Owner's Manual

Arm-type Fully Automatic
Digital Blood Pressure Monitor
Model DBP-6293L

Safety Notice

Thank you for purchasing the DBP-6293L Blood Pressure Monitor. The unit has been constructed using reliable circuitry and durable materials. Used properly, this unit will provide years of satisfactory use.

Indications for use: Arm-type Fully Automatic Digital Blood Pressure monitors are intended to measure blood pressure (systolic and diastolic) and pulse rate of adults and adolescents over 12 years of age.

All functions can be used safely and values can be read out in one LCD DISPLAY. Measurement position is on adult upper arm only. The PATIENT is an intended OPERATOR.

Blood pressure measurement determined with this device are equivalent to those obtained by a trained observer using the cuff/stethoscope auscultation method, within the limits prescribed by the Recognized Consensus Standard (IEC 81060-2-30) for electronic sphygmomanometers.

Precautions to Ensure Safe, Reliable Operation
1. Do not drop the unit. Protect it from sudden jars or shocks.
2. Do not insert foreign objects into any openings.
3. Do not attempt to disassemble the unit.
4. Do not crush the pressure cuff.
5. If the unit has been stored at temperatures below 0 ℃, leave it in a warm place for about 15 minutes before using it. Otherwise, the cuff may not inflate properly.
6. If the unit has been stored at temperatures above 40 ℃, leave it in a cool place for about 15 minutes before using it. Otherwise, the cuff may not inflate properly.
7. Do not store the unit in direct sunlight, high humidity or dust.
8. To avoid any possibility of accidental strangulation, keep this unit away from children and do not drape tubing around your neck.
9. Ensure that children do not use the instrument unsupervised; some parts are small enough to be swallowed.
10. Some may get a skin irritation from the cuff taking frequent readings over the course of the day, but this irritation typically goes away on its own after the monitor is removed.

Contact Information
The lay operator or lay responsible or ganization should contact the manufacturer or the representative of manufacturer:
- for assistance, if needed, in setting up, using or maintaining the product, or
- to report unexpected operation or events.
Manufactured by JOYTECH Healthcare Co., Ltd.
No. 365, Wuzhou Road, Yuhang Economic Development Zone, Hangzhou City, 311100 Zhejiang, China
Email: info@sejoy.com
Telephone: +86-571-81957767
Fax: +86-571-81957750

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Safety Notice

Important Instructions Before Use

1. Do not confuse self-monitoring with self-diagnosis. Blood pressure measurements should only be interpreted by a health professional who is familiar with your medical history.

2. Contact your physician if test results regularly indicate abnormal readings.

3. If you are taking medication, consult with your physician to determine the most appropriate time to measure your blood pressure. NEVER change a prescribed medication without first consulting with your physician.

4. Individuals with serious circulation problems may experience discomfort. Consult your physician prior to use.

5. For persons with irregular or unstable circulation resulting from diabetes, liver disease, arteriosclerosis or other medical conditions, there may be variations in blood pressure values measured at the wrist versus at the upper arm. Monitoring the trends in your blood pressure taken at either the arm or the wrist is nevertheless useful and important.

6. People suffering from vascular constriction, liver disorders or diabetes, people with cardiac pacemakers or a weak pulse, and women who are pregnant should consult their physician before measuring their blood pressure themselves. Different values may be obtained due to their condition.

7. People suffering from arrhythmias such as atrial or ventricular premature beats or atrial fibrillation only use this blood pressure monitor in consultation with your doctor. In certain cases oscillometric measurement method can produce incorrect readings.

8. Too frequent measurements can cause injury to the patient due to blood flow interference.

9. The cuff should not be applied over a wound as this can cause further injury.

10. DO NOT attach the cuff to a limb being used for IV infusions or any other intravascular access, therapy or an arterio-venous (A-V) shunt. The cuff inflation can temporarily block blood flow, potentially causing harm to the patient.

11. The cuff should not be placed on the arm on the side of a mastectomy. In the case of a double mastectomy use the side of the least dominant arm.

12. Pressurization of the cuff can temporarily cause loss of function of simultaneously used monitoring equipment on the same limb.

13. A compressed or kinked connection hose may cause continuous cuff pressure resulting in blood flow interference and potentially harmful injury to the patient.

