

## **INSTRUCTION MANUAL**

# 3 series Upper Arm Blood Pressure Monitor

## **Model BP7100**



# **Table of Contents**

| Intr | roduction                               | 3  |
|------|---|----|
| Imp  | portant Safety Information              | 4  |
| 1.   | Know Your Monitor                       | 8  |
|      | Preparation                             |    |
|      | Using the Monitor                       |    |
|      | Error Messages and Troubleshooting      |    |
| 5.   | Maintenance                             | 23 |
| 6.   | Specifications                          | 25 |
|      | FCC Statement                           |    |
|      | Limited Warranty                        |    |
| 9.   | Guidance and Manufacturer's Declaration | 30 |

## Introduction

Thank you for purchasing the OMRON BP7100 Blood Pressure Monitor. This new 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.

#### **Safety Instructions**

This instruction manual provides you with important information about the OMRON BP7100 Blood Pressure Monitor. To ensure the safe and proper use of this monitor, READ and UNDERSTAND all of these instructions. If you do not understand these instructions or have any questions, contact 1-800-634-4350 before attempting to use this monitor. For specific information about your own blood pressure, consult with your physician.

#### Intended Use

This 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.

Environments of Use: Home Patient Population: Adult

#### **Receiving and Inspection**

Remove this monitor from the packaging and inspect for damage. If this monitor is damaged, DO NOT USE and contact 1-800-634-4350.

#### Symbols Glossary

For symbol information, visit:

OmronHealthcare.com/symbols-glossary

#### Contraindications

- This monitor is contraindicated for use in ambulatory environments.
- This monitor is contraindicated for use on aircraft.

## **Important Safety Information**

Please read the Important Safety Information in this instruction manual before using this monitor.

Please follow this instruction manual thoroughly for your safety.

Please keep for future reference. For specific information about your own blood pressure, CONSULT WITH YOUR PHYSICIAN.

## **A** 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.
- 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 or in oxygen rich environments. This may result in incorrect operation of the monitor and/or cause an inaccurate reading.
- 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 away from infants, toddlers or children.
- This product contains small parts that may cause a choking hazard if swallowed by infants, toddlers or children.

#### **Battery Handling and Usage**

• Keep batteries out of the reach of infants, toddlers or children.

# **⚠** 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 which 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.
- Do not use a mobile phone or other devices that emit electromagnetic fields, near the monitor. This may result in incorrect operation of the monitor.
- 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.
- 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 5.
- Ensure this monitor is not impairing blood circulation by observing the arm while measurement is occurring.

- 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 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 of operating and storage/transport temperature, refer to section 5.
- 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 air 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 arm cuff, batteries and accessories specified for this monitor.
   Use of unsupported 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: please refer to "If your systolic pressure is more than 210 mmHg" in section 3 of this instruction manual for additional information.
- Read and follow the "Important information regarding Electromagnetic Compatibility (EMC)" in section 9 of this manual.

#### **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.
- 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.

## 1. Know Your Monitor

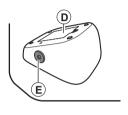
#### Contents:

Monitor, arm cuff, instruction manual, quick start guide

### **Monitor:**

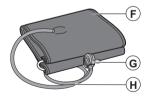


- A. Display
- B. START/STOP button
- C. Memory button



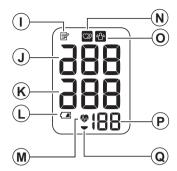
- D. Battery compartment
- E. Air jack

#### Arm cuff:



- F. Arm cuff (Arm circumference 9" 17" (22 42 cm))
- G. Air plug
- H. Air tube

## Display:

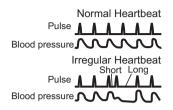


- I. Memory symbol
- J. Systolic blood pressure
- K. Diastolic blood pressure
- L. Low battery symbol
- M. Heartbeat symbol (Flashes during measurement.)
- N. Irregular heartbeat symbol
- O. Movement error symbol
- P. Pulse display / Memory number
- Q. Deflation symbol

## 1.1 Display symbols

#### Irregular Heartbeat Symbol ()

When the monitor detects an irregular rhythm two or more times during the measurement, the irregular heartbeat symbol will appear on the display with the measurement values. An irregular heartbeat rhythm is defined as a rhythm that is 25% less or 25% more than the average rhythm detected while the monitor is measuring the systolic and diastolic blood pressure.



