








US Blood Pressure Validated Device Listing (VDL)




The ultimate judgment regarding whether a BP measurement device meets the requisite VDL Criteria rests with the Independent Review Committee and is not in any way determined or influenced by the AMA. The AMA does not receive funding from any device manufacturer or other third party in relation to the development of the VDL Criteria or VDL process. For more information, visit www.validateBP.org




Filters applied: **Home**




Brand/Device Type	Name And Model Number	Validation Protocol	Cuff Sizes	Populations Served
<div>ForaCare</div> <div>Home</div>	<div>Fora TN'G BP</div> <div>P80</div> <div></div>	<div>ANSI/AAMI/ISO 81060-2:</div> <div>2013</div>	Standard (24-43 cm)	General
<div>Smart Meter</div> <div>Home</div>	<div>iBloodPressure</div> <div>SMBP802-GS-001</div> <div></div>	<div>ANSI/AAMI/ISO 81060-</div> <div>2:2009</div>	Standard (22-42 cm)	General




Brand/Device Type	Name And Model Number	Validation Protocol	Cuff Sizes	Populations Served
Greater Goods Home	Greater Goods BP 0040, 0604, 0664 	ISO 81060-2:2018	Standard (22-42 cm)	General
BodyTrace Home	Blood Pressure Monitor BT105 	ANSI/AAMI/ISO 81060-2:2013	Adult (22-32 cm) Extended (22-42 cm)	General




Brand/Device Type	Name And Model Number	Validation Protocol	Cuff Sizes	Populations Served
CareSimple Home	Blood Pressure Monitor BT105 	ANSI/AAMI/ISO 81060-2:2013	Adult (22-32 cm) Extended (22-42 cm)	General
Transtek Home	Transtek Blood Pressure Monitor LS802-GS 	ISO 81060-2:2018	Extra Large (22-45 cm) Large (22-42 cm) Standard (22-32cm)	General
Transtek Home	Transtek Blood Pressure Monitor LS802-GP 	ISO 81060-2:2018	Extra Large (22-45 cm) Large (22-42 cm) Standard (22-32cm)	General




Brand/Device Type	Name And Model Number	Validation Protocol	Cuff Sizes	Populations Served
Omron Home	Automatic Upper Arm Blood Pressure Monitor HEM-9210T 	ANSI/AAMI/ISO 81060-2:2009	Extra Large (42-50 cm) Small (17-22 cm) Wide Range (22-42 cm)	General
Omron Home	Blood Pressure Monitor HEM-9200T 	ANSI/AAMI/ISO 81060-2:2009	Adult (22-42 cm)	General
Withings Home	Withings BPM Connect WPM05 	ANSI/AAMI/ISO81060-2:2013	Integrated (22-42 cm)	General



Brand/Device Type	Name And Model Number	Validation Protocol	Cuff Sizes	Populations Served
Microlife Home	WatchBP Home A BT BP3MX1-3C 	BHS Revised Protocol: 1993	Large (32-42 cm) Small (14-22 cm) Standard (22-32cm)	General
Microlife Home	WatchBP Home A BP3MX1-3 	BHS Revised Protocol: 1993	Large (32-42 cm) Small (14-22 cm) Standard (22-32cm)	General
Microlife Home	WatchBP Home N BP3MX1-4 	BHS Revised Protocol: 1993	Large (32-42 cm) Small (14-22 cm) Standard (22-42 cm)	General



Brand/Device Type	Name And Model Number	Validation Protocol	Cuff Sizes	Populations Served
Microlife Home	WatchBP Home BP3MX1-1 	BHS Revised Protocol: 1993	Large (32-42 cm) Small (14-22 cm) Standard (22-32cm)	General
Omron Home	Evolv® Wireless Upper Arm Blood Pressure Monitor BP7000 	ANSI/AAMI/ISO 81060-2: 2013	Adult (22-42 cm)	General
Omron Home	5 Series® Wireless BP7250 HEM-7311 	Validation Protocol(s): ANSI/AAMI/ISO 81060-2: 2009	Adult (22-42 cm)	General

Brand/Device Type	Name And Model Number	Validation Protocol	Cuff Sizes	Populations Served
Omron Home	5 Series® BP7200 HEM-7311 	Validation Protocol(s): ANSI/AAMI/ISO 81060-2: 2009	Adult (22-42 cm)	General
Omron Home	3 Series® BP7100 HEM-7311 	Validation Protocol(s): ANSI/AAMI/ISO 81060-2: 2009	Adult (22-42 cm)	General
Omron Home	Platinum Upper Arm BP5450 HEM-7311 	Validation Protocol(s): ANSI/AAMI/ISO 81060-2: 2009	Adult (22-42 cm)	General