14. Check that operation of the unit does not result in prolonged impairment of the circulation of the patient.

15. Product is designed for its intended use only. Do not misuse in any way.

16. Product is not intended for infants or individuals who cannot express their intentions.

17. Prolonged over-inflation of the bladder may cause edema of your arm.

18. Do not disassemble the unit or arm cuff. Do not attempt to repair.

19. Use only the approved arm cuff for this unit. Use of other arm cuffs may result in incorrect measurement results.

20. The system might produce incorrect readings if stored or used outside the manufacturer’s specified temperature and humidity ranges. Make sure to store the blood pressure monitor, children, pets and pets are outside of accessible range.

21. Do not use the device near strong electrical or electromagnetic fields generated by cell phones or other devices, they may cause incorrect readings and interference or become interference source to the device.

22. This product contains lithium battery, do not discard.

Safety Notice

Federla Commulation Commission (FCC) Interference Statement

1. This device complies with part 15 of the FCC Rules. Operation is subject to the condition that this device does not cause harmful interference.

2. This device is verified to comply with part 15 of the FCC Rules for use with cable television service.

3. This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
   (1) This device may not cause harmful interference, and
   (2) this device must accept any interference received, including interference that may cause undesired operation. Please note that changes or modifications not expressly approved by the party responsible for compliance could void the user’s authority to operate the equipment.

4. This equipment has been tested and found to comply with the limits for a Class B digital device, pursuant to part 15 of the FCC Rules. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This equipment generates, uses and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation. If this equipment does cause harmful interference to radio or television reception, which can be determined by turning the equipment off and on, the user is encouraged to try to correct the interference by one or more of the following measures:
   — Retort or relocate the receiving antenna.
   — Increase the separation between the equipment and receiver.
   — Connect the equipment into an outlet on a circuit different from that to which the receiver is connected.
   — Consult the dealer or an experienced radio/TV technician for help.

5. This equipment complies with radio frequency exposure limits set forth by the FCC for an uncontrolled environment.

6. This device must not be co-located or operating in conjunction with any other antenna or transmitter.

7. Essential performance:
   - Electrosurgery interference recovery: Refer 202.6.2.101 IEC 80601-2-30
   - Limits of the error of the manometer: Refer 202.12.1.102 IEC 80601-2-30
   - Reproducibility of the BLOOD PRESSURE DETERMINATION: Refer 201.12.1.107 IEC 80601-2-30

Unit Illustration

Monitor Unit

LCD

"MEM" Button
"ON/OFF" Button
"SET" Button

WARNING SIGNS AND SYMBOLS USED

Keep Dry
Keep off Sunlight
Type BF Equipment
Note, refer to the attached document
The warning symbol on the lithium battery indicates that the lithium battery should not be replaced at will
Instructions For Use MUST be Consulted
Discard the used product to the recycling collection point according to local regulations

IP21
Magnetic Resonance unsafe

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Important Testing Guidelines

1. Avoid eating, exercising, and bathing for 30 minutes prior to testing.
2. Sit in a calm environment for at least 5 minutes prior to testing.
3. Do not stand while testing. Sit in a relaxed position while keeping your arm level with your heart.
4. Avoid speaking or moving body parts while testing.
5. While testing, avoid strong electromagnetic interference such as microwave ovens and cell phones.
6. Wait 3 minutes or longer before re-testing.
7. Try to measure your blood pressure at the same time each day for consistency.
8. Test comparisons should only be made when monitor is used on the same arm, in the same position, and at the same time of day.
9. This blood pressure monitor is not recommended for people with severe arrhythmia.
10. Do not use this blood pressure monitor if the device is damaged.

Any blood pressure recording can be affected by the following factors:
1. The position of the subject, his or her physiologic condition;
2. The performance and accuracy of the device;
3. Cuff size: too small cuff (bladder) will produce a higher blood pressure value than usual, too big cuff (bladder) will produce a lower blood pressure value;
4. Measuring position does not keep level with your heart;
5. Speaking or moving body parts while testing;
6. Not relaxing for about 5 minutes before taking the measurement.

Quick Start

1. Open the packing box and take out the device. (See Figure A)
2. Insert cuff air plug into the behind side of monitor unit. (See Figure B)
3. Remove thick clothing from the arm area.
4. Rest for several minutes prior to testing. Sit down in a quietplace comfortably, back and arm support on a desk or table, with your legs uncrossed, your arm resting on a firm and your feet flat on the floor. (See Figure C)
5. Apply cuff to your left arm and middle of the cuff at the level of your heart. Bottom of cuff should be placed approximately 1-2cm (1/2") above elbow joint. (See Figures D&E)
6. Press "START/STOP" Button to start testing.

