

User Manual

Wrist Blood Pressure Monitor EBP-017



Healthcare-Manager.com

EASY AT HOME MEDICAL,LLC Any questions,please call toll-free: 1-855-822-6999 M-F 9 a.m.-5 p.m. CST E-MAIL:service@healthcare-manager.com

- Thank you very much for selecting easy tione Wrist Blood Pressure Monitor EBP-017.
- Please do read the user manual carefully and thoroughly so as to ensure the safe usage of this product.
- Please keep this manual in order to reference in future.

CATALOGUE

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♥ General Description

Thank you for selecting easy when wrist type blood pressure Monitor (EBP-017). The monitor features blood pressure measurement, pulse rate measurement and the result storage. The design provides you with two years of reliable service.

Readings taken by the EBP-017 are equivalent to those obtained by a trained observer using the cuff and stethoscope auscultation method.

This manual contains important safety and care information, and provides step-bystep instructions for using the product.

Read the manual thoroughly before using the product.

FEATURES:

- · Systolic Blood Pressure
- Pulse Rate
- · Diastolic Blood Pressure
- · Memory: Up to 60 pieces of records

♥ Safety Information

The signs below might be in the user manual, labeling or other components. They are the requirement of standard and using.

requiremen	it or standard and daing.		
(3)	Symbol for "THE OPERATION GUIDE MUST BE READ"	★	Symbol for "TYPE B APPLIED PARTS"
SN	Symbol for "SERIAL NUMBER"	***	Symbol for "ENVIRONMENT PROTECTION - Electrical waste products should not be disposed of
	Symbol for "MANUFACTURER"	X	with household waste. Please recycle where facilities exist. Check with your local authority or retailer for recycling
==	Symbol for "DIRECT CURRENT"		advice"
[[Symbol for "MANUFACTURE DATE"		

2



This device is intended for adult use only.

This device is intended for non-invasive measuring and monitoring of arterial blood pressure. It is not intended for use on extremities other than the wrist or for functions other than obtaining a blood pressure measurement.

Do not confuse self-monitoring with self-diagnosis. This unit allows you to monitor your blood pressure Do not begin or end medical treatment without asking a physician for treatment advice. If you are taking medication, consult your physician to determine the most appropriate time to measure your blood pressure. Never chance a prescribed medication without consulting your treatment of the propriate that the propriate that

priysician.

If the cuff pressure exceeds 40 kPa (300 mmHg),the unit will automatically deflate. Should the cuff not deflate when pressures exceeds 40 kPa (300 mmHg), detach the cuff from the wrist and press

the START/STOP button to stop inflation.

To avoid measurement errors, carefully read this manual before using the product.

The equipment is not AP/APG equipment and not suitable for use in the presence of a flammable anesthetic mixture with air of with oxygen or nitrous oxide.

The operator shall not touch output of batteries and the patient simultaneously.

Do not use the monitor under the conditions of strong electromagnetic field (e.g. medical RF

Do not use the monitor under the conditions of strong electromagnetic field (e.g. medical RF equipment) that radiates interference signal or electrical fast transient / burst signal. The user must check that the equipment functions safely and see that it is in proper working

condition before being used.

Please use ACCESSORIES and detachable partes specified/ authorised by MANUFACTURE.

Otherwise, it may cause damage to the unit or danger to the user/patients.

Manufacturer will make available on request circuit diagrams, component parts list etc.

This unit is not suitable for continuous monitoring during medical emergencies or operations. Otherwise, the patient's wrist and fingers will become anaesthetic, swollen and even purple due to a lack of blood.

Please use the device under the environment which was provided in the user manual. Otherwise, the performance and lifetime of the device will be impacted and reduced.

During use, the patient will be in contact with the cuff. The materials of the cuff have been tested and found to comply with requirements of ISO 10993-5:2009 and ISO 10993-10:2010. It will not cause any potential alergic reaction or contact injury.

The device doesn't need to be calibrated within the two years of reliable service.

Please dispose of ACCESSORIES, detachable parts, and the ME EQUIPMENT according to the

Please dispose of ACCESSORIES, detachable parts, and the ME EQUIPMENT according to the local guidelines.

