RELATED ITEMS



- Replacement Adult Size Cuff #526019
- Large Adult Size Cuff #526020
- Hopkins® Flex Temp Thermometer #579420
- Hopkins[®] Wave[™] Pulse Oximeter #594029

User Manual

Hopkins® IMPACT Digital Arm BP



Thank you for selecting the Hopkins[®] IMPACT Digital Arm BP #526018. To use this BP monitor correctly and safely, please read this entire manual thoroughly and keep it for your reference.





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Thank you for selecting the Hopkins® Digital IMPACT Arm BP (Item #526018). This monitor features blood pressure measurement, pulse rate measurement, and an auto-save function.

Readings taken by the #526018 are equivalent to those obtained by a trained observer using the cuff and stethoscope auscultation method.

This manual contains important safety and care information, and provides step by step instructions for using this product. Read this manual thoroughly before using this product

- Measures Systolic Blood Pressure
- Measures Diastolic Blood Pressure
- Measures Pulse Rate

♥ Safety Information

The following symbols may be used in the user manual, product labeling, or on other components. They are the requirement of standard and use.

$\mathbf{\Lambda}$	Caution: consult accompanying documents	Ŕ	Type B applied part
C €0123	CE Mark: conforms to essential requirements of the Medical Device Directive 93/42/EEC	X	DISPOSAL: Do not dispose this product as unsorted municipal waste. Collection of such waste separately for special treatment is necessary.
***	Manufacturer		Direct current
SN	Specifies serial number	EC REP	Authorized Representative in the European Community

This device is for adult use only.

This device is intended for non-invasive measuring and monitoring of arterial blood pressure. It is not intended for use on extremities other than the arm or for functions other than obtaining a blood pressure measurement and pulse rate.

Do not confuse self-monitoring with self-diagnosis. This unit allows you to monitor your blood pressure. Do not begin or end medical treatment based solely on self-monitoring, consult a physician for treatment advice.

If you are taking medication, consult your physician to determine the most appropriate time to measure your blood pressure. Never change a prescribed medication without consulting your physician. This unit is not suitable for continuous monitoring during medical emergencies or operations. If the cuff pressure exceeds (300 mmHg), the unit will automatically deflate. Should the cuff not deflate when pressures exceeds (300 mmHg), detach the cuff from the arm and press the START/STOP button to stop inflation.

To avoid measurement errors, carefully read this manual before using this device.

This device is not AP/APG equipment and not suitable for use in the presence of a flammable anesthetic mixture with air, or with oxygen, or nitrous oxide.

Please avoid strong electromagnetic fields as they may interfere with or damage this device.

Always check this device to make sure that it is in proper working condition before use.

 Retains historic records of up to 60 previous measurements

• Large 5.5" x 1.5" bright LCD display

♥ LCD Display



SYMBOL	DESCRIPTION	EXPLANATION	
SYS	Systolic BP	High pressure result	
DIA	Diastolic BP	Low pressure result	
Pul/min	Pulse	Pulse per Minute (Beats per Minute)	
lo	Low Battery	Batteries are low and need to be replaced	
mmHg	Millimeters Mercury	Measurement unit of blood pressure	
M <mark>. 18∕80</mark>	Memory	If "M" shows, the displayed measurement value is from the memory of recorded measurements (See page 9 for instructions)	
Inflating Inflating The unit is inflating with air t required level of pressure		The unit is inflating with air to obtain the required level of pressure	
►	Deflating	The unit is deflating and the air is being exhausted from the cuff	
ям 8:59	Time (hour:minute)	The current time (set by user)	
Ţ	Shock Warning	Inaccurate readings may occur if unit is hit, dropped, shaken, or otherwise shocked	
AVG	Average	The average of the blood pressure measurements	
0,	Recalling	The stored records are being retrieved	
	Irregular Heartbeat	Irregular heartbeat (arrhythmia) detected	
Normal BP Category Indicator (Blood pressure level reading (See page 13 for instructions)	

Monitor Components



BATTERY COMPARTMENT



♥ Parts Included









Adult Digital Cuff Cuff Circumference: 8.5" - 12.5" (22cm - 32cm) Four AAA Batteries

BEFORE YOU START



♥ Installing the Batteries

- 1. Slide off the battery cover on the back of the unit.
- **2.** Install the batteries by matching the correct polarity, as shown above. *Always use the correct battery type (4 alkaline AAA size)*
- **3.** Replace the battery cover.

Replace the batteries when any of the following occurs:
•The shows
•The display dims
•The display does not light up

- If this device will be stored for an extended period of time, please remove the batteries in order to avoid corrosion.
- Used batteries are harmful to the environment, so please dispose of appropriately. Remove the old batteries from the device and follow your local recycling guidelines
- Battery corrosion is not covered under your three year limited warranty.

