iHealth®

Wireless Blood Pressure Monitor

Tensiomètre sans fil
Misuratore di pressione wireless
Monitor de presión arterial inalámbrico
Drahtloses blutdruckmessgerät
Monitor sem fios da tensão arterial
Draadloze bloeddrukmeter
Ασύρματο Πιεσόμετρο

OWNER'S MANUAL

MANUEL DE L'UTILISATEUR
MANUALE D'ISTRUZIONI
MANUAL DEL PROPIETARIO
BEDIENUNGSANLEITUNG
MANUAL DO PROPRIETÁRIO
GEBRUIKERSHANDLEIDING
ΟΔΗΓΟΣ ΧΡΗΣΤΗ

EN

iHealth®

Wireless Blood Pressure Monitor (BP5)
OWNER'S MANUAL

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INTRODUCTION

Thank you for selecting the iHealth Wireless Blood Pressure Monitor. The iHealth Wireless Blood Pressure Monitor is a fully automatic arm cuff blood pressure monitor that uses the oscillometric principle to measure your blood pressure and pulse rate. The monitor works with your mobile device to test, track and share vital blood pressure data.

PACKAGE CONTENTS

- 1 iHealth Wireless Blood Pressure Monitor
- 1 Owner's Manual
- 1 Quick Start Guide
- 1 Charging Cable
- 1 Travel Bag

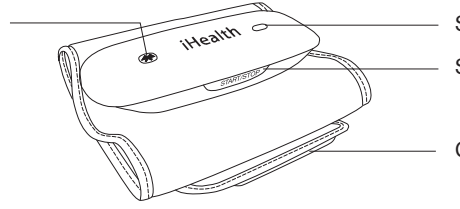
INTENDED USE

The iHealth Wireless Blood Pressure Monitor (Electronic Sphygmomanometer) is intended for use in a professional setting or at home and is a non-invasive blood pressure measurement system. It is designed to measure the systolic and diastolic blood pressures and pulse rate of an adult individual by using a technique in which an inflatable cuff is wrapped around the upper arm. The measurement range of the cuff circumference is 8.6" to 18.9"(22cm-48cm).
Note: *Consult your medical professional for proper interpretation of blood pressure results.*

CONTRAINDICATION

⚠ It is not recommended for people with serious arrhythmia to use this Wireless Blood Pressure Monitor.

PARTS AND DISPLAY INDICATORS



SET UP REQUIREMENTS

The iHealth Wireless Blood Pressure Monitor is designed to be used with the following iPod touch, iPhone and iPad models:
iPhone 4+
iPad mini +
iPad Air
iPad 2+
iPod touch (5th generation)+
The iOS version of these devices should be V6.0 or higher.

The iHealth Wireless Blood Pressure Monitor is also compatible with a number of Android devices, the Android version should be V3.0 or higher, and RAM should be 1.0G or more. For a complete list of compatible devices, please visit the support page on www.ihealthlabs.com

SET UP PROCEDURES

Download the Free iHealth MyVitals App

For iOS device: Prior to first use, download and install “iHealth MyVitals” from the App Store. For Android device: Prior to first use, download and install “iHealth MyVitals” from the Google Play. Follow the on-screen instructions to register and set up your personal account.

Access the iHealth Cloud Account

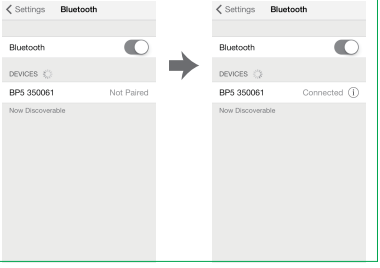
Your iHealth account also gives you access to the free and secure iHealth cloud service. Go to www.ihealthlabs.com and click “Sign In” for access once your account has been created.

Charge Battery Before First Use

Connect the monitor to a USB port using the charging cable provided until the green indicator light stabilizes.

Connect to iOS Device via Bluetooth

- Apply the cuff or press the START/STOP button, the *Bluetooth* indicator will begin flashing.
- Turn *Bluetooth* "On" under the "Settings" menu on the iOS device.
- Wait until the model name printed on the monitor, (i.e. "BP5 xxxxxx") and "Not Paired" appear in the *Bluetooth* menu, and select the model name "BP5



xxxxxx" to pair and connect. The *Bluetooth* indicator will remain steady upon successful connection. When using the monitor for the first time, it may take up to 30 seconds for your iOS device to detect the Bluetooth signal.