When the device was used to measure patients who have common arrhythmias such as atrial or.

When the device was used to measure patients who have common armytimias such as atrail or ventricular premature beats or afrial fibrillation, the best result may occur with deviation. Please consult your physician about the result.

This device is contrainficiated for any female subject who may be suspected of, or is preparant.

Besides providing inaccurate readings, the effects of this device on the fetus are unknown.

The device has been evaluated clinically using manual culfistethoscope auscultation as the reference Blood pressure measurements determined with this device are equivalent to those obtained by a trained observer using the culfistethoscope auscultatory method, within the limits prescribed by the American National Standard, Manual, electronic, or automated sphygmomanometers."

▼ LCD Display Signal



SYMBOL	DESCRIPTION	EXPLANATION
SYS	Systolic blood pressure	High pressure result
DIA	Diastolic blood pressure	Low pressure result
Pul/min	ВРМ	Pulse/minute; heartbeats/minute
88	Memory	The displayed measurement values is from the memory.
611)	Movement error symbol	Shocking will result in inaccurate
□ +Lo	Low battery	Batteries are low and need to be replace
kPa	kPa	Measurement unit of the blood pressure (1kPa=7.5mmHg)
mmHg	mmHg	Measurement unit the blood pressure (1mmHg=0.133kPa)
•	Deflating	CUFF air is exhausting of deflating
88:88	Current time	Year/Month/Day,Hour/Minute
4	Grade	The grade of the blood pressure
₩2	Arrhythmia	Irregular heartbeat
*	Heartbeat	Heartbeat detection during the measurement

BEFORE YOU START

♥ Monitor Components





▼ List

1.Wrist Blood Pressure Monitor (EBP-017)

2. 2×AAA batteries 3.User manual

♥ Installing and Replacing the Batteries

1. Open the battery door.

2. Insert the batteries according to the polarity indications.

(Always select the authorized / specified 2×AAA batteries.)

3. Close the battery door.



Replace the batteries under following circumstances:

- □+Lo displays on the LCD.
- •The LCD display dims.
- . When powering on the monitor, the LCD doesn't light up.

- . Remove batteries if the device is not likely to be used for some time.
- . Worn batteries are harmful to the environment. so please do not dispose with other daily trash.
- Remove the old batteries from the device following your local recycling guidelines. . Do not dispose of batteries in fire. Batteries may explode or leak.

BEFORE YOU START BEFORE YOU START

♥ Setting Date, Time and Measurement Unit

Please proceed to time setting before your initial use so as to ensure each piece of record are labled with a time stamp. (Year Range: 2010-2050)

1. When the monitor is OFF, press the SET button to enter into the clock mode, then press and hold "SET" button for about 3 seconds to enter year setting.



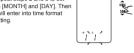
2.Press "MEM" button to change the numeral. Each press will increase the numeral by one in a cycling manner.



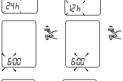
3.Press "SET" button to confirm the [YEAR]. Then the monitor diverts to [MONTH] and [DAY] setting.



4.Repeat steps 2 and 3 to set the [MONTH] and [DAY]. Then it will enter into time format setting.



5.Select the 24 hours time format or the 12 hours time format. Repeat steps 2 and 3 to set the [HOUR] and [MINUTE].



SEE

6.Repeat steps 2 and 3 to set the measurement unit.



7.After confirming the meausrement unit, the LCD will display all the settings you have done one by one and the monitor will shut off.

▼ Tie the Cuff

- Remove all accessories (watch, bracelet, etc) from your left wrist. If your physician has diagnosed you with poor circulation in your left wrist, use your right wrist.
- 2.Roll or push up your sleeve to expose the skin.
- 3.Apply the cuff to your left wrist with your palm facing up.
- 4. Position the edge of the cuff about 1-1.5cm (0.4"-0.6") from wrist joints.
- Fasten the wrist cuff around your wrist, leaving no extra room between the cuff and your skin. If the cuff is too loose, the measurement will be inaccurate.
- · Rest for 5 minutes before measuring.
- Wait at least 3 minutes between measurements. This allows your blood circulation to recover.
- For a meaningful comparison, try to measure under similar conditions. For example, take daily measurement at approximately the same time, on the same wrist, or as directed by a physician.