Lithium Battery Charging

The device has a built-in lithium battery that must be recharged using a specific power adapter. When low battery indicator icon " " appears on the screen Please connect the power adapter to charge. When charging, the screen will be displayed in a cycle from " " to " "

Insert the adapter connector into the adapter interface, then insert the adapter into the socket, and then connect the adapter to the cable end. Insert the adapter interface on the device and press the on/off button to measure normally. (if this model has been packed) When you plug in the adapter, the screen power display changes, indicating that you are using the adapter for power.

Note: Power supply is specified as part of ME EQUIPMENT.
**System Settings**

With power off, press "SET" button to activate System Settings. The Memory Group icon flashes.

1. **Select Memory Group**
   
   While in the System Setting mode, you may accumulate test results into 2 different groups. This allows multiple users to save individual test results (up to 150 memories per group.) Press "MEM" button to choose a group setting. Test results will automatically store in each selected group.

2. **Time/Date setting**
   
   Press "SET" button again to set the Time/Date mode. Set the year first by adjusting the "MEM" button. Press "SET" button again to confirm current month. Continue setting the date, hour and minute in the same way. Every time the "SET" button is pressed, it will lock in your selection and continue in succession (month, day, hour, minute, 12/24 hours)

3. **Time Format setting**
   
   Press "SET" button again to set the time format setting mode. Set the time format by adjusting the "MEM" button. EU means European Time US means U.S Time

4. **Triple measurement setting**
   
   Press "SET" to enter the triple measurement function setting mode, and press "MEM" to turn ON or OFF the triple measurement function.

5. **Saved Settings**
   
   While in any setting mode, press "ON/OFF" button to turn the unit off. All information will be saved.

6. **Voice Setting**
   
   Press "SET" button to enter voice setting mode. Set voice format ON or OFF by pressing the "MEM" button.

7. **Volume Setting**
   
   Press "SET" button to enter volume setting mode. Set the voice volume by adjusting the "MEM" button. There are six volume levels.

Applying the Arm Cuff

1. Pick up the sphygmomanometer, open the arm band, put the sphygmomanometer and arm band on the arm, to firm air leakage is appropriate;

2. With sticky nylon section facing outward, insert end of cuff underneath metal ring of cuff.

3. Fasten cuff about 1-2 cm (0.4-0.8") above the elbow joint. For best results apply cuff to bare arm and keep level with heart while testing.

Note: Do not insert air plug into opening located on right side of monitor unit. This opening is designed for an optional power supply only.
Unit Operation

Testing

1. Power On

Press and hold "START/STOP" button to turn the unit on. The LCD screen will appear for one second as unit performs a quick diagnosis. A voice tone will indicate when unit is ready for testing.

Note: Unit will not function if residual air from previous testing is present in cuff. The LCD will flash " " until pressure is stabilized.

2. Testing

After cuff inflation, air will slowly rise as indicated by the corresponding cuff pressure value. A flashing " " will appear simultaneously on screen signaling heart beat detection.

Note: Keep relaxed during testing. Avoid speaking or moving body parts.

Irregular Heartbeat Indicator

If the monitor detects an irregular heart rhythm two or more times during the measuring process, the Irregular Heartbeat Symbol " " appears on screen along with measurement results. Irregular heartbeat rhythm is defined as rhythm that is either 25% slower or faster than the average rhythm detected while measuring systolic blood pressure and diastolic blood pressure. Consult your physician if the Irregular Heartbeat Symbol " " frequently appears with your test results.

Power Off

The "START/STOP" button can be pressed to turn off the unit in any mode. The unit can turn off the power itself about 3 minutes no operation in any mode.

Note: Refer to Page 23–24 for detail WHO Blood Pressure Classification Information.

Memory Check

You may check past test results by using the "MEM" button. The most recent test result and oldest test result in memory can be viewed by pressing and holding the "MEM" button. Upon activating test results, you can press the "MEM" button to scroll through all test results stored in memory.

Memory Deletion

Memory for a selected group may be deleted while in Memory Check mode. Press and hold the "SET" button for approximately 3 seconds to delete all memory records from the selected group with voice broadcast "Memory Clear" and then transfer into testing mode. Press the "START/STOP" button to turn the unit off.

Note: Memory cannot be recovered once it has been deleted.
Unit Operation

Low Battery Indicator

The unit will broadcast "Low Battery" when battery life is depleting and unable to inflate cuff for testing. The " " appears simultaneously for approximately 5 seconds prior to shutting off. Replace batteries at this time. No memory loss will occur throughout this process.

Static Pressure Measurement

In the power down state, press and hold the "START/STOP" button, and then install the batteries. Until the LCD screen is full, release the "START/STOP" button. When the LCD screen displays the double zero, the blood pressure meter is in static state. Software version is displayed: 1.0 is a software version in the figure.

