click [**Stop**] in the upgrade menu, remove the network cable, and stop and restart the monitor.

For details about the network program upgrade method, refer to the user manual and help of the Mindray monitor network upgrade software or consult the after-sales service personnel of Mindray.

6.6.3 Upgrade by USB Disk

6.6.3.1 Preparing USB Disk Upgrade Directory Structure

Prepare the following required tool:

- USB disk: 2 GB or larger ordinary FAT USB disk, for example, Kingston or Netac.
- 1. Create UPGRADE_AMP\Precise in the root directory of the USB disk.
- 2. Copy the upgrade Bootstrap Precise-Installer.pkg to the UPGRADE_AMP\Precise directory.
- 3. Copy the upgrade file (PKG or MPKG) to the UPGRADE_AMP\Precise directory.

6.6.3.2 Inserting the USB Disk into the USB Port of the Monitor

Insert the prepared USB disk into any USB port of the main control board.

6.6.3.3 Triggering Upgrade Mode

- Method 1. Slide on the touch screen using two or more fingers to enter the upgrade mode during startup.
- Method 2: Connect the monitor with a USB keyboard, and press F2+F3, F4+F5, or * repeatedly during startup to enter the upgrade mode.

6.6.3.4 Selecting Upgrade File





- If there is only one upgrade file, it is selected by default. If there are multiple upgrade files, they are displayed in two columns. A maximum of 16 upgrade files can be displayed, which should not be exceeded. Click the up/down/right/left keys to switch to required upgrade file.
- Click on the touch screen to select an upgrade file downward, or press \checkmark on the keyboard to select one.
- Click UP t on the touch screen to select an upgrade file upward, or press ↑ on the keyboard to select one.
- Click LEFT ★ on the touch screen to select an upgrade file rightward, or press ← on the keyboard to select one.
- Click Click Click Click on the touch screen to select an upgrade file rightward, or press → on the keyboard to select one.
- Click Chter on the touch screen to confirm the selected upgrade file, or press [Enter] on the keyboard to confirm the selection.

6.6.3.5 Completing Upgrade Process

If the following page is displayed, the upgrade is complete. Stop and restart the monitor to validate the upgrade.



Attention

- Before the upgrade, disconnect the monitor from a patient and save important data in the monitor.
- During upgrade of the Bootstrap and FPGA, do not stop or power off the monitor. Otherwise, the device will be corrupted.
- Program upgrade can be performed by professional maintenance personnel only. Failure to avoid any maintenance warning may cause slight personal injury, product fault, or property loss.

Precautions

- After the Bootstrap is upgraded, re-upgrade the system program or other programs to ensure their compatibility.
- Before upgrade, ensure that the version of the upgrade package is the required one. To obtain the latest upgrade package, contact the product after-sales service department of this company.

FOR YOUR NOTES

7 Maintenance and Disassembly

7.1 Tool

Before disassembling or replacing parts, the following tools may be used:

- Small Phillips screwdriver
- Phillips screwdriver
- Tweezers
- Needle-nose pliers
- Cutting pliers

7.2 Preparations

Before disassembling the monitor, make the following preparations:

Stop monitoring the patient, power off the monitor, and disconnect all attachments and external devices.

/ Warnings

- Before disassembly, eliminate static electricity. When disassembling components with electrostatic sensitive mark, wear ESD wristband or gloves to avoid damage to the components.
- During reinstallation, connect and place cables properly to avoid crushing the cables, which may cause a short circuit.
- During reinstallation, select screws with proper model. If you forcibly tighten improper screws, the device may be damaged. And the screws or components may become loose during use, which causes unpredictable product damage or personal injury.
- Disassemble the device based on the disassembly sequence. Otherwise, an irreversible damage may be caused to the device.
- Loosened parts should be categorized and placed properly so that they can be used for reinstallation and will not be dropped, polluted or lost.
- During reinstallation, install the components prior to the host, and connect and place cables and hoses properly.
- The machine has waterproof requirement. During reinstallation, ensure that waterproof case and other waterproof materials are assembled properly.

7.3 ePM 15/ePM 15A/ePM 15C Host Disassembly

7.3.1 Disassembling Front/rear Housing Components of Host

1. Use a Phillips screwdriver to loosen four M4X8 screws.



2. Open the front/rear housings, and remove the display screen connection cable and keypad connection cable.





Note: During reassembly, close the front/rear covers and pull the cables upward using a hand.



7.3.2 Disassembling Keypad and Switch Board

- 1. Remove the keypad connection cable, loosen the three ST3.3X8 screws shown in the figure, and take the keypad out.
- 2. Remove the switch board connection cable, loosen the two ST3.3X8 screws shown in the figure, and take the switch board out.



7.3.3 Disassembling Display Screen and Alarm Indicator

1. Remove the five cable ties from cables.



- 2. Remove the cables connecting with the touch screen, display screen, and alarm indicator.
- 3. Loosen the sticker connecting the touch screen PFC with the sheet metal and the cable stuck to the sheet metal.



4. Loosen the ten ST3.3X8 screws shown in the figure, and remove the display screen component.



5. Loosen the two M3X6 screws, and take the alarm indicator board out.



- 6. Loosen the two M3X6 screws with the mark "b" on the right top of the sheet metal.
- 7. Loosen the two M2X4 screws at the left side of the sheet metal.
- 8. Loosen the two M2X4 screws at the left side of the sheet metal, and take the display screen out.



9. Loosen the sticker connecting the touch screen with the front housing, and tilt and take the touch screen out.



Note 1. During reassembly of the touch screen rubber, follow the requirements below to perform assembly.



Note 2. During reassembly of the touch screen, follow the requirements below to perform assembly.







7.3.4 Disassembling WiFi and Parameter Panel

1. Remove cables.



Note. During reassembly, follow the requirements below to perform binding and fixing.



2. When WiFi is configured, disassemble the WiFi module.

A. Take the WiFi module out.

B. Remove the WiFi cable.

C. Loosen the three M2X4 screws fixing the WiFi module and WiFi load board, and take the WiFi module and WiFi load board out.



Note. During reassembly of the WiFi module, follow the requirements below to perform assembly.



3. When mainstream CO₂ is configured, disassemble the CO₂ module.

Remove the cables pointed by the arrows in the following figure, loosen three ST3.3X8 cross pan head tapping screws using the screwdriver, and take the module out.



4. When microstream CO₂ is configured, disassemble the CO₂ module.

A. Loosen three ST3.3X8 cross pan head tapping screws using the screwdriver, and take the module out.B. Remove the connection between the microstream module exhaust hose and panel exhaust hose.



Note. During reassembly of the microstream CO_2 module, follow the requirements below to perform assembly.



5. When sidestream CO₂ is configured, disassemble the CO₂ module.

A. Loosen the cable ties shown in the following figure, take the three ST3.3X8 cross pan head tapping screws out, and take the sidestream CO_2 module out.



