



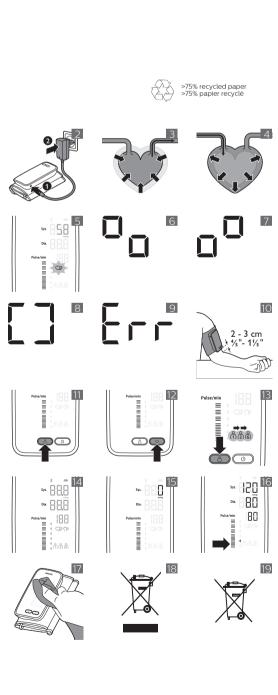
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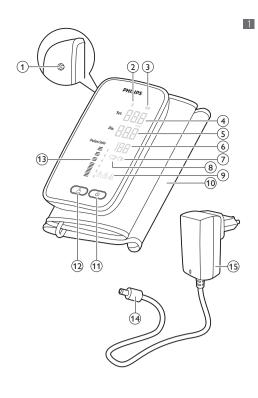
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#### Introduction

Congratulations on your purchase and welcome to Philips! To fully benefit from the support that Philips offers, register your product at

### www.philips.com/welcome

#### General

The Philips upper arm blood pressure monitor with Bluetooth® enables you to perform blood pressure measurements, heart rate (pulse) measurements transmit data via Bluetooth® to your mobile device and display your personal measurement results in the Philips HealthSuite health app. The device can also be used as a standalone device.

This user manual contains important safety information and provides step-by-step instructions for using the blood pressure monitor.

Read this information carefully before you use the device and save it for future reference.

Features:

- 86.5mm×24mm display with white backlight
- Measure-during-inflation technology
- Supports 2 users

### Intended use

The Philips upper arm blood pressure monitor is a digital monitor intended for use in measuring blood pressure and heart rate in adult patient population with arm circumference ranging from 22cm to 42cm (about 9-17 inches). The device is intended to be used in a home environment.

The warning signs and symbols are essential to ensure that you use this product safely and correctly and to protect you and others from injury. Below you find the meaning of the warning signs and symbols on the label and in the user manual

1 Symbol for 'follow instructions for use'.

沈 This symbol means that the part of the device that comes into physical contact with the user (also known as the applied part) is of type BF (Body Floating) according to IEC 60601-1. The applied part is the cuff.

Symbol for 'the device complies with European Medical Device Directive 93/42/EEC requirements'. 0344 refers to **C**€ 0344

the notified body 凤 Symbol for WEEE, waste electrical and electronical equipment. Electrical waste products should not be disposed of with nousehold waste. Please recycle where facilities exist. Check with your local authority or retailer for recycling advice and see chapter 'Battery recycling'.

凉 This symbol means that this product contains batteries which shall not be disposed of with normal household waste (2006/66/EC).

Indicates the manufacturer, as defined in EU Directives 93/42/EEC.

Symbol for 'direct current' Symbol for the 'Bluetooth Combination The device uses Bluetooth for

Indicates manufacturing date.

communication.

LOT Indicates the manufacturer's batch code. SN Indicates the manufacturer's serial number so that a specific medical device

Fuse T1A/250V Φ3.6\*10CCC

can be identified.

Symbol for 'Class II Equipment'. The adapter is double insulated (Class II) and complies with IEC 60601-1.

Symbol for indoor use only.



W

Symbol for 'Including RF transmitter' means that this device emits non-ionizing radiation. All devices with RF transmitters or that use RF electromagnetic energy must have a label with this symbol.



Indicates caution. The user should consult the instructions for use for important cautionary information such as warnings and precautions that cannot, for a variety be presented on the medical of reasons,



IP22

device itself This symbol on the device means protected against access to hazardous parts with a finger and against vertically



falling water drops when tilted up to 15 degrees. Indicates the storage and transportation temperature limits to which the medical device can be safely exposed: -20°C to



Symbol for the 2 year Philips guarantee.



The Green Dot ('Der Grüne Punkt' in German) is the license symbol of a European network of industry-funded systems for recycling the packaging materials of consumer goods.



This symbol on the DC charger indicates that it is TUV certified.



General description (Fig. 1)

## Socket for DC charger plug Bluetooth® symbol 2

- Battery symbol Systolic blood pressure
- 4
- 5 Diastolic blood pressure
- 6 Heart rate
- 7 Movement detector Heart rate/irregular heart rate detector 8
- 9 User IDs 10 Cuff
- On button 11
- User ID button 12

Important safety information

- 13 Blood pressure classification
- DC plug DC charger 14 15

Read this important information carefully before you use the device and save it for future reference.

