User Manual for Vista AVS®

REF
L500VA (with Stand and Basket)
L500VAC (Stand and Basket not included)

WALLACH®
SURGICAL DEVICES
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(Vista AVS)

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SECTION 1: Introduction
Thank you for choosing the Vista AVS® from Wallach Surgical Devices. We believe you have purchased the finest arterial examination system available.

Your total satisfaction is our highest priority. We strive to continually improve our products and services. Please contact us with any suggestions. We look forward to enjoying a long-term relationship with you!

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Important: Please read this manual carefully and become familiar with the features, operation, care, and safety requirements of the Vista AVS prior to use. Please note that while operating the Vista AVS, step-by-step instructions are shown on the display to assist you through the examination.

Wallach Surgical Devices provides general reimbursement information related to the diagnosis of peripheral arterial disease as an overview for our customers. It is important to understand that reimbursement is a complex process and requirements are subject to change without notice. It is the responsibility of the healthcare provider to determine and submit appropriate codes, charges, and modifiers for services that are rendered. Prior to filing any claims, customers are advised to contact their third-party payers for specific coverage, coding and payment information. Wallach Surgical Devices makes no promise or guarantee of reimbursement by Medicare or any other third-party payer.

Package Contents
The Vista AVS unit includes the following:
- User Manual
- Quick Guides
- ABI Chart
- Hand Controller
- Digit Cuff
- 10 cm Cuffs (2)
- 12 cm Cuffs (4)
- 17 cm Cuffs (2)
- Printer Paper
- Tape Measure
- Hose Set
- PPG Probe
- Training Video
- Ultrasound Gel
- Stand with Basket & Knobs (L500VA only)
- 8 MHz Bi-Directional Probe
- AVS Report Software
- USB Cable
- Power Supply (+7 VDC) & cable

The following are optional and sold separately from the Vista AVS.
- 5 MHz Bi-Directional Probe
- Carrying case
SECTION 2: Safety Information

**Intended Use:**
This device is intended for detection of blood flow in veins and arteries and as an aid for the diagnosis of peripheral vascular disease.

**Contraindications:**

**WARNINGS**
- The ultrasound probes are not to be used on or near the eyes.
- This device is for use only on intact skin.
- This device is not intended for use with HF surgical equipment.
- This device is not intended for fetal use.

**General Warnings:**

**WARNINGS**
- The Vista AVS is for use by qualified personnel only. Read the User Manual before use.
- Carefully route all cables and tubing to reduce the possibility of patient entanglement or strangulation.
- Do not allow the patient to operate any portion of the equipment, including the hand-held controller.
- Do not place the equipment in any position that would allow it to roll, fall, or collide with the user or patient.
- Do not use equipment that is damaged or malfunctioning. Seek appropriate service when needed. Inspect equipment regularly for signs of damage. Use alternate equipment if needed.
- Do not connect Luer fittings from Summit Doppler equipment into any other equipment.
- Confirm the setting of the real-time clock prior to saving patient data.
- Any equipment connected to the USB data port must be configured to comply with IEC 60601-1. By connecting additional equipment to the USB data port, the user is configuring a medical system and the user is responsible for ensuring the overall system compliance. Connected equipment must be certified to the applicable IEC standard (i.e. IEC 950 for data processing equipment, IEC 60601-1 for medical equipment). Contact the technical service department for more information.
General Cautions:

CAUTIONS

- Do not plug the probe cables into a telephone system, computer, or any equipment other than the Vista AVS.
- Read the Maintenance and Cleaning Section (Section 13) before cleaning.
- U.S. Federal law restricts this device to sale by or on the order of a licensed practitioner.
- Do not drop or mishandle the Vista AVS main unit, probes, hand controller or other accessories. Damage may occur.

Limitations of Noninvasive Vascular Tests:

In Current Noninvasive Vascular Diagnosis (Chapter 13, Page 208), Ali F. AbuRahma and Edward B. Diethrich note the following limitations of arterial leg Doppler examinations.

- Falsely high segmental pressure readings in areas with calcified arteries
- Artificially elevated high-thigh pressure in very large or obese patients
- Difficult interpretation of segmental pressures in patients with multilevel occlusive disease
- Difficulty in interpretation of high-thigh readings
- False-negative results in patients with mild vascular occlusive disease who have normal resting ankle pressures

Safety of Ultrasound:

The Vista AVS Doppler probes were designed to be safe and effective. However, the risk from some hazards cannot be eliminated completely. Instead, they are reduced to a level that is As Low As Reasonably Achievable (ALARA). Prudent use of this device in accordance with the principle of ALARA includes minimizing the duration of the patient’s exposure to ultrasound to the extent practical.

Clinical Safety

Approved by the American Institute of Ultrasound - March 1997, October 1982

Diagnostic ultrasound has been in use since the late 1950s. Given its known benefits and recognized efficacy for medical diagnosis, including use during human pregnancy, the American Institute of Ultrasound in Medicine herein addresses the clinical safety of such use:

There are no confirmed biological effects on patients or instrument operators caused by exposures from present diagnostic ultrasound instruments. Although the possibility exists that such biological effects may be identified in the future, current data indicates that the benefits to patients of the prudent use of diagnostic ultrasound outweigh the risks, if any, that may be present.
Safety in Training and Research

Approved by the American Institute of Ultrasound - March 1997, March 1983

Diagnostic ultrasound has been in use since the late 1950s. There are no confirmed adverse biological effects on patients resulting from this usage. Although no hazard has been identified that would preclude the prudent and conservative use of diagnostic ultrasound in education and research, experience from normal diagnostic practice may or may not be relevant to extended exposure times and altered exposure conditions. It is therefore considered appropriate to make the following recommendation:

In those special situations in which examinations are to be carried out for purposes other than direct medical benefit to the individual being examined, the subject should be informed of the anticipated exposure conditions, and of how these compare with conditions for normal diagnostic practice.
SECTION 3: Description of Product and Controls

Description of Unit

The Vista AVS is a physiologic exam system designed to aid the clinician in the diagnosis of peripheral arterial disease (P.A.D.). The unit includes a sensitive, bi-directional Doppler system, arterial photoplethysmograph (PPG), and a pressure system that provides inflation, controlled deflation, and pulse volume recording (PVR) capabilities.

The Vista AVS is well suited for the ankle brachial index (ABI) examination, the gold standard for the diagnosis of P.A.D. The ABI compares the systolic blood pressure at the ankles with the systolic pressure at the brachial arteries. A significantly reduced ankle pressure results in a low (<0.9) ABI value, which indicates P.A.D. Systolic ankle pressures are obtained with a pressure cuff and audio Doppler probe. ABI measurements are discussed in detail in Section 6.

A single level, lower extremity arterial exam (CPT® 93922) includes the ABI pressures, calculated index, and arterial physiologic waveforms. Two types of waveform modalities are provided on the Vista AVS: continuous-wave (CW), bi-directional Doppler and PVR. Both of these waveform modalities meet the requirements of CPT 93922. Although both modalities have significant clinical utility, it is generally not necessary to include both PVR and Doppler waveforms in reimbursement documentation for CPT 93922 - either one is sufficient. Waveform analysis is discussed in Section 12.

The Vista AVS is designed to perform segmental studies to compare three or more lower limb pressures to the brachial pressures. This procedure is reimbursable under CPT code 93923 as a non-invasive, physiologic study of upper or lower extremity arteries, multiple levels or with provocative functional maneuvers, complete bilateral study.

The Doppler waveform is a graph with a vertical axis (Doppler frequency shift, or pitch) proportional to the velocity of arterial blood flow. Flow toward the probe is indicated above the baseline. Flow away from the probe is indicated below the baseline. The 8 MHz, bi-directional probe is best for superficial vessels and all-around use. The optional 5 MHz, bi-directional probe is used for deeper vessels and with some obese patients.

Pulse Volume Recording (PVR) is a form of plethysmography, which is an indirect method of limb volume measurement. A pressure cuff is applied to the limb and inflated to 65 mmHg to detect the minute fluctuations in limb volume that occur with each heart beat. The PVR waveform’s contour is a qualitative indicator of presence or absence of peripheral arterial disease. This type of PVR does not permit calibration by injection of a known air volume and is used for arterial waveform analysis.

Photoplethysmography (PPG) is an optoelectronic technique for detecting the small changes of blood volume that occur in the capillary bed. Infrared (IR) light is emitted by the PPG probe into the skin. Light reflected from the underlying tissue is received by a detector and converted to an electrical signal. Since blood attenuates IR light at a higher level than the surrounding tissue, the signal’s pulse contours are determined by the arterial blood supply.
This type of PPG system is primarily for arterial pulse detection. In conjunction with a digit cuff, arterial PPG is quite useful for toe pressure measurement, which is an additional diagnostic tool when the arteries at the ankle are noncompressible (ABI > 1.30) due to calcification. Calcified arteries are prevalent among patients with diabetes or kidney disease, but the small arteries of the toes seldom become calcified. When the Vista AVS is configured for PPG, the system automatically calculates the toe brachial index (TBI). A low TBI (<0.65) indicates an arterial obstruction. The TBI is discussed in Section 7.

The Vista AVS includes everything that is needed to conduct fast, efficient peripheral arterial examinations including display, printer, Doppler probes, PPG probe, pressure hose, limb pressure cuffs, and a digit cuff. The instrument operates from either its internal battery pack or from an external medical grade power supply at 100-240 VAC.

