

SVM APP Course Handouts Table of Contents Sunday, March 17, 2024

PAD Joint Session

Diagnosis PAD? *Bryan Wells, MD, FSVM*

Preserving Life and Limb in PAD, *Aditya Sharma, MBBS, RPVI, FSVM*

I think we need to revasc? *Andrew Klein, MD, FSVM*

CLTI Controversies - Case CLTI – Surgical Revascularization Approach, *Olamide Alabi, MD, RPVI*

CLTI Controversies - Case CLTI – Endovascular Revascularization Approach, *Yulanka Castro Dominguez, MD, RPVI*

Non-Atherosclerotic Arterial Diseases

Large Vessel Vasculitis, *Alexandra Solomon, MD, RPVI*

Fibromuscular Dysplasia & Related Arteriopathies, *Bryan Wells, MD, FSVM*

Lipid Management: Biomarkers and More, *Merry Ellen Barnett, MD*

Hypertension Management: When basics Aren't Working, *Ali Moran Baird, RN, AGACNP, DNP*

PAD: Non-Pharmacologic Therapies, SET, *Diane Treat-Jacobson, PhD, RN, MSVM*

Imaging & Vascular Cases


Post-thrombotic and Post-Intervention Imaging, *Eri Fukaya, MD, FSVM*

Asymptomatic High Grade Carotid Lesion, *Deborah Hornacek, MD, RPVI, FSVM*

Resistant HTN, *Daniella Kadian-Dodov, MD, FSVM*

Intermittent Claudication Case Post-Revasc, *Danielle Vlazny, PA-C, MS*

AAA Pre and Post EVAR Imaging, *Christine Owen, ACNP*




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Diagnosing PAD

March 2024
Bryan J. Wells, MD

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Learning Objectives

- Discuss differential of PAD and other possible diagnoses of LE pain
- Review classic presentation/ physical exam in PAD
- Role of ABI and other vascular testing in diagnosis

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Case

- 57-year-old man with a history of tobacco abuse presents to the clinic c/o calf pain
- States that after he walks in the parking lot, he gets a burning sensation in the back of his legs
- This is worse with hills and resolves completely after 30 seconds of rest
- Past Medical and Social History
 - Tobacco Abuse
 - Tried to quit with Chantix but failed
 - HLD
 - EtOH
 - HTN
 - Works as a bartender and is on his feet most of the day


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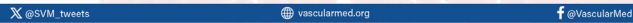
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Case Continued

MEDS	EXAM
<ul style="list-style-type: none">• Aspirin• Atorvastatin• Ibuprofen• Cyclobenzaprine	<ul style="list-style-type: none">• 2+ DP/PT bilaterally• No peripheral edema• No abdominal bruits



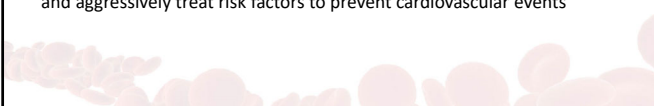
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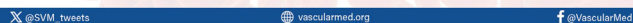
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Why Screen for PAD?

- To identify disease and prevent progression and complications related to PAD
- To identify patients at high risk for other forms of cardiovascular disease and aggressively treat risk factors to prevent cardiovascular events




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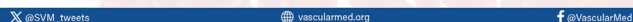
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Question 1

- What is the differential diagnosis for lower extremity pain?



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Lower Extremity Pain

- History and physical is important to determine the etiology
- Differential diagnosis is broad
 - Acute versus Chronic
 - Resting versus exertional
 - Positional, location, associated signs/symptoms
- 3 main categories
 - Vascular
 - Neurological
 - Musculoskeletal

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VASCULAR CAUSES

DVT	Unilateral pain, swelling, discoloration; risk factors include smoking, recent travel, cancer, etc
Peripheral vascular disease	Bilateral claudication; history of smoking, atherosclerosis
Arterial endofibrosis	Associated with repetitive hip flexion, unilateral, most common in cyclists; ischemic pain and loss of power
Cystic adventitial disease	Men in mid-40s, intermittent claudication with activity
Popliteal artery aneurysm	Most common peripheral aneurysm, more common in males, associated with smoking and HTN; acute or chronic ischemic pain or arterial insufficiency, often asx
Popliteal artery entrapment syndrome	Lower limb pain and ischemia with high-intensity exercise associated with excessive dorsiflexion and plantar flexion of ankle

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NEUROLOGICAL CAUSES

Lumbar radiculopathy	Possible association w/ low back pain, sx in distribution of dermatome, often unilateral
Peripheral neuropathy	History of DM, vitamin deficiency, or other systemic disease; pain and sensory loss
Spinal stenosis	Age >/- 50, lower back pain, symptoms worse with activity, relieved with sitting or flexing spine; numbness and tingling from buttocks into legs
Nerve entrapment	Pain and tingling worse with activity in distribution of affected nerve; trauma more likely than overuse

MUSCULOSKELETAL CAUSES

CECS	Consistent bilateral symptoms; numbness and weakness may occur
MTSS	Common in runners, often bilateral, diffuse tenderness
Muscle strain	Immediate onset of symptoms, unilateral, no pain at rest
Stress fracture	Common in athletes, especially female athlete triad; unilateral, focal pain, tenderness
Tendinopathy	Gradual onset of symptoms related to overuse

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True or False

- The majority of patients with peripheral arterial disease will have typical claudication symptoms

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PAD Symptoms

- Classic claudication symptoms
 - Exertional leg pain that resolves within 10 minutes of rest
 - Present in only 32% of patients; 20% have no leg pain, 48% have atypical symptoms
- Rest pain
- Ulcer
- Gangrene
- Leriche's syndrome



McDermott, et al. JAMA. 2001; 286:1599.

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	Claudication	Pseudoclaudication
Characteristic of discomfort	Cramping, tightness, aching, fatigue	Same as claudication + tingling, burning, numbness
Location of discomfort	Buttock, hip, thigh, calf, foot	Same as claudication
Exercise-induced	Yes	Variable
Distance	Consistent	Variable
Occurs with standing	No	Yes
Action for relief	Stand	Sit, change position
Time to relief	<5 minutes	≤30 minutes

Hirsch AT, et al. J Am Coll Cardiol. 2006;47:1239-1312.

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Lower Extremity Peripheral Arterial Disease (PAD)

Iliac artery
Femoral artery
Popliteal artery
Tibial artery

Narrowed artery
Plaque
Artery

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Question 2

- What are the risk factors for peripheral arterial disease?

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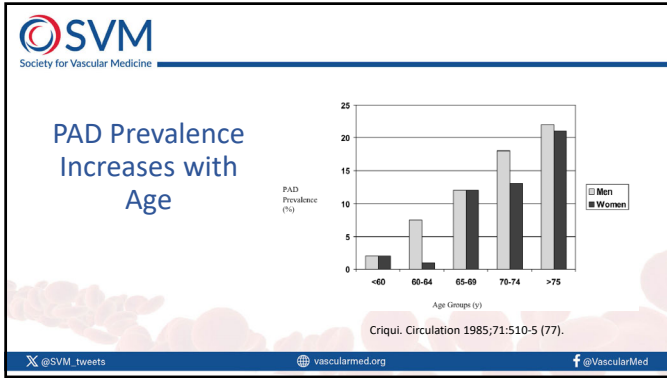
Risk Factors for PAD

Risk Factor	Relative Risk (approx.)
Smoking	3.5
Diabetes	3.0
Hypertension	2.0
Hypercholesterolemia	1.8
Hyperhomocysteinemia	1.5
C-Reactive Protein	1.2

Dormandy JA, et al. J Vasc Surgery, 2000, 31.

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Question 3


- What are the physical examination findings associated with PAD?

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PAD Physical Exam Findings

- Peripheral pulses
- Skin temperature
- Skin color
- Bruits
- Hair loss
- Tissue loss
- Signs of infection



Khan NA, JAMA 2006; 295:536.

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Question 4

- What diagnostic testing is available to evaluate for PAD?

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AHA/ACC 2016 Guidelines

COR	LOE	Recommendations
I	B-NR	In patients with history or physical examination findings suggestive of PAD (Table 4), the resting ABI, with or without segmental pressures and waveforms, is recommended to establish the diagnosis (69-65).
I	C-LD	Resting ABI results should be reported as abnormal (ABI ≤0.90), borderline (ABI 0.91-0.99), normal (1.00-1.40), or noncompressible (ABI >1.40) (46, 63-66).
IIa	B-NR	In patients at increased risk of PAD (Table 3) but without history or physical examination findings suggestive of PAD (Table 4), measurement of the resting ABI is reasonable (41, 42, 67-89).
III: No Benefit	B-NR	In patients not at increased risk of PAD (Table 3) and without history or physical examination findings suggestive of PAD (Table 4), the ABI is not recommended (87, 90).

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Ankle-Brachial Index

$$ABI = \frac{\text{Ankle systolic pressure}}{\text{Brachial systolic pressure}}$$

- ABI has excellent accuracy in detecting stenosis >50%

Right ASBP: Higher right ankle pressures
Right ASBP: Higher arm pressures
Left ASBP: Higher left ankle pressures
Left ASBP: Higher arm pressures

Right brachial systolic pressure (R_{BP})
Left brachial systolic pressure (L_{BP})
Right ankle systolic pressure (R_{ASBP})
Left ankle systolic pressure (L_{ASBP})

Interpretation of ABI:
 >1.30: Noncompressible
 1.00-1.29: Normal
 0.91-0.99: Borderline (borderline) (borderline)
 0.41-0.90: Abnormal (borderline) (borderline)
 0.00-0.40: Severe peripheral arterial disease (borderline)

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Diagnostic Performance of the ABI

TEST	SENSITIVITY	SPECIFICITY
ABI	95-97%	99-100%
PULSE EXAM (DP)	50%	73%
PULSE EXAM (PT)	71%	91%
Rose Claudication Questionnaire	20%	96%

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Diagnostic studies to evaluate for PAD

Ankle-Brachial Index Segmental Pressures & Pulse Volume Recordings Spectral Doppler Waveforms

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Our Patient's Resting ABI

RESTING ABI REPORT

BRACHIAL BLOOD PRESSURE (mmHg)

RIGHT: 120/80 LEFT: 120/80

ANKLE BLOOD PRESSURE (mmHg)

RIGHT: 90/60 LEFT: 90/60

ABI: 0.75 0.75

SPR: 0.80 0.80

SPD: 0.80 0.80

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Exercise ABI

	BRACHIAL	TIME	RIGHT			LEFT		
			ANKLE	ABI	TIME	ANKLE	ABI	TIME
RESTING								
IMMEDIATE	163	02:09	59	0.36	00:55	148	0.91	01:30
PERIOD 1			75		02:53			
PERIOD 2								
PERIOD 3								
PERIOD 4								
PERIOD 5								

Total Exercise Time: 05:14
RIGHT CALF symptomatic at: 01:36

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Arterial Duplex

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Arterial Duplex

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Arterial Duplex

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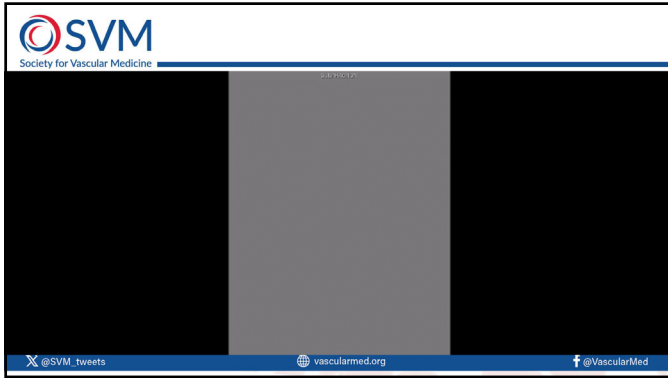
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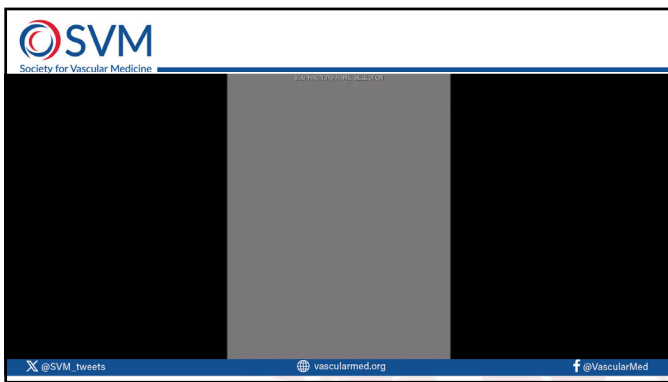
Lower Extremity Angiogram

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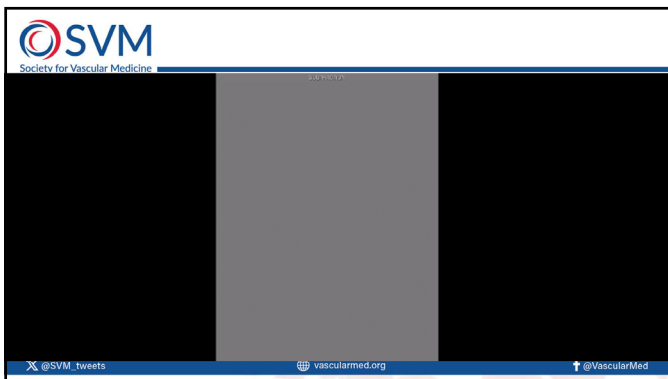
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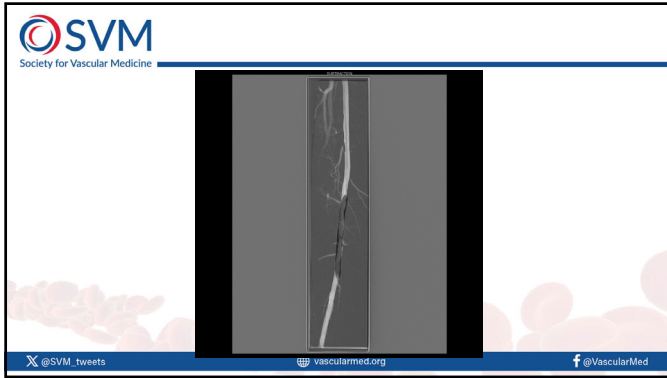
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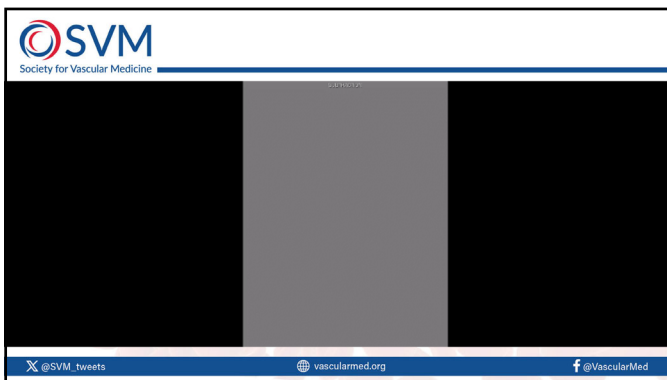
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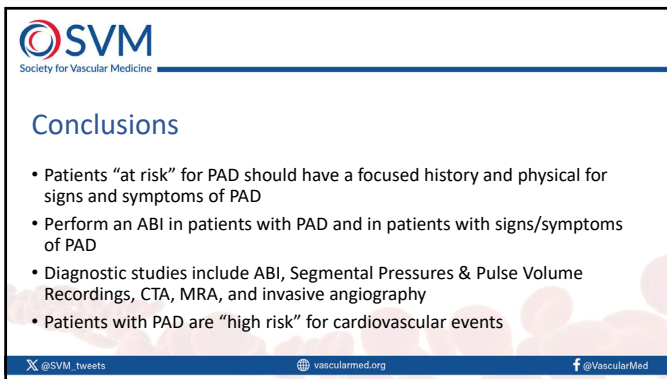
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Preserving Life and Limb in PAD

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SVM Disclosures
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- Research funding:
 - PI: Boston Scientific (HI-PEITHO)
 - PI: Vascular Medcure
 - PI: NIH (C-TRACT)
 - Co-I: Thrombolex (RESCUE) and NIH (ATTRACT)
- Speakers bureau and consulting:
 - Boston Scientific
 - enVveno
- Clinical Excellence Committee:
 - Thrombolex

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SVM PAD: A POLYVASCULAR DISEASE
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> 5000 physician practices in 44 countries
> 65,000 patients 45y+

Proportion of PAD patients with CAD and/or CBVD: 61%

CAD CBVD PAD

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Not ALL PAD is the SAME

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Patients with PAD who have undergone a REVASCLARIZATION:
Very high risk for major adverse cardiovascular events (MACE)
and major adverse limb events (MALE)




Compared to stable PAD
→ 4x higher risk of ALI
→ > 30% increased risk of MI

J Am Coll Cardiol. 2016;67:2719-2728

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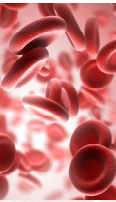
PAD patient

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- Antithrombotics
- Hyperlipidemia
- Diabetes
- Diet and Smoking
- Hypertension
- Exercise therapy

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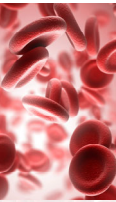
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Antithrombotic Therapy

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- ATT data: Reduced CV event rate by 22% ↓ in large meta-analysis
- CAPRIE trial: Incremental benefit of clopidogrel vs. aspirin (8.7 RR reduction)
- Berger et al: 2009 meta-analysis suggests aspirin not adequately proven to be anti-platelet agent of choice for preventing CV events in PAD patients
- EUCLID trial: ticagrelor monotherapy not superior to clopidogrel
- Path specimens: plaque thrombosis and microembolism
 - 2/3rd of infrapopliteal lesions are thrombosis without atherosclerosis



N Engl J Med 2017; 376:32-40
JACC Cardiovasc Imaging. 2019;12:1501-1513.
BMJ. 2009;339:71.
Lancet. 1996;348:1329
Eur Heart J. 2020;41:1912

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SVM Society for Vascular Medicine **Combination Therapies**

Aspirin+ticagrelor 60 mg twice daily vs aspirin

Pivotal studies: PEGASUS-TIMI 54^{1,26,128}

Population: Prior MI with symptomatic PAD subgroup†

Effects for MACE: 15% relative risk reduction

Effects for MALE: 35% relative risk reduction

Bleeding: 2.32-fold relative excess in TIMI major bleeding

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Enrichment Criteria for Eligibility to COMPASS

- Age < 65 Years Old
- Abdominal Aortic Stenosis >70%
- Diabetes
- Heart Failure
- Chronic Kidney Disease (eGFR <60 mL/min)
- History of Ischemic Stroke
- Peripheral Artery Disease
- Current Smoking Status

of Combined Ischemic Endpoint and Serious Bleeding†

	p Value	OR (95% CI)	p Value
<<0.0001	1.7 (1.3-2.3)	<0.0001	
0.57	1.2 (0.9-1.7)	0.14	
0.007	1.2 (0.9-1.7)	0.16	
<0.0001	1.3 (1.1-1.5)	0.031	
<0.0001	1.2 (1.0-1.6)	0.08	
<0.0001	1.4 (1.1-1.8)	0.003	
<0.0001	1.3 (1.1-1.6)	0.012	
<0.0001	1.4 (1.1-1.8)	0.010	

†including all enrichment criteria. †infarction; OR = odds ratio.

