

Little Doctor

Digital Blood Pressure Monitor

Instruction Manual

Ciśnieniomierz elektroniczny półautomatyczny LD do pomiaru ciśnienia tętniczego krwi i pulsu Instrukcja Obskacj

Instrukcja Obsługi

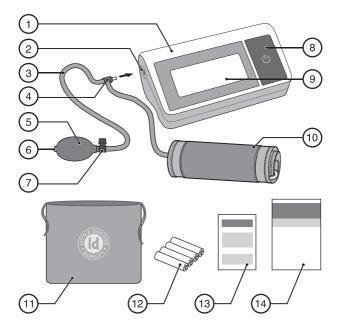


TABLE OF CONTENTS

PARTS AND COMPONENTS	3
GENERAL INFORMATION	4
INDICATIONS FOR USE	4
OPERATION PRINCIPLE	4
APPLIED NEW TECHNOLOGIES LD	4
RECOMMENDATIONS ON CORRECT MEASUREMENTS	5
POWER SUPPLY	6
CORRECT POSITION DURING MEASUREMENT	6
CUFF PREPARATION	7
MEASUREMENT PROCEDURE	8
INFORMATION ABOUT ERRORS	10
CARE, STORAGE, REPAIR AND DISPOSAL	
TROUBLESHOOTING TIPS	
WARRANTY	12
TECHNICAL SPECIFICATIONS	12
CERTIFICATION AND STATE REGISTRATION	13

PARTS AND COMPONENTS

ENG



- 1. Electronic Unit.
- 2. Arm cuff and pump jack.
- 3. Air tube.
- 4. T-piece for connecting the bulb and cuff to the electronic Device.
- 5. Bulb.
- 6. Check valve.
- 7. Pressure release valve.
- 8. () (Power ON/OFF).
- 9. LCD.
- 10. Arm cuff CUFF-LDA.
- 11. Storage Case.
- 12. Power elements.
- 13. Warranty card.
- 14. Instruction Manual.

GENERAL INFORMATION

This Instruction Manual is designed to assist the user with safe and effective operation of the semiautomatic digital Device for measurement of blood pressure and heartbeat rate LD, (modification LD22 (hereinafter - the "Device"). Use this Device according to the rules described in this Manual. Operate the Device only as intended. Do not use the Device for any other purposes. Read and under stand the whole Instruction Manual in particular "Becommendations on Correct stand the whole Instruction Manual, in particular "Recommendations on Correct Measurement".

INDICATIONS FOR USE

Use this Device to measure your systolic and diastolic blood pressure and heartbeat rate in patients aged from 15. This Device is recommended for use by persons with unstable blood pressure or known arterial hypertension at home as an addition to medical surveillance. The cuff is designed for the upper arm with the circumference approximately from 25 to 36 cm.

OPERATION PRINCIPLE

This Device uses the oscillometric method of blood pressure with Fuzzy Algorithm and pulse rate measurement. Wrap the cuff around your upper arm and it starts to be inflated by bulb. The sensitive element of the Device feels the weak pressure oscillations in the cuff generated by widening and contraction of the brachial artery in response to every heartbeat. Pumping in is ceased when cuff is adequately pumped in to determine diastolic and systolic pressure (the amplitude of the pressure waves is measured, converted into millimeters Hg and shown on the display as figures) after which air is released from cuff. Remember that the Device will not maintain the mentioned accuracy of a measurement if it is used or stored at a temperature or humidity other than those specified in Technical Specifications of this Manual. We are warning about possibility of mistakes in blood pressure measurement with this Device in persons with pronounced cardiac arrhythmia. Consult the doctor concerning blood pressure measurement of your child.

APPLIED NEW LD TECHNOLOGIES



Fuzzy Algorithm - algorithm for processing the measurement values with regard to peculiarities of the man's heartbeat, thus, ensuring high measurement accuracy.



Identyfikacja arytmii – specjalny znak «)» na wyświetlaczu urzadzenia, informujący o wykryciu nieregularnego pulsu.

