FT-500(R/L) FULLY AUTOMATED BLOOD PRESSURE MONITOR

OWNER'S MANUAL

- ONE TOUCH operation
- WIDE LED display
- QUICK measure
- HIGH accuracy
- THERMAL HIGH SPEED printer
- VOICE message
- CARD using





The device bears the CE label in accordance with the provisions of Medical Device Directive 93/42/EEC.

THE PERSONS RESPONSIBLE FOR PLACING DEVICES ON THE EC MARKET UNDER MDD 93/42/EEC



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INTRODUCTION

We highly appreciate that you chose our company's product.

You are kindly requested to be familiar with these directions before using this product and always keep it together with the product. In case you are not sure about any directions or problems arising while using the product, please contact our service center.

We will provide you with detailed instructions.

1. INTEDED USE

FT-500 (R/L) Automatic Blood Pressure Monitor is designed to measure systolic and diastolic blood pressure and pulse rate of Persons who are 18 years and older using the oscillometric method on a cuffed arm.

FT-500 L measure the Left arm.

FT-500 R measure the Right arm.

- · Target user: Persons who are 18 years and older
- · This medical device is not for home use

2. WORD DEFINITIONS

To ensure safe operation and long term performance stability, it is essential that you fully understand the functions, operating and maintenance instructions by reading this manual before operating your unit.

Particular attention must be paid to all warnings, cautions and notes incorporated herein.

The following conventions are used throughout the manual to denote information of special emphasis.

Warning



"Warning" indicates important information to the presence of a hazard which may cause severe personal injury, death of substantial property damage if the warning is ignored.

Caution



"Caution" indicates important information to the presence of a hazard which may cause minor personal injury or property damage if the caution is ignored.

Note



"Notice" indicates important information to notify installation, operation or maintenance of this device. "Notice" is important but not hazard-related. Hazard warnings are not included here.

3. CLASSIFICATION AND COMPLIANCE

- 1) This device is classified as;
- Class 1 type-BF against electric shock
- Ordinary equipment without protection against ingress of water
- Equipment not suitable for use in presence of a flammable anesthetic mixture by standard of EN 60601-1: 2006(Basic safety and essential performance of Medical Electrical Equipment)
- 2) This device is complied with Class A for Noise-Emission, Level B for Noise-immunity, by standard of IEC 60601-1-2:2007(Electromagnetic Compatibility Requirements).
- 3) This device is complies with the EN 1060-1: 1995+A2:2009 Non-invasive Sphygmomanometers general requirements as well as EN 1060-3: 1997+A2:2009 supplementary requirements for electromechanical blood pressure measuring systems.

4. SAFETY PRECAUTIONS

This device is designed and manufactured with consideration of safety of the operator and subject and also to the reliability of the unit.

The following precautions must be observed for additional safety;

- 1) The unit must be operated only by, or under supervision of a qualified person with our company or our distributors.
 - 2) This device is specified as Class 1 type BF unit under the standard of IEC 60601-1:2005(Safety of Medical Electrical Equipment).
- O Do not touch or handle inner side of the system at any time.
- The INTERNAL ELECTRICAL POWER SOURCE is to be used if the integrity of the PROTECTIVE EARTH CONDUCTOR or the protective earthing system in the installation is in doubt.
- ▲ 3) Do not modify the unit. If any modification is needed, ask our company or its authorized dealer for service.
 - 4) The unit has previously been adjusted in the factory for optimum performance.
- Do not attempt to adjust switches or any other things except those specified in this manual for operation.
- 5) If you have experienced any trouble with the unit, switch it off immediately, and contact our company or its authorized dealer for assistance.
 - 6) If you plan to connect any device of other manufacturers electrically or mechanically to the unit, contact our company or its authorized dealer for instructions before doing so.
 - When you connect computer or other system to the unit (RS-232C), the attached systems should be those certified by IEC 60950 or equivalent standards for data processing equipment.
 - Configurations shall comply with the system standard IEC 60601-1:2005.
 - Everybody who connects additional equipment to the signal input part or signal output part configures a medical system standard IEC 60601-1:2005.
 - If in doubt, consult the A/S department of local distributor.

