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5th EDITION

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# Vascular Technology An Illustrated Review

# 5th Edition

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# **Contents**

Reviewers vii

Preface to the 5th Edition ix

Acknowledgments xi

# PART I ARTERIAL EVALUATION 1

- Chapter 1 Gross Anatomy of the Central and Peripheral Arterial Systems 3
- Chapter 2 Physiology and Fluid Dynamics 15
- Chapter 3 Testing Considerations, Patient History, Mechanisms of Disease, and Physical Examination 43
- Chapter 4 Doppler Waveform Analysis in the Upper and Lower Extremities 57
- Chapter 5 Doppler Segmental Pressures—Lower Extremities 69
- Chapter 6 Doppler Segmental Pressures—Upper Extremities 81
- Chapter 7 Laser Doppler 89
- Chapter 8 Penile Pressures and Waveforms—Duplex/Color Flow Imaging Evaluation 93
- Chapter 9 Plethysmography—Upper and Lower Extremities 99
- Chapter 10 Digital Pressures and Plethysmography 107
- **Chapter 11** Transcutaneous Oximetry (TcPO<sub>2</sub>) **115**
- Chapter 12 Duplex Scanning and Color Flow Imaging of the Upper Extremities 119

Chapter 13	Duplex Scanning and Color Flow Imaging of the Lower Extremities <b>127</b>
Chapter 14	Duplex Scanning and Color Flow Imaging of the Abdominal Vessels 139
Chapter 15	Preoperative Mapping Procedures 159
Chapter 16	Atypical Vascular Disorders 165
Chapter 17	Alternative Diagnostic Tests and Therapeutic Interventions 173
PART II CEF	REBROVASCULAR EVALUATION 197
Chapter 18	Gross Anatomy, Physiology, and Fluid Dynamics 199
Chapter 19	Testing Considerations, Patient History, Mechanisms of Disease, and Physical Examination <b>207</b>
Chapter 20	Carotid Duplex Scanning and Color Flow Imaging 221
Chapter 21	Transcranial Doppler (TCD) 239
Chapter 22	Atypical Vascular Disorders 249
Chapter 23	Alternative Diagnostic Tests and Therapeutic Interventions 253
PART III VEI	NOUS EVALUATION 263
Chapter 24	Gross Anatomy of the Central and Peripheral Venous Systems 265
Chapter 25	Venous Hemodynamics 277
Chapter 26	Testing Considerations, Patient History, Mechanisms of Disease, and Physical Examination <b>281</b>

Chapter 27 Venous Photoplethysmography 293

- Chapter 28 Venous Air Plethysmography 297
- Chapter 29 Continuous-Wave Doppler 301
- Chapter 30 Duplex Scanning and Color Flow Imaging in Venous Evaluation 307
- Chapter 31 Alternative Diagnostic Tests 331
- Chapter 32 Medical Therapies 337
- Chapter 33 Surgical and Endovascular Therapies and Other Treatment Options 341

PART IV TEST VALIDATION, STATISTICS, AND PATIENT SAFETY 353

- Chapter 34 Statistical Profile and Correlation 355
- Chapter 35 Patient Care and Environmental Issues 363
- Appendix A Case Studies for Self-Assessment 367
- Appendix B Application for CME Credit 401
- Appendix C Bibliography 433
- Index 435

# с нарте r **2**

# Physiology and Fluid Dynamics

Arterial System

Energy

**Blood Flow Characteristics** 

Poiseuille's Law

**Reynolds Number** 

Pressure/Velocity Relationships (Bernoulli Principle)

Steady versus Pulsatile Flow

Peripheral Resistance

**Cardiac Effects** 

**Collateral Effects** 

**Effects of Exercise** 

Effects of Stenosis on Flow

# **ARTERIAL SYSTEM**

The arterial system is a multibranched elastic conduit that carries blood away from the heart and outward to the most distant tissues.

- The arterial tree oscillates with every beat of the heart, each one of which pumps approximately 70 ml of blood into the aorta and causes a blood pressure pulse.
- At the beginning of the cardiac contraction, the pressure in the left ventricle rises rapidly, quickly exceeding that in the aorta so that the aortic valve opens, blood is ejected, and the blood pressure rises. The amount of blood ejected is called the *stroke volume*.
- Increased heart rate delivers an increased blood volume that supplies more nutrients. Conversely, the lower the heart rate, the smaller the volume of pumped blood.
- The patient's cardiac status plays an important role in the movement of blood throughout the vascular system.

The heart pump generates the pressure (*potential energy*) to move the blood. The stroke volume of blood produced by each heartbeat creates a pressure (or *energy*) wave that travels rapidly throughout the arterial system (Figure 2-1):

• The propagation speed, shape, and strength of the pressure wave change as the wave moves through the arterial system.





**Example:** As the arterial pressure wave moves distally, away from the heart and out toward the periphery, the propagation speed—the pulse wave velocity—increases with the growing stiffness of the arterial walls.

- Variations in the characteristics of the vessels influence these alterations in blood flow. Velocity and flow direction also vary with each heartbeat.
- As the pressure wave moves from the large arteries through the highresistance vessels, capillaries, and then into the venous side, the mean pressure gradually declines because of losses in total fluid energy.

The pumping action of the heart maintains a high volume of blood in the arterial side of the system that in turn sustains a high pressure gradient between the arterial and venous sides of the circulation. This pressure gradient is necessary to maintain flow.

Cardiac output governs the amount of blood that enters the arterial system, while arterial pressure and total peripheral resistance (which is controlled by the level of vasoconstriction in the microcirculation) determine the volume of blood that leaves it.

A large portion of the energy created with each left ventricular contraction results in distention of the arteries, producing an arterial "reservoir" that stores some of the blood volume and the potential energy supplied to the system.

It is this store of energy and volume that promotes the flow of blood into the tissue during diastole. That is, potential energy is stored in the distended arterial wall and is released when the wall recoils.

Pressure is greatest at the heart and gradually decreases as the blood moves distally. This pressure difference (or *gradient*) is necessary to maintain blood flow.

# ENERGY

The movement of any fluid medium between two points requires two things: (1) a route along which the fluid can flow and (2) a difference in energy (pressure) levels between the two points. The volume of flow depends on the net energy difference between these two points, a factor that is affected by losses resulting from the movement of the fluid—i.e., friction—and any resistance within the pathway that opposes such movement.

The greater the energy difference (or the lower the resistance), the greater the flow, as illustrated in Figure 2-2.

The greater the pathway resistance and/or energy losses, the lower the flow.

*lower resistance = higher flow rate higher resistance = lower flow rate*  **Figure 2-2.** The greater the pressure gradient, the greater the flow. Example **A** has less pressure difference and therefore less flow than example **B**, which has a larger pressure gradient and greater flow.



The total energy contained in moving fluid is the sum of pressure (potential) energy, kinetic (movement) energy, and gravitational energy, as explained below:

- *Pressure energy* is the main form of energy present in flowing blood (see Figure 2-3A). It is created by the pumping action of the heart, which subsequently distends the arterial vessels. This distention occurs because of elastin, which allows for a rapid increase in capacity. Arteries are more elastic proximally and less distally as they become smaller. Elastin quickly converts kinetic energy to pressure energy. The pressure energy is then converted back to kinetic energy in diastole. Pressure energy is also referred to as *potential energy*. Potential energy has several components. The dominant source is the pumping action of the heart muscle, as well as distention of the arterial wall.
- Kinetic energy is the ability of flowing blood to do work as a result of its velocity; it is the energy of something in motion (see Figure 2-3B). With regard to blood flow, the kinetic energy portion is small compared to the pressure energy. Kinetic energy is also proportional to the density of blood (which is normally stable) and to the square of its velocity. The everyday example of a dam illustrates the difference between potential and kinetic energy: The water behind the dam has *potential* energy (with the height of the dam providing a form of *gravitational* energy), while the water flowing through the dam has *kinetic* energy.
- Gravitational energy (hydrostatic pressure) is expressed in millimeters of mercury (mmHg). Changes in the height of the fluid column introduce the element of gravitational energy, which is hydrostatic pressure. In the circulatory system, hydrostatic pressure is equivalent to the weight of the column of blood extending from the heart, where the right atrium is considered the 0 pressure reference point (i.e., atmospheric pressure), to the level where the pressure is being measured.

