

## Foreword

**Please read the User Manual carefully before using this product. The User Manual which describes the operating procedures should be followed strictly.** This manual detailed introduce the steps must be noted when using the product, operation which may result in abnormal, the risk may cause personal injury and product damage and other contents, refer to the chapters for details. Any anomalies or personal injury and device damage arising from use, maintain, store do not follow requirements of the User Manual, Our company is not responsible for the safety, reliability and performance guarantees! The manufacturer's warranty service does not cover such faults!

Our company has a factory record and user profile for each device, users enjoy free maintenance services for one year from the date of purchase. In order to facilitate us to provide you with a comprehensive and efficient maintenance service, please be sure to return the warranty card when you need repair service.

 **Note: Please read the User Manual carefully before using this product.**

Described in this User Manual is in accordance with practical situation of the product. In case of modifications and software upgrades, the information contained in this document is subject to change without notice.

## **The warning items**

**Before using this product, you should consider the safety and efficacy of the following described:**

- Described each measurement results combined with clinical symptoms by qualified doctors.
- The reliability and operation of using this product whether meets the operation of this manual relate to the maintenance instructions.
- The intended operator of this product may be the patient.
- Do not perform maintenance and service while the device is in use.

## **Responsibility of operator**

- The operator must carefully read the User Manual before use this product, and strictly follow the operating procedure of the User Manual.
- Fully consider the security requirements during product design, but the operator should not ignore the observation for the patient and the state of machine.
- The operator has the responsibility to provide the use condition of the product to our company.

### **Responsibility for our company**

- Our company have the responsibility to provide qualified product which conform to company standard of this product.
- Our company will provide the circuit diagram, calibration method and other information at the request of the user to help the appropriate and qualified technicians to repair those parts designated by our company.
- Our company have the responsibility to complete product maintenance according to the contract.
- Our company have the responsibility to respond the requirements of user in time.
- In the following case, our company is responsible for the impact on the safety, reliability and performance of the device:

Assembly, addition, debugging, modification or repair are carried out by personnel approved by our company.

The electrical facilities in the room are in compliance with the relevant requirements and the device is used in accordance with the User Manual.

**The User Manual is written by our company. All rights reserved.**

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## Chapter1 Introduction

Operators do not need professional training, but should use this product after fully understanding the requirements in this manual.

To prevent users from suffering hurt or damnification due to improper use, please refer to "**Safety Precautions**" and use this product properly.

For an overall introduction to the Blood Pressure Monitor, please refer to **General Information**.

For basic operating instructions, please refer to **Button Function**.

For allocation of interface sockets, please refer to **Interfaces**.

### 1.1 Safety Precautions



- If not use correctly, it exists the possibility of damage for personnel and goods.
- Good damage means the damage of house, property, domestic animal and pet.
- For severe blood circulation disorder or arrhythmia patients, please use the device under the guidance of a doctor. Otherwise it may lead to acute hemorrhage, or measurement error as a result of squeezed arm.
- You must not perform NIBP measurements on patients with sickle-cell disease or under any condition which the skin is damaged or expected to be damaged.

■ **For a thrombasthemia patient, it is important to determine whether measurement of the blood pressure shall be done automatically. The determination should be based on the clinical evaluation.**

 **Contraindication** 

No contraindications.

 **Warning** 

**Do not use the device in the case of there are flammable anesthetic gasses mixing with the air or nitrous oxide.**

Otherwise it may cause risk.

**For children and the person who can't express oneself, please use the device under the guidance of a doctor.**

Otherwise it may cause accident or dissension.

**Self-diagnosis and treatment using measured results may be dangerous. Follow the instructions of your physician.**

Please hand measurement results to the doctor who knows your health and accept diagnosis.

**Please do not use for any other purpose except BP measurement.**

Otherwise it may cause accident or holdback

**Please use special cuff.**

Otherwise it is possible that measurement result is incorrect.

**Please do not keep the cuff in the over-inflated state for a long time.**

Otherwise it may cause risk.

**If liquid splashes on the device or accessories, especially when liquids may enter the pipe or device, stop using and contact the service department.**

Otherwise it may cause risk.

**Dispose of the packaging material, observing the applicable waste control regulations and keeping it out of children's reach.**

Otherwise it may cause harm to the environment or children.

**Please use approved accessories for the device and check that the device and accessories are working properly and safely before use.**

Otherwise the measurement result may be inaccurate or an accident may occur.

**When the device is accidentally damp, it should be placed in a dry and ventilated place for a period of time to dissipate moisture.**

Otherwise the device may be damaged due to moisture.

**Do not store and transport the device outside the specified environment.**

Otherwise it may cause measurement error.

**It is recommended that you check if there is any damage on the device or the accessories regularly, if you find any damage, stop using it, and contact the biomedical engineer of the hospital or our Customer Service immediately. Do not disassemble, repair and modify the device without permission.**

Otherwise it cannot be accurately measured.

**This device can not be used on mobile transport platforms.**

Otherwise it may cause measurement error.

**This device can not be used on a tilted tabletop.**

Otherwise there is a risk of falling.

**Dispose of packaging materials, waste batteries and end-of-life products in accordance with local laws and regulations. The end-of-life products and materials are properly disposed of by the user in accordance with the authority's decree.**

Replace accessories which not provided by our company may lead to the occurrence of errors.

Without our company or other approved maintenance organizations trained service personnel should not try to maintain the product.

**This device can only be used for one test object at a time.**

If the small parts on the device are inhaled or swallowed, please consult a doctor promptly.

The device and accessories are processed with allergenic materials. If you are allergic to it , stop

using this product.

**Do not use a mobile phone near the blood pressure monitor. Excessive radiation fields generated by mobile phones can interfere with the normal use of the blood pressure monitor. The blood pressure monitor has slight electromagnetic radiation to the external environment, but does not affect the normal use of other equipment.**

This device is suitable for occasions with electrosurgical equipment, but when used with electrosurgical equipment, patient safety must be given the highest priority.

**The parts of the device that are in contact with the patient (cuffs, air pipes, enclosure, etc.) are made of insulating material and the device is protected against electric shock. When high frequency or defibrillation devices are applied to the patient, no special precautions need to be taken and the defibrillator discharge will not affect the device.**

If Luer lock connectors are used in the construction of tubing, there is a possibility that they might be inadvertently connected to intravascular fluid systems, allowing air to be pumped into a blood vessel.

**This device is suitable for occasions with electrosurgical equipment, but when used with electrosurgical equipment, patient safety must be given the highest priority.**

When the monitor is wetted, please stop using it and contact us.

**After pressing the power button, if the device has display fault such as white screen,**

blurred screen or no display content, please contact our company.



- The software was developed in accordance with IEC60601-1. The possibility of hazards arising from errors in the software program has been minimized.
- All analog and digital equipment connected to this device must be certified to IEC standards(such as IEC60950: Information technology equipment-Safety and IEC60601-1: Medical electrical equipment-Safety), and all equipment should be connected to in accordance with the requirement of the valid version of the IEC60601-1-1 system standard. The person connecting the additional equipment to the signal input and output port is responsible for whether the system complies with the IEC60601-1 standard.
- Refer to the following chapters for the minimum value of patient physiological signals. Operation of the device below the minimum value may result in inaccurate results.
- The Monitor shall comply with the standard IEC 80601-2-30:Particular requirements for basic safety and essential performance of automated non-invasive sphygmomanometers.

## 1.2 General Information

This device is defibrillator protected,the time of defibrillation recovery is 5 seconds.Note that no

precautions specific to the device is required during defibrillation, and defibrillation discharge has no effect on the monitor. The equipment uses the gray silicone airway, in case of the effect to the equipment when defibrillation device was used on the patient.

**General instruction:**

The device is applied to Blood Pressure(BP) measure and monitor for adult(including pregnant women), pediatric, and neonatal. It most stores 300 records of common user and 350 of ambulatory Blood Pressure data. Every record includes the detailed measure time, systolic blood pressure, diastolic blood pressure, mean blood pressure, pulse rate, error message and record number, etc.

This device has friendly operation interface, and adopts 2.4inch color LCD. It integrates data review function and display function which includes BIG FONT single record data review, data list, BP data trends chart, the current time,date,power and so on.

User can power on/off the monitor,start manual measure, set system parameters and so on with five keys in the front panel. (Please refer to "Button Functions" part for detail)

The monitor does not have an alarm system, but it will prompt when the power is low, the measure is wrong, or the measure data exceeds the set limits. When the power is low or the measure is wrong, the prompt is audible and visual, the device will buzz intermittently and the red light will flash to prompt the user to replace batteries or prompt the reason of the failed

measurement; when the measure data exceeds the set limits, the prompt is audible, the font color of measure results will change to red. Users can open and close the prompt according to needs. The cuff socket is located on the top of the device and the USB socket at the bottom of the device. The stored data can be transferred to computer with the USB interface, and then various operations can be performed by using the PC software. (Please refer to "Software Functions" part for detailed contents)



**In the common user mode, the monitor will periodically turn off backlight if there is no operation, and automatically shuts down if there is no operation for two minutes. When the backlight turn off in the ambulatory blood pressure mode, the blue indicator intermittently flashes to prompt the device in running state.**

### 1.3 Button Functions

All the operations of the Blood Pressure Monitor could be completed with the buttons. The names of button are on them. They are:



Press the button for a long time, then the system will start. When turning on and off the monitor, the red light and the blue both flash once to prompt that the on or off operation is successful.

Press it for a short time to return the boot-strap interface.



The text in the middle bottom of the screen indicate the function of this key. Whatever menus the system is in, press the button and the system immediately executes a certain function.



The text in the left bottom of the screen indicate the function of this key.

Such as: The button is the prompt switch in the boot-strap interface, up key in the "SYSTEM MENU", and left key in the "TREND" chart.



The text in the right bottom of the screen indicate the function of this key.

Such as: the button is the data review key of current user in the boot-strap interface and down key in the "SYSTEM MENU" and right key in the "TREND" chart.



Start/Stop button. If measuring, press this key to cancel the current measurement.



■ **After connecting the USB cable, all of the buttons are disabled. If the BP**

measurement is in progress, this measurement is will be automatically canceled.

■ During measurement,  three buttons are all disabled.

The rectangular mark in the screen moving with the operation of ,  buttons is called “cursor”. Operation can be performed in any position at which the cursor can stay. When the item is not selected, the cursor is yellow; when selected, the cursor becomes red.

#### 1.4 Interfaces

For the convenience of operation, different kinds of interfaces are in different parts of the device. NIBP cuff socket is at the top of the device.



