UPPER ARM AUTOMATIC DIGITAL BLOOD PRESSURE MONITOR

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INSTRUCTION MANUAL Model: LD-575

1.Main Body

4.Tube Plug

9.Button ' (り)

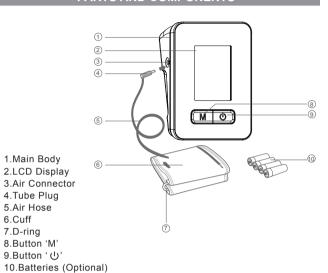
5.Air Hose

6.Cuff

7.D-ring 8.Button 'M



PARTS AND COMPONENTS



SYMBOLS

	1
Symbols	Meaning
<u> </u>	Manufacturer
EC REP	Authorized Representative in the European community
	Symbol for the marking of electrical and electronics devices according to Directive 2002/96/EC. The device, accessories and the packaging have to be disposed of waste correctly at the end of the usage. Please follow Local Ordinances or Regulations for disposal.
C€ 0123	CE marking in conformity with EC directive 93/42/EEC
Ť	Keep dry
Ţ i	Attention, consult accompanying documents
፟	Type BF Applied Part
(h	Stand by

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GENERAL

This instruction manual is intended to assist the user for safe and efficient operation of the automatic digital blood pressure monitor (hereinafter: device) model LD-575. The device must be used in accordance with the procedures described in the manual. It is important to read and understand the entire manual, especially the section <Tips on taking blood pressure measurement>.

This device is intended for the non-invasive measurement of systolic and diastolic arterial blood pressure and pulse rate in adults (age 15 and above) . Consult the physician if measurement is taken in children or persons with arrhythmia as errors

PRINCIPLE OF OPERATION

This device adopts the oscillometric technology with Fuzzy Algorithm measuring the arterial blood pressure and pulse rate. The cuff is wrapped around the arm and automatically inflated by the air pump. The sensor of the device catches weak fluctuation of the pressure in the cuff produced by extension and contraction of the artery of the arm in response to each heartbeat. The amplitude of the pressure waves is measured, converted in millimeters of the mercury column, and is displayed by digital value.

Annotation: This device can not provide reasonable accuracy if used or stored in the temperature or humidity beyond the range stated in the section <SPECIFICATIONS> of this manual.

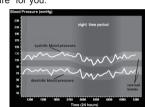
CAUTION: DO NOT USE THE DEVICE OUTDOORS

NEW TECHNOLOGIES USED

Fuzzy Algorithm is the processing algorithm taking into account of the speciality of individual heartbeats, which provides higher accuracy of measurement Software version: V1.1

TIPS ON TAKING BLOOD PRESSURE MEASUREMENT

1.It is necessary to know that arterial blood pressure is subjected to sharp fluctuations. The level of the arterial blood pressure depends on many factors. Generally arterial blood pressure is lower in summer and higher in winter. Arterial blood pressure changes with atmosphere pressure and is affected considerably by many factors, e.g. physical loads, emotional excitability, stress, meals, etc. Medicines, drinking, smoking affects greatly the level of individual blood pressure. When blood pressure is measured in hospital, the value is always higher than that at home. The reason is the tensity and such case is especially serious in given group patients, which is known as 'White coat effect' medically. Blood pressure will raise in low temperature, so it is better to take blood pressure measurement in room temperature (approximately 20°C). If this device was stored in low temperature, it is necessary to leave it in room temperature for at least 1 hour, otherwise the measurement can be inaccurate. Blood pressure does vary with age and individual, and it is recommended to write down the readings in blood pressure record daily, then you can check with your doctor to find out what is "normal blood pressure" for you



The illustration is from British Hypertension Society

2. Take measurement under doctor's instruction for patients with cardio-vascular diseases

Under no circumstances should you alter the dosages of any drugs prescribed

3.Accurate measurement of blood pressure may be difficult in arrhythmia, premature beat, atrial fibrillation atreriosclerosis hypoperfusion diabetes pregnancy nephropathy, weak pulse, or in patients with obvious fluctuation of heart contraction rhythm. Please consult a qualified physician to interpret your blood pressure readings.

4.It is necessary to keep quiet during measurement to get accurate readings. Measurement should be conducted in quiet environment at room temperature. Don't eat or smoke before a measurement.

This device is supplied with the standard cuff and adult large cuff which are fit for the arm size 22-32 cm and 32-42cm. Care should be taken to ensure that the cuff size is appropriate for the person whose blood pressure is being taken. Children and adults with cuff size fall outside the range of the standard cuff size and large adult cuff size should select special size cuffs. Please contact the dealer to get these special size cuffs.