♥ Setting the Date and Time

It is important to set the clock before using your blood pressure monitor so that a time stamp can be assigned to each record stored in the memory. (The year can be set to anywhere between 2000 and 2050.)

1. When the unit is off, press and hold the "SET" button for 3 seconds to enter the mode settings.



2. Press the "MEM" button to advance to the current [YEAR].



3. Press "SET" when you have reached the correct year to save your choice and advance to the next step.



4. Repeat steps 2 and 3 to set the [MONTH] and [DAY]



5. Repeat steps 2 and 3 to set the [HOUR] and [MINUTE]



6. Once you have set the clock, the screen will read "done" and the monitor will then turn off automatically.



♥ Positioning the Cuff

1. Make sure the air tube is connected securely to the unit.

2. Fasten the cuff on your left upper arm and position it so that the tube toward the inside of arm in line with your pinky finger.

3. The cuff should be snug but not too tight. You should be able to insert one finger between the cuff and your arm.

4. Sit comfortably with your left arm resting on a flat surface.

If the index arrows fall outside the area marked "OK", please select a different size cuff.

♥ Tips for Accuracy

- •You should always rest for at least 5 minutes before measuring your blood pressure.
- •Wait at least 3 minutes between measurements. This allows your blood circulation to recover.
- •For a meaningful comparison, try to take blood pressure measurements under similar conditions. For example, take daily measurements at approximately the same time, on the same arm, or as directed by a physician.



♥ Start the Measurement

1. Press the "START/STOP" button to turn on the monitor. The whole measurement will take place automatically.

The LCD Display will power up.

The unit will adjust to zero and calibrate automatically. (The unit will calibrate every time it is turned on.)

The unit will inflate and begin to take the measurement.

Once the measurement has been taken, the unit will deflate, display the reading and the category of the reading, and store the reading in the memory.

2. Press the "START/STOP" button to turn the monitor off, otherwise the monitor will power off automatically within 1 minute.



Recalling Recorded Measurements

1. Press the "MEM" button to display the average of the records.



AVG will display on the right side of the screen



2. Press either the "MEM" button or the "SET" button to toggle between the records.



The order of the record, date, and time will display alternately.



The most recent record M 1/60 is shown first. Each new measurement is assigned to the first (1) record. All other records are pushed back one digit (e.g., 2 becomes 3, and so on), and the last record M 60/60 is dropped from the list.

Deleting a Record from the Memory

You can delete all the measurement results in the memory by following the steps below:

1. Press and hold the "MEM" button for 3 seconds – the display will alternately flash "DEL ALL" and "DEL ONE".

If you don't want to delete any of the records, press the "START/ STOP" button to escape.

2. To delete the most recent record, wait until the display shows "DEL ONE", then press the "SET" button to delete the latest record.

3. To delete all of the records at once, repeat step 1 and when the display shows "DEL ALL" press the "SET" button.

The word "DONE" will appear at the bottom of the screen once the action has been completed.

4. If there is no record. the display will be blank.









Tips for Measurement

Measurements taken under some circumstances may cause the monitor to provide inaccurate readings:



♥ Care and Storage

In order to get the best performance out of your blood pressure monitor, please follow the instructions below:



Avoid direct sunlight.



Avoid shaking or dropping the monitor.



Never use a wet cloth to remove dirt. Always use a clean, dry cloth.



Do not submerge in water or any other liquid.



Store at room temperature in a dry, clean environment.



Do not wash the monitor or the cuff.

♥ What are Systolic and Diastolic Pressure?





When the ventricles contract to pump blood out of the heart, the blood pressure reaches its highest pressure in the cycle. This is known as *systolic pressure*. The lowest blood pressure in the cycle occurs when the heart relaxes between heartbeats, and this is known as *diastolic pressure*.

The systolic pressure is the top number in your blood pressure reading, and the diastolic is the bottom number.

What is a Normal Blood Pressure?

The following charts illustrate the blood pressure classification level established by the World Health Organization (WHO) and International Society of Hypertension (ISH) in 1999.



BP Level (mmHg)	Optimal BP	Normal BP	Mild Hypertension	Moderate Hypertension	Severe Hypertension
SYS	<120	120~139	140~159	160~179	>180
DIA	<80	80~89	90~99	100~109	>110

Only a physician can tell you your normal blood pressure range and the point at which you are at risk. Consult your physician to obtain these values.

If the measurements taken with these products fall outside the range, consult your physician.

ABOUT BLOOD PRESSURE

Why does my blood pressure fluctuate throughout the day?

1. Individual blood pressure readings can naturally vary throughout the day, so for the best accuracy, take your readings the same way each time. (See *Tips for Accuracy* on page 7.)