**Controls and Indicators:**

**Main Unit**

- **ON/OFF** - *Turns the unit on or off with momentary actuation*
- **FEED** - *Printer feeds paper while pressed*
- **CHARGING**
  - On: *Charging*
  - Flashing: *Charging complete (trickle charging while flashing)*
- **LOW BATT** - Flashing: *Low battery*
- **PAPER RELEASE**
  - Blue button under paper dispenser: *Opens printer cover*
Display Screen - Protocol Location and Markers

- Pressure Gauge (mmHg)
- Right & Left ABI
- Right & Left Brachial Sites
- Waveform Sites (Right & Left PT or DP)
- Right & Left Ankle Sites (PT or DP)
- Right & Left Ankle Sites (PT or DP) (Toe Sites when in PPG Mode)
- Date and Time
- Step-By-Step ABI Instructions

Hand-Held Controller

PUMP - Pump runs while pressed

SAVE - During deflation- Stores current cuff pressure and deflates
- Active waveform- Stores waveform data
- Frozen waveform- Stores the frozen waveform
- 2nd button press- Confirms the stored pressure or waveform

FREEZE - Active waveform- Freezes the displayed waveform
- Frozen waveform- Starts new waveform acquisition

SITE - Moves the protocol marker to the next location
- Saves data before leaving old protocol location

SCALE - Changes the waveform scaling
**MUTE** - Enables/disables Doppler speaker output

**PRINT** - Prints out the waveform on adhesive-backed label paper

**ENTER/MENU** - Enters the displayed value or makes a menu selection
- Also used to open the OPTIONS screen

**UP** - Navigates through the menu in the up direction & moves the caliper tool to adjust the pressure values

**DOWN** - Navigates through the menu in the down direction & moves the caliper tool to adjust the pressure values

<table>
<thead>
<tr>
<th>Numeral Mode</th>
<th>Character Mode</th>
</tr>
</thead>
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<tr>
<td></td>
<td>Space</td>
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<tr>
<td></td>
<td>Hyphen</td>
</tr>
</tbody>
</table>

See Section 5 for information on how to change between Character and Numeral Modes.
SECTION 4: Preparation for Use

Tools required: Phillips head screwdriver

1. Unpack the Vista AVS and inspect the unit for external damage.
2. Verify that you have received each of the contents listed on the packing list.
3. Assemble the rolling stand (if applicable) using the instructions provided.
4. Use a Phillips head screwdriver to attach the Vista AVS mounting bracket to the end of the rolling stand’s pole. Be sure to fit the pole’s alignment pin through the alignment hole on the bracket.

5. Use the two stand mounting knobs to attach the mounting bracket to the Vista AVS.

6. Review the unit and locate each of the controls and connectors (see Section 3 and information on the next page).
7. Plug the external power supply into the +7 VDC (Power) connector.
**VISTA TIP™:**

- For first time use, allow the unit to charge for at least one hour before operating the Vista AVS from its battery.

8. Plug the power cord into the external power supply and then into a properly grounded wall outlet.


10. Plug the Doppler cable into the Doppler connector. (Blue port)

11. Plug the PPG probe into the PPG connector. (Black port)

**VISTA TIP:**

- The Doppler and PPG connectors are color coded and physically interchangeable. If you fail to connect the proper probe, no damage will occur.

12. Attach a cuff to the Luer style hose fitting.

13. Plug the hose quick connect style fitting into the CUFF connector on the connector panel.

14. Plug the hand-held controller into the HAND CONTROLLER connector.

**VISTA TIP:**

- To maintain patient safety, it is not possible to conduct an examination while the Vista AVS is connected to a PC via the USB port.

**Connectors**

![Connectors Diagram]

- Doppler
- PPG
- Hand Controller
- USB
- +7 VDC (Power)
- Cuff
SECTION 5: Menu Configuration and Set Up

Loading Paper:

The Vista AVS is shipped with paper pre-loaded. To load a new roll of paper, press the printer release button. Remove any remaining paper from the old roll and drop in the new roll as shown below. Close the paper door and push firmly enough to latch the door into position. Press the FEED button to ensure proper paper alignment.

WARNING

- Printer components become hot during printing. Do not touch the metal pieces inside the paper holder immediately after printing. After loading paper, press the ON/OFF switch to begin using the instrument.

Configuring the Examination

The Vista AVS can be configured to perform P.A.D. testing in four different automated modes. Pressure can be obtained using either the Doppler at the ankle for typical ABI testing described in Section 6 or using a PPG at the toes for TBI testing described in Section 7. Additionally, waveforms used to complete the physiologic study can be performed with either a Doppler or the Pulse Volume Recording (PVR) mode.

1. Press the ENTER/MENU key to open the OPTIONS menu.
2. Press the number 1 key to select the CONFIGURE EXAMINATION.
3. Select the desired configuration for the examination by pressing the appropriate number:

   1 – ABI with PVR Waveform
   2 – ABI with Doppler Waveform
   3 – TBI with PVR Waveform
   4 – TBI with Doppler Waveform
   5 – Segmental with PVR Waveform
   6 – Segmental with Doppler Waveform
**VISTA TIP:**
- The current configuration mode will be displayed at the bottom of the waveform upon returning to the main screen.

**Setting the Date and Time**

1. Press **ENTER/MENU** to open the OPTIONS menu.
2. Press the number 2 key to select SYSTEM SETTINGS.
3. Press the number 1 key to set the clock.
4. The clock setting format is: MM DD YY HH MM SS (Month-Day-Year-Hours-Minutes-Seconds)
5. Use the **UP** key to move the * cursor over the field that needs to be changed.
6. Use the numerals on the hand-held controller to set the date and time.
7. Press **ENTER/MENU** or move the * cursor to the end to exit the CLOCK menu.
8. Press the **ENTER/MENU** key to exit the OPTIONS menu.

**Contrast Adjust**

To modify the contrast level on the Vista AVS screen:

1. Press the **ENTER/MENU** key to open the menu.
2. Press the number 2 key to select the SYSTEM SETTINGS.
3. Press the number 2 key to increase CONTRAST ADJUST LEVEL one level.
4. Continue to press the number 2 key until desired level is reached.

**VISTA TIP:**
- If there is flickering or shadowing on the text, the contrast is adjusted too high. Readjust contrast level by pressing the number 2 key until contrast level starts over at 1 and slowly increase the contrast level.
Setting the Power Down Status

The Vista AVS can be set to turn off automatically to save battery power. If the automatic power down is set to ON, the unit will turn off after 20 minutes automatically after the last key press. If the automatic power down is set to OFF, the unit will not turn off until either the user presses the ON/OFF button or the battery becomes low.

1. Press **ENTER/MENU** to open the OPTIONS menu.
2. Press the number 2 key to select SYSTEM SETTINGS.
3. Press the number 3 key to toggle the power down value to ON and OFF.
4. After setting the desired value, press **ENTER/MENU** to exit the OPTIONS menu.

Setting the Ankle Protocol

The Vista AVS can be set to accommodate protocols that use either one pressure measurement or two pressure measurements from each ankle. The DUAL ANKLE PRESSURE mode will be set to OFF for a single ankle pressure. For protocols that use measurements from both the dorsalis pedis (DP) and the posterior tibia (PT) arteries, select HIGH or LOW to select the desired protocol with second ankle pressure.

1. Press **ENTER/MENU** to open the OPTIONS menu.
2. Press the number 2 key to select SYSTEM SETTINGS.
3. Press the number 5 key to toggle the ankle value between OFF, HIGH or LOW.
4. After setting the desired value, press **ENTER/MENU** to exit the OPTIONS menu.

File Annotation

The File Management SAVE function can use either alpha or numeric characters for saving a patient filename. To configure the system to the desired function:

1. Press the **ENTER/MENU** key to open the OPTIONS menu.
2. Press the number 2 key to select the SYSTEM SETTINGS.
3. Press the number 4 key to toggle between NAME or NUMBER modes.
4. To enter a filename, use the alphanumeric keys on the hand-held controller. Use the **DOWN** key to backspace and the **UP** key to move between different letters on the same button.
5. Press the **SAVE** key to save the filename and begin the exam.
Selecting a Modality for Obtaining Pressures

Choose between the Doppler probe and the photoplethysmography (PPG) probe to obtain pressure values for ankle brachial index (ABI) exams, toe brachial index (TBI) exams or segmental studies:

1. Press the ENTER/MENU key to open the OPTIONS menu.
2. Press the number 2 key to select the SYSTEM SETTINGS.
3. Press the number 6 key to toggle between DOPPLER and PPG.

The selected pressure modality is indicated in the lower right corner of the display (DOP or PPG will appear depending on which is selected) as shown below.

VISTA TIP:
- If you’re trying to obtain pressures or waveforms using the Doppler probe but it is not audible when rubbing the probe tip, the system is probably set in PPG mode. To change the setting, go to the System Settings menu and select Doppler for obtaining pressures and waveforms.
Entering a Filename Prior to Beginning an Exam

If beginning a new exam by pressing the number 6 key for NEW EXAM – CLEAR ALL under the OPTIONS screen, the system will display the ENTER FILENAME message. This message ensures that the user is aware that the patient data will not be automatically saved. After completing the exam, the user can save the data by following the instructions provided in the “Saving a File” paragraph in Section 11.

If a current exam is not saved and the user selects either NEW EXAM - CLEAR ALL or NEW EXAM – KEEP BRACHIALS under the OPTIONS screen, the system will display the UNSAVED DATA message below to let the user know that the previous data was not saved.

Press the ENTER/MENU key to ignore this message and start a new exam (the ENTER FILENAME will appear), or press the SAVE key to save the data and the SAVE FILE menu will appear.
SECTION 6: The Ankle Brachial Index (ABI) Examination

Preparing the Patient

In a warm room, have the patient take off his/her shoes and socks and rest in a supine position for approximately 5 minutes prior to taking pressures. The patient should wear thin, loose fitting clothing. Avoid rolling up sleeves or pant cuffs in such a manner that it obstructs blood flow. Bulky items such as sweaters should be removed.

Wrap the cuffs snugly around the arms and ankles as shown below. The edge of the cuff should be placed approximately 1 to 2 inches above the site of examination. Select the proper cuff width, equivalent to 40% of the patient’s limb circumference. In general, average sized adults use 10 cm cuffs, while larger adults use 12 cm cuffs. The most efficient way to conduct the exam is to wrap all four limbs prior to taking any pressures.

While applying the cuffs, it may be a good time to explain the examination to the patient and answer any questions.

NOTE: It is very important that the patient remain still and in a supine position for the duration of the exam.

VISTA TIP:
- The Vista AVS performs the ABI exam in the following order: Right Brachial Pressure (R-BRA), Right Ankle Waveform (R-WAV), Right Ankle Pressure (R-ANK), Left Ankle Waveform (L-WAV), Left Ankle Pressure (L-ANK), and Left Brachial Pressure (L-BRA). You may override this order by using the SITE key to move between protocol locations.

- Notice that the patient’s right hand side is shown on the left side of the Vista AVS’s screen in order to match what you see when you face the patient.
Getting Started

Press the ENTER/MENU key to open the OPTIONS menu. Press the number 6 key to start a new exam. You may enter a filename at this time by using the keypad. Refer to Section 5 for File Annotation instructions.

**VISTA TIP:**
- To start a new study on the current patient, press the number 5 key to clear the data except for brachial pressures. This is useful when it is determined a TBI will be required following an ABI with noncompressible arteries.

