Instruction Manual 1

OMRON

Automatic Upper Arm Blood Pressure Monitor

M7 Intelli IT (HEM-7361T-EBK)
X7 Smart (HEM-7361T-ESL)

Read Instruction manual 1 and 2 before use.

FR Lire le mode d'emploi 1 et 2 avant l'utilisation.
DE Lesen Sie vor der Verwendung Gebrauchsanweisung 1 und 2.
IT Leggere il manuale di istruzioni 1 e 2 prima dell'uso.
ES Lea el manual de instrucciones 1 y 2 antes del uso.
NL Lees gebruiksaanwijzing 1 en 2 voor gebruik.
RU Прочтите руководства по эксплуатации 1 и 2 перед использованием.
TR Kullanmadan önce, kullanım kılavuzu 1 ve 2'yi okuyun.
AR اقرأ دليل الإرشادات 1 و 2 قبل الاستخدام.

CE 0197

All for Healthcare

EN

1. Introduction

Thank you for purchasing the OMRON Automatic Upper Arm Blood Pressure Monitor. This blood pressure monitor uses the oscillometric method of blood pressure measurement. This means this monitor detects your blood movement through your brachial artery and converts the movements into a digital reading.

1.1 Safety Instructions

This instruction manual provides you with important information about the OMRON Automatic Upper Arm Blood Pressure Monitor. To ensure the safe and proper use of this monitor, READ and UNDERSTAND all of the safety and operating instructions. **If you do not understand these instructions or have any questions, contact your OMRON retail outlet or distributor before attempting to use this monitor. For specific information about your own blood pressure, consult with your physician.**

1.2 Intended Use

The device is a digital monitor intended for use in measuring blood pressure and pulse rate in adult patient population. The device detects the appearance of irregular heartbeats during measurement and gives a warning signal with readings. It is mainly designed for general household use.
The device can detect an irregular pulse suggestive of Atrial Fibrillation (Afib). Please note that the device is not intended to diagnose Afib. A diagnosis of Afib can only be confirmed by Electrocardiogram (ECG). If the Afib symbol appears, consult your physician.

1.3 Receiving and Inspection

Remove this monitor and other components from the packaging and inspect for damage. If this monitor or any other components is damaged, DO NOT USE and consult with your OMRON retail outlet or distributor.

2. Important Safety Information

Read the Important Safety Information in this instruction manual before using this monitor. Follow this instruction manual thoroughly for your safety.
Keep for future reference. For specific information about your own blood pressure, CONSULT WITH YOUR PHYSICIAN.

⚠ 2.1 Warning **Indicates a potentially hazardous situation which, if not avoided, could result in death or serious injury.**

- DO NOT use this monitor on infants, toddlers, children or persons who cannot express themselves.
- DO NOT adjust medication based on readings from this blood pressure monitor. Take medication as prescribed by your physician. ONLY a physician is qualified to diagnose and treat high blood pressure and Afib.
- DO NOT use this monitor on an injured arm or an arm under medical treatment.
- DO NOT apply the arm cuff on your arm while on an intravenous drip or blood transfusion.
- DO NOT use this monitor in areas containing high frequency (HF) surgical equipment, magnetic resonance imaging (MRI) equipment, computerized tomography (CT) scanners. This may result in incorrect operation of the monitor and/or cause an inaccurate reading.
- DO NOT use this monitor in oxygen rich environments or near flammable gas.
- Consult with your physician before using this monitor if you have common arrhythmias such as atrial or ventricular premature beats or atrial fibrillation; arterial sclerosis; poor perfusion; diabetes; pregnancy; pre-eclampsia or renal disease. NOTE that any of these conditions in addition to patient motion, trembling, or shivering may affect the measurement reading.
- NEVER diagnose or treat yourself based on your readings. ALWAYS consult with your physician.
- To help avoid strangulation, keep the air tube and AC adapter cable away from infants, toddlers and children.
- This product contains small parts that may cause a choking hazard if swallowed by infants, toddlers and children.

Data Transmission

- This product emits radio frequencies (RF) in the 2.4 GHz band. DO NOT use this product in locations where RF is restricted, such as on an aircraft or in hospitals. Turn off the **Bluetooth®** feature in this monitor, remove batteries and/or unplug the AC adapter when in RF restricted areas.

AC Adapter (optional accessory) Handling and Usage

- DO NOT use the AC adapter if this monitor or the AC adapter cable is damaged. If this monitor or the cable is damaged, turn off the power and unplug the AC adapter immediately.
- Plug the AC adapter into the appropriate voltage outlet. DO NOT use in a multi-outlet plug.
- NEVER plug in or unplug the AC adapter from the electric outlet with wet hands.
- DO NOT disassemble or attempt to repair the AC adapter.

Battery Handling and Usage

- Keep batteries out of the reach of infants, toddlers and children.

⚠ 2.2 Caution **Indicates a potentially hazardous situation which, if not avoided,may result in minor or moderate injury to the user or patient, or cause damage to the equipment or other property.**

- Stop using this monitor and consult with your physician if you experience skin irritation or discomfort.
- Consult with your physician before using this monitor on an arm where intravascular access or therapy, or an arteriovenous (A-V) shunt, is present because of temporary interference to blood flow and could result in injury.
- Consult with your physician before using this monitor if you have had a mastectomy.

- Consult with your physician before using this monitor if you have severe blood flow problems or blood disorders as cuff inflation can cause bruising.
- DO NOT take measurements more often than necessary because bruising, due to blood flow interference, may occur.
- ONLY inflate the arm cuff when it is applied on your upper arm.
- Remove the arm cuff if it does not start deflating during a measurement.
- DO NOT use this monitor for any purpose other than measuring blood pressure and/or detecting the possibility of Afib.
- During measurement, make sure that no mobile device or any other electrical device that emit electromagnetic fields is within 30 cm of this monitor. This may result in incorrect operation of the monitor and/or cause an inaccurate reading.
- DO NOT disassemble or attempt to repair this monitor or other components. This may cause an inaccurate reading.
- DO NOT use in a location where there is moisture or a risk of water splashing this monitor. This may damage this monitor.
- DO NOT use this monitor in a moving vehicle such as in a car or on an aircraft.
- DO NOT drop or subject this monitor to strong shocks or vibrations.
- DO NOT use this monitor in places with high or low humidity or high or low temperatures. Refer to section 6.
- During measurement, observe the arm to ensure that the monitor is not causing prolonged impairment to blood circulation.
- DO NOT use this monitor in high-use environments such as medical clinics or physician offices.
- DO NOT use this monitor with other medical electrical (ME) equipment simultaneously. This may result in incorrect operation of the monitor and/or cause an inaccurate reading.
- Avoid bathing, drinking alcohol or caffeine, smoking, exercising and eating for at least 30 minutes before taking a measurement.
- Rest for at least 5 minutes before taking a measurement.
- Remove tight-fitting, thick clothing and any accessories from your arm while taking a measurement.
- Remain still and DO NOT talk while taking a measurement.
- ONLY use the arm cuff on persons whose arm circumference is within the specified range of the cuff.
- Ensure that this monitor has acclimated to room temperature before taking a measurement. Taking a measurement after an extreme temperature change could lead to an inaccurate reading. OMRON recommends waiting for approximately 2 hours for the monitor to warm up or cool down when the monitor is used in an environment within the temperature specified as operating conditions after it is stored either at the maximum or at the minimum storage temperature. For additional information on operating and storage / transport temperature, refer to section 6.
- DO NOT use this monitor after the durable period has ended. Refer to section 6.
- DO NOT crease the arm cuff or the air tube excessively.
- DO NOT fold or kink the air tube while taking a measurement. This may cause an injury by interrupting blood flow.
- To unplug the air plug, pull on the plastic air plug at the base of the tube, not the tube itself.
- ONLY use the AC adapter, arm cuff, batteries and accessories specified for this monitor. Use of unsupported AC adapters, arm cuffs and batteries may damage and/or may be hazardous to this monitor.
- ONLY use the approved arm cuff for this monitor. Use of other arm cuffs may result in incorrect readings.
- Inflating to a higher pressure than necessary may result in bruising of the arm where the cuff is applied. NOTE: refer to "If your systolic pressure is more than 210 mmHg" in section 13 of instruction manual (2) for additional information.
- Read and follow the "Correct Disposal of This Product" in section 7 when disposing of the device and any used accessories or optional parts.

Data Transmission

- DO NOT replace batteries or unplug the AC adapter while your readings are being transferred to your smart device. This may result in incorrect operation of this monitor and failure to transfer your blood pressure data.

AC Adapter (optional accessory) Handling and Usage

- Fully insert the AC adapter into the outlet.
- When unplugging the AC adapter from the outlet, be sure to safely pull from the AC adapter. DO NOT pull from the AC adapter cable.
- When handling the AC adapter cable:
  Do not damage it. / Do not break it. / Do not tamper with it. / DO NOT pinch it. / Do not forcibly bend or pull it. / Do not twist it. / DO NOT use it if it is gathered in a bundle. / DO NOT place it under heavy objects.
- Wipe any dust off of the AC adapter.
- Unplug the AC adapter when not in use.
- Unplug the AC adapter before cleaning this monitor.

Battery Handling and Usage

- DO NOT insert batteries with their polarities incorrectly aligned.
- ONLY use 4 "AA" alkaline or manganese batteries with this monitor. DO NOT use other types of batteries. DO NOT use new and used batteries together. DO NOT use different brands of batteries together.
- Remove batteries if this monitor will not be used for a long period of time.
- If battery fluid should get in your eyes, immediately rinse with plenty of clean water. Consult with your physician immediately.

- If battery fluid should get on your skin, wash your skin immediately with plenty of clean, lukewarm water. If irritation, injury or pain persists, consult with your physician.
- DO NOT use batteries after their expiration date.
- Periodically check batteries to ensure they are in good working condition.

2.3 General Precautions

- When you take a measurement on the right arm, the air tube should be at the side of your elbow. Be careful not to rest your arm on the air tube.

- Blood pressure may differ between the right and left arm, and may result in a different measurement value. Always use the same arm for measurements. If the values between both arms differ substantially, check with your physician on which arm to use for your measurements.

Battery Handling and Usage

- Disposal of used batteries should be carried out in accordance with local regulations.

3. Error Messages and Troubleshooting

If any of the below problems occur during measurement, check to make sure that no other electrical device is within 30 cm. If the problem persists, please refer to the table below.

Display/Problem	Possible Cause	Solution
E1 appears or the arm cuff does not inflate.	The [START/STOP] button was pressed while the arm cuff is not applied.	Press the [START/STOP] button again to turn the monitor off. After inserting the air plug securely and applying the arm cuff correctly, press the [START/STOP] button.
	Air plug is not completely plugged into the monitor.	Insert the air plug securely.
	The arm cuff is not applied correctly.	Apply the arm cuff correctly, then take another measurement. Refer to section 7 of instruction manual (2).
	Air is leaking from the arm cuff.	Replace the arm cuff to the new one. Refer to section 14 of instruction manual (2).
E2 appears or a measurement cannot be completed after the arm cuff inflates.	You move or talk during a measurement and the arm cuff does not inflate sufficiently.	Remain still and do not talk during a measurement. If "E2" appears repeatedly, inflate the arm cuff manually until the systolic pressure is 30 to 40 mmHg above your previous readings. Refer to section 13 of instruction manual (2).
	Due to the systolic pressure is above 210 mmHg, a measurement cannot be taken.	
E3 appears	The arm cuff is inflated exceeding the maximum allowable pressure.	Do not touch the arm cuff and/or bend the air tube while taking a measurement. If inflating the arm cuff manually, refer to section 13 of instruction manual (2).
E4 appears	You move or talk during a measurement. Vibrations disrupt a measurement.	Remain still and do not talk during a measurement.
E5 appears	The pulse rate is not detected correctly.	Apply the arm cuff correctly, then take another measurement. Refer to section 7 of instruction manual (2). Remain still and sit correctly during a measurement. If the " " symbol continues to appear, we recommend you to consult with your physician.
/ / appears		
does not flash during a measurement		

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Display/Problem	Possible Cause	Solution
E6 appears	Blood pressure measurements were not taken correctly in an Afib mode measurement.	Apply the arm cuff correctly, then take another measurement. Refer to section 7 of instruction manual (2). Remain still and sit correctly during a measurement. Refer to section 8 of instruction manual (2).
Er appears	The monitor has malfunctioned.	Press the [START/STOP] button again. If "Er" still appears, contact your OMRON retail outlet or distributor.
E rr appears	The montior cannot connect to a smart device or transmit data correctly.	Follow the instructions shown in the "OMRON connect" app. If the "Err" symbol still appears after checking the app, contact your OMRON retail outlet or distributor.
P flashes	The monitor is waiting for pairing with the smart device.	Refer to section 5 of instruction manual (2) for pairing your monitor with your smart device, or press [START/STOP] button to cancel pairing and turn your monitor off.
0 flashes	The monitor is ready to transfer your readings to the smart device.	Open the "OMRON connect" app to transfer your readings.
 flashes	More than 80 readings are not transferred.	Pair or transfer your readings to the "OMRON connect" app so you can keep them in memory in the app, and this error symbol disappears.
	The date and time is not set.	
 appears	100 readings are not transferred.	
 flashes	Batteries are low.	Replacing all 4 batteries with new ones is recommended. Refer to section 4 of instruction manual (2).
 appears or the monitor is turned off unexpectedly during a measurement	Batteries are depleted.	Immediately replace all 4 batteries with new ones. Refer to section 4 of instruction manual (2).
Nothing appears on the display of the monitor.	Battery polarities are not properly aligned.	Check the battery installation for proper placement. Refer to section 4 of instruction manual (2).
Readings appear too high or too low.	Blood pressure varies constantly. Many factors including stress, time of day, and/or how you apply the arm cuff, may affect your blood pressure. Review sections 2 of instruction manual (2).	
Any other communication issue occurs.	Follow the instructions shown in the smart device, or visit the "Help" section in the "OMRON connect" app for further help. If the problem still persists, contact your OMRON retail outlet or distributor.	