- Each subsequent time you use the monitor, "Connected" will be displayed next to "BP5 xxxxxx" in the *Bluetooth* Menu.
- Launch the “iHealth MyVitals” app to start using your monitor.
- Please repeat these steps when you switch to another iOS device with the Monitor.

Connect to Android Device via Bluetooth

- Apply the cuff or press the START/STOP button, the Bluetooth indicator will begin flashing.
- In the setting menu, turn the Bluetooth on.
- When using the monitor for the first time, you should pair the monitor to the Android device. Wait until the model name printed on the monitor, (i.e. "BP5 xxxxxx") appears in the Bluetooth menu, and select the model name "BP5 xxxxxx" to pair. it may take up to 30 seconds for your Android device to detect the Bluetooth signal.
- Launch the “iHealth MyVitals” app to start using your monitor.
- Please repeat these steps when you switch to another Android device with the monitor.

Monitor Status	Bluetooth Indicator
Waiting to connect	Flashing blue light
Connected and measuring	Steady blue light
Measurement completed and ready to disconnect	Gradually extinguishing light

MEASUREMENT PROCEDURES

Blood pressure can be affected by the position of the cuff and your emotional condition. It is very important that the cuff is positioned at your heart level during blood pressure measurements.

Body Posture

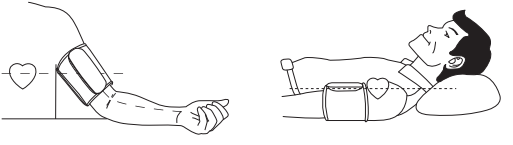
Sitting Comfortably During Measurement

a. Be seated with your feet flat on the floor without crossing your legs.
b. Place your hand palm-side up in front of you on a flat surface such as a desk or a table.
c. The middle of the cuff should be at the level of your heart.

Lying Down During Measurement

a. Lie on your back.
b. Place your arm straight along your side with your hand palm-side up.
c. The cuff should be placed at the same level as your heart.

Note: *Blood pressure can be affected by the position of the cuff and your emotional condition, so it is important that you always measure your blood pressure in the same position.*



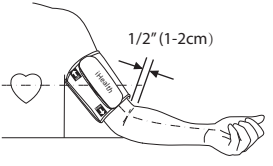
Apply the Cuff

a. Pull the cuff end through the metal loop, positioning it outward (away from your body).
b. Place a bare arm through the cuff and position the cuff 1/2" (1-2cm) above the elbow joint.

c. Tighten the cuff by pulling it towards your body, securing it closed with the Velcro fastener.

d. While seated, place your hand, palm-side up, in front of you on a flat surface such as a desk or table. Position the monitor in the middle of your arm so that it is aligned with your middle finger.

e. The cuff should fit comfortably, yet snugly around your arm. You should be able to insert one finger between your arm and the cuff.



Remember to:

1. Make sure that the appropriate cuff size is used; refer to the cuff circumference range in the Specifications section of this manual.
2. Measure on the same arm each time.
3. Stay still during a measurement. Do not move your arm, body or the monitor.
4. Stay still and calm for one to one and half minutes before taking a blood pressure measure ment.Prolonged over-inflation of the cuff may cause bruises on your arm.
5. Keep the cuff clean. Cleaning the cuff after every 200 measurements is recommended.If the cuff becomes dirty, clean it with a moistened cloth. Do not rinse the monitor or cuff with running water.

Press the “START/STOP” button at any time to interrupt a measurement.
Press the “START/STOP” button for 2 seconds to turn off the monitor manually.
Note: *Physical activity, eating, drinking, smoking, excitement, stress, and many other factors influence blood pressure results.*

Auto Connect Option

The auto connect option allows the monitor to find the last used iOS device and re-establish the connection automatically. The auto connect option can be enabled in the App.

Taking Measurements with Mutiple iOS Devices
Turn off the Bluetooth of the last used iOS device if the Auto Connect option is enabled in your App, then follow the Set Up Instructions in the Quick Start Guide.