Obtaining the Right Brachial Pressure

Connect the pressure hose to the fitting on the right brachial cuff. Apply a small amount of gel over the right brachial artery and place the Doppler probe at approximately 45 degrees, pointing in the direction toward the shoulder as shown below. Slide the probe laterally across the arm to find the brachial artery and obtain the best signal possible. Adjust the volume knob to acquire a proper audio level. Once the best signal is obtained, brace the hand holding the probe against the patient’s arm to ensure a stable position. Be careful not to apply too much pressure against the skin. Excessive pressure could occlude the artery.

Next, press and hold the PUMP key to inflate the cuff. Release the key once the pressure reaches about 20 mmHg above the occlusion pressure. After the pump stops the Vista AVS automatically bleeds the cuff at the correct rate.

Press the SAVE key at the moment flow returns. This will store the brachial pressure value in the R-BRA protocol location. This is the systolic pressure. In general, the audible Doppler signal is a slightly more sensitive indicator of the systolic pressure than the Doppler waveform display.
The cuff deflates automatically when SAVE is pressed. If needed, press PUMP again to repeat the measurement, or use the arrow keys or numeric keypad to adjust the pressure value before confirming the value by pressing the SAVE or SITE key.

Press SITE or SAVE again to confirm the right brachial pressure and move to the next protocol location.

**VISTA TIP:**
- Once a satisfactory pressure has been obtained, pressing SAVE or SITE will confirm the stored value and move to the next protocol location.
- During deflation, pressing PUMP again will re-inflate the cuff and allow a new pressure to be taken.

**OBTAINING PRESSURES USING THE PPG PROBE:**

In the supine position, ABI pressures can also be obtained using the PPG probe. For obtaining pressures, apply the PPG probe to an index finger for brachial pressures or the great toe for ankle pressures. It is important the patient is still and the fingers and toes are reasonably warm. To obtain brachial pressures, wrap the cuffs around the arms at the brachial artery site. Place the PPG on the index finger with the blue side of the probe against the skin. It should be snug to ensure contact, but not too tight to occlude blood flow as shown below.

It will take a few seconds for the waveform to stabilize on the display after applying the probe. You may want to consider changing the scale of the display to get a better view of the waveform by pressing the SCALE key. Once the waveform is stable, press the PUMP key to inflate the cuff to a pressure approximately 20 mmHg above the pressure until the PPG pulsations disappear. The arm cuff will begin to deflate automatically once the pump stops.
After the first consistent pulsation returns press the **FREEZE** key to freeze the waveform on the display. Use the caliper tool by pressing the **UP** or **DOWN** arrow keys to adjust the pressure reading. Move the caliper to the beginning of the first upward slope at the start of the pulsation as shown below. This is the systolic pressure. Press the **SAVE** key to save the pressure and move to the next site to be measured.

Refer to Section 7 for instructions for using the PPG probe to obtain toe pressures.

**Right Ankle Waveform**

Verify that the protocol marker is on R-WAV.

**VISTA TIP:**

- A single level lower extremity arterial examination normally consists of the ABI value and either bi-directional Doppler waveforms or PVR waveforms. The following shows how to set the waveform mode. See Section 12 for information on waveform interpretation. Note that arterial tests documented with PPG waveforms are not reimbursable as opposed to Doppler or PVR waveforms.

The Vista AVS can acquire bi-directional Doppler waveforms, pulse volume recording (PVR) waveforms and photoplethysmography (PPG) waveforms. See Section 5 for configuring the desired waveform mode.

**OBTAINING DOPPLER WAVEFORMS:**

Apply a small amount of coupling gel over the site of the artery, either the PT or DP artery. Place the probe over the artery at an angle of approximately 45 degrees, pointing the probe tip in the direction toward the calf and knee. Slide the probe slowly across the artery until the best signal is obtained.
Once the waveform is optimized, press the SCALE button one or more times as needed to adjust the vertical scaling to obtain a waveform as large as possible without clipping at the top. If the probe is pointed in the correct direction (approximately 45 degrees “upstream”) the majority of the waveform will be above the baseline. Press PRINT to print the waveform or FREEZE to freeze or pause the waveform. When frozen, the UP or DOWN keys will scroll the waveform. Press SAVE to store the waveform, and then use the UP key to adjust the site label (PT, DP, TOE, BRA, ***). Press SITE or SAVE again to store these values and move to the R-ANK (right ankle) location.

**OBTAINING PVR WAVEFORMS:**

**VISTA TIP:**

- If you pump over 75 mmHg pressure, the unit will automatically deflate to 65 mmHg. Note that the audio will be off or muted while in the PVR mode.
- You may want to consider setting the scale to PVR x 2 to view the waveform. This is done by pressing SCALE on the hand-held controller to change the view of the display.

To obtain the PVR waveform from the ankle, wrap the proper size cuff [see the beginning of this section (Preparing the Patient) for proper cuff selection] around the ankle. Move the cursor to the Right WAV site and use the PUMP control to inflate the ankle cuff to 65 mmHg or slightly higher. It will take several seconds for the position of the waveform to “settle out” and return to the center of the display. After the waveform stabilizes, press the FREEZE key. Here, you have two options: if the waveform is acceptable, press SAVE to record it. Or, you can use the UP or DOWN key to scroll to the optimum waveform. Press SITE or SAVE again to move to the R ANK (right ankle) location.

![PVR Waveform](image)

**Right Ankle Pressure**

Verify that the protocol marker is on the R-ANK location.

**VISTA TIP:**

- The system can be configured for exam protocols that use either one or two (DP and PT) ankle pressure measurements. See Section 5 for information on how to set the Ankle Protocol.
Ensure that the pressure hose is connected to the fitting on the right ankle cuff. Position the Doppler probe over the PT or DP artery. Be sure to use ample gel. Obtain an adequate signal, then press and hold the PUMP key to inflate the cuff. Release the key once the pressure reaches about 20 mmHg above the occlusion pressure.

After releasing the PUMP key, the Vista AVS automatically deflates or bleeds the cuff at the correct rate. Press the SAVE key at the moment flow returns to store ankle pressure value in the R-ANK location.

The cuff fully deflates automatically when the pressure is saved. If needed, press PUMP again to repeat the measurement, or use the UP or DOWN keys or numeric keypad to adjust the pressure value before confirming the value by pressing the SAVE or SITE key.

**Protocol for Two Ankle Pressures**

If the protocol configuration is set to HIGH or LOW, press SAVE while deflating to the PT pressure. Press SAVE again to confirm this pressure.

The site will change to DP. Press PUMP to repeat the pressure cycle. Both pressures will be stored and displayed. The displayed ABI will be either the higher or the lower value, depending on the user setting.

**VISTA TIP:**
- When conducting the ABI exam on an amputee, use the SITE key to select the appropriate pressure/waveform locations.
**Left Ankle Waveform**

Repeat the procedure described previously for the right ankle waveform. When the left ankle waveform is stored in the L-WAV site, the site cursor moves to L-ANK (left ankle pressure) site.

**Left Ankle Pressure**

Move the cuff hose to the left ankle cuff and repeat the procedure described for the right ankle on the left ankle. When the left ankle pressure is stored in the L-ANK site, the site cursor moves to L-BRA (left brachial pressure) site.

**Left Brachial Pressure**

Move the cuff hose to the left arm cuff and repeat the procedure described for the right brachial on the left brachial. The greater of the two arm pressures will be used to calculate the ABI. A difference of 20 mmHg or more between the two brachial pressures is a possible indication of upper extremity stenosis. In this case, the arm pressures should be repeated and if the result still indicates a difference of 20 mmHg or more, upper extremity testing should be considered.

After the left brachial pressure is stored in the L-BRA site, the site cursor moves to R-BRA (right brachial pressure) site. At this point, the exam data is ready for saving, printing and/or downloading.

**Assessment of the Ankle Brachial Index (ABI)**

The Vista AVS automatically calculates the ankle brachial index (ABI) for both sides. The right and left ABI are both calculated using the higher of the two brachial pressures.

<table>
<thead>
<tr>
<th>ABI Assessment</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Above 1.30</td>
<td>Noncompressible Artery</td>
</tr>
<tr>
<td>1.00 to 1.29</td>
<td>Normal</td>
</tr>
<tr>
<td>0.91 to 0.99</td>
<td>Borderline (equivocal)</td>
</tr>
<tr>
<td>0.41 to 0.90</td>
<td>Moderate Disease</td>
</tr>
<tr>
<td>0.00 to 0.40</td>
<td>Severe Disease</td>
</tr>
</tbody>
</table>

See Section 11 for File Management information including saving, printing, recalling patient data, and for information on how to reset for new patient data.
SECTION 7: The Toe Brachial Index (TBI) Examination

Toe pressures are useful in the assessment of diabetic patients with noncompressible arteries (ABI > 1.30), and patients with Raynaud’s syndrome, or with advanced peripheral arterial occlusive disease including the smaller distal vessels. Most clinicians do not perform toe pressures on patients with normal ABI results.

Preparing the Patient

Patient preparation for the TBI exam is similar to preparation for the ABI exam; however the need to obtain PPG signals from the toes requires consideration of patient temperature. It may be more difficult to obtain PPG waveforms if the patient is cold. If needed, consider laying a blanket over the patient or wrapping a warm towel around the patient’s foot. Some clinicians prefer to leave the patient’s socks on until they are ready to take the toe pressures.

Brachial Pressures

Brachial pressures are conducted in the manner described in Section 6 for the ABI examination.

Ankle Waveforms

Either Doppler or PVR waveforms may be used accurately on patients with noncompressible arteries. In some cases, calcification may result in reduced ultrasound penetration and weak Doppler signals. Refer to Section 5 for details on switching to Doppler or PVR waveforms. Obtaining ankle waveforms is described in Section 6.

Toe Pressures

Refer to Section 5 for details on configuring the system for TBI using PPG.

Wrap the great toe with a 2.5 cm or 1.9 cm digit cuff and attach the pressure hose. The cuff should be 40% of the toe circumference. Place the cuff at the base of the toe and attach the PPG probe to the distal phalanx with the blue side of the probe against the skin as shown in the photos below. NOTE: The ankle cuff is not used while taking toe pressures.
Wait a few seconds for the PPG waveform to stabilize after applying the probe. Once a stable PPG waveform is obtained, press the PUMP key to inflate the digit cuff. Continue to inflate to a pressure approximately 20 mmHg above the pressure where the PPG pulsations disappear. The digit cuff will begin to deflate automatically once the pump stops. To get a better view of the waveform on the display, press the SCALE key.