2. Blood pressure readings can fluctuate if you are taking medications.

2. If you are taking multiple readings, wait at least 3 minutes between readings for best accuracy.

Why is the reading I get from a healthcare provider different from the ones I take at home?

Your blood pressure can be different even during a 24 hour period because of factors like the weather, emotional state, recent exercise or exertion, etc. Your readings may be elevated in a healthcare setting because of anxiety caused by the "white coat effect" and you may get lower readings at home where you are calmer.

Which arm should I use to take my blood pressure?

Blood pressure readings can be taken on either the right or left arm, but it is recommended to take readings using the *left arm*. Consult your physician if you think you need to use your right arm instead. Whichever arm you use, it is important to take your blood pressure using the same arm every time you take a reading.



When taking your blood pressure at home you should always make sure the cuff is fastened and placed properly, and that the cuff is not too tight or too loose.

Make sure you are not overly anxious or stressed as this will elevate your readings. It is recommended to take 2-3 deep breaths before beginning to take your blood pressure.



This section includes a list of error messages and frequently asked questions for problems you may encounter with your blood pressure monitor. If this product is not operating as you think it should, please check here before arranging for service.

PROBLEM	SYMPTOM	CHECK THIS	REMEDY
	Display is dim or	Are the batteries dead?	Replace with new batteries.
No Power	will not light up	Are the batteries inserted correctly?	Insert the batteries correctly.
Low Batteries	Lo shows on the display	Are the batteries low?	Replace with new batteries.
	"E 1" Shows on the display	The monitor detected that the cuff is not secure	Refasten the cuff and take a new reading.
	"E 2" Shows on the display	The monitor detected that the cuff is too tight	Refasten the cuff and take a new reading.
	"E 3" Shows on the display	The monitor detected that the pressure of the cuff is in excess	Relax for a moment and then take a new reading.
Error Message	"E 10" or "E 11" Shows on the display	The monitor detected motion while measuring	Movement can affect the measurement. Relax for a moment and then take a new reading. Try to remain still for the second reading.
	"EEXX" Shows on the display	A calibration error oc- curred	Take a new reading. If the problem persists, contact customer service for further assistance. Refer to page 21 for warranty and contact information.

¥	Complied	European	Standards	List
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Risk management in the application of medical devices	EN/ISO 14971:2007
Graphical symbols for labeling medical devices	EN 980: 2008
Medical equipment manufacturers to provide information	EN 1041: 2008
Medical equipment Part1-1: General Requirements for Safety Collateral Standard: Safety requirements for medical electrical systems	EN 60601-1: 1990+A1+A2+A13
Non-Invasive blood pressure Part 1: General Requirements	EN 1060-1: 2001/A1:2002
Non-Invasive blood pressure Part 3: Supplementary requirements for electromechanical blood pressure measuring system	EN 1060-3: 1997/A1:2005
Automatic Blood Pressure Monitor overall system interventional accuracy of the testing process	EN 1060-4: 2004
Medical electrical equipment Part 1-2: Basic safety and essential performance of the general requirements Collateral Standard: Electromagnetic compatibility requirements and tests	EN 60601-1-2:2001+A1: 2006

Table 1

Guidance and manufacture's declaration - electromagnetic emissions - for all EQUIPMENT and SYSTEMS

Guidance and manufacture's declaration – electromagnetic emission					
The 526018 is intended for use in the electromagnetic environment specified below. The customer of the user of the 526018 should assure that it is used in such and environment.					
Emission Test	Emission Test Compliance Electromagnetic Environment - Guid				
RF emissions CISPR 11	Group 1	The 526018 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 emission CISPR 11	Group B	The 526018 is suitable for use in all establishments other than domestic 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	Not applicable				
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Not applicable				

Power Supply	Four AAA Alkaline Batteries (included)	
Display Model	Digital LCD V.A. 5.5" x 1.5"	
Measurement Mode	Oscillographic Testing Mode	
Measurement Range	Pressure: 0 - 300mmHg Pulse Value: 40 - 199 beats/minute	
Accuracy	Pressure: Within ± 3mmHg Pulse Value: ± 5%	
Normal Working Conditions	Temperature: Approx. 41°F - 104°F Relative Humidity ≤ 80%	
Cuff Circumference	Approx. 8.5" - 16.5" (22cm - 32cm)	
Weight	Approx. 12.2 oz (excluding batteries)	
External Dimensions	Approx. 7" x 3.75" x 1.5"	
Parts Included	Four AAA Alkaline Batteries, an Adult Cuff, and a User Manual	
Mode of Operation	Continuous Operation	
Degree of Protection	Type B Applied Part	
Protection Against Ingress of Water	IPX-0	