Display/Problem	Possible Cause	Solution
Any other problem occurs.		Press the [START/STOP] button to turn the monitor off, then press it again to take a measurement. If the problem continues, remove all batteries and wait for 30 seconds. Then re-install batteries. If the problem still persists, contact your OMRON retail outlet or distributor.
Troubleshooting for Afib indicator function:		
What is different between the Afib indicator function and ECG?		The Afib indicator function and ECG use completely different technologies. An ECG measures the electrical activity of the heart and can be used to diagnose Afib. The Afib indicator function detects irregular heartbeat and can suggest the possibility of Afib with a sensitivity of 95.5% and specificity of 93.8%. Refer to section 11 for details.
If the " " symbol does not appear, it means there is no possibility of Afib?		Even if the " " symbol does not appear, there is still a possibility of Afib.
Should I consult with my physician if the " " symbol appears?		We recommend you to consult with your physician because there is a possibility of Afib. However, the " " symbol may be displayed for other reasons, such as other heart arrhythmias.
What is different between Afib indicator function and irregular heart beat function?		The irregular heartbeat function detects irregularities in the pulse waves in one measurement. The Afib indicator function suggests the possibility of Afib when blood pressure is measured 3 consecutive times.
What should I do if the " " symbol sometimes appears?		Afib does not always have symptoms. We recommend you to consult with and follow the directions of your physician.
I have been diagnosed with Afib by the physician, but the " " symbol does not appear.		Afib may not occur at the time of specific blood pressure measurements. We recommend you to consult with your physician regularly.
Is the blood pressure reading reliable when the " " symbol appears?		Afib or an irregular heartbeat can influence your blood pressure measurements and make it difficult to get an accurate reading. Repeated measurements may be required to overcome variabilities.* In Afib mode, the blood pressure measurement is taken 3 times, and the average is displayed. The monitor will indicate an error message (E5/E6) if the influence of the irregular heartbeat is too severe to give a measurement result. If this occurs repeatedly, we recommend that you consult with your physician.

* Prof. Roland Asmar et al. European Society of Hypertension Recommendations for Conventional, Ambulatory and Home Blood Pressure Measurement

4. Limited Warranty

Thank you for buying an OMRON product. This product is constructed of high quality materials and great care has been taken in its manufacturing. It is designed to give you every satisfaction, provided that it is properly operated and maintained as described in the instruction manual.

This product is warranted by OMRON for a period of 3 years after the date of purchase. The proper construction, workmanship and materials of this product is warranted by OMRON. During this period of warranty OMRON will, without charge for labour or parts, repair or replace the defect product or any defective parts.

The warranty does not cover any of the following:

A. Transport costs and risks of transport.
B. Costs for repairs and / or defects resulting from repairs done by unauthorised persons.
C. Periodic check-ups and maintenance.
D. Failure or wear of optional parts or other attachments other than the main device itself, unless explicitly warranted above.
E. Costs arising due to non-acceptance of a claim (those will be charged for).
F. Damages of any kind including personal caused accidentally or from misuse.
G. Calibration service is not included within the warranty.
H. Optional parts have a one (1) year warranty from date of purchase. Optional parts include, but are not limited to the following items: cuff and cuff tube.

Should warranty service be required please apply to the dealer whom the product was purchased from or an authorised OMRON distributor. For the address refer to the product packaging / literature or to your specialised retailer. If you have difficulties in finding OMRON customer services, contact us for information:

www.omron-healthcare.com

Repair or replacement under the warranty does not give rise to any extension or renewal of the warranty period.

The warranty will be granted only if the complete product is returned together with the original invoice / cash ticket issued to the consumer by the retailer.

5. Maintenance

5.1 Maintenance

To protect your monitor from damage, follow the directions below:
Changes or modifications not approved by the manufacturer will void the user warranty.

⚠ Caution

DO NOT disassemble or attempt to repair this monitor or other components. This may cause an inaccurate reading.

5.2 Storage

- Keep your monitor in the storage case when not in use.
  1. Remove the arm cuff from the monitor.

⚠ Caution

To unplug the air plug, pull on the plastic air plug at the base of the tube, not the tube itself.

  2. Gently fold the air tube into the arm cuff. Note: Do not bend or crease the air tube excessively.
  3. Place your monitor and other components in the storage case.
- Store your monitor and other components in a clean, safe location.
- Do not store your monitor and other components:
  - If your monitor and other components are wet.
  - In locations exposed to extreme temperatures, humidity, direct sunlight, dust or corrosive vapors such as bleach.
  - In locations exposed to vibrations or shocks.
- To protect your monitor during storage, an optional LCD cover is available as accessory. Refer to section 15 of Instruction Manual (2).

5.3 Cleaning

- Do not use any abrasive or volatile cleaners.
- Use a soft dry cloth or a soft cloth moistened with mild (neutral) detergent to clean your monitor and arm cuff, and then wipe them with a dry cloth.
- Do not wash or immerse your monitor and arm cuff or other components in water.
- Do not use gasoline, thinners or similar solvents to clean your monitor and arm cuff or other components.

5.4 Calibration and Service

- The accuracy of this blood pressure monitor has been carefully tested and is designed for a long service life.
- It is generally recommended to have the unit inspected every two years to ensure correct functioning and accuracy. Please consult your authorised OMRON dealer or the OMRON Customer Service at the address given on the packaging or attached literature.

6. Specifications

Product Category	Electronic Sphygmomanometers
Product description	Automatic Upper Arm Blood Pressure Monitor
Model (Code)	M7 Intelli IT (HEM-7361T-EBK) / X7 Smart (HEM-7361T-ESL)
Display	LCD digital display
Cuff pressure range	0 to 299 mmHg
Blood pressure measurement range	SYS: 60 to 260 mmHg DIA: 40 to 215 mmHg
Pulse measurement range	40 to 180 beats / min.
Accuracy	Pressure: ±3 mmHg Pulse: ±5% of display reading
Inflation	Automatic by electric pump
Deflation	Automatic pressure release valve
Measurement method	Oscillometric method
Transmission method	**Bluetooth**® Low Energy
Wireless communication	Frequency range: 2.4 GHz (2400 - 2483.5 MHz) / Modulation: GFSK Effective radiated power: < 20 dBm
Operation mode	Continuous operation
IP classification	Monitor: IP20 Optional AC adapter: IP21
Rating	DC6 V 4.0 W
Power source	4 "AA" batteries 1.5 V or optional AC adapter (INPUT AC 100 - 240 V 50/60 Hz 0.12 - 0.065 A)
Battery life	Approximately 1000 measurements (using new alkaline batteries) The number of times may decrease when using Afib mode because one Afib indication consists of 3 regular measurements.
Durable period (Service life)	Monitor: 5 years / Cuff: 5 years / Optional AC adapter: 5 years
Operating conditions	+10 to +40°C / 15 to 90% RH (non-condensing) / 800 to 1060 hPa
Storage / Transport conditions	-20 to +60°C / 10 to 90% RH (non-condensing)
Weight	Monitor: approximately 460 g (not including batteries) Arm cuff: approximately 163 g
Dimensions (approximately value)	Monitor: 191 mm (W) × 85 mm (H) × 120 mm (L) / Arm cuff:145 mm × 532 mm (air tube: 750 mm)
Cuff circumference applicable to the monitor	220 to 420 mm
Memory	Stores up to 100 readings per user
Contents	Monitor, arm cuff (HEM-FL31), 4 "AA" batteries, Instruction Manual ① and ②, setup instructions, storage case
Protection against electric shock	Internally powered ME equipment (When using only batteries) Class II ME equipment (Optional AC adapter)
Applied part	Type BF (arm cuff)

Note

- These specifications are subject to change without notice.
- This monitor is clinically investigated according to the requirements of ISO 81060-2:2013. In the clinical validation study, K5 was used on 85 subjects for determination of diastolic blood pressure.
- This device has been validated for use on pregnant and pre-eclampsia patients according to the Modified European Society of Hypertension Protocol*.
- This device has been validated for use on diabetic (Type II) population**.

- IP classification is degrees of protection provided by enclosures in accordance with IEC 60529. This monitor and optional AC adapter are protected against solid foreign objects of 12.5 mm diameter and greater such as a finger. The optional AC adapter is protected against vertically falling water drops which may cause issues during a normal operation.

* Topouchian J et al. Vascular Health and Risk Management 2018:14 189–197
** Chahine M.N. et al. Medical Devices: Evidence and Research 2018:11 11–20

About a wireless communication interference

This product operates in an unlicensed ISM band at 2.4 GHz. In the event this product is used near other wireless devices such as microwave and wireless LAN, which operate on the same frequency band as this product, there is a possibility that interference may occur. If interference occurs, stop the operation of the other devices or relocate this product away from other wireless devices before attempting to use it.

7. Correct Disposal of This Product (Waste Electrical & Electronic Equipment)

This marking shown on the product or its literature, indicates that it should not be disposed of, with other household wastes at the end of its working life.
To prevent possible harm to the environment or human health from uncontrolled waste disposal, please separate this product from other types of wastes and recycle it responsibly to promote the sustainable reuse of material resources.
Household users should contact either the retailer where they purchased this product, or their local government office, for details of where and how they can return this item for environmentally safe recycling.
Business users should contact their supplier and check the terms and conditions of the purchase contract. This product should not be mixed with other commercial waste for disposal.

8. Important Information regarding Electromagnetic Compatibility (EMC)

HEM-7361T-EBK/ESL conforms to the EN60601-1-2:2015 Electromagnetic Compatibility (EMC) standard.
Further documentation in accordance with this EMC standard is available at OMRON HEALTHCARE EUROPE at the address mentioned in this instruction manual or at: www.omron-healthcare.com.

9. Guidance and Manufacturer's Declaration

- This blood pressure monitor is designed according to the European Standard EN1060, Non-invasive sphygmomanometers Part 1: General Requirements and Part 3: Supplementary requirements for electromechanical blood pressure measuring systems.
- Hereby, OMRON HEALTHCARE Co., Ltd., declares that the radio equipments type HEM-7361T-EBK/ESL is in compliance with Directive 2014/53/EU.
- The full text of the EU declaration of conformity is available at the following internet address: www.omron-healthcare.com
- This OMRON product is produced under the strict quality system of OMRON HEALTHCARE Co., Ltd., Japan. The Core component for OMRON blood pressure monitors, which is the Pressure Sensor, is produced in Japan.
- Please report to the manufacturer and the competent authority of the Member State in which you are established about any serious incident that has occurred in relation to this device.

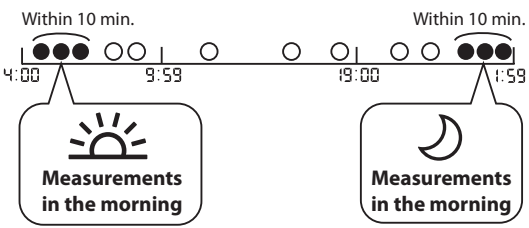
10. How to Calculate Weekly Averages

Morning Weekly Average Calculation

This is the average for the measurements taken during the morning (4:00 - 9:59) between Sunday and Saturday. The 2 or 3 readings taken within the first 10 minute timeframe in the morning between 4:00 - 9:59 will be used to calculate the morning average for each day.

Evening Weekly Average Calculation

This is the average for the measurements taken during the evening (19:00 - 1:59) between Sunday and Saturday. The 2 or 3 readings taken within the last 10 minute timeframe in the evening between 19:00 - 1:59 will be used to calculate the evening average for each day.



11. Useful Information

What is Blood Pressure?

Blood pressure is a measure of the force of blood flowing against the walls of the arteries. Arterial blood pressure is constantly changing during the course of the heart's cycle.
The highest pressure in the cycle is called the Systolic Blood Pressure; the lowest is the Diastolic Blood Pressure. Both pressures, the Systolic and Diastolic, are necessary to enable a physician to evaluate the status of a patient's blood pressure.

What is Arrhythmia?

Arrhythmia is a condition where the heartbeat rhythm is abnormal due to flaws in the bio-electrical system that drives the heartbeat. Typical symptoms are skipped heartbeats, premature contraction, an abnormally rapid (tachycardia) or slow (bradycardia) pulse.

What is Afib?

Atrial fibrillation (also called AFib or AF) is a quivering or irregular heartbeat (arrhythmia) that can lead to blood clots, stroke, heart failure and other heart-related complications. During atrial fibrillation, the heart's two upper chambers (the atria) beat chaotically and irregularly — out of coordination with the two lower chambers (the ventricles) of the heart. Episodes of atrial fibrillation can come and go, or you may develop atrial fibrillation that doesn't go away and may require treatment.
Afib indicator function detects the possibility of Afib with an accuracy of 94.2% (with sensitivity of 95.5% and specificity of 93.8%) as demonstrated in the study* with Single-lead ECG as reference measurement.

*M. Ishizawa, T. Noma, T. Minamino et al., Multiple measurements with automated blood pressure monitor can detect atrial fibrillation with high sensitivity and specificity in general cardiac patients, ESC Congress 2018

Symbols Description

FR Description des symboles
DE Beschreibung der Symbole
IT Descrizione dei simboli
ES Descripción de los símbolos
NL Beschrijving van symbolen
RU Описание символов
TR Simgelerin Açıklaması
AR وصف الرموز

Applied part - Type BF Degree of protection against electric shock (leakage current)

FR Partie appliquée - Type BF Degré de protection contre les chocs électriques (courant de fuite)

DE Anwendungsteil – Typ BF Schutz vor Stromschlägen (Ableitstrom)

IT Parti applicate - Tipo BF Livello di protezione contro le folgorazioni (corrente di dispersione)

ES Partes en contacto: Tipo BF Grado de protección contra descargas eléctricas (corriente de fuga)

NL Toegepast onderdeel - Type BF-beschermingsgraad tegen elektrische schokken (lekstroom)

RU Рабочая часть аппарата - Тип BF Степень защиты от поражения электрическим током (токи утечки)

TR Uygulanan parça - Tip BF Elektrik çarpmasına karşı koruma derecesi (kaçak akım)

AR جزء مطبق من النوع BF - درجة الحماية ضد الصدمات الكهربائية (التيار المتسرب)

Class II equipment. Protection against electric shock

FR Équipement de classe II. Protection contre les chocs électriques

DE Gerät der Klasse II. Schutz vor Stromschlägen

IT Apparecchiatura di Classe II. Protezione contro le folgorazioni

ES Equipo de Clase II. Protección contra descargas eléctricas

NL Apparatuur van Klasse II. Bescherming tegen elektrische schokken

RU Оборудование класса II. Защита от поражения электрическим током

TR Sınıf II ekipman. Elektrik çarpmasına karşı koruma

AR جهاز من الفئة II. الحماية ضد الصدمات الكهربائية

IP XX

Ingress protection degree provided by IEC 60529

FR Degré de protection selon CEI 60529

DE Grad des Eindringschutzes gemäß IEC 60529

IT Livello di protezione IP in base a IEC 60529

ES Grado de protección según la norma internacional IEC 60529

NL Beschermingsklasse volgens IEC 60529

RU Степень защиты, обеспечиваемая оболочкой, в соответствии со стандартом IEC 60529