Figure 2-7. A In a popliteal artery aneurysm, resistance to flow is reduced due to the larger-thannormal diameter. Although the volume of blood flow through this vessel remains constant, velocities are decreased compared to normal. B In this superficial femoral artery (SFA) stenosis, resistance to flow is increased because of the stenotic lumen. Although the volume flow remains constant, velocities are elevated as a result of the stenosis.





# **POISEUILLE'S LAW**

Poiseuille's law defines the relationship between volume flow (Q), pressure (P), and resistance (R) and may be written as Q = P/R. It helps to answer the question of how much fluid is moving through a vessel. When combined with the resistance equation (see page 21), Poiseuille's law may be stated as follows:

$$\mathsf{Q} = \frac{(\mathsf{P}_1 - \mathsf{P}_2) \, \pi \mathsf{r}^2}{8 \eta \mathsf{L}}$$

where Q = volume flow,  $P_1$  = pressure at the proximal end of the vessel,  $P_2$  = pressure at the distal end of the vessel, r = radius of the vessel, L = length of the vessel,  $\pi$  = 3.1416, and  $\eta$  = viscosity of the fluid.

- A change in the diameter (hence, radius) of a vessel affects resistance more dramatically than viscosity or vessel length (see Figures 2-7A and 2-7B).
- The radius of a vessel is directly proportional to the volume flow. Small changes in radius may result in large changes in flow, as demonstrated in Figure 2-8.



**Figure 2-8.** Parts **A** and **B** both illustrate narrowed segments. The reduction in volume flow through segment A will be greater than that through segment B, where the radius of the stenotic segment is somewhat larger.

# **REYNOLDS NUMBER**

After the initial acceleration in systole, blood movement continues and develops into distinct streamline formations. Where P = pressure, note in Figure 2-10A that the streamlines are evenly distributed. When the flow pattern becomes unstable, these continuous streamlines break up and form small circular currents called eddy currents and vortices (swirling patterns of rotational flow) (Figure 2-10B).

Osborne Reynolds sought to determine how viscosity, vessel radius, and the pressure/volume relationship influence the stability of flow through a vessel. Although most of his work applied to straight, rigid tubes, it still provides insight into the physics of blood flow.

Flow volume increases as pressure increases, but only to a point. As flow changes from stable to disturbed, Reynolds found that an increase in pressure no longer increased flow volume. Instead, it increased flow disturbance, contributing to the formation of eddy currents.

The elements that affect the development of turbulent flow are expressed by a "dimensionless" number called the Reynolds number (Re). The factors that affect the development of turbulence are expressed by this number according to the following equation:

$$\text{Re} = \frac{V\rho 2r}{\eta}$$

where Re = Reynolds number, V = velohe density of the fluid, r = theradius of the tube, and  $\eta =$  the viscosity of the fluid. Because the density and viscosity of the blood are fairly constant, the development of turbulence depends mainly on the size of the vessels and the velocity of flow. When the Reynolds number exceeds 2000, laminar flow tends to become disturbed. Flow disturbances also can occur at lower values because of other factors, such as body movement, pulsatility of blood flow, and irregularities of the vessel wall and plaque.

Turbulent flow may cause vessel walls to vibrate. The harmonics of this vibration produce vascular bruits.





ocity, 
$$\rho = th$$

# PRESSURE/VELOCITY RELATIONSHIPS (BERNOULLI PRINCIPLE)

As previously described in the section on energy, the total energy contained in moving fluid is the sum of potential (i.e., pressure), kinetic, and gravitational energies. If one of these variables changes, the others also must change to maintain total fluid energy at the same level.

*Example:* If gravitational energy remains unchanged (that is, there is no change in the height of the fluid) but kinetic energy (velocity) increases, then potential (pressure) energy must decrease to maintain the same total fluid energy.

The Bernoulli equation shows that velocity and pressure are inversely related. Where there is high velocity, there is low pressure; where there is low velocity, there is high pressure. This inverse relationship between pressure and velocity explains why pressure decreases where fluid velocity increases (within the stenotic segment of an artery, for example) and why pressure distal to a stenosis (the region of poststenotic turbulence, where velocity decreases) is higher than that within the lesion itself. (See Figure 2-11.)

In other words, this is the law of conservation of energy:

- 1. In the region proximal to the stenosis (prestenosis), the pressure energy is higher and the kinetic energy lower. This region has the highest total energy sum.
- 2. As blood flows into the area of the stenosis, the pressure energy decreases and the kinetic energy increases (higher). However, total energy in this stenotic segment is less than that in the prestenotic segment because energy is lost (i.e., converted into heat) as it moves through the narrowing.
- 3. Lastly, distal to the stenosis (poststenosis), the kinetic energy decreases and the pressure energy increases. At this point the total energy sum is lowest.

Pressure gradients—the difference in pressure between two points in a vessel—are described as *flow separations*. Flow separations within a vessel may be



**Figure 2-11.** Velocity energy is elevated and pressure energy diminished within a stenosis. Relative velocity energy decreases and pressure energy increases distal to a stenosis. However, overall energy decreases.

Figure 2-12. Flow separation patterns. In both examples A and B, pressure energy is higher and velocity energy lower in the area of the pressure gradient (flow separation). This causes flow direction to move to the area of lower pressure energy. A Flow separation in the carotid bulb. B Flow separation at a curve (note that it is located on the inside of the vessel curve).





**Figure 2-13.** Longitudinal image of the carotid bifurcation in systole. The inset shows the same image in diastole. Note that flow separation (blue) is present during systole, absent in diastole.

**Figure 2-14.** In this reversed saphenous vein graft at the endto-side proximal anastomosis there is an area of flow stagnation (flow separation) on the inside wall. Distally, there are areas of flow separation at the site of a valve cusp. Note the variability of vessel dimension in the bypass graft. CFA = common femoral artery, SFA = superficial femoral artery. caused by changes in the geometry of the vessel (with or without intraluminal disease) or the direction of the vessel, as depicted in Figures 2-12A and B. See also Figure 2-13.

Flow separations leave behind regions of flow reversal, stagnant or little movement. Figure 2-14 demonstrates two different regions within a bypass graft where these flow separations can occur.

Because flow moves from high to low pressure (described as a *pressure gradient*), the direction of flow in the region of flow separation (e.g., carotid bulb, bypass graft anastomosis) changes with respect to the transducer, causing a visible color change in the color flow image at systole.

During diastole, when flow at the vessel wall is stagnant, there is no movement of blood and therefore no color in the color flow image. The flow separation pattern is an ideal one to use to help define whether an image is in systole or diastole.





**Figure 2-22.** An irregular heart rhythm is evident in this spectral analysis of a common carotid artery.

 A severely irregular heart rhythm (Figure 2-23), on the other hand, represents a challenge in obtaining reliable peak systolic velocity (PSV) measurements. Some protocols call for averaging a few cycles to be the most reliable; others consider the averaging of 10 cycles to be the best method. When determining the significance of a stenosis, calculating a velocity ratio may be helpful; e.g., the highest internal carotid artery PSV is divided by the PSV of the more distal common carotid artery.

# Stenosis of the Aortic Valve

- A delay in the systolic upstroke is evident (Figure 2-24).
- Decreased peak systolic velocities (PSVs) are also usually seen. Therefore, PSVs may underestimate a stenosis.



**Figure 2-23.** Severely irregular heart rhythm provides significant challenges in determining a reliable peak systolic velocity measurement.



**Figure 2-24.** A delay in the systolic upstroke is evident with a stenosis of the aortic valve. A decrease in peak systolic velocities is also usually seen.

**Figure 2-25.** Double systolic peak (pulsus bisferiens) in the common carotid artery of a patient with aortic regurgitation/insufficiency.

**Figure 2-26.** Pandiastolic flow reversal in the subclavian artery of a patient with aortic regurgitation/ insufficiency.





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# Regurgitation/Insufficiency of the Aortic Valve

- A double systolic peak, sometimes referred to as *pulsus bisferiens*, may be evident (Figure 2-25). However, a similar finding, i.e., a second systolic peak, may be a normal finding in a young patient secondary to vessel wall compliance. In addition, one author\* suggests that there may be a relationship between a double systolic peak and aortic dissection.
- Diminished diastolic flow or even reversed flow throughout diastole may also be documented on spectral analysis (Figures 2-26 and 2-27).