After the first pulsation returns press the FREEZE key to freeze the waveform on the display. Use the caliper tool by pressing the UP or DOWN arrow keys to adjust the pressure reading. Move the caliper to the beginning of the first upward slope at the start of the pulsation as shown below. This is the toe systolic pressure. Press the SAVE key to save the pressure and move to the next site to be measured.

NOTE: A substantial shift in the baseline may occur when blood flow returns. This is a useful indicator of pulse reappearance.

Press SAVE or SITE to confirm the value and move to the next exam location.

**Assessment of the Toe Brachial Index**

The Vista AVS automatically calculates the toe brachial index (TBI) for both sides. The right and left TBI are both calculated using the higher of the two brachial pressures.

**TBI Assessment**

| Above 0.70 | Normal |
| 0.64-0.70 | Borderline |
| Below 0.64 | Abnormal |
SECTION 8: The Seated ABI Examination

Performing the ABI in the seated position makes it possible to assess patients who are not able to lie in the supine position for ABI determination. These patients may be physically disabled or wheel chair bound, patients with degenerative disease of the spine or severe back pain, patients with advanced cardiopulmonary disease and orthopnea.

The Vista AVS is able to calculate ABI values for seated patients by compensating for the effects of gravity on the lower extremity pressures based on the vertical distance between the arm and ankle cuffs.

To begin the exam, have the patient sit erect in a chair with knees at a 90-degree angle. The feet should rest flat on the floor or on an elevated platform - feet should not be dangling. Wrap the cuffs around the arms and ankles.

To determine the difference in height between the two cuffs, using a tape measure or yardstick, measure the distance in centimeters between the middle of the arm cuff to the floor and the middle of the ankle cuff to the floor as shown below. Next, subtract the ankle distance from the arm distance to get the vertical distance to be entered for the hydrostatic correction.

Press the ENTER/MENU key on the hand-held controller to open the OPTIONS menu. Press the number 1 key to CONFIGURE EXAMINATION. You will press the number 1 key to select “ABI with PVR Waveform”. Press the ENTER/MENU key to open the OPTIONS menu again and press number 2 key for SYSTEM SETTINGS.

Press the number 6 key to use the Doppler probe for obtaining pressures. Press the ENTER/MENU key to exit. Press the ENTER/MENU key again to open the OPTIONS menu. You will then press the number 7 key for HYDROSTATIC CORRECTION and using the keypad, enter the vertical height between the cuffs in centimeters. Press the SAVE key when finished.

Start at the right brachial by applying the gel and placing the Doppler probe over the artery site to listen to the blood flow. While holding the probe steady on the artery, press and hold the PUMP key to about 20 mmHg above the occlusion pressure. Release the PUMP key and the cuff will automatically start to deflate.
Press the **SAVE** key when you hear the first pulsation return. If you were late pressing the **SAVE** key, you can adjust the pressure reading by using the **UP** or **DOWN** key. You will then press the **SAVE** key again to record the systolic pressure and the marker will move to the next protocol location.

Move the hose from the arm cuff to the ankle cuff to obtain the ankle waveform using PVR. Press and hold the **PUMP** key to inflate the cuff to 75 mmHg and release the key when the inflation automatically stops. Pressure will deflate to 65 mmHg for optimum PVR. Wait for the “cuff deflating” message on the display to disappear. If needed, use the **SCALE** key to optimize the waveform.

After the waveform stabilizes, press the **FREEZE** key and if the waveform is acceptable, press the **SAVE** key to record. Or, use the **UP** or **DOWN** key to scroll to the optimum waveform and press the **SAVE** key to record it. Press the **SAVE** key again to move to the next protocol location.

Now, obtain the ankle pressure. The hose should be connected to the ankle cuff. Apply gel to either the dorsalis pedis artery or the posterior tibia artery and use the Doppler probe to listen to the blood flow. While holding the probe steady on the artery, press and hold the **PUMP** key to about 20 mmHg above the occlusion pressure. Release the **PUMP** key and the cuff will automatically start to deflate.

Press the **SAVE** key when you hear the first pulsation return. If you were late pressing the **SAVE** key, adjust the pressure reading by using the **UP** or **DOWN** key. Press the **SAVE** key again to record the systolic pressure and the marker will move to the next protocol location. Continue these steps for the remaining sites to complete the exam. The Vista AVS makes the hydrostatic correction based on the vertical distance entered prior to obtaining pressures.

**VISTA TIP:**

- If the calculation for the hydrostatic correction factor results in an ankle pressure that is equal to or less than zero, the result is displayed as "---" and the associated Seated Index (SI) is "0.0".
- The PPG probe is not suitable for obtaining pressures in the seated position. The Doppler probe is available and recommended.
- The two pressure modes, high or low, are not available while performing the Seated ABI.
SECTION 9: Segmental Studies

Segmental studies are usually performed on patients diagnosed with P.A.D. to localize the occlusion in the lower limbs. A segmental study is an extension of the ABI exam, comparing three or more lower limb pressures to the brachial pressures. It is possible to obtain pressures and waveforms from five lower-limb, arterial sites using the Vista AVS.

For a complete segmental study, you must obtain at least three pressures and waveforms from the lower extremity. A difference of 20 mmHg or greater from one cuff site to the next distal cuff site suggests P.A.D. at the area just proximal to the distal cuff.

To begin, press the ENTER/MENU key to open the OPTIONS menu and press the number 1 key to CONFIGURE EXAMINATION. Here, you have two options: press the number 5 key to select “Segmental with PVR Waveform” or press the number 6 key to select “Segmental with Doppler Waveform”.

Once you have made your selection, the SEGMENTAL EXAMINATION CONFIGURATION screen will be displayed. Press number 1 through 5 keys toggle through the various arterial sites to assess. Press the number 6 key to choose between performing the exam “In-Line” down the leg or “Laterally” moving from one side of the patient to the other. When finished, press the ENTER/MENU key.

Now that you’ve configured the segmental exam, press the ENTER/MENU key to open the OPTIONS menu again and then press the number 2 key for SYSTEM SETTING. Press the number 6 key to choose a modality for obtaining pressures (either PPG or DOPPLER).
After making your selection, press the **ENTER/MENU** key to exit the SYSTEM SETTING menu and the screen should look like the example below.

[Image of a screen displaying segmental interpretation results]

Wrap a cuff at each site as the patient lies supine to prepare for the exam. Usually a 10 cm cuff is used at the ankle, a 12 cm cuff is used at the calf and a special, 17 cm-contoured, thigh cuff is used at the thigh. Be sure to connect the hose to the cuff of the site to be measured. Proceed with obtaining pressures and waveforms from the desired sites. Refer to Sections 6 and 7 for more details for using a Doppler probe, PVR and PPG.

<table>
<thead>
<tr>
<th>Segmental Interpretation:</th>
</tr>
</thead>
</table>

**Normal**
- All segmental pressures, including the thigh pressure, should be equal to or slightly greater than the brachial pressure.

**Abnormal**
- A 20 mmHg or greater drop in pressure generally is considered positive for hemodynamically significant stenosis (> 60% diameter) in the segment(s) leading into the cuff or under the cuff. It may be helpful to compare pressures in one limb to the other, and to consider the PVR (or Doppler) waveforms in addition to the pressures.

**Caution:** It is necessary that an appropriately sized cuff be used for thigh pressures – typically 12 cm, 17 cm or 22 cm in diameter. If the patient is too large for the available cuff, the girth of the limb may cause an artificially high pressure, known as the “high thigh cuff artifact”. In this situation, the following exceptions apply:

**Normal**
- High Thigh Exception: The pressure should be at least 20 mmHg above the brachial pressure.

**Abnormal**
- High Thigh Exception: A pressure drop of 30 mmHg from the high thigh to above knee is suggestive of disease.
SECTION 10: Individual Site Mode

If you would like to quickly obtain a pressure or waveform from an arterial site (NOT following the standard protocol sequence of the exam calculation), use the INDIVIDUAL SITE MODE by pressing the ENTER/MENU key to open the OPTIONS screen and pressing the number 4 key. You can use this mode while in the middle of performing the ABI, TBI and segmental study and it will not interrupt the order of the exam. When you exit the INDIVIDUAL SITE MODE, you will return to the exact site where you left off to complete the exam.

There are three modality options in this mode: Doppler, PVR or PPG. Use the UP or DOWN keys to move the menu cursor to select a modality option.

NOTE: Pressures and waveforms obtained using the INDIVIDUAL SITE MODE can be printed but the data is only temporarily stored on the display screen and cannot be saved or downloaded. Temporarily storing or freezing a waveform enables you to change to another modality to obtain the next pressure.

Use the SITE key to select the desired arterial site. There are many site options to choose from:

<table>
<thead>
<tr>
<th>RIGHT BRACHIA (brachial)</th>
<th>LEFT BRACHIA (brachial)</th>
</tr>
</thead>
<tbody>
<tr>
<td>RIGHT FINGER</td>
<td>LEFT FINGER</td>
</tr>
<tr>
<td>RIGHT THIGH</td>
<td>LEFT THIGH</td>
</tr>
<tr>
<td>RIGHT HI KNEE (above the knee)</td>
<td>LEFT HI KNEE (above the knee)</td>
</tr>
<tr>
<td>RIGHT LO KNEE (below the knee)</td>
<td>LEFT LO KNEE (below the knee)</td>
</tr>
<tr>
<td>RIGHT CALF</td>
<td>LEFT CALF</td>
</tr>
<tr>
<td>RIGHT ANKLE</td>
<td>LEFT ANKLE</td>
</tr>
<tr>
<td>RIGHT PT (posterior tibia)</td>
<td>LEFT PT (posterior tibia)</td>
</tr>
<tr>
<td>RIGHT DP (dorsalis pedis)</td>
<td>LEFT DP (dorsalis pedis)</td>
</tr>
<tr>
<td>RIGHT TOE</td>
<td>LEFT TOE</td>
</tr>
<tr>
<td>PENIS</td>
<td></td>
</tr>
</tbody>
</table>

To obtain a pressure, use the PUMP key to inflate the cuff then use the SAVE key to record the pressure and deflate remaining pressure in the cuff. If system is set to digit (i.e. finger, toe or penis), the cuff will pump slowly. Otherwise, pump is set at normal speed. To obtain a waveform, press the SAVE or FREEZE keys to freeze the waveform on the display screen. Use the PRINT key to print out the waveform, including site and pressure information. To restore waveform activity, press the SAVE or FREEZE key again. Please refer to Sections 6 and 7 for a more detailed explanation of obtaining pressures and waveforms using a Doppler, PVR or PPG.