WARNING! This Device may be used only with cuffs:

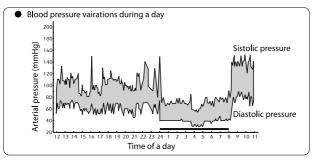
- Cuff-LDA, size 25-36 cm (delivered in a set with the Device).

RECOMMENDATIONS ON CORRECT MEASUREMENTS

1. For correct measurement you should know that THE BLOOD PRESSURE IS SUBJECT TO SHARP VARIATIONS EVEN WITHIN THE SHORT TIME INTERVALS. The blood pressure depends on many factors. It is usually lower in summer and higher in winter. The blood pressure varies together with the atmospheric pressure, depends on physical loads, emotional excitement, stresses and dietary regime. Drugs, drinking alcohol and smoking produce significant effect. Even the very procedure of blood pressure measurement in a polyclinic sends the blood pressure high in many people, thus, the blood pressure measured at home often differs from the values received in a polyclinic. As the blood pressure tends to rise at low temperatures, make measurements at an indoor temperature (approximately 20° C). If this Device stayed under a low temperature, keep it for at least 1 hour at an indoor temperature before use, otherwise the measurement result may be incorrect. During a day the difference in readings for healthy people may be 30-50 mmHg of systolic pressure and to 10 mmHg of diastolic pressure. The dependence of the blood pressure on various factors is individual for each person. Accordingly, it is recommended to keep a special book with blood pressure records, ONLY A CERTIFIED DOCTOR USING YOUR RECORDS IS CAPABLE TO ANA-LYZE THE TENDENCY OF YOUR BLOOD PRESSURE VARIATIONS.

2. At cardiovascular and some other diseases requiring blood pressure monitoring make measurements in the hours fixed by your attending doctor. REMEMBER THAT THE DIAGNOSTIC AND ANY TREATMENT OF HYPERTENSION MAY BE CONDUCTED ONLY BY A CERTIFIED DOCTOR ON THE BASIS OF BLOOD PRESSURE VALUES OBTAINED BY THIS DOCTOR. TAKING OF DRUGS AND THEIR DOSES SHOULD BE PRESCRIBED ONLY BY YOUR ATTENDING DOCTOR.

3. At such disorders as deep vascular sclerosis, weak pulse wave and also in patients with the prominent distortions of cardiac rhythm it may be difficult to measure the blood pressure accurately. IN SUCH CASES CONSULT A CERTIFIED DOCTOR ABOUT APPLICATION OF THE ELECTRONIC DEVICE.

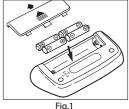


4. KEEP QUIET DURING A MEASUREMENT TO OBTAIN THE ACCURATE VALUES OF YOUR BLOOD PRESSURE WITH THE FLECTRONIC DEVICE, MEASURE YOUR BLOOD PRESSURE IN THE CALM AND COMFORTABLE CONDITIONS AT THE INDOOR TEMPERATURE. NO EATING AN HOUR BEFORE MEASUREMENT; NO SMOKING, TAKING TONIC AGENTS, ALCOHOL 1.5-2 HOURS BEFORE MEASUREMENT.

5. The accuracy of blood pressure measurement depends on whether the cuff matches the size of your arm. THE CUFF SHOULD NOT BE TOO SMALL OR TOO LARGE.
6. Wait 3 minutes between measurements for the blood to restore its circulation. However, the persons with prominent atherosclerosis due to considerable loss of vascular elasticity may need to increase the wait time between measurements (10-15 minutes). This also refers to the patients suffering for long from diabetes. For more accurate determination of blood pressure it is recommended to make a series of 3 consecutive measurements and to use the average value.

POWER SUPPLY OF THE DEVICE

1. Open the cover of the battery compartment and install 4 "AA" size batteries according to polarity marked inside the compartment. Do not use much force to remove the cover of the battery compartment.



2. Close the battery cover.

Replace all batteries when the Low Battery Indicator ", appears on the screen or when there is no any indication on the screen. The Low Battery Indicator does not show the discharge level. Alkali batteries are recommended. When the device operating rules are observed, the batteries serve for approximately 1000 measurements. The batteries supplied with the Device are intended for check of the Device performance at sale and their service life may be shorter than of the recommended batteries. Replace all four batteries at the same time. Do not use the waste batteries. If the Device is unused for a long time, remove all batteries. Do not leave the waste batteries in the Device.