# Warning



- children or pets, since inhalation or swallowing of small parts can be dangerous or even fatal. The device is not suitable for measuring the
- blood pressure of children.
- The device is not suitable for persons who have electrical implants. Do not use this blood pressure monitor on any
- arm where intravascular access or therapy (such as an intravenous drip or a blood transfusion), or an arterio-venous shunt (A-V shunt) is present. The temporary interference to
- blood flow by the blood pressure measurement could result in injury. If you had a mastectomy (breast amputation) do not use this blood pressure monitor on the arm on the side of the mastectomy. The inflating cuff can lead to pain, trauma and
- further injury in the arm on the side of the mastectomy. Consult your doctor if your suffer from illnesses prior to using the device. No modifications of this equipment are allowed.
- This may result in increased emissions or decreased immunity of the device. Do not use the blood pressure monitor during
- charging as this can cause injury. Do not touch the output of the adapter as this can cause injury.
- Do not dispose of batteries in fire. Battery may explode or leak.
- If you experience discomfort during a measurement, such as pain in the arm or other
- complaints, press the 'on' button to release the air immediately from the cuff. Loosen the cuff
- and remove it from your arm. On the rare occasion of a fault causing the cuff to remain fully inflated during measurement, open the cuff immediately. Prolonged high pressure (cuff pressure >300mmHg or constant pressure >15mmHg for more than 3 minutes)
- (ecchymosis). Too frequent and consecutive measurements could cause disturbances in blood circulation injuries and

applied to the arm, may lead to bruises

- Beware of strangulation, particularly for children and infants due to cables.
  - Do not connect the tube to other medical equipment, as this could lead to dangerous iniuries.
  - This device is not intended for patients outside a home environment.
- Never use any accessories or parts from other manufacturers or that Philips does not specifically recommend. Using such accessories or parts could cause a hazardous situation for
- the user or damage to the device. Only use the accessories and detachable parts authorized by the manufacturer. The use of unauthorized parts or accessories may cause damage to the device or injury to the user.

# Caution

- Always check the device before you use it. Do not use the device if it is damaged, as this may cause injury. The effectiveness of this blood pressure
- monitor has not been established in pregnant (including pre-eclamptic) women.

and peripheral artery disease / arteriosclerosis can affect the performance (accuracy) of this blood pressure monitor. Please consult your doctor how to best use this blood pressure monitor if you suffer from any of these conditions. Only use this device for its intended purpose as described in this user manual. Do not confuse self-monitoring with self-

Common arrhythmias (such as atrial or ventricular premature beats or atrial fibrillation)

- diagnosis. This device allows you to monitor your blood pressure. Do not begin or end medical treatment based on the measurement results. Always consult your doctor for treatment advice.
- Do not take any therapeutic measures on the basis of a self-measurement. Never change prescribed medication without consulting your doctor. Consult your doctor if you have any questions about your blood pressure.

  If you are taking medication, consult your
- physician to determine the most appropriate time to measure your blood pressure. This device is not intended for use on extremities other than the arm or for functions
- other than obtaining a blood pressure measurement.
- If the cuff pressure exceeds 40kPa (300mmHg), the unit will deflate automatically. If the cuff does not deflate when pressures exceeds
- 40kPa (300mmHg), detach the cuff from the arm and press the 'on' button to stop inflation. Do not attach the cuff on the same arm on
- which other monitoring medical electrical
- equipment is attached simultaneously, because this could cause temporary loss of function of those simultaneously-used monitoring medical
- electrical equipment. Never attach the cuff on injured skin, an injured arm or an arm under medical treatment as this
- can cause further injury.
  Do not use the device in case of existing polyester or nylon material allergies. This device is not washable. Never immerse the device in water and do not rinse it under the tap. This device is not suitable for continuous
- monitoring during medical emergencies or operations. This device cannot be used with HF (High Frequency) surgical equipment at the same
- Never use compressed air, scouring pads, abrasive cleaning agents or aggressive liquids such as petrol or acetone to clean the device
- Do not use the adapter in or near wall sockets that contain or have contained an electric air freshener to prevent irreparable damage to the
- adapter. If the battery can no longer be recharged or when the device does not function properly anymore (see 'Specifications'), please contact
- the Philips Consumer Care center in your country. Keep the device away from fire and heat
  - sources, as the battery can overheat, causing fire or bursting. The battery could explode causing injury or death.
    After charging, remove the small plug from the device and remove the adapter from the wall
  - socket. The equipment is not AP/APG equipment and is not suitable for use in the presence of a flammable anesthetic mixture with air, with oxygen or nitrous oxide.
  - To avoid measurement errors, do not use the device near strong electrical or magnetic fields, for example magnets, radio transmitters, microwave ovens.
  - To avoid measurement errors, do not use the device near a strong electromagnetic field radiated interference signal or electrical fast
  - transient/burst signal. Use this device under the right environmental conditions as indicated in this user manual. If
  - not, this could affect the performance, lifetime of the device and measurement results.
    Use a soft cloth to clean the whole unit. Do not use any abrasive or volatile cleaners.
    Only use the DC charger supplied (model: AC power adapter KH0601000EW) to charge the
- device. The device does not need to be calibrated for two years of normal use. After two years the measurements may become less accurate.
- If you have any problems with this device, such as setting up, malfunction, maintaining or using, please contact Philips Consumer Care Do not open, disassemble or repair the device yourself. Modification of the device voids your