Am Coll Cardiol @VascularMed

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SVM Society for Vascular Medicine **Individualized Approach of Antithrombotic Therapy in PAD**

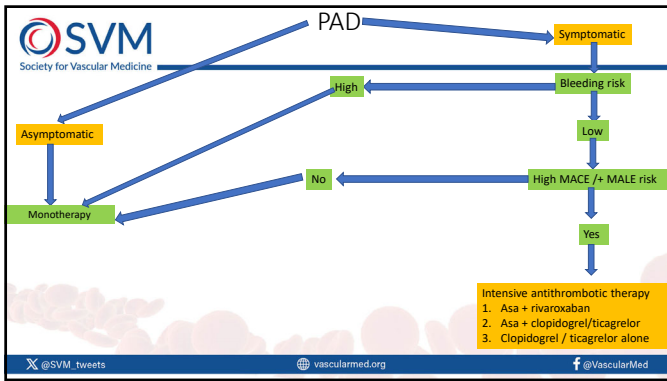
Increase Ischemic Events (MACE)	Increase Limb Events (MALE)	Increase bleeding events
Polyvascular disease (CAD/CVD)	Prior bypass (especially prosthetic or below the knee bypass) or Prior revascularization	Recent major bleeding
Diabetes Mellitus	Prior amputations / tissue loss	Prior Intracranial bleeding
Old age	Below the knee disease/ multilevel disease	Chronic anticoagulation (A fib/VTE)
Active smoking	CLI (ARR higher 5.7% vs. 3.9%)	Anemia
Heart failure / Renal disease	Prior arterial thrombotic events	Fragility / old age

RISK vs BENEFIT

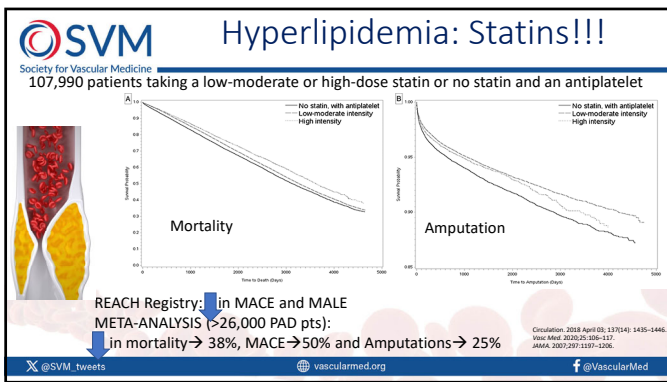
J Am Coll Cardiol. 2018;71(12):2456-67

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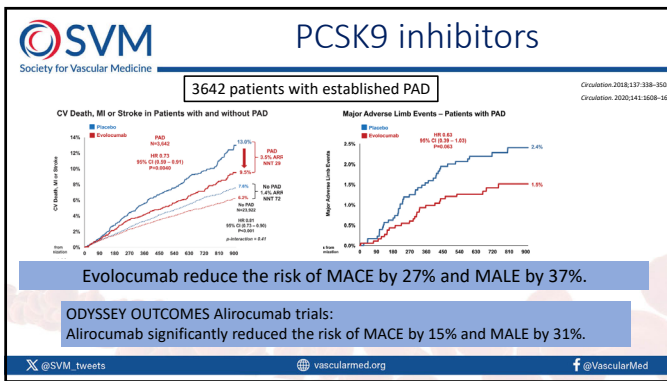
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SVM Approach to Lipid management
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- ALL PATIENTS WITH PAD -HIGH INTENSITY STATINS
 - Atorvastatin 40 to 80 mg or Rosuvastatin 20-40 mg
- PCSK9i considered in some scenarios
 - LDL > 70 or less than 50% reduction from base LDL on high intensity statins
 - Statin intolerance
- It reasonable to have a low LDL even up to 25 mg/dl
- GOAL is to keep < 70 mg/dl

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SVM Diabetes Management: More than LOWERING GLUCOSE !!!
SGLT2 inhibitors and GLP-1 agonists
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- GLP-1 agonists: Liraglutide and Semiglutide
- Liraglutide: LEADER trial (>9000 patients)
 - Lower MACE (HR, 0.87; $p < 0.001$) and CV death (HR, 0.78; $p = 0.007$)
 - Amputation reduction by 35% (HR, 0.65; $p = 0.03$)
- Semiglutide:
 - SUSTAIN-6 trial: (>3000 patients), lowered MACE (HR, 0.74; $p < 0.001$)
 - POST HOC ANALYSIS OF ONLY PAD patients (>1500 patients): MACE is 35% higher in PAD and greater benefits seen compared to non-PAD patients.
 - Taiwan National database: Lower risk of MALE and MACE



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SVM SGLT2 inhibitors
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	Empagliflozin	Canagliflozin	Dapagliflozin
Trial			
CV or renal Outcome			
Limb outcomes			

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Approach to Diabetes

- At risk for amputations (previous amputation, CLI or neuropathy): GLP-1 agonists
- Heart failure and/ renal disease: SGLT2 inhibitors

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Supervised Exercise Therapy and Revascularization for Intermittent Claudication

Network Meta-Analysis of Randomized Controlled Trials

Stenting with supervised exercise therapy alone, supervised exercise therapy alone or stenting alone !!!

Supervised Exercise Therapy (SET)
1,189 patients (25 arms)

Best Medical Therapy (BMT)
688 patients (28 arms)

Angioplasty
511 patients (12 arms)

SET + Angioplasty
395 patients (8 arms)

Compared with SET

Angioplasty + SET	+85 (4 – 170)
Angioplasty	+290 (180 – 390)
SET	+180 (130 – 230)

Mean MWD benefit (95% CrI)

BMT	-180 [(-230) – (-130)]
Angioplasty	-93 [(-170) – (-16)]
Angioplasty + SET	+110 (16 – 200)

JACC Cardiovasc Interv 2023 | @VascularMed

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HOPE: Benefits in CV Risk Subgroups

	No. of Patients	Relative risk in ramipril group	
		Reduced	Increased
History of CAD	7477	Reduced	Increased
No history of CAD	1820	Reduced	Increased
Prior MI	4892	Reduced	Increased
No prior MI	4405	Reduced	Increased
CBV disease	1013	Reduced	Increased
No CBV disease	8284	Reduced	Increased
Peripheral vascular disease	4051	Reduced	Increased
No peripheral vascular disease	5246	Reduced	Increased
Microalbuminuria	1956	Reduced	Increased
No microalbuminuria	7341	Reduced	Increased

N Engl J Med. 2000;342:145-153

- ACEI and ARB are first line anti-hypertensive therapies for hypertension in PAD patients.
- CV event risk reduction seen independent of BP reduction.

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SVM Minimizing Tissue Loss in Patients With PAD
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- Counsel your patients on **daily** self-foot examination
- Prompt diagnosis and treatment of foot infection
- Biannual foot examination by a clinician is reasonable for patients with PAD and diabetes mellitus

Circulation 2017;135:e726

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SVM Take Home Points
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- All patients with PAD require medical therapy and risk factor modification
- Select patients will benefit from intensive antithrombotic therapy
 - These includes those with polyvascular disease, CLI, HF, DM, prior LER

Save Limbs

Save Lives

Hyperlipidemia:

- High intensity statins for ALL
- If intolerant or LDL > 70 mg/dl on statins → PCSK9i


Diabetes:

- Goal HbA1c < 7
- Prior amputations or CLI: GLP1 agonists
- Heart failure or kidney disease: SGLT2i (avoid canaglifazon)

- Exercise therapy, diet and smoking cessation for ALL
- Hypertension: Consider ACEI/ ARB

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CLTI Controversies – Case CLTI –
Surgical Revascularization Approach

Olamide Alabi MD MS
Chief Quality Officer, Division of Vascular Surgery and Endovascular Therapy
Emory University School of Medicine

Sunday May 17, 2024

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Objectives

- Open versus Endo
- “No Stent Zones”
- Angiosome Concept

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Objectives

- ~~Open~~ versus Endo
- “No Stent Zones”
- Angiosome Concept

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3

Objectives

- Open versus Endo Complementary Therapies
- “No Stent Zones”
- Angiosome Concept

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Open Surgical Revascularization: Complementary Therapies

	BEST-CLI	BASIL-2
Trial Endpoint	composite of major adverse limb events or death from any cause	Composite time to above ankle amputation of the trial limb or death from any cause
Study Period	8/2014-10/2019	7/2014-11/2020
Intended N	2100	600
Actual N	1830	345
Differences in inclusion criteria	<ul style="list-style-type: none"> • Life expectancy 2 years • Availability of autogenous bypass conduit 	<ul style="list-style-type: none"> • Life expectancy >6mo • Required an infrapopliteal target
Exclusions	Excessive risk for open vascular surgery	<ul style="list-style-type: none"> • Prior vascular intervention to infrapopliteal target within prior 12mo • Ischemic pain or tissue loss considered not primarily due to PAD

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Open Surgical Revascularization: Complementary Therapies

TABLE 2. Comparison of key patient characteristics comparing BEST-CLI and BASIL-2.

	BEST-CLI COHORT 1	BEST-CLI COHORT 2	BASIL-2
Patients who did not undergo any procedure	4.3% / 1.1%	3.6% / 2.0%	8.7% / 1.2%
Cross-over to other revascularization method	3.5% / 0.4%	1.0% / 2.0%	7% / 3.5%
Median age (years)	66.9 / 67.0	68.4 / 68.8	72.4 / 72.5
Male	72% / 71.1%	71.6% / 72.4%	81% / 82%
Diabetes mellitus	72.1% / 71.6%	62.2% / 58.3%	68% / 69%
Mean eGFR (ml/min)	NA	NA	55.9% / 54%
Chronic hemodialysis	9.4% / 11.8%	12.8% / 10.1%	6% / 3%
Previous stroke	12.8% / 13.9%	19.4% / 12.1%	15% / 10%
Coronary artery disease	42.3% / 44.4%	49.5% / 53.8%	NA
Previous MI	NA	NA	24% / 13%
Previous PCI & CABG	NA	NA	26% / 46%
Previous intervention study leg	5.6% / 3.2%	10.3% / 10.2%	15% / 19%
RBC 4	20.3% / 20%	29.4% / 20.2%	15% / 15%
RBC 5 (and 4)	79.7% / 80%	70.6% / 69.8%	87% / 89% (RBC 5 only)
ASA class 3 or 4	80.8% / 75.9%	83.5% / 80.9%	NA
No. patients completing the trial	560 / 1434 (39%)	226 / 996 (22.7%)	232 / 745 (31.1%)

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Open Surgical Revascularization: Complementary Therapies

Table 1. Comparison of key outcome characteristics comparing BEST-CLI and BASIL-2.

	BEST-CLI Cohort 1	BEST-CLI Cohort 2	BASIL-2
Bypass location			
Femoro-popliteal	40%	47%	2%
Femoro-popliteal AK	NA	NA	0%
Femoro-popliteal BK	NA	NA	0%
Femoro-BTK	15.7%	17.4%	5%
Popliteal-BTK	15.7%	8.4%	40%


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Table 3. Comparison of key outcome characteristics comparing BEST-CLI and BASIL-2.

	BEST-CLI Cohort 1	BEST-CLI Cohort 2	BASIL-2
Endovascular techniques			
Balloon angioplasty	52.7%	47.2%	60%
Atherectomy	13.6%	15.4%	0%
Drug-coated balloon	27.8%	25.1%	0%
Bare-metal stent	39.3%	43.1%	10%
Drug-eluting stent	24.2%	21.5%	0%
Stent graft	8.6%	12.8%	0%
Technical success	98.3% / 84.7%	100%/80.6%	96% / 80%

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Open Surgical Revascularization: Complementary Therapies


Table 3. Comparison of key outcome characteristics comparing BEST-CLI and BASIL-2.

	BEST-CLI Cohort 1	BEST-CLI Cohort 2	BASIL-2
Technical success	98.3% / 84.7%	100%/80.6%	96% / 80%
All cause death	33% / 37.6%	25.9% / 24.1%	53% / 45%
Major amputation (above ankle)	10.4%/14.9%	15.2%/14.1%	20%/18%
AK amputation	NA	NA	NA
BK amputation	NA	NA	NA
Amputation free survival	43.3%/52.4%	41.1%/38.2%	37%/47%
Cross-over intervention during FU	NA	NA	27% / 19%
Reintervention****	9.2% / 33.1%	14.2%/25.6%	5% / 19%

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Open Surgical Revascularization

- BEST-CLI and BASIL-2 are not really comparable
- Neither should be used as an 'end all argument'
- Lessons to learn from both:
 - Nothing beats single segment great saphenous vein....if this is available
 - Get vein mapping prior to the angiogram
 - The toolbox is large – USE IT!
 - Shared decision making
 - There is a small but not insignificant cohort of patients who would benefit from primary amputation

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Open Surgical Revascularization

- ~~Open versus Endo~~ Complementary Therapies
- “No Stent Zones”
- Angiosome Concept

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TH

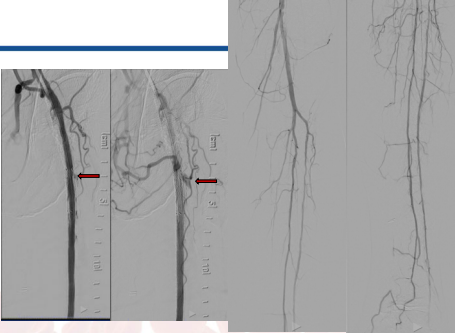
- 3/2012 Left SFA stent
- 8/2012 left SFA atherectomy and BAP of stent
- 11/2012 left SFA laser and BAP of stent
- 7/2013 left SFA revascularization
- 3/2014 BAP Left SFA in-stent stenosis
- 6/2021 shockwave lithotripsy of left CFA and popliteal + DCB BAP
- 6/2023 left SFA stent recanalization
- 8/2023 left SFA stent recanalization

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- 9/18/23 Procedure
- 1) Ultrasound guided access of the right common femoral artery
- 2) Angiogram of abdominal aorta and bilateral iliac arteries
- 3) Angiogram of the left femoral, popliteal, anterior tibial, posterior tibial, and popliteal vessels
- 4) Angiogram of the right femoral, popliteal, anterior tibial, posterior tibial, and popliteal vessels
- 3) PTA of L SFA (stents present)
- 5) Shockwave Lithotripsy
- 6) Laser atherectomy of the L SFA
- 7) Drug coated balloon of the L SFA



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And then this happened...what would you do?

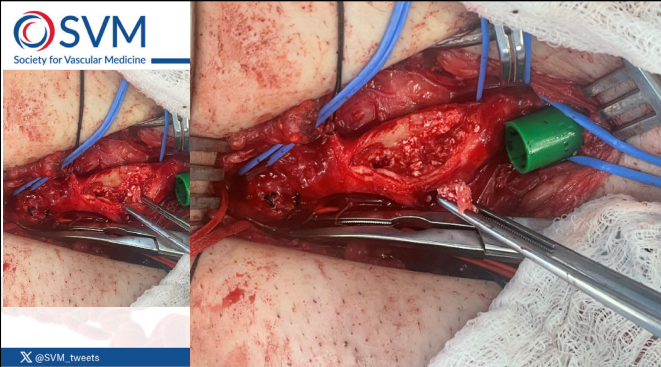


- Abi
- Lt Brachial: 118 mmHg
- Lt Posterior Tibial: 48 mmHg
- Lt Posterior Tibial: 0.41
- Lt Dorsalis Pedis: 50 mmHg
- Lt Dorsalis Pedis: 0.42

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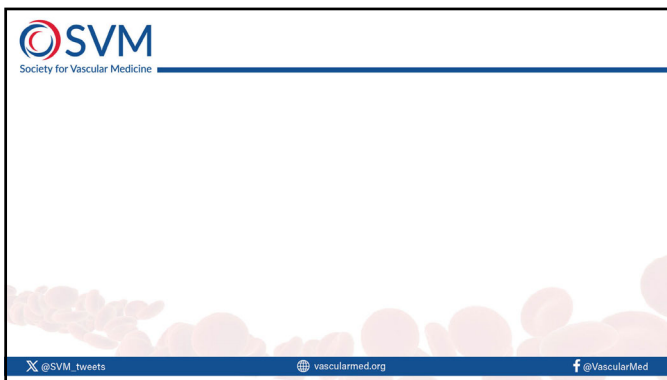


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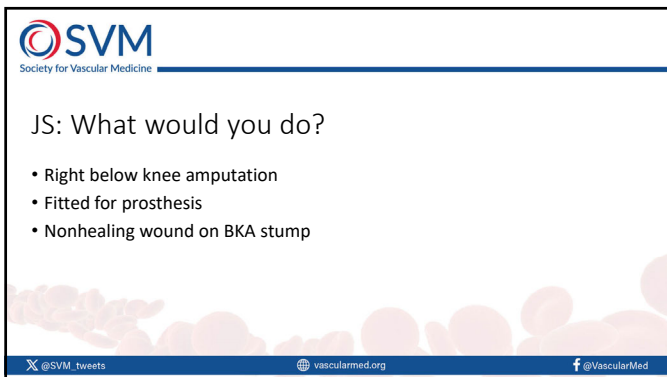
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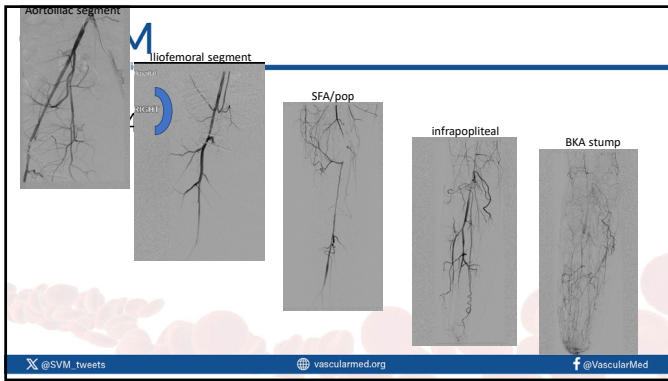
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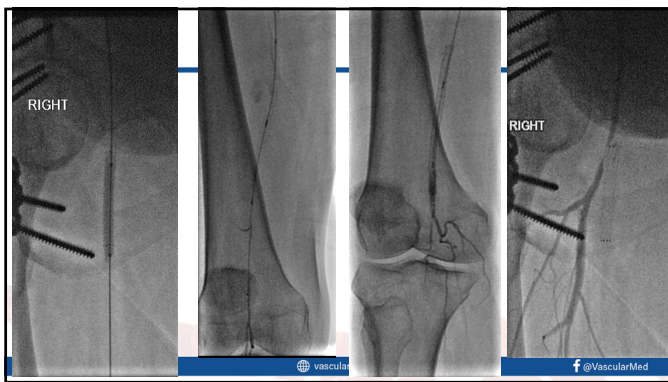
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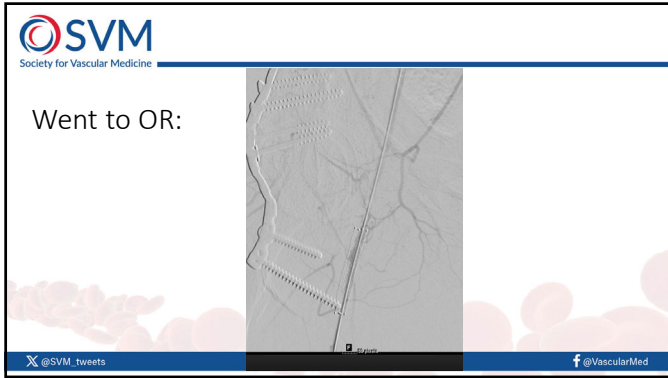
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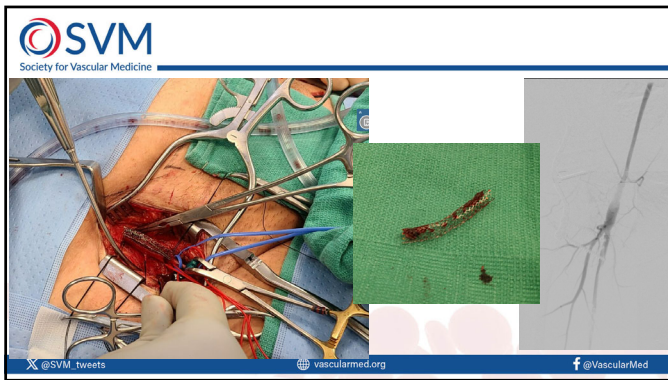
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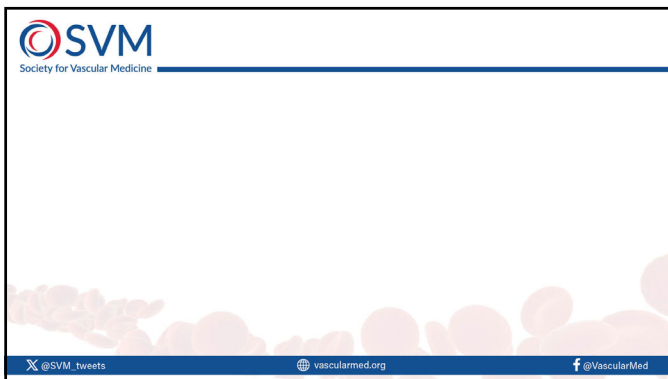
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23



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DH: What would you do?