To exit the INDIVIDUAL SITE MODE, use the UP or DOWN keys to select EXIT and press the ENTER/MENU key.
SECTION 11: File Management

VISTA TIP:
- Exam reports can be configured for either numeric or alpha character filenames. To change the filename configuration, use the SYSTEM SETTINGS option (see Section 5 for details).
- It is not possible to alter exam data after a file is saved.

Saving a File

To save an exam, press the ENTER/MENU key to open the OPTIONS screen. Press the number 3 key to select FILE MANAGEMENT. Press the number 1 key to save the current exam. There are 10 storage locations that retain data if all power is lost. Select the number of an unused file storage location (this location will be highlighted). Use the alphanumeric keys on the hand-held controller to enter a patient filename. Use the DOWN key to backspace. Use the UP key to move between different letters on the same button. Press SAVE to save the file. After saving, the system returns to the main screen.

If the system was powered down prior to saving data, the FILE RESTORE message will appear on the display upon power up (see image below). Pressing the ENTER/MENU key will remove the message and allow the user to start a new exam or recall the previous unsaved patient data by accessing “Restore Data”. Powering down a second time will not repeat the message even if the restored file is still stored.

![FILE RESTORE Message](image)

NOTE: When saving an exam, the SAVE and ENTER/MENU keys perform the same function. Either key can be pressed to save an exam file.

If the user saves an exam under FILE MANAGEMENT by entering a patient name or ID and turns off the power before pressing the SAVE or ENTER/MENU key, the system will automatically complete the save function before the power is turned off.

NOTE: Powering off by unplugging the system will not activate this feature.
Recalling a File

To recall an exam, press the ENTER/MENU key to open the OPTIONS screen. Press the number 3 key to select FILE MANAGEMENT. Press the number 3 key to recall an exam from a file storage location. Select the number of the file to be recalled using the numeric keypad. To return to the existing file without recalling one of the saved files, press the ENTER/MENU key to exit FILE MANAGEMENT.

The Vista AVS automatically stores a backup copy of patient data in the “Restore Data” each time a new name, pressure or waveform is saved. The exam stored in the “Restore Data” is purged upon entering patient data (name, pressure or waveform) for a new exam. In other words, once new patient data is entered for a new exam, the previous exam in the “Restore Data” is deleted and the new exam is stored in the “Restore Data”. It is important to note that the power down and clear exam functions do not erase the previously stored exam in the “Restore Data”.

To access the exam stored in the “Restore Data”, press the ENTER/MENU key to open the OPTIONS screen. Press the number 3 key to open FILE MANAGEMENT and press the number 3 key again to open RECALL EXAM FROM FILE. Next, press the UP key to highlight RESTORE DATA and press the ENTER/MENU key to open the exam.

Printing a File

The PRINT key prints the patient data in the exam screen. Use the RECALL function described above to recall a saved file if needed. When only one waveform has been saved, the saved pressures, ABIs, and one waveform will print. If no waveform has been saved, only the pressures and ABIs will print. The printout will show 4 seconds of recorded waveform data although the display only shows 3 seconds. NOTE: It is not possible to print directly from the FILE menu.
Deleting a File

To delete an exam, press the ENTER/MENU key to open the OPTIONS screen. Press the number 3 key to select FILE MANAGEMENT. Press the number 2 key to delete an exam from a file storage location. Select the file number on the numeric keypad to delete the file, which will be highlighted. Press that number again to confirm or press any other key to cancel. To return to the exam screen, press the ENTER/MENU key to exit. To delete all files, use the UP or DOWN keys to navigate to DELETE ALL and press ENTER/MENU. Press the ENTER/MENU key again to confirm deletion or press any other key to cancel.

Downloading a File to a PC (current exam has NOT been saved)

To download patient data to a PC, open the OPTIONS screen using the ENTER/MENU key. Press the number 3 key to select FILE MANAGEMENT. Press the number 5 key to download the current exam. Current patient data will be transferred automatically, or press the ENTER/MENU key to cancel.

See Section 19 for more information about the AVS Report software.

VISTA TIP:
- When downloading data, the current exam screen is transferred to the PC.
- Downloading is only possible if the USB cable is connected between the Vista AVS and the PC.
- AVS Report software must be installed on the PC in order to transfer data.

Downloading Exam from File (current exam has been saved)

To download patient data to a PC, open the OPTIONS screen using the ENTER/MENU key. Press the number 3 key to select FILE MANAGEMENT. Press the number 4 key to download an exam that has been previously saved. Using the numeric keypad, select the number of the file to recall and download. Saved patient data will be transferred automatically, or press the ENTER/MENU key to cancel. You may also download all of the saved exams at once by using the UP or DOWN arrow keys to select DOWNLOAD ALL and pressing the ENTER/MENU key.

VISTA TIP:
- This mode is useful to help expedite downloads of saved files by eliminating the need to recall the file first.
SECTION 12: Peripheral Arterial Waveform Interpretation

**Bi-Directional Waveform Interpretation:**

Continuous-wave, bi-directional Doppler waveforms are useful for assessment of lower extremity P.A.D. severity and progression and are a valuable addition to the ABI, particularly for patients with noncompressible arteries.

The normal Doppler signal is pulsatile and multi-phasic. Below is a normal waveform from the PT artery. The systolic acceleration and deceleration are rapid. These waveforms are tri-phasic with two diastolic components.

The waveform’s rise and fall times lengthen as disease progresses. A long deceleration is a sensitive indicator of P.A.D. As disease progresses, the diastolic components are lost and the waveform becomes mono-phasic. In addition, the weak signals associated with diminished flow produce waveforms that are often more jagged in appearance.

Above is a waveform with the characteristics of abnormal PT flow. The acceleration and deceleration times are slow, the diastolic components are absent, and the waveforms have a somewhat diminished height. These waveforms are typical of a patient with mild to moderate arterial disease.
The characteristics of waveforms acquired at the DP are similar. Below is an example of normal DP waveforms. This is a bi-phasic waveform from a normal patient. The signal is pulsatile with a pronounced diastolic component. The absence of the second diastolic component without additional evidence is not an indication of disease.

Continuous-wave, Doppler waveforms are often considered to be the most useful type of physiological waveform for detecting P.A.D. Doppler waveforms are given a Recommendation of Class I in the ACC/AHA P.A.D. Guidelines (Hirsch, 2005).

**Pulse Volume Recording (PVR) Waveform Interpretation:**

PVR serves as a qualitative aid in the assessment of peripheral hemodynamics. The waveform contour is useful to confirm the results of the ABI and is a valuable diagnostic tool for patients with noncompressible vessels (ABI > 1.30). Additionally, the PVR can be used to monitor limb perfusion after revascularization.

Normal plethysmographic waveforms resemble the arterial pressure pulse. The normal PVR waveform (shown below) has a sharp systolic upstroke, and a somewhat narrow peak. After peaking, the waveform drops quickly during diastole with its slope bowing in *toward* the baseline. Normal waveforms usually have a dicrotic notch between the peak and baseline. However, the dicrotic notch may become less noticeable with vasodilation or may become more pronounced during vasoconstriction.

Distal to an arterial occlusion, the waveform becomes more rounded with a more gradual up slope, and the down slope may bow *away* from the baseline. Amplitude of the abnormal waveform (shown below) is usually diminished, but this condition is not sufficient for diagnosis because amplitude is influenced by additional factors such as blood pressure, arrhythmia, vasomotor tone, and limb position.
Again, note that the waveform contour may be altered during vasodilation or vasoconstriction. Improper cuff size, over- or under-inflation, low cardiac stroke volume, and improper cuff application may affect the waveform contour.

PVR waveforms are given a Recommendation of Class IIb in the ACC/AHA P.A.D. Guidelines (Hirsch, 2005).

NOTE: Although Doppler waveforms are given a higher recommendation than PVR waveforms, PVR waveforms may be easier to obtain.

**PPG Waveforms:**

PPG waveforms are made available on the *Vista AVS* as an aid for obtaining systolic pressures at the toe. PPG waveform contour interpretation is similar to the interpretation of PVR waveforms. However, because the PPG waveform is obtained at a digit, distortion of the PPG contour due to vasoconstriction or vasodilation is more common.

PPG waveforms are not acceptable for meeting reimbursement criteria for the lower extremity arterial exam; however PPG is useful for obtaining pressures as described in Section 7. Toe pressures obtained with PPG meet reimbursement criteria if accompanied by either Doppler or PVR waveforms.

For these reasons, Wallach Surgical Devices recommends using PVR and Doppler modes for waveform analysis.
SECTION 13: Maintenance and Cleaning

Cleaning Precautions:

WARNINGS

• The Vista AVS is not designed for liquid immersion. Do not soak or drop the probes or main unit in liquids. Do not use solvent cleaners. Do not use products containing bleach. Use only the recommended spray or wipe cleaners and disinfectants (see “Cleaning and Disinfection” below).

• The Vista AVS is not designed for sterilization processes such as autoclaving, irradiation or hydrogen gas.

• The Vista AVS is not intended for use on open skin. If there is evidence of open wound contamination, disinfect the probes before using again as described below.

Cleaning and Disinfection:

Excess gel should be wiped off the ultrasound probes after every examination. Probes and main unit should be cleaned with a cloth dampened with warm water or with pre-saturated isopropyl alcohol wipes. Be sure to clean carefully along any seams, but do not allow liquids to enter the unit through connectors or speaker grill.

To disinfect the unit, use commercially available spray or wipe disinfectants that are registered with the U.S. Environmental Protection Agency (EPA). Clorox® Broad Spectrum Quaternary Disinfectant is the only disinfectant that is Wallach Surgical Devices approved for use with the Vista products. Follow the instructions provided with the disinfectant and be sure to wipe the unit dry when finished.

Vista AVS users should wash hands and change gloves after every exam. Refer to local and hospital policies regarding cleaning and disinfection.

Store the unit in a clean area free of dust and debris. Follow the temperature and humidity storage requirements as specified in Section 15.

Maintenance:

CAUTION

• If the unit is to be stored for longer than 90 days without use, remove the battery pack prior to storage.

Replace the battery pack annually or when its capacity is noticeably diminished.

A qualified technician must verify the accuracy of the pressure readout annually. Return the unit to Wallach Surgical Devices for calibration if needed. See Section 15 for applicable pressure specifications.
Periodically -- not less than annually -- inspect the main unit and probes for signs of cracks or breakage. Inspect cables, tubing, and connectors for signs of wear or failure. Inspect labels for damage. The user should discontinue use of the unit or probes if any loss of housing integrity is detected.