Report to Philips Consumer Care if any unexpected operation or events occur Dispose of accessories, detachable parts, and

the ME equipment according to the local guidelines Do not attempt to replace your blood pressure monitor's battery. It is built-in and not changeable.

warrantv.

Do not clean the blood pressure monitor when it is being charged. Always unplug the charger first before cleaning the blood pressure monitor

'Specifications').

60601-1-2

- Compliance with standards The device meets the relevant standards for this for home use.
  - type of Class IIa electrical medical equipment This Philips device complies with all applicable standards and regulations regarding exposure to electromagnetic fields and complies with EN

Avoid charging your blood pressure monitor in extremely high or low temperatures (see

Sy- m- bol	Description	Explanantion
Sys.	Systolic blood pressure	Maximum blood pressure, see also section systolic and diastolic pressure.
Dia.	Diastolic blood pressure	Minimum blood pressure, see also section systolic and diastolic pressure.
Pulse/min	Heart rate	Number of heartbeats per minute (pulse is typically equivalent to heart rate).
	Battery status	Indicates status of battery during charging.
mmHg	Measure- ment unit	Measurement unit of blood pressure.
(C)))	Irregular heart rate detector	Irregular heart rate detection during the measurement.
ளீவீ®்	User IDs	Start measurement for selected user, and transmit the measurement result.
יול")	Movement detector	Moving during the measurement will result in an inaccurate result.
	Blood pressure classifica- tion	Classification of measured blood pressure following WHO system (see 'Blood pressure classification').

### Battery status indications

Bluetooth® symbol

Heart rate

detection

Battery symbol	Battery status	
	The battery is almost empty.	
□+L[]	The battery is empty.	
When you measure 3 times a day starting with a		

The device uses Bluetooth

Heart rate detection during

for communication.

the measurement.

fully charged battery, the device can be used for about 20 days until a recharge is needed. In case of normal use, the battery can be charged around 300 times

Note: Data will be lost when the battery is completely empty.

# Charging

**Battery** 

\*

The battery of this device is a built-in rechargeable li-polymer battery with a capacity of 1000 mAh. Use the original DC charger supplied to charge the battery When the battery is empty, it takes approx. 2 hours

- to fully charge the battery of the device.

  1 Put the small plug in the socket of the device (Fig. 2).
- 2 Put the adapter in the wall socket.
- Battery charging indications **Battery charging indication**

symbol	
	Battery charging: half full
	Battery charging: almost full

**\_\_\_\_** Battery fully charged

### Using the blood pressure monitor This tubeless device uses the oscillometric method

to measure blood pressure and heart rate. Before every measurement, the unit establishes a "zero point" equivalent to the atmospheric pressure. Then it starts inflating the cuff. During the measurement, the device detects the pressure oscillations in the blood vessels generated by the heart pumping blood through the body. These pressure oscillations are used to determine systolic and diastolic blood pressure as well as heart rate. While measuring heart rate, the device also determines the small variations between the individual heartbeats. If these variations exceed a pre-defined threshold, the irregular heart rate detector symbol lights up. Systolic and diastolic pressure

The heart consists of two large chambers - the ventricles - and two smaller chambers - the atria. The ventricles collect blood from the atria and expel it towards the peripheral beds of blood vessels within the body and the lungs. The atria collect blood from these peripheral beds and prime the ventricles When the ventricles contract and pump blood out

of the heart, the blood pressure reaches its maximum value in the cycle, which is called systolic pressure (Fig. 3).
When the ventricles relax and are filled again with

blood, the blood pressure reaches its minimum value in the cycle, which is called diastolic pressure (Fig. 4).