Unit Operation

Troubleshooting

<table>
<thead>
<tr>
<th>Abnormal phenomenon</th>
<th>Cause analysis</th>
<th>Processing method</th>
</tr>
</thead>
<tbody>
<tr>
<td>er1</td>
<td>The armband is tied too tight or too loose, Or the arm strap is tied incorrectly.</td>
<td>Roll the armband correctly</td>
</tr>
<tr>
<td>er2</td>
<td>Move the arm during measurement or electronic sphygmomanometer</td>
<td>Stay quiet, keep your arm steady, and do not move the monitor</td>
</tr>
<tr>
<td>er3</td>
<td>Speaking, nervous or emotional during measurement</td>
<td>Instead of talking, take deep breaths to calm your mood and relax your body</td>
</tr>
<tr>
<td>er4</td>
<td>Incorrect measurement posture</td>
<td>Adjust posture, see “Blood pressure gauge Wearing”</td>
</tr>
<tr>
<td>Er5</td>
<td>There is interference in charging process or improper operation in measuring process</td>
<td>See operation Instructions.</td>
</tr>
</tbody>
</table>

Bluetooth connection

-Using for the first time
2. Open the App on your phone or tablet. If requested, you should enable Bluetooth on your device. You can enable Bluetooth under the Settings menu on your smart phone or tablet.
3. Create a new user login, or login with your existing user name and password.
4. Selection device “Blood pressure monitor”.

-Pairing your monitor with a Smart Device
1. Long press the "memory" button (or the "M" key) on your monitor until the word "boo" flashes on your monitor.

Transfer your readings

1. As soon as your measurement is complete, open the app on your smart phone to transfer your readings.
2. You can view your blood pressure readings on the app.

Note: If another family member uses it, clean the cuff before use and wait after it dries naturally; Press "SET" to set the memory group, set the memory group. Restart the measurement according to the operation method. If you want to view the measured value, please refer to the memory view description on page 18.
Blood Pressure

Blood pressure is the force of blood pushing against the walls of arteries. It is typically measured in millimeters of mercury (mmHg). Systolic blood pressure is the maximum force exerted against blood vessel walls each time the heart beats. Diastolic blood pressure is the force exerted on blood vessels when the heart is resting between beats.

An individual’s blood pressure frequently changes throughout the course of a day. Excitement and tension can cause blood pressure to rise, while drinking alcohol and bathing can lower blood pressure. Certain hormones like adrenaline (which your body releases under stress) can cause blood vessels to constrict, leading to a rise in blood pressure.

If these measuring numbers become too high, it means the heart is working harder than it should.

**Blood Pressure Q&A**

**Q:** What is the difference between measuring blood pressure at home or at a professional healthcare clinic?

**A:** Blood pressure readings taken at home are now seen to give a more accurate account as they better reflect your daily life. Readings can be elevated when taken in a clinical or medical environment. This is known as White Coat Hypertension and may be caused by feeling anxious or nervous.

**Note:** Abnormal test results may be caused by:
1. Improper cuff placement
   - Make sure cuff is snug—not too tight or too loose.
   - Make sure bottom of the cuff is approximately 1-2cm (1/2'') above the elbow joint.
2. Improper body position
   - Make sure to keep your body in an upright position.
3. Feeling anxious or nervous
   - Take 2-3 deep breaths, wait a few minutes and resume testing.

**Q:** What causes different readings?

**A:** Blood pressure varies throughout the course of a day. Many factors including diet, stress, cuff placement, etc. may affect an individual's blood pressure.

**Q:** Should I apply the cuff to the left or right arm? What is the difference?

**A:** Either arm can be used when testing, however, when comparing results, the same arm should be used. Testing on your left arm may provide more accurate results as it is located closer to your heart.

**Q:** What is the best time of day for testing?

**A:** Morning time or any time you feel relaxed and stress free.
**Maintenance**

1. Avoid dropping, slamming, or throwing the unit.

2. Avoid extreme temperatures. Do not expose unit directly under sunshine.

3. When cleaning the unit, use a soft fabric and lightly wipe with mild detergent. Use a damp cloth to remove dirt and excess detergent. It is recommended to wash once every 1-3 months under normal circumstances. If the previous cleaning instructions are not successful, repeat several times until the visible dirt is removed.