Measuring without an iOS Device
Enable the Offline Measurement function on the App. Apply the cuff, follow the “Measurement Procedures”, and then press the “START/STOP” button to begin measurement. All offline measurements will be uploaded to the App automatically upon the next successful Bluetooth connection.
For answers to frequently asked questions, please visit www.ihealthlabs.com

SPECIFICATIONS

1. Product name: Wireless Blood Pressure Monitor
2. Model: BP5
3. Classification: Internally powered; Type BF applied part; IPX0, No AP or APG; Continuous operation
4. Machine size: approx. 5.7” × 2.3” ×1.2”(145mm × 58mm ×30mm)
5. Cuff circumference: 8.6”-16.5”(22cm-42cm), 16.5”-18.9”(42cm-48cm) (XL size sold separately)
6. Weight: approx. 5.1oz (145g) (including cuff); approx. 5.6oz (160g) (including XL cuff)
7. Memory volume: 120 times with time and date stamp (off-line measurement only)
8. Power: DC:5.0V ⎓ 1.0A, Battery: 1*3.7V ⎓ Li-ion 400mAh
9. Measurement range:
   Cuff pressure: 0-300 mmHg
   Systolic: 60-260 mmHg
   Diastolic: 40-199 mmHg

Pulse rate: 40-180 beats/minute

10. Accuracy:
    Pressure: ±3 mmHg
    Pulse rate: ±5%
11. Environmental temperature for operation: 5°C-40°C(41°F -104°F)
12. Environmental humidity for operation: ≤90%RH
13. Environmental temperature for storage and transport:-20°C-55°C(-4°F-131°F)
14. Environmental humidity for storage and transport: ≤90%RH
15. Environmental pressure: 80kPa-105kPa
16. Battery life: more than 80 measurements on a full charge
17. The blood pressure measurement system includes accessories: pump, valve, cuff, and sensor.

Note: *These specifications are subject to change without notice.*

GENERAL SAFETY AND PRECAUTIONS

1. Read all of the information in the Owner's Manual and other provided instructions before operating the unit.
2. Consult your medical professional for any of the following situations:
   a)The application of the cuff over a wound or inflamed area.
   b)The application of the cuff on any limb with intravascular access or therapy, or an arterio-venous (A-V) shunt.
   c)The application of the cuff on the arm on the side of a mastectomy.
   d)Simultaneous use with other medical monitoring equipment on the same limb.
   e)The blood circulation of the user needs to be checked.
3. Do not use this product in a moving vehicle as this may result in inaccurate measurements.

4. Blood pressure measurements determined by this product are equivalent to those obtained by professional healthcare practitioners using the cuff/stethoscope auscultation method within the limits prescribed by the American National Standard, Electronic or Automated Sphygmomanometer. This device is also clinically validated according to the 2010 Protocol of the European Society of Hypertension(ESH 2010).
5. If you are using a smart phone to operate the device and a phone call comes in during the measurement, the measurement process will be terminated automatically. It is thus recommended that the phone be set in Airplane mode during measurement to avoid interrupting the measurement.
6. If an Irregular Heartbeat (IHB) is detected during the measurement procedure, the IHB symbol will be displayed. Under this condition, the Wireless Blood Pressure Monitor can keep function- ing, but the results may be inaccurate. Please consult your medical professional for an accurate assessment.
   The IHB symbol will be displayed under 2 sets of circumstances:
   1) The coefficient of variation (CV) of the pulse is higher than 25%.
   2) The difference of adjacent pulse periods is higher or equal to 0.14s and the number of such pulse takes more than 53 percent of the total number of pulses.
7. Please do not use any cuff other than that supplied by the manufacturer as this may result in inaccurate measurements.
8. For information regarding potential electromagnetic or other interference between the blood pressure monitor and other devices together with advice regarding avoidance of such interference, please see ELECTROMAGNETIC COMPATIBILITY INFORMATION. It is suggested that the blood pressure monitor should be operated at least 10 metres away from electric or wireless devices, such as routers, microwave oven, etc.
9. If the determined blood pressure (systolic or diastolic) is outside the rated range specified in

part SPECIFICATIONS, the app will immediately display a technical alarm on the screen. In this case, repeat the measurement ensuring that the proper measurement procedures are followed and/or consult with your medical professional. The technical alarm is preset in the factory and cannot be adjusted or inactivated. This technical alarm is assigned as low priority according to IEC 60601-1-8.The technical alarm does not need to be reset.

10. A medical AC adapter with an output of DC 5.0V and complies with IEC 60601-1/UL 60601-1 and IEC 60601-1-2/EN 60601-1-2 is suitable for this monitor, such as ASP5-05010002JU (input: 100-240V, 50/60Hz, 200mA; output: DC 5.0V, 1.0A). Please note that the monitor jack size is USB mini B. The USB jack should be used for charging only.