#### Blood pressure classification Consult a doctor in case of questions about your

blood pressure. Your doctor can inform you: About your normal blood pressure range

- If your measuring result falls out of the range.
- Whether your blood pressure has reached a dangerous level.

The following table shows the classification system for the blood pressure measurements used in this device. This system follows the classification system of the World Health Organisation (WHO).

### Blood pressure classification according to WHO system

Systolic pressure mmHg	Diastolic pressure mmHg		Blood pressure indicator	
≥180	≥110	severe hypertension	red	
160 - 179	100 - 109	moderate hypertension	orange	
140 - 159	90 - 99	mild hypertension	yellow	
130 - 139	85 - 89	high to normal blood pressure	green	
120 - 129	80 - 84	normal blood pressure	green	
< 120	< 80	optimal blood pressure	green	
< 100	< 60	low blood pressure	green	
*Source: Chalmers J et al. WHO-ISH Hypertension				

Guidelines Committee, 1999 World Health Organization - International Society of Hypertension Guidelines for the Management of Hypertension. J Hypertens, 1999, 17:151-185.

#### Irregular heart rate detector The device is equipped with an irregular heart rate

detector. An irregular heart rate is detected when the heart rhythm varies above a pre-defined level while the device is measuring the systolic and diastolic blood pressure. During each measurement, this device records the heartbeat intervals and calculates the standard deviation. If the standard deviation exceeds a pre-defined threshold, the irregular heart rate detector symbol lights up when the measurement results are displayed (Fig. 5).

Caution:The appearance of the irregular heart rate

detector symbol indicates that a heart rate irregularity was detected during measurement Usually this is not a cause for concern. Due to the irregularity in your heart rate the blood pressure measurement might not be accurate, i.e. it might not reflect the 'real' situation in your body. However, if the symbol appears often, we recommend that you seek medical advice. Please note that the device does not replace a cardiac examination.

### Preparing for use Pairing the blood pressure monitor to vour mobile device

### Note: To switch on the device for the first time, press the 'on' button for 3 seconds

Note: Before you use the device for the first time, remove the protective foil from the display.

The blood pressure monitor is equipped with Bluetooth. You can receive your personal health

data on a mobile device that is equipped with the Bluetooth function. Download the Philips

HealthSuite health app from the App store or Google Play. Use the search term 'Philips HealthSuite health app'. The app is available for iOS 8.0+ and Android 4.4+ Note: You can only use the Philips HealthSuite health app to communicate with the device. It is not possible to use third party applications.

Download the Philips HealthSuite health app on your mobile device, start the Setup wizard and follow the steps to create a user profile and

- add the blood pressure monitor. Make sure the app is active and Bluetooth is on when pairing is in progress.
- Keep the mobile device and the blood
- pressure monitor within transmission range (no more than 5 meters from each other, in
- the same room).
- With the device turned off, press the 'on' button for 3 seconds, until it turns on in pairing mode.

  These symbols are shown on the display alternatively, indicating that the connection
- is being established: (Fig. 6) and (Fig. 7). When pairing is successful, the display shows this symbol: (Fig. 8). The app shows which user
- profile is assigned to you. If the connection fails, the display shows this (Fig. svmbol: 9).
  - The blood pressure monitor has 2 user profiles. If both user profiles are in use, choose an existing profile to overwrite
  - You can also delete both user profiles by pressing and holding the user ID button for , 10 seconds. . The display of the approx. device shows 'del'. All stored date is deleted
- and you have to follow step 1-4 to pair and add a new user. The blood pressure monitor shows the Bluetooth icon on the display as soon the

connection has been established and switches off automatically after a few seconds. When the blood pressure monitor is successfully paired with your mobile device, the blood pressure monitor automatically transmits your personal

health data to your mobile device via Bluetooth Note: Only when the Philips HealthSuite health app is active, your personal health data can be transmitted.

# Measuring blood pressure

#### Tips for proper measurement Rest for 5 minutes before you measure your

- blood pressure. Wait at least 3 minutes between measurements.
- This allows your blood circulation to recover.