- 3rd toe wound with underlying osteomyelitis
- No frank cellulitis
- Vein mapping: single segment vein present
- Angiogram

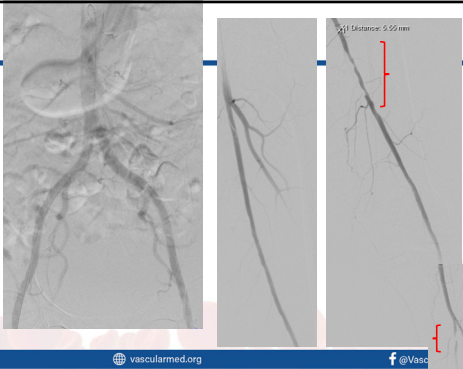
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DH

- Single segment vein present...



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CD: What would you do?


- Chronic emboli from mitral valve vegetation over time
- Developed CLTI with lateral 5th toe wound at base of toe



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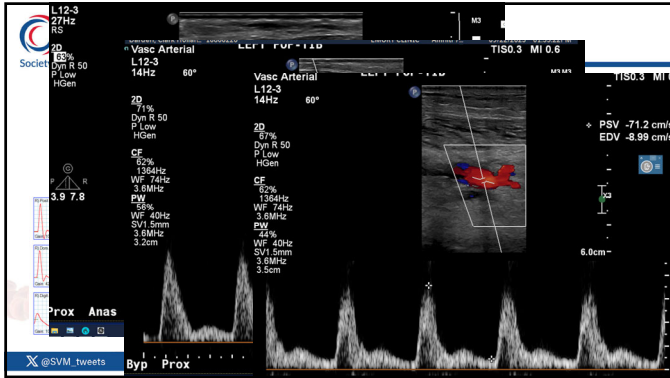
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Open Surgical Revascularization

- ~~Open versus Endo~~ Complementary Therapies
- "No Stent Zones"
- Angiosome Concept

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Angiosomes

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Direct versus Indirect Revascularization

- Meta-analysis, 9 studies
 - 715 direct revascularization
 - 575 indirect revascularization
- Direct revascularization associated with:
 - Lower risk of unhealed wound (HR 0.64, 95% CI 0.52-0.8, I2 0%)
 - Lower risk of major amputation (0.44, 95% CI 0.26-0.75, I2 62%)
- Limb salvage rates
 - 1 year: 86.2% direct; 77.8% indirect
 - 2 years: 84.9% direct; 70.1% indirect

Biancari F, Juvonen T. Angiosome-targeted lower limb revascularization for ischemic foot wounds: systematic review and meta-analysis. *Eur J Vasc Endovasc Surg.* 2014;47(5):517-522.

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Open Surgical Revascularization

- ~~Open versus Endo~~ Complementary Therapies
- “No Stent Zones”
- Angiosome Concept

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Questions


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I think we need to revascularize!

What about medications?
Not just a procedure!!



Andrew J. P. Klein, MD, FACC, FSCAI
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Vascular and Endovascular Medicine
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Atlanta, GA



1

Disclosure

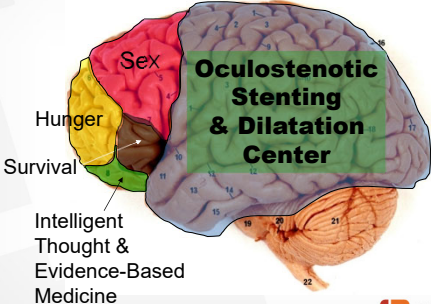

- No Financial Disclosures
- I am
 - An Internist
 - A Cardiologist
 - An Interventional Cardiologist
 - A Vascular and Endovascular Specialist

2


DISCLAIMER

Interventional Cardiology Brain

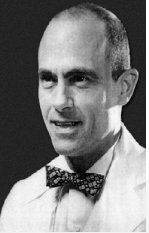




Slide compliments of Jim Hermiller and Chris White

3



“My favorite conceptual trademark is a sketch that I did years ago of a crossed pipe and wrench. It's a gross oversimplification, of course, but what it means to me is that if a plumber can do it to pipes, we can do it to blood vessels”


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Dotter CT, Judkins MP. Transluminal treatment of arteriosclerotic obstruction. Description of a new technic and a preliminary report of its application. *Circulation*. 1964;30:654-70.

4

Objectives

1. PVD care is a medical disease!
2. Revascularization techniques are **complimentary not competitive**
3. “Right procedure for the right patient at the right time”



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5

PAD

Symptom Terminology

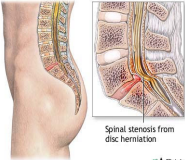
Intermittent Claudication

- Cramping pain
- One or both legs, foot/calf moving up
- Occurs with walking consistent distance
- Does not resolve with continued activity
- Abates with rest (standing) or reduction in walking speed

Pseudoclaudication

- Spinal stenosis
 - Sharp/paresthetic pain, numbness
 - Variable walking distance
 - Relief with sitting or leaning forward
 - Thigh or back moving downward



Spinal stenosis from disc herniation

Miscellaneous

- FMD, Iliac syndrome in cyclist, Buerger's disease, Large vessel vasculitis (Takayasu's, Giant cell arteritis)

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www.sscfund.org/clauidation.html

6

Lower Extremity PAD

Terminology

Critical Limb Threating Ischemia

- Limb pain that occurs at rest or impending limb loss that is caused by severe compromise of blood flow to the affected extremity.
- Chronic versus acute limb ischemia



Hirsch AT, et al. ACC/AHA Guidelines for the Management of Patients with PAD 2005.

7

PAD vs. CAD

Terminology Comparison

CAD		PAD	
STEMI	ACS	CLI	Gangrene Wound
NSTEMI		Rest Pain	
Unstable Angina		Claudication	
Stable Angina		Claudication	

Slide compliments of Ivan Casserly

8

PAD

Risk Factors



- Smoking
- Diabetes
- Age
- Male gender
- Race
- HTN
- Hyperlipidemia
- Hyperhomocystenemia

9

Peripheral Arterial Disease

Prevalence

- PARTNERS Program (PAD Awareness, Risk, and Treatment: New Resources for Survival)
 - 350 Primary care sites
 - Patients (n~7,000)
 - >70 yrs
 - 50-69 yrs with history DM or smoking
 - PVD diagnosis
 - ABI <0.9
 - Previous documentation
 - Abnormal vascular studies
 - Prior revascularization
 - **PAD detected in 1865 patients (29%)**

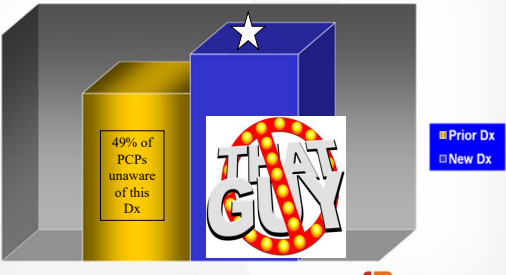




Hirsch AT, JAMA 2001;286:1317-1324

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Peripheral Arterial Disease

Prevalence

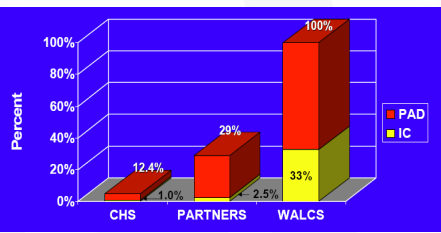



Hirsch AT, JAMA 2001;286:1317-1324


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PAD

Asymptomatic is the rule!



Group	PAD (%)	IC (%)
CHS	12.4%	1.0%
PARTNERS	29%	2.5%
WALCS	100%	33%



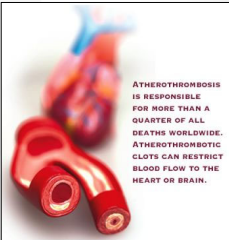
1. Newman A J Clin Epi 2001;54:294-300
2. Hirsch A JAMA 2001;286:1317-1324
3. McDermott M JAMA 2001;286:1599-1606

Graph Courtesy of McDermott, M

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PAD Clinical Impact

- Mortality 2-3x higher than age and sex-matched controls¹
- 5 Year Mortality 30%²
- 15 year Mortality 70%²
- 60-80% of patients with PAD have CAD in at least 1 vessel^{3,4}
- ~25% of PAD patients have a carotid stenosis of >70%⁵



ATHEROTHROMBOSIS IS RESPONSIBLE FOR MORE THAN A QUARTER OF ALL DEATHS WORLDWIDE. ATHEROTHROMBOTIC CLOTS CAN RESTRICT BLOOD FLOW TO THE HEART OR BRAIN.


1. Dermody JJ Cardiovasc Surg (Toron) 1989;36:56-57
2. Watts JT et al. Circulation. 1996;94:3026.
3. Valente RJ, et al. J Vasc Med. 1994;19(4):668.
4. McFalls EO, et al. CMAJ. 2004 Dec 30;171(27):2795.
5. Cheng SW, et al. Cardiovasc Surg. 1999;7(3):183




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PAD Clinical Significance

- Major Risk Factor for Amputation
 - Diabetes
- Quality of Life
 - More severe than that of CHF or recent MI³
 - CLI patients–Terminal Cancer
- Functional impairment and decline is common even in asymptomatic patients^{2,3}



1. Schneider JR et al. Ann Vasc Surg. 1993;7:419.
2. McDermott MM et al. Ann Intern Med. 2002;136:873.
3. McDermott MM, et al. JAMA. 2004; 292:453-461.



14

Ankle-Brachial Index

R brachial
150 mmHg

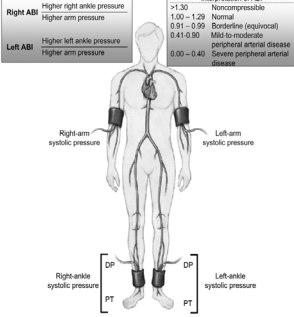
R DP 150
R PT 160

Right ABI
160/150=1.06

L brachial
145


L DP 140
L PT 150

Left ABI
150/150=1.00



Right ABI: Higher right ankle pressure, Higher arm pressure
Left ABI: Higher left ankle pressure, Higher arm pressure

Interpretation of ABI:
>1.30 Noncompressible
1.00 - 1.29 Normal
0.91 - 0.99 Borderline (equivocal)
0.41-0.90 Mild to moderate peripheral arterial disease
0.00 - 0.40 Severe peripheral arterial disease

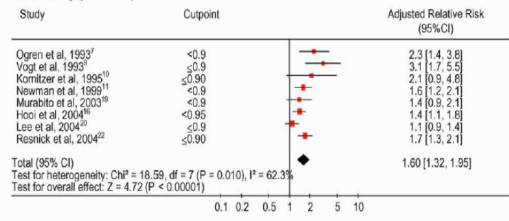


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Abnormal ABI and Mortality



(a) Mortality (all cause)



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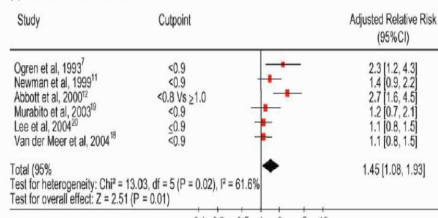
Heald C Atherosclerosis 2006 189:61-69

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Abnormal ABI and CHD



(c): Fatal and non-fatal CHD



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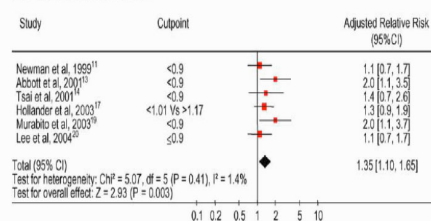
Heald C Atherosclerosis 2006 189:61-69

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Abnormal ABI and CVA



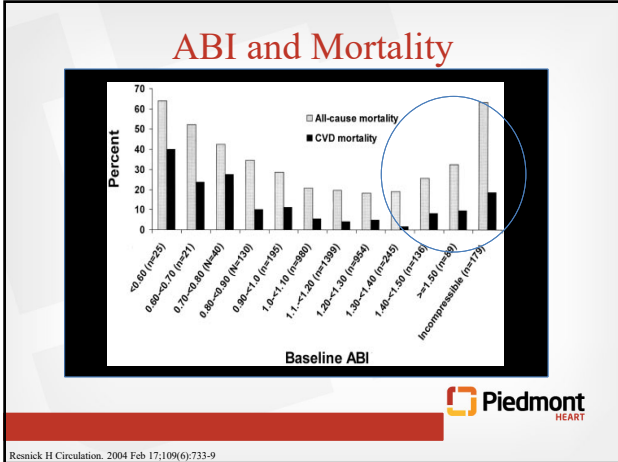
(d) Fatal and non-fatal stroke



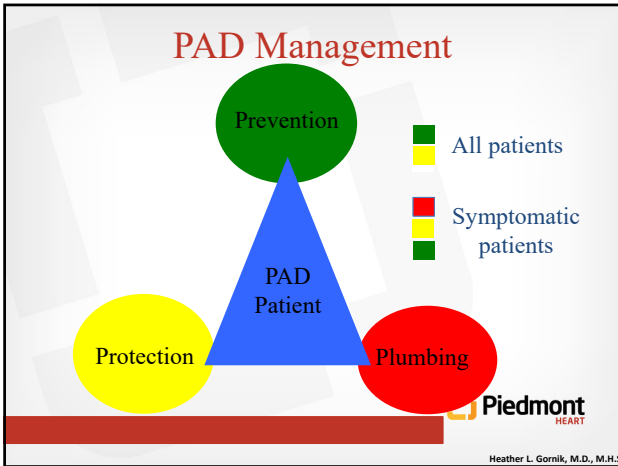
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Heald C Atherosclerosis 2006 189:61-69

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Newer Guidelines

2016 AHA/ACC Guideline on the Management of Patients With Lower Extremity Peripheral Artery Disease

A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines

Developed in Collaboration With the American Association of Cardiovascular and Pulmonary Rehabilitation, Society for Cardiovascular Angiography and Interventions, Society for Clinical Vascular Surgery, Society for Interventional Radiology, Society for Vascular Medicine, Society for Vascular Nursing, Society for Vascular Surgery, Trans-Atlantic Inter-Society Consensus for the Management of Peripheral Arterial Disease, and Vascular and Endovascular Surgery Society

WRITING COMMITTEE MEMBERS*

Marie D. Gerhard-Herman, MD, FACC, FAHA, Chair
Heather L. Gornik, MD, FACC, FAHA, FSVM, Vice Chair*


2016 AHA/ACC Guideline on the Management of Patients with Lower Extremity Peripheral Arterial Disease
Circulation. 2016 Nov 13 And JACC 2016 Nov

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PROTECTION

Foot Care

- Meticulous foot and nail care
- Daily foot self-inspection
- Appropriate footwear
- DM patients
 - Podiatry consultation/collaboration
 - Review warning signs of critical limb ischemia (CLI)
 - Reinforce importance of foot care at each office visit



NEW GUIDELINES

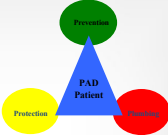

I	C-ID	Patients with PAD and diabetes mellitus should be counseled about self-foot examination and healthy foot behaviors (222, 223).
I	C-ID	In patients with PAD, prompt diagnosis and treatment of foot infection is recommended to avoid amputation (224-228).

2016 AHA/ACC Guideline on the Management of Patients with Lower Extremity Peripheral Arterial Disease

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PREVENTION

- Smoking cessation
- Anti-platelet therapy
- Lipid lowering therapy
 - Statins
- Antihypertensive therapy
- Ace-inhibitors/ARBs
- Glycemic control


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Prevention

Smoking Cessation

I	A	Patients with PAD who smoke cigarettes or use other forms of tobacco should be advised at every visit to quit (170-172).
I	A	Patients with PAD who smoke cigarettes should be assisted in developing a plan for quitting that includes pharmacotherapy (i.e., varenicline, bupropion, and/or nicotine replacement therapy) and/or referral to a smoking cessation program (170, 180-182).
I	B-NR	Patients with PAD should avoid exposure to environmental tobacco smoke at work, home, and public places (185, 186).




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2016 AHA/ACC Guideline on the Management of Patients with Lower Extremity Peripheral Arterial Disease

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Prevention Anti-platelet Therapy



- Meta-analysis of anti-platelet therapy for cardiovascular disease
- 42 clinical trials enrolled patients with PAD
 - Major reductions in Vascular Event rate with Aspirin therapy
 - Benefits similar among PAD subtypes (intermittent claudication, peripheral grafting, and peripheral angioplasty)
- CAPRIE Trial
 - Benefit of clopidogrel over ASA in symptomatic PAD patients with respect to CV risk reduction




Antithrombotic Trialists' Collaboration. BMJ. 2002;324:71.

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Updated Anti-Platelet Statements

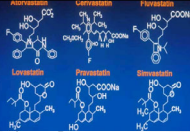
Recommendations for Antiplatelet Agents		
COR	LOE	Recommendations
I	A	Antiplatelet therapy with aspirin alone (range 75–325 mg per day) or clopidogrel alone (75 mg per day) is recommended to reduce MI, stroke, and vascular death in patients with symptomatic PAD (139-142).
IIa	C-EO	In asymptomatic patients with PAD (ABI ≤ 0.90), antiplatelet therapy is reasonable to reduce the risk of MI, stroke, or vascular death.
IIb	B-R	In asymptomatic patients with borderline ABI (0.91 to 0.99), the usefulness of antiplatelet therapy to reduce the risk of MI, stroke, or vascular death is uncertain (75, 76, 139, 142).
IIb	B-R	The effectiveness of dual antiplatelet therapy (DAPT) (aspirin and clopidogrel) to reduce the risk of cardiovascular ischemic events in patients with symptomatic PAD is not well established (143, 144).
IIb	C-LD	DAPT (aspirin and clopidogrel) may be reasonable to reduce the risk of limb-related events in patients with symptomatic PAD following lower extremity revascularization (145-148).
IIb	B-R	The overall clinical benefit of vorapaxar in patients with symptomatic PAD is uncertain (149-152).