4. Cuff Cleaning and Disinfection:
   - A) Spread the cuff (skin-contact surface) upwards onto a clean table. Use a damp clean cloth (water-based) to wipe the skin-contact surface with a force.
   - B) Soak the cloth clean with drinking water and wring it dry. Repeat A) with the damp cloth.
   - C) Apply 70%-80% alcohol to a new cloth (or 75% alcohol cotton-ball), use it to wipe the skin-contact surface with a force. Then soak the cloth with the alcohol again (or change a new 75% alcohol cotton-ball), repeat the disinfection procedure for 3 times.
   - D) When the disinfection towards the skin-contact surface is finished, wipe then the skin-contact surface with a cloth (alcohol-based) or alcohol cotton-ball thoroughly for 3 times.
   - E) Leave the cuff naturally dry, then it is ready for reuse.

   Notice: Do not soak in water or splash water on it.

5. Do not use petrol, thinners or similar solvents.

6. Remove batteries when not in operation for an extended period of time.

7. Do not disassemble product.

8. It is recommended the performance should be checked every 2 years.

9. Expected service life: Approximately three years at 10 tests per day.

10. No service and maintenance while it is in use and maintenance only be performed by service personnel. Service and maintenance require parts, repair, technical support will be provided.

11. If it is not used for a long time, it is recommended to charge the lithium battery within 3-4 months to ensure that it does not lose power. Users can not replace the lithium battery by themselves if it is confirmed that the lithium battery is faulty, please do not repair it by yourself. Please contact the manufacturer or dealer in time.

12. If the lithium battery is not charged in time, or the lithium battery is placed for a long time, there may be the situation that the lithium battery is about to run out. In this case, the device cannot be turned on. In this case, the lithium battery should be charged immediately.

   The following rules must be observed during use:
   1. Prevent lithium batteries from entering water. Do not heat lithium batteries, do not throw lithium batteries into fire.
   2. Lithium batteries should not be used for a long time in an environment of more than 60°C.
   3. The device can only use lithium batteries specified by the manufacturer.
   4. Use the specified AC adapter to charge the device.
   5. In order to avoid the normal use of lithium batteries due to self-discharge in the storage process, please charge the lithium batteries every 3-4 months to ensure the normal operation of lithium batteries.

**Specifications**

<table>
<thead>
<tr>
<th>Product Description</th>
<th>Arm-type Fully Automatic Blood Pressure Monitor</th>
</tr>
</thead>
<tbody>
<tr>
<td>Model</td>
<td>DBP-6293L</td>
</tr>
<tr>
<td>Display</td>
<td>LCD Digital Display  Size:100mm×37mm (3.94&quot;×1.46&quot;)</td>
</tr>
<tr>
<td>Measurement Method</td>
<td>Oscillometric Method</td>
</tr>
<tr>
<td>Measurement Range</td>
<td>Systolic Pressure 60mmHg – 260mmHg</td>
</tr>
<tr>
<td></td>
<td>Diastolic Pressure 30mmHg – 200mmHg</td>
</tr>
<tr>
<td></td>
<td>Pressure 0mmHg – 290mmHg</td>
</tr>
<tr>
<td></td>
<td>Pressure ±3mmHg</td>
</tr>
<tr>
<td></td>
<td>Pulse 50~180 Beats/Minute</td>
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<tr>
<td></td>
<td>Pulse ±5%</td>
</tr>
<tr>
<td>Pressurization</td>
<td>Automatic Pressurization</td>
</tr>
<tr>
<td>Memory</td>
<td>2x150 Memories in Two Groups with Date and Time</td>
</tr>
<tr>
<td>Function</td>
<td>Irregular Heartbeat Detection</td>
</tr>
<tr>
<td></td>
<td>WHO Classification Indicator</td>
</tr>
<tr>
<td></td>
<td>Last 3 Tests Average</td>
</tr>
<tr>
<td></td>
<td>Low Battery Detection</td>
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<tr>
<td></td>
<td>Automatic Power-Off</td>
</tr>
<tr>
<td></td>
<td>Voice</td>
</tr>
<tr>
<td></td>
<td>Backlight</td>
</tr>
<tr>
<td></td>
<td>Bluetooth</td>
</tr>
<tr>
<td>Power Source</td>
<td>Lithium battery 3.7V or DC5.0v 1000mA medical external power supply (DC5.0V, 1000mA)(recommended, not provided)</td>
</tr>
<tr>
<td>Battery Life</td>
<td>Approximately 2 months at 3 tests per day</td>
</tr>
<tr>
<td>Unit Weight</td>
<td>Approx. 1.5kg (3.5lb.)</td>
</tr>
<tr>
<td>Unit Dimensions</td>
<td>Approx. 124 x 71.5 x 24.5mm (4.88&quot; x 2.81&quot; x 0.96&quot;)</td>
</tr>
<tr>
<td>Cuff Circumference</td>
<td>Medium cuff: Fits arm circumference 22-42 cm</td>
</tr>
<tr>
<td>Operating Environment</td>
<td>Temperature 10°C ~ 40°C (50°F ~ 104°F)</td>
</tr>
<tr>
<td></td>
<td>Humidity 15% ~ 93% RH</td>
</tr>
</tbody>
</table>