ATTENTION: Do not use cuffs other than the original cuff contained in this kit!

5.Repeated measurements with interval at 3 minutes are recommended, so you can calculate the average to get more accurate measurement. Atherosclerosis patients are required longer interval (10-15 minutes) as elasticity of patients' vessels decreased significantly in these diseases. 10-15 minutes interval is also applicable for patients suffering from diabetes for a long time.

CLASSIFICATION

- ME EQUIPMENT not intended for use in an oxygen rich environment or in the presence of flammable mixers.
- Internally powered equipment (without adapter), Class II equipment
- (with adapter) Type BF applied part, recognize the cuff as applied part.

BATTERY INSTALLATION

- 1. Open the battery cover and install four 'AA' type batteries into the battery compartment as indicated. Make sure that the polarity is correct; 2. Close the battery compartment cover.
- display or nothing after " U " button is pressed;
- Batteries in this kit are intended to check work capacity of the device and the
- life-span of the batteries can be shorter than the recommended; • Use R6,LR6 or AA alkaline batteries, do not use rechargeable batteries;
- Only same type batteries are allowed to use together. Replace all batteries simultaneously;
- If the device is to be unused for long time, please take out the batteries;
- Don't leave the worn batteries in the device.

USE THE DEVICE WITH AC POWER ADAPTER Besides batteries you can use AC power adapter as the power supply. AC power adapter is optional for the device for sale

Insert the AC adapter cord into the jack on the right side of the monitor

Insert the AC adapter plug into the outlet.

To remove the AC adapter, disconnect the adapter plug from the AC outlet first and then disconnect the cord from the monitor's jack.

CAUTION

- When using optional AC adapter, the AC adapter must comply with the requirements of standard IEC60601-1.
- To avoid possible damage to the monitor, use only the exclusive AC adapter that can be purchased from authorized dealers. Other adapter may damage the blood pressure monitor.

- The AC adapter is used as an isolating means, the AC adapter plug shall insert into the outlet nearby the operator, make it easy to disconnection the
- If long time work, remove the plug after the adapter cools, and prevent bums.

Note: The monitor is designed not to draw power from the batteries when the AC adapter in use.

> DC 6V \oplus \ominus

Optional AC adapter technical feature: Output voltage: 6V±5% Max. output current: At least 600 mA Output plug polarity: <-> inner

CORRECT POSTURE FOR MEASUREMENT

1. Sit beside the table and let the table support your arm as you take the measurement. Make sure that the cuff on the upper arm no cross, and is at approximately the same level as the heart, make sure that your feet lie on the ground and no cross.

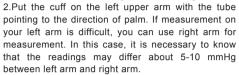


2. You may lie on your back and take measurement. Look at the ceiling, keep calm, and don't move your neck or body during the measurement. Make sure that the cuff on the upper arm is at approximately the same level as the heart.



ASSEMBLY THE CUFF

1.Insert the edge of the cuff approximately 5 centimeters into the D-ring as shown in figure.



3. Wrap cuff around your upper arm with the lower edge of the cuff approximately 2-3 centimeters above the elbow. The mark <ARTERY> must be over the artery of the arm.

4. Press the cuff to make sure that it is attached securely. The cuff should not be too tight or too loose. Two fingers should be easily put in between cuff and

5.The mark <INDEX> on the cuff must point to area <NORMAL> or <LARGE CUFF>. This means the cuff size is correct. If mark <INDEX> points to the area beyond area <NORMAL> or <LARGE CUFF>, please consult your dealer whether you need another size cuff.

6. Sometimes it is difficult to make the cuff regular owning to the shape of the user's upper arm, the cone-shape assembly of cuff is also acceptable.

7.If your clothes restrict blood circulation of your upper arm, or you roll your sleeve up so as to result in such restriction. Please take off your clothes to get accurate measurement if necessary.







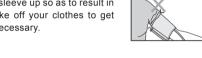






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<section-header> 188 🕾 Fig.1



CARRY OUT A MEASUREMENT

1. Insert the tube plug into the air connector. Before the measurement, take 3~5 times deep breath and relax yourself. Don't talk or move your arm;

2.Press button ' U ', and all symbols will appear on display in 2 seconds as Fig.1.