**The connection of the NIBP external air pipe is as shown:**

- ① Cuff extension tube metal nozzle
- ② the socket for air pipe

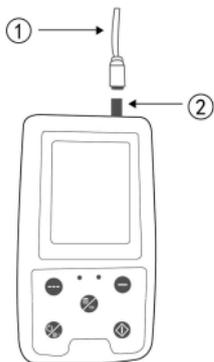


Figure 1.4.1 The Top External airway

**At the bottom is the socket for USB:**

①The Socket for USB, connect the data line to upload data.

### 1.5 Accessories

- 1) A cuff for adult
- 2) An USB data line
- 3) A disk (PC software)
- 4) BP extending tube

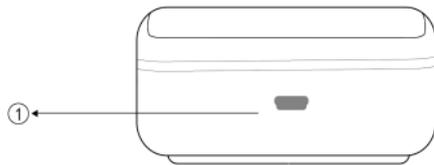


Figure 1.4.2 Bottom

5) A pouch



The monitor also can be equipped with pediatric and neonatal cuff, if necessary, please contact our company or its representatives.

**The width of the cuff should be 40% of the limb circumference(50% for the newborn) or 2/3 of the length of the upper arm.The length of the inflated part of the cuff should be sufficient to surround 50% to 80% of the limb. Unsuitable cuffs can produce erroneous readings. If there is a problem with the size of the cuff, use a larger cuff to reduce the error.**

Reusable Cuff of adult/ pediatric/ neonatal

Patient type	Limb circumference	Width of the cuff	Length of inflatable tube
Neonatal	10~19 cm	8 cm	1.5 m or 3 m
Pediatric	18~26 cm	10.6 cm	
Adult 1	25~35 cm	14 cm	
Adult 2	33~47 cm	17 cm	

Disposable cuff of neonatal

Size	Limb circumference	Width of the cuff	Length of inflatable tube
1	3.1~5.7 cm	2.5 cm	1.5 m or 3 m

2	4.3~8.0 cm	3.2 cm	
3	5.8~10.9 cm	4.3 cm	
4	7.1~13.1 cm	5.1 cm	



**Warning**



**Please use the special accessories supplied by the manufacturer or replace the accessories according to the requirements of the manufacturer in order to avoid making harms to patients.**



**Note**



- The cuff is a consumable. In order to correctly measure blood pressure, please replace the cuff in time.
- If the cuff leaks, please contact our company to buy a new one. The cuff purchased separately does not include the BP extension tube. Please give an explanation if you need to buy a BP extension tube at the same time. If you do not want to buy a BP extension tube, please do not throw the BP extension tube away when replacing the cuff, install it on the new cuff.
- Pouch is convenient for patients to carry the monitor. It is not necessary to replace it when the backpack has a slight wear. Patients can according to the actual situation, contact our company to buy a new backpack when the original backpack can not carry the monitor.

 **Note** 

When the product and accessories described in this manual are about to exceed the period of use, they must be disposed according to relevant product handling specification. If you want to know more information, please contact our company or representative organization.

## Chapter2 Getting Started

### 2.1 Open the Package and Check

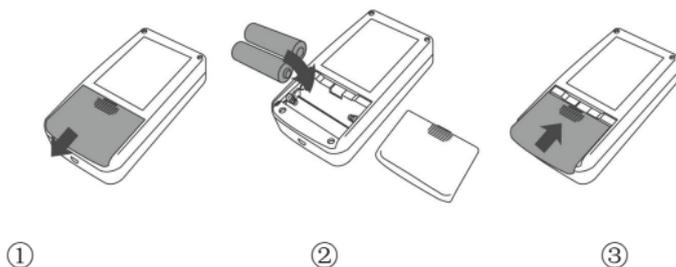
Open the package and take out the equipment and accessories carefully. Keep the package material for possible future transportation or storage. Check the components according to the packing list.

- Check for any mechanical damage.
- Check all the cables, modules and accessories.

If there is any problem, contact the distributor immediately.

### 2.2 Battery Installation

The instrument will be supplied with two 'AA' alkaline batteries or high capacity. Before using the instrument, you shall put the battery in the battery box in the back of the Monitor .



- ① Demount the battery cover in the direction of the arrow.
- ② Install "AA" batteries according to  $\oplus$   $\ominus$  polarities.
- ③ Slide to close the battery cover.

**⚠ Note ⚠**

Icon“

**⚠ Cautions ⚠**

- Turn the unit off before replacing the batteries.
- Please use 2 "AA" size manganese or alkaline batteries, do not use batteries of other types. Otherwise it may cause fire.
- New and old batteries, different kinds batteries can not be put off. Otherwise it may cause battery leakage, heat, rupture, and damage to the monitor.
- "+" and "-" polarities of the batteries must match the polarities of the battery compartment as indicated. When the batteries power exhausts, replace with 2 new batteries at the same time.
- Please take out the batteries when you do not use the device for a long time (more than ten days). Otherwise it may cause battery leakage, heat, rupture, and damage to the monitor.
- If electrolyte of battery gets in your eyes, immediately rinse with plenty of clean

water. Contact a physician immediately. Otherwise it will cause blindness or other hazards.

- If electrolyte of the batteries immodestly glues on the skin or the clothes, please immediately flush with plenty of clean water. Otherwise it may hurt the skin.
- Dispose of the exhausted batteries according to applicable local regulations about environmental. Otherwise it will cause environmental pollution.
- The monitor is internally powered equipment, can be connected to the public grid.

### 2.3 Power on the Instrument

Hold the power button , the indicator will flash once, which shows the boot-strap is successful, then release the button, and the system will enter the main interface.

Hold the power button  after power on, the indicator will flash once, which shows the shutdown is successful, and the device can be safely terminated.



**If any sign of damage is detected, or the instrument displays some error messages, do not use it on any patient. Contact biomedical engineer in the hospital or our Customer Service Center immediately.**

The device can be used normally after it is turned on ,without waiting for the device to be prepared.

⚠ Note ⚠

Check all the functions that possibly be used and make sure that the device is in good status.

#### 2.4 Connect Sensor

⚠ Note ⚠

For information on correct connection of NIBP cuff, refer to Figure 2.4.1

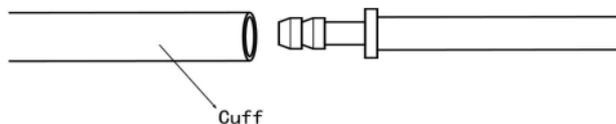


Figure 2.4.1 Connection Method

Connect the sensor between the Monitor and the measure part of the patient.

## Chapter3 Function Interface

### 3.1 Main Interface

Press  to power on the instrument. The indicator will circularly flash once, which show the boot-strap is success, then end pressing, the system will enter into the main interface.

In common user mode, if there is no key-press operation during the time which system sets, the device will turn off LCD and enter into standby mode, if there is no any operation in the standby mode, the device will automatically turn off; the "RUN" indicator flashes once every 3 seconds to prompt the device in working state.

When the power is low, the battery progress bar is empty, at the same time the prompt sound occurs, and the red indicator flashes in fixed time.

#### **In the Main Interface:**

Prompt-switch status is displayed in the left top of the screen,  button can switch the prompt status shortly.

User bar displays the current patient type (adult, pediatric, neonatal), and the amount of the common user's data record.

Current date and time is displayed in the middle top of the screen.



- All interfaces except the trend retain power icon, prompt switch, as well as a small font of the current time.
- The oldest record will be overwritten after the memory overflows. “Overflow” message is shown in the main interface.

### 3.2 Measuring Interface

Measuring interface displays real-time cuff pressure and the current measurement information. In the measurement process, except the  and the  buttons, other buttons are disabled.



In any interface except the measurement, press  key to exit current interface and back to the boot-strap interface.

### 3.3 Measure Result Interface

#### The measure result includes:

SYS: systolic blood pressure (mmHg/kPa)

DIA: diastolic blood pressure (mmHg/kPa)

PR: pulse rate (bpm)

If there is an error during the measurement, an error message text will appear on the screen. If the PROMPT SOUND is set to be on, the sound would occur. Press the SILENCE key to stop the sound and press it once more to continue

### 3.4 System Menu

In the main interface, according to the text in the middle bottom of the screen, press  button, then enter the system menu and execute different option operations by using  and  buttons.



Figure 3.4.1 System Menu

### 3.4.1 System Setup

Enter "SYSTEM SET" item in the [SYSTEM MENU], the "SYSTEM SET" menu includes:

"LANGUAGE" item: switch the current system language;

"UNIT" item has two choice: mmHg, kPa;

"MEASURE MODE" item has three options: adult, pediatric, neonatal;

"ABPM SET" item: set ABPM parameters:

"BACKLIGHT TIME(s)" item: 15, 30, 60, 120

"CALIBRATION": is only used when it is necessary to detect the real-time pressure of the

device.



**Note**

**"BACKLIGHT TIME" in the "SYSTEM SETUP" is used by the common user, ambulatory blood pressure backlight time is a fixed value of 5s.**

To perform ambulatory blood pressure monitoring, first select "ABPM SETUP" item in [SYSTEM SETUP] menu, the pop-up menu is shown in the Figure3.4.2:

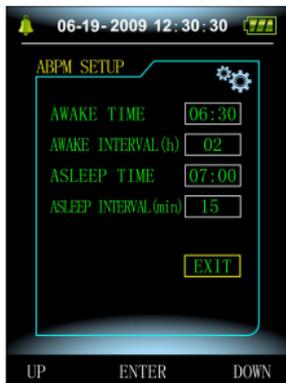


Figure 3.4.2 ABPM Setup

Options for "AWAKE INTERVAL(min)" and "ASLEEP INTERVAL(min)": 15, 20, 30, 40, 60, 90, 120, 180, 240;

The step of each adjustment for "AWAKE TIME" and "ASLEEP TIME" is 30 minutes. The adjustment range: 00:00~23:30.



**The set measurement interval in "AWAKE INTERVAL" and "ASLEEP INTERVAL" is time interval when automatically start the measurement under the mode of ambulatory blood pressure, not including manual starting. For example: set "AWAKE TIME" to 7:00, set "AWAKE INTERVAL" to 15min, then, the device will make the first blood pressure measurement at 7:15; if the user start a blood pressure measurement by pressing measuring button between 7:00-7:15, the device will also automatically start the measurement at 7:15, and not affected by manual measurement.**

After each item of this interface is set, the ambulatory blood pressure menu also need to be correctly set to start the ABPM function. Refer to 3.4.5 ambulatory blood pressure menu for details.

### 3.4.2 System Time

Select "SYSTEM TIME" item in [SYSTEM MENU], the following menu will pop up:



Figure 3.4.3 System Time

Select "SAVE" after completing time setup, time change is successful and exit system time setup and return the previous menu. Select "CANCEL" to cancel the setting and return to the previous menu.