If the irregular heartbeat symbol displays with your measurement results, we recommend you consult your physician. Follow the directions of your physician.

### Movement Error Symbol (M)

The movement error symbol is displayed if you move your body during the measurement. Please remove the arm cuff. and wait 2 - 3 minutes.

Take another measurement, remain still during measurement.

#### Classification of BP (Blood Pressure)

ACC/AHA 2017 Hypertension guidelines define categories of BP in adults as below.

#### Categories of BP in Adults\*

| BP Category  | Systolic BP  |       | Diastolic BP |            |
|--------------|--------------|-------|--------------|------------|
| Normal       | < 120 mmHg   | and   |              | < 80 mmHg  |
| Elevated     | 120-129 mmHg | and < |              | < 80 mmHg  |
| Hypertension |              |       |              |            |
| Stage 1      | 130-139 mmHg | 0     | r            | 80-89 mmHg |
| Stage 2      | ≥ 140 mmHg   | 0     | r            | ≥ 90 mmHg  |

<sup>\*</sup> Individuals with Systolic BP and Diastolic BP in 2 categories should be designated to the higher BP category.

BP indicates blood pressure (based on an average of  $\geq 2$  careful readings obtained on  $\geq 2$  occasions).

Source: ACC/AHA 2017 High Blood Pressure Clinical Practice Guideline



NEVER diagnose or treat yourself based on your readings. ALWAYS consult with your physician.

## 1.2 Blood Pressure Measurement Tips

To help ensure an accurate reading, follow these directions:

- Stress raises blood pressure. Avoid taking measurements during stressful times.
- · Measurements should be taken in a quiet place.
- It is important to take measurements at the same times each day.
   Taking measurements in the morning and in the evening is recommended.
- Remember to have a record of your blood pressure and pulse readings for your physician. A single measurement does not provide an accurate indication of your true blood pressure. You need to take and record several readings over a period of time.

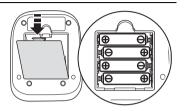


- Avoid bathing, drinking alcohol or caffeine, smoking, exercising and eating for at least 30 minutes before taking a measurement.
- Rest for at least 5 minutes before taking a measurement.

## 2. Preparation

## 2.1 Battery Installation

- 1. Remove the battery cover.
- 2. Insert 4 "AA" batteries as indicated in the battery compartment.



3. Replace the battery cover.

#### Notes:

- When the low battery symbol (r) appears on the display, turn the monitor
  off, then replace all batteries at the same time. Long life alkaline batteries
  are recommended.
- The measurement values continue to be stored in memory even after the batteries are replaced.
- Disposal of used batteries should be carried out in accordance with local regulations.

# 3. Using the Monitor

## 3.1 Applying the Arm Cuff

**↑** Caution

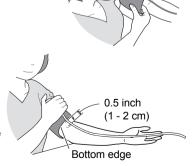
Remove tight-fitting, thick clothing from your arm while taking a measurement.

1. Insert the air plug into the air jack securely.



Wrap the arm cuff firmly in place around your left upper arm.

The bottom edge of the arm cuff should be 0.5 inch (1 to 2 cm) above the elbow. Air tube is on the inside of your arm and aligned with your middle finger.



# **3.** Secure closed with the fabric fastener.



#### Notes:

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



The blood pressure can differ between the right arm and the left arm, and
the measured blood pressure values can be different. OMRON recommends
to always use the same arm for measurement. If the values between both
arms differ substantially, please check with your physician which arm to use
for your measurements.

## 3.2 How to Sit Correctly

To take a measurement, you need to be relaxed and comfortably seated, under comfortable room temperature. Avoid bathing, drinking alcohol or caffeine, smoking, exercising or eating 30 minutes before taking a measurement.

- Sit in a chair with your legs uncrossed and feet flat on the floor.
- · Sit with your back and arm supported.
- The arm cuff should be placed on your arm at the same level as your heart, with the arm resting comfortably on a table.