- 7) Avoid the following environments for storage;
 - Where the ambient temperature falls -20°C or exceeds 60°C.
 - Where the atmospheric pressure falls below 70kPa (700mbar) or exceeds 106kPa (1060mbar).
 - Where the humidity is over 95% non-condensing.
 - Where the unit is exposed to spray or splashing water.
 - Where the unit is exposed to dust.
 - Where the unit is exposed to water vapor.
 - Where the unit is exposed to salty atmosphere.
 - Where the unit is exposed to explosive gas.
 - Where the unit is exposed to excessive shocks or vibrations.
 - Where the angle of inclination of mounting surface exceeds 10 degrees.
 - Where the unit is exposed to direct sunlight.
 - 1 This equipment has been tested and found to comply with the limits for medical devices to the IEC 60601-1-2:2007. These limits are designed to provide reasonable protection against harmful interference in a typical medical 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 other devices in the vicinity. However, there is no guarantee that interference will not occur in a particular installation. If this equipment does cause harmful interference to other devices, 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:
 - Reorient or relocate the receiving device.
 - Increase the separation between the equipment.
 - Connect the equipment into an outlet on a circuit different from that to which the other device(s) are connected.
 - Consult the manufacturer or field service technician for help.
 - 9) Do not to touch signal input, signal output or other connectors, and the patient simultaneously.
 - ⚠10) a statement that MEDICAL ELECTRICAL EQUIPMENT needs special precautions regarding
 EMC and needs to be installed and put into service according to the EMC information provided
 in the ACCOMPANYING DOCUMENTS;
 - 11) a statement that portable and mobile RF communications equipment can affect MEDICAL ELECTRICAL EQUIPMENT.
 - 12) Please consult a physician or a trained health professional for interpretation of measurement results.
 - ▲ 13) No phthalates are used for this product and its container.
 - 14) The cuff is not made with natural rubber latex

Caution



- 1. Measurements may be impaired if this device is used near televisions, microwave ovens, X-ray equipment or other devices with strong electrical fields. To prevent such interference, use the meter at a sufficient distance from such devices or turn them off.
- 2. Incorrect operation or failure of user to maintain the unit spares the manufacturer or his agent of the responsibility for system's non-compliance with specifications or responsibility for any damage or injury.

Caution



This manual is made for informational purpose and this manual and product are not meant to be a substitute for the advice provided by your own physician or other medical problem. You should not use the information contained in the product for diagnosis or treatment of health problem or prescription of medication by yourself.

If you have or suspect that you have a medical problem, consult with your physician promptly.

Defective unit or accessories must be packed in the replacement cartons to be shipped off from you to our company.

Shipping and insurance costs for return of defective unit must be prepaid by the users.

5. SAFETY SYMBOLS AND INFORMATION

The International Electrotechnical Commission (IEC) has established a set of symbols for medical electrical equipment which classifies a connection or warning of any potential hazard.

The classifications and symbols are shown below. Save these instructions for your safety.

†	Degree of protection against electric shock: TYPE BF
	Please observe operating instructions
	General warning sign
	General prohibition sign
0	General mandatory action sign
Ţ	Caution
	Waste Electrical and Electronic Equipment (WEEE) The device could be sent back to the manufacturer for recycling or proper disposal after their useful lives. Alternatively the device shall be disposed in accordance with national laws after their useful lives.
Ċ	"OFF" (only for a part of equipment)

•	"ON" (only for a part of equipment)
	This symbol is used inside system. Identifies the point where the safety ground of the system is fastened to
	the chassis.
CAL	Do not open. This is for factory only.
\sim	Alternating current
===	Direct current
$\qquad \qquad \square$	Date of manufacture
	Manufacturer
(((•)))	Non-ionizing radiation
(6 0197	CE mark
SN	Serial No.
EC REP	Authorized representative in the European community.
	Keep dry
RoHS2	RoHS2

6. Guidance for Electromagnetic compatibility (EMC)

Details about the electromagnetic compatibility (EMC) of the FT-500 are given below. Before using the FT-500, be sure to read and understand the following information.

1) Guidance and manufacturer's declaration - electromagnetic emissions

The FT-500 is intended for use in the electromagnetic environment specified below. The customer or the user of the FT-500 should assure that it is used in such an environment.

Emissions test	Compliance	Electromagnetic environment – guidance		
		The FT-500 uses RF energy only for its internal		
RF emissions	Croup 1	function. Therefore, its RF emissions are very low and		
CISPR 11	Group 1	are not likely to cause any interference in nearby		
		electronic equipment.		
RF emissions	Class B			
CISPR 11	Class B			
Harmonic		The FT-500 is suitable for use in all establishments,		
emissions	Class A	including domestic establishments and those directly		
IEC 61000-3-2		connected to the public low-voltage power supply		
Voltage		network that supplies buildings used for domestic		
fluctuations/	Compliance	purposes.		
flicker emissions	Compliance			
IEC 61000-3-3				

2) Guidance and manufacturer's declaration - electromagnetic immunity

The FT-500 is intended for use in the electromagnetic environment specified below. The customer or the user of the FT-500 should assure that it is used in such an environment.

Immunity toot	IEC 60601 test	Compliance	Electromagnetic environment-
Immunity test	level	level	guidance
Electrostatic			Floors should be wood, concrete or
	±6kV: Contact	±6kV: Contact	ceramic tile. If floors are covered with
discharge(ESD)	+8kV: Air	+8kV: Air	synthetic material, the relative
IEC 61000-4-2	TOKV. All	TORV. All	
			humidity should be at least 30 %.
Electrical fast	±2kV: Power	±2kV: Power	Mains power quality should be that of
transition/burst	supply lines	supply lines	a typical commercial or hospital
IEC 61000-4-4	±1kV:	±1kV:	environment.