2016 AHA/ACC Guideline on the Management of Patients with Lower Extremity Peripheral Arterial Disease

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Prevention Statins




CLASS I

- **Treatment with an HMG coenzyme-A reductase inhibitor (statin) medication is indicated for all patients with peripheral arterial disease to achieve a target LDL cholesterol of less than 100 mg/dl.**

NEW GUIDELINES

COR	LOE	Recommendations
I	A	Treatment with a statin medication is indicated for all patients with PAD (96, 153-157).



Hirsch AT, et al. ACC/AHA Guidelines for the Management of Patients with PAD 2005. 2016 AHA/ACC Guideline on the Management of Patients with Lower Extremity Peripheral Arterial Disease

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Prevention HTN Treatment

Class I
Benefit >>> Risk
Procedure/Treatment SHOULD be performed/administered

Class I

- Antihypertensive therapy should be administered to hypertensive patients with lower extremity PAD
- Beta-adrenergic blocking drugs are effective antihypertensive agents and are not contraindicated in patients with PAD.

NEW GUIDELINES

I	A	Antihypertensive therapy should be administered to patients with hypertension and PAD to reduce the risk of MI, stroke, heart failure, and cardiovascular death (158-162).
---	---	--

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Prevention HTN Treatment

ACE-Inhibitors and ARBS

- HOPE trial: Beneficial in PAD patients
- ONTARGET Trial: Benefit of Telmisartan in PAD
- Also have been shown to increase walking distance versus placebo

NEW GUIDELINES

IIa	A	The use of angiotensin-converting enzyme inhibitors or angiotensin-receptor blockers can be effective to reduce the risk of cardiovascular ischemic events in patients with PAD (161, 168, 169).
-----	---	--

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Ahimastos AA, et al. Ann Intern Med. 2006;144:660 AND JAMA. 2013; Feb 6:309(5):453-60.
2016 AHA/ACC Guideline on the Management of Patients with Lower Extremity Peripheral Arterial Disease

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Prevention Diabetes Therapies

Class I

- Proper foot care, including use of appropriate footwear, chiropody/podiatric medicine, daily foot inspection, skin cleansing, and use of topical moisturizing creams, should be encouraged and skin lesions and ulcerations should be addressed urgently in all diabetic patients with lower extremity PAD. (Level of Evidence: B)

NEW GUIDELINES

I	C-LD	Patients with PAD and diabetes mellitus should be counseled about self-foot examination and healthy foot behaviors (222, 223).
I	C-EO	Management of diabetes mellitus in the patient with PAD should be coordinated between members of the healthcare team.

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Hirsch AT, et al. ACC/AHA Guidelines for the Management of Patients with PAD 2005.
2016 AHA/ACC Guideline on the Management of Patients with Lower Extremity Peripheral Arterial Disease

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Prevention Diabetes Therapies

Class IIa

- Treatment of diabetes in individuals with lower extremity PAD by administration of glucose control therapies to reduce the hemoglobin A1C to less than 7% can be effective to reduce microvascular complications and potentially improve cardiovascular outcomes. *(Level of Evidence: C)*

NEW GUIDELINES

IIa	B-NR	Glycemic control can be beneficial for patients with CLI to reduce limb-related outcomes (191, 192).
-----	------	--

Hirsch AT, et al. ACC/AHA Guidelines for the Management of Patients with PAD 2005.
2016 AHA/ACC Guidelines for the Management of Patients with Lower Extremity Peripheral Arterial Disease.

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Guideline Concordance Saves Limbs

Optimally managed				
N	31	23	17	8
N Major amputation/death	0	2	1	0
% Survival	100	92.7	88.3	88.3
SE	0	0.05	0.06	0.06
Suboptimally managed				
N	67	43	39	23
N Major amputation/death	0	19	4	2
% Survival	100	71.5	64.9	60.4
SE	0	0.05	0.06	0.06

Chung et al. JVS 2013 38:792-80

32

Guideline Concordance Saves Lives

A

Hazard ratio, 0.64; 95% CI, 0.45-0.89; P=0.009

Number at risk

<4 Guideline	502	450	399	355	322	288	256
4 Guideline	237	222	207	180	156	143	123

B

Hazard ratio, 0.55; 95% CI, 0.37-0.83; P=0.005

Number at risk

<4 Guideline	202	306	240	201	175	142	125
4 Guideline	237	155	133	102	94	76	64

ASA Rx 88%; Smoking Abstinence 71%; ACE 60%; STATIN 67%

Armstrong EJ et al. J Am Heart Assoc 2014;3

33

Plumbing Symptomatic Patient

- Medical Therapy
- Exercise Therapy
- Endovascular Therapy
- Surgical Therapy

The Right Time

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Plumbing Symptomatic Patient Medical Treatment

Cilostazol

- Mechanism of Action: Phosphodiesterase inhibitor
- Extensive metabolism issues
 - Diltiazem, Fluoxetine, Omeprazole
 - Consider 50 mg BID starting dose
- Side effects of cilostazol are common; 20% DC Rate
 - Headaches (25-35%)
 - Abnormal stools or diarrhea (~15-20%)
 - Palpitations or tachycardia (~15%)
- **Cilostazol is contraindicated in patients with congestive heart failure of any severity**

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Plumbing Symptomatic Patient Medical Treatment

CLASS IA

- **Cilostazol (100 mg orally 2 times per day) is indicated as an effective therapy to improve symptoms and increase walking distance in patients with lower extremity PAD and IC.**
- **A therapeutic trial of cilostazol should be considered in all patients with lifestyle-limiting claudication (in the absence of heart failure).**

NEW GUIDELINES

COR	LOE	Recommendation
I	A	Cilostazol is an effective therapy to improve symptoms and increase walking distance in patients with claudication (199, 200).

Hirsch AT, et al. ACC/AHA Guidelines for the Management of Patients with PAD 2005.

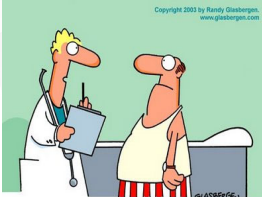
36

Plumbing

Symptomatic Patient

Exercise Treatment

- Improves
 - Exercise performance
 - Walking ability
 - Physical functioning
 - Quality of Life
- Highly cost-effective
- Supervised exercise




Copyright 2003 by Randy Steinberg, www.gaborgen.com

“What fits your busy schedule better, exercising one hour a day or being dead 24 hours a day?”

NEW GUIDELINES

I	A	<p>In patients with claudication, a supervised exercise program is recommended to improve functional status and QoL, and to reduce leg symptoms (36-38, 40-46, 48, 210, 211).</p>
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2016 AHA/ACC Guideline on the Management of Patients with Lower Extremity Peripheral Arterial Disease
Stewart KJ, et al. N Engl J Med. 2007;357:1941.

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Plumbing

Symptomatic Patient

Exercise Treatment

CLEVER

- RCT of aorto-iliac claudicants to Endovascular Therapy (EVT) vs. Optimal Medical Therapy (OMT) vs. Supervised Exercise Therapy (SET)
- At 18 months, the peak-walking time improved for both EVT and SET, but not OMT
- QOL was better for EVT compared with SET or OMT

ERASE

- RCT of aorto-iliac and femoral-popliteal claudicants to EVT + SET vs. SET alone
- EVT + SET had greater improvement in walking distance and health-related QOL vs. SET alone**

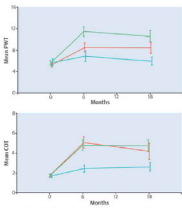



FIGURE 1 Upper panel: Peak Walking Time (PWT); Patients with 18-month follow-up visit only. Lower panel: Claudication Onset Time (COT); Patients with 18-month follow-up visit only (Figure 1 reproduced with permission) [56]



Murphy T et al. J Am Coll Cardiol 2015;65:999-1009.
Fakhry F et al. JAMA 2015;314:1936-1944.

CONCLUSIONS

- SET is an effective alternative to revascularization
- SET + EVT in the presence of GDMT = best option**



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Plumbing

Symptomatic Patient

Exercise Treatment Guidelines

- Warm-up and cool down for 5-10 minutes
- Types of Exercise
 - Treadmill and track walking most effective
 - Resistance training complementary only
- Intensity
 - Initial workload to elicit sx within 3-5 minutes
 - Walk until moderate severity occurs, rest for brief period till sx abate
- Duration
 - Exercise-rest-exercise; 35 minutes and increasing 5 minutes each session until 50 minutes of intermittent walking can be accomplished
- Frequency
 - Treadmill or track 3-5x per week
- Supervision
 - As ability improves, workload should be increased by increasing grade or speed to ensure stimulus of claudication pain always occurs

Stewart KJ, et al. N Engl J Med. 2002;347:1941.

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Plumbing

Symptomatic Patient

Exercise Treatment


Supervised exercise program (COR I, LOE A)

- Takes place in a hospital or outpatient facility.
- Uses intermittent walking exercise as the treatment modality.
- Program can be stand alone or within a cardiac rehabilitation program.
- Program is directly supervised by qualified health-care provider(s)
- Training performed for a minimum of 30-45 min/session; sessions performed at least 3 times/wk for a minimum of 12 wk (36-46).
- Training involves intermittent bouts of walking to moderate-to-maximum claudication alternating with periods of rest.
- Warm-up and cool-down periods precede and follow each session of walking.

Structured community or home-based exercise program (COR IIa, LOE A)

- Takes place in the personal setting of the patient rather than in a clinical setting (41, 47-51).
- Self-directed program with guidance of health-care providers.
- Health-care providers prescribe an exercise prescription similar to that of a supervised program.
- Patient counseling ensures understanding of how to begin and maintain the program and how to progress the difficulty of the walking (by increasing distance or speed).
- Program may incorporate behavioral change techniques such as health coaching and/or use of activity monitors.

COR indicates Class of Recommendation; LOE, Level of Evidence; and PAD, peripheral artery disease.



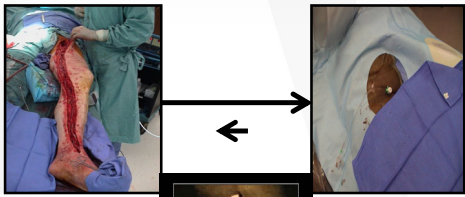

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Plumbing

Symptomatic Patient

Paradigm is Changing

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Plumbing


Symptomatic Patient

- Should you revascularize?
- What is the Indication?
- Where is the disease?
- What are the patient’s co-morbidities?
- What are you trying to achieve?
- What is the disease like?
- What is the renal function?
- What is the realistic long term patency?
- Who is your operator?
- What is your personal threshold?
- What would you do for your family or yourself?

6 CRITICAL THINKING QUESTIONS FOR ANY SITUATION

- What’s happening?
- Why is it important?
- What don’t I see?
- How do I know?
- Who is saying it?
- What else? What if?

BOTTOM LINE



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Plumbing

Symptomatic Patient

EVT Strategy Selection Based on TASC Class

Previously - surgery
Today - "EVT-first" c

TASC A
Endovascular

TASC D
Surgery

Aorto-iliac
Femoral-popliteal

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TASC II Working Group JVS January 2007

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Plumbing

Symptomatic Patient

Surgical therapy: Claudication

8.1.2. Surgical Revascularization for Claudication: Recommendations

COR	LOE	Recommendations
I	A	When surgical revascularization is performed, bypass to the popliteal artery with autogenous vein is recommended in preference to prosthetic graft material (263-271).
IIa	B-NR	Surgical procedures are reasonable as a revascularization option for patients with lifestyle-limiting claudication with inadequate response to GDEM, acceptable perioperative risk, and technical factors suggesting advantages over endovascular procedures (232, 265, 275-277).
III: Harm	B-R	Femoral-tibial artery bypasses with prosthetic graft material should not be used for the treatment of claudication (287-289).
III: Harm	B-NR	Surgical procedures should not be performed in patients with PAD solely to prevent progression to CLI (234-237, 262).

Where are you going to suture?
Is there a suitable vein?
Perioperative risk
Wound infection risk

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Symptomatic Patient

Endovascular therapy: CLAUDICATION

COR	LOE	Recommendations
I	A	Endovascular procedures are effective as a revascularization option for patients with lifestyle limiting claudication and hemodynamically-significant aortoiliac occlusive disease (12, 37, 38, 252, 240, 242, 246).
IIa	B-R	Endovascular procedures are reasonable as a revascularization option for patients with lifestyle limiting claudication and hemodynamically-significant femoropopliteal disease (217, 232, 243-245, 250, 251).
IIb	C-LD	The usefulness of endovascular procedures as a revascularization option for patients with claudication due to isolated infrapopliteal artery disease is unknown (256-258).
III: Harm	B-NR	Endovascular procedures should not be performed in patients with PAD solely to prevent progression to CLI (234-237, 259-261).

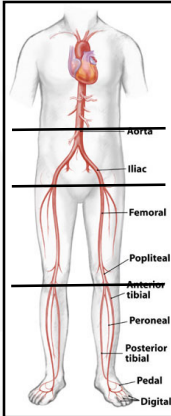
Where is the disease?
Who is your operator?
Periprocedural risk?

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Plumbing
Symptomatic Patient
EVT Treatment

- **Aorto-Iliac**
 - Endovascular approach unless AAA to be also repaired or failure of EVT
- **Femoral-Popliteal**
 - Depends on type of disease (focal vs. diffuse), patient risk factors and comorbidities, claudicant vs. CLI, long term patency, renal function
- **Infra-popliteal**
 - Medical therapy for most, unless CLI
 - DES consideration
 - May change with bioabsorbable scaffolds



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Plumbing
Symptomatic Patient
Aortoiliac Revascularization Indications

Aorto-iliac disease *with symptoms*

- Relieve claudication
- Wound healing in CLI
- Improve functional status and Quality of Life (QOL)

Aorto-iliac disease *without symptoms*

- Situations where large-bore arterial access is required for hemodynamic support devices (e.g., intra-aortic balloon pumps (IABP) or other catheter-based ventricular assist devices), for structural, valvular (e.g., TAVR), and vascular (e.g., EVAR) procedures



Klein A, Jaff M, Gray B et al. Catheter Cardiovasc Interv. 2017;90:E90-E110.

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Plumbing
Symptomatic Patient
EVT Treatment
2005 ACC/AHA Guidelines
Class I

- Endovascular procedures are indicated for individuals with a vocational or lifestyle-limiting disability due to intermittent claudication when clinical features suggest a reasonable likelihood of symptomatic improvement with endovascular intervention and (a) there has been an inadequate response to exercise or pharmacological therapy and/or (b) there is a very favorable risk-benefit ratio (e.g., focal aortoiliac occlusive disease). (Level of Evidence: A)

2016 ACC/AHA GUIDELINES

I A Endovascular procedures are effective as a revascularization option for patients with lifestyle limiting claudication and hemodynamically-significant aortoiliac occlusive disease (12, 37, 38, 232, 240, 242, 246).

Hirsch AT, et al. ACC/AHA Guidelines for the Management of Patients with PAD 2005. 2016 AHA/ACC Guideline on the Management of Patients with Lower Extremity Peripheral Arterial Disease



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Plumbing

Symptomatic Patient EVT in CFA

Registry Data

- Supports EVT-first approach, 5 year f/u data on CFA stenting
 - 79% freedom from TLR

TECCO Trial

- 117 pts RCT of **common femoral endarterectomy vs. EVT** for isolated CFA disease
- 1st outcome: M&M within 30 days
 - 16 of 61 patients (26%) in the CFE group and 7 of 56 patients (12.5%) in the EVT group (odds ratio, 2.5; 95% CI, 0.9 to 6.6; p<0.05).
- The mean duration of hospitalization was significantly lower in the EVT group (3.2±2.9 days vs 6.3±3 days; p<0.0001).
- At 24-months: No difference in the sustained clinical improvement, the primary patency rate, and the target lesion and extremity revascularization rates**

Time (months)	Stenting (%)	Surgery (%)
0	100	100
6	~95	~95
12	~90	~95
18	~85	~95
24	~80	~95

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Goueffe Y et al. JACC Cardiovasc Interv. 2017 Jul 10;10(13):1344-1354
Azema L et al. J Vasc Endovasc Surg 2011;41:787-793.

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Plumbing

Symptomatic Patient

EVT in FP Disease: TASC and ACC/AHA guideline

- TASC 2015 update recommends
 - "endovascular first" recommendation for experienced operators and teams
- 2016 ACC/AHA guidelines on PAD provide a class IIA recommendation (Level of Evidence B) for EVT of FP disease

CONCLUSION

"the choice of EVT as a revascularization approach for claudication due to femoral-popliteal disease should include a **discussion of outcomes, addressing the risk of restenosis and repeat intervention, particularly for lesions with a poor likelihood of long-term durability**"

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Jaff MR et al. Catheter Cardiovasc Intervent Off J Soc Cardiac Angiogr Intervent 2015;86:611-625
Gerhard-Herman MD et al. J Am Coll Cardiol 2016;69:1465-1508

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Plumbing

Symptomatic Patient

EVT in Infrapopliteal Dz

- Generally limited to Critical Limb Ischemia (CLI) patients
- Small vessels, diffuse and long disease, high rates of restenosis
- Intervention to provide straight line flow to the foot, angiosome-based approach
- For claudicants, only moderate to severe (>50% diameter stenosis) lesions and multivessel tibial disease (2 tibial vessels) should be considered for revascularization.
- Prior to considering infra-popliteal intervention, all hemodynamically significant inflow disease should be treated to normalize inflow to the infra-popliteal circulation.

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Plumbing Symptomatic Patient CTLI

9.1. Revascularization for CLI: Recommendations

Recommendation for Revascularization for CLI

COR	LOE	Recommendation
I	B-NR	In patients with CLI, revascularization should be performed when possible to <i>optimize leg care</i> (290a). An evaluation for revascularization options should be performed by an interdisciplinary care team (Table 9) prior to amputation in the patient with CLI.
I	C-EO	

9.1.1. Endovascular Revascularization for CLI: Recommendations

Recommendations for Endovascular Revascularization for CLI

COR	LOE	Recommendation
I	B-R	Endovascular procedures are recommended to establish in-line blood flow to the foot in patients with nonhealing wounds or gangrene (292, 293).

9.1.2. Surgical Revascularization for CLI: Recommendations

Recommendations for Surgical Revascularization for CLI

COR	LOE	Recommendation
I	A	When surgery is performed for CLI, bypass to the popliteal or infrapopliteal arteries (i.e., tibial, pedal) should be constructed with suitable autogenous vein (263, 266, 269, 272).

Guidelines

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- Who is at your center?
 - Culture may drive things but don't let it
- Who is doing the procedure?
 - Experience
 - Back-up
 - Thoughtful
 - Collaborative

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PTA vs. Bypass

Figure 2: Amputation-free survival after bypass surgery and balloon angioplasty
Bars show 95% CIs for survival up to 1, 2, 3, and 4 years of follow-up, which were calculated from the cumulative hazards.

Number at risk	0	1	2	3	4	5
Angioplasty	224	149	100	51	19	2
Surgery	228	148	108	64	23	7

Figure 3: All-cause mortality after bypass surgery and balloon angioplasty
Bars show 95% CIs for survival up to 1, 2, 3, and 4 years of follow-up, which were calculated from the cumulative hazards.

