Fully Automatic Upper Arm

Blood Pressure Monitor

Model Number: C02 ------

USER'S MANUAL



Shenzhen Jamr Technology Co., Ltd.
A101-301, D101-201, Jamr Science & Technology Park, No. 2 Guiyuan Road, Guixiang Community, Guanlan Street, Longhua District, 518100 Shenzhen, PEOPLE'S REPUBLIC OF CHINA

LEGAL DISCLAIMER

Disclaimer regarding the blood pressure monitor, the device is Over-the-Counter (OTC), and the measurements are for informational purposes only and are not intended to diganose, treat, cure or prevent any disease or health condition.

The model number of this device is: CO2

Has been approved by FDA, certification 510K number: K200437 In order to get more accurate measurement results, please pay attention to the correct measurement requirements.

SAFETY INFORMATION AND PRECAUTIONSINTRODUCTION

- Read all instructions.
- 2. This device is not intended to be a diagnostic device.
- 3. To avoid any possibility of accidental strangulation, do not drape tubing around your neck. Keep this unit away from children and pets.
- 4. This device is intended for use on upper arm to obtain a blood pressure measurement. This device is not suitable for continuous monitoring or frequently measuring your blood pressure. Do not use the device for any other purpose
- 5. Consult your physician before measuring blood pressure, especially if you have any of following conditions: hypertension, diabetes, arteriosclerosis, vascular disease, or any conditions affecting circulation.
- 6. Do not change medication or diagnose based on the measurement from the device. Only a physician is auglified to diagnose and treat high blood pressure.
- 7. Do not put the cuff on wounds. If you feel uncomfortable when the cuff inflates. press START/STOP button to stop inflation.
- 8. If heart arrhythmias occur during a measurement, it may cause inaccurate readings or fail to detect blood pressure. Consult your physician to evaluate vour blood pressure.
- 9. Power the device with either 3 AA batteries or an AC adapter. Do not use both. Remove batteries if the device is not likely to be used some time. Avoid strong electromagnetic field radiated interference signals and electrical burst signals.
- 10. Do not use this device near flammable gases or flammable liquids are present
- 11. Use this device under Operating Conditions specified in the SPECIFICATIONS.
- 12. Do not disassemble or attempt to repair this device or any of its components.
- 13. Use only Jamr authorized parts and accessories under Operating Conditions specified in the SPECIFICATIONS. Consult your physician to evaluate yourblood pressure.

WARRANTY

Your Blood Pressure Monitor is covered by a one-year manufacturer's warranty against defects from the original purchaser from the date of purchase. Proof of purchase is required.

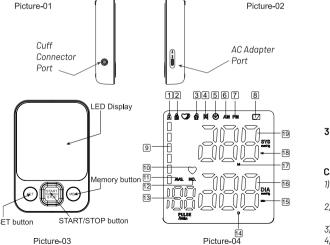
Type-C charaina cable is excluded from the warranty. The warranty is void if the product has been subjected to mechanical damage or mistreatment, such as immersion. This warranty is in lieu of all other warranties, and limits the liability of the manufacturer.

INTRODUCTION

The device is a fully automatic blood pressure monitoring device it is intended to measure the systolic and diastolic blood pressure as well as the pulse rate by using the oscillometric method.

The device is used for adult that gaes are more than 12 years old and used for over-the-counter it can be used in medical facilities and at home.

DEVICE



The symbols on the LED display

3.Movement error symbol

5.Cuff self-detecting function 6.AM symbol

8.Low battery symbol 9.WHÓ function symbol

10.Heartbeat symbol (Flashes during measurement

12.NO. symbol 13.Pulse symbol 14. Date symbol 15.Minute symbol

16.Diastolic blood pressure

17.Month sýmbol 18. Hour symbol 19.Systolic blood pressure

Features of Model C02

1.Average value symbol

Cuff self-checking function 2. Double users: 2 x 120 sets memory 3. Average value function 4. Low battery display

2 User R

4. Mute symbol

5. WHO function 6. Auto power-off 7. External power adapter support 8. Time stamp in memory

1. Connect the Cuff Tube to the Monitor.

Insert the tube into the opening on the left side of the monitor.