B. Take the exhaust hose connected with the sidestream CO_2 module out.



B. Loosen the cable connection, and take the module out.



C. Disassemble the two PT2.0X6 screws on the panel water tank, and take the water tank out.



Note. During reassembly of the sidestream CO₂ module, follow the requirements below to perform assembly.



- 6. Remove the panel cable from the mainboard, and remove the NIBP exhaust hose.
- 7. Loosen two ST3.3X8 cross pan head tapping screws using the screwdriver, and take the panel fixing pin and panel component out.



- 8. When a mainstream module is configured, disassemble the mainstream panel port.
- A. Use pliers or tweezers to jack the fastener on the panel, and take the interface board out.
- B. Rotate the cable anticlockwise, and remove the CO₂ cable.
- C. Remove the plug from the panel.
- D. Loosen the M5 screw from the upper right corner of the panel, and take the nozzle out.



Note 1. During reassembly, the interface board of the mainstream panel should be in the following direction.



Note 2. During reassembly of the cable to the interface board, tighten it firmly anticlockwise.



9. When an microstream module is configured, disassemble the microstream water tank.

A. Use pliers or tweezers to jack the fastener on the panel, and take the microstream water tank base out.

B. Loosen the M5 screw from the upper right corner of the panel, and take the exhaust nozzle out.

C. Use tweezers to loosen the fastener on the water tank base, and take the microstream connector out.



10. When a sidestream module is configured, disassemble the sidestream water tank.

A. Remove the cable or hose from the water tank.

B. Remove the exhaust hose from the panel, loosen the M4 screw on the exhaust nozzle, and take the nozzle out.

C. Loosen the M5 screw/M4 nut on the panel, and take the exhaust nozzle out.

D. Loosen the four PT2.0X6 screws on the panel, and take the bracket of the sidestream water tank out.



- 11. Disassemble the panel cable.
- A. According to the figure, rotate different parameter cables anticlockwise, and remove them.
- B. Take the arrival reminding shrapnel out.


Note. During reassembly, follow the requirements below to perform cable assembling.



12. Disassemble the NIBP nozzle: Rotate the NIBP nozzle anticlockwise, and disassemble the NIBP nozzle.



7.3.5 Disassembling Gas Module

1. When mainstream CO₂ is configured:

A. Loosen the three M3X6 screws in the figure, and take the mainstream CO_2 isolation power board out.

B. Remove the cable connecting the mainstream isolation power board with the mainboard.



- 2. When microstream CO₂ is configured:
- A. Remove the cable connecting the microstream CO₂ module with the adapter.
- B. Remove the cable connecting the mainboard.
- C. Loosen the four M3X6 screws in the figure, and take the microstream CO_2 module out.
- D. Loosen the three M3X6 screws, and take the adapter out.



3. When sidestream CO₂ is configured:

A. Loosen the two M3X6 screws in the figure, and take the sidestream gas module out.



- B. Take the air filter and short-circuited hose out.
- C. Loosen the three M2.5X4 countersunk screws, and take the cover out.
- D. Take the silicone case out.



7.3.6 Disassembling Recorder/Recorder Bracket

1. When the recorder is configured, disassemble the recorder:

A. Loosen the two M3X6 screws of the recorder, loosen the two fasteners of the recorder, and take the recorder out.

B. Remove the cable connecting the two sockets of the recorder.



2. Take the recorder bracket out: Loosen the two ST3.3X8 screws on the recorder bracket, and take the recorder bracket out.



7.3.7 Disassembling Main Bracket Component

- 1. Loosen the battery cover.
- 2. Loosen the five ST3.3X8 screws shown in the figure, and remove the main bracket component.



Note 1. Before reassembling the main bracket component, insert the connection belt of the battery cover to the locating post of the rear housing shown in the following figure.



Note 2. Before closing the battery cover, switch the battery to the vertical position shown in the figure.



7.3.8 Disassembling Speaker

1. Loosen the two ST3.3X8 screws of the speaker component shown in the figure, and take the speaker component out.



Note. During reassembly of the speaker, follow the requirements below to perform assembly.



7.3.9 Disassembling Rear Alarm Indicator (Configured)

- 1. Loosen the eight ST3.3X8 screws on the cover component shown in the figure, and take the top cover component out.
- 2. Loosen the one ST3.3X8 screw on the rear alarm indicator shown in the figure, and take the rear alarm indicator component out.

7.3.10 Disassembling Power Module

- 1. Remove the AC input cable, and remove the cable connecting the power module with mainboard out.
- 2. Loosen the four M3X6 screws of the power module, and take the power module out.



7.3.11 Disassembling SpO₂ Module (When Nellcor/Massimo SpO₂ Is Configured)

1. When Nellcor SpO₂ is configured:

Loosen the one M2X4 screw on the Nellcor SpO_2 , and take the Nellcor SpO_2 board out.



2. When Massimo SpO₂ is configured:

Loosen the two M2X4 screws on the Massimo SpO₂, and take the Massimo SpO₂ board and insulation sheet out.



7.3.12 Disassembling C.O. Board (Configured)

- 1. Remove the cable connecting the C.O. board with the mainboard.
- 2. Loosen the two M3X6 screws on the Massimo SpO₂, and take the C.O. board and insulation sheet out.



7.3.13 Disassembling Mainboard

- 1. Remove the pump/valve connection cables from the mainboard.
- 2. Take the two interfaces of the NIBP hose from the mainboard sensor.



3. Remove the connection cable of the battery adapter from the mainboard.



4. When two batteries are configured, remove the connection cable of the 2# battery adapter from the mainboard in the figure.



- 5. Loosen the six M3X6 screws from the main bracket in the figure.
- 6. Loosen the two screws on the rear of the main bracket sheet metal, and take the mainboard out.



7. As shown in the following figure, loosen the nuts or screws on the rear of the mainboard, and take the studs out.



7.3.14 Disassembling Power Adapter and NIBP Pump/Valve

- 1. As shown in the following figure, loosen the two ST3.3X8 screws, and take the 1# battery adapter out.
- 2. When 2# battery adapter is configured, loosen the two ST3.3X8 screws, and take the 2# battery adapter out.



Note. During reassembly of 2# battery adapter, follow the requirements below to perform cable assembly.



- 3. Loosen the hose connecting the NIBP pump/valve.
- 4. Loosen the fastener fixing the NIBP valve, and take the NIBP valve out.
- 5. Loosen the two binding straps fixing the NIBP pump, and take the NIBP pump out.



Note: During reassembly, ensure that the hose is correctly connected with the quick/slow release valve.



7.4 ePM 12/ePM 12A/ePM 12C Host Disassembly

7.4.1 Disassembling Battery Box (Configured)

1. As shown in the following figure, use a Phillips screwdriver to loosen four M3X6 screws, and separate the battery adapter cable from the extension cable.





- 2. Remove the battery box component.
- 3. Pull the connection belt of the battery cover outward to open the battery cover, and put the screwdriver near the shaft of the battery cover to level the battery cover, and take the battery cover out.