TR Su girmesine karşı koruma derecesi IEC 60529 tarafından verilmiştir

AR درجة الحماية من التسرب وفقًا لمعيار IEC 60529

CE

CE Marking

FR Marquage CE
DE CE-Kennzeichnung
IT Contrassegno CE
ES Marcado CE
NL CE-merkteken
RU Знак соответствия директиве ЕС
TR CE İşareti
AR علامة التوافق مع اللجنة الأوروبية (CE)

SN

Serial number

FR Numéro de série
DE Seriennummer
IT Numero di serie
ES Número de serie
NL Serienummer
RU Серийный номер
TR Seri numarası
AR الرقم المتسلسل

LOT

LOT number

FR Numéro de LOT
DE LOT-Nummer
IT Numero di lotto
ES Número de lote
NL Partijnummer
RU Номер ПАРТИИ
TR Parti numarası
AR رقم التشغيلة

MD

Medical device

FR Dispositif médical
DE Medizinprodukt
IT Dispositivo medico
ES Producto sanitario
NL Medisch apparaat
RU Медицинский прибор
TR Tıbbi cihaz
AR جهاز طبي

Indicates the manufacturer's catalogue number

FR Indique le numéro de catalogue du fabricant
DE Angabe der Hersteller-Katalognummer
IT Indica il numero di catalogo del produttore
ES Indica el número de catálogo del fabricante
NL Geeft het catalogusnummer van de fabrikant aan
RU Указывает номер в каталоге производителя
TR Üreticinin katalog numarasını belirtir
AR للإشارة إلى رقم القائمة الخاص بالشركة المصنعة

Temperature limitation

FR Limitation de température
DE Temperaturbegrenzung
IT Limite di temperatura
ES Limitación de la temperatura
NL Temperatuurbegrenzing
RU Температурный диапазон
TR Sıcaklık sınırlaması
AR حدود درجة الحرارة المناسبة

Humidity limitation

FR Limitation d'humidité
DE Luftfeuchtigkeits-begrenzung
IT Limite di umidità
ES Limitación de la humedad
NL Vochtigheidsbegrenzing
RU Диапазон влажности
TR Nem sınırlaması
AR حدود الرطوبة المناسبة

Atmospheric pressure limitation

FR Limitation de pression atmosphérique
DE Luftdruckbegrenzung
IT Limite di pressione atmosferica
ES Limitación de la presión atmosférica
NL Luchtdrukbegrenzing
RU Диапазон атмосферного давления
TR Atmosferik basınç sınırlaması
AR حدود الضغط الجوي المناسب

Indication of connector polarity

FR Indication de la polarité des connecteurs
DE Anzeige der Steckerpolarität
IT Indicazione della polarità dei connettori
ES Indicación de la polaridad del conector
NL Indicatie van polariteit van aansluiting
RU Индикация полярности разъема
TR Bağlantı polarite göstergesi
AR علامة تشير إلى قطبية الموصل

For indoor use only

FR Pour un usage à l'intérieur uniquement
DE Nur für die Nutzung in Innenbereichen
IT Solo per uso in interni
ES Para uso solo en interiores
NL Alleen voor gebruik binnenshuis
RU Для использования только внутри помещений
TR Sadece iç mekanda kullanım için
AR صالح للاستخدام في الأماكن المغلقة فقط

OMRON's trademarked technology for blood pressure measurement

FR Technologie brevetée OMRON pour la mesure de la pression artérielle
DE Markenrechtlich geschützte Technologie von OMRON zur Blutdruckmessung
IT Tecnologia brevettata OMRON per la misurazione della pressione arteriosa
ES La tecnología de OMRON para medir la presión arterial
NL Technologie voor bloeddrukmeting onder handelsmerk van OMRON
RU Зарегистрированная технология измерения артериального давления OMRON
TR OMRON'un kan basıncı ölçümü için ticari markalı teknolojisidir
AR تقنية العلامة التجارية OMRON لقياس ضغط الدم

Identifier of cuffs compatible for the device

FR Identificateur des brassards compatibles avec l'appareil

DE Kennzeichnung der mit dem Gerät kompatiblen Manschetten

IT Identifica i bracciali compatibili con il dispositivo

ES Identificador para manguitos compatibles con el dispositivo

NL Identificatie van manchetten die compatibel zijn met het apparaat

RU Манжеты, совместимые с устройством

TR Cihaz ile uyumlu kollukların tanıtım işareti

AR علامات تحديد الشرائط الضاغطة المتوافقة مع الجهاز

Marker on the cuff to be positioned above the artery

FR Repère sur le brassard, à positionner au-dessus de l'artère

DE Markierung auf der Manschette, die oberhalb der Arterie liegen muss

IT Contrassegno sul bracciale da posizionare al di sopra dell'arteria

ES La marca del manguito debe colocarse sobre la arteria

NL Markering op de manchet die boven de slagader moet worden geplaatst

RU Указатель на манжете для расположения над артерией

TR Kolluk üzerindeki işaretin konumu arterin üzerine gelmelidir

AR علامة بالشريط الضاغط للإشارة إلى وجوب وضعه فوق الشريان

Manufacturer's quality control mark

FR Marque de contrôle de la qualité du fabricant

DE Qualitätskontrollzeichen des Herstellers

IT Contrassegno controllo qualità del produttore

ES Marca del control de calidad del fabricante

NL Symbool voor kwaliteitscontrole van fabrikant

RU Отметка производителя о контроле качества

TR Üreticinin kalite kontrol işareti

AR علامة التحكم في الجودة الخاصة بالشركة المصنعة

LATEX FREE

Not made with natural rubber latex

FR Ne contient pas de latex de caoutchouc naturel

DE Enthält kein Naturlatex

IT Non contiene lattice di gomma naturale

ES No contiene látex de caucho natural

NL Bevat geen natuurrubberlatex

RU Не содержит натуральный латекс

TR Doğal kauçuk lateksten üretilmemiştir

AR تم التصنيع دون استخدام لاتيكس المطاط الطبيعي

Arm circumference

FR Circonférence du bras

DE Armumfang

IT Circonferenza del braccio

ES Perímetro de brazo

NL Armomtrek

RU Окружность плеча

TR Kol çevresi

AR محيط الذراع

Necessity for the user to consult this instruction manual

FR L'utilisateur doit consulter le mode d'emploi

DE Der Benutzer muss diese Gebrauchsanweisung lesen

IT L'utente deve consultare il presente manuale di istruzioni

ES Es necesario que el usuario consulte este manual de instrucciones

NL De gebruiker dient deze gebruiksaanwijzing te raadplegen

RU Необходимость для пользователя обратиться к данному руководству по эксплуатации

TR Kullanıcı, bu kullanım kılavuzuna başvurmalıdır

AR ضرورة رجوع المستخدم إلى دليل الإرشادات هذا

Need for the user to follow this instruction manual thoroughly for your safety.

FR L'utilisateur doit suivre attentivement ce mode d'emploi pour votre sécurité.

DE Damit die Sicherheit gewährleistet ist, muss der Benutzer diese Gebrauchsanweisung sorgfältig befolgen.

IT Per la propria sicurezza, l'utente deve seguire attentamente il presente manuale di istruzioni.

ES Es necesario que el usuario siga rigurosamente este manual de instrucciones para su seguridad.

NL Voor de eigen veiligheid dient de gebruiker zich zorgvuldig aan deze gebruiksaanwijzing te houden.

RU В целях обеспечения безопасности строго следуйте указаниям в данном руководстве по эксплуатации.

TR Güvenlik açısından kullanıcının bu kullanım kılavuzuna dikkatle uyması gerekir.

AR حاجة المستخدم إلى اتباع دليل الإرشادات بالكامل للحفاظ على السلامة.

Direct current

FR Courant continu
DE Gleichstrom
IT Corrente diretta
ES Corriente directa
NL Gelijkstroom
RU Постоянный ток
TR Doğru akım
AR التيار المباشر

Alternating current

FR Courant alternatif
DE Wechselstrom
IT Corrente alternata
ES Corriente alterna
NL Wisselstroom
RU Переменный ток
TR Alternatif akım
AR التيار المتناوب

Date of manufacture

FR Date de fabrication
DE Herstellungsdatum
IT Data di fabbricazione
ES Fecha de fabricación
NL Productiedatum
RU Дата изготовления
TR Üretim tarihi
AR تاريخ التصنيع

Prohibited action

FR Action interdite
DE Verbotene Aktion
IT Operazione proibita
ES Acción prohibida
NL Verboden handeling
RU Запрещенные действия
TR Yasaklanmış eylem
AR اجراء محظور

To indicate generally elevated, potentially hazardous, levels of non-ionizing radiation, or to indicate equipment or systems. e.g. in the medical electrical area that include RF transmitters or that intentionally apply RF electromagnetic energy for diagnosis or treatment.

FR Pour indiquer des niveaux généralement élevés, potentiellement dangereux, de rayonnement non ionisant, ou pour indiquer l'équipement ou les systèmes, par exemple dans le domaine de l'électricité médicale qui comprennent des émetteurs RF ou qui utilisent intentionnellement l'énergie électromagnétique RF pour le diagnostic ou le traitement.

DE Als Hinweis auf allgemein erhöhte, potenziell gefährliche Stufen nicht-ionisierender Strahlung oder als Hinweis auf Geräte oder Systeme zum Beispiel im medizinisch-elektrischen Bereich, etwa HF-Übertragungsgeräte, bzw. auf solche, die elektromagnetische HF-Strahlung zur Diagnose oder Behandlung verwenden.

IT Indica livelli generalmente elevati, potenzialmente pericolosi, di radiazioni non ionizzanti oppure indica apparecchiature o sistemi (ad esempio per le aree elettromedicali in cui sono presenti trasmettitori RF o in cui viene intenzionalmente applicata energia elettromagnetica a radiofrequenza per la diagnosi o il trattamento).

ES Para indicar niveles de radiación no ionizante generalmente elevados y potencialmente peligrosos, o bien para indicar equipos o sistemas, como los usados en el ámbito electro médico, que incorporen transmisores de radiofrecuencia o que apliquen energía electromagnética de radiofrecuencia intencionadamente para diagnósticos o tratamientos.

NL Geeft in het algemeen verhoogde, potentieel gevaarlijke niveaus aan van niet-ioniserende straling of duidt op apparatuur of systemen, bijvoorbeeld in de medische elektrische omgeving, die RF-zenders bevatten of die opzettelijk elektromagnetische RF-energie toepassen voor diagnose of behandeling.

RU Для указания повышенных и потенциально опасных уровней неионизирующей радиации или же оборудования или систем (например, в зоне размещения медицинского электрооборудования), включающих в себя радиопередатчики или устройства, использующие радиочастотную электромагнитную энергию в целях диагностики или терапии.

TR Genellikle yüksek ve zararlı olabilecek iyonlaşmayan radyasyon seviyelerini belirtir veya RF vericileri içeren veya tanı ya da tedavi amacıyla bilinçli olarak RF elektromanyetik enerji uygulayan (örneğin medikal elektrik alanında bulunan) ekipman ve sistemleri belirtir.

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Issue Date:
Date de publication :
Ausgabedatum:
Data di pubblicazione:
Fecha de publicación: 2019-09-04
Uitgiftedatum
Дата выпуска:
Teslim Tarihi:
تاريخ الإصدار :

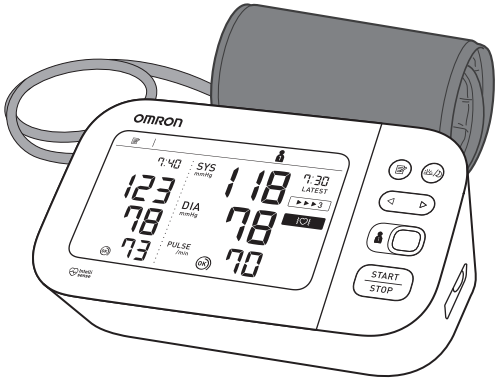
IM1-HEM-7361T-E-02-09/2019
2895864-6B

Instruction Manual 2

OMRON

Automatic Upper Arm Blood Pressure Monitor

M7 Intelli IT (HEM-7361T-EBK)
X7 Smart (HEM-7361T-ESL)



Intellisense

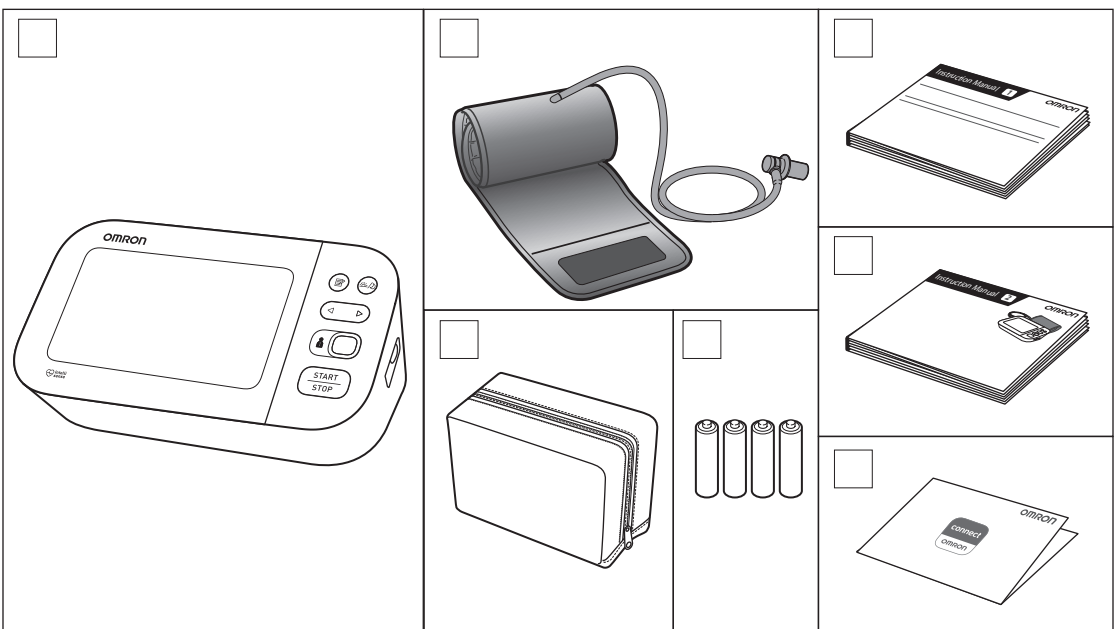
All for Healthcare

Read Instruction manual (1) and (2) before use.