Then two short beep will sound and '0' will appear on the screen. Pump begins to inflate with display showing the reading of pressure. Generally the pressure will reach 190mmHg as Fig.2;

3. The pump stop inflating and pressure begins to decrease gradually, during which the user's blood pressure and pulse will be calculated as Fig.3;

4. There will be a long beep following the accomplishment of measurement. The air in the cuff will deflate quickly and the blood pressure reading, pulse reading will show in the display as Fig.4;

5. Press the button '(1) 'to turn off the device. Please rest for at least 3 minutes for another measurement. If the power supply is not switched off and the device keeps unused for 3 minutes, the device will be switched off automatically.

AUTOMATIC INFLATION

There are 4 given levels of given inflation pressure for this device: 190mmHg, 230mmHg, 270mmHg and 300mmHg. When 190mmHg is not enough or movement of arm occurs, the device will automatically inflate to reasonable pressure level to ensure a successful measurement. It is not a fault.



RAPID DEFLATION DURING MEASUREMENT

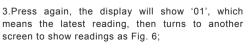
If you do not feel well during measurement or want to stop the measurement for some reason, you can press the " மு " button. The device will quickly release the air in cuff and the device will be switched off.

FUNCTION OF MEMORY

MEMORY RECALL

1.LD-575 can store 90 sets of readings and automatically calculate the average value of the latest 3 readings. When the memory is full (90 sets of readings are stored), the oldest reading will be replaced by new one automatically. Memory will not clear away even if power supply is removed;

2. After a measurement or when the device stands by, the user can press button Memory to recall memory. Press button Memory, the display will show the average value of the latest 3 readings as Fig. 5;



4. Press again, the display will show '02', which means the second to the latest reading...

MEMORY CLEARANCE

After a measurement is finished or when the device stands by, hold down button Memory for at least 5 seconds, the display will show 'CLR' which means all the stored reading are removed as Fig.7.



WHO BLOOD PRESSURE CLASSIFICATION INDICATION

Standards for assessment of high or low blood pressure, regardless of age, have been established by World Health Organization(WHO) as show in the chart as right:

The indicator displays a segment, based on the current data, corresponding to the WHO classification.

For example, if your blood pressure is 135mmHg (Systolic Pressure), 78mmHg (Diastolic Pressure), according to the world health organization standard, your blood pressure level is High Normal.

Note:

- 1.If the systolic blood pressure and diastolic blood pressure fall into different categories, the higher value should be taken for classification.
- 2. The WHO blood pressure classification indication in the device is only a reminder, it can not be regarded as the final diagnosis

ERROR AND LOW BATTERY INFORMATION

INDICATION POSSIBLE REASON		CORRECTION METHODS
Err	The cuff is put on wrongly or the tube plug is inserted too loosely. Movement of arm/hand or talking during measurement. The cuff is not inflated to necessary pressure. Arrhythmia.	Make sure that cuff is put on correctly and the tube plug is inserted tightly and repeat the measurement. Repeat the measurement with following completely recommendations of manual. Repeat the measurement with pumping cuff to higher pressure Consult your personal physician.
	The batteries are weak	Replace all 4 batteries with new-ones.

CARE, STORING, REPAIR AND RECYCLING

- 1. It's necessary to protect this device against high moisture, direct sunlight, shock, solvent, alcohol and gasoline.
- 2. Remove the batteries if the device is to be stored for a long time and keep the batteries far from the children.
- 3. Keep the cuff from sharp subject and don't extend or twist the cuff.
- 4. Use only soft and dry cloth to clean the device.
- 5. The wrist cuff are sensitive and must be handled with care. You can clean the cuff cover with damp cloth as daily maintenance.

To avoid across infection when share the cuff, you can sanitary treatment of inner side of fabrics cover of the cuff and contacting with help of cotton wool tampons, moistened by 3% solution of hydrogen dioxide. After long using, it is allowed partial discoloration of fabrics covering of the cuff. It is not allowed the laundry of the cuff, as well as ironing by hot flatiron.

WARNING: Under no circumstances may you wash the inner bladder! 6. Since neither the device nor batteries are household waste, follow your

local recycling rules and dispose them at appropriate collection sites. 7. Do not open the device. It is delicate electrical components and an intricate air unit that could be damaged. If you can not fix the problem

using the troubleshooting instruction, request service from your dealer. 8. It is generally recommended to have the monitor inspected every 2 years, to ensure proper functioning and accuracy and safety. Please contact your dealer for maintenance