4. Use the Phillips screwdriver to loosen two M3X6 screws, take the bracket fixing the battery adapter out, use the Phillips screwdriver to loosen two M3X6 screws, and take the battery adapter out.



7.4.2 Disassembling Front/rear Housing Components of Host

1. Use a long Phillips screwdriver to loosen four M4X8 screws.



2. Open the front/rear housings, and remove the display screen connection cable and keypad connection cable.





Note: During reassembly, close the front/rear covers and pull the cables upward using a hand.



7.4.3 Disassembling Front Housing Component

- 1. Remove the cable connecting the keypad.
- 2. Loosen the five ST3.3X8 screws shown in the figure, and take the keypad out.



7.4.4 Disassembling Display Screen and Alarm Indicator

1. Remove the four cable ties from the cables.



- 2. Remove the cables connecting with the touch screen, display screen, and alarm indicator.
- 3. Loosen the sticker connecting the touch screen PFC with the sheet metal and the cable stuck to the sheet metal.



4. Loosen the eight ST3.3X8 screws shown in the figure, and remove the display screen component.



5. Loosen the two M3X6 screws, and take the alarm indicator board out.



- 6. Loosen the two M3X6 screws with the mark "b" on the right top of the sheet metal.
- 7. Loosen the two M2X4 screws at the left side of the sheet metal.
- 8. Loosen the two M2X4 screws at the left side of the sheet metal, and take the display screen out.



9. Loosen the sticker connecting the touch screen with the front housing, and tilt and take the touch screen out.



10. Use cutting pliers to cut the binding straps fixing the display screen connection cable and keypad connection cable, remove the cable from the keypad socket, and remove the display screen connection cable and keypad connection cable.



Note 1. During reassembly of the touch screen rubber, follow the requirements below to perform assembly.



Note 2. During reassembly of the touch screen, follow the requirements below to perform assembly.







7.4.5 Disassembling WiFi and Parameter Panel

1. Remove cables.



Note. During reassembly, follow the requirements below to perform binding and fixing.



2. When WiFi is configured, disassemble the WiFi module.

A. Take the WiFi module out.

B. Remove the WiFi cable.

C. Loosen the three M2X4 screws fixing the WiFi module and WiFi load board, and take the WiFi module and WiFi load board out.



Note. During reassembly of the WiFi module, follow the requirements below to perform assembly.



3. When mainstream CO₂ is configured, disassemble the CO₂ module.

Remove the cables pointed by the arrows in the following figure, loosen three ST3.3X8 cross pan head tapping screws using the screwdriver, and take the module out.



4. When microstream CO₂ is configured, disassemble the CO₂ module.

Loosen three ST3.3X8 cross pan head tapping screws using the screwdriver, and take the module out.
Remove the connection between the microstream module exhaust hose and panel exhaust hose.



Note. During reassembly of the microstream CO_2 module, follow the requirements below to perform assembly.



5. When sidestream CO_2 is configured, disassemble the CO_2 module.

A. Loosen three ST3.3X8 cross pan head tapping screws using the screwdriver, and loosen the hose of the sidestream CO₂.



B. Loosen the cable connection, and take the module out.



C. Disassemble the two PT2.0X6 screws on the panel water tank, and take the water tank out.



Note. During reassembly of the sidestream CO₂ module, follow the requirements below to perform assembly.



- 6. Remove the panel cable from the mainboard, and remove the NIBP exhaust hose.
- 7. Loosen two ST3.3X8 cross pan head tapping screws using the screwdriver, and take the panel fixing pin and panel component out.



- 8. When a mainstream module is configured, disassemble the mainstream panel port.
- A. Use pliers or tweezers to jack the fastener on the panel, and take the interface board out.
- B. Rotate the cable anticlockwise, and remove the CO₂ cable.
- C. Remove the plug from the panel.
- D. Loosen the M5 screw from the upper right corner of the panel, and take the nozzle out.



Note 1. During reassembly, the interface board of the mainstream panel should be in the following direction.



Note 2. During reassembly of the cable to the interface board, tighten it firmly anticlockwise.



9. When an microstream module is configured, disassemble the microstream water tank.

A. Use pliers or tweezers to jack the fastener on the panel, and take the microstream water tank base out.

- B. Loosen the M5 screw from the upper right corner of the panel, and take the exhaust nozzle out.
- C. Use tweezers to loosen the fastener on the water tank base, and take the microstream connector out.



10. When a sidestream module is configured, disassemble the sidestream water tank.

A. Remove the cable or hose from the water tank.

B. Remove the exhaust hose from the panel, loosen the M4 screw on the exhaust nozzle, and take the nozzle out.

C. Loosen the M5 screw/M4 nut on the panel, and take the exhaust nozzle out.

D. Loosen the four PT2.0X6 screws on the panel, and take the bracket of the sidestream water tank out.



- 11. Disassemble the panel cable.
- A. According to the figure, rotate different parameter cables anticlockwise, and remove them.
- B. Take the arrival reminding shrapnel out.



Note. During reassembly, follow the requirements below to perform cable assembling.



12. Disassemble the NIBP nozzle: Rotate the NIBP nozzle anticlockwise, and disassemble the NIBP nozzle.



7.4.6 Disassembling Gas Module

1. When mainstream CO₂ is configured:

A. Loosen the three M3X6 screws in the figure, and take the mainstream CO_2 isolation power board out.

B. Remove the cable connecting the mainstream isolation power board with the mainboard.



- 2. When microstream CO₂ is configured:
- A. Remove the cable connecting the microstream CO_2 module with the adapter.
- B. Remove the cable connecting the mainboard.
- C. Loosen the four M3X6 screws in the figure, and take the microstream CO_2 module out.
- D. Loosen the three M3X6 screws, and take the adapter out.



3. When sidestream CO₂ is configured:

A. Loosen the two M3X6 screws in the figure, and take the sidestream gas module out.



B. Take the air filter and short-circuited hose out.

C. Loosen the three M2.5X4 countersunk screws, and take the cover out.

D. Take the silicone case out.









7.4.7 Disassembling Recorder/Recorder Bracket

1. When the recorder is configured, disassemble the recorder:

A. Loosen the two M3X6 screws of the recorder, loosen the two fasteners of the recorder, and take the recorder out.

B. Remove the cable connecting the two sockets of the recorder.



2. Take the recorder bracket out: Loosen the two ST3.3X8 screws on the recorder bracket, and take the recorder bracket out.



7.4.8 Disassembling Main Bracket Component

1. When the battery box is configured, remove the cable connecting the battery box.



2. Loosen the battery cover.

3. Loosen the five ST3.3X8 screws shown in the figure, and remove the main bracket component.



Note 1. Before reassembling the main bracket component, insert the connection belt of the battery cover to the locating post of the rear housing shown in the following figure.



Note 2. Before closing the battery cover, switch the battery to the vertical position shown in the figure.