⚠ This Wireless Blood Pressure Monitor is designed for adults and should never be used on infants, young children, pregnant or pre-eclamptic patients. Consult your medical professional before use on children under the age of 16 years.

⚠ This product might not meet its performance specifications if stored or used outside the specified temperature and humidity ranges.

⚠ Please do not share the cuff or apply the cuff on a wound to avoid cross-infection.

BATTERY HANDLING AND USAGE

- When the monitor is connected to an iOS device, the battery volume will be displayed on the iOS device. If the power is less than 25%, please charge the battery. The monitor will not work until the battery has enough power.
- When charging is needed, please connect the monitor to a power source. The monitor can work normally while charging.
- It is suggested that you charge the battery when the battery is less than 25%. Overcharging the battery may reduce its lifetime.

• When in charging mode, the LED on the device will be displayed with different colous indicating the charging status. See the table below for details.

Monitor Status	Status Indicator
Charging	Flashing green light
Fully charged	Steady green light
Low battery	Flashing red light (for a few seconds)
Abnormal state	Steady red light

⚠ The battery in this device is a fixed battery and can only be changed by an authorized iHealth After Sales Service agent. Should your battery be defective, please contact the iHealth Customer Service for a replacement.
⚠ Overcharging the battery may reduce its lifetime.
⚠ Lithium battery replacement by inadequately trained personnel could result in a hazard such as a fire or explosion.
⚠ Do not plug or unplug the power cord into the electrical outlet with wet hands.
⚠ If the AC adapter is abnormal, please change the adapter.
⚠ Do not pull out the adapter when you are using the monitor.
⚠ Do not use any other type of AC adapter as it may harm the monitor.

The monitor, cable, battery and cuff must be disposed of according to local regulations at the end of their usage.

Note: *Battery life and charge cycles vary by use and settings.*

TROUBLESHOOTING

PROBLEM	POSSIBLE CAUSE	SOLUTION
Low Battery	Battery is less than 25%	Charge the battery
Display reads "ERROR"	Blood pressure is outside of measurement range	Retest and contact your health professional if blood pressure measurement is still outside of normal range
	Arm or monitor was moved during test	Retest, make sure not to move your arm or the monitor
	The cuff does not inflate properly or pressure falls quickly during test	Review the cuff application instructions and retest
	Irregular heartbeat (arrhythmia)	It is inappropriate for people with serious arrhythmia to use this monitor. Check with your medical professional
	The cuff was not properly applied	Review the cuff application instructions and retest
Display reads an abnormal result	The cuff position was not correct or it was not properly tightened	Review the cuff application instructions and retest
	Body posture was not correct during testing	Review body posture instructions and retest
	Speaking, moving arm or body, being angry, excited or nervous during test	Retest when calm; avoid speaking or movement during the test
Bluetooth connection unstable	**Bluetooth** connection unsuccessful, monitor is abnormal, or strong electromagnetic interference is present	Reset iOS/Android device. Reset monitor by pressing the START/STOP button and holding for about 10 seconds. Make sure the monitor and iOS/Android device are away from other electrical equipment. Please see GENERAL SAFETY AND PRECAUTIONS
No response	Incorrect operation or strong electromagnetic interference	Press the START/STOP button and hold for about 10 seconds to reset the device. Relaunch app, and reconnect the iOS/Android device to the monitor

CARE AND MAINTENANCE

1. If this monitor is stored at a temperature under 5°C, allow it to acclimatise to room temperature before use (environmental temperature for operation: 5°C~40°C).
2. If the monitor is not used for a long time, please be sure to fully charge it every month.
3. It is recommended that product performance be checked every 2 years or after each repair. Please contact the iHealth Customer Service Center to do so.
4. No monitor component needs to be maintained by the user. The circuit diagrams, component part lists, descriptions, calibration instructions, or other information which will assist the user's appropriately qualified technical personnel to repair those parts of the equipment which are designated for repair can be supplied by the iHealth technical department.
5. Clean the monitor with a dry, soft cloth or a moistened and well wrung soft cloth using water, diluted disinfectant alcohol, or diluted detergent.
6. The monitor can maintain the safety and performance characteristics for a minimum of 10,000 measurements or three years of usage, and the cuff integrity is maintained after 1,000 open-close cycles.
7. The battery can maintain the performance characteristics for a minimum of 300 charge cycles.
8. It is recommended that if the cuff is used, for example, in a hospital or a clinic, it be disinfected twice a week. Wipe the inner side (the side that contacts skin) of the cuff with a soft cloth lightly moistened with Ethyl alcohol (75-90%). Then air dry the cuff.