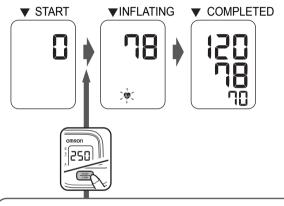
## 3.3 Taking a Measurement

#### Notes:

- To stop the measurement, press the START/STOP button once to deflate the arm cuff.
- · Remain still and guiet while taking a measurement.

#### 1. Press the START/STOP button.

The arm cuff will start to inflate automatically.



## 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.

#### Notes:

•The monitor will not inflate above 299 mmHg.

**⚠** Caution

Inflating to a higher pressure than necessary may result in bruising where the arm cuff is applied.

### 2. Remove the arm cuff.

3. Press the START/STOP button to turn the monitor off.

The monitor automatically stores the measurement result in its memory. It will automatically turn off after 2 minutes.

**Note:** Wait 2-3 minutes before taking another measurement. Waiting between measurements allows the arteries to return to the condition prior to taking a measurement.

## 3.4 Using the Memory Function

The monitor automatically stores the results up to 14 sets.

Note: If the memory is full, the monitor will delete the oldest value.

## To View the Measurement Values Stored in Memory

1. Press the button.

The Memory number appears for a second before the pulse rate is displayed.

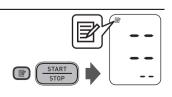
The newest set is numbered "1".



Press the button repeatedly to view the values stored in memory.

## To Delete All the Values Stored in Memory

- **1.** Press the button, while the memory symbol appears.
- While holding the button down, press the START/STOP button for more than 3 seconds.



Note: You cannot partially delete the values stored in memory.

# 4. Error Messages and Troubleshooting

## 4.1 Error Messages

| Error Display                | Cause                              | Solution   |  |
|------------------------------|------------------------------------|--|--|
| (3))                         | Irregular heartbeats are detected. | Remove the arm cuff. Wait 2 - 3 minutes and then take another measurement. Repeat the steps in section 3.3. If this error continues to appear, contact your physician. |  |
| The batteries are exhausted. |                                    | Carefully read and repeat the steps in section 3.3.  |  |
|                              |                                    | You should replace the batteries with new ones ahead of time. Refer to section 2.1.  |  |
|                              |                                    | You should replace the batteries with new ones at once. Refer to section 2.1.  |  |
|                              | Air plug disconnected.             | Insert the plug securely.<br>Refer to section 3.1.   |  |
| E 1                          | Arm cuff is applied too loosely.   | Apply the arm cuff tighter.<br>Refer to section 3.1.   |  |
|                              | Air is leaking from the arm cuff.  | Replace the arm cuff with a new one. Refer to section 5.5.   |  |

| Error Display Cause |   | Solution  |  |
|---------------------|---|---|--|
|                     | Movement during measurement   | Repeat measurement. Remain still and do not talk during measurement. Refer to section 3.3.  |  |
| E2                  | and the arm cuff has not been inflated sufficiently.  | If "E2" appears repeatedly, inflate the arm cuff manually until it is 30 to 40 mmHg above your previous measurement result. Refer to section 3.3.     |  |
| E3                  | The arm cuff was inflated exceeding the maximum allowable pressure, and then deflated automatically when inflating the arm cuff manually. | Do not touch the arm cuff and/or bend the air tube while taking a measurement. Do not inflate the arm cuff more than necessary. Refer to section 3.3. |  |
| E4                  | Movement during measurement.  | Repeat measurement. Remain still and do not talk during measurement. Refer to section 3.3.  |  |
| E5                  | Clothing is interfering with the arm cuff.  | Remove any clothing interfering with the arm cuff. Refer to section 3.1.  |  |
| Er                  | Device error.   | Contact Customer Service.   |  |

# 4.2 Troubleshooting

| Problem   | Cause and Solution   |
|---|--|
| No power.<br>No display appears on the monitor. | Replace all batteries with new ones. Check the battery installation for proper placement of the battery polarities. Refer to section 2.1.                                    |
| Measurement values appear too high or too low.  | Blood pressure varies constantly. Many factors including stress, time of day, and how you wrap the cuff, may affect your blood pressure. Review section 1.2 and section 3.3. |

## 5. Maintenance

#### 5.1 Maintenance

To protect your monitor from damage, please follow the directions below:

- Changes or modifications not approved by the manufacturer will void the user warranty.
- DO NOT disassemble or attempt to repair this monitor or other components. This may cause an inaccurate reading.