### 3.4.3 Data Delete

Select "YES" in "DELETE DATA" menu of [SYSTEM MENU], after you press certain key, the following menu will pop up:



Figure 3.4.4 Data Delete

If press "CONFIRM", the common user data will be deleted, press "CANCEL", the operation will be canceled.

### **3.4.4 Prompt Setup**

Select "PROMPT SET" item in [SYSTEM MENU] to enter its setup interface, then make corresponding settings according to the following procedure:

"SYS PROMPT" and "DIA PROMPT" can control closing or unsealing of the SYS and DIA prompt separately.

The prompt is on or off according to the high and low limits which have been set up. When the measure result is higher than the high limit or lower than the low limit, and meanwhile, the "PROMPT SOUND" is on, "SYS PROMPT" or "DIA PROMPT" accordingly on, the prompt will occur.

The adjustable ranges of the high and low limits of the adult mode prompt are as follows:

SYS PROMPT: 40~270 mmHg

DIA PROMPT: 10~215 mmHg

The adjustable ranges of the high and low limits of the pediatric mode prompt are as follows:

SYS PROMPT: 40~200 mmHg

DIA PROMPT: 10~150 mmHg

The adjustable ranges of the high and low limits of the neonatal mode prompt are as follows:

SYS PROMPT: 40~135 mmHg

DIA PROMPT: 10~100 mmHg

"DEFAULT" includes the main content:

Measure mode: adult;

Parameter prompting limit:

User mode	High pressure high limit	High pressure low limit	Low pressure high limit	Low pressure low limit
Adult	140	90	90	40
Pediatric	120	70	70	40
Neonatal	100	60	60	20

PROMPT SOUND switch: ON;

Measure unit: mmHg;

Ordinary user backlight time: 15s;

ABPM switch: END;

Asleep time: 22:00;

Asleep measurement interval: 30minutes;

Awake measurement interval: 15minutes;

Awake time: 7:00;

SYS PROMPT switch: ON;

DIA PROMPT switch: ON.

**Note: The monitor does not have an alarm system, the above-mentioned "ALARM"**

**represent audible/visual prompt.**

### **3.4.5 ABPM Menu**

#### **1. ABPM mode**

After the ambulatory blood pressure menu is operated correctly(refer to 3.4.1), select "ABPM" menu in [SYSTEM MENU] to enter its interface.

Switch the "ABPM ON-OFF" to "BEGIN", then a prompt message for ABPM of current user will pop up, such as:

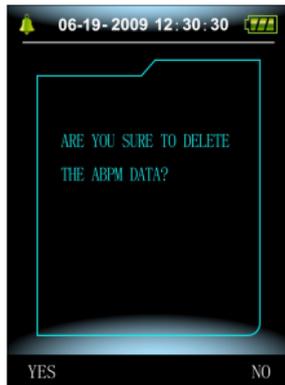


Figure 3.4.5 ABPM Prompt Menu

Press  button, clear the ambulatory blood pressure measure data, enter ambulatory blood pressure mode, and start the ambulatory blood pressure monitoring. Refer to Figure 3.4.6 for the ABPM interface.

Press  button, save ambulatory blood pressure measure data, enter ambulatory blood pressure mode, and start the ambulatory blood pressure monitoring. Ambulatory blood pressure measurement record includes previous data. Refer to Figure 3.4.6 for the ABPM interface.

Press  button, give up the choice, return the previous menu, and the ambulatory blood pressure monitoring is not turned on.

## **2. ABPM Working Interface**

In ABPM working environment, backlight is only for 5 seconds, except the , press any key to wake the backlight, ABPM working interface is as shown:



Figure 3.4.6 ABPM Working Interface

If PROMPT SOUND occurs, press SILENCE key to stop it and press it again to continue.

In ABPM working interface, long press  button, the exit ABPM hint interface will pop up. In this interface, press  button to exit ABPM working environment, and enter the ordinary user working environment, the boot-strap interface will be displayed. In the exit ABPM hint interface, press  button to exit the interface, and return the ABPM working interface.

In the ABPM working interface, to turn off the device, exit the ABPM mode first, then long press the power switch to turn it off.

### 3. ABPM Data Review

Select "ABPM DATA" item in "ABPM" menu to enter the data review interface.

- "BIG FONT" display interface: Every record is an interface, and display contents include: the current user, total of the current user record data, serial number of the record, stored time of the record, high pressure, low pressure, mean pressure, Pulse Rate.
- In the ABPM "BIG FONT" data review interface, press  button to select "LIST", the data table interface will pop up. Each interface contains 5 records, every record includes time, high pressure, low pressure, mean pressure, Pulse Rate.
- In the ABPM "LIST" data review interface, press  button to select "TREND", the data trend interface will pop up. Trend interface can trace 100 trend records, if measuring data are more than 100 items, press ,  buttons to glide trend curve for left and right, the scale of the vertical axis and the starting point, end point automatically adjust according to the width of

the stored data. The date displayed at the bottom of trends are the data recording time of the first point and last point respectively for current trend.

#### **3.4.6 PROMPT SOUND**

After selecting "ON", the loudspeaker turns on. The symbol  will display in the main interface. After selecting "OFF", the loudspeaker turns off,  will appear. When you change the settings, the password input box will appear, enter the correct password "8015" to change.

The inputting method of password: move the cursor to the password display area, press the middle button, when the rectangle frame turns to red selected state, adjust the number by the "Up" and "Down" button, then press the middle button again to exit the selected state after adjusting. After entering the 4-bit password, move the cursor to "CONFIRM", then press the middle button, the prompt sound setting can be changed if the password is correct.

### **3.5 Ordinary User Data Review**

- Ordinary User "BIG FONT" Data Review

Press  button to enter the ordinary user "BIG FONT" data review in boot-strap interface.

Display content is similar to ambulatory blood pressure BIG FONT data review.

- Ordinary User “LIST” Data Review

Press  button to pop up the ordinary user data “LIST” in the ordinary user BIG FONT data review interface. Display content is similar to ambulatory blood pressure data list.

- Ordinary User “TREND” Data Review

Press  button to pop up the ordinary user data “TREND” in the ordinary user LIST data review interface. Display content is similar to ambulatory blood pressure trend.

the device. Press  button to exit the interface, and return the ABPM working interface.

## Chapter4 NIBP Measuring

### 4.1 General

- The Non-invasive Blood Pressure (NIBP) module measures the blood pressure using the oscillometric method .It is that: using the blade to block artery blood, checking the oscillometric wave during degassing for sure that it was not affected by the operator's subjective factors or the disruption of the environmental noise.
- There are two modes of measurement available: manual and automatic .Each mode displays the diastolic, systolic and MAP blood pressure and pulse rate.
- It is applicable for adult, pediatric, and neonatal usage.

#### **Warning**

Prolonged non-invasive blood pressure measurements in Auto mode may be associated with purport, ischemia and neuropathy in the limb wearing the cuff. When monitoring a patient, examine the extremities of the limb frequently for normal color, warmth and sensitivity. If any abnormality is observed, stop the blood pressure measurements.

#### **Warning**

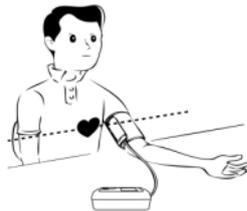
You must not perform NIBP measurements on patients with sickle-cell disease or under any

condition which the skin is damaged or expected to be damaged.

**For a thrombasthenia patient, it is important to determine whether measurement of the blood pressure shall be done automatically. The determination should be based on the clinical evaluation.**

#### 4.1.1 Accurate Measurement Way

1. Adopt a comfortable sitting position, use back and arms to support the body.
2. Place your elbow on a table, the palm faces up.
3. The cuff is level with your heart.
4. Feet flat on the floor, and do not cross your legs.



#### **Note**

- Do not speak or move during measuring.
- Do not use mobile devices such as cellphone near the device when measuring.
- Measurement results may be different due to different cuff position.
- Do not touch the device, cuff or extension tube during measuring.
- Refer to Section 1.1 for the contraindications of NIBP measurement.
- When measuring on pediatric or neonatal patients, be sure to select the correct

measurement mode (refer to the measurement mode setting) and use specified cuff for pediatric or neonate. Using incorrect measurement mode may cause danger to the patient, because the adult pressure level is relative high and is not suitable for pediatric or neonatal patients.

■ In neonatal mode, the initial inflation value of cuff is 9.3kPa (70mmHg). The cuff can be inflated to 20kPa (150mmHg), if the inflating value exceeds this value, the device will reset and deflate automatically.

■ The device has double overpressure protection for hardware and software. If over-inflating occurs, the device will reset and deflate immediately. If the device keeps over-inflating status, please disconnect the cuff from the device, and cut off the power or turn off the device.

■ Please use the device under proper temperature and humidity (refer to Specification), otherwise measured result may not accurate.



**Measure should be taken in a quiet place, and relax the body.**

Remain still 4~5 minutes before measure.

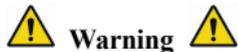
Relax the body, do not let the muscle function.

Do not talk and move during the measure.

Wait 4~5 minutes when measuring in succession.

Do not use mobile equipment such as mobile telephone near the device.

## 4.2 Applying the Cuff and NIBP Measuring



**Before starting a measurement, verify that you have selected a setting appropriate for your patient. (adult, pediatric or neonate). Do not apply the cuff to a limb that has an intravenous infusion or catheter. This could cause tissue damage around the catheter when infusion is slowed or blocked during cuff inflation.**

**The minimum value of the patient's physiological signal is the lower limit that the device can measure. The measured result may inaccurate if the device running below the minimum amplitude or minimum value of patient's physiological signal.**

**Do not twist or tangle the airway tube, otherwise it will cause continuous pressure in the cuff, then causing blocked blood flow and serious injury to the patient.**

**Do not use the cuff on the injured area, otherwise it will cause more serious damage to the injured area.**

**Do not use the cuff on the site where intravascular treatment is being performed or with catheter connection, otherwise it may cause temporary blockage of blood flow and then cause injury to the patient.**

**Do not use the cuff on the side of the mastectomy;**

**The pressure by cuff may cause temporary weakness of some functions of the body. So do**

**not use monitoring medical electrical equipment on corresponding arm.**

**Do not move during measuring, because it will have a delayed effect on the patient's blood flow. The device needs 2 hours recovery time to reach its performance of intended use after taking out from the lowest storage temperature.**

**The device needs 4 hours recovery time to reach its performance of intended use after taking out from the highest storage temperature.**

1. Plug the air hose to the cuff socket on the device, and connect the device to power supply.
2. Apply the cuff to the patient's upper arm following the instructions below (Figure4.2.1).
  - Ensure the cuff is completely deflated.
  - Apply the appropriate size cuff to the patient, and make sure that the symbol " $\phi$ " is over the appropriate artery. Ensure that the cuff is not wrapped too tightly around the limb. Excessive tightness may cause discoloration and eventual ischemia of the extremities.
  - Make sure that the cuff edge falls within the range of mark <->. If it does not, use a larger or smaller cuff that fits better.