- 2. Use only AA batteries. (For eco-friendly and transport safe, batteries and adapters are not included in the product package.)
- 1) Open battery cover.
- 2) Install the 3AA batteries by matching the correct polarity.
- Close the cover.
- 4) Replace the batteries when Low Battery Indicator a shows, the display dims or the display does not light up.





3. Option: External AC Adapter (Not Included)

You may also operate this monitor using an external AC adapter (output 5V DC/1A with Type-C Cable).

- 1) Remove batteries if the device is not likely to be used for over three months. Battery leakage can damage your device.
- 2) Do not dispose of batteries with other ordinary trash. They are harmful to the environment.
- 3) Do not dispose of batteries in fire. They may explode.
- 4) Follow your local recycle guidelines to properly dispose of used batteries.

SYSTEM SETUP

Setting the User:

Press the SET button to select User A or User B. When display A (/B) on the screen press the SET button to switch to user B (/A). Press the START/STOP button to

Setting the Year/Month&Date/Time:

Long press the SET button for more than 6s, and then you can start to set.

Setting the Year:

Initial year is 2022, when the year display is flashing, press the MEM button, the vear will increase by I year each, hold the MEM button and it will increase continuously 1 by 1, until 2049, and then rollover to 2022, once the year set is OK, press SET button to confirm.

Setting Month/Date:

Initial Month/Date is 01/01, when the Month display is flashing, press the MEM button, the month will increase by 1, press SET button to confirm, and do in the same way to set the date. Press SET button to confirm.

When the hour display is flashing, press the MEM button, the hour will increase by 1, press SET button to confirm, and do in the same way to set the minute. Press SET button to confirm.

Record Delete:

When you checking the memory data, long press MEM button to delete existing user measurement data.

If you decide to delete the all memory, please keep the memory in another way. incase you need it some days later. Take the battery out won't lead to a memory

TIPS FOR MEASUREMENT

1. Take measurements under the similar conditions to have consistent and comparable measurements.

• within an hour of getting up in the morning, after emptying bladder and before

Position Yourself

2. Use the same arm each time to measure your blood pressure.

4 Wait at least three minutes between measurements is recommended

5. Avoid taking measurements under following conditions that may cause

1) Fasten the cuff on your left upper arm about one inch (2-3 cm/1") away from

your elbow. Position the tube off-center toward the inner side of the arm. The

artery mark Φ (on the edge of the cuff) is close to the artery upper arm in line

2) The cuff should snug but not too tight. You should be able to insert one finger

3) The middle of the cuff should be at the level of your heart.

3. The hest time to take measurement

within one hour of eating or drinking

when talking or moving your fingers

when needing to use the restroom

in a very cold or hot environment

within 20 minutes after taking a bath or a shower

before bed at night.

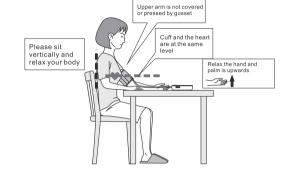
MEASUREMENT

with the little finaer.

between the cuff and your arm.

Fasten the Cuff

Sit comfortably with your arm resting on a flat surface, palm up, feet flat on the floor and leas uncrossed.



Start the Measurement

- 1) Select your user ID. Press "SET" button to toggle which user you want to measure. Press "START/STOP" button to turn off the monitor.
- 2) Press "START/STOP" button to turn on the blood pressure monitor and the measurement starts. It will display Systolic/Diastolic blood pressures and pulse rate when the measurement is done.
- 3) The monitor will display 🕽 if irregular heartbeat is detected during the measurement.
- 4) Press "START/STOP" button to power off, otherwise the monitor will
- automatically turn off within one minute of the reading.

Note: If you feel uncomfortable when the cuff inflates, press START/STOP button to stop inflation.

SPECIFICATIONS Technical alterations reserved.

Size(mm): 126 (W)×85 (L)×28 (H) Size(inch):4.84"(W)×3.23"(L)×0.98"(H)

Display: LED Digital Display Display(mm): 62.5×63.5mm Display(inch): 2.46"x2.5"

Weight: 261a (Batteries and AC adapter are not included): Accessories: 1×Main Device. 1×Users manual, 1×Type-c Cable, 1×Wide range rigid cuff 8.7" - 15.7" (22 - 40 cm)

Automatically power off: 60 seconds Measuring method: Oscillometric Pressure sensor: Resistive Measuring resolution: 1 mmHa