Keep the BP monitor the same level of your heart

♥ Start Measurement

 After correctly positioning the cuff, press START/STOP button to turn on the monitor, and it will complete the measurement process.



LCD Display.



Adjust to zero.



Inflating and measuring.



Display and save the measuring result.



 Press START/STOP button to turn off the monitor. Otherwise, the monitor will shut off within 1 minute.



♥ Recall the Records

1. When the blood pressure monitor is off. press "MEM" button to access the memory, it will display the latest record first when the records are less than three groups. When there are three or more than three groups ,it will display the average value of the latest three records first.



2. Press "MEM" button again, it will display the latest measurement record. Press "MEM" button to rotate the history records. Press SET button to look back the record. You can check 60 groups of record totally. When it displays the record, it will shutdown in 1minute if has no operation.



The exact time and date of the measurement will display alternatively.



3. When there has no record, the LCD displays "---".



The most recent record (1) is shown first. Each new measurement is assigned to the first (1) record. All other records are pushed back one digit (e.g., 2 becomes 3, and so on), and the last record (60) is dropped from the list.

INFORMATION FOR USER

DATA MANAGEMENT

♥ Delete the Records

When you did not obtain the accurate measurement, you can clear all the measuring results by following steps below.

 Press and hold both the "MEM" button for 3 seconds when the monitor is in the memory recall mode, the flash display "dEL ALL" will show.



2.When the LCD display blinking "dEL ALL", press "SET" button to confirm clearing. The LCD will display "dEL dONE", indicating that memory clearing is complete.



 If you wish to give up clearing, press the START/STOP button to turn off the monitor.

♥ Tips for Measurement

Measurements may be inaccurate if taken in the following circumstances.



♥ Maintenance

To obtain the best performance, please follow the instructions below.



Put in a dry place and avoid the sunshine



Avoid touching water, clean it with a dry cloth in case.



Avoid dusty and unstable temperature environment



and collisions

Use wet cloths to remove dirt

Avoid intense shaking



O ALLITION

- If you have any problems with this device, such as setting up, maintaining or using, please contact the SERVICE PERSONNEL of service Don't open or repair the device by yourself.
- Please report to several from if any unexpected operation or events occur.

♥ What are systolic pressure and diastolic pressure?

When ventricles contract and pump blood out of the heart, the blood pressure reaches its maximum value in the cycle, which is called systolic pressure. When the ventricles relax, the blood pressure reaches its minimum value in the cycle, which is called diastolic pressure.





♥ What is the standard blood pressure classification?

The chart on the right is the standard blood pressure classification published by American Heart Association (AHA).

AHA Home Guideline for Jpper Limit of Normal B			
SYS	135 mm Hg		
DIA	85 mm Hg		

Blood Pressure Category	Systolic mmHg (upper#)		Diastolic mmHg (lower#)
Normal	less than 120	and	less than 80
Prehypertension	120-139	or	80-89
High Blood Pressure (Hypertension) Stage 1	140-159	or	90-99
High Blood Pressure (Hypertension) Stage 2	160 or higher	or	100 or higher
Hypertensive Crisis (Emergency care needed)	Higher than 180	or	Higher than 110

Please consult a physician if your measuring result falls outside the range. Please note that only a physician can tell whether your blood pressure value has reached a dangerous point.

▼ Irregular Heartbeat Detector

An irregular heartheat is detected when a heartheat rhythm varies while the unit is measuring the systolic and diastolic blood pressure During each measurement, this equipment records the heartheat intervals and works out the standard deviation. If the calculated value is larger than or equal to 15, the irregular heartheat symbol appears on the symbol when the measurement results are displayed.