CORRECT POSITION DURING MEASUREMENT

1. Sit at a table so that during blood pressure measurement your hand rests on its surface. Be sure that the cuff is placed approximately at the level of your heart and that your arm lies freely on the table and does not move.



You can measure the blood pressure lying on the back. Look at the ceiling, keep quiet and do not move during measurement. Be sure that the cuff is placed approximately at the level of your heart.





ENG

CUFF PREPARATION

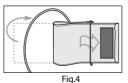
1. Insert the cuff end for about 5 cm into a metal ring as shown in the figure.

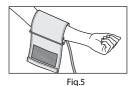
2. Apply the cuff to your left upper arm so that the air tube is directed to your palm. If the measurement on your left arm is difficult, you may use your right arm. In this case remember that the readings may differ by 5-10 mmHg and even more.

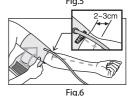
3. Wrap the cuff around your upper arm so that the bottom of the cuff is approximately 2-3 cm above your elbow. The sign "ARTERY" should be over the arm artery.

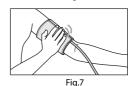
4. Fix the cuff so that it fits tightly to the arm, but see that it is not overtight. Too tight or too free placement of the cuff may give inaccurate readings.

5. On the fixed cuff the sign «INDEX» should point to the area «NORMAL (25-36 cm)». It means that the cuff is chosen correctly and fits the size of your upper arm. if the sign points to the area marked «IIII» the cuff is too small and the readings will be higher. If the sign points to the area marked (III)» the cuff is too large and the readings will be lower.













6. If the arm has a conic form, the cuff should be put on with a spiral movement as shown in the figure.

ENG C

7. If the rolled-up sleeve squeezes the arm interfering with free blood flow the Device may give inaccurate figures not corresponding to your actual blood pressure.

MEASUREMENT PROCEDURE

1. Connect the air hoses of the cuff and bulb to the T-piece. Insert the T-piece into the socket on the device casing (fig. 11).

Make 3-5 deep inhales and exhales before taking a measurement and relax. Do not move, do not speak and do not toughen your arm.

2. Press (¹).

3. All symbols will appear on the display screen for a short time, two short sound signals will be given (fig. 12). After the signal, the screen shows "0" and marker "A" starts blinking. It means that the devise is ready for measurement (fig. 13).

4. Pump the cuff by pressing on the bulb up to the pressure 30-40 mmHg higher than your expected systolic (upper) pressure.

The value of the pressure in the cuff is constantly displayed on the device's screen. If the sum of the expected systolic (upper) pressure plus

30-40 mmHg is less or equal to 190 mmHg, you may, for convenience, pump the cuff until the signal (the device produces a sound when the cuff pressure is 190 mmHg).

IF THIS SUM IS GREATER THAN 190 MMHG, CONTINUE TO PUMP THE CUFF EVEN AFTER THE SIGNAL UP TO THE DESIRED PRESSURE, WATCHING THE DISPLAY INDICATIONS.

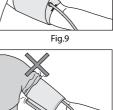






Fig.11



Fig.12



Fig.13

DO NOT PUMP THE CUFF PRESSURE HIGHER THAN 300 MMHG SINCE IT EXCEEDS THE DEVICE MEASUREMENT MAXIMUM. IF THE PRESSURE HIGHER THAN 300 MMHG IS REACHED, AN INTERMITTENT SIGNAL WILL BE HEARD.

5. When the desired cuff pressure is reached, stop pressing on the bulb and put it on the table accurately.

6. The cuff pressure then starts automatically decreasing and the marker " \P " appears on the screen.

7. If symbol " \blacktriangle ", appears during pressure release, it means that the cuff has not been pumped up to the required level. Pump the cuff up by additional 30-40 mmHg for 4 seconds above the previous value, otherwise the display will show the error message "Err".

8. The pressure measurement continues all the time when symbol " \P " is blinking. Try to stay motionless and not to move your arm during the measurement.