Number at risk	0	1	2	3	4	5
Angioplasty	224	173	116	63	25	6
Surgery	228	169	120	71	26	7

Rates of MI, wound complications, Pulmonary comps increased in Surgery group vs. repeat revasc in PTA group

Basal Trial Lancet 2005; 366: 1925-34

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BEST-CLI

The **NEW ENGLAND**
JOURNAL of MEDICINE

ESTABLISHED IN 1812 DECEMBER 22, 2022 VOL. 387 NO. 25

Surgery or Endovascular Therapy for Chronic Limb-Threatening Ischemia

A. Farber, M.T. Menard, M.S. Conte, J.A. Kaufman, R.J. Powell, N.K. Choudhry, T.H. Hamza, S.F. Assmann,* M.A. Creager, M.J. Czraky, M.D. Dake, M.R. Jaff, D. Reid, F.S. Siami, G. Sopko, C.J. White, M. van Oyer, M.B. Strong, M.F. Villarreal, M. McKean, E. Azene, A. Azarbal, A. Barleben, D.K. Chew, L.C. Clavijo, Y. Douville, L. Findtiss, N. Garg, W. Gasper, K.A. Giles, P.P. Goodney, B.M. Hawkins, C.R. Herman, J.A. Kalish, M.C. Koopmann, I.A. Laskowski, C. Mena-Hurtado, R. Motaganahalli, V.L. Rowe, A. Schanzer, P.A. Schneider, J.J. Siracuse, M. Venermo, and K. Rosenfield, for the BEST-CLI Investigators†



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BEST-CLI

•~30% of patients were dead at end of trial (3 years) no matter what intervention





You want your leg filleted open when you have 1/3 chance of being dead soon?




56

BEST-CLI: Cohort 1

- Comparison of the BEST Surgical intervention (GSV) vs. "Best Endo"
- What is 'best' Endo?
 - DCB/DES
 - Trial started in 2014
 - Trial Data
 - 52% PTA only
 - 15% Atherectomy
 - 25% DCB
 - 22% DES





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BEST-CLI: WHO?

•Endovascular interventions were performed by:

- **Vascular surgeons: 73%**
- Interventional cardiologists: 15%
- Interventional radiologists: 13%



• The **technical success** of the index procedure was **98% in the surgical** group and **85% in the endovascular** group

15% failure rate in ENDO
Is this 'Best Endo'?



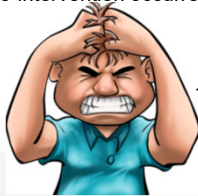
JVS 2016;63:958-965 Large Series report Endovascular Success rates >90%

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BEST-CLI: Cohort 1



- **'BEST' Endo: Success Rate equal to BASIL 1: 17 years ago !!!!!**
- 108 cases of early technical failure in the endovascular group → 66 were treated with a bypass operation within 30 days.
- 42.5% re-intervention occurred within 30 days.



15% failure rate in ENDO
? Shocked that this led to 42.5% reintervention rate



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BEST-CLI

CONCLUSIONS

Among patients with CLTI who had an adequate great saphenous vein for surgical revascularization (cohort 1), the incidence of a major adverse limb event or death was significantly lower in the surgical group than in the endovascular group. Among the patients who lacked an adequate saphenous vein conduit (cohort 2), the outcomes in the two groups were similar. (Funded by the National Heart, Lung, and Blood Institute; BEST-CLI ClinicalTrials.gov number, NCT02060630.)



IF


1. You are at a Center with a HIGH Endo failure rate
2. Endo therapy that is given is not based on the most current evidence




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BEST-CLI: Limitations (Listed)

- White male patients
- **Majority of pts with CLTI do not have SSGSV**
- Trial ran out of funds so follow up limited on cohort 2
- MAJOR DROP IN DCB use because of Paclitaxel debate (which is now been settled) but started in 2014 so.....BS
- 66% infrapopliteal disease but lots of fem-pop bypasses?
- Angiographic analysis pending






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BEST-CLI: Limitations (not listed)




- **Majority of operators were VS**
 - Procedure 1: 98-100% Success rate
 - Procedure 2: 80-85% Success rate
 - Which one were people better at ?
- **No real DCB/DES use**
- Major endpoint driven by re-intervention (not CD-TLR)
- Low enrollment of women and Black patients
- **High burden of CV disease but Medical therapy Awwful:**
 - 65-70% only on ASA and/or statins




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BEST-CLI: Limitations (not listed)

- 75% of sites had some combo of IR, VS, IC
- Only 13% had all 3 and 28% of sites with only VS performing surgery and endo procedures
- *High mortality for endo procedures ??GETA for VS*
- POBA was used most of the time vs DCB/DES/atherectomy
- Enrollment was very very slow
- SUBGROUP ANALYSIS
 - COHORT 1: NO difference in
 - Age>80
 - CKD
 - Black Patients








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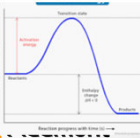
BEST-CLI Take Home

- GSV + Cohort did well with surgery
 - **Low periop Death rate: 1.9%**
- Comparing the **best surgical** procedure to **crappy endo** procedures
 - **80-85% Failure rate**
 - Heterogenous procedure w >50% POBA alone
- QOL showed Endo to be better
- Need better and more meds
- Cannot generalize results
 - What is your center like?
 - Who are your operators
 - What is your threshold?
 - If you have no out to the OR would you work harder at endo? Activation Energy



Best Endovascular versus Best Surgical Therapy in Patients with CLI (BEST-CLI) Trial: A Misleading Trial Name

From Edwin A. Takahashi, MD
Robert A. Lockstein, MD
Sergey Aliev, MD
Division of Vascular and Interventional Radiology (E.A.T., S.M.),
Department of Radiology, Mayo Clinic, 200 1st Street SW,
Rochester, MN 55905, and Division of Interventional Radiology
(R.A.L.), St. Mount Sinai Hospital, New York, NY



HEART

Simons JP, Schanzer A, Flahive JM, et al. Survival prediction in patients with chronic limb-threatening ischemia who undergo infrainguinal revascularization. J Vasc Med Biol. 2019;31(5):1515-1523.

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Which one do you want?



VS.





65

How should we best treat PAD?

Team Approach

TEAM

T TOGETHER

E EVERYONE

A ACHIEVES

M MORE



Who is on your team?



66

Bottom Line: TEAMWORK


Table 9. Interdisciplinary Care Team for PAD


A team of professionals representing different disciplines to assist in the evaluation and management of the patient with PAD. For the care of patients with CLI, the interdisciplinary care team should include individuals who are skilled in endovascular revascularization, surgical revascularization, wound healing therapies and foot surgery, and medical evaluation and care.

Interdisciplinary care team members may include:

- Vascular medical and surgical specialists (i.e., vascular medicine, vascular surgery, interventional radiology, interventional cardiology)
- Nurses
- Orthopedic surgeons and podiatrists
- Endocrinologists
- Internal medicine specialists
- Infectious disease specialists
- Radiology and vascular imaging specialists
- Physical medicine and rehabilitation clinicians
- Orthotics and prosthetics specialists
- Social workers
- Exercise physiologists
- Physical and occupational therapists
- Nutritionists/dieticians

Faster. Stronger. Farther.







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Conclusions

- PAD is common
- Patients with PAD die from CV complications
- Prevention, Protection, Plumbing
- Endo and Surgery are not competing but complimentary therapies
- “Right procedure for the right patient at the right time” on top of solid medical and exercise therapy






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Get the word out!!!

- 70% of Americans are not familiar with PAD and its devastating risks.
- Approximately all (91%) of the survey respondents would dismiss pain as just part of getting older, although pain in the leg when walking that goes away with rest is one of the first symptoms of PAD.
- More than half (53%) of respondents would wait more than a week with ongoing leg pain before calling their doctor.
- 8 in 10 Black and Hispanic respondents never had a doctor or healthcare provider talk with them about PAD.
- Amputations are 4-5x high in African Americans compared to Caucasians
- Despite 71% of Black adults having one or more risk factors for PAD or knowing someone with one or more risk factors, 65% report they are at little to no risk at all for developing PAD.
- Three-quarters of Hispanic adults have one or more risk factors for PAD or know someone with one or more risk factors but 70% think they are not at risk for developing PAD.

PADPulse.org



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Get a Pulse on PAD
Kick Off The Conversation

PAD Pulse Alliance Founding Partners


PAD Pulse Alliance Supporting Partners

PADPulse.org

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Thank you!

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Society for Vascular Medicine


CLTI Controversies – Case CLTI –
Surgical Revascularization Approach

Olamide Alabi MD MS
Chief Quality Officer, Division of Vascular Surgery and Endovascular Therapy
Emory University School of Medicine

Sunday May 17, 2024

[@SVM_tweets](#) [vascularmed.org](#) [@VascularMed](#)

1




Society for Vascular Medicine

Objectives

- Open versus Endo
- “No Stent Zones”
- Angiosome Concept

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Objectives

- ~~Open~~ versus Endo
- “No Stent Zones”
- Angiosome Concept

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Objectives

- Open versus Endo Complementary Therapies
- “No Stent Zones”
- Angiosome Concept

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Open Surgical Revascularization: Complementary Therapies

	BEST-CLI	BASIL-2
Trial Endpoint	composite of major adverse limb events or death from any cause	Composite time to above ankle amputation of the trial limb or death from any cause
Study Period	8/2014-10/2019	7/2014-11/2020
Intended N	2100	600
Actual N	1830	345
Differences in inclusion criteria	<ul style="list-style-type: none"> • Life expectancy 2 years • Availability of autogenous bypass conduit 	<ul style="list-style-type: none"> • Life expectancy >6mo • Required an infrapopliteal target
Exclusions	Excessive risk for open vascular surgery	<ul style="list-style-type: none"> • Prior vascular intervention to infrapopliteal target within prior 12mo • Ischemic pain or tissue loss considered not primarily due to PAD

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Open Surgical Revascularization: Complementary Therapies

TABLE 2. Comparison of key patient characteristics comparing BEST-CLI and BASIL-2.

	BEST-CLI COHORT 1	BEST-CLI COHORT 2	BASIL-2
Patients who did not undergo any procedure	4.3% / 1.1%	3.6% / 2.0%	8.7% / 1.2%
Cross-over to other revascularization method	3.5% / 0.4%	1.0% / 2.0%	7% / 3.5%
Median age (years)	66.9 / 67.0	68.4 / 68.8	72.4 / 72.5
Male	72% / 71.1%	71.6% / 72.4%	81% / 82%
Diabetes mellitus	72.1% / 71.6%	62.2% / 58.3%	68% / 69%
Mean eGFR (mL/min)	NA	NA	55.9% / 54%
Chronic hemodialysis	9.4% / 11.8%	12.8% / 10.1%	6% / 3%
Previous stroke	12.8% / 13.9%	19.4% / 12.1%	15% / 10%
Coronary artery disease	42.3% / 44.4%	49.5% / 53.8%	NA
Previous MI	NA	NA	24% / 13%
Previous PCI & CABG	NA	NA	26% / 46%
Previous intervention study leg	5.6% / 3.2%	10.3% / 10.2%	15% / 19%
RBC 4	20.3% / 20%	29.4% / 20.2%	15% / 15%
RBC 5 (and 4)	79.7% / 80%	70.6% / 69.8%	87% / 89% (RBC 5 only)
ASA class 3 or 4	80.8% / 75.9%	83.5% / 80.9%	NA
No. patients completing the trial	560 / 1434 (39%)	226 / 996 (22.7%)	232 / 745 (31.1%)

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Open Surgical Revascularization: Complementary Therapies

Table 1. Comparison of key outcome characteristics comparing BEST-CLI and BASIL-2.

	BEST-CLI Cohort 1	BEST-CLI Cohort 2	BASIL-2
Bypass location			
Femoro-popliteal	40%	47%	2%
Femoro-popliteal AK	NA	NA	0%
Femoro-popliteal BK	NA	NA	0%
Femoro-BTK	35.7%	37.4%	59%
Popliteal-BTK	15.7%	6.4%	40%


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Table 3. Comparison of key outcome characteristics comparing BEST-CLI and BASIL-2.

	BEST-CLI Cohort 1	BEST-CLI Cohort 2	BASIL-2
Endovascular techniques			
Balloon angioplasty	52.7%	47.2%	60%
Atherectomy	13.6%	15.4%	0%
Drug-coated balloon	27.8%	25.1%	0%
Bare-metal stent	39.3%	43.1%	10%
Drug-eluting stent	24.2%	21.5%	0%
Stent graft	8.6%	12.8%	0%
Technical success	98.3% / 84.7%	100%/80.6%	96% / 80%

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Open Surgical Revascularization: Complementary Therapies


Table 3. Comparison of key outcome characteristics comparing BEST-CLI and BASIL-2.

	BEST-CLI Cohort 1	BEST-CLI Cohort 2	BASIL-2
Technical success	98.3% / 84.7%	100%/80.6%	96% / 80%
All cause death	33% / 37.6%	25.9% / 24.1%	53% / 45%
Major amputation (above ankle)	10.4%/14.9%	15.2%/14.1%	20%/18%
AK amputation	NA	NA	NA
BK amputation	NA	NA	NA
Amputation free survival	43.3%/52.4%	41.1%/38.2%	37%/47%
Cross-over intervention during FU	NA	NA	27% / 19%
Reintervention****	9.2% / 33.1%	14.2%/25.6%	5% / 19%

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Open Surgical Revascularization

- BEST-CLI and BASIL-2 are not really comparable
- Neither should be used as an 'end all argument'
- Lessons to learn from both:
 - Nothing beats single segment great saphenous vein....if this is available
 - Get vein mapping prior to the angiogram
 - The toolbox is large – USE IT!
 - Shared decision making
 - There is a small but not insignificant cohort of patients who would benefit from primary amputation

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Open Surgical Revascularization

- Open versus Endo Complementary Therapies
- “No Stent Zones”
- Angiosome Concept

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TH

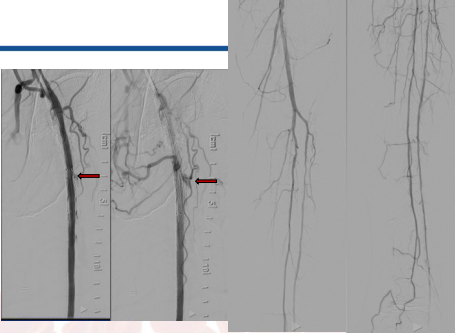
- 3/2012 Left SFA stent
- 8/2012 left SFA atherectomy and BAP of stent
- 11/2012 left SFA laser and BAP of stent
- 7/2013 left SFA revascularization
- 3/2014 BAP Left SFA in-stent stenosis
- 6/2021 shockwave lithotripsy of left CFA and popliteal + DCB BAP
- 6/2023 left SFA stent recanalization
- 8/2023 left SFA stent recanalization

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- 9/18/23 Procedure
- 1) Ultrasound guided access of the right common femoral artery
- 2) Angiogram of abdominal aorta and bilateral iliac arteries
- 3) Angiogram of the left femoral, popliteal, anterior tibial, posterior tibial, and popliteal vessels
- 4) Angiogram of the right femoral, popliteal, anterior tibial, posterior tibial, and popliteal vessels
- 3) PTA of L SFA (stents present)
- 5) Shockwave Lithotripsy
- 6) Laser atherectomy of the L SFA
- 7) Drug coated balloon of the L SFA



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And then this happened...what would you do?

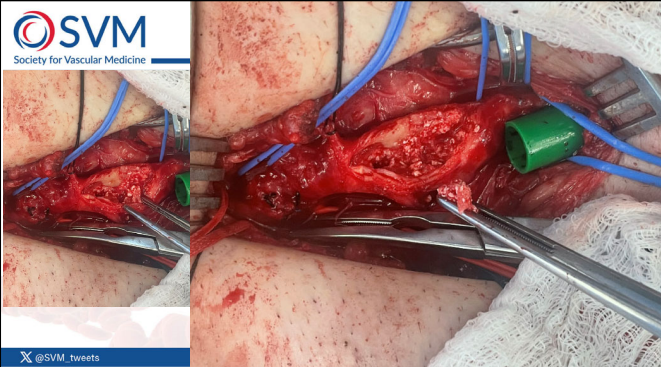


- Abi
- Lt Brachial: 118 mmHg
- Lt Posterior Tibial: 48 mmHg
- Lt Posterior Tibial: 0.41
- Lt Dorsalis Pedis: 50 mmHg
- Lt Dorsalis Pedis: 0.42

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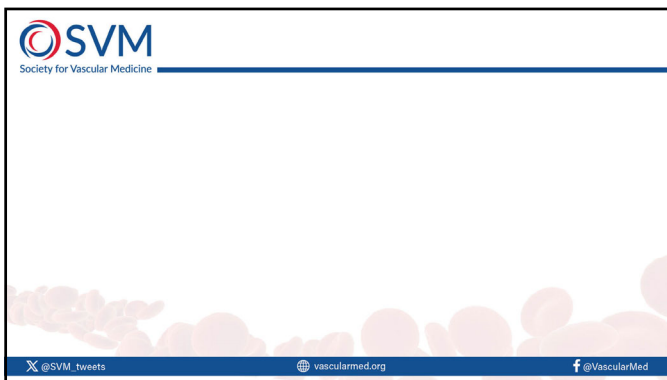


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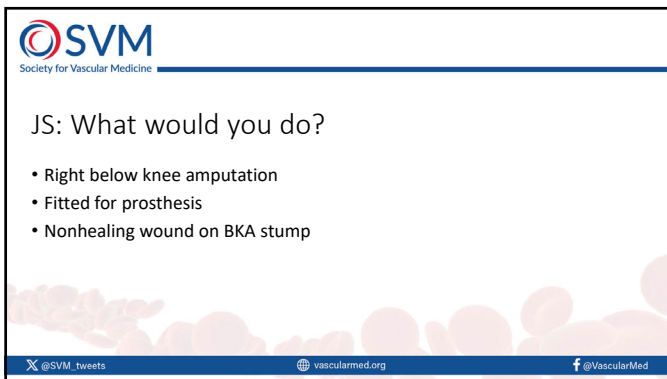
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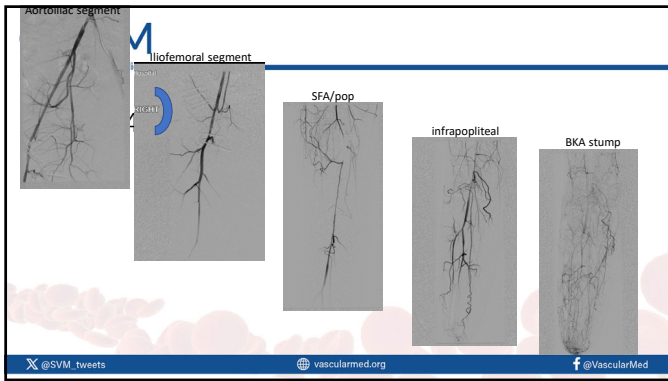
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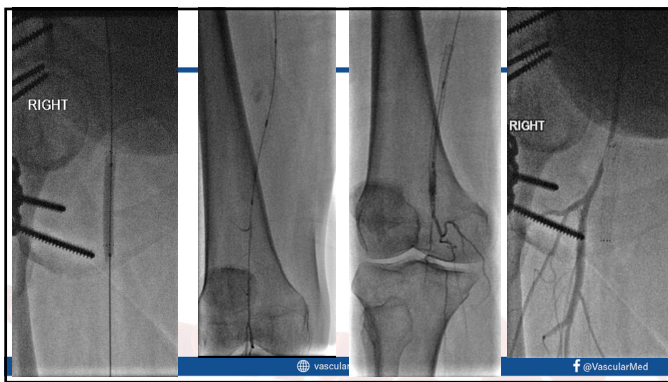
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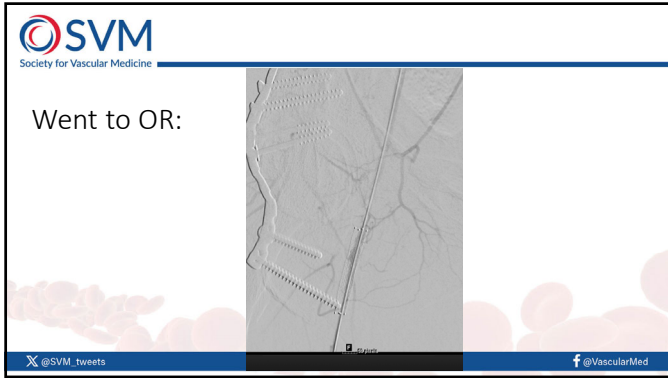
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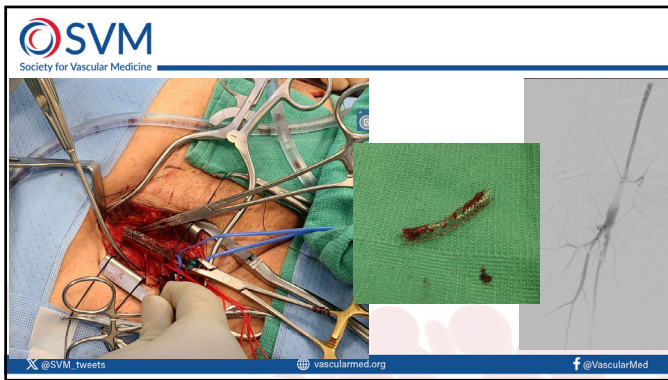
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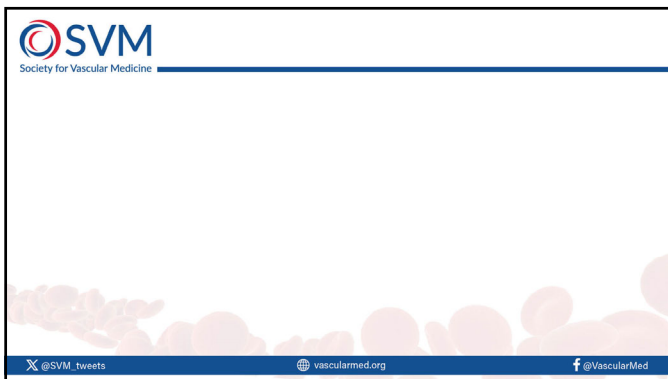
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DH: What would you do?