# High Cardiac Output

 A systemic increase in peak systolic velocities may normally be evident in younger, athletic individuals or physically fit adults (Figure 2-28). A decrease in end-diastolic velocities may also be evident in a low-resistance vessel, e.g., the internal carotid artery. As previously mentioned, a double systolic peak may also be a normal finding in a young patient secondary to vessel wall compliance (Figure 2-25).

\*Burgess W: Recommended protocol for duplex ultrasound in common carotid artery dissection extending from the aortic arch. Paper presented at the Annual Conference of the Society for Vascular Ultrasound, San Francisco, California, May 2013.

**Figure 2-27.** Significant flow reversal in the proximal aorta of a patient with severe aortic regurgitation/insufficiency.



**Figure 2-40.** Because exercise produces a demand for blood to the muscles, a normally high-resistance arterial Doppler signal becomes low-resistance. In this analog waveform, the usual reversal below the baseline is seen as a forward reflection instead. This finding is quite normal after exercise.

# **EFFECTS OF EXERCISE**

Exercise should induce peripheral vasodilation in the microcirculation so that distal peripheral resistance diminishes and blood flow markedly increases.

Peripheral resistance—the resistance to blood flow caused by the ever-decreasing size of the vessels, especially in the microcirculation—changes in response to a variety of stimuli such as heat, cold, tobacco use, and emotional stress.

Vasoconstriction and vasodilation of the blood vessels within skeletal muscles are also influenced by sympathetic innervation fibers that function primarily to regulate body temperature.

Exercise is probably the best single vasodilator of high-resistance vessels within skeletal muscle.

Autoregulation also controls vasoconstriction and vasodilation. Autoregulation accounts for the ability of most vascular beds to maintain a constant level of blood flow over a wide range of perfusion pressures.

- Autoregulation does not function normally when perfusion pressure drops below a critical level.
- High-resistance vessels constrict in response to increased blood pressure and dilate in response to decreased blood pressure.

By decreasing resistance in the working muscle, exercise normally decreases reflection (flow reversal) of the Doppler flow signal in the exercising extremity. *Example:* A low-resistance, monophasic Doppler flow signal (Figure 2-40) may be present normally in an extremity artery after vigorous exercise because the exercise causes peripheral dilatation and reduced flow resistance. This same low-resistance monophasic pattern is also seen pathologically when peripheral dilatation occurs in response to proximal arterial obstruction (see Figure 2-38).

On the other hand, a high-resistance signal (Figure 2-41) may occur from normal (physiologic) vasoconstriction at the arteriolar level or from distal arterial obstruction.

**Figure 2-41.** This high-resistance Doppler signal can occur with normal vasoconstriction at the arteriolar level. It can also occur proximal to distal arterial obstruction.



Additional notes:

- Proper characterization of velocity waveforms requires an understanding of both the normal flow characteristics of a particular artery and the physiologic status of the circulation supplied by the vessel.
- Questions need to be asked: Was the extremity cooled or warmed? Has it been exercised prior to the exam? Flow to a cool, vasoconstricted extremity will have pulsatile signals. Flow to a warm, vasodilated extremity will have continuous, steady signals.
- Proximal and distal pulsatility changes do not precisely differentiate between occlusion and severe stenosis.
- If good collateralization is present, proximal or distal Doppler velocity waveform qualities may not be altered.
- The distal effects of obstructive disease may be detectable in the presence of exercise or hyperemic evaluation.

# **EFFECTS OF STENOSIS ON FLOW**

Laminar flow review:

- Laminar flow has an even distribution of frequencies at systole, with the lower frequencies distributed at the walls (the boundary layer) and the higher frequencies in center stream.
- Stable flow through a relatively straight vessel is usually laminar, the layers of fluid slipping over one another with minimal friction normally.

A hemodynamically significant stenosis causes a major reduction in volume flow and pressure. A stenosis usually becomes hemodynamically significant when the cross-sectional area of the arterial lumen is reduced 75%, which corresponds to a diameter reduction of 50%.

- Diameter reduction is a one-dimensional measurement (Figure 2-42).
- Area reduction is a two-dimensional measurement (Figure 2-43).
- In hemodynamically significant stenoses, both pressure and flow volume decrease.



**Figure 2-42.** The nondiseased arterial segment is measured from wall to wall in a longitudinal approach (**B**) and then compared to the residual flow channel at the area of stenosis (**A**). A percentage diameter reduction is calculated.



**Figure 2-43.** The original lumen (**B**) is measured and compared to the residual lumen (**A**) when the vessel is in a transverse approach. A percentage area reduction is calculated.

# снартек **12**

# Duplex Scanning and Color Flow Imaging of the Upper Extremities

**Capabilities and Limitations** 

**Physical Principles** 

Technique

Interpretation

Brachial Artery Reactivity Testing

# **CAPABILITIES AND LIMITATIONS**

# **Capabilities**

- Localizes arterial stenosis or occlusion and evaluates degree of stenosis.
- Determines the presence or absence of aneurysm.
- Evaluates hemodialysis access graft or arterial bypass graft postoperatively.
- Detects arteriovenous fistulas or other unusual abnormalities.
- Evaluates medical treatment or surgery on a follow-up basis.

# Limitations

- Cannot be used in the presence of dressings, skin staples, sutures, or open wounds.
- Imaging is difficult around IV site.
- Hemodialysis access grafts:
  - Difficult to assess the anastomotic sites because of graft angulation.
  - Difficult to adequately evaluate the outflow vein secondary to increased collateral development.
- For diagnosis of Raynaud's phenomenon, previously described in Chapter 10 (page 113), physiologic studies combined with the clinical presentation (Chapter 3, pages 52–53) constitute the most useful diagnostic approach.

# **PHYSICAL PRINCIPLES**

- Duplex scanning combines real-time B-mode imaging (gray-scale evaluation) and Doppler spectral analysis (which is an analysis and display of the Dopplershifted frequencies).
- Color flow imaging continues to provide the duplex information described above and in addition evaluates the Doppler flow information for its *phase* (direction toward or away from the transducer, on which basis color is assigned) and its *frequency content* (which determines the hue or shade of the assigned color).
- More in-depth coverage of this topic should be gleaned from textbooks focusing on physical principles of ultrasound scanning.

# **TECHNIQUE**

- The patient is supine with a small pillow under the head.
- The extremity is positioned close to the examiner.
- The arm to be evaluated is externally rotated and positioned at approximately a 45-degree angle from the body in what has been called the "pledge position."

For the patient with a dialysis graft, auscultate for a bruit and/or palpate for a "thrill" (vibration), bearing in mind that both graft stenosis and the high volume of blood flow through a patent dialysis graft will produce this effect.

- A multifrequency (e.g., 5–7 MHz) linear array transducer is used.
- The neck vessels are identified, with attention given to the innominate artery on the right. The left common carotid artery arises from the aortic arch.
- Duplex scanning (with or without color flow imaging) is performed at the following anatomic sites in this order:
  - 1. Subclavian artery
  - 2. Axillary artery



**Figure 12-1.** Spectral Doppler of the brachial artery, accompanied by color flow. Courtesy of Steve Bernhardt, BS, RDMS, RVT, RPhS.

- 3. Brachial artery
- 4. Radial artery
- 5. Ulnar artery
- 6. Palmar arch if necessary

Gray-scale imaging is used to observe vessel walls and to identify the presence of plaque and its morphology. Spectral analysis of these vessels, including evaluation of waveforms and measurement of peak systolic velocities, is used to assess blood flow. (See Figure 12-1.)

Color flow imaging is useful to discriminate flow deviations, flow channel narrowing within the vessel, and absence of flow where it should be, as well as assessing whether there is flow where there should not be (e.g., pseudo- or true aneurysm or inflamed lymph node).

- It is rather uncommon for arteries in the upper extremities to become stenotic. The main use for duplex and color flow imaging in the upper extremity is to evaluate hemodialysis access grafts, although other applications exist as well, including vein mapping and evaluation for thrombosis.
- Hemodialysis grafts are evaluated in the following order:
  - 1. Identify and evaluate inflow artery.
  - 2. Identify and evaluate arterial anastomosis.
  - 3. Identify and evaluate body of the graft.
  - 4. Observe for abnormalities, e.g., aneurysm, puncture site leaks, perigraft fluid collections.
  - If color flow imaging is available, observe the image for frequency increases, turbulence, flow channel changes, and other deviations from normal.
  - 6. Identify and evaluate venous anastomoses.
  - 7. Identify and evaluate outflow vein.