After the battery is switched to the vertical position, close the battery cover.

7.4.9 Disassembling Speaker

1. Loosen the two ST3.3X8 screws of the speaker component shown in the figure, and take the speaker component out.





7.4.10 Disassembling Rear Alarm Indicator (Configured)

- 1. Loosen the eight ST3.3X8 screws on the cover component shown in the figure, and take the top cover component out.
- 2. Loosen the one ST3.3X8 screw on the rear alarm indicator shown in the figure, and take the rear alarm indicator component out.

7.4.11 Disassembling Power Module

- 1. Remove the AC input cable, and remove the cable connecting the power module with mainboard out.
- 2. Loosen the four M3X6 screws of the power module, and take the power module out.





7.4.12 Disassembling SpO₂ Module (When Nellcor/Massimo SpO₂ Is Configured)

1. When Nellcor SpO₂ is configured:

Loosen the one M2X4 screw on the Nellcor SpO_{2r} and take the Nellcor SpO_2 board out.



2. When Massimo SpO₂ is configured:

Loosen the two M2X4 screws on the Massimo SpO $_2$, and take the Massimo SpO $_2$ board and insulation sheet out.



7.4.13 Disassembling C.O. Board (Configured)

- 1. Remove the cable connecting the C.O. board with the mainboard.
- 2. Loosen the two M3X6 screws on the Massimo SpO $_2$, and take the C.O. board and insulation sheet out.



7.4.14 Disassembling Mainboard

- 1. Remove the pump/valve connection cables from the mainboard.
- 2. Take the two interfaces of the NIBP hose from the mainboard sensor.



3. Remove the connection cable of the battery adapter from the mainboard.



- 4. Loosen the six M3X6 screws from the main bracket in the figure.
- 5. Loosen the two screws on the rear of the main bracket sheet metal, and take the mainboard out.



6. As shown in the following figure, loosen the nuts or screws on the rear of the mainboard, and take the studs out.



7.4.15 Disassembling Power Adapter and NIBP Pump/Valve

1. As shown in the following figure, loosen the two ST3.3X8 screws, and take the battery adapter out.



- 2. Loosen the hose connecting the NIBP pump/valve.
- 3. Loosen the fastener fixing the NIBP valve, and take the NIBP valve out.
- 4. Loosen the two binding straps fixing the NIBP pump, and take the NIBP pump out.



Note: During reassembly, ensure that the hose is correctly connected with the quick/slow release valve.



7.5 ePM 10/ePM 10A/ePM 10C Host Disassembly

7.5.1 Disassembling Front/rear Housing Components of Host

1. Use a Phillips screwdriver to loosen four M4X8 screws.



2. Open the front/rear housings, and remove the display screen connection cable and keypad connection cable.





Note: During reassembly, close the front/rear covers and pull the cables upward using a hand.


7.5.2 Disassembling Keypad

- 1. Remove the cable connecting the keypad.
- 2. Loosen the five ST3.3X8 screws shown in the figure, and take the keypad out.



7.5.3 Disassembling Display Screen and Alarm Indicator

1. Remove the cables connecting with the touch screen, display screen, and alarm indicator.



- 2. Remove the five cable ties from cables, and loosen the cables stuck to the sheet metal.
- 3. Loosen the sticker connecting the touch screen PFC with the sheet metal.



4. Loosen the eight ST3.3X8 screws shown in the figure, and remove the display screen component.



5. Loosen the two M3X6 screws, and take the alarm indicator board out.



- 6. Loosen the two M3X6 screws with the mark "b" on the right top of the sheet metal.
- 7. Loosen the two M2X4 screws at the left side of the sheet metal.
- 8. Loosen the two M2X4 screws at the left side of the sheet metal, and take the display screen out.



9. Loosen the sticker connecting the touch screen with the front housing, and tilt and take the touch screen out.



Note 1. During reassembly of the touch screen rubber, follow the requirements below to perform assembly.



Note 2. During reassembly of the touch screen, follow the requirements below to perform assembly.







7.5.4 Disassembling WiFi and Parameter Panel

1. Remove cables.



Note. During reassembly, follow the requirements below to perform binding and fixing.



2. When WiFi is configured, disassemble the WiFi module.

A. Take the WiFi module out.

B. Remove the WiFi cable.

C. Loosen the three M2X4 screws fixing the WiFi module and WiFi load board, and take the WiFi module and WiFi load board out.



Note. During reassembly of the WiFi module, follow the requirements below to perform assembly.



3. When mainstream CO₂ is configured, disassemble the CO₂ module.

Remove the cables pointed by the arrows in the following figure, loosen three ST3.3X8 cross pan head tapping screws using the screwdriver, and take the module out.



4. When microstream CO₂ is configured, disassemble the CO₂ module.

A. Loosen three ST3.3X8 cross pan head tapping screws using the screwdriver, and take the module out.B. Remove the connection between the microstream module exhaust hose and panel exhaust hose.





Note. During reassembly of the microstream CO_2 module, follow the requirements below to perform assembly.



5. When sidestream CO_2 is configured, disassemble the CO_2 module.

A. Loosen three ST3.3X8 cross pan head tapping screws using the screwdriver, and loosen the hose of the sidestream CO₂.





B. Loosen the cable connection, and take the module out.



C. Disassemble the two PT2.0X6 screws on the panel water tank, and take the water tank out.



Note. During reassembly of the sidestream CO2 module, follow the requirements below to perform



- 6. Remove the panel cable from the mainboard, and remove the NIBP exhaust hose.
- 7. Loosen two ST3.3X8 cross pan head tapping screws using the screwdriver, and take the panel fixing pin and panel component out.



- 8. When a mainstream module is configured, disassemble the mainstream panel port.
- A. Use pliers or tweezers to jack the fastener on the panel, and take the interface board out.
- B. Rotate the cable anticlockwise, and remove the CO_2 cable.
- C. Remove the plug from the panel.
- D. Loosen the M5 screw from the upper right corner of the panel, and take the nozzle out.



Note 1. During reassembly, the interface board of the mainstream panel should be in the following direction.



Note 2. During reassembly of the cable to the interface board, tighten it firmly anticlockwise.



9. When an microstream module is configured, disassemble the microstream water tank.

A. Use pliers or tweezers to jack the fastener on the panel, and take the microstream water tank base out.B. Loosen the M5 screw from the upper right corner of the panel, and take the exhaust nozzle out.C. Use tweezers to loosen the fastener on the water tank base, and take the microstream connector out.





- 10. When a sidestream module is configured, disassemble the sidestream water tank.
- A. Remove the cable or hose from the water tank.

B. Remove the exhaust hose from the panel, loosen the M4 screw on the exhaust nozzle, and take the nozzle out.

- C. Loosen the M5 screw/M4 nut on the panel, and take the exhaust nozzle out.
- D. Loosen the four PT2.0X6 screws on the panel, and take the bracket of the sidestream water tank out.