Check the tightness of the screw that attaches the mounting bracket to the rolling stand’s pole (if applicable) at least annually.

In order to preserve, protect and improve the quality of the environment, protect human health and utilize natural resources prudently and rationally – do not dispose of waste electrical or electronic equipment (WEEE) as unsorted municipal waste. Contact local WEEE disposal sites.
SECTION 14: Battery Recharging and Replacement

The external power supply (Product Number B200), available from Wallach Surgical Devices, becomes the power source when the Vista AVS is operated while the supply is connected. When the external power supply is connected while Vista AVS is off, the supply provides power for recharging. Plug the supply into the + 7 VDC (POWER) connector.

The green CHARGING indicator is held ON during charging. The indicator flashes during trickle charge. The CHARGING indicator will turn off when charging is complete.

The Vista AVS uses a 7.2 V six-cell Nickel-Metal Hydride (NiMH) battery pack. This battery pack has sufficient capacity to allow the product to be used for 12 ABI exams. The system charges a fully depleted battery in 3 hours. After the charging is completed, a maintenance charge (sometimes called a “trickle charge”) is applied.

When the remaining battery life is less than 15 minutes, the yellow LOW BATT indicator on the front panel will flash. When the battery life is severely depleted, the flashing rate will increase. At this point, plug in the external power supply immediately to avoid losing any unsaved exam data.

NOTE: The Vista AVS will not print when the battery is low to ensure that there is no loss of data.

VISTA TIP: 
- Once the LOW BATT indicator begins to flash, the battery may become depleted rapidly if a mode that requires additional power is selected such as PUMP. Plug in the external power supply as soon as possible once the LOW BATT indicator begins to flash.

When operating from the external power supply, the Vista AVS will switch over to battery power automatically if there is a power outage.

NOTE: Charging will not begin if the battery temperature is less than 0 °C (32 °F).

NOTE: It is important not to unplug the Vista AVS from the wall during an exam. This may result in system reset.
Replacing the battery:

*Tool required: Phillips head screwdriver*

To replace the battery, use a Phillips head screwdriver to remove the battery compartment screw. Press the latch on the side of the battery connector and unplug the old battery. Plug in the new battery and close the battery door. Tighten the screw firmly but do not over tighten. Charge the new battery for 2 hours before use.

**CAUTIONS**

- The battery pack contains a temperature measurement component that is matched to the cell type and to the product requirements. Use only replacement battery packs as specified by Wallach Surgical Devices.

- Unauthorized battery substitutions will void the warranty and may cause overheating.

- Used NiMH batteries must be recycled or disposed of properly. Do not incinerate.
SECTION 15: Specifications

Note: Unless otherwise indicated, all specifications are subject to change without notice.

System Functionality:

Ankle Brachial Index
- Ankle and Arm Systolic Pressures
- Automatic ABI Calculation
- Automatic Selection of Arm Pressure (higher of the two)
- Configurable for one or two Ankle Pressures

Toe Brachial Index
- Arm and Toe Systolic Pressures
- Automatic TBI Calculation
- Automatic Selection of Arm Pressure (higher of the two)

Seated ABI
- Ankle and Arm Systolic Pressures
- Automatic Hydrostatic Correction and ABI Calculation
- Automatic Selection of Arm Pressure (higher of the two)
- Configurable for one or two Ankle Pressures

Segmental Studies
- Arm and 5 Lower Limb Systolic Pressures
- Automatic Index Calculation
- Automatic Selection of Arm Pressure (higher of the two)
- Ability to customize study by selecting the number and location of arterial sites

Waveforms
- Bi-Directional Doppler
- Pulse Volume Recording (PVR)
- Photoplethysmography (PPG)

Reporting
- Index and Waveform Printing on Label Paper
- Optional Download Capability to PC
- Ten patient examination storage locations

Doppler:

Doppler Type: Bi-Directional, Continuous-Wave (CW)
Audio Bandwidth: >4 KHz
Waveform Display: Digital Mean Frequency Estimator
Separate Forward and Reverse Flow Waveforms
Available Probe Frequencies: 8.43 MHz, 5.5 MHz
Displayed Waveform Amplitude Accuracy: ±10%
Speaker Output: > 0.25 W
Speaker Type: 2”, 5 W
Cable length: 6.5 ft
Volume Control: Continuously variable with mute

**Photoplethysmography (PPG):**

Wavelength: 940 nm
Artifact Rejection: Synchronous demodulation, AC coupled output
Operating Frequency: 1 kHz
Cable length: 6.5 ft

**Pressure:**

Range: 0 – 265 mmHg
Accuracy: 0 – 200 mmHg: ±3 mmHg
above 200 mmHg : ±2%
Safety: Pressure released if greater than 280 mmHg
Pressure released if held above 100 mmHg beyond 3 minutes
Pressure released if held between 30-100 mmHg beyond 5 minutes*
(*Operator may over-ride for exam purposes)
Redundant Control
Bleed Rate: 2.5 mmHg/sec nominal
PVR: 0.16 to 12.5 Hz bandwidth, AC Coupled

**Display:**

Size: 5.7”, ¼ VGA (320 x 240 dot)
Backlight: LED Backlight
Type: Transflective Monochrome

**Printer:**

Type High Speed Thermal Array
Paper Width: 58 mm
Print Width: 48 mm
Resolution: 8 dots/mm
Paper Type: Thermal, Product Number K180, Adhesive Backed Label or Thermal, Archival, 58 mm

**CAUTION**

- Use only the specified paper supplied by Wallach Surgical Devices. Use of improper paper may damage the print mechanism.
**Data Storage and Transfer:**

Storage: 10 storage locations, 16 numeric or 24 alpha character length filenames

Type: USB 2.0

Connector: Peripheral

Output: AVS Report Software – Adobe PDF or comma delimited files

**Power (External Supply):**

Type: Medical Grade Supply

Rated Mains Supply Voltage: 100-240 VAC, 50-60 Hz, 1.2 A

Output: +7 V, 5 A

**Battery:**

Type: 6-cell NiMH pack

Rating: 7.2 V, 2000 mAH

Replacement Part No: B200

Battery Life: 3 hours continuous use or minimum of 12 ABI Exams

Replacement Interval: 1 year

Charge Time: 3 hours

**Mechanical and Environmental:**

Dimensions: 10” H x 12” W x 3.6” D

Weight: 4.75 lbs.

Stand height: 47”

Operating Conditions: Temp. 10 to 30 °C, Humidity 30 to 75% non-condensing

Transport & Storage Conditions: Temp. –20 to 50 °C, Humidity 5 to 90% non-condensing
Safety

Degree of protection against electric shock:

- Type B Applied part

Classification:

- Class II Equipment - when externally connected
- Internally Powered Equipment - when not externally connected (battery powered)

Degree of protection against ingress of water: Ordinary (not rated)

Designed and tested to meet: IEC601-1, IEC60601-1-2, IEC60601-1-4, IEC60601-2-37. EMC Classification: Class A

This equipment was tested to applicable standards for electromagnetic interference. If interference occurs, attempt to eliminate the source of the interference or increase the distance between the Vista AVS and the source of the interference.

WARNING

- Connect the Vista AVS only to equipment that meets the appropriate standards. The Vista AVS and its External Power Supply are a medical system. The External Power Supply must meet IEC60601-1 construction requirements for Reinforced Insulation at minimum in order for the system to comply with the appropriate safety standards.

⚠️ Attention: Consult Accompanying Documents
Transducer Model: LifeDop® 5 MHz Bi-Dir
Application(s): Peripheral Vascular

**Operating Mode:** Continuous-Wave (cw)

### ACOUSTIC OUTPUT

<table>
<thead>
<tr>
<th>Global Maximum Value</th>
<th>MI</th>
<th>ISPTA.3 (mW/cm²)</th>
<th>ISPPA.3 (W/cm²)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0.05</td>
<td>500</td>
<td>0.5</td>
</tr>
</tbody>
</table>

**Associated Acoustic Parameter**

- \( P_{3.1} \) (Mpa)
- \( w_o \) (mW)
- \( f_c \) (MHz)
- \( z_{sp} \) (cm)
- Beam Dimensions \( x_6 \) (cm)
- Beam Dimensions \( y_6 \) (cm)
- EBD

**Pr.3** the peak rarefractional pressure (megapascals) associated with the transmit pattern giving rise to the value reported for MI.

**Wo** the total time-average ultrasonic power (mwatts).

**EBD** the entrance beam dimensions (cm). These dimensions are the same as the dimensions of the transmit crystal.

**Measurement Uncertainties:**
- Power: ±34, -42%
- Pressure: +11, -16%
- Intensity (ISPTA): ±23, -26%
- Frequency: ±5%

Acoustic Output Parameters are measured in water. Derated values, denoted by the subscript “.3”, take into account a conservative level of attenuation that would be encountered in the human body. The derated intensity values (\( I_{.3} \)) are obtained from water values of intensity (\( I_w \)) at a depth of \( z \) calculated by:

\[
I_{.3} = \exp(-0.23\times0.3\times f\times z) \times I_w
\]

(where \( f \) is the probe frequency in MHz and \( z \) is the depth in centimeters)

The derated peak rarefractional pressure is calculated from the value of measure water (\( P_r \)) by:

\[
P_{r.3} = \exp(-0.115\times0.3\times f\times z) \times P_r
\]

(Where pressure is given in megapascals)

Additional Output Reporting Information for IEC 61157

8 MHz: \( I_{ob} < 112 \text{ mW/cm}^2 \)

Note that parameter \( z_{sp} \) (in the probe reporting tables) is the same parameter as \( I_p \) in IEC 61157.

**Operating Conditions:** There are no user controls which affect the ultrasound output.
SECTION 16: Accessories

The following accessories and replacement items are available from Wallach Surgical Devices. Contact us at 1-800-243-2463 or (203) 799-2000, or order on-line at www.wallachsurgical.com.