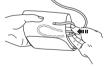
## 5.2 Storage

Store your monitor and other components in a clean, safe location.

- 1. Unplug the air plug from the air jack.
- 2. Gently fold the air tube into the arm cuff.

#### Note:

• Do not bend or crease the air tube excessively.



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.

## 5.3 Cleaning

- · Do not use any abrasive or volatile cleaners.
- Use a soft dry cloth or a soft cloth moistened with neutral soap 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 Disposal

- Dispose of your monitor, other components and optional accessories according to applicable local regulations.
- Unlawful disposal may cause environmental pollution.

## 5.5 Optional Medical Accessories

Arm circumference 7" - 9" (17 - 22 cm) Arm cuff

Arm circumference 9" - 17" (22 - 42 cm)



CD-CS9 (Model: HEM-CS24)



CD-WR17 (Model: HEM-RML31)

## 6. Specifications

Model REF HEM-7121-Z2 BP7100

Display LCD digital display

Cuff pressure range Pressure: 0 to 299 mmHa Pressure: 20 to 280 mmHa Measurement range

Pulse: 40 to 180 beats / min.

Accuracy Pressure: +3 mmHa

Pulse: ± 5% of display reading

Inflation Fuzzy-logic controlled by electric pump Deflation Automatic pressure release valve

Measurement method Oscillometric method

IP classification IP 20

Rating DC6 V 4 W

4 "AA" batteries 1.5 V Power source

Battery life Approximately 1000 measurements (using new alkaline batteries)

Durable period Monitor: 5 years (Service life) Cuff: 5 years

+50 °F to +104 °F (+10 °C to + 40 °C) / 15 to 90% RH Operating conditions

(non-condensing) / 800 to 1060 hPa

Storage / Transport -4 °F to +140 °F (-20 °C to +60 °C) / 10 to 90% RH (non-condensing) conditions

Weight Monitor: Approximately 8.8 oz. (250 g) not including batteries

Arm cuff: Approximately 6 oz. (170 g)

Monitor: Approximately 4.0" (w)  $\times$  3.1" (h)  $\times$  5.1" (l) Dimensions  $(103 \text{ mm} \times 80 \text{ mm} \times 129 \text{ mm})$ 

> Arm cuff: Approximately 5.7" × 23.4" (air tube: 29.5")  $(145 \text{ mm} \times 594 \text{ mm (air tube: } 750 \text{ mm)})$ 7" to 17" (170 to 420 mm) (included arm cuff 9" to 17" (220 to 420

Cuff circumference applicable to the

Memory

mm)) monitor

Up to 14 Readings

Monitor, arm cuff, instruction manual, quick start quide Contents

Applied part Type BF (arm cuff)

Protection against Internally powered ME equipment

electric shock

#### Notes:

- 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 monitor has not been validated for use on pregnant patients.
- IP classification is degrees of protection provided by enclosures in accordance with IEC 60529. This monitor is protected against solid foreign objects of 12.5 mm diameter and greater such as a finger.

#### Pictogram on the Product

| ART. O | Type of marks used to indicate the correct positioning of the artery for the upper arm |  |  |
|--------|--|--|--|
|        | Identifies the type of cuff that is compatible with the blood Pressure Monitor         |  |  |

## 7. FCC Statement

#### FCC CAUTION

Changes or modifications not expressly approved by the party responsible for compliance could void the user's authority to operate the equipment.

#### Note:

This equipment 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 equipment generates, uses and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation. If this equipment 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.

## 8. Limited Warranty

Your BP7100 Automatic Blood Pressure Monitor, excluding the arm cuff, is warranted to be free from defects in materials and workmanship appearing within 2 years from the date of purchase, when used in accordance with the instructions provided with the monitor. The arm cuff is warranted to be free from defects in materials and workmanship appearing within 1 year from the date of purchase when the monitor is used in accordance with the instructions provided with the monitor. The above warranty extends only to the original retail purchaser, and only to products purchased from an Omron authorized seller who is subject to and follows Omron's quality control standards, unless otherwise prohibited by law. We will, at our option, replace without charge any monitor or arm cuff covered by the above warranty. Replacement is our only responsibility and your only remedy under the above warranty.