Brand/Device Type	Name And Model Number	Validation Protocol	Cuff Sizes	Populations Served
Omron Home	Gold Upper Arm BP5350 HEM-7311 	Validation Protocol(s): ANSI/AAMI/ISO 81060-2: 2009	Adult (22-42 cm)	General
Omron Home	10 Series® Wireless BP7450 HEM-7320 	Validation Protocol(s): ANSI/AAMI/ISO 81060-2: 2009	Adult (22-42 cm)	General
Omron Home	7 Series® Wireless BP7350 HEM-7320 	Validation Protocol(s): ANSI/AAMI/ISO 81060-2: 2009	Adult (22-42 cm)	General

Brand/Device Type	Name And Model Number	Validation Protocol	Cuff Sizes	Populations Served
Omron Home	Bronze Upper Arm BP5100 HEM-7311 	Validation Protocol(s): ANSI/AAMI/ISO 81060-2: 2009	Adult (22-42 cm)	General
Omron Home	Silver Wireless BP5250 HEM-7320 	Validation Protocol(s): ANSI/AAMI/ISO 81060-2: 2009	Adult (22-42 cm)	General
Omron Home	Complete™ Wireless BP7900 HEM-7311 	Validation Protocol(s): ANSI/AAMI/ISO 81060-2: 2009	Adult (22-42 cm)	General

Brand/Device Type	Name And Model Number	Validation Protocol	Cuff Sizes	Populations Served
<div>A&D Medical</div> <div>Home</div>	<div>ULTRACONNECT Wireless Blood Pressure Monitor</div> <div>UA-1200BLE</div> <div></div>	<div>Validation Protocol(s): ANSI/AAMI/ISO 81060-2: 2009</div>	<div>Integrated (22-42 cm)</div>	<div>General</div>
<div>Hillrom-Welch Allyn</div> <div>Home</div>	<div>Welch Allyn Home® Blood Pressure Monitor, 1700 Series</div> <div>H-BP100SBP</div> <div></div>	<div>Validation Protocol(s): ANSI/AAMI/ISO 81060-2: 2009</div>	<div>Standard (22-42 cm)</div> <div>XL (40-54 cm)</div> <div>XS (15-24 cm)</div>	<div>General</div>

Brand/Device Type	Name And Model Number	Validation Protocol	Cuff Sizes	Populations Served
A&D Medical Home	Talking+ Blood Pressure Monitor UA-1030T UA-1020 	Validation Protocol(s): BHS Revised Protocol: 1993	Large (31-45 cm) Medium (23-37 cm) Small (16-24 cm) Smooth Fit (23-37 cm)	General
A&D Medical Home	Advanced Manual Inflate Blood Pressure Monitor UA-705 UA-705V, UA-704 	Validation Protocol(s): BHS Revised Protocol: 1993	Large (36-45 cm) Medium (23.8-36 cm)	General

NOTE: Additional devices may be added to the VDL in the future through successive application and review processes as more devices are submitted and reviewed by the Independent Review Committee. The listing currently contains devices that measure other biometrics, it is important to note that for those devices, only the BP measurement components have been reviewed for validation.

Clinical Disclaimer: Devices on this list for use at home are validated for clinical accuracy for the population tested and may not always provide accurate BP measurements for a specific individual. Whenever practical, patients are encouraged to bring their home BP devices to the physician's office or some other qualified health care facility for comparison with an accurate device.

Wrist Cuff Device Disclaimer:

Clinical guidelines call for the user of an upper arm blood pressure (BP) measurement device that has been validated for clinical accuracy for self-measured blood pressure (SMBP). However, a validated wrist cuff blood pressure (BP) device may be used in place of an upper arm device for certain clinical indications, specific patient needs, or additional circumstances (eg, upper arm cuff not suitable for patient arm circumference, unavailability of devices with the appropriate small, large or extra large size cuffs, medical conditions where upper arm devices cannot or should not be used).

Proper technique is especially important when using a wrist cuff device to obtain accurate readings. The device must be correctly placed over the radial artery and held at heart level when readings are taken, with limited movement or wrist flexion.

***Protocol Disclaimer:** Protocol did not require detailed reporting on arm and cuff sizes. Device met all requirements, but it is unclear if valid with larger arm circumferences.

****Welch-Allyn ProBP3400 recall information:** <https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfRes/res.cfm?ID=186903>

*The American Medical Association (“AMA”) does not receive funding from any third-party in relation to the development of the Validated Device Listing Criteria (“VDL™ Criteria”), and does not receive funding from any blood pressure (“BP”) device manufacturer or other third-party in relation to the development of the VDL process. No recommendation, promotion, or endorsement is implied or intended by the AMA (or any of the AMA’s affiliated or partner organizations) of any third-party organization, product, device, policy, or service. BP measurement devices are selected to appear on the VDL through an independent review process that determines which available BP measurement devices meet established VDL Criteria for the validation of clinical accuracy. An Independent Review Committee, composed of members who are experts in the BP field, assess whether a BP measurement device satisfies the VDL Criteria. The ultimate judgment regarding whether a BP measurement device meets the requisite VDL Criteria rests with the Independent Review Committee, and is not in any way determined or influenced by the AMA.