- Types of dialysis access include Brescia-Cimino fistulas (i.e., radial artery and cephalic vein) (Figure 12-2), straight synthetic grafts (see Figure 12-3), and looped synthetic grafts.
- Longitudinal and transverse approaches are used to evaluate the gray-scale image for thrombus, wall thickening, and other changes.
- Doppler peak systolic velocities (PSVs) are measured at the aforementioned sites (depending on which study is being performed) and also at other locations of interest (e.g., for stenosis, preocclusion).
- Documentation consists of storing the digital images (e.g., via the Picture Archiving and Communication System [PACS]) or making hard-copy prints.

# **INTERPRETATION**

# **Stenosis**

- Currently, there are no criteria for classifying upper extremity disease as there are for the lower extremities.
- Peak systolic velocities vary widely with changes in skin temperature. The Doppler signal quality is usually triphasic (Figure 12-4A), but when the hand is quite warm or the arm has been exercised, the Doppler signal quality becomes lower-resistance (continuous through diastole). (See Figure 12-4B.) The effects of cooling were previously discussed in Chapter 10, page 111.

**Figure 14-2.** In this transverse image of an abdominal aortic aneurysm, at least 7 cm in diameter, the A/P measured distance is depicted with black X's, while the lateral (also referred to as transverse) measured distance is depicted with white X's. It contains a mixture of echoes that are more heterogeneous than homogeneous. Peak systolic velocities vary according to the extent of thrombus. Large lumen (less thrombus) = lower velocities; small lumen (more thrombus) = higher velocities; preocclusive = lower velocities.



- The examiner should note the type of aneurysm—true (i.e., fusiform or saccular), false, or dissecting—and the presence of thrombus, if any. See the example of a thrombotic aneurysm in Figure 14-2.
- The most frequent complication and danger of an AAA is rupture, but it can also embolize. Both thrombosis and embolization are considered primary complications of peripheral arterial aneurysms. It is not uncommon for both abdominal and peripheral aneurysms to contain varying amounts of thrombus.

### **Renal Artery and Kidney—Technique**

See "General Remarks" on page 140. It is helpful to know that many of the patients undergoing this study present with hypertension (controlled or not well controlled) and that many of these hypertensive patients have *renovascular* hypertension. Renovascular hypertension, a secondary form of high blood pressure, is often caused by renal artery stenosis (which can be secondary to atherosclerosis or fibromuscular dysplasia) or occlusion. The narrowing of the artery reduces blood flow to the kidney and, in response, the kidney produces the enzyme renin. Through a series of complicated interactions, there is a conversion of angiotensinogen to angiotensin II, which results in constriction of the blood vessels. Through another process, sodium and fluid retention also occur.

- The examiner begins in the longitudinal plane, acquiring velocity data from the celiac and superior mesenteric arteries.
- The peak systolic velocity of the aorta is obtained just distal to the superior mesenteric artery.
- Using a transverse approach, the examiner locates the renal arteries. The left renal vein is a good landmark for identifying the left renal artery, as depicted in Figure 14-3. See also Figures 14-4, 14-5, and 14-6, which show not only the left renal vein but also the renal artery branches.
- Kidney size and morphology are evaluated bilaterally. Figure 14-7 shows a normal-sized kidney with color flow clearly demonstrating the segmental arteries, as well as the renal artery.



**Figure 14-3.** Transverse view of the abdomen showing the position of the left renal vein in relation to the left renal artery.



**Figure 14-4.** Transverse view of the abdominal aorta also showing the superior mesenteric artery (SMA), left renal vein, and splenic vein. This image is a good reminder to watch the color bars for flow direction. The veins, in this example, are not blue.



**Figure 14-5.** Transverse image of the abdominal aorta, showing two left renal arteries and a single right renal artery.



**Figure 14-6.** Transverse image of the abdominal aorta (Ao), showing the following vessels with an accompanying arrow: right (RRA) and left (LRA) renal arteries and left renal vein (LRV), which can be insonated with an appropriate Doppler angle. Note also the inferior vena cava (IVC), to the left of the aorta and the left renal vein. Courtesy of UC Davis Medical Center.



**Figure 14-7.** Normal-sized kidney with color flow clearly showing the segmental arteries, as well as the right renal artery (RRA) depicted from its origin at the aorta (Ao) into the kidney. The X's shown on the ends of the kidney depict the start and end points for length measurement. Courtesy of UC Davis Medical Center.



The results are interpreted as follows:

- Normal: ≥0.2
- Abnormal: <0.2</li>

An EDR or PRR of <0.2 indicates an increase in resistance within the kidney parenchyma. An example of the type of waveform obtained in this situation appears in Figure 14-14.\*

 Another ratio used (to a lesser degree) to determine whether resistance is increasing in the kidney (especially the transplanted kidney) is Pourcelot's ratio/resistivity index (RI), which is calculated as follows:

$$\frac{\text{PSV} - \text{EDV}}{\text{PSV}}$$

- Normal: <0.7</li>
- Abnormal (increased resistance): ≥0.7\*\*

\*Neumyer MM: Native renal artery and kidney parenchymal blood flow duplex evaluation with/without color flow imaging. In Rumwell CB, McPharlin MM, Strandness DE, et al (eds): *Vascular Laboratory Policies & Procedures Manual*. Pasadena, Davies Publishing, 2008.

\*\*Rifkin MD, Needleman L, Pasto ME, et al: Evaluation of renal transplant rejection by duplex Doppler examination: value of the resistive index. Am J Roentgenol 148:759–762, 1987.

Figure 14-13. Longitudinal view of a normal kidney. The X's located at the ends of the kidney show how the kidney length is measured.

**Figure 14-14.** Transverse view of the right kidney, using a coronal approach. Doppler signals are obtained from a segmental artery within the kidney. Normally, the signals are low-resistance, as noted on the image with the first set of waveforms. In an abnormal (e.g., nephrosclerotic) kidney, the Doppler signal is high-resistance, as seen to the right. Example of velocity findings in a kidney with abnormal arterial flow patterns:

- Doppler findings: PSV = 70, EDV = 10
- End-diastolic ratio (EDR): 10/70 = 0.14
- *Resistivity index (RI):*  $\frac{70 10}{70} = 0.86$
- The renal resistive-index value (RRIV)\* also uses the peak systolic velocity and the end-diastolic velocity obtained from the kidneys' segmental arteries. The purpose of using this value is to help prospectively identify patients whose renal function or blood pressure will improve after correction of the renal artery stenosis. A lower-resistance RRIV value of <0.75 is associated with improvement in both blood pressure and renal function after the correction of renal artery stenosis. Conversely, an RRIV of ≥0.75 is a strong predictor of worsening renal function and lack of blood pressure improvement, despite the correction of renal artery stenosis. Calculation of the RRIV is as follows: 1 – EDV/PSV. When you compare this to the EDR or PRR, described above, note that the sum of the PRR and the RRIV is 1; that is, subtract the PRR from 1 and you will get the RRIV, and vice versa.
- Proximal high-grade stenosis or occlusion of the renal artery may result in a dampened (prolonged-upstroke) but still low-resistance waveform (see Figure 14-15A). The angiographic result (see Figure 14-15B) is an example of what can cause the poor flow depicted in Figure 14-15A.
- Some references describe the usefulness of evaluating the kidney arterial flow by determining the acceleration time (AT) and acceleration index (AI).
  - The AT is the time interval from the onset of systole to the initial peak and is reported in milliseconds (msec). A proximal stenosis of ≥60% diameter reduction is most likely to produce an AT of ≥100 msec.
  - The AI describes the slope of the Doppler velocity waveform. It is calculated as the change in velocity between the onset of systole and the





\*Yuksel UC, Anabtawi AG, Cam A, et al: Predictive value of renal resistive index in percutaneous renal interventions for atherosclerotic renal artery stenosis. J Invasive Cardiol 24:504–509, 2012.