	Input/output lines	Input/output	
		lines	
Surge IEC 61000-4-5	±1 kV differential mode ±2 kV common mode	±1 kV differential mode ±2 kV common mode	Mains power quality should be that of a typical commercial or hospital environment.
	<5 % <i>U</i> T (>95 % dip in <i>U</i> T) for 0,5 cycle	<5 % <i>U</i> T (>95 % dip in <i>U</i> T) for 0,5 cycle	Mains power quality should be that of
Voltage drops,	40 % <i>U</i> T	40 % <i>U</i> T	a typical commercial or hospital
dips, and	(60 % dip in <i>U</i> T)	(60 % dip in <i>U</i> T)	environment. If the user of the FT-500
fluctuations of	for 5 cycles	for 5 cycles	requires continued operation during
input power	70 % <i>U</i> T	70 % <i>U</i> T	power mains interruptions, it is
supply line IEC	(30 % dip in <i>U</i> T)	(30 % dip in <i>U</i> T)	recommended that the FT-500 be
61000-4-11	for 25 cycles	for 25 cycles	powered from an uninterruptible
	<5 % <i>U</i> T	<5 % <i>U</i> T	power supply or a battery.
	(>95 % dip in <i>U</i> T)	(>95 % dip in	
	for 5 sec	<i>U</i> T)	
		for 5 sec	
Magnetic field of			
commercial			Power frequency magnetic fields
frequency	3 A/m	3 A/m	should be at levels characteristic of a
(50/60Hz)			typical location in a typical
IEC 61000-4-8			commercial or hospital environment.

Note



*U*T is the a.c. mains voltage prior to application of the test level.

3) Guidance and manufacturer's declaration – electromagnetic immunity 2

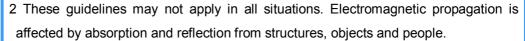
The FT-500 is intended for use in the electromagnetic environment specified below. The customer or the user of the FT-500 should assure that it is used in such an environment.

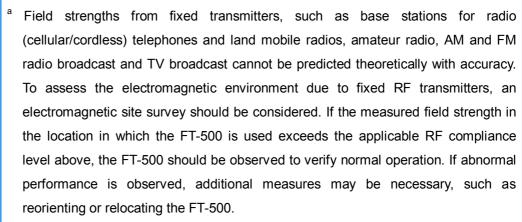
Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment-guidance
			Portable and mobile RF communications equipment should be used no closer to any part of the FT-500, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.
			Recommended separation distance
			$d = 1.2\sqrt{P}$
Conducted RF	3 Vrms	3 Vrms	$d = 1.2\sqrt{P}$ 80 MHz to 900 MHz
IEC 61000-4-6	-4-6 150 kHz to 80 MHz		$d = 2.3\sqrt{P}$ 900 MHz to 2,5 GHz
Radiated RF IEC 61000-4-3	3 V/m 80 MHz to 2,5 GHz	3 V/m	where <i>P</i> is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and <i>d</i> is the recommended separation distance in meters (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:



Caution

1. At 80 MHz and 900 MHz, the higher frequency range applies.





^b Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.

4) Recommended separation distances between portable and mobile RF communications equipment and the FT-500

The FT-500 is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the FT-500 can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the FT-500 as recommended below, according to the maximum output power of the communications equipment.

Rated maximum	Separation distance according to frequency of transmitter			
output power	m			
of transmitter	150 kHz to 80 MHz	80 MHz to 900 MHz	900 MHz to 2,5 GHz	
W	$d = 1.2\sqrt{P}$	$d = 1.2\sqrt{P}$	$d = 1.2\sqrt{P}$	
0.01	0.12	0.12	0.23	
0.1	0.38	0.38	0.73	
1	1.2	1.2	2.3	
10	3.8	3.8	7.3	
100	12	12	23	

For transmitters rated at a maximum output power not listed above, the recommended separation distance d in meters (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.

Caution

1. At 80 MHz and 900 MHz, the separation distance for the higher frequency range applies.



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

TERMS OF EACH PART AND FUNCTIONS

1. FRONT PART

1 START BUTTON

Press START button after being ready to measure, the cuff will be wrapped automatically and begins to pressurize.

② STOP BUTTON

Press STOP button if you want to stop it during measurement. Pressurizing will stop and the air will exhaust from the cuff.

③ SYSTOLIC DISPLAY

It indicates systolic blood pressure values.

4 DIASTOLIC DISPLAY

It indicates diastolic blood pressure values.

5 PULSE DISPLAY

It indicates pulse rate.

6 TIMER

It indicates time.

(7) EMERGENCY STOP BUTTON

When your arm is oppressed due to high pressurizing or irregular operation is done, press this button then the cuff will be exhausted rapidly.

8 PRINTER COVER

It protects the printer.

9 AUTOMATIC CUTTER (printing paper let-out slot)

Printing paper is automatically cut off when it comes out through the slot.

10 CUFF

It wraps and releases the arm automatically for measurement.

(1) ARM REST

When the arm is placed on the cuff, the arm supporter sustains the arm and makes the right position.

(2) HUMAN SENSOR

When a user approaches, power is automatically turned on and vice versa.