Cuffs and Tubing:
- Cuff, 10 cm        CUF0002
- Cuff, 12 cm        CUF0003
- Cuff, 17 cm        CUF0008
- Cuff, 1.9 cm        CUF0004
- Cuff, 2.5 cm        CUF0005
- Cuff, 1.9 cm        CUF0009
- Hose Set        ASY0008

Software:
- AVS Report Software with DICOM install (by Special Order only) MKT0210

Cable:
- USB, 6 ft        CBL0024

Printer Paper:
- ABI Printer Label Paper, Box of 5 rolls    K180

Gel:
- Ultrasound Gel, 250 g        G155
- Ultrasound Gel, 60 g        G150

Probes: (serial numbers required)
- Bi-Directional Probe, 8 MHz (for Vista product line) SD8V
- Bi-Directional Probe, 5 MHz (for Vista product line) SD5V
- PPG Probe        SPPG

Battery Pack:
- Battery, Vista AVS, 7.2 V        B200

Rolling Cart:
- Roll-stand with Basket        K270

Carrying Case (16” x 12” x 4”):
- For the Vista AVS only; (not for the pole, roll-stand and basket) K280
SECTION 17: Troubleshooting Guide

Troubleshooting:

WARNINGS

- Use alternate equipment in case of unit failure. Call Wallach Surgical Devices’ Service Department if the unit or probes malfunction.

- **Risk of Electric Shock** - the Vista AVS contains no user serviceable components. Only qualified personnel should attempt to service the unit.

CAUTION

- Do not drop or mishandle the Vista AVS probes. Damage may occur.

### Vista AVS Troubleshooting Guide

<table>
<thead>
<tr>
<th>Symptom:</th>
<th>Check for:</th>
</tr>
</thead>
</table>
| Doppler is not audible when user rubs probe tip | - Doppler mode is not selected (see pg. 14 for selecting modalities)  
- Mute is selected  
- Probe cable is not plugged into the main unit or plugged into an incorrect part  
- USB cable is connected to PC |
| Cannot obtain PPG signal at toe | - TBI mode is not selected  
- Probe cable is not plugged into main unit  
- Verify PPG signal can be obtained on a finger – if yes, try warming the foot |
| Unit will not print | - Printer out of paper  
- Paper coming off roll too loosely – tighten roll  
- Incorrect paper type  
- Paper roll inserted improperly |
| Unit will not pump | - Verify that the site cursor is at a BRA or ANK position or at WAV with PVR (i.e. not in Doppler waveform mode) – also ensure that the hose connection is secure |
| Pump runs but cuff does not inflate | - Verify that both ends of the cuff hose are plugged in correctly to the Vista AVS and the cuff |
| Unit will not download | - Verify that Host PC is running AVS Report Software, Software key has been entered properly, USB cable is connected |
| Cannot save 2nd ankle pressure | - Verify that DUAL ANKLE PRESSURES is not set to OFF. |
| Unit shuts off | - Power Down is set to ON or the battery is low |
| Faint Doppler signal audible, but waveform is not displayed | - There is normally a slight noise threshold that must be overcome prior to a waveform being displayed |
| Printed waveform differs from displayed waveform | - The printed waveform always includes two additional seconds of waveform data |
| Doppler Audio cuts out when attempting to take an ankle pressure or waveform | - Verify proper exam configuration. |
| Unit pumps too slowly | - PPG using digit cuffs automatically inflate slowly – switch to a digit cuff |
| If you press “ENTER/MENU” during a segmental study, you will return to the Main Menu | - If you did not want to go to the Main Menu, press “ENTER/MENU” again to return to the segmental study. |
SECTION 18: Clinical References

Clinical References:


General Information about P.A.D:

National Institute of Health (NIH) Stay in Circulation Campaign: www.aboutpad.org

P.A.D. Coalition: www.padcoalition.org

The Vascular Disease Foundation (VDF): www.vdf.org

The AVS Report software allows the clinician to download examination results from the Vista AVS and annotate patient data, add clinician interpretation and comments, store and print a patient examination report.

WARNINGS

• User-supplied external hardware used to operate the AVS Report software should conform to recognized standards for safety such as IEC601-1, IEC60065, UL2601-1 or UL544.

• The Vista AVS shall not be connected to the user-supplied external hardware during an examination.

• The Vista AVS and AVS Report software shall be operated by trained personnel who have read all manufacturer-supplied documentation.

• The AVS Report software does not provide an interpretation of the examination results. Results should be verified prior to performing additional procedures on the patient.

CAUTIONS

• In the event of AVS Report software transmission error, the data is invalid and shall be re-transmitted.

• To ensure integrity of data on the user-supplied external hardware, the equipment should be backed up frequently.

• To ensure proper operation of the user-supplied external hardware, the equipment should run virus checking software and system diagnostics frequently.

• When the AVS Report application is running the Windows® Standby and Hibernate functions are disabled.

Getting Started:

Minimum Requirements:

Operating system supported only with a 64 or 32-bit Windows® 7 Home Premium or better, a 32-bit Windows® XP Professional, and a 32-bit Windows® Vista Home Premium or better.
Installation instructions:

Insert the **AVS Report** installing disk into the CD ROM Drive. Installation will start automatically and the following screen will appear:

**NOTE:** The standard installation does not include DICOM formatted file option. To install the DICOM feature, contact Wallach Surgical Devices.

1. When you begin to install the **AVS Report** software, this screen will appear.

2. Select “Next” to continue the installation.

3. Select “Install” to load to the **AVS Report** software program to your computer.
4. This screen will appear as the installation is in process.

5. This screen will be displayed briefly just before the Installshield Wizard is complete.

6. When this screen appears, the AVS Report software has been successfully installed. Select “Finish” to complete the installation process.

Connecting to Vista AVS:
Using the USB cable provided, connect the equipment end of the cable to the Vista AVS and the computer end of the cable to an unused USB Port on the PC. Once the cable is connected to both devices, turn on the Vista AVS. When the computer successfully recognizes the device, the following message will appear:
Starting **AVS Report**:
If the installation was completed properly, an application icon is already on your computer desktop. Double click on the icon to start the software.

**Generating a Report:**

USB Communication:
When the **Vista AVS** is on and connected to the computer while the **AVS Report** software is running, the bottom of the screen will indicate “Connected”. The “Read Success” indicates that the communication channel is open and operating properly.

<table>
<thead>
<tr>
<th>12/26/2006</th>
<th>1:35 PM</th>
<th>Connected</th>
<th>Read Success</th>
</tr>
</thead>
</table>

If the indication is “Disconnected” then the software was started without the **Vista AVS** connected. PLEASE NOTE: The software can be operated without the device connected for opening previously generated files, printing or importing as described below.

<table>
<thead>
<tr>
<th>12/26/2006</th>
<th>1:37 PM</th>
<th>Disconnected</th>
</tr>
</thead>
</table>

If the indication is “Write Call Failed” then the **AVS Report** has lost communication with the **Vista AVS**. Check to ensure the cables are connected at both ends and ensure the unit is on.

<table>
<thead>
<tr>
<th>12/26/2006</th>
<th>1:36 PM</th>
<th>Write Call Failed</th>
<th>-- POLL --</th>
</tr>
</thead>
</table>

**Data Download:**
On the **Vista AVS** OPTIONS menu, select FILE and DOWNLOAD to transfer data that is currently active. The **AVS Report** software will briefly indicate “Upload Success” to verify that the data was accepted.

<table>
<thead>
<tr>
<th>12/26/2006</th>
<th>2:05 PM</th>
<th>Connected</th>
<th>Upload Success</th>
</tr>
</thead>
</table>
If a transmission error occurs, the report software screen will indicate that the data verification failed. Check to ensure the USB cable is securely connected on both ends and download the file again. Contact Wallach Surgical Devices if the error continues to occur.

Report data is segmented into 4 Sections of the software and separated by tabs at the top of the application. Select the appropriate tab to access or enter information.

Patient Info – User editable information regarding the patient, general comments and the examiner identification.
Pressures/Waveforms – Non-editable pressure, waveform and ABI information from the Vista AVS downloaded file.
CPT-ICD9 Codes – User editable billing information to appear on the report.

Patient Information (“Patient Info” Tab):
Patient Name or ID will be transferred from the Vista AVS file information and are shown in the Patient Info Tab. Other information can be annotated at this time such as:

Patient Name -
Date of Birth -
Age -
Sex -
I.D. Number -
Date/Time -
Interpreting Physician -
Referring Physician -
Patient Name or Patient ID will be downloaded directly from the Vista AVS file information if it was saved on the device.

Risk Factors –
Check the appropriate boxes to enter the patient’s P.A.D. risk factors. The “Other” section can be used for extra details such as pack years of smoking or date of previous CV event.

Current Symptoms –
Check appropriate boxes to enter the patient’s P.A.D. symptoms and the location of the symptoms. The “Other” section can be used for extra details such as when or how often symptoms occur.

Examiner Comments –
General comments and interpretation sections are provided for extended comments from the interpreting clinician or from the examiner.

Pressures, Waveforms and ABI Values (“Pressures/Waveforms” Tab):
The ankle and brachial pressures download from the Vista AVS are included in the Pressure/ABI Tab along with the modality used to take the pressures, Doppler or PPG. A reference table is also displayed for either ABI or TBI clinically accepted values as an aid for the clinician in determining a diagnosis.
Segmental Study

CPT - ICD-9 Codes (“CPT - ICD-9 Codes” Tab):
The CPT and ICD-9 Codes Tab allows the user to select a billing option to be displayed on the report. Select the tabs at the bottom of the page to choose either CPT or ICD-9. Select the appropriate code on each screen or type a code in the “Other” section if the desired code is not available. Only one selection can be valid on any given report.
Managing a Report:

File Menu:

Open:
Allows the user to open a previously saved report.

Save/Save As:
Allows the user to select the report name and location for later recovery and backup.

Close:
Allows the user to close out of the currently opened file.

Export:
Allows the user to generate an Adobe® PDF file, DICOM file or Comma Delimited file for export. See below for further explanation on exporting files.

Print:
Allows the user to print the report on the selected printer.

Print Preview:
Allows the user to view the completed report prior to printing. Zoom in and out, scroll pages or print directly from the preview window.

Printer Select:
Allows the user to change the default printer.

Exit:
Allows the user to exit the application.
Preferences Menu:

Allows the user to enter facility information that will be included in the report such as facility name and address. This information will be stored in the application and used on each subsequent report that is generated until the information is changed.

The File Paths menu allows the user to select a default directory for each file format to save exams.

The above screen allows the user to enter facility information that will be included in the report such as the facility name and address. This information will be stored in the application and used on each subsequent report that is generated until the information is changed.

The File Paths menu allows the user to select a default directory for each file format to save exams.

The above screen allows the user to enter physician names to be used for referring or interpreting physician on the patient information tab. Select “Add” to enter a new name.

Enter the information in the fields and click “OK”.
This screen allows a user to customize the severity table shown in the report. The default is from the ACC/AHA guidelines. To customize, select “Custom”.