To obtain warranty service contact Customer Service by calling **1-800-634-4350** for the address of the inspection center and the return shipping and handling fee.

Enclose the original printed receipt. Include a letter, with your name, address, phone number, and description of the specific problem. Pack the product carefully to prevent damage in transit. Because of possible loss in transit, we recommend insuring the product with return receipt requested.

THE FOREGOING IS THE SOLE WARRANTY PROVIDED BY OMRON IN CONNECTION WITH THIS PRODUCT, AND OMRON HEREBY DISCLAIMS ANY OTHER WARRANTIES, EXPRESS OR IMPLIED, INCLUDING IMPLIED WARRANTIES OF MERCHANTABILITY AND FITNESS FOR A PARTICULAR PURPOSE. IMPLIED WARRANTIES AND OTHER TERMS THAT MAY BE IMPOSED BY LAW, IF ANY, ARE LIMITED IN DURATION TO THE PERIOD OF THE ABOVE EXPRESS WARRANTY.

OMRON SHALL NOT BE LIABLE FOR LOSS OF USE OR ANY OTHER SPECIAL, INCIDENTAL, CONSEQUENTIAL OR INDIRECT COSTS, EXPENSES OR DAMAGES.

This warranty provides you with specific legal rights, and you may have other rights that vary by jurisdiction. Because of special local requirements, some of the above limitations and exclusions may not apply to you.

#### FOR CUSTOMER SERVICE

Visit our web site at: www OmronHealthcare com Call toll free:

1-800-634-4350

## 9. Guidance and Manufacturer's Declaration

OMRON Automatic Blood Pressure Monitor
Information for accompanying documents in the scope of IEC60601-1-2:2007
Model: BP7100

#### Important information regarding Electromagnetic Compatibility (EMC)

With the increased number of electronic devices such as PCs and mobile (cellular) telephones, medical devices in use may be susceptible to electromagnetic interference from other devices. Electromagnetic interference may result in incorrect operation of the medical device and create a potentially unsafe situation. Medical devices should also not interfere with other devices.

In order to regulate the requirements for EMC (Electromagnetic Compatibility) with the aim to prevent unsafe product situations, the IEC60601-1-2 standard has been implemented. This standard defines the levels of immunity to electromagnetic interferences as well as maximum levels of electromagnetic emissions for medical devices.

Medical devices manufactured by OMRON Healthcare conform to this IEC60601-1-2:2007 standard for both immunity and emissions.

Nevertheless, special precautions need to be observed:

- The use of accessories and cables other than those specified by OMRON, with the exception
  of cables sold by OMRON as replacement parts for internal components, may result in
  increased emission or decreased immunity of the device.
- The medical devices should not be used adjacent to or stacked with other equipment.
   In case adjacent or stacked use is necessary, the medical device should be observed to verify normal operation in the configuration in which it will be used.
- Refer to further guidance below regarding the EMC environment in which the device should be used.
- The MEDICAL ELECTRICAL EQUIPMENT BP7100 needs special precautions regarding EMC and needs to be installed and put into service according to the EMC information provided in this documentations.
- The Essential Performance of the BP7100 is to measure a blood pressure and a pulse rate and using the memory function.

The BP7100 may be interfered with by other equipment, even if that other equipment complies with CISPR EMISSION requirements.

#### Guidance and manufacturer's declaration - electromagnetic emissions

OMRON BP7100 is intended for use in the electromagnetic environment specified below. The customer or the user of this OMRON BP7100 should assure that it is used in such environment.

| Emissions test  | Compliance      | Electromagnetic environment - guidance  |
|---|-----------------|---|
| RF emissions<br>CISPR 11                                      | Group 1         | The OMRON BP7100 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<br>CISPR 11                                      | Class B         |   |
| Harmonic<br>emissions<br>IEC 61000-3-2                        | Not Applicable. | The OMRON BP7100 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 |
| Voltage<br>fluctuations/<br>flicker emissions<br>IEC61000-3-3 | Not Applicable. | purposes.   |

#### Guidance and manufacturer's declaration - electromagnetic immunity

OMRON BP7100 is intended for use in the electromagnetic environment specified below. The customer or the user of this OMRON BP7100 should assure that it is used in such environment.