(3) CARD-READER

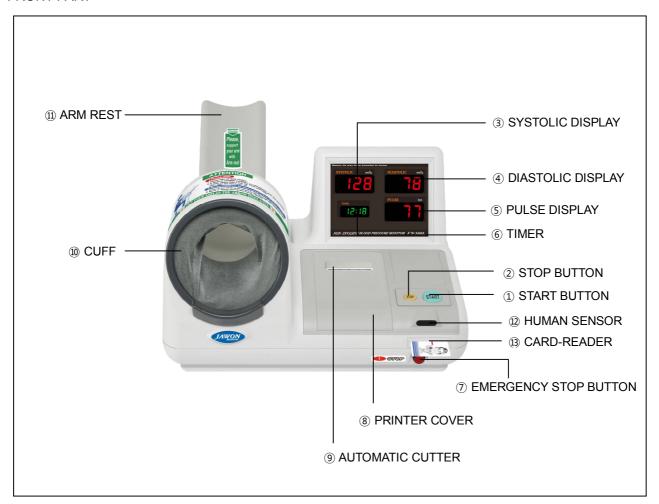
When card is inserted, it reads information in it and stores the measured results then the cuff will be exhausted rapidly.

Note



The cuff and the buttons (START and STOP button) of this device are located at reverse side by R and L type.

FRONT PART



Note

Printer, card reader and human sensor are optional.

ID card can be issued either by the machine manager or by the manufacturer of the model.



The card stores 2 previous measured results and can contain 3 measured results with the current one altogether.

When the model manager writes the card, please refer to the manual and specifications for operation and programs attached to the card writing device at purchase.

2. PRINTER

1 PRINT button

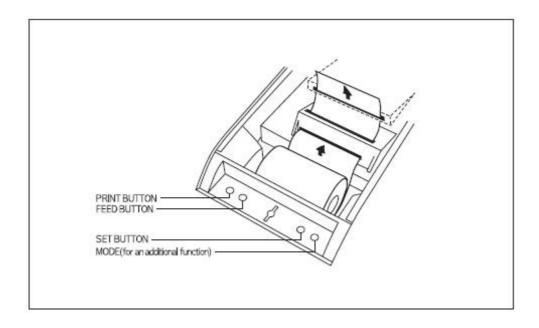
- Use it when you print out the data.
- If you set [ON] at the rear (PRINT ON/OFF switch), the data is printed automatically even when you do not press PRINT button.
- Normally, when you press this button, all the data output after you turn the power on will be printed. (If you turn it off, all memorized data would be deleted.)
- When you set the date and time, the number goes up with this button pressed.

2 FEED button

- Use this button for setting the paper
- When you set the date and time, the number goes down with this button pressed.

③ SET button

- Set the date and time
- The functions are as follows when pressing this before or after measurement. (It does not work during measurement)
- Sequence is HOUR \rightarrow MIN. \rightarrow MON. \rightarrow DAY \rightarrow YEAR
- Within 5seconds, Setting of the date and time finished.
- See the page '23' for detailed method



3. REAR PART

1 POWER

It is used to turn the power on and off.

2 AC INPUT

It is used to connect with the power code.

③ FUSE

Two (250V/1.6AL) fuses are installed to prevent overload.

(4) CAL

This is only for inspection. Never open it.

5 EARTH (POTENTIAL EQUALIZATION TERMINAL)

Please make sure for safety.

6 COMMUNICATION PORTS (RS-232C)

It is for connecting between the main body and a computer or other equipment with cable (RS-232C) to transfer the data collected or measured.

And it is for connecting between the main body and the coin slot with RS-232C cable to transfer the data, too.

(7) SENSOR ON/OFF

Human sensor is switched on and off.

8 SOUND ON/OFF

Music and voice output functions are activated with the switch [ON], and vice versa.

9 CARD ON/OFF

ID card is usable when the switch [ON], and vice versa.

① PRINT ON/OFF

Measured results are printed out when switched [ON], and vice versa.

11 VOLUME (ANNOUNCE ON/OFF)

It controls volume output when switch is on while all volume is [OFF].

(2) INFORMATION BOARD FIXER

Fix the information board here.

Note



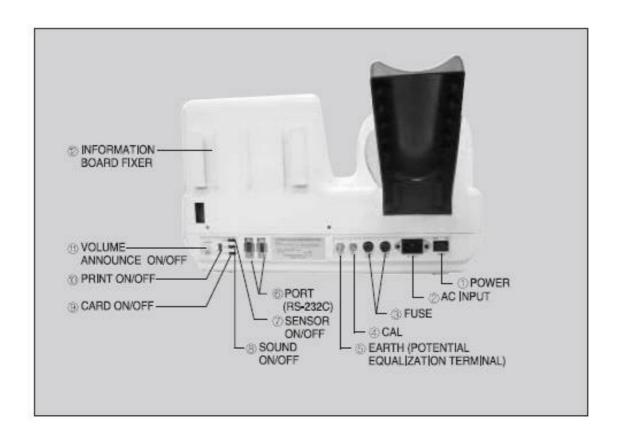
- While sound switch is activated on (so sound is functioning), place the VOLUME ON/OFF switch to [OFF] for deactivation of voice message and music play.
- 2. Printer, card reader, human sensor and reverse monitor is optional.