- 3rd toe wound with underlying osteomyelitis
- No frank cellulitis
- Vein mapping: single segment vein present
- Angiogram

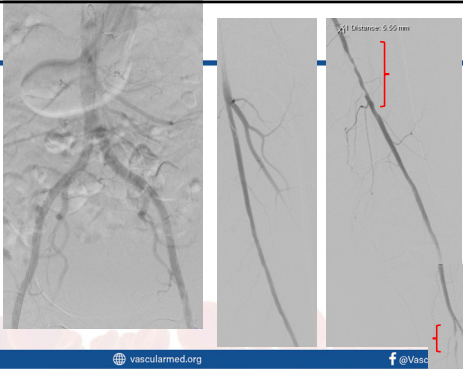
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DH

- Single segment vein present...



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CD: What would you do?


- Chronic emboli from mitral valve vegetation over time
- Developed CLTI with lateral 5th toe wound at base of toe



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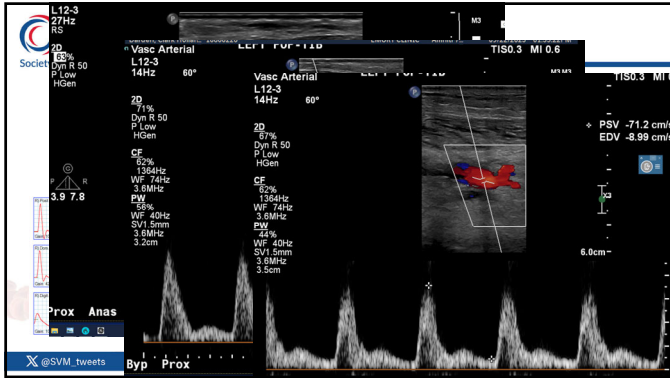
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Open Surgical Revascularization

- ~~Open versus Endo~~ Complementary Therapies
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Angiosomes


Anterior tibial angiosome Posterior tibial angiosome Peroneal angiosome

Anterior tibial artery Posterior tibial artery Peroneal artery

Medial plantar branch Lateral plantar branch Calcaneal branch

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Direct versus Indirect Revascularization

- Meta-analysis, 9 studies
 - 715 direct revascularization
 - 575 indirect revascularization
- Direct revascularization associated with:
 - Lower risk of unhealed wound (HR 0.64, 95% CI 0.52-0.8, I2 0%)
 - Lower risk of major amputation (0.44, 95% CI 0.26-0.75, I2 62%)
 - Limb salvage rates
 - 1 year: 86.2% direct; 77.8% indirect
 - 2 years: 84.9% direct; 70.1% indirect

Biancari F, Juvonen T. Angiosome-targeted lower limb revascularization for ischemic foot wounds: systematic review and meta-analysis. *Eur J Vasc Endovasc Surg.* 2014;47(5):517-522.

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Open Surgical Revascularization

- ~~Open versus Endo~~ Complementary Therapies
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Questions

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Harrington Heart & Vascular Institute
Cleveland, Ohio

CLTI Management Endovascular Revascularization

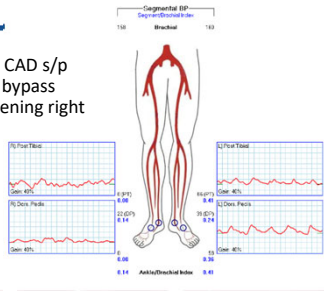

Yulanka Castro, MD
Clinical Assistant Professor of Medicine
University Hospitals Harrington Heart and Vascular Institute
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@YSCastroMD

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SVM Case
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- 79-year-old woman with history of CAD s/p CABG, PAD s/p PTFE right fem-pop bypass presenting with rest pain and worsening right posterior ankle ulcer



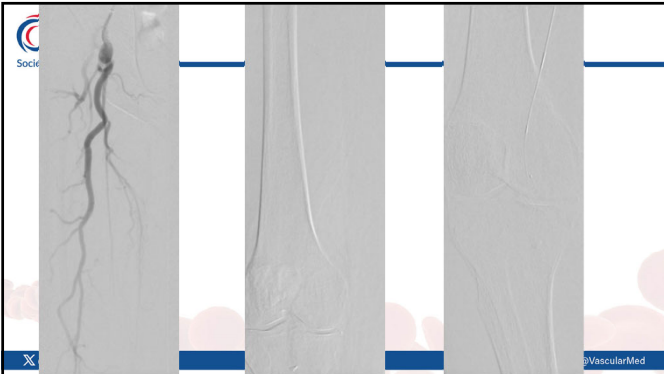
Right Foot Pulse: 0.91% (0.91)
Left Foot Pulse: 0.91% (0.91)
Right Ankle Pulse: 0.91% (0.91)
Left Ankle Pulse: 0.91% (0.91)

Right Foot Pulse: 0.91% (0.91)
Left Foot Pulse: 0.91% (0.91)
Right Ankle Pulse: 0.91% (0.91)
Left Ankle Pulse: 0.91% (0.91)

Right Ankle/Brachial Index: 0.91
Left Ankle/Brachial Index: 0.91

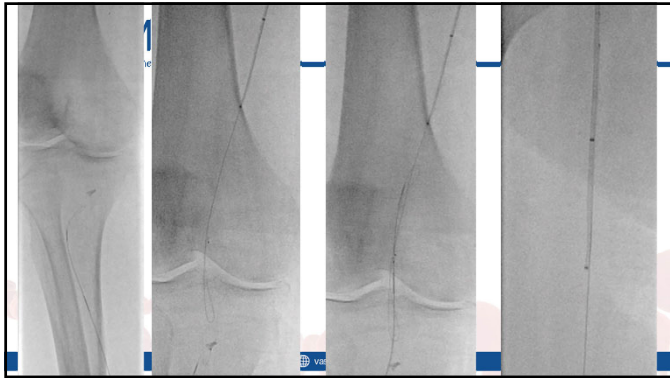
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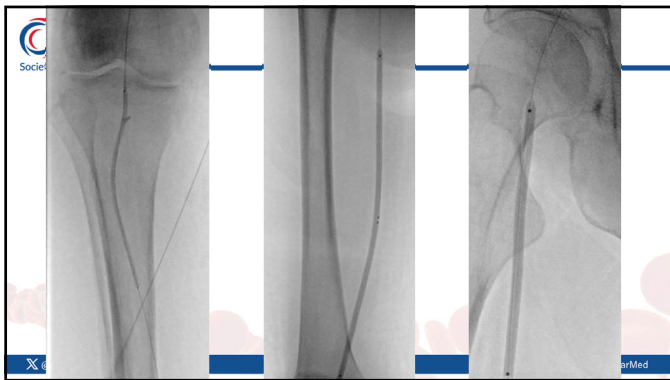


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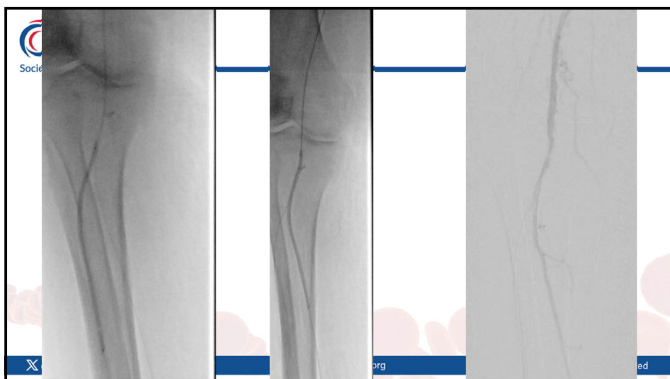
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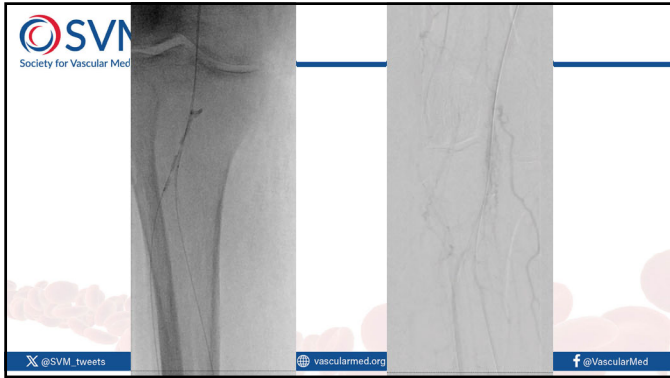
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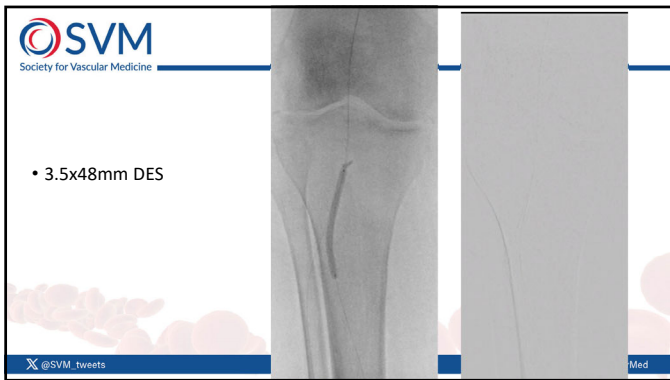
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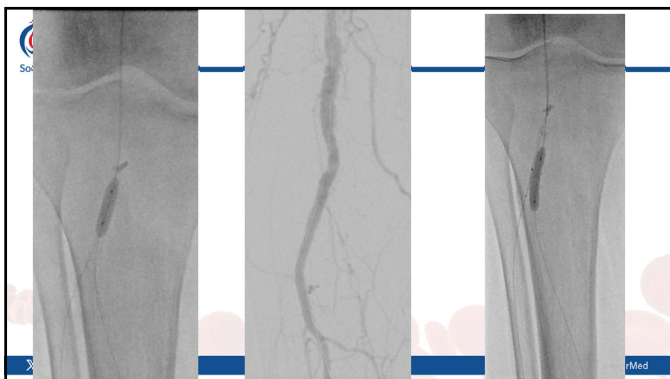
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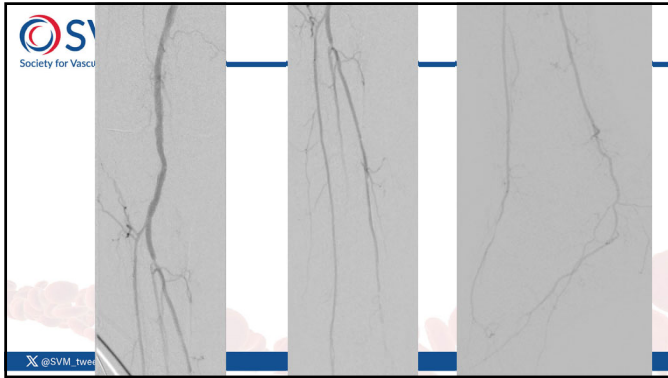
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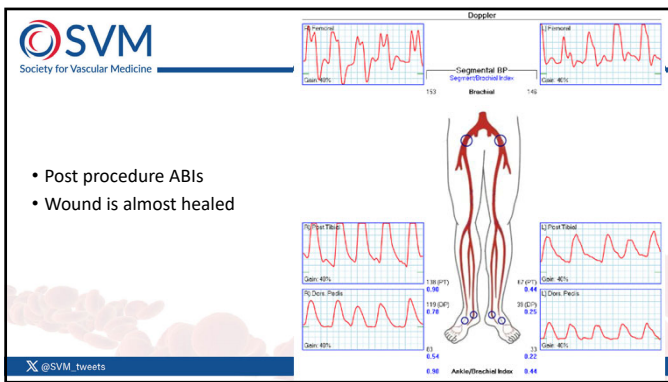
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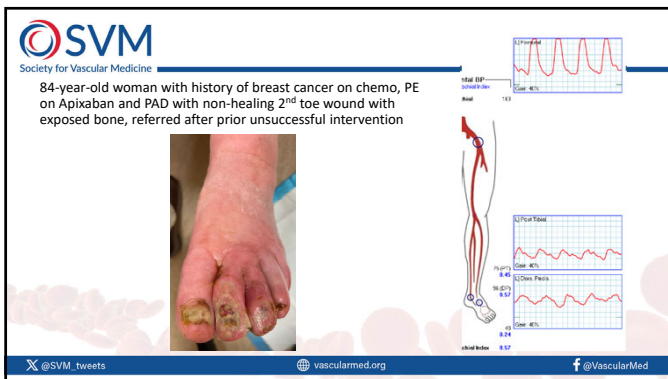
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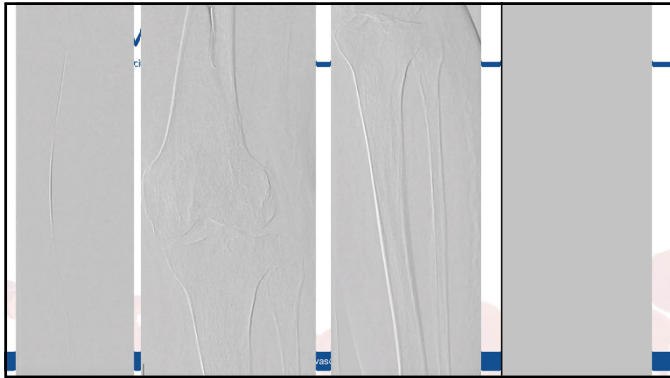
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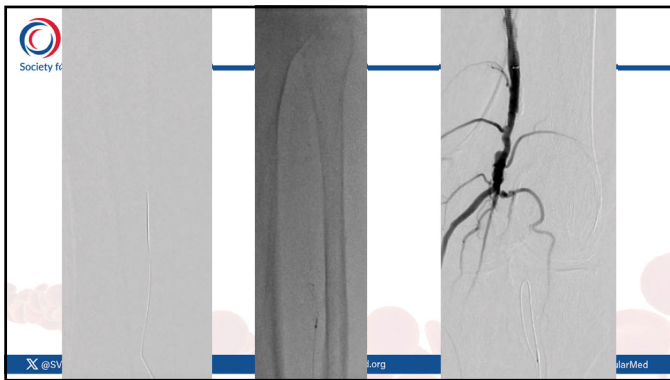
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SVM BASIL-2 Trial
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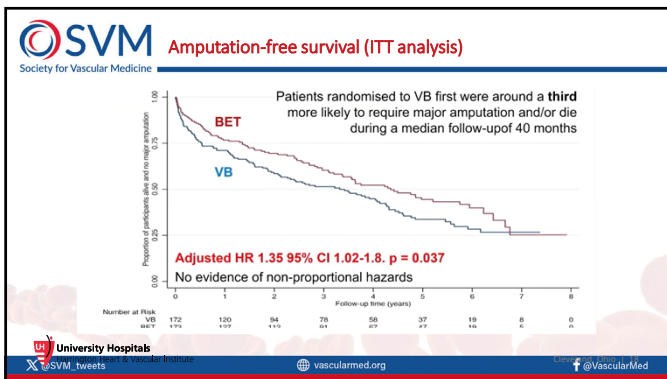
A vein bypass first versus a best endovascular treatment first revascularisation strategy for patients with chronic limb threatening ischaemia who required an infra-popliteal, with or without an additional more proximal infra-inguinal revascularisation procedure to restore limb perfusion (BASIL-2): an open-label, randomised, multicentre, phase 3 trial

Andrew W Bradbury, Catherine A Moolke, Matthew Peggelwell, Lewis Meechem, Gareth R Batz, Lisa Kelly, Ian Chetter, Athanasios Diamantopoulos, Anil Ganeshan, Jack Hall, Simon Hobbs, Kim Houldred, Hugh Jarrett, Suzanne Lockyer, James Malmstadt, Jai V Patel, Smitas Patel, S Touqeer Rashid, Athanasios Saratzis, Gemma Siles, D Julian A Scott, Harry Zayed, Jonathan J Deeks, on behalf of the BASIL-2 investigators

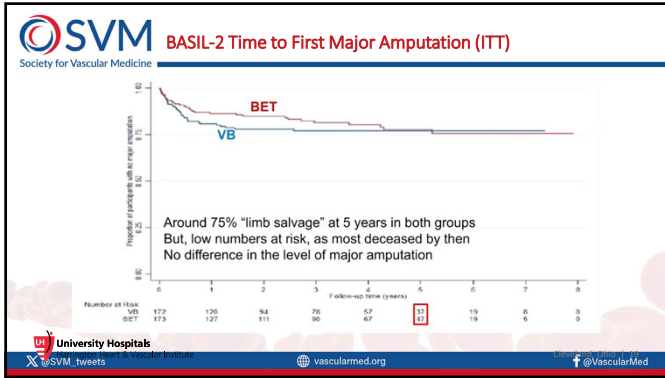
THE LANCET

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Outcomes

- Number of re-interventions was higher in BET (19%) than in the VB (5%) (0.27 [0.13-0.55])
- Crossover interventions – more common in VB (27%) than in the BET (19%) (1.43 [0.94-2.18])
- NO differences in 30-day morbidity and death, MALE, MACE, relief of ischemic pain, or QoL