FR Lire le mode d'emploi (1) et (2) vant l'utilisation.
DE Lesen Sie vor der Verwendung Gebrauchsanweisung (1) und (2).
IT Leggere il manuale di istruzioni (1) e (2) prima dell'uso.
ES Lea el manual de instrucciones (1) y (2) antes del uso.
NL Lees gebruiksaanwijzing (1) en (2) voor gebruik.
RU Прочтите руководства по эксплуатации (1) и (2) перед использованием.
TR Kullanmadan önce, kullanım kılavuzu (1) ve (2)'yi okuyun.
AR اقرأ دليل الإرشادات (1) و(2) قبل الاستخدام.

1 Package Contents

FR Contenu de l'emballage
DE Packungsinhalt
IT Contenuto della confezione
ES Contenido del envase
NL Inhoud van de verpakking
RU Комплект поставки
TR Paketin İçindekiler
AR محتويات العلبة



2 Preparing for a Measurement

FR Préparation pour une prise de mesure
DE Vorbereiten einer Messung
IT Preparazione per la misurazione
ES Preparación para una medición
NL Een meting voorbereiden
RU Подготовка к измерению
TR Ölçüm Hazırlığı
AR التجهيز لعملية القياس

30 minutes before

FR 30 minutes avant

DE 30 Minuten vorher

IT 30 minuti prima

ES 30 minutos antes

NL 30 minuten ervoor

RU За 30 минут до

TR 30 dakika önce

AR لق إلى مع. لفة في تصمم واعيد التصمم والمحتوى مثل يدية لأو

5 minutes before: Relax and rest.

FR 5 minutes avant : détente et repos.

DE 5 Minuten vorher: ruhig hinsetzen.

IT 5 minuti prima: rilassarsi e stare a riposo.

ES 5 minutos antes: relájese y descanse.

NL 5 minuten ervoor: ontspan en rust.

RU За 5 минут до: расслабьтесь и отдохните.

TR 5 dakika önce: Gevşeyin ve dinlenin.

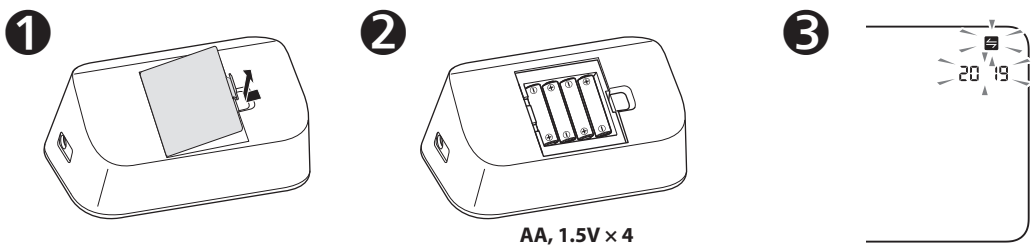
AR قبل القياس بمدة تبلغ ٥ دقائق: استرح واسترخ

3 Downloading the "OMRON connect" App

FR Téléchargement de l'application « OMRON connect »
DE Herunterladen der App „OMRON connect“
IT Download dell'app “OMRON connect”
ES Descarga de la aplicación “OMRON connect”
NL De app “OMRON connect“ downloaden
RU Загрузка приложения «OMRON connect»
TR "OMRON Connect" Uygulamasını İndirme
AR تنزيل تطبيق "اتصال OMRON"

4 Inserting Batteries

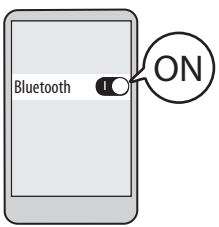
FR Mise en place des piles
DE Einlegen von Batterien
IT Inserimento delle batterie
ES Introducción de las pilas
NL De batterijen plaatsen
RU Установка элементов питания
TR Pilleri Takma
AR إدخال البطاريات



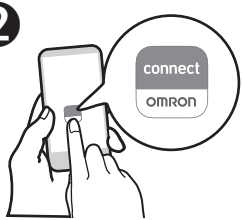
5 Pairing Your Smart Device

FR Jumelage de votre appareil intelligent
DE Koppeln mit Smartphone oder Tablet
IT Associazione del dispositivo smart
ES Sincronización con un dispositivo inteligente
NL Uw smartapparaat koppelen
RU Синхронизация со смарт-устройством
TR Akıllı Cihazınızın Eşleştirilmesi
AR إقران هاتفك الذكي

1



2



3

Follow the instructions.

FR Suivez les instructions.
DE Befolgen Sie die Anweisungen.
IT Attenersi alle istruzioni.
ES Siga las instrucciones.
NL Volg de instructies.
RU Выполняйте инструкции.
TR Talimatları izleyin.
AR اتبع الإرشادات.

The date and time will automatically be set when your monitor is paired with the app.

FR La date et l'heure seront automatiquement réglées lorsque votre tensiomètre sera jumelé avec l'application.
DE Wenn Ihr Messgerät mit der App verknüpft ist, werden Datum und Uhrzeit automatisch eingestellt.
IT La data e l'ora verranno impostate automaticamente una volta che il misuratore verrà associato alla app.
ES La fecha y la hora se ajustarán automáticamente cuando el monitor se sincronice con la aplicación.
NL De datum en tijd worden automatisch ingesteld wanneer uw bloeddrukmeter is gekoppeld aan de app.
RU Дата и время установятся автоматически после синхронизации с приложением.
TR Ölçüm cihazınız uygulamayla eşleştirildiğinde tarih ve saat otomatik olarak ayarlanır.
AR سيتم ضبط الوقت والتاريخ تلقائيًا عند إقران جهاز القياس الخاص بك بالتطبيق.

6 Setting Date and Time Manually

FR Réglage manuel de la date et de l'heure
DE Manuelles Einstellen von Datum und Uhrzeit
IT Impostazione manuale di data e ora
ES Ajuste manual de la fecha y la hora
NL Datum en tijd handmatig instellen
RU Установка даты и времени вручную
TR Tarih ve Saatin Manuel Olarak Ayarlanması
AR ضبط الوقت والتاريخ يدويًا

If your monitor is paired with your smart device, date and time is set automatically. When you need to set them manually, set year > month > day > hour > minute.

FR Si votre tensiomètre est jumelé avec votre appareil intelligent, la date et l'heure sont réglées automatiquement. Si vous devez les régler manuellement, réglez dans l'ordre année > mois > jour > heure > minute.

DE Wenn Ihr Messgerät mit Ihrem Smartphone oder Tablet gekoppelt ist, werden Datum und Uhrzeit automatisch eingestellt. Bei manueller Einstellung müssen Sie Jahr > Monat > Tag > Stunde > Minute einstellen.

IT Se il misuratore viene accoppiato con un dispositivo smart, la data e l'ora verranno impostate automaticamente. Se occorre impostarle manualmente, impostare anno > mese > giorno > ore > minuti.

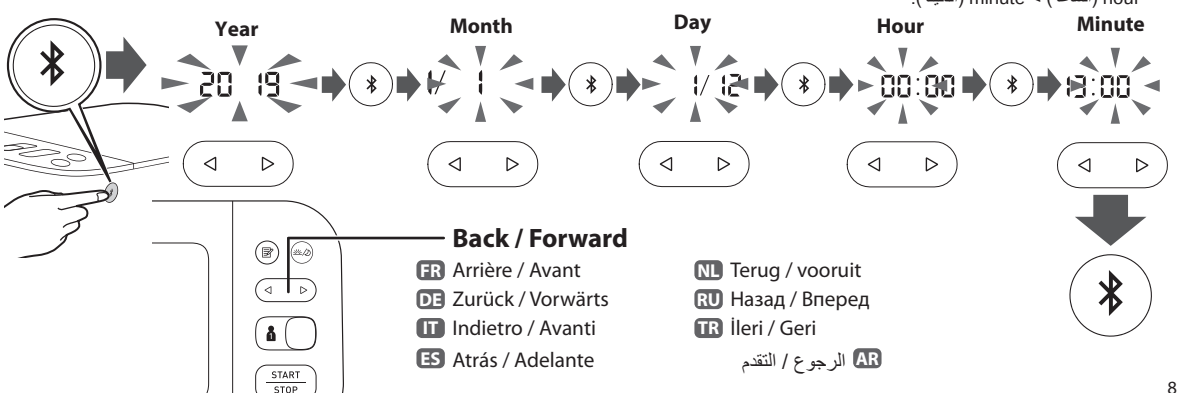
ES Si el monitor está sincronizado con su dispositivo inteligente, la fecha y la hora se ajustarán automáticamente. Cuando tenga que configurarlas manualmente, ajuste año > mes > día > hora > minuto.

NL Als uw bloeddrukmeter gekoppeld is met uw smartapparaat, worden de datum en tijd automatisch ingesteld. Wanneer u ze handmatig moet instellen, stelt u jaar > maand > dag > uur > minuut in.

RU Если прибор синхронизирован со смарт-устройством, дата и время устанавливаются автоматически. Если необходимо установить их вручную, выберите год > месяц > день > час > минута.

TR Ölçüm cihazınız akıllı cihazınızla eşleştirilmişse tarih ve saat otomatik olarak ayarlanır. Bunları manuel olarak ayarlamanız gerektiğinde, yıl > ay > gün > saat > dakika değerini ayarlayın.

AR في حال إقران جهاز القياس الخاص بك بالجهاز الذكي، يتم ضبط الوقت والتاريخ تلقائيًا. عند الحاجة إلى ضبطهما يدويًا، اضبط year (السنة) > month (الشهر) > day (اليوم) > hour (الساعة) > minute (الدقيقة).

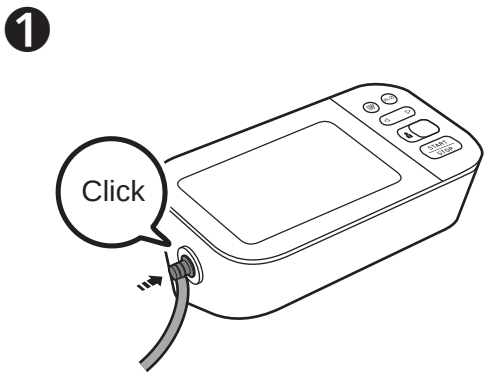


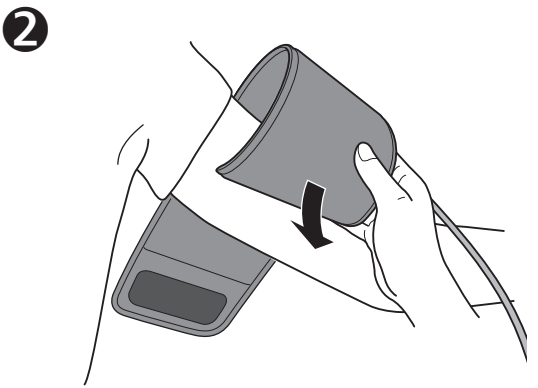
Back / Forward

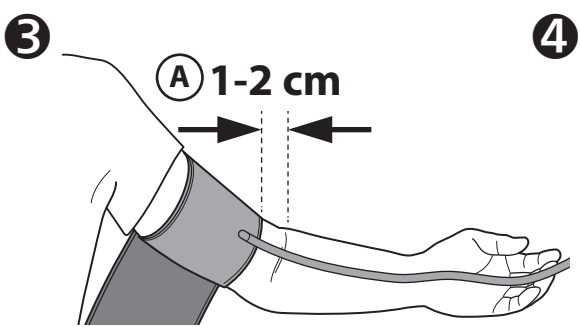
FR Arrière / Avant
DE Zurück / Vorwärts
IT Indietro / Avanti
ES Atrás / Adelante
NL Terug / vooruit
RU Назад / Вперед
TR İleri / Geri
AR الرجوع / التقدم

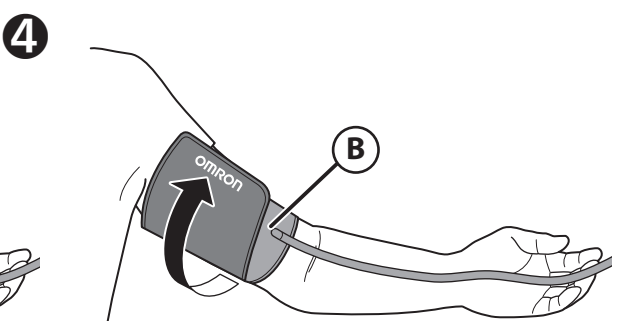
7 Applying the Cuff on the Left Arm

FR Pose du brassard sur le bras gauche
DE Anbringen der Manschette am linken Arm
IT Applicazione del bracciale sul braccio sinistro
ES Colocación del manguito en el brazo izquierdo
NL De manchet op de linkerarm aanbrengen
RU Расположение манжеты на левой руке
TR Kolluğun Sol Kola Takılması
AR لف الشريط الضاغط على الذراع الأيسر









(A) Tube side of the cuff should be 1 - 2 cm above the inside elbow.

FR Le côté tuyau du brassard doit être positionné 1 à 2 cm au-dessus de l'intérieur du coude.

DE Das Manschettenstück mit dem Schlauch muss 1 bis 2 cm oberhalb des Ellbogens liegen.

IT Il lato del bracciale con il tubo deve trovarsi al di sopra del gomito interno, a una distanza di circa 1 o 2 cm.

ES El lado del tubo del manguito deberá quedar 1 o 2 cm por encima de la parte interna del codo.

NL De kant met de slang van de manchet moet 1 - 2 cm boven de binnenkant van de elleboog liggen.

RU Край манжеты с трубкой должен находиться на 1 - 2 см выше локтевого сгиба.

TR Kolluğun boru tarafı, dirsek içinin 1-2 cm üstünde olmalıdır.

AR يجب أن يرتفع جانب الخرطوم من الشريط الضاغط من ١ إلى ٢ سم عن المرفق.

If taking measurements on the right arm, refer to:

FR Pour la prise de mesures au bras droit, voir :

DE Bei Messungen am rechten Arm siehe:

IT Se la misurazione viene eseguita al braccio destro, fare riferimento a:

ES Si va a realizar mediciones en el brazo derecho, consulte:

NL Als u metingen aan de rechterarm uitvoert, raadpleeg dan:

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☞ 2.3

RU При измерении на правой руке см.

TR Sağ koldan ölçüm yapıyorsanız aşağıdakilere bakın:

AR في حال إجراء القياس على الذراع الأيمن، ارجع إلى:

(B) Make sure that air tube is on the inside of your arm and wrap the cuff securely so it can no longer slip round.

FR Assurez-vous que le tuyau à air se trouve du côté intérieur de votre bras et enroulez fermement le brassard de manière qu'il ne puisse plus tourner.

DE Stellen Sie sicher, dass der Luftschlauch an der Arminnenseite sitzt, und befestigen Sie die Manschette sicher, so dass sie nicht verrutscht.