- To edit the text line, double-click the text line.
- To delete the text line, right-click the text line and select Delete.
- To add a new line, click on “New Line”, left-click and drag to a new location.
- Press “OK” when finished editing a line of text.

This screen will appear next. Click “OK” when complete.

**Exporting a Report:**

There are three file export types that are supported: Adobe PDF, DICOM and Comma Delimited files (.csv). NOTE: The saved files will default directly into the folder initially created in the File Path menu to store the AVS Report software.
Adobe PDF (2 Options):

1. Select **File>Export>Adobe Acrobat (.pdf)>Export and Launch Acrobat** to immediately view the PDF file on your computer screen.

2. Select **File>Export>Adobe Acrobat (.pdf)>Export Report File Only** to save/store the file to view at a later time.

   - Enter a file name and save it to convert the current file to this format and the Adobe application (if it is currently loaded on the PC) will automatically start.

   - If the application is not loaded, a free reader version can be downloaded from the website: www.adobe.com

   - Once the file has been converted and opened into the Adobe application, all features and functions of that application are supported. The file can be saved, printed, e-mailed, etc.

Comma Delimited Files:
Select **File>Export>Comma Delimited (.csv)**. Enter a file name and save it to convert the current file to a delimited format that can be read using any software application for viewing text files, such as Microsoft Word®, Microsoft Excel® and Notepad. These applications do not automatically open once the file has been saved. This file can be used to configure data into EMR systems. Special programming may be required by the system administrator.

DICOM (Digital Imaging and Communications in Medicine)
Select **File>Export>DICOM (.dcm)**. Enter a file name and save it to convert the current file to a DICOM format that can be connected to a DICOM system (i.e. PACS). A DICOM viewer must be used to open and review these files.

**AVS Report DICOM Conformance Statement:** The DICOM portions of the **AVS Report** application were developed using software tools supplied by Lead Technologies, Inc., a leading supplier of industrial medical imaging applications. DICOM files created by **AVS Report** use the Secondary Capture Image Storage SOP Class. These files have been evaluated using the conformance test supplied by Lead Technologies and meet all applicable criteria of this SOP Class. Additionally, **AVS Report** files have been successfully evaluated using several independent DICOM viewing applications. As these standards are constantly evolving, all efforts have been made to verify the DICOM files and data elements conform to the latest published DICOM standards for this SOP Class.
Completed Sample ABI Report in PDF File Format

Lower Extremity Physiologic Study
Single Level (ABI Exam)

Patient Name: Mary Johnson
Patient DOB: 05/31/1942
Patient Sex: F
Patient ID: 1234567890
Age: 60
Exam Date: 03/01/10 09:31:06

Risk Factors
- Tobacco Use
- Diabetes
- Heart Disease
- Previous CV Event
- 1 pack per smoker

Current Symptoms
- Hypertension
- Hyperlipidemia
- Stroke/TIA
- Prev. Vascular Surgery
- Intermittent Claudication
- Ulcerations
- Rest Pain
- Gangrene
- Resting pain at night

Location
- Femoral
- Thigh
- Calf
- Foot

ABI Severity of Disease
- > 1.3 Non-compressible
- 1.0 - 1.29 Normal
- 0.91 - 0.99 Borderline
- 0.41 - 0.90 Moderate to Mild
- 0.06 - 0.40 Severe

Interpreting Physician: Dr. John M. Smith
Referring Physician: Dr. John M. Smith
Comments: No evidence of PAD

Right:
- Brachial: 123 mmHg
- PT: 135 mmHg
- DP: 132 mmHg
- Index: 1.09

Left:
- Brachial: 120 mmHg
- PT: 138 mmHg
- DP: 143 mmHg
- Index: 1.16

Higher ABI: 1.16

Interpretation:

CPT Code: 93922 Non-invasive physiologic studies of upper or lower extremity arteries, single level, bilateral.
ICD-9 Code: 440.22 Atherosclerosis of native arteries of the extremities with rest pain
Completed Sample Seated ABI Report in PDF File Format
Completed Sample Segmental Study Report in PDF File Format
Example of data from a comma-delimited file:

<table>
<thead>
<tr>
<th>SubTitle</th>
<th>Ankle Brachial Index Assessment Form</th>
</tr>
</thead>
<tbody>
<tr>
<td>PatientName</td>
<td>John Smith</td>
</tr>
<tr>
<td>PatientID</td>
<td>123-45-6789</td>
</tr>
<tr>
<td>Severity</td>
<td>ABI Severity</td>
</tr>
<tr>
<td>Severity1</td>
<td>0.90 - 1.30 - Normal</td>
</tr>
<tr>
<td>Severity2</td>
<td>0.70 - 0.89 - Mild</td>
</tr>
<tr>
<td>Severity3</td>
<td>0.40 - 0.69 - Moderate</td>
</tr>
<tr>
<td>Severity4</td>
<td>0.0 - 0.39 - Severe</td>
</tr>
<tr>
<td>Results</td>
<td>ABI Results</td>
</tr>
<tr>
<td>Comments</td>
<td>Right side P.A.D. Spoke to patient re: life style modifications Recommend further evaluation</td>
</tr>
<tr>
<td>CPTCode</td>
<td>93922 Non-invasive physiologic studies of upper or lower extremity arteries, single level, bilateral</td>
</tr>
<tr>
<td>ICD9Code</td>
<td>440.21 Atherosclerosis of native arteries of the extremities with intermittent claudication</td>
</tr>
<tr>
<td>RightSite</td>
<td>PT</td>
</tr>
<tr>
<td>LeftSite</td>
<td>PT</td>
</tr>
<tr>
<td>RightProbe</td>
<td>8 MHZ</td>
</tr>
<tr>
<td>LeftProbe</td>
<td>8 MHZ</td>
</tr>
<tr>
<td>ABIRightArm</td>
<td>154</td>
</tr>
<tr>
<td>ABILeftArm</td>
<td>148</td>
</tr>
<tr>
<td>ABIRightToeAnkle</td>
<td>Right Ankle</td>
</tr>
<tr>
<td>ABIRightTA</td>
<td>96</td>
</tr>
<tr>
<td>ABILeftToeAnkle</td>
<td>Left Ankle</td>
</tr>
<tr>
<td>ABILeftTA</td>
<td>158</td>
</tr>
<tr>
<td>LeftTAP</td>
<td>Left Ankle Pressure</td>
</tr>
<tr>
<td>RightTAP</td>
<td>Right Ankle Pressure</td>
</tr>
<tr>
<td>LeftTAPVal</td>
<td>158</td>
</tr>
<tr>
<td>RightTAPVal</td>
<td>96</td>
</tr>
<tr>
<td>HAPValue1</td>
<td>154</td>
</tr>
<tr>
<td>HAPValue2</td>
<td>154</td>
</tr>
<tr>
<td>LeftABIValue</td>
<td>1.02</td>
</tr>
<tr>
<td>RightABIValue</td>
<td>0.62</td>
</tr>
<tr>
<td>PatientAge</td>
<td>67</td>
</tr>
<tr>
<td>Date</td>
<td>12/27/06 10:02:54</td>
</tr>
<tr>
<td>RiskOther</td>
<td>2 pack per day smoker</td>
</tr>
<tr>
<td>SymptomOther</td>
<td>Max walking distance - 1 block</td>
</tr>
<tr>
<td>HospitalName</td>
<td>St. Joseph</td>
</tr>
<tr>
<td>HospitalAddr</td>
<td>1234 Main Golden</td>
</tr>
<tr>
<td>SummitAddr</td>
<td>4680 Table Mountain Drive #150 Golden, CO 80403</td>
</tr>
<tr>
<td>Phone:</td>
<td>303.423.7572</td>
</tr>
<tr>
<td>Fax:</td>
<td>303.940.7165</td>
</tr>
<tr>
<td>Toll Free:</td>
<td>800.554.5090</td>
</tr>
<tr>
<td>Examiner</td>
<td>Kathy C. - RVT</td>
</tr>
<tr>
<td>Modality</td>
<td>Pressures – Doppler</td>
</tr>
</tbody>
</table>
SECTION 20: Warranty

The Vista AVS is warranted against defects in material and workmanship for 12 months from the original sale of the device. Product life is specified to be 5 years from manufacture, though the device may be repairable beyond this timeframe. This includes all parts and labor required to repair or replace the unit to original specifications and shipping costs associated with sending the product back to the customer. The customer is responsible for providing adequate packaging materials and shipping costs to Wallach Surgical Devices. Products shall be repaired or replaced in a reasonable amount of time.

Wallach Surgical Devices’ liability for any claim is limited to materials and labor associated with repair or replacement. In no event shall Wallach Surgical Devices be liable for incidental or consequential losses or damages in connection with the purchase of this product.

Wallach Surgical Devices disclaims all expressed or implied warranties, agreements or arrangements other than issued in this warranty.

Wallach Surgical Devices is not responsible for damages to the device that occur as a result of the inadequate packaging on return shipments to Wallach Surgical Devices, improper maintenance or cleaning as described in the user manual, misuse, abuse, alteration of the equipment from its original specifications, or dismantling the unit (other than by Wallach Surgical Devices approved service technicians).

If you need to return the Vista AVS or one of its components or accessories for repair:

1) Call Wallach Surgical Devices to obtain a Return Authorization (RA) and any final instructions prior to shipping.
2) Clean the product before shipping.
3) Ensure the product is well packaged and suitable for shipment.

Note: The customer is responsible for any damage incurred during return shipment associated with improper packaging.

Send the product to:
   Repair Department
   Wallach Surgical Devices
   95 Corporate Drive
   Trumbull, CT 06611 USA

For customer service, technical service, cleaning or maintenance or shipping questions, please call (203) 799-2000 or 1-800-243-2463.
SECTION 21: Explanation of Symbols

<table>
<thead>
<tr>
<th>REF</th>
<th>Reorder Number</th>
</tr>
</thead>
<tbody>
<tr>
<td>SN</td>
<td>Serial Number</td>
</tr>
<tr>
<td>Latex Free</td>
<td>Class II Equipment</td>
</tr>
<tr>
<td>![Symbol] Keep Dry</td>
<td></td>
</tr>
<tr>
<td>![Symbol] Type B Applied Part</td>
<td></td>
</tr>
</tbody>
</table>

ATTENTION:
See instructions for use.

Manufacturer

Date of Manufacture

Symbol indicates that the device should be sent to the special agencies according to local regulations for separate collection after its useful life.

Product conforms to the Medical Device Directive 93/42/EEC

Authorized Representative in the European Community.

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