Figure 14-15. A Spectral Doppler arterial signal obtained from the kidney in the presence of a proximal high-grade stenosis or occlusion of the renal artery. This waveform pattern is termed *tardus parvus*. B Angiogram showing an occlusion (arrow) of the left renal artery. Collateral vessels are providing poorquality arterial flow to the kidney. Α

systolic peak (cm/sec) divided by the AT. The units of measure for AI are cm/sec<sup>2</sup>. A positive result (consistent with a  $\geq$ 60% diameter reduction) is defined as AI  $\leq$ 291 cm/sec<sup>2</sup>.\*, \*\*

# Mesenteric Arteries—Technique

- Patient history: Patients who present with a history of dull, achy, or crampy abdominal pain 15–30 minutes after meals may suffer from *mesenteric ischemia*. Mesenteric ischemic pain is also known as *mesenteric angina*. This condition may be caused by a stenosis or occlusion of vessels such as the superior mesenteric, celiac, or inferior mesenteric arteries.
- Mesenteric ischemia may be acute or chronic. Both conditions are difficult to diagnose. The study described here assists with the diagnostic process, but an arteriogram is essential for diagnosis.
- It is essential that this study always be performed on a fasting patient.
- Peak systolic and end-diastolic velocities are obtained in the longitudinal plane from the following arteries:
  - Celiac trunk (artery), hepatic artery, and splenic artery, all of which often require a transverse approach to the aorta (see Figure 14-16)
  - Proximal, mid, and distal superior mesenteric artery (see Figure 14-17)



**Figure 14-16.** Celiac artery (CA) i.e., celiac trunk—with branches. The hepatic artery (HA) takes blood to the liver; the splenic artery (SA) takes blood to the spleen. All Doppler flow is low-resistance because organs have a constant need for blood supply. Courtesy of Steve Bernhardt, BS, RDMS, RVT, RPhS.



**Figure 14-17.** Color flow Doppler image using a low-frequency transducer in the longitudinal (sagittal) approach demonstrates the celiac artery followed by the superior mesenteric artery (SMA) branching off the aorta. Courtesty of Steve Bernhardt, BS, RDMS, RVT, RPhS.

\*Issacson JA, Zierler RE, Spittell PC, et al: Noninvasive screening for renal artery stenosis: comparison of renal artery and renal hilar duplex scanning. J Vasc Technol 19:105–110, 1995.

\*\*Martin RL, Nanra RS, Wlodarczyk J, et al: Renal hilar Doppler analysis in the detection of renal artery stenosis. J Vasc Technol 15:173–180, 1991.

# Table 14-1Summary criteria for interpreting pre- and postprandial<br/>findings in the superior mesenteric and celiac arteries.

Description	SMA	Celiac Artery
Preprandial (fasting):		
PSV	High	High
EDV	Low	High
Flow reversal	Yes	No
Postprandial (food challenge):		
PSV	Marked increase	No change
EDV	Marked increase	No change
Loss of flow reversal	Yes	N/A
Velocity criteria:		
Normal PSV*	110-177 cm/sec	50–160 cm/sec
Stenosis criteria* (includes poststenotic turbulence)	$PSV \ge 275 \text{ cm/sec}$ predicts $\ge 70\%$ diameter reduction	$PSV \ge 200 \text{ cm/sec}$ predicts $\ge 70\%$ diameter reduction

\*Criteria and other information from various publications by Gregory L. Moneta, MD, Oregon Health & Science University Hospital, Portland, Oregon.

- Celiac band syndrome, which is an extrinsic compression of the celiac artery origin by the median arcuate ligament of the diaphragm, is a fairly frequent anatomic finding, especially among young, athletic women. A reversible celiac artery stenosis occurs and is rarely the cause of clinical symptoms. In such cases the high-velocity signals indicative of stenosis are substantially improved with deep inspiration and return with expiration. See Figure 14-23. An audible bruit can be auscultated, and during duplex/color flow imaging a "color bruit" may appear during exhalation.
- The celiac artery branches into the hepatic and splenic arteries (see Figure 14-16). The liver and spleen have fixed metabolic requirements and are not likely to be influenced by the postprandial state. Therefore, high peak systolic and end-diastolic velocities are normally evident in these vessels at all times.



**Figure 14-23.** Longitudinal image of the abdominal aorta and celiac artery. The Doppler spectral waveforms were recorded at both phases of respiration. Extrinsic compression of the celiac artery during expiration is a frequent finding. In this example, it causes peak systolic velocities and end-diastolic velocities to increase (>284/130 cm/sec) with aliasing noted. During inspiration, velocities decrease to 188/34 cm/sec. Courtesy of Steve Bernhardt, BS, RDMS, RVT, RPhS.



**Figure 14-24.** Inferior mesenteric artery (IMA) easily observed in this image because of a superior mesenteric artery (SMA) occlusion (not seen on this image).

- The hepatic artery and splenic artery Doppler signals should be low-resistance in quality because each provides arterial inflow to a particular organ. There are no specific peak systolic velocity criteria for what suggests stenosis in either of these vessels (other than the stenosis profile described on pages 40–41), but if the celiac artery is occluded, retrograde flow may occur in the hepatic artery and, if so, should be identified during the duplex evaluation.
- Because of its small caliber, it is quite difficult to locate the inferior mesenteric artery (IMA), which branches off the distal aorta. Easy detection and ultrasono-graphic dominance of the inferior mesenteric artery (see Figure 14-24) may suggest occlusion of the superior mesenteric artery.
- Of interest is the ability of the IMA to provide collateralization in response to an occlusion of the SMA. Two possible collateral connections between the SMA and IMA include the arc of Riolan and the marginal artery of the colon (also known as the marginal artery of Drummond).
- Chronic mesenteric ischemia is suggested when there are abnormal findings for at least two of the three mesenteric vessels (celiac, superior mesenteric, and inferior mesenteric arteries) in symptomatic patients.

# **ORGAN TRANSPLANTS**

# *Liver Transplant (Allograft)*

- Liver transplantation is becoming a more frequently utilized procedure to treat patients with end-stage liver disease. In the pre- and postoperative assessment of these patients, the examiner uses a multifrequency (e.g., 3–5 MHz) linear or phased array transducer.
- The preoperative duplex evaluation of candidates for liver transplantation includes documenting the patency of the portal vein, splenic vein, superior mesenteric vein, hepatic veins, inferior vena cava, and hepatic artery while observing for abnormalities (i.e., tumors in the liver and surrounding tissue) and/or other vasculature (see Figure 14-25). Doppler waveforms are evaluated according to the appropriate criteria. It is also important to determine the status of the biliary tree. Abnormalities such as thrombosis of the portal vein can make transplantation difficult, if not impossible.
- Postoperatively, duplex ultrasonography is used to document patency of the portal vein (Figure 14-26), splenic vein, superior mesenteric vein, hepatic veins, inferior vena cava, and hepatic artery, as well as portal vein flow direction and vessel size. Normal portal vein size should be ≤1 up to 1.5 cm.
- Postoperative complications include allograft rejection, pseudoaneurysm, hepatic infarction, and thrombosis of the portal vein, inferior vena cava, and/ or hepatic artery. Obviously, the hepatic artery inflow becomes crucial to the viability of the liver.
  - The location of this artery provides a challenge to obtaining a proper Doppler angle. If the Doppler angle is high (i.e., 80–90 degrees), no flow





**Figure 14-25.** Sagittal/oblique view of the liver revealing ascites and a dilated paraumbilical vein. It is not unusual to observe one or both of these findings in patients being evaluated for liver transplantation. A sudden increase of ascites often precedes portal vein thrombosis, especially in cirrhotic patients. Dilatation of the paraumbilical vein is a specific finding for portal hypertension, in which normally hepatopetal flow becomes hepatofugal as blood flow is diverted to the systemic venous circulation through a collateral pathway.

**Figure 14-26.** Duplex scan of the liver with a patent, normal portal vein. Flow direction (red color) is toward the liver (hepatopetal).

may be detected because of the poor or nonexistent Doppler shift. Before concluding that the hepatic artery is occluded, the examiner must be certain that an appropriate Doppler angle has been used. If possible, the proximal hepatic artery is examined where it branches off the celiac artery to determine the presence or absence of flow.

 Acute rejection will cause liver dysfunction. It has been thought by some that rejection would cause arterial resistance (measured by the resistivity index) to increase. Others, however, suggest that rejection is a cellular process that does not affect the peripheral vessels.