	Vain bypass group (n=172)	Best endovascular treatment group (n=173)	Estimate (95% CI)
Subsequent intervention	55 (29%)	58 (24%)	RR 0.94 (0.68 to 1.28); uRR -0.03 (-0.13 to 0.06)
Reintervention	9 (5%)	33 (19%)	RR 0.27 (0.13 to 0.55); uRR -0.14 (-0.21 to -0.07)
Crossover intervention	46 (27%)	33 (19%)	RR 1.43 (0.94 to 2.18); uRR 0.08 (-0.01 to 0.17)

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Patient Characteristics: BASIL-2 and BEST-CLI

	BASIL-2	BEST-CLI (cohort 1)	BEST-CLI (cohort 2)
Enrolled participants	345	1434	396
Median age (yrs)	73	67	68
Diabetes	68%/69%	72%/72%	62%/58%
ESRD on HD	6%/3%	13%/14%	13%/10%
Previous intervention study leg	31%/39%	10%/10%	6%/5%
Antiplatelet use	76%/80%	ASA – 67%/67% Clopidogrel 19%/25%	ASA – 71%/71% Clopidogrel 25%/28%
Cholesterol lowering agent	75%/80%	71%/70%	79%/77%
Tissue loss	87%/89%	80%/80%	71%/70%
Tibial disease treated	100%	51%/51%	46%/43%

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Primary Outcomes: BASIL-2 and BEST-CLI

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- BEST-CLI
 - MALE – major amputation above ankle, major limb reintervention (new bypass graft, graft revision, thrombectomy, or thrombolysis) or death from any cause
- BASIL-2
 - Amputation-free survival – time to major amputation above ankle or death from any cause
 - What ultimately matters to patients – to avoid amputation and prolong life
 - Does not reflect burden of major reinterventions

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Key Comparisons: BASIL-2 and BEST-CLI


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	BASIL-2	BEST-CLI (cohort 1)
Endo technical success	87%	85%
Bypass technical success	96%	98%
Crossover (endo to bypass)	19%	23%
Major amputation/Death		
Bypass	63%	43%
Endovascular	53%	53%

Outcome	Surgery	Endovascular Therapy	Hazard Ratio (95% CI)
Efficacy			
Primary outcome: major adverse limb event or death from any cause — no, total no. (%)	302/709 (42.6)	408/711 (57.4)	0.68 (0.59-0.79)
Secondary outcomes — no, total no. (%)			
Death from any cause	234/709 (33.0)	267/711 (37.6)	0.98 (0.82-1.17)
Above-ankle amputation of the index limb	74/709 (10.4)	106/711 (14.9)	0.73 (0.54-0.98)
Intervention in index limb			
Major	65/709 (9.2)	167/711 (23.5)	0.35 (0.27-0.47)

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Data we need from BEST-CLI and BASIL-2

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- Few patients with CLTI are deemed suitable and have an optimal vein for infra-popliteal bypass
 - Risk profiles of patients deemed not appropriate for surgery
- Angiographic lesion-level data and severity of disease morphology in both arms
- Registry data of patients that were treated during trial period but not enrolled – understand patients who BEST-CLI and BASIL-2 results does not apply to
 - No equipoise when disease morphology is straightforward for endo or patient unfit for open surgery

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Individual Assessment

- Involvement of CFA/PFA
- Younger, low risk patient
- Suitable vein
- Extensive tibial disease with poor targets
- No suitable vein
- Older patient, poor surgical candidate

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1 C-ID An evaluation for revascularization options should be performed by an interdisciplinary care team (Table 9) before amputation in the patient with CLI.

Impact of Interdisciplinary System-Wide Limb Salvage Advisory Council on Lower Extremity Major Amputation

Mehdi H. Shishebor, DO, MPH, PhD, Tarek A. Hammad, MD, Tonia J. Rhone, MS, Ahmad Younes, MD, Norman Kumins, MD, Abdullah Abdullah, MD, Jun Li, MD, Karen Harth, MD, Teresa L. Carman, MD, Heather L. Gornik, MD, Peter J. Pronovost, MD, PhD, and Vikram S. Kashyap, MD

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SVM Learning Points

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- Reminder poor prognosis of patients with CLTI. ~ 50% of patients died by 5 years
- Primary prevention – early detection of PAD and institution of GDMT and lifestyle interventions
- Saphenous vein mapping and surgical risk assessment should be more regularly considered
- Management of patients with CLTI requires multidisciplinary expertise in limb salvage programs

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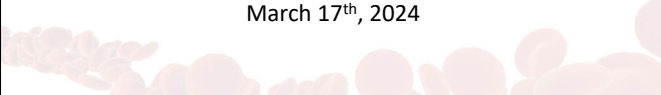
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Large Vessel Vasculitis

Alexandra Solomon, MD, RPVI
March 17th, 2024



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Disclosures

- None, but any are welcome!



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Learning Objectives

At the end of this session, participants will be able to:

1. Devise a differential diagnosis for large vessel vasculitis (LVV).
2. Complete a basic diagnostic workup for LVV.
3. Discuss the basics of LVV management.



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Our Patient

- 32 yo woman
- No significant past medical history
- 1 year of progressive right arm and bilateral leg cramping with activity; 1 month of left neck tenderness, and intermittent dizziness



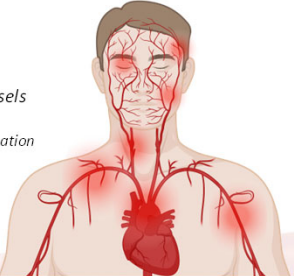
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Large Vessel Vasculitis

- **Vasculitis:** inflammation of blood vessels
 - WBC infiltration into vessel wall
 - Narrowing, obstruction, aneurysm formation
- Large vessels (mostly)
 - Aorta and its branches
 1. Giant cell arteritis
 2. Takayasu arteritis

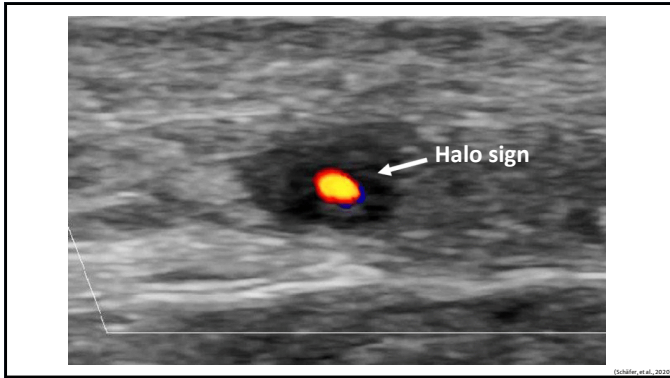


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Infectious causes (eg, endocarditis, HBV, HCV, HIV)
Atherosclerosis
Thromboembolic disease
Congenital causes (eg, aortic coarctation, middle aortic syndrome)
Hereditary disorders (eg, Marfan syndrome, Ehlers-Danlos syndrome)
Fibromuscular dysplasia
Hypercoagulable states (eg, APS, TTP)
Vasospastic disorders (eg, RCVS, drug exposures)
Other multisystem inflammatory disorders (eg, sarcoidosis, Susac syndrome)
Malignancy (eg, lymphoma, leukemia)
Iatrogenic (eg, postradiation therapy)
IgG4-related disease

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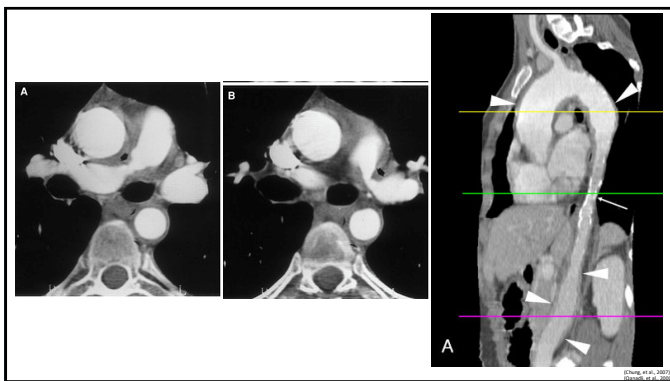
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GCA Diagnostic Criteria (score ≥ 6)

Criteria	Score
ABSOLUTE REQUIREMENT	
Age ≥ 50 years at time of diagnosis	
ADDITIONAL CLINICAL CRITERIA	
Morning stiffness in shoulders/neck	+2
Sudden visual loss	+3
Jaw or tongue claudication	+2
New temporal headache	+2
Scalp tenderness	+2
Abnormal examination of the temporal artery ¹	+2
LABORATORY, IMAGING, AND BIOPSY CRITERIA	
Maximum ESR ≥ 50 mm/hour or maximum CRP ≥ 10 mg/liter ²	+3
Positive temporal artery biopsy or halo sign on temporal artery ultrasound ¹	+5
Bilateral axillary involvement ³	+2
FDG-PET activity throughout aorta ⁴	+2

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Takayasu Diagnostic Criteria (score ≥5)

ABSOLUTE REQUIREMENTS

- Age ≥ 60 years at time of diagnosis
- Evidence of vasculitis on imaging*

ADDITIONAL CLINICAL CRITERIA

Female sex	+1
Angina or ischemic cardiac pain	+2
Arm or leg claudication	+2
Vascular bruit†	+2
Reduced pulse in upper extremity‡	+2
Carotid artery abnormality†	+2
Systemic blood pressure difference in arms ≥ 20 mm Hg	+1

ADDITIONAL IMAGING CRITERIA

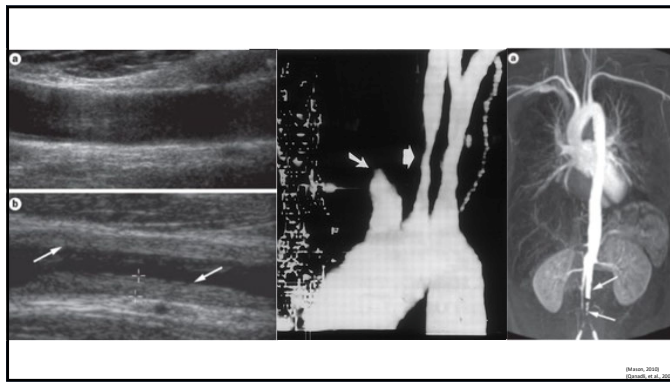
Number of affected arterial territories (select one)†

One arterial territory	+1
Two arterial territories	+2
Three or more arterial territories	+3
Symmetric involvement of paired arteries†	+1
Abdominal aorta involvement with renal or mesenteric involvement†	+3

*MRA, CT angiography, or digital subtraction angiography
†Physical exam or Doppler ultrasonography
‡At least two arteries

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Additional Management Considerations

- Blood pressure management: Always use the **HIGHEST** blood pressure
- Long-term effects of steroids:
 - Osteoporosis
 - Infections
 - Weight gain/metabolic syndrome
 - Gastritis
 - Psychiatric symptoms
- Pregnancy
- Medicine is a team sport...

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Medicine is a team sport!

- Rheumatology
- Internal Medicine
- Cardiology
- Neurology
- Ophthalmology
- Vascular Medicine
- Vascular Surgery
- Dermatology
- Gynecology (high-risk OB)

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Take Home Points

- Vasculitides have overlapping clinical features; making a diagnosis can be difficult!
- Evaluation includes history, physical exam, labs, vascular imaging (CTA, MRA, US), and sometimes biopsy (GCA).
- Steroids are generally the mainstay of treatment for LVV (significant side effects). Steroid-sparing agents are used as well. Surgical intervention sometimes needed.
- Take blood pressures in all 4 limbs; use **HIGHEST** reading and correlate with anatomy.
- The expertise of multiple specialties is often needed to care for these complex patients!

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Thank you!

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- Jennette JC, Falk RJ, Bacon PA, et al. 2012 revised International Chapel Hill Consensus Conference Nomenclature of Vasculitides. *Arthritis Rheum.* 2013; 65:1. Reproduced with permission from John Wiley & Sons, Inc. Copyright © 2013 by the American College of Rheumatology.
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- Qanadli SD, Sisakian JF, Rocha P, Piette AM, Lacombe P. Takayasu's arteritis : spiral CT angiography findings. *Circulation.* 2000 Jan 25;101(3):345-7. doi: 10.1161/01.cir.101.3.345. PMID: 10645393.
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- UpToDate. (n.d.). https://www.uptodate.com/contents/overview-of-and-approach-to-the-vasculitides-in-adults?search=large-vessel-vasculitis&source=search_result&selectedTitle=1~150&usage_type=default&display_rank=1#H17

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Fibromuscular Dysplasia

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1

Disclosures

- None

2

Objectives

- Identify under recognized vascular disorders
- Understand the evaluation and treatment of patients with known or suspected FMD/SCAD and other related arteriopathies
- Know the differential diagnosis for nonatherosclerotic vascular diseases

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Case

- 43-year-old woman presents to clinic with severe HTN and headaches
- Past medical history significant for active smoking
- Family history of ? aneurysm in family
- Blood pressure remains >160's despite two medications

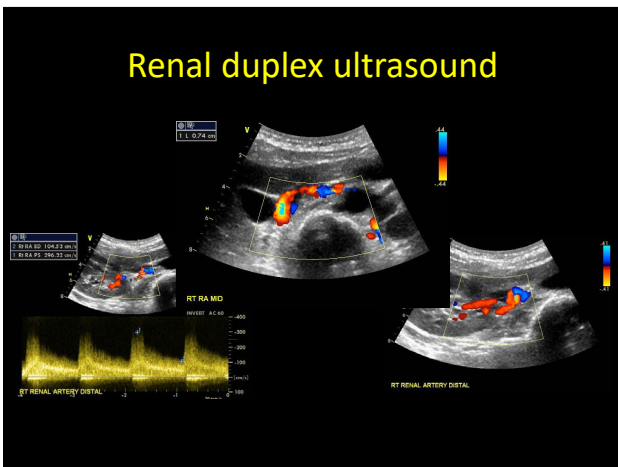
4

(continued)

- Referred to Emory vascular medicine for further evaluation
- Office blood pressure: 211/93 mmHg
- Renal duplex ultrasound was performed

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Renal duplex ultrasound



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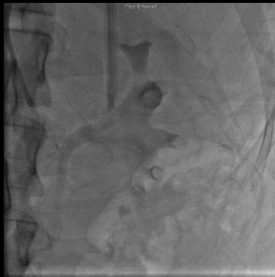
Right Renal Artery



Pd/Pa=0.51

7

Left Renal Artery



Pd/Pa=0.92

8

Right Renal Artery Angioplasty



9

Right Renal Artery Final Result



Pd/Pa=0.98

10

Clinical Follow-up

- Dramatic reduction in medication requirement
- Blood pressure stable on one medication
- Headaches resolved
- Follow-up renal duplex ultrasound demonstrates normal velocities

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Fibromuscular dysplasia

- Nonatherosclerotic, noninflammatory disease
- Etiology is unknown, genetics unknown
- Causes stenosis, occlusion, aneurysm, dissection
- Most commonly involves renal and extracranial carotid arteries (65% of cases)
- More common in women and younger patients
- Average delay in first symptom to diagnosis of 4-9 years
- Hypertension, headache, pulsatile tinnitus

Olin, JW. *Circulation*, 2012.

12

Epidemiology

- The prevalence of FMD is unknown
- Renal FMD
 - 1% with autopsy data
 - 3-6% with angiographic data
- Carotid FMD
 - 4/20,244 in one autopsy series
 - 0.3-3.2% with angiographic data

Heffelfinger MK. *Am J Clin Pathol*, 1970.
 Olin JW. *Circulation*, 2012.
 Touze E. *Int J Stroke*, 2010.
 Schievink WI. *N Engl J Med*, 2001.

13

Epidemiology

- The cause of FMD is unknown
- More common in women by a ratio of 9:1
- Dose dependent relationship with smoking
 - 30-37% of FMD patients are smokers
 - Compared with 18% of age/sex matched peers

Sang CN. *Hypertension*, 1989.
 Savard S. *Hypertension*, 2013.

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PHACTR1 and COL5A1 Variants

- *PHACTR1* rs9349379-A variant
 - Associated with cervical artery dissection (CeAD), hypertension, migraine headache, and FMD
 - First genetic susceptibility locus for FMD

PLOS Genet 2016; 12: e1006367

- *COL5A1* variant associated with an arteriopathy in 4 unrelated families
 - Associated with arterial aneurysms, dissections, tortuosity, and FMD

Arteriosclerosis, Thrombosis, and Vascular Biology. 2020;40:2686–2699

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Classification of FMD

Table 1. Classification of Fibromuscular Dysplasia

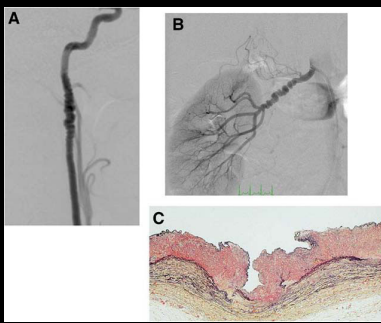
Histological	Angiographic		
	Harrison and McCormack (1971) ¹³	French/Belgian Consensus (2012) ²	American Heart Association (2014)
Medial Medial fibroplasia (60%–70%) Perimedial fibroplasia (15%–25%) Medial hyperplasia (5%–15%)		Multifocal	Multifocal
Intimal fibroplasia (1%–2%) Adventitial (<1%)		Unifocal (<1 cm) Tubular (≥1 cm)	Focal*

*There may be multiple areas of focal disease (eg, renal artery and carotid artery in the same patient). Focal and multifocal disease can occur in the same patient.

Olin JW. *Circulation*, 2014.

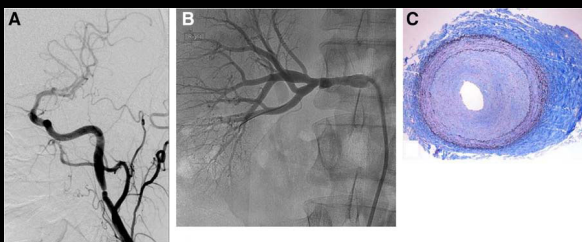
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Multifocal FMD



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Focal FMD



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Presenting Signs and Symptoms from US FMD Registry (447 patients)

- Hypertension 285 (63.8)
- Headache 234 (52.4)
- Current headache 135 (30.2)
- History of headache 173 (38.7)
- **Pulsatile tinnitus 123 (27.5)**
- Dizziness 116 (26)
- Cervical bruit 99 (22.2)
- Neck pain 99 (22.2)
- Tinnitus 84 (18.8)
- Chest pain or shortness of breath 72 (16.1)
- Flank/abdominal pain 70 (15.7)
- Aneurysm 63 (14.1)
- Cervical dissection 54 (12.1)
- Epigastric bruit 42 (9.4)
- Hemispheric transient ischemic attack 39 (8.7)
- Postprandial abdominal pain 35 (7.8)
- Stroke 31 (6.9)
- Claudication 23 (5.2)
- Amaurosis fugax 23 (5.2)
- Weight loss 23 (5.2)
- Horner syndrome 21 (4.7)
- Renal artery dissection 14 (3.1)

Olin JW. *Circulation*, 2012.

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Renal FMD presentation

- Most commonly presents with hypertension in a younger patient
 - Average age at presentation is 43 years
- Abdominal bruit is presenting sign in 9%
- Abdominal pain with renal artery dissection
- CKD and progression to ESRD is uncommon
- Headaches are common

Olin JW. *Circulation*, 2012.