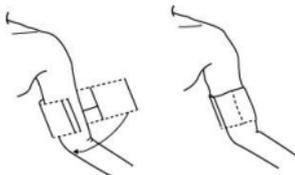


Figure 4.2.1 Cuff applying

3. Connect the cuff to the airway tube. The cuff should be placed at the same level as the patient's heart. Otherwise amend the measurement results by the following methods

- If the cuff is placed higher than the heart level, add 0.75 mmHg (0.10 kPa) for each inch of difference.
- If it is placed lower than the heart level, minus 0.75 mmHg (0.10 kPa) for each inch of difference.

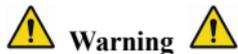
4. Check whether the measure mode is appropriately selected. (the measure mode displays in the information area of the main interface ).

5. Press  button on the front panel to start inflating and measuring.

### 4.3 Operation Hints

1. To start auto measuring:

In ABPM SETUP menu, select the "ASLEEP INTERVAL" item and "AWAKE INTERVAL" item, in which the user may select the time interval for auto measurement. After that, enter "ABPM" menu and select to enter the ABPM working environment, and the system will start inflating and measuring automatically according to the set time interval.



**Prolonged non-invasive blood pressure measurements in Auto mode may be associated**

**with purport, ischemia and neuropathy in the limb wearing the cuff. When monitoring a patient, examine the extremities of the limb frequently for normal color, warmth and sensitivity. If any abnormality is observed, stop the blood pressure measurements.**

2. To stop auto measuring:

During auto measuring, press  button at any time to stop auto measurement.

3. To start a manual measuring:

■ Press  button to start a manual measuring in the ordinary user working environment.

■ During the idle period of auto measuring process, press  button at any time to start a manual measurement. Then press  button to stop manual measurement and the system continues executing auto-measuring program.



**If you are in doubt about the accuracy of any reading(s), check the patient's vital signs by an alternative method before checking the functioning of the monitor.**



**Warning**



**If liquid is inadvertently splashed on the equipment or its accessories, or may enter the conduit or inside the monitor, contact local Customer Service Center.**

### **Measurement Limitations**

The oscillometry method has some limitations depending on the patient's condition. This measure is based on the regular pulse wave generated by arterial pressure. In the case where the patient condition makes such a detection method difficult, the measured value becomes unreliable and the measuring time increases. The user should be aware that the following conditions will make the measurement unreliable or measurement time extended. In this case, the patient's condition will make the measurement impossible:

- **Patient Movement**

Measurements will be unreliable or can not perform if the patient is moving, shivering or having convulsions. These motions may interfere with the detection of the arterial pressure pulses. In addition, the measurement time will be prolonged.

- **Cardiac Arrhythmia's**

Measurements will be unreliable and may not be possible if the patient's cardiac arrhythmia has caused an irregular heartbeat. The measuring time thus will be prolonged.

- **Heart-lung Machine**

Measurements will not be possible if the patient is connected to a heart-lung machine.

- Pressure Changes

Measurements will be unreliable and may not be possible if the patient's blood pressure is changing rapidly over the period of time during which the arterial pressure pulses are being analyzed to obtain the measurement.

- Severe Shock

If the patient is in severe shock or hypothermia, measurements will be unreliable since reduced blood flow to the peripheries will cause reduced pulsation of the arteries.

- Heart Rate Extremes

Measurements can not be made at a heart rate of less than 40 bpm and greater than 240 bpm.

- Obesity patient

The thick fat layer of body will reduce the measurement accuracy, because the fat that come from the shock of arteries can not access the cuffs due to the damping.

**The following conditions may also cause changes in the blood pressure measurement value**

- After eating (within 1h), or having drinks containing alcohol or caffeine, or after smoking, taking exercises or bathing;
- Using incorrect posture such as standing or lying down, etc.;
- The patient speaks or moves his body during measurement;

- When measuring, the patient is nervous, excited, or in unstable emotion;
- The room temperature rise or fall sharply, or the environment of measurement often changes;
- Measuring in a moving vehicle;
- The position of cuff applied (higher or lower than the heart level);
- Continuous measurement for a long time;

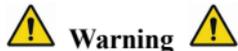
#### 4.4 NIBP Error Messages and Solutions

Display message	Cause	Solution
Low battery	Device battery is low.	Replace the battery. If the problem still exists, please contact us.
Loose cuff	Cuff is not connected correctly.	Reconnect the cuff. If the problem still exists, please contact us.
Atmospheric pressure error	Valve can not be open.	Restart the device. If the problem still exists, please contact us.
Signal is too weak	Object measuring the pulse is too weak or the cuff is loose.	Check the cuff connection, tighten the cuff if it is loose.
It is over the range	Object measuring blood pressure is over the measurement range.	Take another measurement. If the problem still exists, please contact us.

Excessive movement	Movement may result in too much interference in the signal during measuring process.	Be sure to keep still during measuring process.
Overpressure	Cuff pressure is over the scope, ADU 300 mmHg, Newborn: 150mmHg.	Check the cuff to make sure it is not blocked or squeezed.
Saturated signal	Movement or other factors may lead to too big signal amplitude.	Check the connection of air tube to make sure it is not squeezed. Patient should keep quiet and then take a new measurement.
Air leakage	Possible air leakage in the valve or airway	Check the air tube and the cuff.
System failure	Possible failure caused by pump, air valve or pressure sensor.	Please contact us.
It spends too much time	The time for a single measurement exceeds the maximum measurement time (adult: 180s, neonate: 90s).	Check the connection of air tube and tighten the cuff.

## 4.5 Maintenance and Cleaning

**\*Please do obey the precautions and correct operating methods in this user manual. Otherwise, we will not responsible for any fault.**



- Remove the batteries before cleaning the device or peripheral equipment. The accessories and main unit must be separated for cleaning.
- Do not squeeze the rubber tube on the cuff.

### **Cleaning:**

- Do not soak the device and accessories in liquid.
- If any damage or deterioration of the device and accessories is found, please do not use it.
- Do not let water or cleaning agent flow into the sockets to avoid device damage.
- Do not use gasoline, volatile oil, diluent, etc. to wipe the device.

### **Maintenance:**

- Clean the device and accessories regularly. It is recommended to clean them every one month. When it gets dirty, use a dry and soft cloth to wipe. If the device, accessory or the peripheral equipment is very dirty, it is available to dip the soft cloth into water or mild detergent, and wring out, then use the cloth for cleaning. Do not clean the inner parts.

■ The device should be inspected and calibrated periodically (or obey the requirements of the hospital). It is available to inspect in the state specified inspection institution or by professional personal, or you can contact our company.

### **Reusable Blood Pressure Cuff**

The cuff can be sterilized by means of conventional autoclaving, gas, or radiation sterilization in hot air ovens or disinfected by immersion in decontamination solutions, but remember to remove the rubber bag if you use these methods. The cuff should not be dry-cleaned. It can be machine-washed or hand-washed, the latter method may prolong the service life of the cuff. Before washing, remove the latex rubber bag. Allow the cuff to dry thoroughly after washing, then reinsert the rubber bag in the cuff.

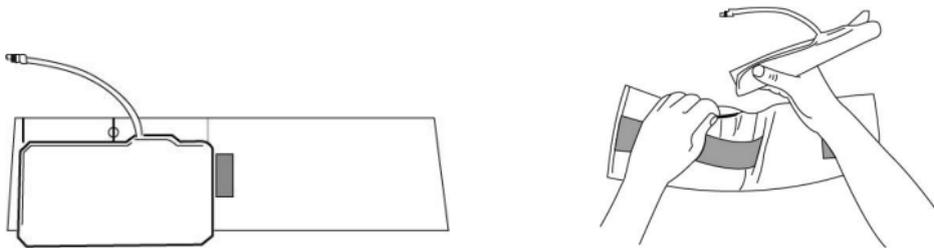


Figure 5.1 Replace the rubber bag

To insert the rubber bag in the cuff, first place the bag on top of the cuff so that the rubber tubes

line up with the large opening on the long side of the cuff. Now roll the bag lengthwise and insert it into the opening on the long side of the cuff. Hold the tubes and the cuff and shake the cuff until the bag is in position. Thread the rubber tubes from inside the cuff, and out through the small hole under the internal flap.

### **Disposable Blood Pressure Cuffs**

Disposable cuff is intended for one-patient use only. Do not use the same cuff on any other patient. No disinfection or high-pressure steam sterilizing to the disposable cuff. Disposable cuff can be cleaned using soap solution to prevent infection.



**Considering environmental protection, the disposable blood pressure cuffs must be recycled or disposed properly.**

### **Storage:**



Do not expose the device in direct sunlight for long time, otherwise the display screen maybe damaged.

The basic performance and safety of the device are not affected by the dust or cotton wool in home environment, while the device shall not be placed where with high temperature, humidity,

dusty or corrosive gases.

Aged cuff may result in inaccurate measurement, please replace the cuff periodically according to the user manual.

To avoid device damage, keep the device out the reach of children and pets.

Avoid the device close to extreme high temperature such as fireplace, otherwise the device performance may be affected.

Do not store the device with chemical medicine or corrosive gas.

Do not place the device where there is water.

Do not place the device where with slope, vibration or impact

#### **4.6 Transportation and Storage**

❖ The packaged device can be transport by general vehicle or according to the order contract. Do not transport the device mixed with toxic, harmful or corrosive materials.

❖ The device after packaged should be stored in a well ventilated room without any corrosive gas, temperature range:  $-20^{\circ}\text{C}\sim+55^{\circ}\text{C}$ , relative humidity no more than 95%.

## 4.7 Key and Symbols

Your device may not contain all the following symbols.