11. Disassemble the panel cable.

A. According to the figure, rotate different parameter cables anticlockwise, and remove them.

B. Take the arrival reminding shrapnel out.





Note. During reassembly, follow the requirements below to perform cable assembling.



12. Disassemble the NIBP nozzle: Rotate the NIBP nozzle anticlockwise, and disassemble the NIBP nozzle.



7.5.5 Disassembling Gas Module

- 1. When mainstream CO₂ is configured:
- A. Loosen the three M3X6 screws in the figure, and take the mainstream CO₂ isolation power board out.
- B. Remove the cable connecting the mainstream isolation power board with the mainboard.



- 2. When microstream CO₂ is configured:
- A. Remove the cable connecting the microstream CO_2 module with the adapter.
- B. Remove the cable connecting the mainboard.
- C. Loosen the four M3X6 screws in the figure, and take the microstream CO_2 module out.
- D. Loosen the three M3X6 screws, and take the adapter out.



3. When sidestream CO₂ is configured:

A. Loosen the two M3X6 screws in the figure, and take the sidestream gas module out.



B. Take the air filter and short-circuited hose out.

C. Loosen the three M2.5X4 countersunk screws, and take the cover out.

D. Take the silicone case out.



7.5.6 Disassembling Recorder/Recorder Bracket

1. When the recorder is configured, disassemble the recorder:

A. Loosen the two M3X6 screws of the recorder, loosen the two fasteners of the recorder, and take the recorder out.

B. Remove the cable connecting the two sockets of the recorder.



2. Take the recorder bracket out: Loosen the two ST3.3X8 screws on the recorder bracket, and take the recorder bracket out.



7.5.7 Disassembling Main Bracket Component

- 1. Loosen the battery cover.
- 2. Loosen the five ST3.3X8 screws shown in the figure, and remove the main bracket component.





Note 1. Before reassembling the main bracket component, insert the connection belt of the battery cover to the locating post of the rear housing shown in the following figure.



Note 2. Before closing the battery cover, switch the battery to the vertical position shown in the figure.



7.5.8 Disassembling Speaker

1. Loosen the two ST3.3X8 screws of the speaker component shown in the figure, and take the speaker component out.





7.5.9 Disassembling Rear Alarm Indicator (Configured)

- 1. Loosen the six ST3.3X8 screws on the cover component shown in the figure, and take the top cover component out.
- 2. Loosen the one ST3.3X8 screw on the rear alarm indicator shown in the figure, and take the rear alarm indicator component out.

7.5.10 Disassembling Power Module

- 1. Remove the AC input cable, and remove the cable connecting the power module with mainboard out.
- 2. Loosen the four M3X6 screws of the power module, and take the power module out.



7.5.11 Disassembling SpO₂ Module (When Nellcor/Massimo SpO₂ Is Configured)

1. When Nellcor SpO₂ is configured:

Loosen the one M2X4 screw on the Nellcor SpO $_2$, and take the Nellcor SpO $_2$ board out.



2. When Massimo SpO₂ is configured:

Loosen the two M2X4 screws on the Massimo SpO₂, and take the Massimo SpO₂ board and insulation sheet out.



7.5.12 Disassembling C.O. Board (Configured)

- 1. Remove the cable connecting the C.O. board with the mainboard.
- 2. Loosen the two M3X6 screws on the Massimo SpO $_2$, and take the C.O. board and insulation sheet out.



7.5.13 Disassembling Mainboard

- 1. Remove the pump/valve connection cables from the mainboard.
- 2. Take the two interfaces of the NIBP hose from the mainboard sensor.



3. Remove the connection cable of the battery adapter from the mainboard.



- 4. Loosen the six M3X6 screws from the main bracket in the figure.
- 5. Loosen the two screws on the rear of the main bracket sheet metal, and take the mainboard out.



6. As shown in the following figure, loosen the nuts or screws on the rear of the mainboard, and take the studs out.



7.5.14 Disassembling Power Adapter and NIBP Pump/Valve

1. As shown in the following figure, loosen the two ST3.3X8 screws, and take the battery adapter out.



- 2. Loosen the hose connecting the NIBP pump/valve.
- 3. Loosen the fastener fixing the NIBP valve, and take the NIBP valve out.
- 4. Loosen the two binding straps fixing the NIBP pump, and take the NIBP pump out.



Note: During reassembly, ensure that the hose is correctly connected with the quick/slow release valve.



7.6 ePM T10 Vehicle-mounted Charger Base Disassembly

7.6.1 Disassembling Transfer Base

1. Push the release handle towards the arrow direction marked on the handle, and take the adapter component out upward.



2. Loosen the four M4X12 combination screws using the Phillips screwdriver, and separate the transfer base from the installation base.



3. Use tweezers to peel the waterproof tape off. In case of reinstallation, use a new waterproof tape.



4. Remove the output cable from the PCBA base, and take the cable out. Remove the input cable terminal from the PCBA, loosen the four M3X8 screws, and take the power input connector out from the base.



5. Loose the six M3X6 screws using a Phillips screwdriver, remove the socket of the LED indicator from the PCBA, and take the PCBA out.



6. Use tweezers to take the conductive rubber of the PCBA from the cooling block, and pull the LED indicator out from the base hole.



7.6.2 Disassembling Installation Base

1. Use tweezers to peel the sealing tape off, and loosen the three M3X6 pan head screws with pad using the screwdriver.



2. Take the release board and release handle out upward, loosen the two M3X6 countersunk screws using the screwdriver, and separate the release board from the release handle.



3. Take the two locating springs out, and use the tweezers to remove the two pads.



4. Loosen the twelve M3X6 countersunk screws using the screwdriver, and remove the four locking block press boards.



5. Take the four pressing springs and slide blocks out upward.



7.6.3 Disassembling Adapter Component

Use tweezers to take the four foot pads out from the adapter component, and loosen the four M4X10 screws using the screwdriver, and separate the installation block from the adapter.





FOR YOUR NOTES

This part lists the exploded view and PN of the monitor, auxiliary plugin box and parameter module, so that maintenance personnel can recognize the names of different parts when they disassemble or replace the parts.