DIA Measurina range: 40-200mmHa SYS Measurina range: 60-255mmHa Pulse Measuring range: 40 to 170 per minute Cuff pressure display range:<295mmHq Atmospheric pressure range: 70kPa~106kPa

Operating Conditions: Temperature: 5°C to 40°C: Humidity: 15% to 93% RH: Storage And Shipping Conditions:Temperature: -25 °C to +70 °C; Humidity:≤93% RH; Memory: Automatically stores the last 120 measurements for 2 users (total 240) Accuracy: Pressure within ± 3 mmHa / pulse ± 5 % of the reading

Power source: a) 3×AA batteries, 4.5 V b) AC adapter INPUT: 100-240VAC 50/60HZ OUTPUT: 5V DC 1A

Expected service life of the device and accessories: 5 years

CERTIFICATIONS

This device is manufactured to meet the American blood pressure monitors: IFC 80601-2-30 • IFC60601-1-11 • IFC60601-1

Electromagnetic compatibility:

Device fulfills the stipulations of the International standard: IEC60601-1-2

CARE AND MAINTENANCE

- 1. Store the monitor kit in a dry place.
- 2. Clean the monitor with a dry cloth as needed. Avoid contact with water.
- 3. Clean the cuff with a damp cloth as needed. Do not wash the cuff.
- 4. Handle the tube carefully. Do not pull or twist the tube.
- 5. Remove batteries if the device is not likely to be used for some time.

MEMORY

1. View Records

- 1) Press "SET" button to togale which user you want to view history for.
- 2) Press the "MEM" button to display the average value of the latest three records. Continue to press the "MEM". The monitor will display previous records, starting with the latest record first.

2. Delete Records

SYMBOL CAUSE

per minute)

Warning: ALL records associated with the user will be deleted! Please write down as needed before operating the steps below.

- 1) Press "SET" button to select the user.
- 2) Press "MEM" button to view the user's history.
- 3) Hold down the "MEM" button for three seconds. All records associated with the user will be erased.

CORRECTION

TROUBLE SHOOTING

			WITH
No display appears	Weak battery or improper placement	Replace both batteries with new ones. Check the battery installation for proper placement of the battery polarities.	systoli The blo
Er1	Sensor abnormal	Please make sure the cuff pressure is drained and then measure again. If the error is still displayed, please send it to local distributor	two for
Er 2	Monitor could not detect pulse wave or cannot calculate the blood pressure data	Tie the cuff correctly and make the measurement again. If the error is still displayed, please send it to local distributor	its max as the
Er 3	Measurement result is abnormal	Occasionally-measure for one more time/ Always - send it to local distributor	cycle.
Er 4	Too loose cuff or air leakage	Tie the cuff correctly and make sure the air plug is properly inserted in the unit	Blood The fiv
Er 5	The air tube is crimped or the cuff is tied too tight.	Correct it and make the measurement again	Blood press
Er 6	The sensor is sensing great fluctuation in the pressure	Please keep quiet and don't move	Best
Er7	The pressure that the sensor sensing is over the limit	Please send back to the local distributor	Normal Elevated
Er 8	The demarcation is incorrect or the device has not been demarcated	Please send back to the local distributor	High Blood
HI	The pulse rate exceeds the upper limit (>170 per minute)	Beyond the measurement range, normal reminder	High Blood
L0	The pulse rate is less than the lower limit (<40	Beyond the measurement range, normal reminder	27

The following symbol will appear on the display when measuring abnormal Blood Pressure Fluctuations

olem	Check	Cause and solutions
oower	Check the battery power	Replace new one
	Check the polarity position	Installation for proper
		placement of the batteries
		polarities
nflation	Whether the plug insert	Insert into the air socket tightly
	Whether the plug broken or leak	Change a new cuff
ind stop working	Whether move the arm when inflate	Keep the body peaceful
	Check if chatting when measured	Keep quite when measure
leak	Whether the cuff wrap too loose	Wrap the cuff tightly
	Whether the cuff is broken	Change a new cuff
\ Please contact	the distributor if you can't solve the prob	olem, do not disassemble the unit
ourself!		

BASICS ABOUT BLOOD PRESSURE

What is the Blood Pressure? What is the difference between olic pressure and diastolic pressure?

blood pressure is the pressure of the blood within the arteries, the result of forces. The first force (systolic pressure) occurs as blood pumps out of the t and into the arteries that are part of the circulatory system when it reaches aximum value in the cycle. The second force (diastolic pressure) is created heart rests between heart beats when it reaches its minimum value in the

d Pressure Categories

ive blood pressure ranges are categorized by the WHO Heart Association.