- For a meaningful comparison, try to measure under similar conditions. For example, take daily measurements at approximately the same time, on the same arm, or as directed by your doctor For a good Bluetooth connection between the blood pressure monitor and your mobile device, make sure the two are in close distance
- and there are no obstacles between the two devices. We recommend not to have the two
- devices further than 5 meters apart. We do not advise you to take a measurement under the following circumstances, as this may cause inaccurate measurements:
  - Within 1 hour after eating or drinking
  - Immediately after smoking
- Within 20 minutes after taking a bath While you are talking or moving your arm, hand or fingers In a very cold environment
- When you need to urinate Attaching the cuff

- Remove all jewelry, such as watches and bracelets from your left arm. Note: If your doctor has diagnosed you with poor circulation in your left arm, use your right
- 2
- Roll or push up your sleeve to expose the skin. Make sure your sleeve is not too tight.
- Hold your arm with your palm facing up and slide the cuff onto your left upper arm (Fig. 10). Position the lower edge of the cuff about 2-3cm
- above the crease of the elbow.
- Fasten the cuff around your arm, leaving no extra room between the cuff and your skin. If
- the cuff is too loose, the measurement will not be accurate.
- The cuff will not cause any potential sensitization or irritation of the skin. The
- materials of the cuff have been tested and
- found to comply with requirements of ISO 10993-5:2009, ISO 10993-1:2009 and ISO 10993-10:2010.
- Correct posture for measurement: Make sure you do not wear tight clothing
- during measurement.
- Sit comfortably with legs uncrossed, feet flat on the floor. Make sure that you sit upright with your back straight.
- The center of the cuff should be at the same
- level as the heart. Relax your wrist and hand. Do not bend your wrist back, clench your fist, or bend your
- wrist forward. Start measurement
- Press the user ID button (Fig. 11) or 'on' button (Fig. 12) once, to switch on the device. The
- device automatically selects the previous user.
- To change the user profile, press the user ID button (Fig. 11) and select a different user (Fig. 13). Make sure the correct user is

- selected, so the measurement data is properly transmitted and stored. It is not possible to change a user profile after a

- measurement.
- When the health app is open, the app

- automatically selects the correct user profile. In this case, the user profile can be changed
- by either closing the app and reopening it
- again with the correct user profile, or by
- closing the app and using the user ID button. Also a guest user can be selected. The guest
- user is for performing a measurement on
- other persons without a user profile in the
- health app. Measurements performed when
- using the guest user are not stored in the
- memory nor transmitted to the app
- Attach the cuff to your arm (see 'Attaching the

- cuff') and make sure your posture is correct (see 'Tips for proper measurement').

  Press the 'on' button to start the measurement 3
- (Fig. 12). All display characters are briefly shown on the display (Fig. 14). The device is ready for measurement and the number 0 appears (Fig. 15). Inflation of the cuff starts automatically.
- - During inflation, the unit determines the systolic pressure and diastolic pressure as well as heart rate. This is shown by the
    - heart rate detection symbol. The movement detector will light up when movement is detected. This may result in
      - inaccurate measurement results. When the measurement is finalized, the cuff

4

5

- deflates and the measurement results are shown on the display (Fig. 16). To transmit the measurement results to the app, see section 'Transmit and store personal health data in the
- app'
- Press the 'on' button to switch off the device.
- Note: after 1 minute, the device will turn off
- automatically
- If, after finishing the first measurement, another measurement is required, press the user ID button to select the correct user profile and follow steps
- 2-7 Note: Wait at least 3 minutes between measurements. This allows your blood circulation
- to recover.
- The device can store results of 60 blood pressure measurements for both user 1 and 2. However, only your current measurement will be visible on
- the device display. Your previous measurements will be visible in the app after syncing.

health app.

- Transmit and store personal health data in the app Note: Your personal measurement data is only stored and displayed in the Philips HealthSuite
- Activate the Philips HealthSuite health app and Bluetooth on your mobile device directly after a measurement.
- Keep the mobile device and the blood pressure monitor at transmission distance (no more than 5 meters from each other, in the same room).