Note

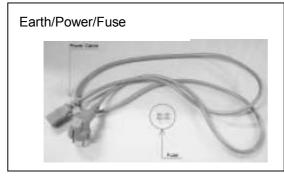


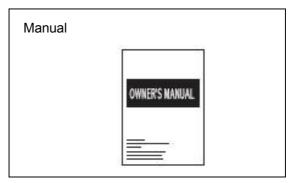
The operator shall not contact the parts (SIP/SOP) and the patient simultaneously and "SIP/SOP shall be available to operator only"

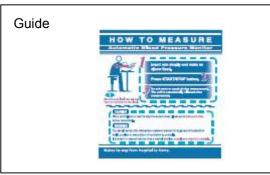
REAR PART

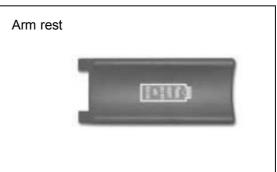


4. ACCESSORIES



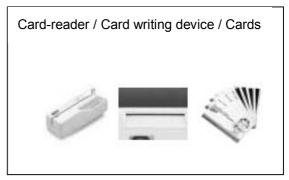




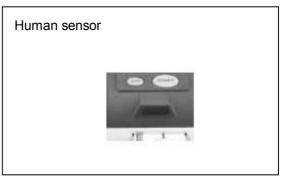


5. OPTIONS





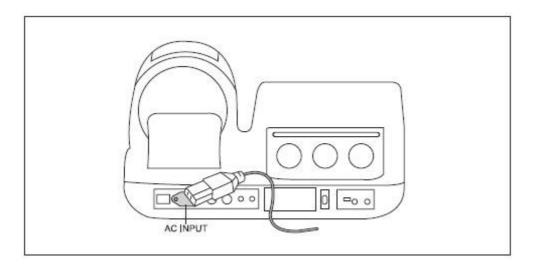




INSTALLATION

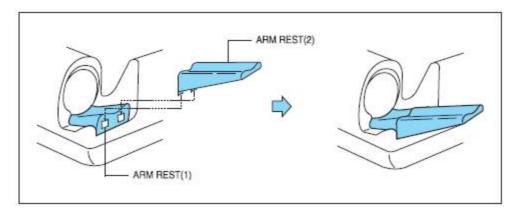
1. CONNECTING POWER CABLE

Just connect the power cable to the AC INPUT slot on the rear and turn the POWER ON/OFF switch (O/I) on the lower part of the rear (See the picture).



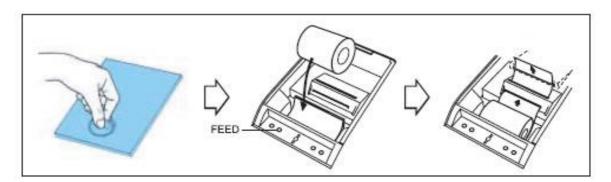
2. ASSEMBLING OF ARM SUPPORTER

The arm supporter consists of two parts. The first part (1) is attached to the main body when it is manufactured in the factory while the second part (2) is assembled to the first part as shown in the picture.



3. LOADING THE PRINT PAPER

- ① Check and see if power is turned on.
- Turn the nut (with a driver on the groove in the middle) on the lower printer cover clockwise to 90 degrees and open the cover.
- 3 Load the print paper as shown in the picture.
- Insert the paper edge deep under the black roll, then it comes out above the CUTTER.
- S Balance the paper in the right place.
- © Cut the paper by pressing the FEED button.
- O Close the cover and turn the nut counterclockwise back.



Note

Being thermal type, printing is photocopied on one side of the paper (slippery side), without using printing ink.

Please check remainder of the paper always and then replace it.

Please use exclusive paper (58mm).

Keep paper rolls in a dark and ventilated place.



Avoid any dust on the paper.

Do not pull the paper during printing. It could cause jam.

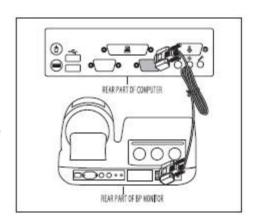
When printing paper is not loaded in correct place, it may cause the malfunction of the printer or paper will be shoved out.

After the exchange of paper to the printer cover does not close properly, the alarm sounds, LED on the 'Err' is displayed. Please check the status of the printer cover.

4. CONNECTING PORT

To transmit the data, connect a computer or other external options to the unit.

Connect the USB cable both to port of the unit and to the computer jack or other external options. (See the picture)



5. SETTING TIME AND DATE

- Turn the power of the unit on.
- Open the printer cover.
- Sequence is HOUR ightarrow MINUTE ightarrow MONTH ightarrow DAY ightarrow YEAR

HOUR

- ① Press SET button, then indicator says its counts that have been measured since keeping button ON.
- ② Press SET button one more. First 2 figures will blink.
- To set the current hour, press PRINT button to make the number goes up or press FEED button to make the number goes down.