**Operating Environment**

<table>
<thead>
<tr>
<th>Pressure</th>
<th>80kPa~106kPa</th>
</tr>
</thead>
</table>

**Storage Environment**

<table>
<thead>
<tr>
<th>Temperature</th>
<th>-25°C<del>55°C (-13°F</del>131°F)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Humidity</td>
<td>≤93% RH</td>
</tr>
</tbody>
</table>

**Transport Environment**

<table>
<thead>
<tr>
<th>Temperature</th>
<th>-25°C<del>55°C (-13°F</del>131°F)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Humidity</td>
<td>≤93% RH</td>
</tr>
</tbody>
</table>

**Bluetooth**

<table>
<thead>
<tr>
<th>Modulation Type</th>
<th>GFSK</th>
</tr>
</thead>
<tbody>
<tr>
<td>Version</td>
<td>5.0</td>
</tr>
<tr>
<td>Operation frequency</td>
<td>2.4GHz (2400-2483.5MHz)</td>
</tr>
<tr>
<td>Antenna gain</td>
<td>0.5</td>
</tr>
<tr>
<td>Bandwidth</td>
<td>2.0</td>
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</tbody>
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**Classification**

<table>
<thead>
<tr>
<th>Internal Powered Equipment, Type BF</th>
<th>Cuff is the Applied Part</th>
</tr>
</thead>
</table>

**Ingress Protection rating**

| IP 22, Indoor Used Only |

**Battery Shelf life**

| 60 months |

**Battery Storage Temperature**

| -25°C~35°C (-13°F~95°F) |

Specifications are subject to change without notice.

Safety Standard (included but not limited):
1. IEC 60601-2-30, medical electrical equipment – part 2-30: particular requirements for the basic safety and essential performance of automated noninvasive sphygmomanometers. (Cardiovascular)
2. ISO 81060-2, non-invasive sphygmomanometers – part 2: clinical validation of automated measurement type. (Cardiovascular)
5. IEC 60601-1-11, medical electrical equipment – part 1-11: general requirements for basic safety and essential performance – collatral standard: requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment.
Warranty

The Blood Pressure Monitor is guaranteed for 2-year from the date of purchase. If the Blood Pressure Monitor does not function properly due to defective components or poor workmanship, we will repair or replace it freely. Please contact local retailer for details.

Electromagnetic Compatibility Information

The device satisfies the EMC requirements of the international standard IEC 60601-1-2. The requirements are satisfied under the conditions described in the table below. The device is an electrical medical product and is subject to special precautionary measures with regard to EMC which must be published in the instructions for use. Portable and mobile HF communications equipment can affect the device. Use of the unit in conjunction with non-approved accessories can affect the device negatively and alter the electromagnetic compatibility. The device should not be used directly adjacent to or between other electrical equipment.

Table 1

<table>
<thead>
<tr>
<th>Guidance and declaration of manufacturer-electromagnetic immunity</th>
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<tbody>
<tr>
<td><strong>Emissions test</strong></td>
</tr>
<tr>
<td>Radiated emission CISPR 11</td>
</tr>
<tr>
<td>Conducted emission CISPR 11</td>
</tr>
<tr>
<td>Harmonic emissions IEC 61000-3-2</td>
</tr>
<tr>
<td>Voltage fluctuations/ flicker emissions IEC 61000-3-3</td>
</tr>
</tbody>
</table>

Table 2

<table>
<thead>
<tr>
<th>IMMUNITY test</th>
<th>IEC 60601 test level</th>
<th>Compliance level</th>
<th>Electromagnetic environment -guidance</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Surge</strong> IEC 61000-4-5</td>
<td>±0.5 kV, ±1 kV (differential mode)</td>
<td>-</td>
<td>Mains power quality should be that of a typical commercial or hospital environment.</td>
</tr>
<tr>
<td><strong>Voltage dips, short interruptions and voltage variations on power supply input lines</strong> IEC 61000-4-11</td>
<td>0 % UT; 0.5 cycle at 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315°; 0.5 cycle at 0° and 70 % UT; 25/30 cycles at 0°</td>
<td>0 % UT; 1 cycle and 70 % UT; 25/30 cycles at 0°</td>
<td>Mains power quality should be that of a typical commercial or hospital environment.</td>
</tr>
<tr>
<td><strong>Power frequency (50/60 Hz) magnetic field</strong> IEC 61000-4-8</td>
<td>30 A/m; 50 Hz or 60 Hz; 30 A/m; 50 Hz or 60 Hz</td>
<td>-</td>
<td>Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.</td>
</tr>
</tbody>
</table>