- Once successfully connected, the measurement results are being transmitted to the health app 2 and the Bluetooth symbol lights up. If the data transmission is successful, the
- measurement results are displayed in
- the health app. If the data transmission fails, the Bluetooth symbol together with 'Err' is shown. The pending measurement data will be transmitted to your mobile device the next time it connects with your blood pressure monitor. You can also try to resend the data:
  - Activate the health app on your mobile device.
  - Press the user ID button or 'on' button to switch on the blood pressure monitor. The measurement results will be
    - automatically sent to your mobile device if a user profile has been connected. When the blood pressure monitor connects via Bluetooth to the app of a user, the device will automatically select that user and measurements can only be done for that user

### Cleaning and storage Caution: This device is not washable. Never

immerse the device in water and do not rinse it under running water. Caution: Avoid violent movements and hard

- contacts with objects. Switch off the device and unplug the adapter
- from the wall socket.
- Use a slightly damp or dry cloth to wipe the surface of the display (Fig. 17). Store the device in a cool, dry, and ventilated 2 3

environment.. For further information please refer to the transport and storage specifications

## detailed in this manual. Ordering accessories

To buy accessories or spare parts, visit www.shop.philips.com/service or go to your Philips dealer. You can also contact the Philips Consumer Care Center in your country (see the international warranty leaflet for contact details).

Recycling This symbol means that this product shall not be disposed of with normal household waste (2012/19/EU) (Fig. 18).

- This symbol means that this product contains a built-in rechargeable battery which shall not be disposed of with normal household waste (Fig. 19) (2006/66/EC). We strongly advise you to take your product to an official collection point or a Philips service center to have a professional remove the rechargeable battery. Follow your country's rules for the separate collection of electrical and electronic products
- helps prevent negative consequences for the environment and human health. Removing the rechargeable battery Warning: This procedure is irreversible. You

and rechargeable batteries. Correct disposal

#### cannot use the device anymore after this procedure.

Note: We strongly advise you to take your product to an official collection point or a Philips service centre to have a professional remove the battery

Caution: Observe basic safety precautions when you follow the procedure described below. Be sure to protect your eyes, hands, fingers, and the surface on which you work. Make sure the rechargeable battery is empty.

Remove the battery with appropriate tools. Warranty and support

- If you need information or support, please visit **www.philips.com/support** or read the

international warranty leaflet. Recalibration can be carried out by an appropriate authority or authorized service center. This

Open the device.

calibration will be charged for by said authority.
If you need more information about the app, please visit www.philips.com/healthprograms Troubleshooting

This chapter summarizes the most common problems you could encounter with the device. If you are unable to solve the problem with the information below, visit www.philips.com/support

# the Consumer Care Center in your country.

list of frequently

nt

**Troubleshooting** Possible Solution **Problem** cause

asked questions or

My blood pressure fluctuates throughout the day.	Your measurement position, the conditions under which you measure or the time of measurement, are different during each measureme-	For a meaningful comparison, try to measure under similar conditions. For example, take daily measurements at approximately the same time, on the same arm, or as directed by a doctor.

Problem	Possible cause	Solution	
	Fluctuations of blood pressure during the day are normal.	Blood pressure fluctuates from minute to minute and normally shows a circadian rhythm over a 24-hour period, with highest readings in the afternoons and lowest readings at night. That is why, for comparable measurements, the measurements should be taken at approx, the same time of day.	
	You are using medication.	The variations in blood pressure can be greater if you are using medication.	
	You performed multiple measure- ments directly after each other.	Wait at least 3 minutes between measurements. This allows your blood circulation to recover.	
My blood pressure measurement from the hospital is different from the measurement at home.	Multiple variables may affect your blood pressure such as the weather, emotions and exercise.	Pay attention when you measure your blood pressure at home. Check for instance: If the cuff is attached properly. If the cuff is not too tight or too loose. If the cuff is attached on the upper arm. If you feel anxious or stressed, try to relax. Take a deep breath 2-3 times before you start a measurement. Advice: Rest for 5 minutes before you measure your blood pressure	
The result is different when I perform measure-ments on my right arm.	The blood pressure monitor is suitable to be used on both arms, but the measurement results on the right arm and left arm will differ.	For a meaningful comparison, try to measure under similar conditions and measure on the same arm every time.	
The blood pressure monitor does not work when I press the 'on' button	rechargeable battery (see 'Charging'). s not k when I ss the button  light of display s and a sery abol-Lo nowing  display  Communica-  Check if the a		
The light of the display dims and a battery symbol+Lo is showing			
The display shows Err			
The display shows E3	The cuff is not properly secured.	Refasten the cuff, wait 3 minutes and then measure again.	
The display shows E10 or E11	The device detected motion, talking or the heart rate is too weak during the measurement.	Wait for 3 minutes and then measure again. Do not move during measurement.	
The display shows E20	The device does not detect the heart rate signal.	Make sure the device is in contact with the skin. Loosen the clothing on the arm and measure again.	
The display show E21	The measurement failed.	Wait for 3 minutes and then measure again.	
The display shows EExx	A system error occurred.	Retake the measurement. If the problem persists, contact the Philips Consumer Care Center in your country.	
Data transmission or pairing failed.	Bluetooth is off.	Turn on Bluetooth on your mobile device.	
	The Philips HealthSuite health app is off.	Press the icon on your mobile device to activate the health app.	