# Renal Transplant (Allograft)

- Both living related-donor and cadaveric kidneys are implanted most commonly in the right iliac fossa.
- Postoperative examination is facilitated if the examiner knows what type of anastomosis was used. The transplanted renal artery vessels are anastomosed either end-to-side with the external iliac artery or end-to-end with the internal iliac artery. The venous anastomosis connects the renal vein to the external iliac vein in an end-to-side fashion. Most often, the donor ureter is implanted into the bladder directly.

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DAVIES

# Introduction to Vascular Ultrasonography



# Pellerito Polak



7th edition

# Introduction to Vascular Ultrasonography

7th Edition

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# Preface

The seventh edition of *Introduction to Vascular Ultrasonography* is a significant update to our previous editions. My co-editor, Joseph F. Polak, and I have pulled together the topics and authors we thought represented the best of vascular ultrasound imaging. Having collaborated together for more than two decades with our vascular course and the sixth edition of this book, we feel that this edition has met our goal to present the definitive text in vascular ultrasound. Jo and I have contributed to 22 of the 35 chapters. Not only have all the chapters in this edition been revised to deliver the newest techniques, protocols, and topics in vascular ultrasound, we also invited several new experts to provide their perspectives and experience in significant areas. For example, the physics section has been completely revamped by Dr. Fred Kremkau, the favored speaker and authority in ultrasound physics. We welcome Heather Gornik, MD, the current president of IAC vascular testing, for her contribution on credentialing, accreditation, and quality in the vascular laboratory. We also are excited to have several world-renowned experts provide us the state of the art in ultrasound contrast applications for vascular imaging. Of course, we brought back many of our favorite authors who received positive reviews from our previous edition.

In addition to multiple new figures and illustrations, a major enhancement to this version of *Introduction to Vascular Ultrasonography* is the integration of Practical Tips sections throughout all the chapters. These Tips are intended to focus the reader on major teaching points and pearls for the successful performance and interpretation of vascular studies.

We are extremely proud to present the seventh edition of *Introduction to Vascular Ultrasonography*. We hope that this book will provide guidance to students, technologists, sonographers, and all practitioners of vascular ultrasound to improve patient diagnosis and management.

John S. Pellerito, MD, FACR, FSRU, FAIUM

# **Table of Contents**

# Section 1–Basics

# 1 The Hemodynamics of Vascular Disease

Abstract

Overview

Introduction

Physiologic Factors Governing Blood Flow and Its Characteristics

Effects of Arterial Obstruction Venous Hemodynamics Summary References

2 Principles and Instruments of Ultrasonography

Introduction Ultrasound Principles Transducers Sonographic Instruments Advanced Features Doppler Principles References

# 3 Doppler Flow Imaging and Spectral Analysis

Introduction Blood Flow Patterns Listening to the Auditory Frequency Spectrum What You Need to Know About Waveform Analysis Diagnosis of Arterial Obstruction Color Flow Ultrasound Imaging Three-Dimensional Vascular Imaging B-Mode Flow Imaging References

# Section 2-Cerebral Vessels

- 4 Anatomy of the Cerebral Structure
  - Abstract Introduction Vascular Anatomy Cerebral Hemodynamics Collateralization Summary References

# 5 Carotid Sonography

# Abstract

B-Mode Ultrasound of the Normal Carotid Artery Wall Normal Blood Flow Characteristics Vessel Identity Examination Protocol The Examination Sequence References

- 6 Evaluating Carotid Plaque and Carotid Intima-Media Thickness
  - Abstract Introduction Pathogenesis of Carotid Atherosclerosis Imaging Protocol: IMT and Plaque Plaque Characterization Plaque Classification Schemes Plaque Surface Features Plaque Neovascularity Perspective on Plaque Characterization Paradigm Shift: Intima-Media Thickness Carotid Plaques: Extent, Severity, and Follow-Up References
- 7 Ultrasound Assessment of Carotid Stenosis

Abstract Introduction Clinical Background Imaging of Stenosis Diagnostic Criteria Plaque Characterization Technical Considerations and Pitfalls The Common Carotid Artery The External Carotid Artery The Innominate Artery Summary References

 8 How to Assess Difficult and Uncommon Carotid Cases
 Introduction
 Carotid Occlusion
 Carotid Artery Dissection Carotid Pseudoaneurysm Carotid Arteriovenous Fistula Fibromuscular Dysplasia Carotid Body Tumor (Glomus Tumor) Difficult Carotid Cases

9 Ultrasound Assessment of the Vertebral Arteries

Abstract Introduction Examination Technique Vertebral Artery Hemodynamics: Qualitative Assessment Vertebral Artery Hemodynamics: Quantitative Assessment Correlative Imaging With Magnetic Resonance Angiography and Computed Tomographic Angiography Treatment of Disease Summary References

10 Ultrasound Assessment of the Intracranial Arteries Introduction Examination Techniques Diagnostic Parameters for Specific Clinical Applications

Summary References

# Section 3–Extremity Arteries

11 Anatomy of the Upper and Lower Extremity Arteries

Abstract Introduction Upper Extremity Lower Extremity References

12 Physiologic Testing of Lower Extremity Arterial Disease

Abstract Introduction Instrumentation Physiologic Testing Procedures Diagnostic Algorithm Clinical Applications Summary References

13 Assessment of Upper Extremity Arterial Disease

Abstract Introduction Basic Anatomy Obstructive Arterial Disease Thoracic Outlet Evaluation Digital Artery Evaluation Raynaud Disease/Phenomenon Color Doppler Imaging and Duplex Ultrasound of Vasculitis Ultrasound Imaging of Upper Extremity Arterial Aneurysms Upper Extremity Arterial Access Radial Artery Harvest Summary References 14 Ultrasound Evaluation Before and After Hemodialysis Access Abstract Introduction

Basic Concepts of Hemodialysis Access Descriptive Terminology Vascular Mapping Before Hemodialysis Access Placement Arteriovenous Fistula Maturity Assessment Graft Evaluation Acknowledgments References

15 Ultrasound Assessment of Lower Extremity Arteries Abstract

Introduction Instrumentation Duplex Ultrasound Technique Classification of Disease Clinical Applications Summary References

16 Ultrasound Assessment During and After Carotid Peripheral Interventions

Introduction Intraprocedural Duplex Ultrasound Assessment Surveillance After Carotid Intervention Surveillance After Peripheral Intervention Infrainguinal Bypass Assessment Synthetic Grafts Duplex-Monitored Peripheral Angioplasty/Stent Placement Post-Procedural Follow-Up Summary References

- 17 Ultrasound in the Assessment and Management of Arterial Emergencies
  - Abstract Introduction Ruptured Abdominal Aortic Aneurysm Carotid Artery Stenosis/Thrombosis Carotid Artery Dissection Acute Lower Extremity Ischemia Femoral Pseudoaneurysm Traumatic Arteriovenous Fistula Penetrating Arterial Fistula Summary References

# Section 4–Extremity Veins

18 Extremity Venous Anatomy and Technique for Ultrasound Examination

Introduction Anatomy of the Lower Extremity Venous Duplex Imaging Examination Technique and Protocol The Venous Anatomy and Examination Protocol Anatomy of the Upper Extremity Upper Extremity Examination Protocol Characterization of Thrombus Summary References

# 19 Ultrasound Diagnosis of Lower Extremity Venous Thrombosis

Abstract Introduction Prevalence, Etiology, and Risk Factors Venous Anatomy of the Lower Extremities Technique Ultrasound Findings Pitfalls May-Thurner Syndrome Alternative Diagnoses/Incidental Findings Summary References

# 20 Risk Factors and the Role of Ultrasound in the Management of Extremity Venous Disease

Abstract Introduction

Acute Deep Vein Thrombosis Etiology and Risk Factors Anticoagulation and Thrombolysis in the Management of

Venous Thrombosis Acute Deep Vein Thrombosis of Specific Extremity Veins

Lower Extremity Duplex Scanning in Suspected Pulmonary Embolism

Lower Extremity Superficial Thrombophlebitis Upper Extremity Venous Thrombosis Sequelae of Deep Vein Thrombosis Primary Varicose Veins Preprocedure Venous Mapping Summary References

21 Ultrasound Diagnosis of Venous Insufficiency

Abstract Introduction Lower Limb Venous Anatomy Pathophysiology of Venous Insufficiency Duplex Diagnosis of Venous Insufficiency Quantitative Measurement of Venous Incompetence The Role of Sonography in the Treatment of Chronic Venous