8.1 ePM 15/ePM 15A/ePM 15C Parts

8.1.1 System Structure

Exploded View



No.	Name and Specification	Quantity	PN
1	Front Housing Assembly	1	115-060001-00
2	Repair kit for integrated 15-inch rear housing component (rear alarm light is not supported)	1	115-059949-00
2	Repair kit for integrated 15-inch rear housing component (rear alarm light is supported)	1	115-059950-00

8.1.2 Front Housing

Exploded View



No.	Name and Specification	Quantity	PN
1	TFT displayer assembly (AUO 15-inch screen)	1	115-059826-00
2	Repair kit for front housing and touch screen	1	115-060001-00
3	9202 C15 function keypad PCBA	1	115-059768-00

8.1.3 Rear Housing

Exploded View



No.	Name and Specification	Quantity	PN
1	TR6F recorder	1	115-059807-00
2	Integrated 15-inch standard configuration mainboard FRU (with software)	1	115-059942-00
	Integrated 15-inch full configuration 3/5-lead MR SpO ₂ 1 mainboard FRU (with software)		115-059943-00
	Integrated 15-inch full configuration 3/5-lead OEM SpO ₂ mainboard FRU (with software)	1	115-059944-00
	Integrated 15-inch full configuration 12-lead MR SpO ₂ mainboard FRU (with software)	1	115-059945-00
	Integrated 15-inch full configuration 12-lead OEM SpO ₂ mainboard FRU (with software)	1	115-059946-00
3	Mindray sidestream CO ₂ module kit	1	115-059955-00
4	Speaker 2W40hm 500Hz	1	115-059960-00
5	Repair kit for integrated panel component (CO2 is not supported)	1	115-059951-00
	Repair kit for integrated panel component (CO ₂ is supported)	1	115-059952-00

8.2 ePM 12/ePM 12A/ePM 12C Parts

8.2.1 System Structure

Exploded View



No.	Name and Specification	Quantity	PN
1	Front Housing Assembly	1	115-059976-00
2	Repair kit for integrated 12-inch rear housing component (rear alarm light is not supported)	1	115-059972-00
2	Repair kit for integrated 12-inch rear housing component (rear alarm light is supported)	1	115-059973-00

8.2.2 Front Housing

Exploded View



No.	Name and Specification	Quantity	PN
1	Front Housing repair kit	1	115-059976-00
2	TFT display assembly (Innolux 12-inch screen)	1	115-059827-00

8.2.3 Rear Housing

Exploded View



No.	Name and Specification	Quantity	PN
	Repair kit for integrated panel component (CO ₂ is not		115 050051 00
1	supported)		112-029921-00
	Repair kit for integrated panel component (CO ₂ is supported)		115-059952-00
2	CO ₂ module (M02D)	1	115-059974-00
3	Integrated 12-inch standard configuration mainboard FRU	1	115-050065-00
	(with software)		115-059965-00
	Integrated 12-inch full configuration $3/5$ -lead MR SpO ₂	1	115-050066-00
	mainboard FRU (with software)		115-059900-00
	Integrated 12-inch full configuration $3/5$ -lead OEM SpO ₂	1	115 050067 00
	mainboard FRU (with software)		113-039907-00
	Integrated 12-inch full configuration 12-lead MR SpO $_{\rm 2}$	1	115 050069 00
	mainboard FRU (with software)		113-03996-00
	Integrated 12-inch full configuration 12-lead OEM SpO ₂	1	115 050060 00
	mainboard FRU (with software)		112-029909-00
4	Speaker repair kit	1	115-059830-00
5	TR6F recorder	1	115-059807-00
6	9202 rear alarm indicator PCBA	1	115-059941-00

8.3 ePM 10/ePM 10A/ePM 10C Parts

8.3.1 System Structure Exploded View



No.	Name and Specification	Quantity	PN
1	Repair kit for front housing and touch screen (DC-in is not supported)	1	115-059997-00
	Repair kit for front housing and touch screen (DC-in is supported)	1	115-059998-00
2	Repair kit for integrated 10-inch rear housing component (rear alarm light is not supported)	1	115-059993-00
	Repair kit for integrated 10-inch rear housing component (rear alarm light is supported)	1	115-059994-00

8.3.2 Front Housing

Exploded View



No.	Name and Specification	Quantity	PN
1	Repair kit for front housing and touch screen	1	115-059997-00
2	TFT display assembly (Innolux 10-inch screen)	1	115-059828-00

8.3.3 Rear Housing

Exploded View



No.	Name and Specification	Quantity	PN
1	Repair kit for front alarm light board (light sensor is not supported)	1	115-059746-00
	Repair kit for front alarm light board (light sensor is supported)	1	115-059747-00
2	ePM battery cover connection belt	1	043-010075-00
3	TR6F recorder	1	115-059807-00
4	Speaker	1	115-059830-00
5	CO ₂ module (M02D)	1	115-059974-00
6	Integrated 10-inch standard configuration mainboard FRU (with software)	1	115-059985-00
	Integrated 10-inch full configuration 3/5-lead MR SpO ₂ mainboard FRU (with software, DC-in is not supported)	1	115-059986-00
	Integrated 10-inch full configuration 3/5-lead OEM SpO ₂ mainboard FRU (with software, DC-in is not supported)	1	115-059987-00

No.	Name and Specification	Quantity	PN
	Integrated 10-inch full configuration 3/5-lead	1	
	MR SpO ₂ mainboard FRU (with software, DC-in		115-059988-00
	is supported)		
	Integrated 10-inch full configuration 3/5-lead	1	
	$OEM SpO_2$ mainboard FRU (with software,		115-059989-00
	DC-in is supported)		
The following electrical safety tests are recommended as part of a comprehensive preventive maintenance program. They are a proven means of detecting abnormalities that, if undetected, could prove dangerous to either the patient or the operator. Additional tests may be required according to local regulations.

All tests can be performed using commercially available safety analyzer test equipment. These procedures assume the use of a 601PROXL International Safety Analyzer or equivalent safety analyzer. Other popular testers complying with IEC 60601-1 used in Europe such as Fluke, Metron, or Gerb may require modifications to the procedure. Follow the instructions of the analyzer manufacturer.

The consistent use of a safety analyzer as a routine step in closing a repair or upgrade is emphasized as a mandatory step if an approved agency status is to be maintained. The safety analyzer also proves to be an excellent troubleshooting tool to detect abnormalities of line voltage and grounding, as well as total current loads.

A. 1	Power	Cord	Plug
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A.1.1 The Power Plug

Test Item		Acceptance Criteria		
	The power plug pins	No broken or bent pin. No discolored pins.		
	The plug body	No physical damage to the plug body.		
The power plug	The strain relief	No physical damage to the strain relief. No plug warmth for device in use.		
	The power plug	No loose connections.		
		No physical damage to the cord. No deterioration to the cord.		
The power cord		For devices with detachable power cords, inspect the connection at the device.		
		For devices with non-detachable power cords, inspect the strain relief at the device.		

A.2 Device Enclosure and Accessories

A.2.1 Visual Inspection

Test Item	Acceptance Criteria
	No physical damage to the enclosure and accessories.
	No physical damage to meters, switches, connectors, etc.
The enclosure and accessories	No residue of fluid spillage (e.g., water, coffee, chemicals, etc.).
	No loose or missing parts (e.g., knobs, dials, terminals, etc.).

A.2.2 Contextual Inspection

Test Item	Acceptance Criteria		
	No unusual noises (e.g., a rattle inside the case).		
The enclosure and accessories	No unusual smells (e.g., burning or smoky smells, particularly from ventilation holes).		
	No taped notes that may suggest device deficiencies		
	or operator concerns.		