Signal	Description	Signal	Description
	Attention! Please refer to the accompanying document (the user manual).		Attention! Please refer to the accompanying document (the user manual).
SYS	Systolic pressure	DIA	Diastolic pressure
MAP	Mean blood pressure	PR	Pulse rate (bpm)
SN	Serial number	EMC	Electromagnetic compatibility
IPXX	The degree of protection against ingress of water	P/N	Material code of manufacturer
ADU	Adult	NEO	Neonatal
PED	Pediatric	INFO	Information
ABPM	Ambulatory Blood Pressure Monitor		Type BF defibrillator proofed applied parts

	Class II equipment		Silence
	Close the prompt sound indication		Open the prompt sound indication
	Batch code		Use by date
	This way up		Fragile, handle with care
	Keep dry		Storage atmospheric pressure limitation
	Storage temperature limitation		Storage humidity limitation
	Manufacturer		Date of manufacture

	Batteries Power		Pulse rate (bpm)
	1.No finger inserted to SpO2 probe 2.No NIBP data to review 3.An indicator of signal inadequacy		1.No Pulse rate 2.An indicator of signal inadequacy
	Waste disposal mark, this symbol indicates that the waste of electrical and electronic equipment can not be disposed as an unclassified municipal waste and must be recovered separately.		This item is compliant with Medical Device Directive 93/42/EEC of June 14, 1993,a directive of the European Economic Community.
	Recyclable		European Representative
	Free of natural rubber latex		

## **Chapter5 Software installation**

### **5.1 Requirements of Hardware**

Processor: Intel Celeron 2.5G or more

Operation System:Windows XP/Win7/Win8

EMS memory: 1GB and more

Hard Disk: 250G or more

Display: 17 inch or more

CD-ROM

USB: 2 or more

Resolution of printer: 600 DPI

### **5.2 Software Installation**

1. Place the CD in the CD-ROM compartment located on your computer.
2. If Auto Play for CDs is enabled, please install the software following the instructions below:  
Under Windows operating system:  
Open Windows Explorer.  
Click on the root CD-ROM directory.  
Double click the file "ABPM\_Setup.exe".  
Operate according to the instructions in the screen.

## Chapter 6 Software Functions

### 6.1 User Register

Double-click the software icon, then the dialog box shown as below appears.

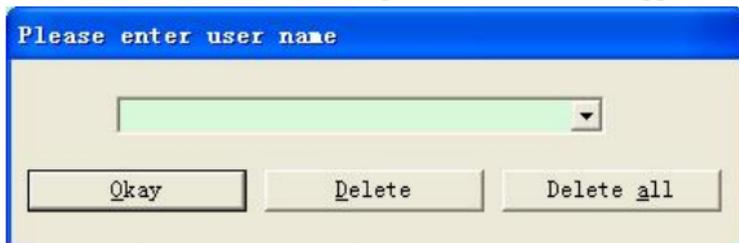


Figure 6.1.1 User Register

Enter the user name, click "Okay", then the "Configuration Set" dialog box shown as Figure 6.1.2 will appear. Click "Delete" to delete the configuration information of the user.

"Delete all" is used to delete the configuration information of all users.

If you are a new user, then the following dialog box will appear.

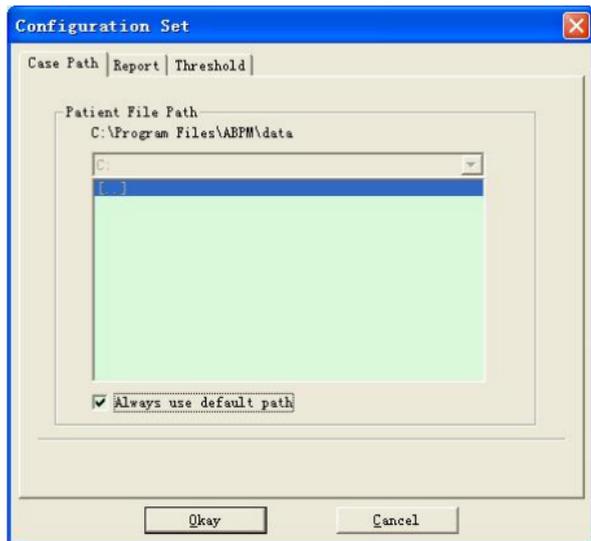


Figure 6.1.2 Configuration Set

"Case path": select the default storage path for case, after getting the data from the device, the case file will be saved in this path.

If you select "Always use default path", then the case file will be automatically saved in the installation path.

## 6.2 Main Interface

The device will enter to the main interface(shown as below) after setting the configuration information.

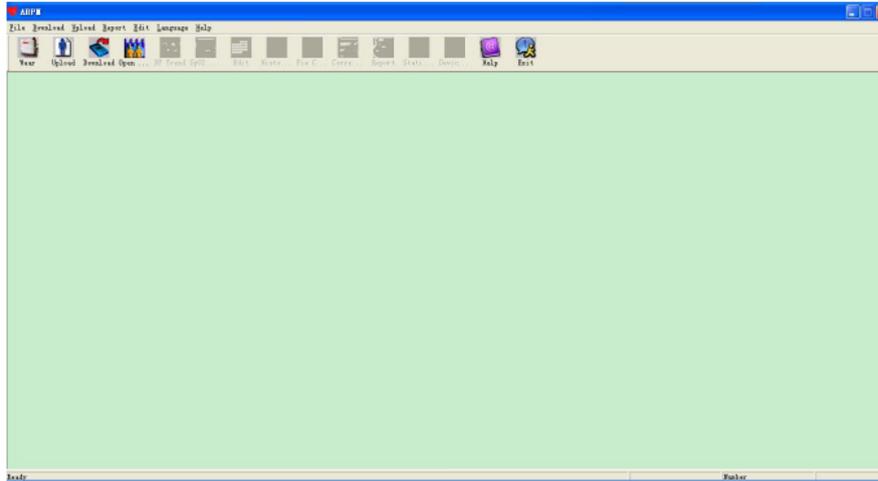


Figure 6.2 Main Interface

### 6.3 Wear

After clicking the shortcut key  , the following figure appears. Before using the device, please read "Matters need" carefully, and wear the device according the following figure.

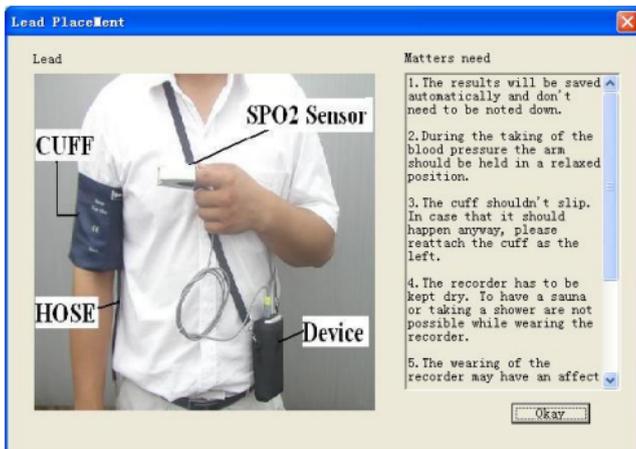
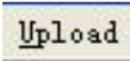
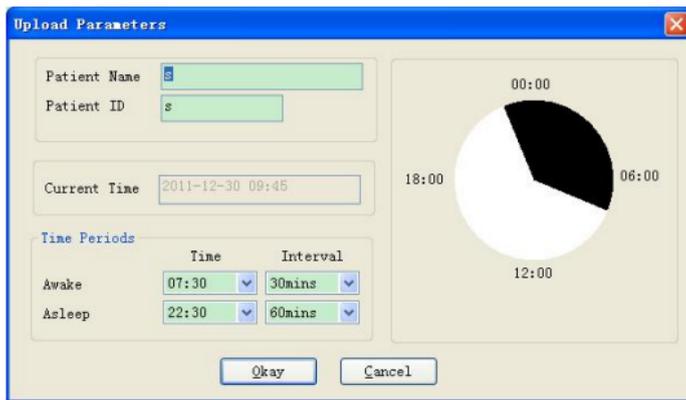


Figure 6.3 Wear

## 6.4 Setting for Collection Plan

Click shortcut key , or click menu bar  item, and the " Upload parameters " dialog box will appear:



Time Periods	Time	Interval
Awake	07:30	30mins
Asleep	22:30	60mins

Figure 6.4 Set Collection Parameter

As above figure, the doctor could set parameter according to the patient status and diagnosis

requirement, then the monitor could finish the collection according to the setting. Parameter explanation is as follows:

**Patient Name:** the patient's name

**Patient ID:** the patient's ID number. It is used for marking patient, and it is exclusive in order to avoid homonymy patient arisen state

**Current Time:** Current system display time

**Time Periods:**

**Awake Time:** the patient is awake state

**Asleep Time:** the patient is asleep state

**Interval:** collection Interval. In order to reduce the effect to the patient sleep, the asleep collection interval should be longer.

For example as above figure: the awake time area is 7:30-22:30, and the asleep time area is 22:30-tomorrow 7:30. The awake collection interval is 30 minutes, and the asleep collection interval is 60 minutes.

The asleep time area and awake time area will be displayed at the right side.

When the parameter setting is finished, click " Okay " to upload the project to the monitor.

## 6.5 Data Download

Before you download the measurement data from device, please make sure that:

1. The device is correctly connected to the computer properly.
2. The device is turned on.
3. Disconnect the device from patient before connecting it to the computer.

The downloaded patient data will be saved in case storage path set. If you want to change the storage path, select "Set file path", the dialogue box (Figure 6.1.2) will appear, then you can change the path.



Click the shortcut key **Download** or "Download" from menu to select the data in which status is to be obtained, then start to download the data.

## 6.6 Open Data File

Click "Open Data" to open the case interface shown as below:

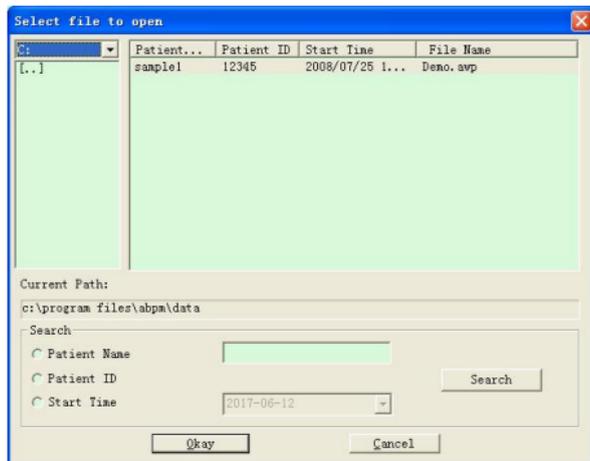


Figure 6.6 Case Selection

In this interface, you can operate the drive and folder selection on top left part to load the specified disk and folder content, if case file exists in this folder, the basic information of these case files will be displayed in the form of list, contents including: patient name, patient ID, start

time and file name. Click to select the case file to be opened, then click "Okay" to open and load the case file information.

When there are many case data, select one inquiry item, enter the key information, then click "Search" to query.

## 6.7 Delete Data File

If you feel some patient data are not necessary, you can delete them. Select "Delete Data" from menu to enter its sub-menu shown as below.