OF THE MARKER "Err", APPEARS ON THE SCREEN, STOP THE MEASUREMENT AND FULLY RELEASE THE CUFF PRESSURE BY PRESSING THE PRESSURE RELEASE VALVE. REPEAT THE MEASUREMENT FOLLOWING STRICTLY THE RECOMMENDATIONS OF THIS MANUAL.

9. In the end of the measurement the display will show the values of your pressure and pulse, and the symbol " Ψ " (fig. 14) will start blinking.

If irregular pulse rhythm is detected during measurement, symbol of arrhythmia «**Q**», will appear upon measurement end. Appearance

of arrhythmia indicator may also be caused by body movement during measurement. During periodical appearance of this indication apply to You attending doctor.

10. Release the remaining pressure in the cuff using the pressure release valve.

11. For taking a new measurement repeat all steps described in this paragraph.

TO OBTAIN THE ACCURATE RESULT MAKE INTERVAL BETWEEN MEASUREMENTS TO RESTORE THE BLOOD CIRCULATION. WAIT FOR AT LEAST 3 MINUTES BEFORE MAKING A NEW MEASUREMENT.

If no actions are done in mode of date and time setting for more than 3 minute, the Device will be switched off automatically.

12.Press () to switch off.



ENG



CN1

	Indication	Likely cause	Methods of correction
-	ξm	The arm cuff is applied incorrectly or the air tube T-piece is inserted not tightly enough.	Be sure that the arm cuff is applied correctly and the plug is inserted tightly. Repeat the whole measurement proce- dure.
		Measurements cannot be made due to hand movement or talking during measurements.	Repeat the measurement fol- lowing strictly the recommen- dations of this Manual.
		The cuff has not been pumped up to the required pressure.	Repeat the measurement after pumping the cuff up 30-40 mmHg above the expected systolic pressure.
		At such disorders as deep vascular sclerosis, weak pulse wave and also in patients with the prominent distortions of cardiac rhythm it may be difficult to measure the blood pressure accurately.	In such cases consult a certified doctor about application of the electronic Device.
-		Batteries are discharged.	Replace the batteries for new ones.

CARE, STORAGE, REPAIR AND DISPOSAL

1. Keep this Device from exposure to higher humidity, direct sunlight, shocks, vibration. THIS Device IS NOT WATERTIGHT!

2. Do not keep and use this Device near heating installations and open fire.

3. Remove the batteries if the Device will be unused for a long time. Battery leaking may damage the Device. KEEP BATTERIES OUT OF REACH OF CHILDREN!

4. Keep the Device clean and protect it from dust. Use the dry soft cloth to clean the Device.

5. Keep the Device and its components away from water, solvents, spirit and petrol.

6. Protect the arm cuff from contacting on sharp things; do not stretch or fold tightly the arm cuff.

7. Do not subject the Device to strong shocks, such as dropping on the floor.

8. This Device does not contain special controls to adjust the measurement accuracy. It is prohibited to open individually the electronic block. Repair the Device only in authorized organizations.

9. On expiration of the warranted service life apply from time to time to authorized repair organizations to check the technical condition of the Device.

ENG

10. Dispose of the Device and its components according to the application local regulations. No special requirements to disposal of this Device are defined by the manufacturer.

11. The arm cuff may withstand multiple sanitary treatments. The internal tissue surface of the arm cuff (contacting on arm) may be cleaned with cotton ball soaked in 3%-solution of hydrogen peroxide. At long use the partial color fading of the tissue coating of the arm cuff is possible. Washing and ironing of the arm cuff are not allowed.

TROUBLESHOOTING TIPS		
PROBLEM	LIKELY CAUSE	METHOD OF CORRECTION
After pressing the ① Button no indication on the display.	Discharge of batteries. Polarity of batteries is not observed. Battery terminals are contaminated.	Replace all batteries for new ones. Install batteries correctly. Clean the terminals with dry cloth.
Every time the blood pressure is different. Measurements are too low/ high.	Check that the arm cuff is level with your heart? Check that the arm cuff is applied correctly? Perhaps your arm muscles are tough? Perhaps you talk or move your arm during the measurement?	Take the correct position for measurement. Take the correct position for measurement. Apply the arm cuff correctly. Keep silence and quiet during measurement.
Measurements of the pulse rate are too high/low.	Perhaps you talk or move your arm during the measurement? Perhaps you make measurement directly after physical load?	Keep silence and quiet during measurement. Repeat the measurement at least in 5 min.