	perminate			
Accuracy	Pressure: 5°C-40°C within ±0.4kPa (3mmHg) heart rate: ±5% of measurement result on display			
Normal operating condition	Temperature: 5°C to 40°C. Relative humidity: ≤85%RH. Atmospheric pressure: 86kPa to 106kPa			
Storage and transportation conditions	Temperature: -20°C to 60°C. Relative humidity: 10% to 93%. Atmospheric pressure: 50kPa to 106kPa			
Measurement perimeter of the upper arm	About 22cm-42cm			
Net weight	Approx. 265g			
External dimensions	Approx. 130.9mm×73mm×29.4m- m			
Accessories	DC charger, user manual			
Mode of operation	Continuous operation			
Degree of protection	Type BF applied part			
Protection against ingress of water	IP22, This means: protected against access to hazardous parts with a finger and against vertically falling water drops when tilted up to 15 degrees.			
Device classification	Battery Powered Mode: Internally Powered ME Equipment. DC charger charged mode: Class II ME Equipment.			
Frequency band	2402-2480 MHz			
Modulation	GFSK			
Effective radiated power	Max. +5dBm			
Caution: No modification of this equipment is allowed.				
standard EN 60601-1-2. typical domestic enviror Hereby, Philips Consum that the radio equipmen compliance with Directiv The full text of the EU davailable at the followin address: www.philips.co	according to EMC safety It is designed to be used in ments. er Lifestyle BV declares at type Bluetooth LE is in ve 2014/53/EU. eclaration of conformity is g internet			
	Monitor needs special			

precautions regarding EMC and needs to be installed and put into service according to the EMC information provided in the accompanying

Wireless communications equipment such as wireless home network devices, mobile phones cordless telephones and their base stations walkie-talkies can affect this equipment and should be kept at least a distance d = 3.3 m

Note: As indicated in IEC 60601-1-2:2007 for ME equipment, a typical cell phone with a maximum output power of 2 W yields d = 3.3 m at an

documents.

away from the equipment.

immunity level of 3V/m.

Possible

The blood pressure

monitor and

You selected

the wrong profile on the

blood

Specifications

Measurement method

Measurement range

Power supply

Display

pressure

monitor.

device are more than 5 meters away from each other

cause

mobile

**Problem** 

Solution

monitor.

Place your mobile

Select the correct

user profile on the blood pressure

your measurement

Otherwise the data cannot be transmitted to your app. Repeat the measurement with the correct profile selected

monitor before

3.7V 1000mAH built-in

Display with white LED backlight

x 24mm (W)

0kPa-40kPa (OmmHg-300mmHg). Measurement pressure: 3kPa-30.7kPa (40mmHg-230mmHg). Heart rate: 40-199 beats

per minute

rechargeable li-polymer battery, 6V 1A DC charger

Visible area = 86.1mm (L)

Oscillometric method Rated cuff pressure:

blood pressure

device closer to the

### Table 1 Guidance and manufacturer's declaration – electromagnetic emissions - for all ME equipment and ME systems Guidance and manufacturer's declaration -

electromagnetic emissions The device is intended for use in the electromagnetic environment specified below. The customer or the user of the device should assure

that it is used in such an environm		actice bilouta abbaic
Emissions test	Com- pli- ance	Electromagnetic environment - guidance
RF emissions CISPR 11	Group 1	The device must emit electromagnetic energy in order to perform its intended function. Nearby electronic equipment may be affected.
RF emissions CISPR 11	Class B	
Harmonic	Not	

applica ble

applica

Not

emissions IEC 61000-3-2

Voltage

fluctua-

tions/flicker

emissions IEC 61000-3-3	
	manufacturer's declaration letic immunity – for all ME d ME systems