| Immunity test                                       | IEC 60601 test<br>level  | Compliance level           | Electromagnetic environment - guidance   |
|---|--|----------------------------|--|
| Electrostatic discharge<br>(ESD)<br>IEC 61000-4-2   | ±6 kV contact<br>±8 kV air   | ±6 kV contact<br>±8 kV air | Floor should be wood, concrete, or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30 %. |
| Electrical fast<br>transient/burst<br>IEC 61000-4-4 | ±2 kV for power<br>supply lines<br>±1 kV for<br>input/output lines | Not Applicable.            | Not Applicable.  |
| Surge<br>IEC 61000-4-5                              | ±1 kV line(s) to<br>line(s)<br>±2 kV line(s) to<br>earth           | Not Applicable.            | Not Applicable.  |

| Voltage dips, short<br>interruptions and<br>voltage variations on<br>power supply inputlines<br>IEC 61000-4-11 | <5 % Uτ (>95 % dip<br>in Uτ)<br>for 0.5 cycle<br>40 % Uτ (60 % dip<br>in Uτ)<br>for 5 cycles<br>70 % Uτ (30 % dip<br>in Uτ)<br>for 25 cycles<br><5 % Uτ (95 % dip<br>in Uτ) | Not Applicable. | Not Applicable.   |
|--|---|-----------------|---|
|  | for 5 sec.  |                 |   |
| Power frequency<br>(50/60 Hz) magnetic<br>field<br>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.                                  |   |                 |   |

#### Guidance and manufacturer's declaration - electromagnetic immunity

OMRON BP7100 is intended for use in the electromagnetic environment specified below. The customer or the user of this OMRON BP7100 should assure that it is used in such environment.

| Immunity test                 | IEC 60601 test<br>level         | Compli<br>ance<br>level | Electromagnetic environment - guidance  |
|-------------------------------|---------------------------------|-------------------------|---|
|                               |                                 |                         | Portable and mobile RF communications equipment should be used no closer to any part of the OMRON BP7100 including cables, than the recommended separation distance calculated from the equation appropriate to the frequency of the transmitter.   |
|                               |                                 |                         | Recommend separation distance   |
| Conducted RF<br>IEC 61000-4-6 | 3 V rms<br>150 kHz to 80<br>MHz | Not<br>Applica<br>ble.  | Not Applicable.   |
| Radiated RF<br>IEC 61000-4-3  | 3 V/m<br>80 MHz to 2.5<br>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. Interference may occur in the vicinity of equipment marked with the following symbol: |

Note1: At 80 MHz and 800 MHz, the higher frequency range applies.

Note2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects, and people.

<sup>&</sup>lt;sup>a</sup> Field strengths from fixed transmitters, such as base stations for radio (cellular/ cordless) telephones and land mobile 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 OMRON BP7100 is used exceeds the applicable RF compliance level above, the OMRON BP7100 should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as reorienting or relocating the OMRON BP7100.
<sup>b</sup> Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.

# Recommended separation distance between portable and mobile RF communications equipment and the OMRON BP7100

OMRON BP7100 is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of this OMRON BP7100 can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the OMRON BP7100 as recommended below, according to the maximum output power of the communications equipment.

| Output Power of Transmitter in | Separation distance according to frequency of transmitter in meter |                      |                    |  |
|--------------------------------|--|----------------------|--------------------|--|
| Watt                           | 150 kHz to 80 MHz<br>Not Applicable.                               | 80 MHz to 800 MHz    | 800 MHz to 2.5GHz  |  |
|                                |  | $d = 1.2 \ \sqrt{P}$ | $d = 2.3 \sqrt{P}$ |  |
| 0.01                           | Not Applicable.  | 0.12                 | 0.23               |  |
| 0.1                            |  | 0.38                 | 0.73               |  |
| 1                              |  | 1.2                  | 2.3                |  |
| 10                             |  | 3.8                  | 7.3                |  |
| 100                            |  | 12                   | 23                 |  |

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

Note: At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies. Note: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects, and people.



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Distributed by: OMRON HEALTHCARE, INC. 2895 Greenspoint Pkwy. Hoffman Estates, IL 60169 USA

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