IT Assicurarsi che il tubo dell'aria si trovi all'interno del braccio e avvolgere il bracciale saldamente in modo che non possa ruotare.

ES Asegúrese de que el tubo de aire se encuentra en la cara interna del brazo y enrolle el manguito con firmeza para que no pueda deslizarse.

NL Zorg ervoor dat de luchtslang zich aan de binnenkant van uw arm bevindt en wikkel de manchet stevig rond uw arm zodat deze niet meer kan wegglijden.

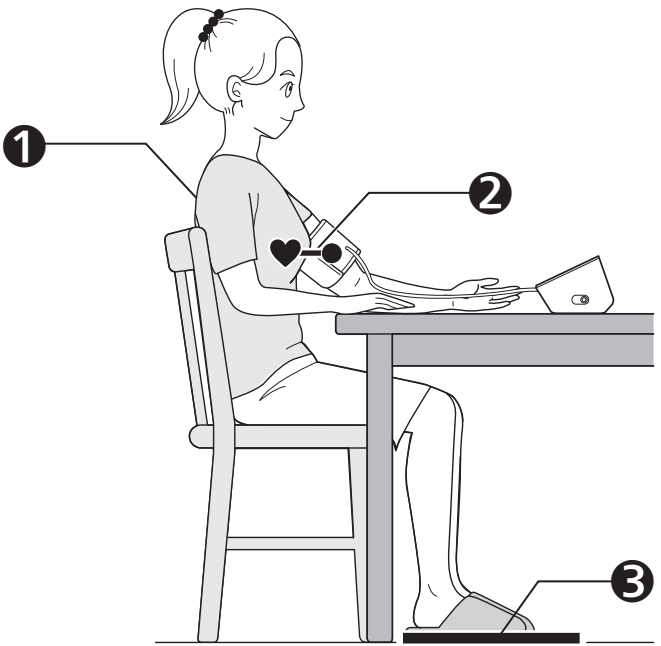
RU Убедитесь, что воздуховодная трубка находится на внутренней поверхности плеча и надежно оберните манжету, чтобы она не прокручивалась вокруг руки.

TR Hava borusunun kolunuzun iç tarafında olduğundan emin olun ve kolluğu kaymayacak şekilde sabit şekilde sarın.

AR تأكد من أن خرطوم الهواء على الجانب الداخلي من ذراعك ولف الشريط الضاغط بإحكام بحيث لا ينزلق حول الذراع.

8 Sitting Correctly

FR Position assise correcte
DE Korrekte Körperhaltung
IT Come sedersi nel modo corretto
ES Cómo sentarse correctamente
NL Correct zitten
RU Сядьте правильно
TR Düzgün Oturma
AR الجلوس بشكل صحيح



❶ Sit with your back and arm supported.

FR S'asseoir de façon à ce que le dos et le bras soient bien soutenus.

DE Mit dem Rücken anlehnen und den Arm auf eine Unterlage legen.

IT Sedere con la schiena e il braccio ben sostenuti.

ES Siéntese de modo que la espalda y el brazo estén bien apoyados.

NL Zit met uw rug en arm ondersteund.

RU Сядьте так, чтобы спина и рука опирались на что-либо.

TR Sırtınız ve kolunuz desteklenecek şekilde oturun.

AR اجلس بحيث يكون الظهر والذراع في وضع استناد.

❷ Place the arm cuff at the same level as your heart.

FR Le brassard doit se trouver au même niveau que votre cœur.

DE Die Manschette auf Herzhöhe platzieren.

IT Posizionare il bracciale allo stesso livello del cuore.

ES Coloque el manguito al mismo nivel que el corazón.

NL Plaats de armmanchet op hetzelfde niveau als uw hart.

RU Манжета должна находиться на уровне сердца.

TR Kolluğu kalbinizle aynı düzeye getirin.

AR ضع الشريط الضاغط للذراع في مستوى القلب نفسه.

❸ Keep feet flat, legs uncrossed, remain still and do not talk.

FR Garder les pieds à plat, les jambes non croisées, ne pas bouger et ne pas parler.

DE Füße flach auf den Boden stellen, die Beine nicht kreuzen, still halten und nicht sprechen.

IT Tenere i piedi ben poggiati, non incrociare le gambe, rimanere fermi e non parlare.

ES Mantenga los pies planos, las piernas sin cruzar, quédese quieto y no hable.

NL Houd de voeten plat, benen niet gekruist, zit stil en praat niet.

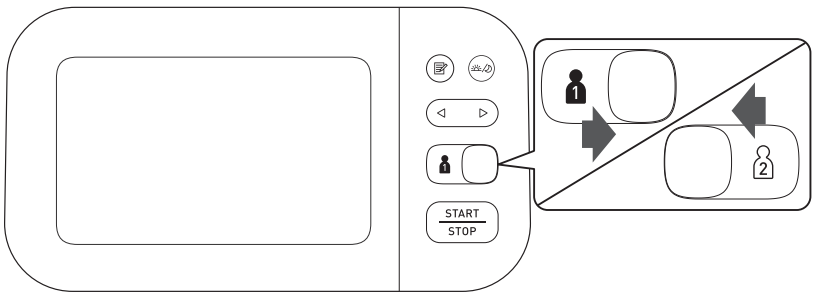
RU Поставьте ноги ровно, не перекрещивая их, не двигайтесь и не разговаривайте.

TR Ayaklarınızı düz, bacaklarınızı açık tutun, hareketsiz kalın ve konuşmayın.

AR اجعل قدميك في وضع مسطّح وساقيك في وضع متباعد وحافظ على ثباتك ولا تتحدث.

9 Selecting User ID (1 or 2)

FR Sélection de l'ID Utilisateur (1 ou 2)
DE Auswahl der Benutzer-ID (1 oder 2)
IT Selezione dell'ID utente (1 o 2)
ES Selección de ID de usuario (1 o 2)
NL Gebruikers-ID selecteren (1 of 2)
RU Выберите идентификатор пользователя (1 или 2)
TR Kullanıcı Kimliğini Seçme (1 veya 2)
AR تحديد هوية المستخدم (١ أو ٢)

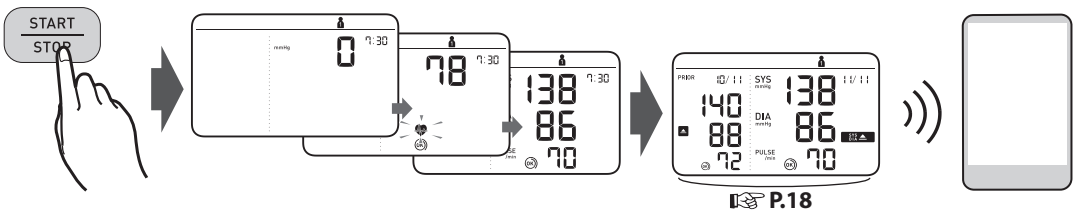


Switching user ID enables you to save readings for 2 people.

FR Changer d'ID Utilisateur permet d'enregistrer les résultats pour 2 personnes.
DE Mit verschiedenen Benutzer-IDs lassen sich Messwerte für 2 Personen speichern.
IT Cambiare ID utente permette di registrare i risultati relativi a 2 persone.
ES Cambiar de ID de usuario le permite guardar las lecturas de 2 personas.
NL Door het wisselen van gebruikers-ID kunt u metingen voor 2 personen opslaan.
RU Переключение идентификатора пользователя позволяет сохранять результаты для 2-х людей.
TR Kullanıcı kimlikleri arasında geçiş yapma, 2 kişi için ölçüm değerlerini kaydetmenizi sağlar.
AR تتيح لك إمكانية تبديل معرف المستخدم حفظ القراءات لشخصين.

10 Taking a Measurement

FR Réalisation d'une mesure
DE Eine Messung vornehmen
IT Misurazione
ES Obtención de una lectura
NL Een meting verrichten
RU Выполнение измерений
TR Bir Ölçüm Yapma
AR إجراء قياس



☞ P.18

When the [START/STOP] button is pressed, the measurement is taken and saved automatically. Open the app to transfer the reading.

FR Après une pression sur le bouton [START/STOP], la mesure démarre et le résultat est enregistré automatiquement. Ouvrir l'application pour transférer le résultat.

DE Beim Drücken der Taste [START/STOP] erfolgt die Messung. Die Messwerte werden automatisch gespeichert. Öffnen Sie die App, um die Messwerte zu übertragen.

IT Quando si preme il pulsante [START/STOP], la misurazione viene eseguita e salvata automaticamente. Aprire la app per trasferire i risultati.

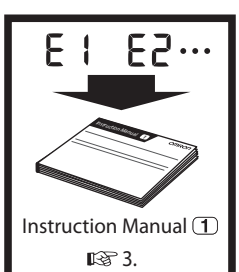
ES Al pulsar el botón [START/STOP], se obtiene la medición, que se guarda automáticamente. Abra la aplicación para transferir la lectura.

NL Wanneer op de knop [START/STOP] wordt gedrukt, wordt de meting automatisch uitgevoerd en opgeslagen. Open de app om de meting te versturen.

RU После того, как кнопка [START/STOP] нажата, измерение выполняется и сохраняется автоматически. Откройте приложение, чтобы передать результаты.

TR [START/STOP] düğmesine basıldığında ölçüm yapılır ve otomatik olarak kaydedilir. Ölçüm değerini aktarmak için uygulamayı açın.

AR عند الضغط على زر [START/STOP]، يتم إجراء القياس وحفظه تلقائيًا. افتح التطبيق لنقل القراءة.



Instruction Manual (1)

☞ 3.

Taking a measurement in Afib mode

FR Réalisation d'une mesure en mode fibrillation auriculaire

DE Vornehmen einer Messung im Afib-Modus

IT Misurazione in modalità Afib

ES Obtención de una lectura en modo fibrilación auricular

NL Een meting in Afib-modus afnemen

RU Выполнение измерений в режиме ФП

TR Afib modunda ölçüm yapma

AR إجراء القياس باستخدام وضع الرجفان الأذيني

In the Afib mode, your monitor automatically takes 3 consecutive readings at 30-second intervals and displays the average. If there is a possibility of Afib, "Afib indicator symbol" ( ) will appear.

This is not a diagnosis, it is only a potential finding for Afib. You should contact your physician to discuss the findings. If you are experiencing any symptoms, contact a medical professional.

FR En mode fibrillation auriculaire, votre moniteur effectue automatiquement 3 mesures consécutives à 30 secondes d'intervalle et affiche la moyenne. S'il y a une possibilité de fibrillation auriculaire, le « symbole indicateur de fibrillation auriculaire » ( ) s'affiche.
Il ne s'agit pas d'un diagnostic mais seulement de l'indication d'une fibrillation auriculaire possible. Vous devriez consulter votre médecin pour discuter des résultats. Si vous ressentez un symptôme quelconque, consultez un professionnel de la santé.

DE Im Afib-Modus nimmt das Messgerät automatisch 3 Messungen hintereinander im Abstand von je 30 Sekunden vor und zeigt den Mittelwert an. Wenn die Möglichkeit von Vorhofflimmern besteht, erscheint das „Afib-Hinweissymbol" ( ).
Dabei handelt es sich nicht um eine Diagnose, sondern lediglich um einen Befund, dass Vorhofflimmern vorliegen könnte. Wenden Sie sich an Ihren Arzt, um den Befund weiter abzuklären. Konsultieren Sie bei Auftreten von Symptomen eine medizinische Fachkraft.

IT Nella modalità Afib, il misuratore esegue automaticamente 3 misurazioni consecutive a intervalli di 30 secondi e visualizza la media. In caso di potenziale presenza di fibrillazione atriale (Afib) viene visualizzato il "simbolo dell'indicatore Afib" ( ).
Questa non è una diagnosi ma è soltanto un rilevamento della potenziale presenza di fibrillazione atriale (Afib). È opportuno rivolgersi al proprio medico curante per discutere questo risultato. Se si dovessero riscontrare eventuali sintomi, rivolgersi a un medico.

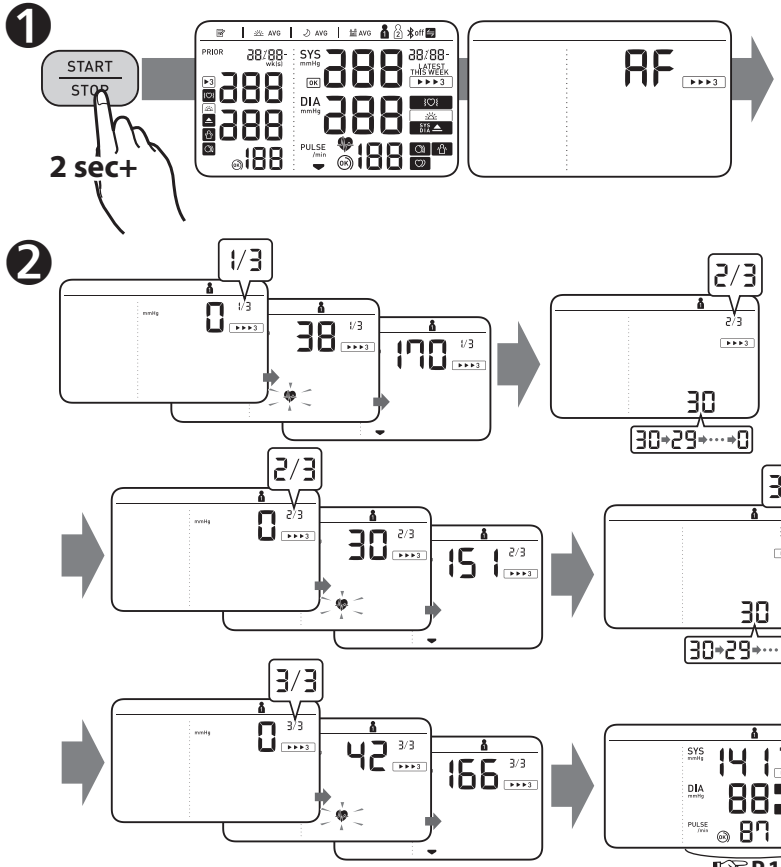
ES En el modo fibrilación auricular, el monitor realiza 3 mediciones consecutivas automáticamente a intervalos de 30 segundos y muestra el valor promedio. Si existe una posible fibrilación auricular, aparecerá el "símbolo indicador de fibrilación auricular" ( ).
Esto no es un diagnóstico, simplemente la detección de una posible fibrilación auricular. Deberá ponerse en contacto con su médico para hablar de los resultados. En caso de padecer síntomas, póngase en contacto con un profesional sanitario.