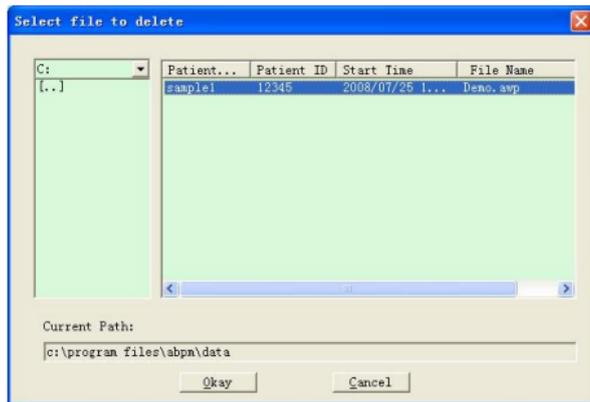


Figure 6.7 Data File Delete

Many files can be deleted at the same time. Push “Ctrl” and click the file you want to delete at the same time, click "Okay", to delete the case file selected. Click "Cancel" to cancel deleting.

## 6.8 Data File Backup

The software has the function of case backup. Select "Copy data" from menu, then the following figure will appear.

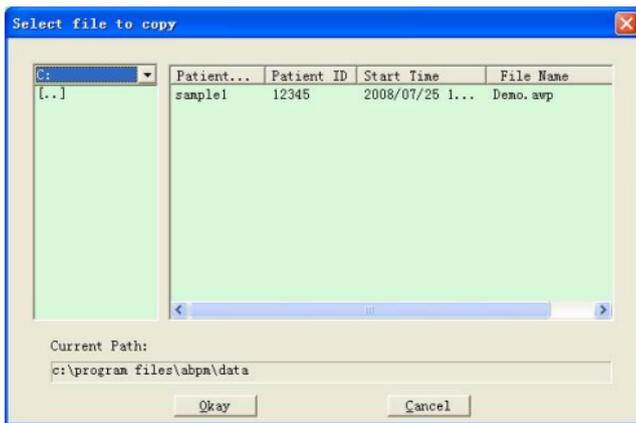


Figure 6.8.1 Data File Copy

After selecting files, click "Okay", then a dialog box which is used to set the storage files

of backup files appears. After setting, click "Okay" to save. The destination directory interface is shown as below:

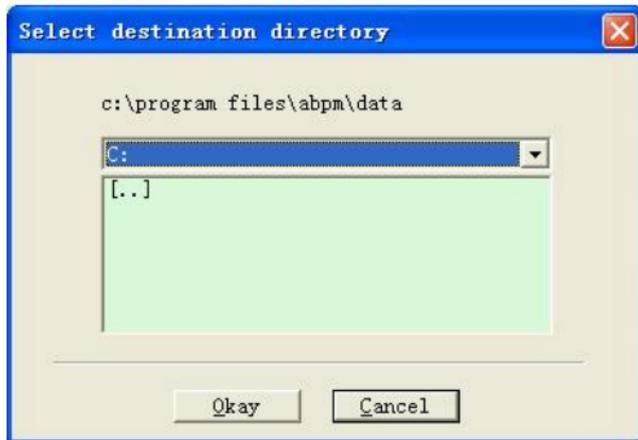
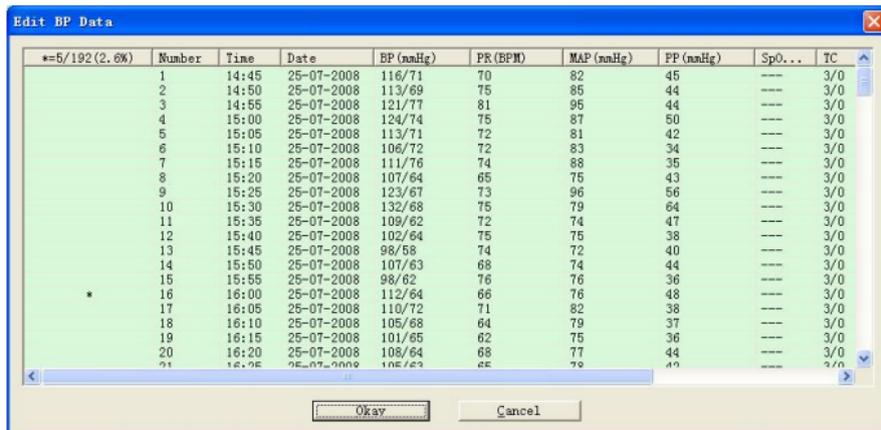


Figure 6.8.2 Backup Path Settings

## 6.9 Edit IP Data

After opening the case file, blood pressure data can be edited. Click the shortcut key  or

select "Bp data" from menu to enter the interface shown as below:



The screenshot shows a dialog box titled "Edit BP Data" with a table of data. The table has columns for Number, Time, Date, BP (mmHg), PR (BPM), MAP (mmHg), PP (mmHg), SpO... (likely SpO2), and TC. The data rows show a sequence of readings from 14:45 to 16:20 on 25-07-2008. At the bottom of the dialog are "Okay" and "Cancel" buttons.

*=5/192 (2.6%)	Number	Time	Date	BP (mmHg)	PR (BPM)	MAP (mmHg)	PP (mmHg)	SpO...	TC
	1	14:45	25-07-2008	116/71	70	82	45	---	3/0
	2	14:50	25-07-2008	113/69	75	85	44	---	3/0
	3	14:55	25-07-2008	121/77	81	96	44	---	3/0
	4	15:00	25-07-2008	124/74	75	87	50	---	3/0
	5	15:05	25-07-2008	113/71	72	81	42	---	3/0
	6	15:10	25-07-2008	106/72	72	83	34	---	3/0
	7	15:15	25-07-2008	111/76	74	88	35	---	3/0
	8	15:20	25-07-2008	107/64	65	75	43	---	3/0
	9	15:25	25-07-2008	123/67	73	96	56	---	3/0
	10	15:30	25-07-2008	132/68	75	79	64	---	3/0
	11	15:35	25-07-2008	109/62	72	74	47	---	3/0
	12	15:40	25-07-2008	102/64	75	75	38	---	3/0
	13	15:45	25-07-2008	98/58	74	72	40	---	3/0
	14	15:50	25-07-2008	107/63	68	74	44	---	3/0
	15	15:55	25-07-2008	98/62	76	76	36	---	3/0
*	16	16:00	25-07-2008	112/64	66	76	48	---	3/0
	17	16:05	25-07-2008	110/72	71	82	38	---	3/0
	18	16:10	25-07-2008	105/68	64	79	37	---	3/0
	19	16:15	25-07-2008	101/65	62	75	36	---	3/0
	20	16:20	25-07-2008	108/64	68	77	44	---	3/0
	21	16:25	25-07-2008	105/63	65	78	49	---	3/0

Figure 6.9 Data Edit Interface

All the BP readings are shown in the above dialog box.

\*=5/192(2.6 %): 192 represents the data sum, 5 represents the data amount deleted, 2.6 % is the percentage of data deleted in all collection data.

Number: stands for data collection serial number.

Time: stands for collection time.

Date: stands for collection date.

BP(mmHg): systolic pressure/diastolic pressure, unit is mmHg.

PR: pulse rate, unit is BPM

MAP: mean pressure, unit is mmHg.

PP: pressure difference between systolic pressure and diastolic pressure, unit is mmHg.

SpO<sub>2</sub>(%): oxygen saturation, unit is %.

TC: error code /measurement mode(refer to chapter 4)

Comment: add comment information to the BP data.

These data can also be performed exclusion operation. The symbol "\*" indicates to delete the data(not be displayed in the trend graph, and not be recorded in statistics). You can click the location area of the first column to add or delete "\*". And in the comment field, you can annotate the data, and the comment information will be displayed in the trend graph and report.

## 6.10 BP Trend Graph

After selecting case file, the BP trend curve will be displayed in the screen automatically.



Click the shortcut key **Bp trend** to its sub-menu. Two graph types: color filling trend and

dotted line trend. The trend graph is shown as below.

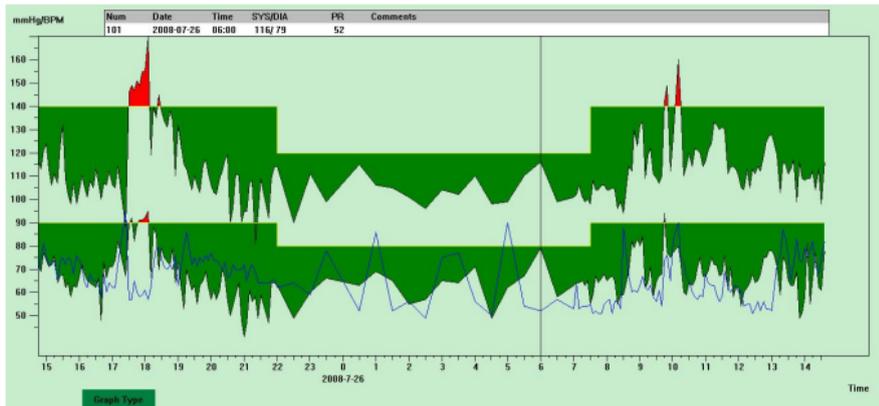


Figure 6.10.1 Color Filling Trend Ggraph

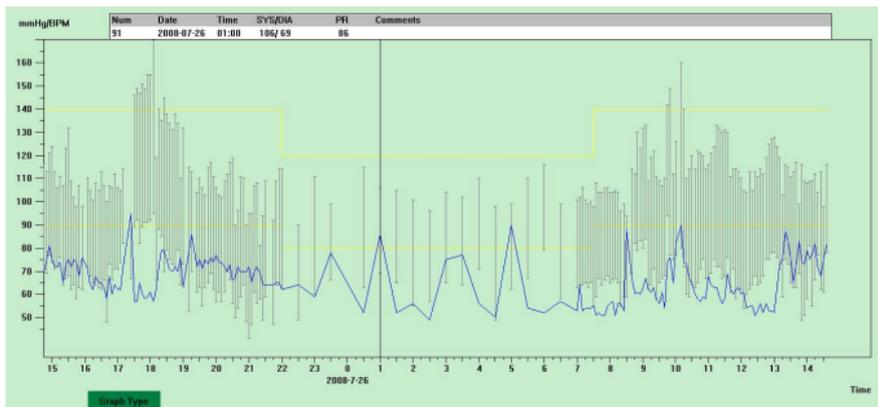


Figure 6.10.2 Dotted Line Trend Graph

You can switch the two trend graph types by "Graph Type" button in the bottom of the software interface. When you move the mouse on the trend area, the detail data information on this location will display in the top of the trend area, including the data serial number, collection time and collection date, high/low blood pressure value, pulse rate, comment, etc. Press mouse' left button to delete or add the data point to be shown.

## 6.11 Display of Statistics Information

Press the shortcut key  or select "Report" from menu to enter its sub-menu shown as below.



