SenseWear®

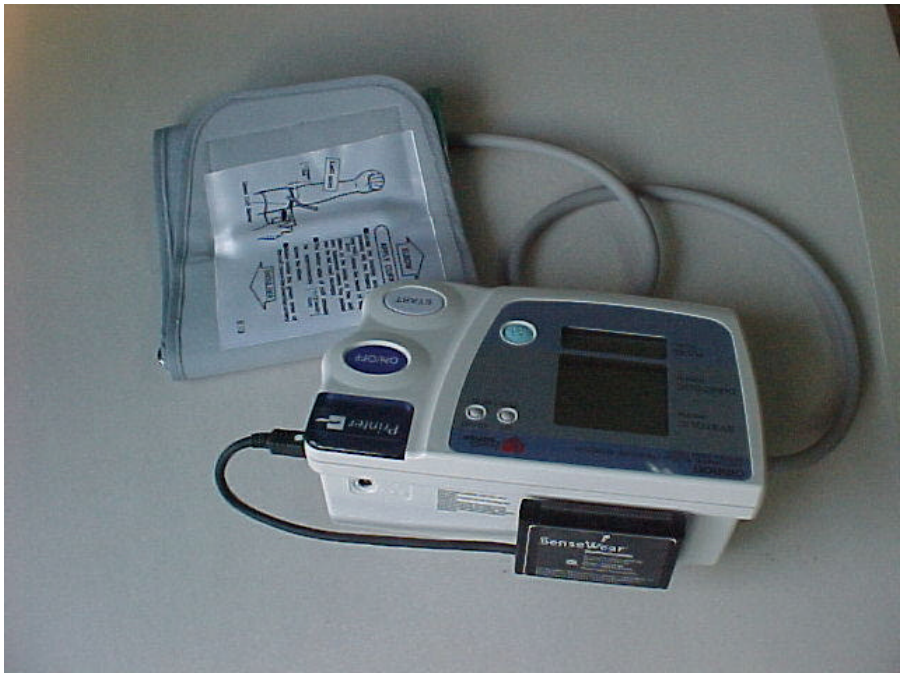
BioTransceiver for Blood Pressure Monitoring

Supplemental Instructions

Please consult the Omron® Instruction Manual for instructions on setting up the monitor and taking a reading.

NOTE: The printer shown in the Omron manual has been replaced with the BioTransceiver which stores the readings taken and forwards them to any nearby Wireless Gateway. This enables your monitoring service to immediately receive and review your readings.

The BioTransceiver should remain plugged into the Omron® Automatic Blood Pressure Monitor at all times as shown below.



Certifications

Disclaimer

This product is not intended to be a medical device or a substitute for proper medical diagnosis, care, or treatment related to your condition. Seek professional consultation if you have any questions related to the readings provided by this product.

FCC statement

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 separate from the receiver.
- Consult the dealer or an experienced radio/TV technician for help.

This device complies with part 15 of the FCC Rules. 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.

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

FCC 47CFR 15C TCB - 47 CFR Part 15 Subpart C Intentional Radiator Certification Test

FCC 47CFR 15B clB - 47 CFR Part 15 Subpart B Unintentional Radiator Class B Verification

IEC 60601-1-1 (2000-12) - Medical Electrical Equipment - Part 1-1: General Requirements for Safety - Collateral Standard: Safety Requirements for Medical Electrical Systems Second Edition