MINUTE

- ① After setting the hour, press the SET button again.
- ② In this time, last 2 figures will blink.
- ③ As the same way as above, set the current minute with PRINT and FEED button.



MONTH

- ① After setting the minute, press SET button again.
- ② First 2 figures will blink.
- 3 Set the current month with PRINT and FEED button.



DAY

- ① After setting the month, press SET button again.
- 2 Last 2 figures will blink.
- 3 Set the current day with PRINT and FEED button.buttons.



YEAR

- ① After setting the day, press SET button again.
- ② First 2 figures will blink.
- 3 Set the current year with PRINT and FEED button.



Note



If you want to measure blood pressure during setting the date and time, press STOP button. Then you can measure again immediately.

Note

You should set all data at once (hour, minute, month, day and year).

In case of stopping setting, the values return to previous ones which you have done before.

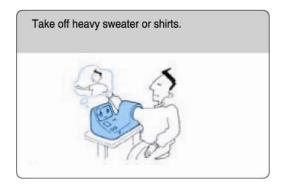


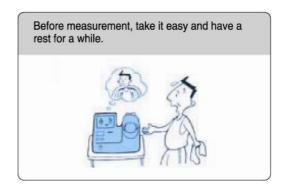
The calendar and time functions work without plugging power cord in.

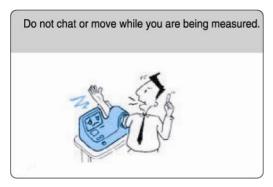
Calendar program is inputted for 100 years, and it would be adjusted automatically even at a leap year.

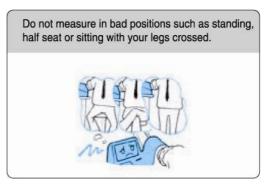
MEASUREMENT

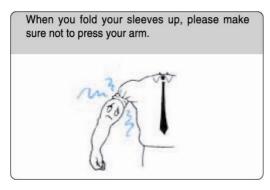
1. CAUTIONS FOR MEASUREMENT

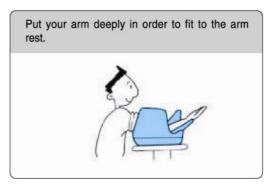


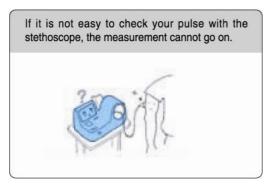


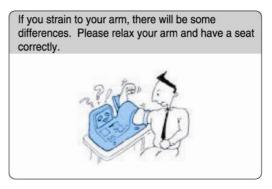






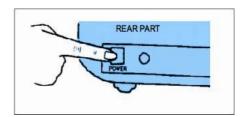






2. MEASUREMENT

① Check the voltage and turn the power on.



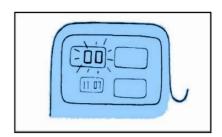
Note



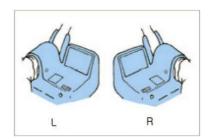
With the human sensor attached optionally, the LED screen is automatically turned on as it detects a user coming to the range.

The sensor should be placed [ON].

② For non-ID card users, a background music flows as power is turned on, and there appears the stand-by scene on the LED screen.



3 Please put either the right arm or left arm into the cuff.



Note



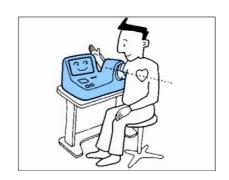
An optimal arm circumference for this equipment is 9" to 14".

Caution



Place your arm on the arm supporter with the palm facing up through the cuff deeply. Adjust the height of the chair so that the arm is leveled off with the heart.

When the arm is placed lower than the heart, blood pressure will become higher than actual value, and vice versa.



4 Press the START button.

Checking blood pressure begins as the cuff is automatically pressurized with the LED screen displaying animations of the checking in process.



Caution



When the measurement is started, the voice message is announced as "Starting measurement, don't move or speak please." When you feel painful and want to stop the measurement, press EMERGENCY BUTTON.



⑤ The LED screen displays the results of checking upon finishing along with a voice message of "Measurement completed, pull your arm out please. Thank you."

The cuff will automatically release your arm and resume the original position.



Note



When the measurement is not satisfactory, the voice message comes out as "Cannot measure, we will try again." At this time, let your arm stay into the cuff and start over again from the beginning.

- ⑥ The results are printed out, and a voice message also announces the results as "Your blood pressure is systolic 000, diastolic 000 and pulse 000."
- Pull your arm out from the cuff.
- When you want to measure one more time, press STOP button and start all procedures from the beginning.

Note



When the PRINT ON/OFF switch on the rear is set as $\lceil \mathsf{OFF} \rfloor$, the result will not be printed even if the measurement is completed.

Note



For the card users, six previous results stored in the card can be recalled to compare with the current ones newly checked.

Seven results altogether could be printed out.