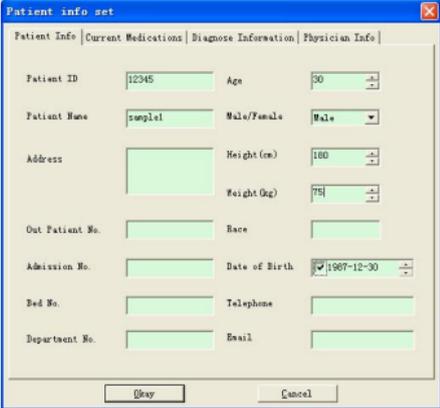
Figure 6.11 BP Statistics Information

The upper half part of the figure shows the average of blood pressure data and the measurement

number under "Awake" and "Asleep" state. The lower part shows the percentage of warning value data, 140/90, 120/80 represent blood pressure warning value of the systolic and diastolic pressure under "Awake" and "Asleep" state, the unit is mmHg.

## 6.12 Patient Information Settings

Select "Patient Data" from the menu to enter its sub-menu shown as below. Patient information including: patient information, current medications, diagnose information and physician information.



The screenshot shows a dialog box titled "Patient info set" with a blue title bar and a close button (X) in the top right corner. The dialog has four tabs: "Patient Info", "Current Medications", "Diagnose Information", and "Physician Info". The "Patient Info" tab is selected. The form contains the following fields:

Patient ID	<input type="text" value="12345"/>	Age	<input type="text" value="30"/>
Patient Name	<input type="text" value="sample1"/>	Male/Female	<input type="text" value="Male"/>
Address	<input type="text"/>	Height (cm)	<input type="text" value="180"/>
		Height (kg)	<input type="text" value="75"/>
Out Patient No.	<input type="text"/>	Race	<input type="text"/>
Admission No.	<input type="text"/>	Date of Birth	<input type="text" value="1987-12-30"/>
Bed No.	<input type="text"/>	Telephone	<input type="text"/>
Department No.	<input type="text"/>	Email	<input type="text"/>

At the bottom of the dialog, there are two buttons: "Okay" and "Cancel".

Figure 6.12 Edit Patient's Information

Recent medication information of patient can be entered in "Current Medications" column. Blood pressure data description and diagnosis information can be entered in "Diagnose Information" column.

Doctor name and doctor advice can be entered in "Physician Info" column.

### 6.13 Sleep Time Setting

Awake and Asleep time can be set by manual mode, after setting, the software will calculate the data again under "Awake" and "Asleep" state, then update the trend graph and calculate the statistic data automatically. The interface shown as below will appear after selecting "Sleep Period" from menu.

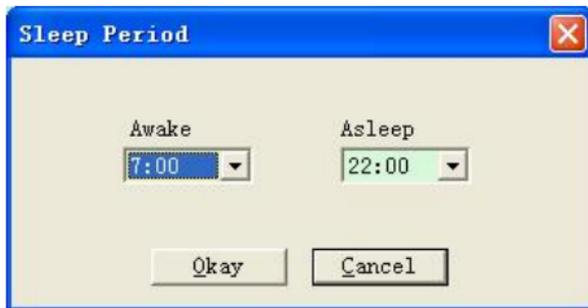


Figure 6.13 Sleep Time Setting

## 6.14 BP Threshold Setting

The BP threshold can be changed by manual mode, after changing, the corresponding trend graph and analysis data will be automatically renewed. Select "Threshold" to enter its sub-menu shown as below.

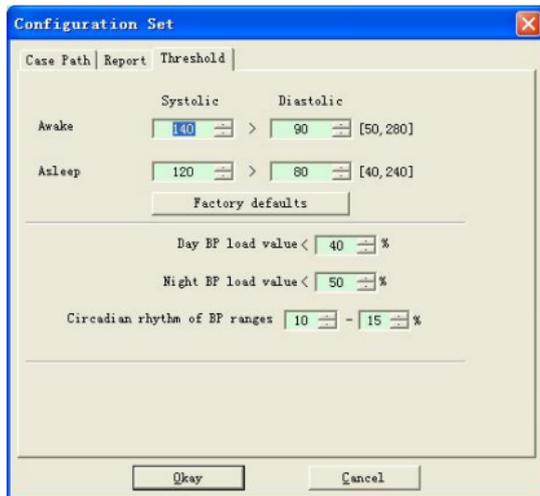


Figure 6.14 BP Threshold Setting

The default recommended thresholds for calculating Blood Pressure Load are 140/90 for wake

periods and 120/80 for sleep periods. These are the default values when you select the Factory Defaults button.

## 6.15 Histogram



Press the shortcut key **Histo...**, the following interface will appear.

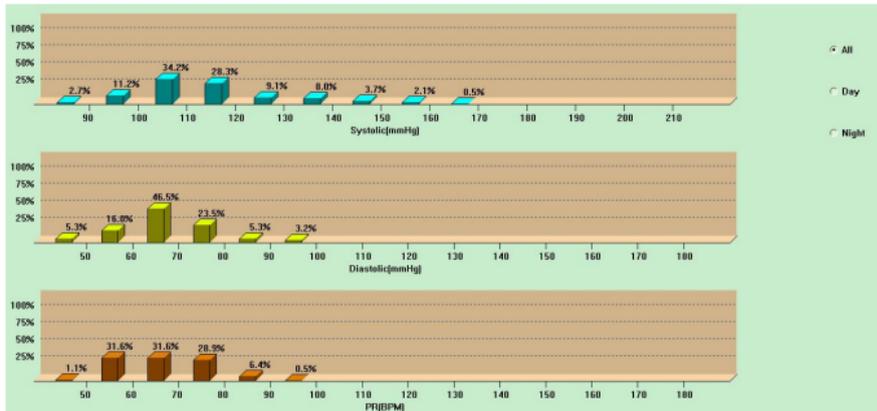


Figure 6.15 Histogram

"All", "Day" and "Night" can respectively display the analysis values in each period.

## 6.16 Pie Chart



Press the shortcut key `Pie c...`, the following interface will appear:

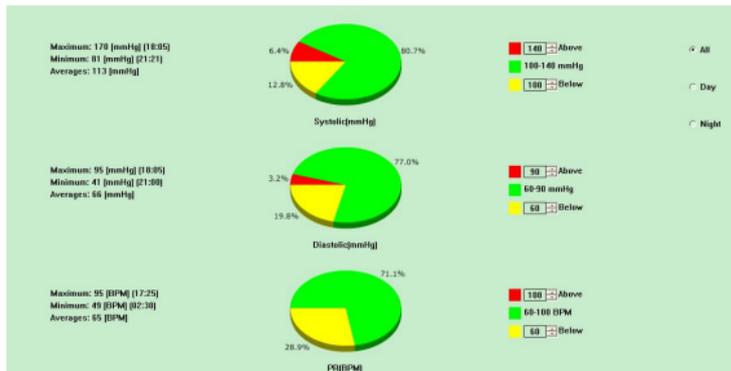


Figure 6.16 Pie Chart

The pie chart interface is divided into four regions, from left to right, the first region is value display area which displays the Maximum, Minimum and Average values among the measurement values, the second region is pie chart display area, the third is the setting area for

pie chart color and values, and the last is the time display area, it has three options: "All", "Day" and "Night", which can respectively display the analysis values in each period.

## 6.17 Correlation Line



Press the shortcut key **Corre...**, the following interface will appear:

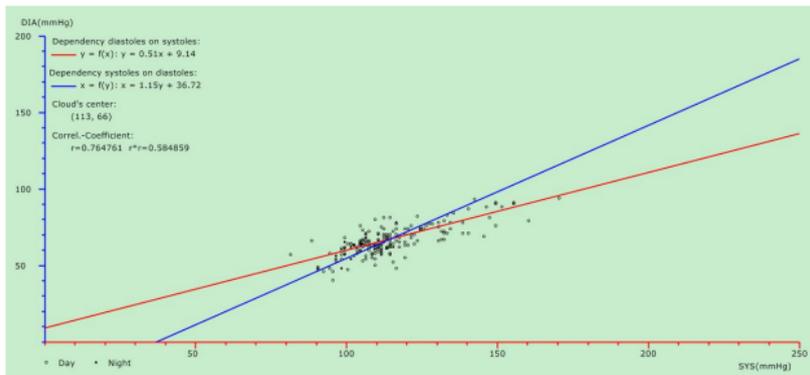


Figure 6.17 Correlation Line

The horizontal axis is the systolic pressure axis, the vertical axis is the diastolic pressure axis. Red represents the dependence for diastolic pressure to systolic pressure; blue represents the

dependence for systolic pressure to diastolic pressure. The hollow circle is the BP value measured in the day, and the solid circle is the BP value measured at night.

## 6.18 Print Report

After editing the BP data and diagnosis information, click "Report", the software will create a series of diagnosis reports, you can select all pages or some of them for printing. Select "Configure Report" in "Report", then the following figure will appear.

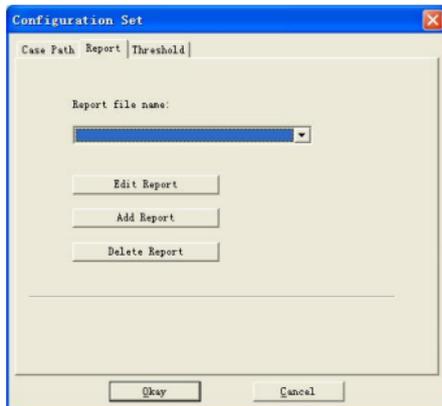


Figure 6.18.1 Configure Report

You can select a report configured for printing, or click "Edit Report" to edit the selected report.

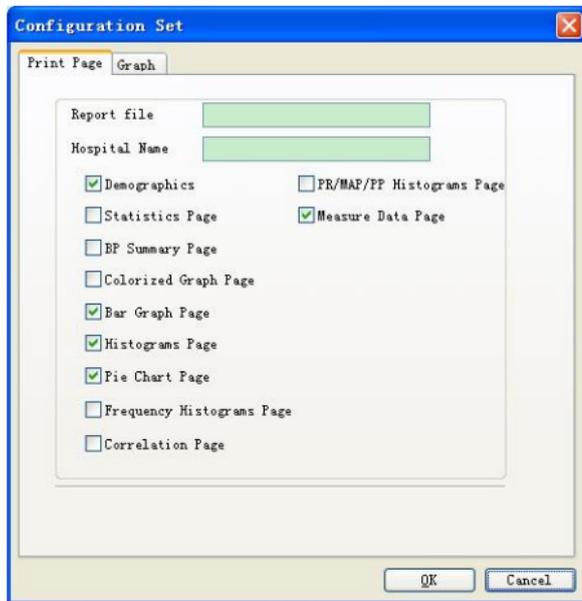


Figure 6.18.2 Edit Report

Click "Add Report" to add a new report. If you don't need the current report, you can also click

"Delete Report" to delete it.



Click the shortcut key  or select "Report" from menu to preview the report, then select "Print" to print the report.