A.3 Device Labeling

Check the labels provided by the manufacturer or the healthcare facility are present and legible.

- Main unit label
- Integrated warning labels

A.4 Protective Earth Resistance

Protective Earth Resistance is measured using the RED test lead attached to the DUT Protective Earth terminal or enclosure. Select the test current by pressing SOFT KEY 3 to toggle between 1AMP, 10AMP, and 25AMP. The front panel outlet power is turned off for this test.

The following conditions apply: L1 and L2 Open.

Preparation

- 1. First select the test current that will be used for performing the Protective Earth Resistance test by pressing AMPERES (SOFT KEY 3).
- 2. Connect the test lead(s) between the RED input jack and the GREEN input jack.
- 3. Press CAL LEADS. The 601PRO will measure the lead resistance, and if less than 0.150 Ohms, it will store the reading and subtract it from all earth resistance readings taken at the calibrated current.



If the calibration fails, the previously stored readings will be used until a passing calibration has occurred.:

• During Earth Resistance testing, the DUT must be plugged into the 601PRO front outlet. If the DUT fails Earth Resistance, discontinue tests and label the device defective.

To Perform the Test

- 1. From the MAIN MENU, or with the outlet unpowered, plug the DUT into the 601PRO front panel outlet.
- 2. Attach the 601PRO RED input lead to the device's Protective Earth terminal or an exposed metal area.
- 3. Press shortcut key 3. The Protective Earth Resistance test is displayed.
- 4. Press SOFT KEY 3 to select a test current (1AMP, 10AMP, or 25AMP). The selected test current is displayed in the upper right corner of the display.



- 5. Press START TEST to start the test. The test current is applied while resistance and current readings are taken. This takes approximately 5 seconds.
- 6. Press the print data key at any time to generate a printout of the latest measurement(s).

NOTE

• When "Over" is displayed for Ohms, this signifies that a valid measurement was not obtained because either an open connection was detected or that the measurement was not within range. Readings greater than 9.999 Ohms will be displayed as Over.

In Case of Failure

Once it reaches the limitation, stop using and inform the Customer Service Engineer for analysis and disposal.

LIMITS

ALL COUNTRIES $R = 0.2 \Omega$ Maximum

A.5 Earth Leakage Test

Run an Earth Leakage test on the device being tested before performing any other leakage tests.

Leakage current is measured the following ways:

- Earth Leakage Current, leakage current measured through DUT outlet Earth
- Earth Leakage Current AP-EARTH (ALL Applied Parts connected to Earth), leakage current measured through DUT outlet Earth

There is no need to attach a test lead; the 601PRO automatically connects the measuring device internally.

To Perform the Test

- 1. From the MAIN MENU, or with the outlet unpowered, plug the DUT into the 601PRO front panel outlet, and turn on the device.
- 2. Attach the device's applied parts to the 601PRO applied part terminals if applicable.
- 3. Press shortcut key 4.The Earth Leakage test appears on the display, and the test begins immediately:



SOFT KEY 1 toggles the DUT outlet Polarity from Normal to Off to Reverse.

- SOFT KEY 2 toggles the DUT outlet from Earth to No Earth.
- SOFT KEY 3 toggles the DUT outlet from L2 to No L2.
- SOFT KEY 4 toggles the AP to Earth to No AP to Earth.
- 4. Press the print data key at any time to generate a printout of the latest measurement.

In Case of Failure

- Check any broken of the enclosure. Replace any defective part.
- Inspect wiring for bad crimps, poor connections, or damage.
- Test the wall outlet; verify it is grounded and is free of other wiring abnormalities. Notify the user or owner to correct any deviations. As a work around, check the other outlets to see if they could be used instead.
- Change another probe to confirm if the fail is caused by console.
- If the leakage current measurement tests fail on a new unit and if situation can not be corrected, submit a Safety Failure Report to document the system problem. Remove unit from operation.
- If all else fails, stop using and inform the Customer Service Engineer for analysis and disposal.

LIMITS

For IEC60601-1,

- 5mA in Normal Condition
- 10mA in Single Fault Condition

A.6 Patient Leakage Current

Patient leakage currents are measured between a selected applied part and mains earth. All measurements have a true RMS only response.

Preparation

Perform a calibration from the Mains on Applied Part menu.

The following outlet conditions apply when performing this test:

- Normal Polarity, Earth Open, Outlet ON Normal Polarity, Outlet ON
- Normal Polarity, L2 Open, Outlet ON Reversed Polarity, Outlet ON
- Reversed Polarity, Earth Open, Outlet ON Reversed Polarity, L2 Open, Outlet ON

• If all of the applied parts correspond to the instrument type, the applied parts will be tied together and one reading will be taken. If any of the applied parts differ from the instrument type, all applied parts will be tested individually, based on the type of applied part. This applies to Auto and Step modes only.

To Perform the Test

- 1. From the MAIN MENU, or with the outlet unpowered, plug the DUT into the 601PRO front panel outlet, and turn on the device.
- 2. Attach the applied parts to the 601PRO's applied part terminals.
- 3. Press shortcut key 6. The Patient Leakage test is displayed, and the test begins immediately.



- 4. Press APPLIED PART (SOFT KEY 4) at any time to select the desired applied part leakage current.
- 5. Modify the configuration of the front panel outlet by pressing the appropriate SOFT KEY on the 601PRO.
- 6. Press the print data key at any time to generate a printout of the latest measurement.

In Case of Failure

- Check any broken of the enclosure. Replace any defective part.
- Inspect wiring for bad crimps, poor connections, or damage.
- Test the wall outlet; verify it is grounded and is free of other wiring abnormalities. Notify the user or owner to correct any deviations. As a work around, check the other outlets to see if they could be used instead.
- Change another probe to confirm if the fail is caused by console.
- If the leakage current measurement tests fail on a new unit and if situation can not be corrected, submit a Safety Failure Report to document the system problem. Remove unit from operation.
- If all else fails, stop using and inform the Customer Service Engineer for analysis and disposal.

LIMITS

For CF Dapplied parts

- 10µA in Normal Condition
- 50µA in Single Fault Condition

For BF **A** applied parts

- 100µA in Normal Condition
- 500µA in Single Fault Condition

A.7 Mains on Applied Part Leakage

The Mains on Applied Part test applies a test voltage, which is 110% of the mains voltage, through a limiting resistance, to selected applied part terminals. Current measurements are then taken between the selected applied part and earth. Measurements are taken with the test voltage (110% of mains) to applied parts in the normal and reverse polarity conditions as indicated on the display.

The following outlet conditions apply when performing the Mains on Applied Part test.

- Normal Polarity;
- Reversed Polarity

Preparation

To perform a calibration from the Mains on Applied Part test, press CAL (SOFT KEY 2).

- 1. Disconnect ALL patient leads, test leads, and DUT outlet connections.
- 2. Press CAL to begin calibration, as shown:



If the calibration fails, the previously stored readings will be used until a passing calibration has occurred. Also, the esc/stop key has no effect during calibration.