Insufficiency Summary Acknowledgments References

# 22 Nonvascular Findings Encountered During Venous Sonography

Introduction Soft Tissue Edema Lymphedema Hematoma Muscle Injury Lymph Nodes Popliteal (Baker) Cysts Joint Effusion Infection Soft Tissue Tumors References

# Section 5-Abdomen and Pelvis

- 23 Anatomy and Normal Doppler Signatures of Abdominal Vessels
  Abstract
  Introduction
  Abdominal Aorta
  Celiac Artery
  Splenic Artery
  Splenic Artery
  Superior and Inferior Mesenteric Arteries
  Renal Arteries
  Portal Venous System
  Hepatic Veins
  Inferior Vena Cava
  Renal Veins
  References
- 24 Ultrasound Assessment of the Abdominal Aorta
  - Abstract Introduction Anatomy Normal Sizes Normal Doppler Velocity Profiles Pathologic States Abdominal Aortic Aneurysms Ultrasound Examination Definition of an Aneurysm: Size Thresholds Growth Rate of Aneurysms Common Iliac Artery Aneurysms Aneurysm Complications Varied Pathologies Postsurgical Assessment Summary References
- 25 Ultrasound Assessment Following Endovascular Aortic Aneurysm Repair

Abstract Introduction Overview of EVAR Ultrasound Examination Endoleak Detection Aneurysm Size Endovascular Graft Deformity and Native Artery Complications Contrast-Enhanced Ultrasound Aneurysm Sac Pressure Measurement Surveillance Intervals Summary References

26 Doppler Ultrasound of the Mesenteric Vasculature

Abstract Introduction Anatomy, Physiology, and Natural History of Bowel Ischemia Technique Examination Protocol Diagnostic Criteria Roles of Duplex Ultrasound in Surveillance Following Mesenteric Revascularization (Stent Angioplasty and Bypass Graft Assessment) Keys to a Successful Examination Pitfalls Other Mesenteric Artery Pathologies Summary Acknowledgments References

# 27 Ultrasound Assessment of the Hepatic Vasculature

Abstract Introduction Technique and Normal Hemodynamics Portal Hypertension Portal Vein Thrombosis Portal Vein Stenosis Arterioportal Fistulas Portal Vein Gas Hepatic Vein Obstruction Transjugular Intrahepatic Portosystemic Shunts Congenital Intrahepatic Portosystemic Shunts Hepatic Artery Stenosis/Occlusion Hepatic Artery Pseudoaneurysm References

# 28 Duplex Ultrasound of Native Renal Vasculature

Abstract Introduction Anatomy Principles of Examination Technique Protocol Vascular Disorders Summary Acknowledgments References

# 29 Duplex Ultrasound Evaluation of the Uterus and Ovaries

Abstract Introduction Technical Issues Normal Anatomy and Hemodynamics Current Applications Ectopic Pregnancy Pseudogestation Value of Doppler for Evaluation of Ectopic Pregnancy Pitfalls Summary Acknowledgments References

30 Duplex Ultrasound Evaluation of the Male Genitalia Abstract Introduction The Scrotum The Penis Summary References

31 Evaluation of Organ Transplants

Abstract Introduction Renal Transplantation Hepatic Transplantation References

# Section 6—Trends in Ultrasound Vascular Imaging

32 Credentialing, Accreditation, and Quality in the Vascular Laboratory

Abstract Introduction Credentialing Accreditation Quality Improvement Appropriateness of Vascular Testing Efforts to Standardize Vascular Testing and Improve Quality at the National Level Summary References

- 33 Ultrasound Screening for Vascular Disease
  - Abstract Introduction Definition and Types of Screening Screening for Asymptomatic Carotid Stenosis Abdominal Aneurysm Screening Screening for Cardiovascular Disease: Risk and Subclinical Cardiovascular Disease Summary References

# 34 Correlative Imaging

Abstract Overview Catheter-Based Arteriography Magnetic Resonance Angiography Computed Tomographic Angiography Overview and Correlative Findings Summary References

35 Ultrasound Contrast Agents in Vascular Disease

Abstract Introduction Technical Aspects of Contrast-Enhanced Ultrasound Safety of Ultrasound Contrast Agents Clinical Applications of Ultrasound Contrast Agents Trauma Future Considerations Summary References

# Index

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### Welcome!

Thanks for choosing the Davies CD-ROM Mock Exam for Vascular Technology. This powerful learning resource will make a significant difference in your performance on the registry exam.

### With this CD you can:

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- Test yourself by selecting Test Mode
- · Score yourself by viewing a complete, Results Analysis at the end of your test
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- Earn CME credit
- The ARDMS exam content outline
- System requirements
- Legal notices

# Design My Own Test

Create your own test by selecting as many as three ARDMS exam topics and specifying the number of questions you want in each topic area.

About This CD

Quit

This is an excellent way to focus on topics you need to work on.

### Click below to select one or more exam topics ...

Help

Total number of questions on the topic you selected.



0 Enter the number of questions you want.

Total number of questions on the topic you selected.

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**Create My Test** 

**Clear My Selections** 

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D)	AVIES Start Page Lai	am Mode	Test Mode	Results Anal	ysts	Help	About This CD	Quit	
160	The ultrasound image be	elow shows an	internal carotid	artery with:					4
A	A calcified plaque					-			-
В	An ulcerated lesion					-			
С	A normal arterial wall					4			
D	An intraplaque hemorrha	ige				*			
E	A homogeneous plaque					4			
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160	The ultrasound image below shows an i	nternal carotid a	artery with:	aiyala	neip	About This		
A	A calcified plaque					RIGHT!	_	÷
в	An ulcerated lesion	You chose the correct answer, A: A calcified plaque		: A				
с	A normal arterial wall	Note the brightly echogenic plaque and acoustic shadowing.						
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	eriarge				Enlarge	More Q&A Inform This question belo ARDMS exam topio I. Cerebrovascular exam] To focus only on the exam topics:	ation ngs to the follo c: [25–35% of A nis or other spe	wing RDMS ecific
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24	The great saphenou	s vein:					2.64		
A	Originates along the	medial dorsum of	the foot			4	Tutorial		
в	Passes superiorly, anterior to the medial malleolus						Anatomy of the Lower Extremity Veins		
С	Is accompanied by the saphenous nerve						David S. Sumner, MD Unlike their arterial counterparts, the veir of the leg are divided into two systems: the superficial and the deep (Figures 1 and 2		
D	Receives tributaries from all surfaces of the lower extremity							arts, the veins o systems: the ures 1 and 2).	
E	All are correct					4	[Click "More Images" on the purple navigation bar below to view figures.] The	purple figures.] The	
Next	24 of 665	Prior Bookma	N/Skip Pausa	First	Last	Finish	system are the or	eater (or lo	ng)
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xam Top	bic I-IV. Anatomy, Physiolog	y & Hemodynamics (4%	-18% of ARDMS exa	m)					



#### **RESULTS FOR CURRENT TEST**

Date: 5/12/2014 Time: 2:45:57 PM

Your results appear below in three categories:

- (1) Overall results.
- (2) Results by exam topic.
- (3) Results by individual questions.

This analysis also tells you how many questions (if any) you failed to answer and for which questions you changed your first answer. From this page you can also print your results, review wrong answers in Learn Mode, repeat the same test again, return to the Start Page for another activity, or Quit.

### **OVERALL RESULTS**

Total Questions = 60 Total percentage correct = 18.33% / 11 of 60 Target time / actual time used = 60 minutes / 1.02 minutes Number of unanswered questions = 0

#### SUBJECT AREA RESULTS

I-IV. Anatomy, Physiology & Hemodynamics = 33.33% / 1 of 3 (Unanswered 0)

- I. Cerebrovascular = 15.38% / 2 of 13 (Unanswered 0)
- II. Venous = 15.38% / 2 of 13 (Unanswered 0)

III. Peripheral Arterial = 18.18% / 2 of 11 (Unanswered 0)

**Review Missed Questions** Browse

**Review Bookmarked Questions** 

**Review All Questions** 



- II. Venous = 15.38% / 2 of 13 (Unanswered 0)
- III. Peripheral Arterial = 18.18% / 2 of 11 (Unanswered 0)
- IV. Abdomen and Visceral = 33.33% / 1 of 3 (Unanswered 0)
- V. Miscellaneous Conditions & Tests = 33.33% / 1 of 3 (Unanswered 0)
- VI. Quality Assurance = 0% / 0 of 2 (Unanswered 0)
- VII. Image-Based Questions = 10% / 1 of 10 (Unanswered 0)
- VIII. Physiology & Fluid Dynamics = 50% / 1 of 2 (Unanswered 0)

#### INDIVIDUAL QUESTION RESULTS

For your convenience, the following question numbers correspond to those in the book version of this mock exam. Use the CD and the book together for best results.