CAUTION -

The appearance of the IHB icon indicates that a pulse irregularity consistent with an irregular heartbeat was detected during measurement. Usually this is NOT a cause for concern. However, if the symbol appears often, we recommend you seek medical advice. Please note that the device does not replace a cardiac examination, but serves to detect pulse irregularities at an early stage. ABOUT BLOOD PRESSURE TROUBLESHOOTING

Why does my blood pressure fluctuate throughout the day?

Individual blood pressure varies multiple times everyday, it is also affected by the way you tie your cuff and measurement position, so please take the measurement under the same conditions.

2.If the person takes medicine, the

pressure will vary more.

3.Wait at least 3 minutes for

another measurement.

▼ Why do I get a different

blood pressure at home compared to the hospital?

The blood pressure is different even throughout the day due to weather emotion, exercise etc. Also, there is the "white coat" effect, which means blood pressure usually increases in clinical settings.

▼ Is the result the same if measuring on the right wrist?

It is ok for both wrists, but there will be some different results for different people. We suggest you measure the same wrist every time.



What you need to pay attention to when you measure your blood pressure at home:

If the cuff is tied properly.
If the cuff is too tight or too loose.
If the cuff is tied on the wrist.
If you feel anxious.
Taking 2-3 deep breaths before
beginning will be better for measuring.
Advice:Relax yourself for 4-5
minutes until you calm down.



This section includes a list of error messages and frequently asked questions for problems you may encounter with your wrist blood pressure monitor. If the products not operating as you think it should, check here before arranging for servicing.

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PROBLEM	SYMPTOM	CHECK THIS	REMEDY
	Display is dim or	Batteries are exhausted.	Replace with new batteries
No power	will not light up.	Batteries are inserted incorrectly.	Insert the batteries correctly
Low batteries	CI+Lo Show on the display	Batteries are low.	Replace with new batteries
	E 1 shows	The cuff is not secure.	Refasten the cuff and then measure again.
	E 2 shows	The cuff is very tight	Refasten the cuff and then measure again.
	E 3 shows The pressure of the cuff is excess.		Relax for a moment and then measure again.
Error massage	E 10 or E 11 shows	The monitor detected motion while measuring.	Movement can affect the measurement.Relax for a moment and then measure again.
	E 20 shows	The measurement process does not detect the pulse signal.	Loosen the clothing on the wrist and then measure again.
	E 21 shows	Measure incorrectly.	Relax for a moment and then measure again.
	EExx,shows on the display.	A calibration error occurred.	Retake the measurement. If the problem persists, contact the retailer or our customer service department for further assistance. Refer to the warranty for contact information and return instructions.

SPECIFICATIONS

Power Supply	Battery powered mode: 2×AAA batteries (3V)
Display Mode	Digital LCD V.A 31.5mm×44mm(1.25"×1.73"
Measurement Mode	Oscillographic testing mode
Measurement Range	Pressure: 0kPa-40kPa (0mmHg-300mmHg) pulse value:(40-199)beat/minute
Accuracy	Pressure: 5°C-40°C(41°F-104°F)within±0.4kPa(3mmHg) pulse value:±5%
Normal working condition	Temperature:5℃ to 40℃(41℉ to 104℉) Relative humidity: ≤85%RH Atmospheric pressure: 86kPa to 106kPa
Storage & transportation condition	Temperature:-20 C-60 C (-4 T - 140 T) Relative Humidity: 10%RH-93%RH Atmospheric Pressure: 50kPa-106 kPa
Measurement perimeter of the wrist	About 13.5cm-21.5cm (5%-8%)
Weight	Approx.112g(3.95oz)(Excluding batteries)
External dimensions	Approx. 68mm×75mm×31mm(2.68"×2.95"×1.22 (Excluding the cuff)
Attachment	2×AAA batteries,one storage,user manual
Mode of operation	Continuous operation
Degree of protection	Type B applied part
IP Classification	IP×0
Software version	V01
Device classification	Internally Powered ME Equipment

♥ Contact Information

For more information about our products, please visit .//inathcare-//funager.com, or call easy//ine Customer 1-866-822-6999 M-F 9 a.m.-5 p.m.CST, usual problems and customer download, easy//ine will serve you anytime.