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

9. Do not serve or maintain the cuff when in use with patient

TROUBLESHOOTING

SYMPTOM	CHECK POINT	REMEDY
No display when connect the power.	The batteries have run down. The polarity of battery is wrong. The contact of battery compartment is polluted.	Replace all the batteries with new ones. Install the batteries correctly. Clean the battery terminals with dry cloth.
Inflation stops and re-inflate later.	The automatic inflation for ensuring correct measurement. Did you talk or move your arm (or hand) during measurement?	See <automatic INFLATION> Keep quiet and silent during the measurement.</automatic
The reading is extremely low or high.	Is the cuff at the same level as the heart? Is the cuff wrapped right? Did you strain your arm during measurement? Did you talk or move your arm (or hand) during measurement?	Make sure that your posture is right. Wrap the cuff correctly. Relax during measurement. Keep quiet and silent during the measurement.
Pulse rate is too low or too high.	Did you talk or move your arm (or hand) during measurement? Did you make measurement right after exercise?	Keep quiet and silent during the measurement. Take measurement again after resting for more than 5 minutes.
The batteries are run down soon.	Faulty batteries are used.	Use alkaline batteries of known manufacturers.

WARRANTY OBLIGATIONS

- 1. Warranty for this automatic digital blood pressure monitor is 24 months since the date of purchase. The 24 months warranty excludes the monitor cuff. The cuff is warranted for 12 months.
- 2. The warranty obligations are prescribed by warranty certificate for buyer. 3. The addresses of organizations for guarantee maintenance are present in the warranty certificate

WARNING

• Do not modify this equipment without authorization of the manufacturer. · All major maintains on the device must be performed by an authorized service center or distributor. No use-serviceable parts inside, before servicing to authorized representative or manufacturer!

DECLARATION:

When technical information for user requirements is not in the scope of confidentiality of the Company, the Company committed to provide information disclosure in accordance with procedure, including circuit diagrams and parts lists, and other related type technology information that do not involve commercial secrets may be disclosed. Access to information channels and procedures, please contact your dealer or manufacturer.

SPECIFICATIONS

Model	LD-575
Size	134mm(L) ×98mm(W) ×57(H)mm
Weight	Approximately 236g without batteries
Measuring method	Oscillometry
Extreme Pressure	300mmHg
Measuring range	40 to 260 mmHg (blood pressure) 40 to 160 beats/minute (pulse rate)
Measuring accuracy	± 3 mmHg for static pressure ± 5% of the reading for the pulse rate
Inflation	Automatic by the pump
Rapid deflation	Automatic electronic valve
Batteries	4"AA"×1.5V

Adapter	Optional component, 6V, 600mA
Memory	90 sets of memory
Operation temperature and humidity, air pressure	+10°C to + 40°C, 85% and below 700hPa to 1060hPa
Transport and storage temperature and humidity, air pressure	-20°C to + 50°C, 85% and below 500hPa to 1060hPa
Upper arm circumference	Applicable for arm circumference 22-32cm (standard cuff); 32-42cm(large adult cuff)
Complete kit	Main body, storage bag(Optional), standard cuff, large adult cuff (Optional), 4×AA batteries (Optional), instruction manual, warranty card
Pollution Degrees	Degrees 2
Overvoltage category	Category II
High Altitudes (m)	≤2000m
Fuse	Adapter: T3.15AH250V Main Unit: T630mAl250V



HONSUN (NANTONG) Co., Ltd.

Address: No.8, Tongxing Road, Economic&Technical Development Area, Nantong City, Jiangsu, P.R.CHINA

REP

SHANGHAI INTERNATIONAL HOLDING CORP. GmbH(EUROPE)

BLOOD PRESSURE RECORD

DATE	SYSTOLIC (mmHg)	DIASTOLIC (mmHg)	PULSE (beats/minute)

MANUFACTURER'S DECLARATION

Guidance and manufacturer's declaration - electromagnetic immunity

The model LD-575 Digital Blood Pressure Monitor is intended for use in the electromagnetic environment specified below. The customer or the user of the model LD-575 Digital Blood Pressure Monitor should assure that is used in such an environment.

Emission test	Compliance level	Electromagnetic environment-guidance	
RF emissions CISPR 11	Group 1	The model LD-575 Digital Blood Pressure Monitor uses RF energy only for its internal function. Therefore, its RF emissions are very low and aren't likely to cause any interference in nearby electronic equipment.	
RF emissions CISPR 11	Class B	The model LD-575 Digital Blood Pressure	
Harmonic emission IEC 61000-3-2	Class A	Monitor is suitable for use in all establishments, including domestic establishments and those directly connected to the public low-voltage powe supply network that supplies buildings use for domestic purposes.	
Voltage fluctuations/ flickeremissions IEC 61000-3-3	Complies		

Guidance and manufacturer's declaration – electromagnetic immunity

The model LD-575 Digital Blood Pressure Monitor is intended for use in the electromagnetic environment specified below. The customer or the user of the model LD-575 Digital Blood Pressure Monitor should assure that is used in such an environment.