Spontaneous failure of power supply.	Actuation of automatic de- energizing system	This is not a defect. The Device is disconnected automatically in 3 minutes after the last operation of the Device.
--	---	---

U If regardless of the above recommendation you are unable to get correct measurement results, stop the use of this Device and apply to a maintenance organization (addresses and telephones of authorized organizations may be found in the warranty card). Do not try to adjust the internal mechanism by yourself.

WARRANTY

- 1. The following LD product is covered by warranty for the period specified in the warranty card.
- 2. The warranty liabilities are contained in the warranty card given at the sale of this Device to a purchaser.
- 3. The addresses of organizations for warranty maintenance are given in the warranty card.

TECHNICAL SPECIFICATIONS

Measurement method	oscillometric with Fuzzy Algorithm
Display	LCD
Pressure indication range in an arm cuff, mmHg	from 0 to 300
Measurement range: pressure in an arm cuff, mmHg pulse rate, 1/min	from 40 to 260 from 40 to 160
Range of admissible absolute er- ror at measurement of air pressure in an arm cuff, mmHg. pulse rate measurement, %	±3 ±5
Power source, V	6
Type of power supply	4 "AA" size batteries (LR6)
Max power intake, W	0,1
Operation conditions: Temperature, °C Relative humidity, % Rh	from 10 to 40 85 and lower

Storage and transportation conditions: Temperature, °C Relative humidity, % Rh	from -20 to 50 85 and lower	
Cuff size:	larger for adults (upper arm circumference 25-36 cm)	(7)
Dimensions: Size (electronic block), mm Weight (without package, case, batteries and adapter), g	129 x 68 x 54 245	ENG
Completeness	Electronic block, Cuff-LDA (with pipes and T-piece), bulb model LD-S035 (with pressure release valve and check valve), 4 batteries, a case, instruction manual, warranty card, package	
Year of manufacture	Year the manufacture is given in the bottom of the Device body in a serial number after symbols "AA"	
Symbols	Type BF Important: Read the instructions Type II	

CERTIFICATION AND STATE REGISTRATION

This Device manufacturing is certified according to international standard ISO 13485:2003.

Devices LD22 comply with the requirements of European Directive MDD 93/42/ EEC, international standards, EN980, EN1041, EN1060-1, EN1060-3, EN10601-1-2, ISO 14971.

Complaints and requests should be addressed to:

Little Doctor Europe Sp. z o.o.

57G Zawila Street, 30-390, Krakow, Poland

Service phone: +48 12 2684748, 2684749

Manufactured under control and for Little Doctor International (S) Pte. Ltd., 35 Selegie Road #09-02 Parklane Shopping Mall, Singapore 188307, Singapore. Postal address: Yishun Central P.O. Box 9293 Singapore 917699.

Manufacturer:

Little Doctor Electronic (Nantong) Co. Ltd., No.8, Tongxing Road Economic & Technical Development Area, Nantong 226010, Jiangsu, PEOPLE'S REPUBLIC OF CHINA

Distributor in Europe:

Little Doctor Europe Sp. z o.o. 57G Zawila Street, 30-390, Kraków, Poland Sales Office phone: +48 12 2684746, 12 2684747, fax: +48 12 268 47 53 E-mail: biuro@littledoctor.pl www.LittleDoctor.pl

Authorized Representative in the EU:

Little Doctor Europe Sp. z o.o. 57G Zawila Street Krakow 30-390 Poland

www.LittleDoctor.sg



EC



LITTLE DOCTOR INTERNATIONAL (S) PTE. LTD. Yishun Central P.O. Box 9293 Singapore 917699, Fax: 65-62342197, E-mail: Id@singaporemail.com

> REP Little Doctor Europe Sp. z o.o. 57G Zawila Street Krakow 30-390 Poland

Registered trade marks of Little Doctor International (S) Pte. Ltd.

© Little Doctor International (S) Pte. Ltd., 2016