WARNING: No modification of this equipment is allowed

Table 2

Guidance and manufacture's declaration - electromagnetic immunity - for all ME EQUIPMENT and ME SYSTEMS

Guidance and manufacture's declaration – electromagnetic immunity						
The 526018 is intended for use in the electromagnetic environment specified below. The customer or the user of 526018 should assure that it is used in such an environment.						
Immunity Test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance			
Electrostatic discharge (ESD) IEC 601000-4-2	±6 kV contact ±8 kV air	±6 kV contact ±8 kV air	Floors should be wood, concrete or ceramic tile. If floor 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	Mains power quality should be that of a typical commercial or hospital environment.			
Surge IEC 6000-4-5	±1 kV line(s) to line(s)	±1 kV differential mode	Mains 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	<5% U _T (>95% dip in U _T) for 0.5 cycle	Mains power quality should be that of a typical commercial			
	40% U _T (>60% dip in U _T) for 5 cycles	40% U _T (>60% dip in U _T) for 5 cycles	or hospital environment. If the user of the 526018 requires continued operation during			
	70% U _T (>30% dip in U _T) for 25 cycle	70% U _T (>30% dip in U _T) for 25 cycle	power mains interruptions, it is recommended that the 526018 be powered from an			
	5% U _T (>95% dip in U _T) for 5 seconds	5% UT (>95% dip in UT) for 5 seconds	uninterruptible power supply or a battery.			
Power frequency (50Hz) magnetic field IEC 61000-4-8	3A/m	3A/m	Power frequency magnetic fields should be at levels char- acteristic 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 4

Guidance and manufacture's declaration – electromagnetic immunity – for ME EQUIPMENT and ME SYSTEMS that are not LIFE-SUPPORTING

Guidance and man	ufacture's declaration – el	lectromagnetic immu	nity		
The 526018 is intended for use in the electromagnetic environment specified below. The customer or the user should assure that it is used in such an environment.					
Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance		
			Portable and mobile RF communications equipment should be used no closer to any part of the YS-6100, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.		
Conducted RF IEC 61000-4-6	3 Vrms 150 kHz to 80 MHz	3 Vrms	Portable and mobile RF communications equipment should be used no closer to any part of the including cables, than the recom- mended separation distance calculated from the equation applicable to the frequency of the transmitter.		
Radiated RF IEC 61000-4-3	3 V/m 80 MHz to 2.5 GHz	3 V/m	Recommended separation distance d = 1.167 \sqrt{P} d = 1.167 \sqrt{P} 80 MHz to 800 MHz d = 2.333 \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 recom- mended 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.		
			Interference may occur in the vicinity of equip- ment 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 the ELE007839V1 is used exceeds the applicable RF compliance level above, the ELE007839V1 should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as reorienting or relocating the ELE007839V1.

b Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.

Table 6

Recommended separation distances between portable and mobile RF communications equipment and the EQUIPMENT or SYSTEM – for ME EQUIPMENT or ME SYSTEM that are not LIFE-SUPPORTING

Recommended separation distances between portable and mobile RF communications equipment and the ELE007839V1 Fitness Equipment.

The 526018 is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the 526018 can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the 526018 as recommended below, according to the maximum output power of the communications equipment.

Rated maximum output power of transmitter W	Separation distance according to frequency of transmitter m				
	150 kHz to 80MHz d = 1.167 √P	80MHz to 800 MHz d = 1.167 √P	800 MHz to 2,5 GHz d = 2.333 √P		
0,01	0.167	0.167	0.233		
0,1	0.369	0.369	0.738		
1	1.167	1.167	2.333		
10	3.690	3.690	7.388		
100	11.67	11.67	23.330		

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 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.

Warranty & Registration Information

Your Hopkins[®] Digital BP comes with a 3 year limited warranty. This BP monitor is warranted against any defects in material and workmanship for 3 years from the date of purchase. If you experience a material or manufacturing defect at any time, repairs will be made without charge upon the return of the instrument to Hopkins Medical Products, except in cases of obvious abuse or accidental damage.

NOTE If your BP monitor will be stored for an extended period of time, please remove the batteries in order to avoid corrosion. Battery corrosion is not covered under your 3 year limited warranty.



To register your new Hopkins® Digital BP for its warranty with Hopkins Medical Products, please visit **www.hmponline.com/registration** or scan this QR code with your smartphone and complete the registration process. (*Reorder #526018*)

Contact Information

For more information about our products, please visit our website: *www.hmponline.com*

For customer service, please call us at 1-800-835-1995 M-F 8:30 AM - 5 PM EST

Hopkins Medical Products

5 Greenwood Place, Baltimore, MD 21208