Systolic mmhg	Diastolic mmhg
≤119	≤79
120-129	80-84
130-139	85-89
140-159	90-99
160-179	100-109
≥180	≥110
	≤119 120-129 130-139 140-159 160-179

EMC DECLARATION

1) Individual blood pressure varies throughout the day. It is normally lower at night while you are sleeping. Your blood pressure starts to rise a few hours before you wake up, continues to rise during the day, usually peak in the middle of the afternoon. Then in the late afternoon and evening, your blood pressure begins dropping again. 2) Certain medications can affect your blood pressure.

- ME system for magnetic resonance imaging, where the intensity of EM 3) Strong emotions, particularly stress and anxiety, can cause your blood pressure to disturbances is high. Warning: Use of this equipment adjacent to or stacked with other equipment
- 4) Weather is also a factor to affect your blood pressure, usually higher in the cold and lower in the heat.
- 5) It is normal to see temporary increases in blood pressure during exercise.

Different Blood Pressures from the Hospital and from Home Possible causes

- 1) Your blood pressure can vary throughout the day due to many factors listed in
- 2) Check the following situations that could be causing a false measurement:
- The cuff is too loose or too tight.
- The cuff is not secured properly
- The cuff is not placed correctly around the arm.
- Your arm is not rest correctly.
- Your leas are crossed.
- You measure too quickly after a previous reading.

Irregular Heartbeat Detector

No Sterilize requirement

Mode of operation: continuous

During each measurement, if the device has detected irregular heart beats, the Irregular Heart beat symbol will be displayed.

The following symbols may appear in this manual, on the Digital Blood Pressure Monitor CO2.

Not category AP / APG equipment

\sim	Date of manufacture.			Manufacturer
SN	Specifies serial number	er – 🥰 Keep dry	†	Type BF applie
===	Direct current	Handle gently	<u>††</u>	Put up
I	Fragile [Avoid the sun
Z	DISPOSAL: Do not disp waste. Collection of su is necessary.	pose this product as unso uch waste separately for s	rted mi special	unicipal treatment
(3)	Follow instructions for	ruse	№	MR unsafe.

IP21 The degree of avoid ingress of water or particulate matter into ME

and so on.

that they are operating normally.

this equipment could result.

OR TYPE REFERENCE).

equipment and result in improper operation."

1. All necessary instructions for maintaining BASIC SAFETY and ESSENTIAL PERFORMANCE with regard to electromagnetic disturbances for the excepted service life.

The ME EOUIPMENT or ME SYSTEM is suitable for home healthcare environments

Warning: Don't near active HF surgical equipment and the RF shielded room of an

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

Warning: Use of accessories, transducers and cables other than those specified

or provided by the manufacturer of this equipment could result in increased

electromagnetic emissions or decreased electromagnetic immunity of this

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 the Blood Pressure Monitor (CO2), including cables

specified by the manufacturer. Otherwise, degradation of the performance of

If any: a list of all cables and maximum lengths of cables (if applicable),

transducers and other ACCESSORIES that are replaceable by the RESPONSIBLE

ORGANIZATION and that are likely to affect compliance of the ME EQUIPMENT or

ME SYSTEM with the requirements of Clause 7 (EMISSIONS) and Clause 8

(IMMUNITY). ACCESSORIES may be specified either generically (e.g. shielded

cable, load impedance) or specifically (e.g. by MANUFACTURER and EQUIPMENT

If any: the performance of the ME EQUIPMENT or ME SYSTEM that was

determined to be ESSENTIAL PERFORMANCE and a description of what the OPERATOR can expect if the ESSENTIAL PERFORMANCE is lost or degraded due

to EM DISTURBANCES (the defined term "ESSENTIAL PERFORMANCE" need not

2. Guidance and manufacturer's declaration -electromagnetic emissions and

	Guidance and manufacturer's declaration - electromagnetic emissions			
,	Emissions test	Compliance		
	RF emissions CISPR 11	Group 1		1
,	RF emissions CISPR 11	Class B		fo
1	Harmonic emissions IEC 61000-3-2	Class A		
	Voltage fluctuations/ flicker emissions IEC 61000-3-3	Applied		
+	Table 2		_	С