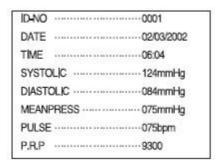
Caution



This device is only for adult.

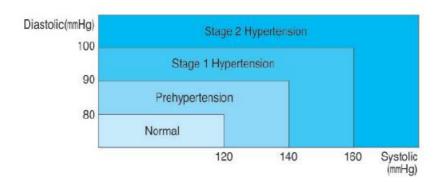
3. DATA ON MEASUREMENT BY PRINTER

▼ Results on Printing Paper



▼ Classification of the blood pressure

: National High Blood Pressure Education Program, National Heart, Lung and Blood institute, NIH (JNC7, 2003)



Note

- When pressure is high with the jammed air hose, message appears on the printer as ERROR PRESSURE.

When the message is repeated, call for maintenance service.



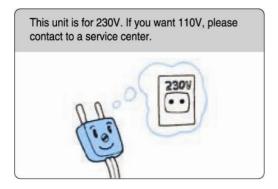
- When pressure is low as air leaks, message appears on the printer as ERROR CUFF.

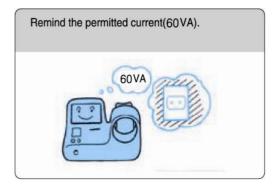
When the message is repeated, call for maintenance service.

- When the subject moves or speaks while in testing, message appears on the printer as ERROR MEASURE.

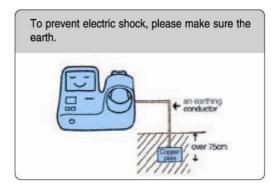
Try to retest after a while. If the message is repeated, call for maintenance service.

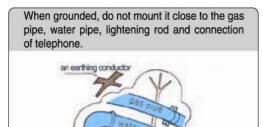
MAINTENANCE

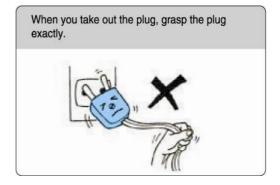




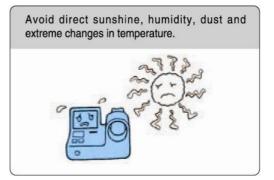


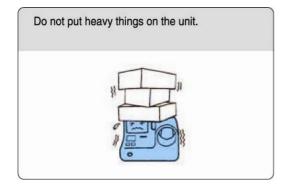


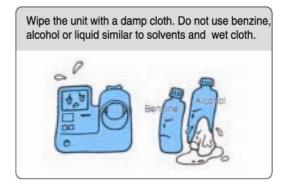


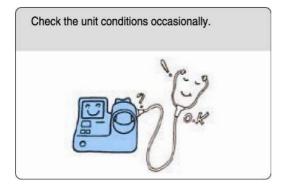


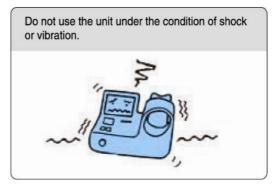


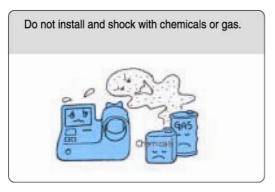




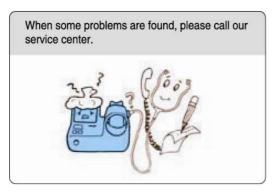












ERROR & REPAIR

Error	Cause	Repair
ERROR PRESSURE	pressure is high with the	When the message is
	jammed air hose	repeated, call for
		maintenance service.
ERROR CUFF	pressure is low as air leaks	When the message is
		repeated, call for
		maintenance service.
ERROR MEASURE	subject moves or speaks	- Don't move or speak.
	while in testing	- When the message is
		repeated, call for
		maintenance service.

AFTER SERVICE

1. AFTER SERVICE

- If there is any problem with the unit, please follow the steps below;
 - ** Contact our company's Overseas Service Department immediately.

 After gathering the model name, Serial Number, date of purchase and description of the problem, contact our company with information shown below.
 - * Try to solve the problem over the phone with the personnel of local service department.

 If the problem cannot be solved over the phone, just return to service department directly.
 - ** Our company or local distributor will make available on-request circuit diagrams, component part list, descriptions, calibration or other information which will assist your appropriately qualified technical personnel to repair those parts of unit which are designated by our company as repairable.

How to contact our company

Write us at:

JAWON MEDICAL CO., LTD

29, Gongdan 4-ro, Jillyang-eup, Gyeongsan-si, Gyeongsangbuk-do, Korea

TEL: 82-53-856-0993 FAX: 82-53-856-0995

(You can also contact the following representative or your local distributor)

2. PACKING AND TRANSPORT

Our company follows his packing ways to protect any impact during transporting etc. So please do not transport or move the unit without our company's packing condition as your wishes.

The normal storage environment; -20°C~ 60°C of temperature, Humidity is less than 95% non-condensing.