♥ Complied Standards List

Risk management	ISO 14971:2007
Labeling	EN 980:2008
User manual	EN 1041:2008
General Requirements for Safety	IEC 60601-1:2005
Electromagnetic compatibility	IEC 60601-1-2:2007
Non-invasive Sphygmomanometers General Requirements	AASI/AAMI SP10:2002/A1:2003/A2:2006/(R)2008

▼ FCC Statement

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.

▼ EMC Guidance

Table 1 Guidance and MANUFACTURER's declaration - ELECTROMAGNETIC EMISSIONS- for all ME EQUIPMENT and ME SYSTEMS

Guidance and manufacturer's declaration - electromagnetic emissions

The device is intended below. The customer such an environment.	for use in the ele or the user of the	ctromagnetic environment specified device should assure that it is used in		
Emissions test	Compliance	Electromagnetic environment - guidance		
RF emissions CISPR 11	Group 1	The device 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 emissions CISPR 11	Class B			
Harmonic emissions IEC 61000-3-2	Not applicable			
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Not applicable			

Table 2 Guidance and MANUFACTURER's declaration – electromagnetic IMMUNITY – for all ME EQUIPMENT and ME SYSTEMS

	nded for use in the element user of the device s		ment specified below. s used in such an
IMMUNITY 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/burst IEC 61000-4-4	±2 kV for power supply lines ±1 kV for input/output lines	Not applicable	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	±1 kV line(s) to line(s) ±2 kV line(s) to earth	Not applicable	Mains power quality should be that of a typical commercial or hospital environment.
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	45% U7 (>95% dip in U _T) for 0.5 cycle 40% U7 (60% dip in U _T) for 5 cycles 70% U7 (30% dip in U _T) for 25 cycles 45% U7 (>95% dip in U _T) for 5 s	Not applicable	Mains power quality should be that of a typical commercial or nospital environment. If the user of the device requires continued operation during power mains interruptions, it is recommended that the device be powered from an uninterruptible power supply or a battery.
Power frequency (50/60Hz) magnetic field IEC 61000-4-8	3A/m	3A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment

Table 4 Guidance and MANUFACTURER's declaration – electromagnetic IMMUNITY – for ME EQUIPMENT and ME SYSTEMS that are not LIFE-SUPPORTING

The customer or to	he user of the device	e should assure th	vironment specified below. at it is used in such an environment.
IMMUNITY test	IEC 60601 TEST LEVEL	Compliance level	Electromagnetic environment - guidance
			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.
Conducted RF IEC 61000-4-6	3 Vrms 150 kHz to 80 MHz	Not applicable	Recommended separation distance $d = [\frac{3.5}{V_1}]\sqrt{P}$
Radiated RF IEC 61000-4-3	3 V/m 80 MHz to 2.5 GHz	3 V/m	$d=1.167\sqrt{P}$ 80 MHz to 800 MHz $d=2.333\sqrt{P}$ 800 MHz to 2,5 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. ^b
			Interference may occur in the vicinity of equipment marked with the following symbol:

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 penale.

Field shrengths from fixed transmitters, such as base stations for radio (cellular / cordiess) feedprone and and mobile radio, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted the restercibly with accuracy. To assess the extinctionage filed environment due for bear FP ammanties, an electromage filed the extent of the rest of the res

Table 6 Recommended separation distances between portable and mobile RF communications equipment and the ME EQUIPMENT or ME SYSTEM – for ME EQUIPMENT and ME SYSTEMS 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)		
	150 kHz to 80 MHz	80 MHz to 800 MHz	800 MHz to 2.5 GHz
	$d = \left[\frac{3,5}{V_1}\right]\sqrt{P}$	$d = 1.167 \sqrt{P}$	$d=2.333\sqrt{P}$
0.01	Not applicable	0.117	0.233
0.1	Not applicable	0.369	0.738
1	Not applicable	1.167	2.333
10	Not applicable	3.690	7.378
100	Not applicable	11.67	23.33

For transmitters rated at a maximum output power not listed above, the recommended separation distance of 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 80MHz and 800MHz, 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.