Emission test	IEC 60601 test level	Compliance level	Electromagnetic environment-guidance
Electrostatic discharge (ESD) IEC 61000-4-2	±6 kV contact ±8 kV air	±6 kV contact ±8 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%
Electrical fast transient/burs IEC 61000-4-4		±2 kV for power supply lines	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	±1 kV differential mode	±1 kV differential mode	Mains power quality should be that of a typical commercial or hospital environment.
Voltage dips, short	(>95% dip in UT) for 0.5 cycle 40% UT (60% dip in UT)	<5% UT (>95% dip in UT) for 0.5 cycle	Mains power quality should be that of a typical commercial or hospital environment. If the user of the LD-575 Digital Blood Pressure Monitor Equipment requires continued operation during power mains interruptions, it is recommended that the LD-575 Digital Blood Pressure Monitor Equipment be powered from an uninterruptible power supply or a battery.
Power frequency magnetic field IEC 61000-4-8		3 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.
Conducted RF IEC 61000-4-6		3 Vrms	Portable and mobile RF communications equipment should be used no closer to any part of the LD-575 Digital Blood Pressure Monitor Equipment, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance $d = \begin{bmatrix} 3.5 \\ V_I \end{bmatrix} \sqrt{P}$ $d = \begin{bmatrix} 3.5 \\ V_I \end{bmatrix} \sqrt{P}$ 80 MHz to 800 MHz $d = \begin{bmatrix} 7 \\ E_I \end{bmatrix} \sqrt{P}$ 800 MHz to 2.5 GHz

	Radiated RF IEC 61000-4-3	10 V/m 80 MHz 2.5 GHz	to
	Note 1 At 801	MHz and	80

3 V/m Where P is the maximum output powe rating of the transmitter in watts (W) according to the manufacturer and and d

recommended separation distance in metres (m). b Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, should be less than the compliance level in each frequency range.d Interference may occur in the vicinity of equipment marked with the $((\bullet))$

transmitte

30MHz and 800MHz, 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 strength from fixed transmitters, such as base stations for radio 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 Model LD-575 Digital Blood Pressure Monitor is used exceeds the applicable RF compliance level above, the Model LD-575 Digital Blood Pressure Monitor should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the Model LD-575 Digital Blood Pressure Monitor.

^b Over the frequency range 150KHz to 80MHz, field strength should be

less than 3 V/m

Recommended separation distances between portable and mobile RF communications equipment and the LD-575 Digital Blood Pressure Monitor

The LD-575 Digital Blood Pressure Monitor is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the LD-575 Digital Blood Pressure Monitor can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the LD-575 Digital Blood Pressure Monitor as recommended below, according to the maximum output power of the communications equipment.

Rated maximum	Separation distance according to frequency of transmitter (m)		
output power of transmitter	150 kHz to 80 MHz	80 MHz to 800 MHz	800 MHz to 2.5 GHz
(W)	$d = \left[\frac{3.5}{V_1}\right] \sqrt{P}$	$d = \left[\frac{3.5}{E_1}\right] \sqrt{P}$	$d = \left[\frac{7}{E_1}\right] \sqrt{P}$
0.01	0.117	0.117	0.233
0.1	0.369	0.369	0.738
1	1.167	1.167	2.333
10	3.689	3.689	7.379
100	11.667	11.667	23.333

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.

QUALITY GUARANTEE

MODEL			
Warranty period	Two years from purchasing date		
Purchasing date			
	Name:	Telephone:	
Purchasing shop	Address:		
Customer	Name:	Telephone:	
Odstorner	Address:		

REQUIRING RECORD

Date	TROUBLE	SERVICE MAN
	1.During warranty period the repair could be made at any BPM repair department.	
	2. the following things not belong to warranty range:	
Guarantee Regulation	(1) Operating BPM different from procedures or instructions of the manual.	
	(2) The body is damaged artificially.	
	(3) Self-repairing or modifying the monitor construction in any way.	
	(4) Breakdown due to corrosion of battery leakage.	
	(5) Problem which occurs under natural calamity and other force	
	majeures.	

PERIODIC SAFETRY CHECKS

If you use the device with power adapter, preventive inspection and maintenance to be performed including the frequency of sunch maintenance

• Every time before use, please check the adapter, once damaged, never to use. Please clean the plug of adapter plug at least once a year. Too much

The manufacturer reserves the right to make technical changes without notice in the interest of progress.

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