NL In de Afib-modus neemt uw meter automatisch 3 achtereenvolgende metingen met intervallen van 30 seconden en toont vervolgens het gemiddelde. Als er een mogelijkheid is op Afib verschijnt het "Afib-indicatorsymbool" ( ).
Dit is geen diagnose, het is slechts een vaststelling dat er mogelijk sprake is van Afib. Neem contact op met uw arts om de bevindingen te bespreken. Neem contact op met een medische professional als u symptomen ondervindt.

RU В режиме ФП автоматически выполняется 3 последовательных измерения с 30-секундным интервалом, после чего отображается среднее значение. В случае вероятности ФП отображается «Значок индикатора ФП» ( ). Это не диагноз, а только вероятность обнаружения ФП. Для обсуждения полученных результатов необходимо обратиться к лечащему врачу. Если у Вас существуют какие-либо жалобы на самочувствие, обратитесь к врачу.

TR Cihazınız Afib modunda iken, otomatik olarak 30 saniyelik aralıklarda ardışık 3 ölçüm yapar ve ortalamayı gösterir. Afib olasılığı varsa "Afib gösterge sembolü" ( ) görünür.
Bu bir tanı değildir, yalnızca olası Afib bulgusudur. Doktorunuzla iletişime geçerek bulguları konuşmanız gerekir. Semptom yaşıyorsanız bir tıp uzmanıyla iletişime geçin.

AR في وضع الرجفان الأذيني، يقوم جهاز القياس بالتقاط ٣ قراءات متتابعة تلقائيًا على فترات منتظمة مدتها ٣٠ ثانية ومن ثم يعرض المعدل. إذا كان الرجفان الأذيني أمرًا محتملًا، فسوف يظهر "رمز مؤشر الرجفان الأذيني" ( ).
لا يُعد هذا الإجراء تشخيصًا، فما هو إلا اكتشاف محتمل لحدوث رجفان أذيني. ومن ثمّ، يجب عليك الاتصال بالطبيب لمناقشة النتائج. إذا تعرضت لأي أعراض، فيجب عليك الاتصال بطبيب مختص.



E6

Instruction Manual 1

☞ 3.

Taking a measurement in guest mode

FR Réalisation d'une mesure en mode Invité

DE Vornehmen einer Messung im Gast-Modus

IT Misurazione in modalità Ospite

ES Obtención de una lectura en modo de invitado

NL Een meting in gastmodus afnemen

RU Выполнение измерений в гостевом режиме

TR Konuk modunda ölçüm yapma

AR إجراء القياس باستخدام وضع الضيف.

The guest mode can be used to take a single measurement for another user. No readings are stored in the memory, and the Afib mode is not available when the guest mode is selected.

FR Le mode Invité permet de mesurer la pression artérielle d'une autre personne une seule fois. Les résultats ne sont pas enregistrés dans la mémoire et le mode fibrillation auriculaire n'est pas disponible lorsque le mode Invité est sélectionné.

DE Soll eine einzelne Messung bei einem anderen Benutzer durchgeführt werden, kann dafür der Gast-Modus aktiviert werden. Im Gast-Modus werden keine Messwerte gespeichert und der Afib-Modus steht nicht zur Verfügung.

IT La modalità Ospite può essere utilizzata per eseguire una singola misurazione su un diverso utente. I risultati ottenuti non vengono memorizzati e la modalità Afib non è disponibile quando si seleziona la modalità Ospite.

ES Para realizar una única medición para otro usuario, se puede utilizar el modo de invitado. Cuando se selecciona el modo de invitado, no se guardan lecturas en la memoria y el modo de fibrilación auricular no está disponible.

NL De gastmodus kan worden gebruikt om één enkele meting uit te voeren voor een andere gebruiker. Er worden geen metingen opgeslagen in het geheugen en de Afib-modus is niet beschikbaar wanneer de gastmodus is geselecteerd.

RU Гостевой режим используется для единичного измерения артериального давления у другого лица. Если выбран гостевой режим, то измерения не сохраняются в памяти, и недоступен режим ФП.

TR Konuk modu, başka bir kişi için tek bir ölçüm yapmak üzere kullanılabilir. Konuk modu seçildiğinde, ölçüm değerleri hafızada saklanmaz ve Afib modu kullanılamaz.

AR ويمكن استخدام وضع الضيف لإجراء عملية قياس واحدة لمستخدم آخر. لا يتم تخزين أي قراءة داخل الذاكرة كما أن وضع الرجفان الأذيني لا يكون متاحًا عند تحديد وضع الضيف.

1 **While holding the ▷ button down, press the [START/STOP] button.**

FR Tout en maintenant le bouton ▷ enfoncé, appuyer sur le bouton [START/STOP].

DE Halten Sie die Taste ▷ gedrückt und drücken Sie gleichzeitig die Taste [START/STOP].

IT Mentre si tiene premuto il pulsante ▷, premere il pulsante [START/STOP].

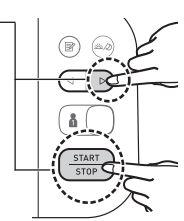
ES Mientras mantiene pulsado el botón ▷, pulse el botón [START/STOP].

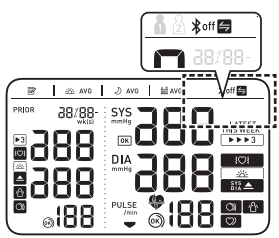
NL Houd de knop ▷ ingedrukt en druk tegelijkertijd op de knop [START/STOP].

RU Удерживая кнопку ▷ в нажатом положении, нажмите кнопку [START/STOP].

TR ▷ düğmesini basılı tutarken, [START/STOP] düğmesine basın.

AR أثناء الاستمرار في الضغط على الزر ▷، اضغط على الزر [START/STOP] (بدء/إيقاف).





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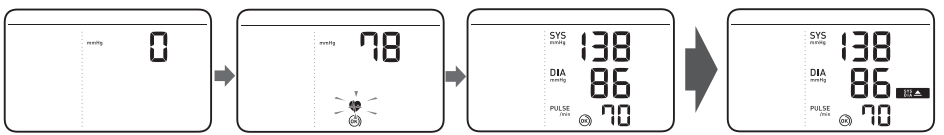


11 Checking Readings in Comparison Mode

FR Vérification des mesures en mode de comparaison

DE Überprüfen von Messwerten im Vergleichsmodus

IT Controllo dei risultati in modalità Confronto

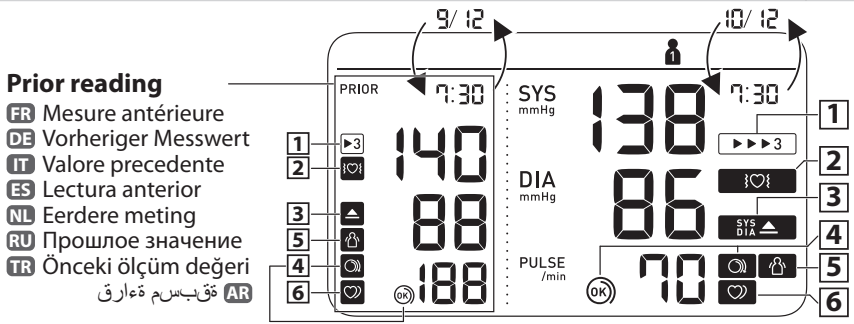
ES Comprobación de las lecturas en modo comparativo

NL Metingen bekijken in vergelijkingsmodus

RU Проверка результатов измерений в режиме сравнения

TR Karşılaştırma Modunda Ölçüm Değerlerini Kontrol Etme

AR التحقق من القراءات في وضع المقارنة



Prior reading

FR Mesure antérieure
DE Vorheriger Messwert
IT Valore precedente
ES Lectura anterior
NL Eerdere meting
RU Прошлое значение
TR Önceki ölçüm değeri
AR ةقباسم ةءارق

1 ▶▶▶3 ▶3

Appears when the reading was taken in Afib mode.

FR S'affiche lorsque la mesure a été effectuée en mode Fibrillation auriculaire.
DE Erscheint, wenn der Messwert im AFib-Modus ermittelt wurde.
IT Appare quando la misurazione è stata effettuata in modalità Afib.
ES Aparece cuando la lectura fue realizada en el modo de fibrilación auricular.
NL Verschijnt wanneer de meting in de Afib-modus werd verricht.
RU Отображается, если значение получено в режиме ФП.
TR Ölçüm Afib moduna alınınca görünür.
AR يظهر عندما يتم الحصول على القراءة في وضع الرجفان الأذيني.

2

Appears if an atrial fibrillation (Afib)* was detected during the Afib mode measurement. If it continues to appear, we recommend you to consult with and follow the directions of your physician.

FR S'affiche si une fibrillation auriculaire* a été détectée durant une mesure en mode fibrillation auriculaire. S'il continue d'apparaître, nous vous recommandons de consulter votre médecin et de suivre ses recommandations.

DE Wird angezeigt, wenn bei einer Messung im Afib-Modus Vorhofflimmern (Afib)* erkannt wurde. Wird das Symbol weiterhin angezeigt, sollten Sie sich an einen Arzt wenden und dessen Anweisungen befolgen.

IT Viene visualizzato se durante una misurazione in modalità Afib è stata rilevata la presenza di fibrillazione atriale (Afib)*. Se il simbolo continua ad apparire, è consigliabile rivolgersi al proprio medico curante e seguirne le indicazioni.

ES Aparece si se detecta fibrilación auricular (Afib)* durante la medición en modo fibrilación auricular. Si esto sigue apareciendo, le recomendamos que consulte a su médico y siga sus recomendaciones.

NL Verschijnt als er een atriumfibrillatie (Afib)* is gedetecteerd tijdens de Afib-modusmeting. Als dit blijft verschijnen raden we u aan contact op te nemen met uw arts en zijn of haar aanwijzingen op te volgen.

RU Отображается, если в режиме измерения ФП была определена фибрилляция предсердий (ФП)*. Если эти нарушения ритма продолжаются, рекомендуем обратиться к лечащему врачу и следовать его указаниям.

TR Afib modunda ölçüm esnasında bir atrial fibrilasyon (Afib)* saptandığında görünür. Görünmeye devam ediyorsa doktorunuza danışmanızı ve kendisinin yönergelerini izlemenizi öneririz.

AR تظهر في حالة اكتشاف الرجفان الأذيني (Afib)* خلال إجراء القياس باستخدام وضع الرجفان الأذيني. وإذا استمرت في الظهور، فإننا نوصيك باستشارة الطبيب الخاص بك واتباع تعليماته.

3

Appears if "SYS" is 135 mmHg or above and/or "DIA" is 85 mmHg** or above.

FR S'affiche si « SYS » est égale ou supérieure à 135 mmHg et/ou « DIA » égale ou supérieure à 85 mmHg**.

DE Wird angezeigt, wenn „SYS" 135 mmHg oder mehr beträgt und/ oder wenn „DIA" 85 mmHg** oder mehr beträgt.

IT Viene visualizzato se la pressione sistolica "SYS" è pari o superiore a 135 mmHg e/o la pressione diastolica "DIA" è pari o superiore a 85 mmHg**.

ES Aparece si "SYS" es 135 mmHg o superior y/o "DIA" es 85 mmHg** o superior.

NL Verschijnt als "SYS" 135 mmHg of hoger is en/of "DIA" 85 mmHg** of hoger is.

RU Отображается, если «SYS» 135 мм.рт.ст. и выше и/или «DIA» 85 мм.рт.ст.** и выше.

TR "SYS" 135 mmHg ya da üstünde olduğunda ve/veya "DIA" 85 mmHg** ya da üstünde olduğunda görünür.

AR تظهر إذا كان "SYS" يبلغ ١٣٥ ملليمترًا زئبقيًا أو أكثر و/ أو "DIA" ٨٥ ملليمترًا زئبقيًا** أو أكثر.

4

Cuff is tight enough.

FR Le brassard est suffisamment serré.

DE Manschette sitzt ausreichend straff.

IT Il bracciale è stretto a sufficienza.

ES El manguito está lo suficientemente prieto.

NL Manchet zit strak genoeg.

RU Манжета затянута достаточно туго.

TR Kolluk yeterince sıkıdır.

AR الشريط الضاغط محكم بما يكفي.

4

Apply cuff again MORE TIGHTLY.

FR Poser le brassard en le serrant davantage.

DE Manschette STRAFFER ziehen.

IT Applicare di nuovo il bracciale STRINGENDOLO DI PIÙ.

ES Vuelva a poner el manguito MÁS PRIETO.

NL Breng de manchet STRAKKER aan.

RU Наложите манжету еще раз БОЛЕЕ ТУГО.

TR Kolluğu tekrar, DAHA SIKI bir şekilde takın.

AR لف الشريط الضاغط مرة أخرى بشكل أكثر إحكامًا.

5 Appears when your body moves during a measurement. Remove the arm cuff, wait 2-3 minutes and try again.

(It does not appear when the possibility of Afib is detected during a Afib mode measurement)

FR S'affiche si vous bougez pendant une mesure. Retirer le brassard, attendre 2 à 3 minutes et essayer à nouveau.
(Ne s'affiche pas si la possibilité d'une fibrillation auriculaire est détectée pendant une mesure en mode fibrillation auriculaire.)

DE Wird angezeigt, wenn Sie sich während der Messung bewegen. Nehmen Sie die Manschette ab, warten Sie 2–3 Minuten und versuchen Sie es erneut.
(Wird nicht angezeigt, wenn bei einer Messung im Afib-Modus mögliches Vorhofflimmern erkannt wird)

IT Viene visualizzato se l'utilizzatore si muove durante la misurazione. Rimuovere il bracciale, attendere 2-3 minuti e riprovare.
(Il simbolo non appare quando viene rilevata una potenziale presenza di fibrillazione atriale durante una misurazione in modalità Afib)

ES Aparece cuando se mueve el cuerpo durante una medición. Retire el manguito, espere unos 2 o 3 minutos e inténtelo de nuevo.
(No aparece cuando se detecta una posible fibrilación auricular durante una medición en modo fibrilación auricular)

NL Verschijnt wanneer uw lichaam tijdens een meting beweegt. Verwijder de armmanchet, wacht 2-3 minuten en probeer het opnieuw.
(Het verschijnt niet wanneer de mogelijkheid op Afib is gedetecteerd tijdens de Afib-modusmeting)

RU Отображается при движении тела во время измерения. Снимите манжету, подождите 2-3 минуты и попробуйте еще раз.
(Не отображается, если при измерении в режиме ФП выявлена вероятность ФП)

TR Bir ölçüm esnasında vücudunuz hareket ettiğinde görünür. Kolluğu çıkarın, 2-3 dakika bekleyip tekrar deneyin.
(Afib modundaki bir ölçümde Afib olasılığı tespit edildiğinde görünmez)

AR تظهر عند تحرك جسمك في أثناء عملية القياس. قم بإزالة الشريط الضاغط للذراع وانتظر من ٢ إلى ٣ دقائق وحاول مرة أخرى.
(لا تظهر عند اكتشاف احتمالية الرجفان الأذيني أثناء القياس باستخدام وضع الرجفان الأذيني)

6 Appears when an irregular rhythm* is detected during a measurement. If it continues to appear, it is recommended to consult your physician.