equipment and ME systems Guidance and manufacturer's declaration — electromagnetic immunity The device is intended for use in the electromagnetic environment specified below. The customer or the user of the device should assure that it is used in such an environment.			
Immu- nity test	IEC 60601 test level	Com- pli- ance level	Electromagnetic environment - guidance
Electro- static dis- charge (ESD) IEC 61000 4-2	±6 kV contact ±8 kV air	±6 kV contact ±8 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%.
Electri- cal fast tran- sient/b- urst IEC 61000 4-4	±2 kV for power supply lines ±1 kV for input/o- utput lines	±2 kV for power supply lines	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000 4-5	±1 kV line(s) to line(s) ±2 kV line(s) to earth	±1 kV line(s) to line(s)	Mains power quality should be that of a typical commercial or hospital environment.
Voltage dips, short interrup- tions and voltage varia- tions on power supply input lines IEC 61000 4-11	<5% UT (>95% dip in UT) for 0.5 cycle 40% UT (60% dip in UT) for 5 cycles 70% UT (30% dip in UT) for 25 cycles <5% UT (>95% dip in UT) for 5 s	<5% UT (>95% dip in UT) for 0.5 cycle 40% UT (60% dip in UT) for 5 cycles 70% UT (30% dip in UT) for 25 cycles <5% UT (>95% dip in UT) for 5	Mains power quality should be that of a typical commercial or hospital environment. If the user of the device requires continued operation during power mains interruptions, it is recommended that the device be powered from an uninterruptible power supply or a battery.
Power fre- quency (50/60- Hz)	3A/m	3A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical

IEC 61000 4-8	hospital environment.
Note: UT is the AC mains application of the test lev	<u> </u>
Table 4 Guidance an	nd manufacturer's

magnetic field

a typical

commercial or

declaration – electromagnetic immunity –for ME equipment and ME systems that are not life supporting Guidance and manufacturer's declaration -

electromagnetic immunity .The device is intended for use in the electromagnetic environment specified below. The customer or the user of the

device should assure that it is used in such an environment.				
IMMUNITY test	IEC 60601 TEST LEVEL	Compliance level		
Conducted RF IEC 61000-4-6	3 Vrms 150 kHz to 80 MHz	3 Vrms		
Radiated RF	3 V/m	3 V/m		

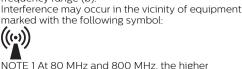
3 V/m 80 MHz to 2.5 IEC 61000-4-3 GHz Electromagnetic environment - guidance Portable and mobile RF communications

equipment should be used no closer to any part of the device, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter

Recommended separation distance:

d =  $1.2 \sqrt{P}$ d =  $1.2 \sqrt{P}$  80 MHz to 800MHz d =  $2.3 \sqrt{P}$  800 MHz to 2.5 GHz

where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in metres (m). Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey (a), should be less than the compliance level in each frequency range (b).



NOTE 1 At 80 MHz and 800 MHz, the higher

frequency range applies. NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures,

objects and people. (a) Field strengths from fixed transmitters, such as

base stations for radio (cellular/cordless)

telephones and land mobile radios, amateur radio AM and FM radio broadcast and TV broadcast

cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site

survey should be considered. If the measured field strength in the location in which the device is used exceeds the applicable RF compliance level above, the device should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the device (b) Over the frequency range 150 kHz to 80 MHz,

Table 6 Recommended separation distances between portable and mobile RF communications equipment and the ME equipment or ME system – for ME equipment and ME systems that are not life supporting

field strengths should be less than 3V/m

Recommended separation distances between portable and mobile RF communications

equipment and the device. The device is intended for use in an electromagnetic environment in which radiated RF

disturbances are controlled. The customer or the user of the device can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile

RF communications equipment (transmitters) and the device as recommended below, according to

the maximum output power of the communications equipment Separation distance according to frequency of transmitter (m)

Rated 150 kHz to 80 MHz 800 MHz 80 MHz d = 1.2 √ P to 800 maximum to 2.5 GHz output d = 2.3MHzpower of transmitter d = 1.2 √ P

(W)			
0.01	0.12	0.12	0.2
O.1	0.37	0.37	0.74
1	1.2	1.2	2.3
10	3.7	3.7	7.4
100	12	12	23
not listed distance the equa	l above, the re d in metres (r tion applicabl	ecommende n) can be es le to the frec	timated using

rating of the transmitter in watts (W) according to the transmitter manufacturer NOTE 1 At 80MHz and 800MHz, the separation distance for the higher frequency range applies. NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.