Question ID VTR583 = WRONG VII. Image-Based Questions Question ID VTR426 = WRONG III. Peripheral Arterial (20%-30% of ARDMS exam) Question ID VTR570 = WRONG VII. Image-Based Questions Question ID VTR11 = WRONG I-IV. Anatomy, Physiology & Hemodynamics (4%-18% of ARDMS exam) Question ID VTR446 = WRONG IV. Abdomen and Visceral (5%-15% of ARDMS exam) Question ID VTR400 = WRONG III. Peripheral Arterial (20%-30% of ARDMS exam) Question ID VTR403 = WRONG III. Peripheral Arterial (20%-30% of ARDMS exam) Question ID VTR502 = WRONG VI. Quality Assurance (3%-5% of ARDMS exam) Question ID VTR615 = WRONG VII. Image-Based Questions Question ID VTR531 = WRONG VII. Image-Based Questions Question ID VTR537 = WRONG VII. Image-Based Questions Question ID VTR137 = WRONG I. Cerebrovascular (25%-35% of ARDMS exam)

**Review Bookmarked Questions** 

**Review All Questions** 



Exam for Vascular Technology. This powerful learning resource will make a significant difference in your performance on the registry exam.

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### **IMPORTANT NOTE:**

Please double check all of your information to ensure smooth, error free processing of your CME application and quiz.

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# An Interactive Q&A Review for the ARDMS SPI Exam

**CD-ROM Version 3.1** 

# Cindy A. Owen, RT, RVT, RDMS James A. Zagzebski, PhD



150 - - - Browse





Thanks for choosing the Davies CD-ROM Mock Exam for Ultrasound Physics. This powerful learning resource will make a significant difference in your performance on the registry exam.

With this CD you can:

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- Test yourself by selecting Test Mode
- Score yourself by viewing a complete Results Analysis at the end of your test
- Focus on image-based questions by selecting Image Library
- Earn 15 hours of SDMS-approved CME credit

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- Earn CME credit
- The ARDMS exam content outline
- System requirements
- Legal notices

This program automatically randomizes questions on all ARDMS exam topics in the same ratio as on the real exam.

### Select the number of questions you want ...



### Need to focus on specific exam topics?

You can create your own test by selecting as many as three ARDMS exam topics and specifying the number of questions you want in each topic area. To do so, click the purple button below.

This is an excellent way to focus on topics you need to work on.

Click here to design your own custom test.



#### Welcome!

Thanks for choosing the Davies CD-ROM Mock Exam for Ultrasound Physics. This powerful learning resource will make a significant difference in your performance on the registry exam.

### With this CD you can:

- Study and learn by selecting Learn Mode
- Test yourself by selecting Test Mode
- Score yourself by viewing a complete Results Analysis at the end of your test
- Focus on image-based questions by selecting Image Library
- Earn 15 hours of SDMS-approved CME credit

To start immediately, choose any of the activities on the right. To learn more about this powerful learning tool from Davies, click any of the following topics:

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- 1-2-3 Step Ultrasound Education & Test Prep
- Earn CME credit
- The ARDMS exam content outline
- System requirements
- Legal notices

# **Design My Own Test**

Create your own test by selecting as many as three ARDMS exam topics and specifying the number of questions you want in each topic area.

**About This CD** 

Quit

This is an excellent way to focus on topics you need to work on.

### Click below to select one or more exam topics ...

Total number of questions on the topic you selected.



0 Enter the number of questions you want.

Total number of questions on the topic you selected.



Enter the number of questions you want.

Total number of questions on the topic you selected.

0 Enter the number of questions you want.

**Create My Test** 

**Clear My Selections** 

**Return To Start** 



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48	Which of the follo	wing sound interact	tions produced t	he shadows in	dicated by th	ne arrows o	on this image?		
A	Reflection					-	RIGHT!		
						-	You chose the correct an	swer, E:	
В	Diffraction		Refraction						
-	Difference						The edge shadowing seen with curved		
C	Diffusion					-	interfaces is caused by re sound beam. This is the	efraction of the	
D	Rayleigh scatterin	Ig				4	principle that can be obse a pencil in a partially filled	erved by placing d glass of water.	
E	Refraction				201		When viewed from the side appears bent.	de, the pencil	
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b0	1.000	100 m 100					II. Physical Principles (20	% of ARDMS	

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#### **RESULTS FOR CURRENT TEST**

Date: 5/12/2014 Time: 3:40:07 PM

Your results appear below in three categories:

- (1) Overall results.
- (2) Results by exam topic.
- (3) Results by individual questions.

This analysis also tells you how many questions (if any) you failed to answer and for which questions you changed your first answer. From this page you can also print your results, review wrong answers in Learn Mode, repeat the same test again, return to the Start Page for another activity, or Quit.

### **OVERALL RESULTS**

Total Questions = 60 Total percentage correct = 13.33% / 8 of 60 Target time / actual time used = 60 minutes / 1.41 minutes Number of unanswered questions = 0

#### SUBJECT AREA RESULTS

- I. Patient Care, Safety, and Communication = 0% / 0 of 3 (Unanswered 0)
- II. Physical Principles = 16.67% / 2 of 12 (Unanswered 0)
- III. Ultrasound Transducers = 9.09% / 1 of 11 (Unanswered 0)
- IV. Pulse-Echo Instrumentation = 16.67% / 3 of 18 (Unanswered 0)



#### SUBJECT AREA RESULTS

I. Patient Care, Safety, and Communication = 0% / 0 of 3 (Unanswered 0)

II. Physical Principles = 16.67% / 2 of 12 (Unanswered 0)

III. Ultrasound Transducers = 9.09% / 1 of 11 (Unanswered 0)

IV. Pulse-Echo Instrumentation = 16.67% / 3 of 18 (Unanswered 0)

V. Doppler Instrumentation and Hemodynamics = 8.33% / 1 of 12 (Unanswered 0)

VI. Quality Assurance/Quality Control of Equipment = 25% / 1 of 4 (Unanswered 0)

#### INDIVIDUAL QUESTION RESULTS

For your convenience, the following question numbers correspond to those in the book version of this mock exam. Use the CD and the book together for best results.

Question ID SPI211 = WRONG III. Ultrasound Transducers (20% of ARDMS exam) Question ID SPI87 = WRONG II. Physical Principles (20% of ARDMS exam) Question ID SPI113 = WRONG II. Physical Principles (20% of ARDMS exam) Question ID SPI110 = WRONG II. Physical Principles (20% of ARDMS exam) Question ID SPI588 = RIGHT VI. Quality Assurance/Quality Control of Equipment (5% of ARDMS exam) Question ID SPI250 = WRONG III. Ultrasound Transducers (20% of ARDMS exam) Question ID SPI121 = WRONG II. Physical Principles (20% of ARDMS exam) Question ID SPI242 = RIGHT III. Ultrasound Transducers (20% of ARDMS exam) Question ID SPI590 = WRONG VI. Quality Assurance/Quality Control of Equipment (5% of ARDMS exam) Question ID SPI463 = WRONG V. Doppler Instrumentation and Hemodynamics (20% of ARDMS exam) Question ID SPI507 = WRONG V. Doppler Instrumentation and Hemodynamics (20% of ARDMS exam) Question ID SPI407 = WRONG IV. Pulse-Echo Instrumentation (30% of ARDMS exam) Question ID SPI553 = RIGHT V. Doppler Instrumentation and Hemodynamics (20% of ARDMS exam) Question ID SPI539 = WRONG V. Doppler Instrumentation and Hemodynamics (20% of ARDMS exam) Question ID SPI348 = WRONG IV. Pulse-Echo Instrumentation (30% of ARDMS exam)

Review Missed Questions	Review Bookmarked Questions	Review All Questions	Copy Results To Clipboard
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Phone

ARDMS #

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IMPORTANT NOTE:

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ARRT #

Please double check all of your information to ensure smooth, error free processing of your CME application and quiz.

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