(It does not appear during a Afib mode measurement)

FR S'affiche lorsqu'un rythme irrégulier* est détecté pendant une mesure. S'il continue d'apparaître, il est recommandé de consulter votre médecin.
(Ne s'affiche pas pendant une mesure en mode fibrillation auriculaire)

DE Wird angezeigt, wenn während einer Messung ein unregelmäßiger Herzschlag* erkannt wird. Wird das Symbol weiterhin angezeigt, sollten Sie sich an Ihren Arzt wenden.
(Wird bei einer Messung im Afib-Modus nicht angezeigt)

IT Viene visualizzato se nel corso di una misurazione viene rilevato un ritmo cardiaco irregolare*. Se il simbolo continua ad apparire è consigliabile rivolgersi al proprio medico curante.
(Il simbolo non viene visualizzato durante le misurazioni in modalità Afib)

ES Aparece cuando se detecta un ritmo irregular* durante una medición. Si esto sigue apareciendo, le recomendamos que consulte a su médico.
(No aparece durante una medición en modo fibrilación auricular)

NL Verschijnt wanneer tijdens een meting een onregelmatig ritme* wordt gedetecteerd. Neem contact op met uw arts als dit blijft verschijnen.
(Het verschijnt niet tijdens een Afib-modusmeting)

RU Отображается, если при измерении определяется нерегулярный ритм*. Если этот значок продолжает отображаться, рекомендуем обратиться к лечащему врачу.
(Не отображается при измерении в режиме ФП)

TR Bir ölçüm esnasında düzensiz ritim* saptandığında görünür. Görünmeye devam ederse doktorunuza danışmanız önerilir.
(Afib modundaki ölçüm esnasında görünmez)

AR تظهر عند اكتشاف عدم انتظام ضربات القلب* أثناء القياس. إذا استمرت في الظهور، يُوصى باستشارة الطبيب المختص.
(لا تظهر أثناء إجراء عملية القياس باستخدام وضع الرجفان الأذيني)

*** Afib and an irregular heartbeat rhythm are defined as a rhythm that is 25 % less or 25 % more than the average rhythm detected while your monitor is measuring blood pressure. The difference between the Afib indicator function and irregular heartbeat function is:**
Afib indicator function: detects Afib possibility in 3-times measurement.
Irregular heartbeat function: detects irregular heartbeat including Afib in 1 measurement.

FR * La fibrillation auriculaire et les pulsations cardiaques irrégulières sont des pulsations dont la fréquence est supérieure ou inférieure de 25 % par rapport à la moyenne détectée lorsque l'appareil mesure la pression artérielle. La différence entre la fonction indicateur de fibrillation auriculaire et la fonction pulsations cardiaques irrégulières est la suivante :
Fonction indicateur de fibrillation auriculaire : détecte la possibilité d'une fibrillation auriculaire par une mesure répétée 3 fois.
Fonction pulsations cardiaques irrégulières : détecte des pulsations cardiaques irrégulières, y compris une fibrillation auriculaire, en 1 mesure.

DE * Vorhofflimmern (Afib) und unregelmäßiger Herzschlag sind definiert als ein Herzrhythmus, der 25 % unter oder 25 % über dem mittleren Herzrhythmus liegt, der während der Blutdruckmessung erkannt wird. Der Unterschied zwischen Afib-Hinweisfunktion und Erkennung unregelmäßiger Herzschläge ist:
Afib-Hinweisfunktion: Erkennt bei 3-maliger Messung möglicherweise vorliegendes Vorhofflimmern.
Erkennung unregelmäßiger Herzschläge: Erkennt unregelmäßigen Herzschlag einschließlich Vorhofflimmern bei 1-maliger Messung.

IT * Per Afib e battito cardiaco irregolare si intende la presenza di variazioni inferiori del 25% o superiori del 25% nel ritmo rispetto al ritmo medio rilevato dall'apparecchio durante la misurazione della pressione arteriosa. La differenza tra la funzione dell'indicatore Afib e la funzione battito cardiaco irregolare è la seguente:
la funzione dell'indicatore Afib rileva la potenziale presenza di fibrillazione atriale mediante 3 misurazioni ripetute.
La funzione battito cardiaco irregolare rileva il battito irregolare inclusa la fibrillazione atriale in 1 misurazione.

ES * La fibrilación auricular y un ritmo de latido cardíaco irregular se definen como aquel ritmo que es el 25% inferior o el 25% superior al ritmo cardíaco medio detectado mientras el monitor mide la presión arterial. La diferencia entre la función de indicador de fibrilación auricular y la función de latido cardíaco arrítmico es:
Función de indicador de fibrilación auricular: detecta una posible fibrilación auricular en 3 mediciones.
Función de latido cardíaco irregular: detecta un latido irregular incluyendo fibrilación auricular en 1 medición.

NL * Afib en onregelmatige hartslag zijn gedefinieerd als een hartritme dat meer dan 25% lager of 25% hoger is dan het gemiddelde hartritme tijdens het meten van de bloeddruk door de meter. Het verschil tussen de Afib-indicatorfunctie en onregelmatige hartslagfunctie is: Afib-indicatorfunctie: detecteert de mogelijkheid op Afib in een driemalige meting.
Onregelmatige hartslagfunctie: detecteert onregelmatige hartslag inclusief Afib in één meting.

RU * ФП и нерегулярный ритм сердцебиения — это ритм, который на 25 % медленнее или на 25 % быстрее среднего ритма, определенного прибором при измерении артериального давления. Различия между функцией индикатора ФП и функцией нерегулярного сердцебиения:
Функция индикатора ФП: определяет вероятность ФП при трехкратном измерении.
Функция нерегулярного сердцебиения: определяет нерегулярные сердечные сокращения, включая ФП, при однократном измерении.

TR * Afib ve düzensiz kalp atışı ritmi, ölçüm cihazınız tansiyon ölçümü yaparken saptanan ortalama ritme göre %25'ten az ya da %25'ten fazla sapma gösteren ritim olarak tanımlanır. Afib göstergesi fonksiyonu ve düzensiz kalp atışı fonksiyonu arasındaki fark şu şekildedir:
Afib göstergesi fonksiyonu: 3 kez yapılan ölçümde Afib olasılığını tespit eder.
Düzensiz kalp atışı fonksiyonu: 1 ölçümde, Afib de dahil olmak üzere düzensiz kalp atışını tespit eder.

AR * يعبر كل من الرجفان الأذيني واضطراب ضربات القلب عن إيقاع يقل بنسبة ٢٥٪ أو يزيد بنسبة ٢٥٪ عن الإيقاع المتوسط الذي يتم اكتشافه أثناء قياس الجهاز لضغط الدم. الاختلاف بين وظيفة مؤشر الرجفان الأذيني ووظيفة اضطراب ضربات القلب هو أن:
وظيفة مؤشر الرجفان الأذيني: تكتشف احتمالية حدوث رجفان أذيني عند إجراء عمليات القياس ٣ مرات.
وظيفة اضطراب ضربات القلب: تشير إلى ضربات القلب غير المنتظمة بما في ذلك الرجفان الأذيني وذلك عند إجراء عملية قياس واحدة.

**** The high blood pressure definition is based on the 2018 ESH/ESC Guidelines.**

FR ** La définition de l'hypertension est basée sur les recommandations ESH/ESC 2018.

DE ** Die Definition für Bluthochdruck basiert auf den 2018 ESH/ESC-Richtlinien.

IT ** La definizione di alta pressione arteriosa si basa sulle linee guida 2018 di ESH/ESC.

ES ** La definición de presión arterial alta se basa en las guías 2018 ESH/ESC.

NL ** De definitie van hoge bloeddruk is gebaseerd op de ESH/ESC-richtlijnen uit 2018.

RU ** Определение высокого артериального давления основано на Рекомендациях 2018 ESH/ESC.

TR ** Yüksek tansiyon tanımı, 2018 ESH/ESC Kılavuzlarını temel alır.

AR ** تعريف ضغط الدم المرتفع يستند إلى إرشادات الجمعية الأوروبية لفرط ضغط الدم (ESH)/ الجمعية الأوروبية لطب القلب (ESC) لعام ٢٠١٨.

E1
E2

Error messages or other problems? Refer to:

FR Messages d'erreur ou autres problèmes ? Voir :

DE Weitere Fehlermeldungen oder Probleme siehe:

IT Messaggi di errore o altri problemi? Fare riferimento a:

ES ¿Hay mensajes de error u otros problemas? Consulte:

NL Foutmeldingen of andere problemen? Raadpleeg:

RU Сообщения об ошибках или другие неисправности? См.

TR Hata mesajları veya başka sorunlar mı var? Bkz:

AR رسائل الخطأ أو المشكلات الأخرى؟ ارجع إلى:

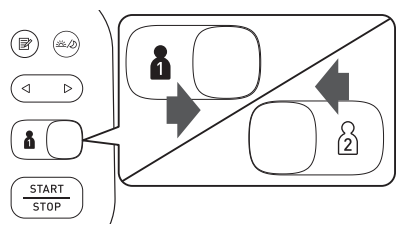
Instruction Manual 1
☞ 3.

12 Using Memory Functions

FR Utilisation des fonctions de mémoire
DE Verwendung der Speicherfunktion
IT Uso delle funzioni di memoria
ES Uso de las funciones de memoria
NL Geheugenfuncties gebruiken
RU Использование функции памяти
TR Hafıza Fonksiyonunun Kullanılması
AR استخدام وظائف الذاكرة

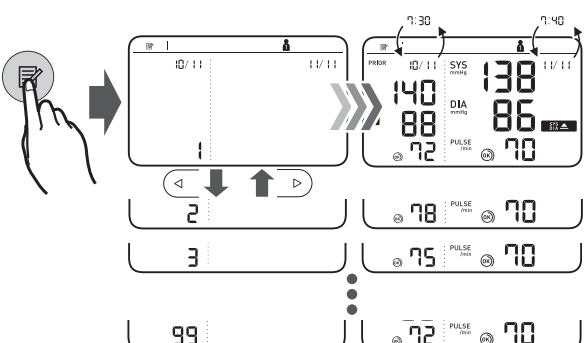
Before using memory functions, select your user ID.

FR Avant d'utiliser les fonctions de mémoire, sélectionner votre ID Utilisateur.
DE Wählen Sie vor dem Verwenden der Speicherfunktionen Ihre Benutzer-ID aus.
IT Prima di utilizzare le funzioni di memoria selezionare il proprio ID utente.
ES Antes de usar las funciones de memoria, seleccione su ID de usuario.
NL Selecteer uw gebruikers-ID voordat u geheugenfuncties gebruikt.
RU Перед использованием функции памяти выберите идентификатор пользователя.
TR Hafıza fonksiyonunu kullanmadan önce kullanıcı kimliğinizi seçin.
AR قبل استخدام وظائف الذاكرة، حدد معرف المستخدم الخاص بك.



12.1 Readings Stored in Memory

FR Mesures stockées en mémoire
DE Gespeicherte Messungen
IT Risultati conservati in memoria
ES Lecturas guardadas en la memoria
NL Meetwaarden opgeslagen in het geheugen
RU Сохранение результатов в памяти
TR Hafızada Saklanan Ölçüm Değerleri
AR القراءات المخزنة في الذاكرة



Stores up to 100 readings.

FR Mémorise jusqu'à 100 résultats.
DE Es werden bis zu 100 Messwerte gespeichert.
IT Conserva fino a 100 risultati.
ES Almacena hasta 100 lecturas.
NL Slaat tot maximaal 100 metingen op.
RU Сохраняется до 100 результатов.
TR 100 adede kadar ölçüm değeri saklar.
AR تقوم بتخزين ما يصل إلى ١٠٠ قراءة.

Morning/Evening Weekly Averages

FR Moyennes hebdomadaires matin/soir

DE Wöchentliche Morgen-/Abendmittelwerte

IT Media settimanale mattutina e serale

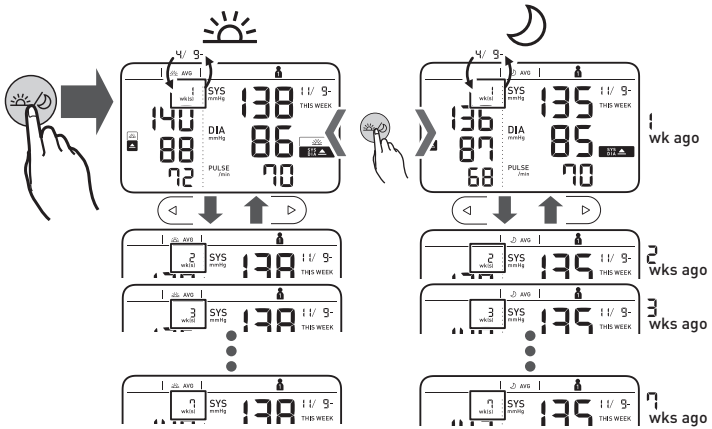
ES Valor promedio semanal de mañana y noche

NL Weekgemiddelden van ochtend- en avondwaarden

RU Утренние/вечерние средние значения за неделю

TR Sabah/Akşam Haftalık Ortalamaları

AR المعدلات الأسبوعية الصباحية/المسائية لضغط الدم



Appears if "SYS" is 135 mmHg or above and/or "DIA" is 85 mmHg or above in the morning weekly average.

FR S'affiche si « SYS » est égale ou supérieure à 135 mmHg et/ou « DIA » égale ou supérieure à 85 mmHg dans la moyenne hebdomadaire du matin.

DE Wird angezeigt, wenn im wöchentlichen Morgenmittelwert „SYS" 135 mmHg oder mehr und/oder „DIA" 85 mmHg oder mehr beträgt.

IT Viene visualizzato se la pressione sistolica "SYS" è pari o superiore a 135 mmHg e/o la pressione diastolica "DIA" è pari o superiore a 85 mmHg nella media mattutina settimanale.

ES Aparece si "SYS" es 135 mmHg o superior y/o "DIA" es 85 mmHg o superior en el promedio semanal de mañana.

NL Verschijnt als "SYS" 135 mmHg of hoger is en/of "DIA" 85 mmHg of hoger is in het ochtendweekgemiddelde.