3. When the calibration is finished, the Mains on Applied Part test will reappear.

- A 2-beep-per-second signal indicates high voltage present at the applied part terminals while a calibration is being performed.
- High voltage is present at applied part terminals while measurements are being taken.

To Perform the Test

- 1. From the MAIN MENU, or with the outlet unpowered, plug the DUT into the 601
- 2. Attach the applied parts to the 601PRO applied part terminals.
- 3. Attach the red terminal lead to a conductive part on the DUT enclosure.
- 4. Press shortcut key 7. The Mains on Applied Part test is displayed.

Mains on Applied Outlet: Norm Pol Norm uA	Part: All-Ear , Earth, L2 Rev u	th A [Limit Inv	/]	
START TEST	CAL	DUT OFF	APPLIED PART	
				ENT

- 5. Select the desired outlet configuration and applied part to test using the appropriate SOFT KEYS:
- 6. Press START TEST (SOFT KEY 1) to begin the test.
- 7. Press the print data key to generate a printout of the latest measurement.

NOTE

• If all of the applied parts correspond to the instrument type, the applied parts will be tied together and one reading will be taken. If any of the applied parts differ from the instrument type, all applied parts will be tested individually, based on the type of applied part. This applies to Auto and Step modes only.

In Case of Failure

- Check any broken of the enclosure. Replace any defective part.
- Inspect wiring for bad crimps, poor connections, or damage.
- Test the wall outlet; verify it is grounded and is free of other wiring abnormalities. Notify the user or owner to correct any deviations. As a work around, check the other outlets to see if they could be used instead.
- Change another probe to confirm if the fail is caused by console.
- If the leakage current measurement tests fail on a new unit and if situation can not be corrected, submit a Safety Failure Report to document the system problem. Remove unit from operation.
- If all else fails, stop using and inform the Customer Service Engineer for analysis and disposal.

LIMITS



A.8 Patient Auxiliary Current

Patient Auxiliary currents are measured between any selected ECG jack and the remaining selected ECG jacks. All measurements may have a true RMS only response.

Preparation

- 1. From the MAIN MENU, or with the outlet unpowered, plug the DUT into the 601PRO front panel outlet, and turn on the device.
- 2. Attach the patient leads to the 601PRO ECG jacks.
- 3. Define the Lead Types from the View Settings Option (refer to: Lead Type Definitions in Section 5 of this chapter).
- 4. Press shortcut key 8. The Patient Auxiliary Current test is displayed, and the test begins immediately. Display values are continuously updated until another test is selected.



- 5. Press SOFT KEYS 1-4 to select leakage tests
- 6. Press APPLIED PART (SOFT KEY 4) at any time to select the desired applied part leakage current:
- 7. Modify the configuration of the front panel outlet by pressing the appropriate SOFT KEY on the 601PRO:
- 8. Press the print data key at any time to generate a printout of the latest measurement.

In Case of Failure

- Check any broken of the enclosure. Replace any defective part.
- Inspect wiring for bad crimps, poor connections, or damage.

- Test the wall outlet; verify it is grounded and is free of other wiring abnormalities. Notify the user or owner to correct any deviations. As a work around, check the other outlets to see if they could be used instead.
- Change another probe to confirm if the fail is caused by console.
- If the leakage current measurement tests fail on a new unit and if situation can not be corrected, submit a Safety Failure Report to document the system problem. Remove unit from operation.
- If all else fails, stop using and inform the Customer Service Engineer for analysis and disposal.

LIMITS

For CF D applied parts,

- 10µA in Normal Condition
- 50µA in Single Fault Condition

For BF 🛣 applied parts,

- 100μA in Normal Condition
- 500µA in Single Fault Condition

ELECTRICAL SAFETY INSPECTION FORM

Overall assessment:

Scheduled inspection

Test item: 1, 2, 3, 4, 5, 6, 7, 8

Location:					Technician:		
Equipment:					Control Number:		
Manufacturer: Model:					SN:		
Measu	urement equip	oment /SN:	Date of Calibration:				
INSPE	CTION AND T	ESTING			Pass/Fail	Limit	
1	Power Cord	Plug					
2	Device Enclo	osure and Acces	sories				
3	Device Labe	ling					
4	Protective Ea	arth Resistance		Ω		Max 0.2 Ω	
_	Farth Leakar	Normal conditio	n(NC)	μΑ		Max:	
		Single Fa	ault n(SFC)	μΑ		SFC: 10mA	
	Patient Leak	Normal conditio	n(NC)	□BFμA □CFμA	_	Max: CF applied part:	
6	Current	Single Fa	ault n(SFC)	□BFμA □CFμA	NC:10µA, SFC: 50µA BF applied part: NC:100µA, SFC: 500µA		
7	Mains on Applied Part Leakage			□BFμA □CFμA	-	Max: CF applied part: 50µA BF applied part: 5000µA	
8	Patient Auxiliary Current	Normal cond	ition(NC)	□BFµА □CFµА	_	Max: CF applied part:	
		Current Single Fault condition(SFC)		□BFµА □CFµА		- NC:10µA, SFC: 50µA BF applied part: NC:100µA, SFC: 500µA	

Name/ Signature: _____ Date: _____

Unopened repair type	Test item: 1, 2, 3
Opened repair type, not replace the power part including transformer or patient circuit board	Test item: 1, 2, 3, 4
Opened repair type, replace the power part including transformer	Test item: 1, 2, 3, 4, 5
Opened repair type, replace patient circuit board	Test item: 1, 2, 3, 4, 6, 7, 8

Location:					Technician:		
Equipment:					Control Number:		
Manufacturer: Model:			SN:				
Meas	Measurement equipment /SN:				Date of Calibration:		
INSPE	INSPECTION AND TESTING				Pass/Fail	Limit	
1	Power Cord	Plug					
2	Device Enclo	osure and Acces	sories				
3	Device Labe	ling					
4	Protective E	arth Resistance		Ω		Max 0.2 Ω	
5			n(NC)	μΑ		Max:	
5	Lurti Leuka	Single Fa	ault n(SFC)	μΑ		SFC: 10mA	
6	Patient Leak	Normal conditio	n(NC)	□BFμА □CFμА	_	Max: CF applied part: NC:10uA, SFC: 50uA	
0	Current	Single Fa	ault n(SFC)	□BFμA □CFμA	_	BF applied part: NC:100μA, SFC: 500μA	
7	Mains on Applied Part Leakage			_BFμA	_	Max: CF applied part: 50μA BF applied part: 5000μA	
8	Patient Auxiliary	Normal cond	ition(NC)	□BFµА □CFµА	_	Max: CF applied part: NC:10µA, SFC: 50µA	
	Current	Single Fault condition(SFG	_)	□BFμA □CFμA	_	BF applied part: NC:100μA, SFC: 500μA	
N	Name/ Signature: Date:						