Table 2 (continued)

<table>
<thead>
<tr>
<th>IMMUNITY test</th>
<th>IEC 60601 test level</th>
<th>Compliance level</th>
<th>Electromagnetic environment -guidance</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Radiated RF EM fields</strong> IEC 61000-4-3</td>
<td>10 V/m, 80 MHz-2.7 Ghz; 80% of AM at 1 kHz</td>
<td>-</td>
<td>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. Recommended separation distance 80 MHz to 800 MHz 800 MHz to 2.7 Ghz 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. Interference may occur in the vicinity of equipment marked with the following symbol:</td>
</tr>
<tr>
<td><strong>Conducted disturbances induced by RF fields</strong> IEC 61000-4-6</td>
<td>3 V in 0.15 MHz; 80 MHz 6 V in ISM and/or amateur radio bands between 0.15 MHz and 80 MHz 80 % AM at 1 kHz</td>
<td>-</td>
<td>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. Recommended separation distance 80 MHz to 800 MHz 800 MHz to 2.7 Ghz 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. Interference may occur in the vicinity of equipment marked with the following symbol:</td>
</tr>
</tbody>
</table>
### Electromagnetic Compatibility Information

#### Table 3

<table>
<thead>
<tr>
<th>Test frequency (MHz)</th>
<th>Band (MHz)</th>
<th>Service</th>
<th>Modulation</th>
<th>Maximum power (W)</th>
<th>Distance (m)</th>
<th>Immunity test level (V/m)</th>
</tr>
</thead>
<tbody>
<tr>
<td>385</td>
<td>380-390</td>
<td>TETRA 400</td>
<td>Pulse modulation 18kHz</td>
<td>1.8</td>
<td>0.3</td>
<td>27</td>
</tr>
<tr>
<td>450</td>
<td>430-470</td>
<td>GSM 800</td>
<td>Pulse modulation 18kHz</td>
<td>2</td>
<td>0.3</td>
<td>28</td>
</tr>
<tr>
<td>710</td>
<td>704-787</td>
<td>LTE Band 13, 17</td>
<td>Pulse modulation 217Hz</td>
<td>0.2</td>
<td>0.3</td>
<td>9</td>
</tr>
<tr>
<td>745</td>
<td>780</td>
<td>GSM 1800, CDMA 1900, GSM 1900, DECT; LTE Band 1, 3, 5, 7, 8, 20</td>
<td>Pulse modulation 18kHz</td>
<td>2</td>
<td>0.3</td>
<td>28</td>
</tr>
<tr>
<td>810</td>
<td>800-960</td>
<td>GSM 400/450, TETRA 450, CDMA 850, LTE Band 5</td>
<td>Pulse modulation 217Hz</td>
<td>0.2</td>
<td>0.3</td>
<td>9</td>
</tr>
<tr>
<td>930</td>
<td>1700-1900</td>
<td>GSM 1800; CDMA 1900; GSM 1900; DECT; LTE Band 1, 3, 5, 7, 8, 20, UMTS</td>
<td>Pulse modulation 18kHz</td>
<td>2</td>
<td>0.3</td>
<td>28</td>
</tr>
<tr>
<td>1275</td>
<td>10</td>
<td>3.8</td>
<td>7.3</td>
<td>12</td>
<td>23</td>
<td></td>
</tr>
<tr>
<td>2450</td>
<td>2400-2570</td>
<td>5.24 GHz</td>
<td>Pulse modulation 217Hz</td>
<td>2</td>
<td>0.3</td>
<td>28</td>
</tr>
<tr>
<td>5240</td>
<td>2100-2300</td>
<td>WLAN 802.11</td>
<td>Pulse modulation 217Hz</td>
<td>0.2</td>
<td>0.3</td>
<td>9</td>
</tr>
<tr>
<td>5785</td>
<td>5100-5800</td>
<td>a/n</td>
<td>Pulse modulation 217Hz</td>
<td>0.2</td>
<td>0.3</td>
<td>9</td>
</tr>
</tbody>
</table>