RU Отображается, если средние утренние значения за неделю «SYS» 135 мм.рт.ст. и выше и/или «DIA» 85 мм.рт.ст. и выше.

TR Sabah haftalık ortalamasında "SYS" 135 mmHg ya da üstünde olduğunda ve/veya "DIA" 85 mmHg** ya da üstünde olduğunda görünür.

AR تظهر إذا كان "SYS" يبلغ ١٣٥ ملليمترًا زئبقيًا أو أكثر و/ أو "DIA" ٨٥ ملليمترًا زئبقيًا أو أكثر في المعدل الأسبوعي الصباحي.

To know how to calculate weekly averages, refer to section 10 of the Instruction Manual (1).

FR Pour savoir comment calculer les moyennes hebdomadaires, se reporter à la section 10 du Mode d'emploi (1).

DE Wie die wöchentlichen Mittelwerte berechnet werden, ist in Abschnitt 10 der Gebrauchsanweisung (1) beschrieben.

IT Per sapere come calcolare le medie settimanali, consultare la sezione 10 del manuale di istruzioni (1).

ES Para saber cómo calcular mediciones semanales, consulte la sección 10 del manual de instrucciones (1).

NL Zie paragraaf 10 van de gebruiksaanwijzing (1) voor aanwijzingen hoe u het wekelijkse gemiddelde kunt berekenen.

RU О том, как рассчитываются средние значения за неделю, см. раздел 10 этого руководства по эксплуатации (1).

TR Haftalık ortalamaların nasıl hesaplanacağını öğrenmek için, Kullanım Kılavuzu (1)'de bölüm 10'a bakın.

AR لمعرفة كيفية حساب المعدلات الأسبوعية، ارجع إلى القسم ١٠ من دليل الإرشادات (1).

12.3 Average of the Latest 2 or 3 Readings Taken within a 10 Minute Span

FR Moyenne des 2 ou 3 dernières mesures prises en l'espace de 10 minutes

DE Mittelwert der letzten 2 oder 3 Messwerte, die in einem 10-Minuten-Zeitrahmen erfasst wurden

IT Media degli ultimi 2 o 3 risultati ottenuti nell'arco di 10 minuti

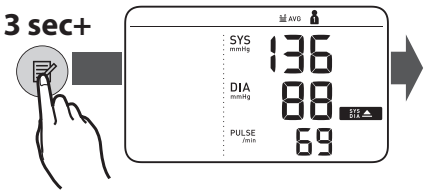
ES Promedio de las 2 o 3 últimas lecturas realizadas en un intervalo de 10 minutos

NL Gemiddelde van de laatste 2 of 3 metingen uitgevoerd binnen een tijdspanne van 10 minuten

RU Среднее значение последних 2-х или 3-х результатов, полученных в течение 10 минут

TR 10 Dakika Aralıkta Alınan Son 2 veya 3 Ölçüm Değerinin Ortalaması

AR معدل آخر قراءتين أو ٣ قراءات مسجلة خلال مدة زمنية تبلغ ١٠ دقائق



12.4 Deleting All Readings for 1 User

FR Suppression de toutes les mesures pour 1 utilisateur

DE Löschen aller Messwerte für Benutzer 1

IT Eliminazione di tutti i risultati relativi a 1 utente

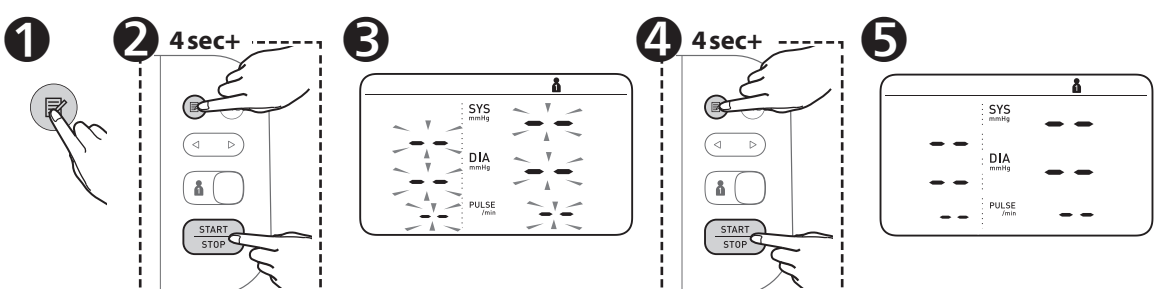
ES Eliminación de todas las lecturas de un usuario

NL Alle meetresultaten voor 1 gebruiker wissen

RU Удаление всех измерений одного пользователя

TR 1 Kullanıcı için Tüm Ölçüm Değerlerini Silme

AR مسح جميع القراءات الخاصة بمستخدم واحد

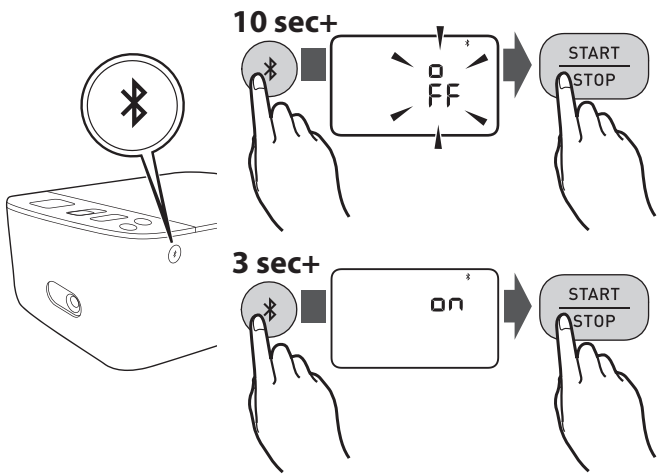


13 Other Settings

FR Autres réglages
DE Weitere Einstellungen
IT Altre impostazioni
ES Otros ajustes
NL Andere instellingen
RU Другие настройки
TR Diğer Ayarlar
AR الإعدادات الأخرى

13.1 Disabling/Enabling Bluetooth

FR Désactivation/activation de Bluetooth
DE Deaktivieren/Aktivieren der Bluetooth-Funktion
IT Disattivazione/attivazione della funzione Bluetooth
ES Activar o desactivar el Bluetooth
NL Bluetooth uitschakelen/inschakelen
RU Выключение/включение Bluetooth
TR Bluetooth'u Devre Dışı Bırakma/Etkinleştirme
AR تمكين/ تعطيل خاصية Bluetooth

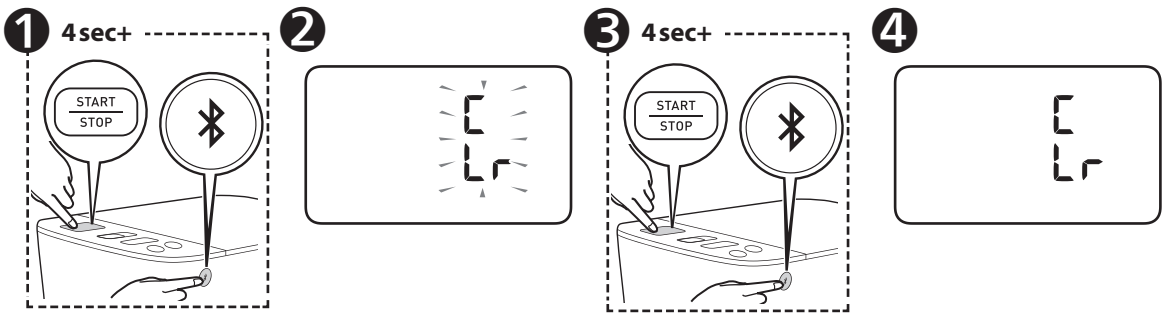


Bluetooth is enabled by default.

FR Bluetooth est activé par défaut.
DE Die Bluetooth-Funktion ist standardmäßig aktiviert.
IT La funzione Bluetooth è attiva per impostazione predefinita.
ES El Bluetooth está activado por defecto.
NL Bluetooth is standaard ingeschakeld.
RU Bluetooth включен по умолчанию.
TR Bluetooth varsayılan olarak etkindir.
AR يتم تمكين خاصية Bluetooth بشكل افتراضي.

13.2 Restoring to the Default Settings

FR Réinitialisation aux réglages par défaut
DE Wiederherstellen der Standardeinstellungen
IT Ripristino delle impostazioni predefinite
ES Restablecimiento a los ajustes de fábrica
NL De standaardinstellingen herstellen
RU Восстановление настроек по умолчанию
TR Varsayılan Ayarları Geri Yükleme
AR استعادة الإعدادات الافتراضية



If your systolic pressure is more than 210 mmHg:

After the arm cuff starts to inflate, press and hold the [START/STOP] button until the monitor inflates 30 to 40 mmHg higher than your expected systolic pressure.

FR Si votre pression systolique est supérieure à 210 mmHg :
Lorsque le brassard commence à se gonfler, appuyer sur le bouton [START/STOP] et le maintenir enfoncé jusqu'à ce que le tensiomètre atteigne une pression de gonflage supérieure de 30 à 40 mmHg à votre pression systolique attendue.

DE Nachdem die Manschette begonnen hat, sich aufzupumpen, drücken Sie die [START/STOP]-Taste, und halten Sie sie gedrückt, bis ein Druck von 30 bis 40 mmHg über Ihrem erwarteten systolischen Druck erreicht ist.

IT Se la pressione sistolica è superiore a 210 mmHg:
Quando il bracciale inizia a gonfiarsi, premere e mantenere premuto il pulsante [START/STOP] finché il misuratore non raggiunge una pressione da 30 a 40 mmHg superiore rispetto al valore di pressione sistolica atteso.

ES En caso de que su presión arterial sistólica esté por encima de 210 mmHg:
Una vez que el manguito comience a inflarse, mantenga pulsado el botón [START/STOP] hasta que el monitor indique que el inflado está entre 30 y 40 mmHg por encima de la presión arterial sistólica estimada.

NL Als uw systolische druk hoger is dan 210 mmHg:
Zodra de armmanchet wordt opgepompt, houdt u de knop [START/STOP] ingedrukt totdat de meter 30 tot 40 mmHg hoger aangeeft dan uw verwachte systolische druk.

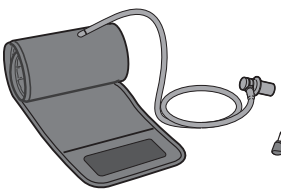
RU После того, как началось автоматическое наполнение манжеты воздухом, нажмите и удерживайте кнопку [START/STOP] до тех пор, пока прибор не поднимет давление до значения, превышающего ожидаемое систолическое давление на 30-40 мм.рт.ст.

TR Sistolik basıncınız 210 mmHg'den fazlaysa:
Manşet şişmeye başladıktan sonra, [START/STOP] düğmesine basın ve ölçüm cihazı beklediğiniz sistolik basınç değerinden 30 ila 40 mmHg daha fazla şişene kadar basılı utun.

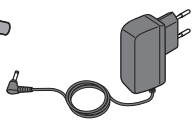
AR إذا كان ضغط الدم الانقباضي أكثر من ٢١٠ ملليمترات زئبقية:
بعد بدء انتفاخ الشريط الضاغط للذراع، اضغط مع الاستمرار على الزر [START/STOP] حتى يقوم الجهاز بالنفخ من ٣٠ إلى ٤٠ ملليمترًا زئبقيًا أعلى من قيمة ضغط الدم الانقباضي المتوقعة.

14 Optional Medical Accessories

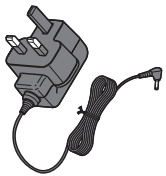
FR Accessoires médicaux en option
DE Medizinisches optionales Zubehör
IT Accessori medicali opzionali
ES Accesorios médicos opcionales
NL Optionele medische accessoires
RU Дополнительно принадлежности
TR Opsiyonel Tıbbi Aksesuarlar
AR الملحقات الطبية الاختيارية



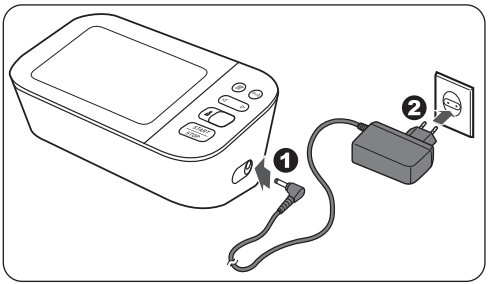
Arm Cuff
(HEM-FL31)
22 - 42 cm



AC Adapter
(HHP-CM01)



(HHP-BFH01)

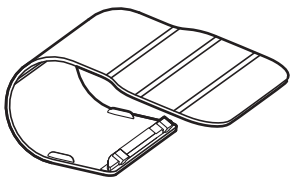


Do not throw the air plug away. The air plug can be applicable to the optional cuff.

FR Ne pas jeter la prise de gonflage. La prise de gonflage peut être utilisée pour le brassard en option.
DE Entsorgen Sie den Luftschlauchstecker nicht. Der Luftschlauchstecker wird für die optionale Manschette verwendet.
IT Non gettare via l'attacco del tubo dell'aria. L'attacco del tubo dell'aria può essere applicato al bracciale opzionale.
ES No tire el conector para tubo de aire. El conector para tubo de aire puede ser utilizado con el manguito opcional.
NL Gooi de plug van de luchtslang niet weg. De plug van de luchtslang kan worden gebruikt op de optionele manchet.
RU Не выбрасывайте воздушный штекер. Он может подойти к дополнительной манжете.
TR Hava tıpasını atmayın. Hava tıpası isteğe bağlı kolluğa uygulanabilir.
AR لا تتخلص من سدادة الهواء. يمكن استعمال سدادة الهواء مع الشريط الضاغط الاختياري.

15 Other Optional Parts

FR Autres pièces en option
DE Weitere optionale Teile
IT Altri componenti opzionali
ES Otras piezas opcionales
NL Overige optionele onderdelen
RU Другие дополнительно приобретаемые запасные части
TR Diğer Aksesuarlar
AR أجزاء اختيارية أخرى



Protective LCD Cover
(HEM-CACO-734)

https://www.omron-healthcare.com/

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Made in Vietnam / Fabriqué en Vietnam / Hergestellt in Vietnam / Prodotto in Vietnam / Fabricado en Vietnam / Geproduceerd in Vietnam / Сделано во Вьетнаме / Vietnam'da Üretilmiştir / صنع في فيتنام

Issue Date / Date de publication / Ausgabedatum / Data di pubblicazione / Fecha de publicación / Uitgiftedatum / Дата выпуска / Teslim Tarihi / تاريخ الإصدار: 2019-09-05

IM2-HEM-7361T-E-02-09/2019
2895865-4B