SPECIFICATION

Model	FT-500(R) FT-500(L)			
Measuring Method	Oscillometric			
Display mode	LED display			
Result Contents	Systolic/Diastolic/Mean blood pressure, Pulse, P.R.P.			
Measuring ranges	Pressure 30~300mmHg, Pulse 30	0~200beats/minute		
Accuracy	Pressure ±2mmHg, Pulse ±5%			
Resolving Power	1mmHg			
Pressurizing method	DC Motor			
Cuff type	Belt type			
Pressurizing time	Approx. 10 seconds			
Measuring time	Approx. 33 seconds			
Printer	Thermal printer (optional)			
Power supply	AC 230V, 50/60Hz			
Power consumption	40VA			
Operation ambient	Temperature 10~40 ℃, Humidity 30~75%			
Storage ambient	Temperature -20~60 ℃, Humidity Less than 95%			
Data transmission	RS-232C			
Dimension	480(W) × 310(D) × 290(H) mm			
Weight	Approx. 11kg			
Measuring parts	FT-500(R): Right arm FT-500(L): Left arm			

WARRANTY

Item	Automatic Blood Pressure Monitor	Warranty period
Model	FT-500(R/L)	1year (main unit only)
Serial NO.		1year (main unit only)

Date of purchase		Month	Day	Year
Customer	Name:			TEL:
Customer	Address:			
Dealer	Name:			TEL:
Dealei	Address:			

Date	Defection	Confirmation			

Note



- When you receive this warranty, make sure that the name of the dealer and the month, day and year of purchase are all completed.
- This warranty will not be reissued, please keep it in a safe place.

Periodic Check List

Manag	ement No.	

Item		Inspection Subject		Requirements			Judgment	Remarks
Visual Check								
Mainframe	1	Enclosure		No scratch, crack,			Pass/Fail	
				defo	rmation and ru	st		
	2	Labels and pa	anels	No peel	ing and dust		Pass/Fail	
	3	Keys		No damage			Pass/Fail	
	4	Cuffs		No scratch and damage			Pass/Fail	
Accessories	1	Power cord		No scratch and damage		Pass/Fail		
	2	User manual		Kept in proper place			Pass/Fail	
Mechanical Check								
Mainframe	1	Keys		Smooth operation			Pass/Fail	
	2	Recorder		Smooth	operation with	no	Pass/Fail	
				abno	ormal sound			
	3	Cuffs		Smooth operation			Pass/Fail	
Accessories 1 Power cor		Power cord	er cord Smooth operation and			Pass/Fail		
				removal				
Electrical Ch	eck							
Performance	1	Power supply		Screen display upon			Pass/Fail	
				power-on				
	2	Display		No abnormality and			Pass/Fail	
				flickering				
	3	Printing	Printing		printing possible		Pass/Fail	
	4	Measurement		Proper measurement			Pass/Fail	
General Judgment					Pass/Fail			
Model FT-500				Serial No.				
Installation place			Date			of purchase		
Check date			Checked by			Approved by		

Copy this sheet for use

If repair is required, write down so in the Remarks column.

Daily Check List

Management No.

Item		Inspection Su	bject	Requirements			Judgment	Remarks
Visual Check			'			J		
Mainframe	1	Enclosure		No scratch, crack, deformation and rust			Pass/Fail	
	2	Labels and pa	anels	No peeling and dust		Pass/Fail		
	3	Keys		No damage			Pass/Fail	
	4	Cuffs		No scratch and damage			Pass/Fail	
A	1	Power cord		No scratch and damage			Pass/Fail	
Accessories	2	User manual		Kept in proper place			Pass/Fail	
Mechanical Check								
	1	Keys		Smooth operation			Pass/Fail	
Mainframe	2	Recorder		Smooth operation with no abnormal sound			Pass/Fail	
Accessories	1	Power cord		Smooth operation and removal			Pass/Fail	
Electrical Check								
Performance	1	Power supply		Screen display upon power-on			Pass/Fail	
	2	Display		No abnormality and flickering		Pass/Fail		
	3	Printing	Printing		Waveform printing possible			
	4	Measurement		Proper measurement			Pass/Fail	
Other	1	Clock		Present date/time		Pass/Fail		
General Judgment					Pass/Fail			
Model FT-500		FT-500					Serial No.	
Installation pla	Installation place					Date	e of purchase	
Check date			Check	Checked by App		roved by		

Copy this sheet for use
If repair is required, write down so in the Remarks column.

Service center TEL: 82 - 2 - 587 - 4056 82 - 53 - 856 - 0993

If the problems continue, call the service center. When you ask for service, the manufacturer's label, serial number, date of original purchase and explanation of malfunction will be required.

For purpose of improvement, specifications and subject to change without notice.



HEAD OFFICE29, Gongdan 4-ro, Jillyang-eup, Gyeongsan-si, Gyeongsangbuk-do, 38470, Republic of Korea

TEL: 82-53-856-0993, FAX: 82-53-856-0995

EUROPEAN REPRESENTATIVE: VITAKO Sp. z o.o.

UL. MALEJ SYRENKI 2 71-790 SZCZECIN, POLAND TEL: 91 8522900 / 901 FAX: 91 8522910