#### Table 4

<table>
<thead>
<tr>
<th>Rated maximum output power of transmitter</th>
<th>Separation distance according to frequency of transmitter (m)</th>
</tr>
</thead>
<tbody>
<tr>
<td>W</td>
<td>80 MHz to 800 MHz</td>
</tr>
<tr>
<td></td>
<td>( d = \frac{10}{P} )</td>
</tr>
<tr>
<td></td>
<td>800 MHz to 2.7 GHz</td>
</tr>
<tr>
<td></td>
<td>( d = \frac{1}{P} )</td>
</tr>
<tr>
<td>0.01</td>
<td>0.12</td>
</tr>
<tr>
<td>0.1</td>
<td>0.38</td>
</tr>
<tr>
<td>1</td>
<td>1.2</td>
</tr>
<tr>
<td>10</td>
<td>3.8</td>
</tr>
<tr>
<td>100</td>
<td>12</td>
</tr>
</tbody>
</table>

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 highest 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.

### Additional Notes

**Important Instructions Before Use**

1. **WARNING:** Use of this equipment adjacent to or stacked with other equipment should be avoided because it could result in improper operation. If such use is necessary, this equipment and the other equipment should be observed to verify that they are operating normally.

2. **WARNING:** PORTABLE RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 30 cm (12 inches) to any part of Arm-type Fully Automatic Digital Blood Pressure Monitor, including cables specified by the MANUFACTURER. Otherwise, degradation of the performance of this equipment could result.

3. The software identifier refer to the software verification and validation report, and the file code is "xxxxxxxxxxx."  
4. **verify manometer pressure accuracy:** In the power down state, press and hold the "START/STOP" button, and then install the batteries. Until the LCD screen is full, release the "START/STOP" button. When the LCD screen displays the double zero, the bloodpressure meter is in static state. At this point, 500ml gas capacity, calibrated standard pressure gauge and manual pressure device can be connected to the sphygmomanometer through the sleeve interface of the sphygmomanometer, and then the difference between the reading of the sphygmomanometer and that of the standard pressure gauge can be compared. This mode can be used to verify manometer pressure accuracy.

5. **Contraindications:** Product is not intended for infants or children who cannot express their intentions.

6. **Intended Use:** The digital blood pressure monitor is reusable for clinical and home use and are non-invasive blood pressure measurement systems designed to measure the systolic and diastolic blood pressure and pulse rate of adolescents and adults individual by using a non-invasive technique, which is a well-known technique in the market called the "oscillometric method," it can measure the systolic blood pressure, diastolic blood pressure and pulse rate on up-arm, and the device is reusable for clinical or home use.

7. The patient is the operator.  
8. **WARNING:** The PATIENT is an intended OPERATOR. Do not carry out other maintenance operations except to replace the battery.

9. **WARNING:** Do not modify this equipment without authorization of the manufacturer.

10. **ESSENTIAL PERFORMANCE Maintenance advice:** Pressure calibration will be carried out when this product leaves the factory. Patients can use the method described in the section "Verify Manometer Pressure Accuracy" to verify the accuracy. If the accuracy deviation is large, please contact the manufacturer to recalibration.

11. **Mechanical strength and resistance to heat:** The resistance to heat will be retained by device during the EXPECTED SERVICE LIFE of the ME EQUIPMENT.

12. The cuff and the case of the blood pressure monitor have been tested for biocompatibility and do not contain allergenic or harmful materials. Please stop using it if allergy occurs during use.

13. **WARNING:** Non-professionals do not modify the equipment, otherwise it will make the equipment measurement is not accurate.

14. **WARNING:** Do not use a damaged cuff for blood pressure measurement.

15. When measuring with the cuff, if the tester feels seriously uncomfortable, press the button of the blood pressure monitor to deflate the cuff, or remove the cuff directly from the arm.

16. **WARNING:** If an unexpected reading occurs, the operator can take several more measurements and consult a doctor.

17. **WARNING:** This equipment is used outside the specified environment, may damage the equipment, and may be inaccurate measurement.

18. **ME equipment not intended for use in conjunction with flammable agents "ME equipment not intended for use in oxygen rich environment"

**Correct Disposal of This Product**

(Waste Electrical & Electronic Equipment)

This marking shown on the product indicates that it should not be disposed with other household waste at the end of its life. To prevent potential harm to the environment or to human health, please separate this product from other types of wastes and recycle it responsibly. When disposing this type of product, contact the retailer where product was purchased or contact your local government office for details regarding how this item can be disposed in an environmentally safe recycling center. Business users should contact their supplier and check the terms and conditions of the purchasing agreement. This product should not be mixed with other commercial wastes for disposal. This product is free of hazardous materials.