⚠ Do not drop this monitor or subject it to strong impacts.

⚠ Avoid high temperature and direct sunlight. Do not immerse the monitor in water as this will result in damage to the monitor.

⚠ Do not attempt to disassemble this monitor.

⚠ Battery replacement should only be performed by a qualified iHealth technician. To do

otherwise will void your warranty and possibly damage your unit.

⚠ Cuff replacement should only be performed by a qualified iHealth technician. To do otherwise will possibly damage your unit.

WARRANTY INFORMATION

The iHealth Wireless Blood Pressure Monitor is warranted to be free from defects in materials and workmanship within one year from the date of purchase when used in accordance with the provided instructions. The warranty extends only to the end user. We will, at our option, repair or replace without charge the iHealth Wireless Blood Pressure Monitor covered by the warranty. Repair or replacement is our only responsibility and your only remedy under the warranty.

EXPLANATION OF SYMBOLS

Symbol for "TYPE BF APPLIED PARTS" (cuff only)

Symbol for "THE OPERATION GUIDE MUST BE READ"
The sign background color: blue The sign graphical symbol: white

Symbol for "ENVIRONMENT PROTECTION – Waste electrical products should not be disposed of with household waste. Please recycle where facilities exist. Check with your local authority or retailer for recycling advice".

Symbol for "KEEP DRY"

 Symbol for "WARNING"

 Symbol for "MANUFACTURER"

SN Symbol for "SERIAL NUMBER"

 Symbol for "EUROPEAN REPRESENTATIVE"

CE 0197 Symbol for "COMPILES WITH MDD93/42/EEC REQUIREMENTS"

iHealth is a trademark of iHealth Lab Inc.

"Made for iPod", "Made for iPhone", and "Made for iPad" mean that an electronic accessory has been designed to connect specifically to iPod, iPhone, or iPad, respectively, and has been certified by the developer to meet Apple performance standards. Apple is not responsible for the operation of this device or its compliance with safety and regulatory standards. Please note that the use of this accessory with iPod, iPhone, or iPad may affect wireless performance. iPad, iPhone, and iPod touch are trademarks of Apple Inc., registered in the U.S. and other countries.

CONTACT AND CUSTOMER SERVICE

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IMPORTANT INFORMATION REQUIRED BY THE FCC

This device complies with Part 15 of the FCC Rules. Its operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and
(2) this device must accept any interference received, including interference that may cause undesired operation.
Changes or modifications not expressly approved by iHealth Lab Inc. would void the user's authority to operate the product.
Note: *This product has been tested and found to comply with the limits for a Class B digital device, pursuant to Part 15 of the FCC Rules. These limits are designed to provide reasonable protection*

against harmful interference in a residential installation. This product generates, uses, and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation. If this product does cause harmful interference to radio or television reception, which can be determined by turning the equipment off and on, the user is encouraged to try to correct the interference by one or more of the following measures:
—Reorient or relocate the receiving antenna.
—Increase the separation between the equipment and receiver.
—Connect the equipment into an outlet on a circuit different from that to which the receiver is connected.
—Consult the dealer or an experienced radio/TV technician for help.
This product complies with Industry Canada. IC: RSS-210
IC NOTICE
This device complies with Industry Canada licence-exempt RSS standard(s). Operation is subject to the following two conditions:
(1) this device may not cause interference, and
(2) this device must accept any interference, including interference that may cause undesired operation of the device.
This product is approved in accordance to R&TTE directive transmitter.

OTHER STANDARDS AND COMPLIANCES

The Wireless Blood Pressure Wrist Monitor corresponds to the following standards:
IEC 60601-1:2005+C1:2006+C2:2007(Medical electrical equipment – Part 1: General requirements for safety);

IEC 60601-1-2:2007 (Medical electrical equipment – Part 1: General requirements for safety; Collateral Standard-Electromagnetic compatibility - Requirements and tests);
EN 1060-1: 1995 + A1: 2002 + A2: 2009 (Non-invasive sphygmomanometers - Part 1: General requirements);
EN 1060-3: 1997 + A1: 2005 + A2: 2009 (Non-invasive sphygmomanometers - Part 3: Supplementary requirements for electro-mechanical blood pressure measuring systems);
AAMI/ANSI 80601-2-30：2009/IEC 80601-2-30：2009+Cor.2010/EN 80601-2-30:2010(Medical electrical equipment –Part 2-30: Particular requirements for the basic safety and essential performance of automated non-invasive sphygmomanometers).