### 6.19 Help



Click the shortcut key  to its sub-menu, which gives a brief description for each program function. In addition, you will find "Help" button in each operation interface, click it to check the description for this function, which is convenient for you to quick know the use of software.

## Specification

<b>Name</b>	Ambulatory Blood Pressure Monitor	
<b>The degree of protection against ingress of water</b>	IP22	
<b>Display</b>	2.4" color LCD Display	
<b>Operating mode</b>	Continuous operating	
<b>NIBP Specifications</b>		
Measurement Method	Oscillometric method	
Working modes	Automatic	
Cuff pressure range	adult	0~297 mmHg(0~39.6 kPa)
	pediatric	0~235 mmHg(0~31.3 kPa)
	neonatal	0~147 mmHg(0~19.6 kPa)
Blood pressure measurement Range	adult	SYS: 40~270 mmHg(5.3~36.0 kPa) DIA: 10~215 mmHg(1.3~28.7 kPa)
	pediatric	SYS: 40~200 mmHg(5.3~26.7 kPa) DIA: 10~150 mmHg(1.3~20.0 kPa)
	neonatal	SYS: 40~135 mmHg(5.3~18.0 kPa) DIA: 10~100 mmHg(1.3~13.3 kPa)
Pulse measurement	40~240/min	

range		
Inflation	adult	160mmHg(21.33 kPa)
	pediatric	120mmHg(16kPa)
	neonatal	70mmHg(9.33 kPa)
Prompt Range	adult mode	SYS PROMPT: 40~270 mmHg(5.3~36.0 kPa) DIA PROMPT: 10~215 mmHg(1.3~28.7 kPa)
	pediatric mode	SYS PROMPT: 40~200 mmHg(5.3~26.7 kPa) DIA PROMPT: 10~150 mmHg(1.3~20.0 kPa)
	neonatal mode	SYS PROMPT: 40~135 mmHg(5.3~18.0 kPa) DIA PROMPT: 10~100 mmHg(1.3~13.3 kPa)
Overpressure protect	adult mode	$297 \pm 3$ mmHg ( $39.6 \pm 0.4$ kPa)
	pediatric mode	$240 \pm 5$ mmHg ( $32 \pm 0.67$ kPa)
	neonatal mode	$147 \pm 3$ mmHg ( $19.6 \pm 0.4$ kPa)
<b>Resolution</b>		
Pressure	1 mmHg (0.133 kPa)	
Pulse Rate	$\pm 5$ %	
<b>Measurement Accuracy</b>		
Cuff	Pressure	Static pressure: $\pm 3$ mmHg( $\pm 0.4$ kPa)

<b>Accuracy</b>	
<b>Error</b>	<p>The BP value measured by the device is equivalent with the measurement value of Stethoscopy, perform clinical verification in accordance with the requirements in ISO 81060-2: 2013, whose error meets the followings:</p> <p>Maximum mean error: <math>\pm 5</math> mmHg  Maximum Standard deviation: 8 mmHg</p>
<b>Operating temperature/humidity</b>	<p>+5 °C~40 °C                      15 %RH~85 %RH(Non-condensing)</p>
<b>Transport</b>	<p>Transport by general vehicle or according to the order contract, avoid pounded, shake and splash by rain and snow in transportation.</p>
<b>Storage</b>	<p>Temperature: -20 °C~+55 °C; Relative humidity: <math>\leq 95</math> %; No corrosive gas and drafty.</p>
<b>Atmospheric pressure</b>	<p>700 hPa~1060 hPa</p>
<b>Power Supply</b>	<p>DC 3 V</p>
<b>Battery service life</b>	<p>When the temperature is 23 °C, limb circumference is 270 mm, the measured blood pressure is normal, 2 "AA" alkaline batteries can</p>

	be used about 150 times.
<b>Rated Power</b>	≤ 3.0 VA
<b>Dimensions</b>	128(L)*69(W)*36 mm(H)
<b>Unit Weight</b>	240 gram(without batteries)
<b>Safety classification</b>	Internally powered equipment Type BF defibrillation-proof applied par
<b>Service life</b>	The service life of the device is five years or 10000 times of BP measurement.
<b>Date of manufacturer</b>	See the label
<b>Accessories</b>	Standard Configure: Adult Cuff: limb circumference 25-35 cm (middle of upper arm) Software CD, User Manual, USB data line, BP extending tube, A pouch. Separate Sale: Pediatric Cuff: limb circumference 10-19 cm (middle of upper arm) Neonatal Cuff: limb circumference 6-11 cm (middle of upper arm)

## Appendix

### Guidance and manufacturer's declaration – electromagnetic emissions- for all EQUIPMENT and SYSTEMS

<b>Guidance and manufacturer's declaration – electromagnetic emission</b>		
The device is intended for use in the electromagnetic environment specified below. The customer of the user of the <i>device</i> should assure that it is used in such and environment.		
<b>Emission test</b>	<b>Compliance</b>	<b>Electromagnetic environment – guidance</b>
RF emissions CISPR 11	Group 1	The <i>device</i> uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emission CISPR 11	Class B	The <i>device</i> is suitable for use in all establishments other than domestic and those directly connected to a low voltage power supply network which supplies buildings used for domestic purposes.

**Guidance and manufacturer's declaration – electromagnetic immunity –  
for all EQUIPMENT and SYSTEMS**

<b>Guidance and manufacturer's declaration – electromagnetic immunity</b>			
The <i>device</i> is intended for use in the electromagnetic environment specified below. The customer or the user of <i>device</i> should assure that it is used in such an environment.			
<b>Immunity test</b>	<b>IEC60601 test level</b>	<b>Compliance level</b>	<b>Electromagnetic environment - guidance</b>
Electrostatic discharge (ESD) IEC61000-4-2	±6 kV contact ±8 kV air	±6 kV contact ±8 kV air	Floors should be wood, concrete or ceramic tile. If floor are covered with synthetic material, the relative humidity should be at least 30%.
Power frequency (50/60Hz) magnetic field IEC61000-4-8	3A/m	3A/m	Mains power quality should be that of a typical commercial or hospital environment.
NOTE: $U_T$ is the a.c. mains voltage prior to application of the test level.			

**Guidance and manufacturer's declaration – electromagnetic immunity –  
for EQUIPMENT and SYSTEMS that are not LIFE-SUPPORTING**

<b>Guidance and manufacturer's declaration – electromagnetic immunity</b>			
The device is intended for use in the electromagnetic environment specified below. The customer or the user of device should assure that it is used in such an environment.			
<b>Immunity test</b>	<b>IEC60601 test level</b>	<b>Compliance level</b>	<b>Electromagnetic environment - guidance</b>
Radiated RF	3 V/m 80 MHz to	3 V/m	<p>Portable and mobile RF communications equipment should be used no closer to any part of the device, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.</p> <p>Recommended separation distance</p> $d = \left[ \frac{3.5}{E_1} \right] \sqrt{P}$ <p align="right">80 MHz to 800 MHz</p>

IEC 61000-4- 3	2.5 GHz		$d = \left[ \frac{7}{E_1} \right] \sqrt{P}$ <p style="text-align: right;">800 MHz to 2.5 GHz</p> <p>Where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in metres (m). Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, a should be less than the compliance level in each frequency range.b</p> <p>Interference may occur in the vicinity of equipment marked with the following symbol:</p> 
NOTE 1	At 80 MHz and 800 MHz, the higher frequency range applies.		
NOTE 2	These guidelines may not apply in all situations. Electromagnetic propagation is		

affected by absorption and reflection from structures, objects and people.

a Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the device is used exceeds the applicable RF compliance level above, the device should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as reorienting or relocating the device.

**Recommended separation distances between portable and mobile RF communications equipment and the EQUIPMENT or SYSTEM – for EQUIPMENT or SYSTEM that are not LIFE-SUPPORTING**

**Recommended separation distances between portable and mobile RF communications equipment and the device**

The *device* is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the *device* can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the *device* as recommended below,

according to the maximum output power of the communications equipment.

Rated maximum output power of transmitter (W)	Separation distance according to frequency of transmitter (m)	
	80 MHz to 800 MHz	800 MHz to 2.5 GHz
	$d = \left[ \frac{3.5}{E_1} \right] \sqrt{P}$	$d = \left[ \frac{7}{E_1} \right] \sqrt{P}$
0.01	0.12	0.23
0.1	0.37	0.74
1	1.17	2.33
10	3.69	7.38
100	11.67	23.33

For transmitters rated at a maximum output power not listed above, the recommended separation distance  $d$  in metres (m) can be estimated using the equation applicable to the frequency of the transmitter, where  $P$  is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer.

NOTE 1 At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.

NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

## **Warning**

- Active medical devices are subject to special EMC precautions and they must be installed and used in accordance with these guidelines.
- Electromagnetic fields can affect the performance of the device, so other equipment used near the equipment must meet the appropriate EMC requirements. Mobile phones, X-rays, or MRI devices are possible interference sources, as they emit high-intensity electromagnetic radiation.
- The use of ACCESSORIES, transducers and cables other than those specified, with the exception of transducers and cables sold by the MANUFACTURER of the the device as replacement parts for internal components, may result in increased EMISSIONS or decreased IMMUNITY of the ME EQUIPMENT or ME SYSTEM.
- The device should not be used when they are close to or stacked with other equipment, if necessary, please observe and verify that they can operate normally in the configurations.
- Devices or systems may still be interfered by other equipment, even if other equipment meets the requirements of the corresponding national standard.
- The device requires special precautions for electromagnetic compatibility (EMC) and requires qualified personnel to install and use in accordance with the EMC information provided below.
- The device should not contact the pins of connectors marked with an ESD warning symbol, unless electrostatic discharge precautions are used, the device should not connect to these

connectors.

- In order to avoid the accumulation of electrostatic charge, it is recommended to store, maintain and use the equipment at a relative humidity of 30% or more. The floor should be covered with ESD dissipated carpets or similar materials. In the use of the components, non-synthetic clothing should be worn.

- In order to prevent electrostatic discharging to the ESD-sensitive parts of the device, the personnel should contact the metal frame of the components or the large metal objects near the device. When using the device, especially when it is possible to contact the ESD-sensitive parts of the device, the operator should wear a grounded bracelet designed for ESD-sensitive devices. For more information on proper use, please refer to the instructions provided with the bracelet.

- All potential users are advised to understand the ESD warning symbols and receive training on ESD precautions.

- The most basic content of the ESD precautionary procedure training should include an introduction to electrostatic charge physics, voltage level in the conventional case, and damage to the electronic components when the operator with electrostatic charge contacts them. In addition, the methods for preventing electrostatic buildup, and the manner and reasons for the release of human body static electricity to the ground or equipment frame or the use of a bracelet to connect the human body to the equipment or the ground before establishing the connection should be described.