Outdoors and manufactured declaration also accompanies anticional

Guidance and manufacturer's declaration - electromagnetic Immunity				
Test frequency	Modulation	IMMUNITY TEST LEVEL (A/m)		
30 kHz	CW	8		
134,2 kHz	Pulse modulation b) 2,1 kHz	65 °)		
13,56 MHz	Pulse modulation b) 50 kHz	75 °)		

a) This test is applicable only to ME EQUIPMENT and ME SYSTEMS intended for use in the HOME HEALTHCARE ENVIRONMENT. h) The carrier shall be modulated using a 50 % duty cycle square wave signal. c) r.m.s., before modulation is applied.

Radiated RF

IEC61000-4-3

Proximity magnetic

IFC 61000-4-39

Guidance and manufacturer's declaration - electromagnetic Immunity			
Immunity Test	IEC 60601-1-2 Test level	Compliance level	
Electrostatic discharge (ESD) IEC 61000-4-2	±8 kV contact ±2 kV, ±4 kV, ±8 kV, ±15 kV air	±8 kV contact ±2 kV, ±4 kV, ±8 kV, ±15 kV air	
Electrical fast transient/burst IEC 61000-4-4	Power supply lines: ±2 kV input/output lines: ±1 kV 100 kHz repetition frequency	Power supply lines: ±2 kV	
Surge IEC 61000-4-5	line(s) to line(s): ±0.5 kV line(s) to earth: ±2 kV line(s) to lines(s): ±1 kV	line(s) to line(s): ±0.5 kV line(s) to line(s): ±1 kV.	
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	0% 0.5 cycle At 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° 0% 1 cycle And 70% 25/30 cycles Single phase: at 0 0% 300 cycle	0% 0.5 cycle At 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° 0% 1 cycle And 70% 25/30 cycles Single phase: at 0 0% 300 cycle	
Power frequency magnetic field IEC 61000-4-8	30 A/m 50Hz/60Hz	30 A/m 50Hz/60Hz	
Conduced RF IEC61000-4-6	150KHz to 80MHz: 3Vrms 6Vrms (ISM and amateur radio bands) 80% Am at 1kHz	150KHz to 80MHz: 3Vrms 6Vrms (ISM and amateur radio bands) 80% Am at 1kHz	

80 MHz - 27 GHz

80 % AM at 1 kHz

134.2 kHz: 65A/m

30 kHz: 8A/m

80 MHz - 2.7 GHz

80 % AM at 1 kHz

134.2 kHz: 65A/m

13.56 MHz: 7.5A/m

30 kHz: 8A/m

NOTE 1: UT is the a.c. mians voltage prior to application of the test level

Guidance and manufacturer's declaration - electromagnetic Immunity						
ted RF 100-4-3 est cations LOSURE INIT UNITY o reless nication oment)	Test Frequency (MHz)	Band ^{a)} (MHz)	Service a)	Modulation ^{a)}		
	385	380-390	TETRA 400	Pulse modulation ^{b)} 18 Hz	_	
	450	430-470	GMRS 460, FRS 460	FM ^{c)} ±5 kHz deviation 1 kHz sine		
	710	704-787	LTE Band 13, 17	Pulse		
	745			Modulation ^{b)} 217 Hz		
	780			211112		
	810		GSM 800/900,	Pulse Modulation ^{b)}		
	870	800-960	TETRA 800,	19 Uz		

LTE Band 13. GSM 800/900 TETRA 800. iDFN 820. CDMA 850. LTE Band 5 GSM 1800: CDMA 1900: 1845 GSM 1900: ITE Band 1.3. 4. 25: UMTS Bluetooth. 2450 802 11 b/a/n. RFID 2450. ITF Band 7 5500

EOUIPMENT or ME SYSTEM may be reduced to 1 m. The 1 m test distance is permitted by IEC 61000-4-3.

18 Hz

Modulationb

Pulse

Modulation

TEST LEVEL

a) For some services, only the uplink frequencies are included.

b) The carrier shall be modulated using a 50 % duty cycle square wave signal

c) As an alternative to FM modulation, the carrier may be pulse modulated using a 50 % duty cycle square wave signal at 18 Hz. While it does not represent actual modulation, it would be worst case.