ELECTROMAGNETIC COMPATIBILITY INFORMATION

Table 1

For all ME EQUIPMENT and ME SYSTEMS

Guidance and manufacture's declaration - electromagnetic emissions		
BP5 is intended for use in the electromagnetic environment specified below. The user of BP5 should ensure that it is used in such an environment.		
Emissions test	**Compliance**	**Electromagnetic environment - guidance**
RF emissions CISPR 11	Group 1	BP5 uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emissions CISPR 11	Class B	BP5 is suitable for use in all establishments, including domestic establishments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.
Harmonic emissions IEC 61000-3-2	Class A	
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Complies	

Table 2

For all ME EQUIPMENT and ME SYSTEMS

Guidance and manufacturer's declaration - electromagnetic immunity			
BP5 is intended for use in the electromagnetic environment specified below. The user of BP5 should ensure that it is used in such an environment.			
IMMUNITY test	**IEC 60601test level**	**Compliance level**	**Electromagnetic environment - guidance**
Electrostatic discharge (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 %.
Electrical fast transient/burst IEC 61000-4-4	± 2 kV for power supply lines	± 2 kV for power supply lines	Main 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) ± 2 kV line(s) to earth	Main power quality should be that of a typical commercial or hospital environment.
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	<5 % U_T (>95 % dip in U_T) for 0.5 cycle 40 % U_T (60 % dip in U_T) for 5 cycles 70 % U_T (30 % dip in U_T) for 25 cycles <5 % U_T (>95 % dip in U_T) for 5 s	<5 % U_T (>95 % dip in U_T) for 0.5 cycle 40 % U_T (60 % dip in U_T) for 5 cycles 70 % U_T (30 % dip in U_T) for 25 cycles <5 % U_T (>95 % dip in U_T) for 5 s	Main power quality should be that of a typical commercial or hospital environment. If the user of BP5requires continued operation during power main interruptions, it is recommended that BP5 be powered from an uninterruptible power supply or a battery.

Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	3 A/m	3 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.
NOTE: U_T is the a.c. mains voltage prior to application of the test level.			

Table 3

For ME EQUIPMENT and ME SYSTEMS that are not LIFE-SUPPORTING

Guidance and manufacturer's declaration - electromagnetic immunity			
BP5 is intended for use in the electromagnetic environment specified below. The user of BP5 should ensure that it is used in such an environment.			
IMMUNITY test	**IEC 60601test level**	**Compliance level**	**Electromagnetic environment - guidance**
			Portable and mobile RF communications equipment should be used no closer to any part of BP5, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance:
Conducted RF IEC 61000-4-6	3 Vrms 150 kHz to 80 MHz	3 V	$d = 1.2\sqrt{P}$
Radiated RF IEC 61000-4-3	3 V/m 80 MHz to 2.5 GHz	3 V/m	$d = 1.2\sqrt{P}$ 80 MHz to 800 MHz $d = 2.3\sqrt{P}$ 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 meters (m). Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey,[a] should be less than the compliance level in each frequency range.[b] Interference may occur in the vicinity of equipment marked with the following symbol: 

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 BP5 is used exceeds the applicable RF compliance level above, BP5 should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating BP5.
b) Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.

Table 4

For ME EQUIPMENT and ME SYSTEMS that are not LIFE-SUPPORTING

Recommended separation distances between portable and mobile RF communications equipment and the Wireless Blood Pressure Monitor

BP5 is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The user of BP5 can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and BP5 as recommended below, according to the maximum output power of the communication equipment.

Rated maximum output power of transmitter W	Separation distance according to frequency of transmitter m		
	150 kHz to 80 MHz $d = 1.2\sqrt{P}$	80 MHz to 800 MHz $d = 1.2\sqrt{P}$	800 MHz to 2,5 GHz $d = 2.3\sqrt{P}$
0.01	0.12	0.12	0.23
0.1	0.38	0.38	0.73
1	1.2	1.2	2.3
10	3.8	3.8	7.3
100	12	12	23

For transmitters rated at a maximum output power not listed above, the recommended separation distance d in meters (m) can be determined using the equation applicable to the frequency of the transmitter, where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer.

NOTE 1 At 80 MHz and 800 MHz, 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.