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Renal FMD diagnosis

- Renal duplex ultrasound
 - Vessel tortuosity, turbulent flow, and elevated velocities in the mid to distal renal arteries
 - Cannot see beading or quantify stenosis
- CTA
 - 100% correlation with angiogram in 1 study
 - May not see mild FMD or branch vessel FMD
- MRA
 - Good at detecting beading, less accurate with stenosis
 - Worse spatial resolution
 - Motion artifact
- Catheter angiography
 - Gold standard
 - <10 mmHg gradient is normal

Sabharwal R. *Eur J Radiol*, 2007.
Willoteaux S. *Radiology*, 2006.

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Renal FMD revascularization


<p>Indications</p> <ul style="list-style-type: none"> • Resistant hypertension • Hypertension of short duration • Renal artery dissection • Renal artery aneurysm • Branch renal artery disease and hypertension • Preservation of renal function 	<p>Modality</p> <ul style="list-style-type: none"> • PTA <ul style="list-style-type: none"> – Avoid stent – Approximately 50% cure, 30% improved, 10-20% not improved • Surgery <ul style="list-style-type: none"> – Small arteries, branch disease, aneurysms – Bypass with reversed SVG – 30-70% cure rate
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Olin JW. *Circulation*, 2014.

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Carotid FMD presentation

- Bruit (22%), headache (60%), **pulsatile tinnitus (presenting symptom in 37%)**, neck pain, dizziness
- TIA (13%), cervical artery dissection (12%), stroke (10%)
 - Stenosis, embolization, thrombosis, dissection, aneurysm
- Cerebral, carotid, vertebral and basilar aneurysms present in 7-10%
 - Intracranial hemorrhage in 1%



Olin JW. *Circulation*, 2012.

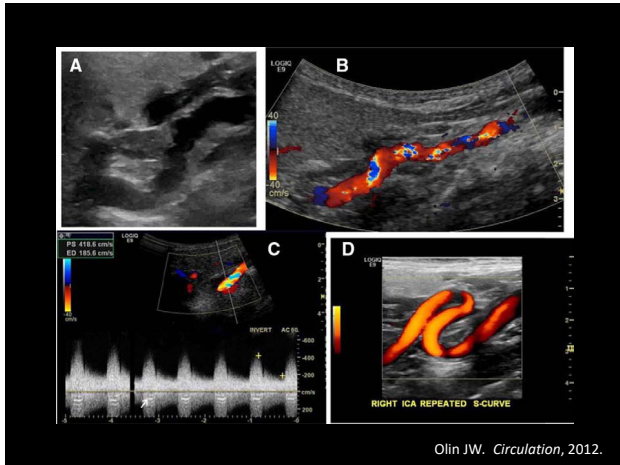
23

Carotid FMD carotid duplex

- Diagnosis
 - Vessel tortuosity, turbulent flow, and elevated velocities in the mid to distal carotid arteries
 - **“S curve”**, marked tortuosity and elongation of vessels in 34% of FMD patients versus 3% general population
 - Older patients may have both atherosclerosis and FMD
 - No data on diagnostic accuracy
 - No velocity criteria for stenosis
- Surveillance
 - Duplex annually when stable (class 2a)
 - Medial FMD generally not progressive

Sethi S. *J Am Coll Cardiol*, 2012.
Brott TG. *Circulation*, 2011.

24



Olin JW. *Circulation*, 2012.

25

Carotid FMD CTA

- Image extracranial and intracranial vessels
- Accurately identify FMD, dissections, aneurysms (>3mm), atherosclerosis
- Difficult to quantify stenosis in beading segments
- Invasive angiogram reserved for intervention/diagnostic uncertainty

Donmez H. *Eur J Radiol*, 2011.

26

FMD and Aneurysms

- **21.6%** of US Registrants have at least 1 aortic or arterial aneurysm
 - Renal artery 32.1%
 - EC carotid 27.9%
 - Mesenteric/Celiac 16.4%
 - Aorta 12.1%
 - Cerebral 12.1%
 - Vert/basilar 11.4%
- 20-40% will have aneurysm and/or dissection
- One time brain to pelvis cross sectional imaging for *all* FMD patients is recommended

41.7%
N = 384

Aneurysm
N = 147

Both A & D
N = 53

Dissection
N = 184

No Aneurysm or Dissection
N = 537

J Am Coll Cardiol. 2016 Jul 12;68(2):176-85.

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FMD Management

- Avoid neck manipulation and heavy lifting
- Avoid tobacco, hormones, and stimulants
- No data on statins or OCPs
- Antiplatelet therapy
- Anti-hypertensives if needed
- Revascularization when necessary, typically not needed
- CTA/MRA brain to pelvis to evaluate for occult FMD, aneurysm, dissection
- Patients diagnosed as having FMD at an older age have a more benign disease process and less severe symptoms

JAMA Cardiol. 2018 Aug; 3(8): 756-760.

Olin JW. *Circulation*, 2012.

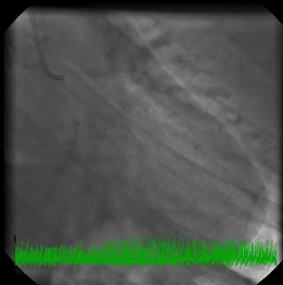
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Case

- 42-year-old woman with no medical history notes new chest pain for 2-3 days
- She presented to the ED where abnormal troponins prompt urgent heart catheterization
- She denies any known family hx of heart disease, smoking history, oral contraceptives
- Pregnancy test negative

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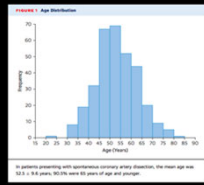
Left coronary angiography



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Spontaneous Coronary Artery Dissection

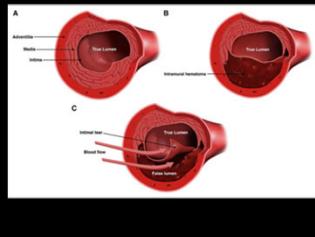
- Nontraumatic/non-iatrogenic
- Estimated prevalence:
 - Autopsy 0.4%
 - Coronary angiography 0.2%-1.1%
 - ACS 0.1%-4%
- Average age 43 to 53 years
- Female
- Most common cause of pregnancy associated MI
- Recurrence 20-25% at 5 years



Circ J. 2014 Aug 25;78(9):2099-110. E2014
 J Am Coll Cardiol. 2017 Aug 29;70(9):1148-1158.

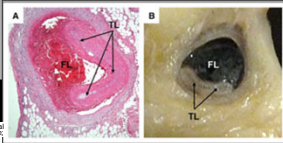
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SCAD Pathophysiology



"Inside out" hypothesis
 (intimal tear → blood enters sub-intimal space from true lumen → generates false lumen)

vs.
"Outside in" hypothesis
 (spontaneous hematoma forms from vasa vasorum within vessel wall)



Rayns RN, et al. American Heart Association Council on Peripheral Vascular Disease, Council on Clinical Cardiology, Council on Cardiovascular and Stroke Nursing, Council on Genomics and Precision Medicine, and Stroke Council. Spontaneous Coronary Artery Dissection: Current State of the Science. A Scientific Statement From the American Heart Association. Circulation. 2018 May 8;137(9):e232-e247.

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SCAD Precipitating Factors

Precipitating factors	>50% Patients recall a precipitating factor ^{1,2}
Intense exercise (isometric or aerobic)	
Intense Valsalva	
Retching, vomiting, bowel movement, coughing, lifting heavy objects	
Intense emotional stress	
Labor and delivery	
Recreational drugs (cocaine, methamphetamines)	
Exogenous hormones/hormone modulators	
β-blocker injections, corticosteroid injections, clomiphene	

Table 4. Precipitating stressors and potential predisposing conditions

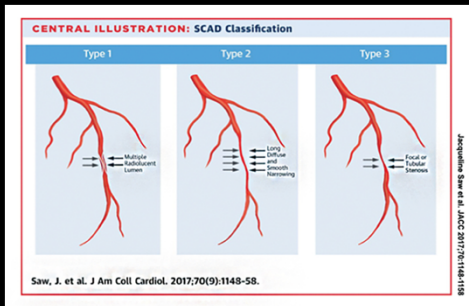
N (%)	N = 750
Precipitating stressors	
Emotional stress (rated high or severe)	377 (50.3)
Perceived stress scale >20	288 (41.2)
Unusually intense physical stress	216 (28.9)
Isometric stress >50 lb	74 (9.8)
Cocaine/amphetamine use	2 (0.3)
Valsalva-type stress	90 (12.0)
No precipitating factor	252 (33.6)

Rayns RN, et al. American Heart Association Council on Peripheral Vascular Disease, Council on Clinical Cardiology, Council on Cardiovascular and Stroke Nursing, Council on Genomics and Precision Medicine, and Stroke Council. Spontaneous Coronary Artery Dissection: Current State of the Science. A Scientific Statement From the American Heart Association. Circulation. 2018 May 8;137(9):e232-e247.

Law J, et al. Canadian spontaneous coronary artery dissection cohort study: incidence and 30-day outcomes. Eur Heart J. 2019 Jun 14;40(11):1185-1192.

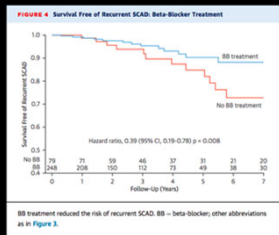
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SCAD Classification



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Acute SCAD Management



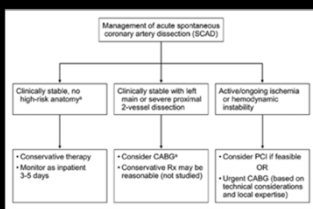
Heparin and DAPT: reduces thrombus burden vs theoretical risk of bleeding and intramural hematoma

Beta blockers – likely decreases risk of SCAD recurrence

Saw, J, Humphries K, Ayimong E, Seifak T, Prakash B, Starovoytov A, Mancini GBJ. Spontaneous Coronary Artery Dissection: Clinical Outcomes and Risk of Recurrence. J Am Coll Cardiol. 2017;70(9):1148-58.

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Acute SCAD Management



“Healing” of SCAD lesions in 70-97% patients managed conservatively - within 1 month of index event

Prolonged in hospital monitoring (3-5 days) to observe for early dissection extension or new recurrent SCAD (in 5-10%)

Stays BN, et al. American Heart Association Council on Peripheral Vascular Disease, Council on Clinical Cardiology, Council on Cardiovascular and Stroke Nursing, Council on Sports and Prevention Medicine, and Stroke Council. Spontaneous Coronary Artery Dissection: Current Role of the Specialist. A Scientific Statement From the American Heart Association. Circulation. 2018 May 8;137(19):e529-e557.

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SCAD Revascularization

- Catheter based angioplasty/stenting carries high risk of complications
 - 36% technical failure (including residual stenosis)
 - Iatrogenic dissection
 - False lumen propagation
 - Stent strut mal-apposition
 - 6% stent thrombosis
- 12% emergency CABG (high risk of graft failure)
- May be necessary for ongoing ischemia



Saw J, et al. Spontaneous coronary artery dissection: association with predisposing arteriopathies and precipitating stressors and cardiovascular outcomes. *Circ Cardiovasc Interv.* 2014.

Adam D, Alfonso F, Maas A, White C, Writing Committee. European Society of Cardiology, acute cardiovascular care association. SCAD: 15th update: a position paper on spontaneous coronary artery dissection. *Heart J.* 2018 Sep 21; 39(8):1325-1336.

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Outpatient Management after SCAD

- Aspirin
- Beta blocker, HR 0.36
- Cardiac rehab, target HR ~70% max HR
- Lift less than 30 pounds
- Avoid isometrics / Valsalva
- Avoid stimulants, exogenous hormones
- Counseling about pregnancy / contraception
- CTA/MRA brain to pelvis
- Vasodilators for angina

J Am Coll Cardiol. 2017;70(9):1148.

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Spontaneous Coronary Artery Dissection: Prevalence of Predisposing Conditions Including Fibromuscular Dysplasia in a Tertiary Center Cohort

J Am Coll Cardiol Interv. 2015;6(1):44-52. doi:10.1016/j.jcin.2012.08.017

FMD in ≥ 1 noncoronary territories	86.0% (43)
FMD in ≥ 2 noncoronary territories	42.0% (21)
FMD not observed	14.0% (7)
Incomplete screening	10.0% (5)
Screened cerebral, renal, iliac	4.0% (2)
FMD vascular involvement (n = 43)	
Renal arteries	58.1% (25)
Iliac arteries	48.8% (21)
Cerebrovasculature	46.5% (19)
Cerebral aneurysm	16.3% (7)

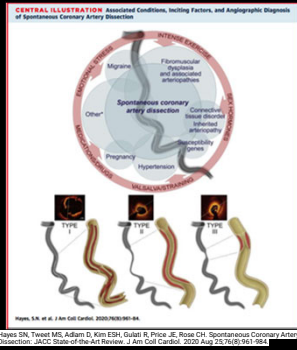
Involvement With Noncoronary FMD Among These Patients With SCAD (N = 50)

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SCAD Associated conditions

Table 2. Conditions and Factors Associated With SCAD

Associated Condition or Factor	Reported Prevalence in Cohort Studies, %
Fibromuscular dysplasia	25-40 ^{10,11}
Pregnancy	2-8 ^{11,12}
Multiparity (at birth)	8.9-10 ¹¹
Inherited arrhythmia and connective tissue disorder (see Table 4)	1.2-3.0 ¹¹
Marfan syndrome, Loeys-Dietz syndrome, vascular Ehlers-Danlos syndrome, α ₁ -antitrypsin deficiency, polycystic kidney disease	
Exogenous hormones	10.7-12.8 ¹¹
Oral contraceptives, postmenopausal therapy, infertility treatments, testosterone, corticosteroids	
Systemic inflammatory disease	<1-8.9 ¹¹
Systemic lupus erythematosus, Crohn disease, ulcerative colitis, polyarteritis nodosa, sarcoidosis, Churg-Strauss syndrome, Wegener granulomatosis, Rheumatoid arthritis, Kawasaki disease, celiac disease	
Migraine headache	NR
Coronary artery spasm	NR



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Thank you

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- 404-686-8203

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Lipid Management: Biomarkers & More

Merry Ellen Barnett, MD
2024 SVM Fellows/APP Course
University of Virginia
3/17/2024

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Traditional Lipid Biomarkers

Cholesterol: Normal range 120-200 mg/dL, Value 120
 HDL: Normal range above 40 mg/dL, Value 58
 Non-HDL Cholesterol: Normal range below 130 mg/dL, Value 71
 LDL CALCULATION: Normal range 0-130 mg/dL, Value 62

Triglyceride: Normal range 0-150 mg/dL, Value 45
 Cholesterol/HDL: Normal value 0.3-0.6, Value 2.2
 LDL CALCULATION: Normal range 0-130 mg/dL, Value 62
 LDL/HDL Ratio: Normal value 0.3-0.6, Value 1.1

Grundy et al. Circ 2019;139(6):e1143.
 Moonik et al. J Am Coll Cardiol 2013; 24(3):2237.
 Fungo et al. Endocrinol 2013; 153:2000.
 Daneshmandi. Endocr 2016; 44:470-483.

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2

Traditional Lipid Biomarkers

- LDL-c is primary lipid target

Formula for LDL-C calculation	r	Reference
$LDL-C = TC - HDL-C - TG/5$	0.954	Friedewald et al. (1972)
$LDL-C = 0.9TC - 0.9TG/5 - 28$	0.898	Anandaraja et al. (2005)
$LDL-C = TC - HDL-C - TG/6$	0.970	Puavilai et al. (2009)
$LDL-C = 90\% \text{Non-HDL-C} - 10\% \text{TG}$	0.824	Chen et al. (2010)
$LDL-C = TC - HDL-C - TG/3$	0.954	Vujovic et al. (2010)
$LDL-C = \frac{1}{4}(TC - HDL-C)$	0.785	de Cordova and de Cordova (2013)

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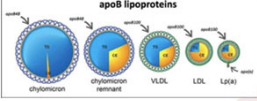
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Grundy et al. *Circ*. 2019;139:e1082-1143.
Macknik et al. *J Mol Sci*. 2020; 24(3):2237.
Feingold. *Endocrinol*. 2020;3:3000.
Dassanahally. *End* 2015; 14:478-483.

New Lipid Biomarkers

- **LDL size** - small, dense LDL is more proatherogenic than large LDL
- **Apolipoprotein B - apo B**
 - Likely a better predictor of ASCVD
 - Consider measuring if TG > 200
 - Level > 130 mg/dL corresponds to LDL-C > 160 mg/dL
 - Carries extra expense, not always reliable
- **Lipoprotein (a) - Lp(a)**
 - Consider if unexplained premature ASCVD, fam hx premature ASCVD
 - Lp(a) > 50 mg/dL or > 125 nmol/L is considered risk enhancing
- **LDL Particle Number**




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Grundy et al. *Circ*. 2019;139:e1082-1143.
Jones et al. *J Am Coll Cardiol*. 2012;80(4):1366-1418.

Statin Adjuncts



- Ezetimibe - 10 mg po daily
- MOA: inhibits NPC1L1 protein & reduces cholesterol absorption in small intestine
- Trials: IMPROVE-IT & SHARP


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Grundy et al. *Circ*. 2019;139:e1082-1143.
Jones et al. *J Am Coll Cardiol*. 2012;80(4):1366-1418.

Statin Adjuncts



- **PCSK9i**
 - Alirocumab - start 75 mg sq q14 days, can increase to 150 mg sq q14 days
 - Evolocumab - start 140 mg sq q14 days; can increase to 420 mg sq q14 days
- Trials: ODYSSEY, FOURIER

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Grundy et al. *Circ*. 2019;139:e1082-e1143.
Jones et al. *J Am Coll Cardiol*. 2022;80(4):1365-1413.

New Agents -

- Bempedoic Acid
- Inclisiran
- Evinacumab
- Lomitapide
- LDL apheresis




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Questions?






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Hypertension Management When the Basics Aren't Working

Ali Moran Baird, DNP, AGACNP
HCA Healthcare
Director of Cardiac and Vascular Service Line
Nashville, TN

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1



Hypertension

1 Billion people with high BP in low-income regions




Leading cause of Death Globally

Hypertension


10.4 million deaths per year

Associated with CV outcomes and mortality

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Hypertension Definition(s)

Table 6. Categories of BP in Adults*




BP Category	SBP	and	DBP
Normal	<120 mm Hg	and	<80 mm Hg
Elevated	120–129 mm Hg	and	<80 mm Hg
Hypertension			
Stage 1	130–139 mm Hg	or	80–89 mm Hg
Stage 2	≥140 mm Hg	or	≥90 mm Hg

Table 2. Criteria for Hypertension Based on Office-, Ambulatory (ABPM)-, and Home Blood Pressure (HBPM) Measurement

	SBP/DBP, mm Hg
Office BP	≥140 and/or ≥90
ABPM	
24-h average	≥130 and/or ≥80
Day time (or awake) average	≥135 and/or ≥85
Night time (or asleep) average	≥120 and/or ≥70
HBPM	≥135 and/or ≥85

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Accurate BP Measurement

Accurate BP Measurement

- Quiet room, comfortable temperature
- No smoking, coffee, exercise for 30min
- Empty bladder 30 min
- Rest for 5 min
- Take 3 measurements at 1-min intervals
- Use the average of the last 2 measurements

Arm Circumference	Usual Cuff Size
29-34 cm	Small adult
34-38 cm	Adult
38-42 cm	Large adult
42-46 cm	Adult thigh

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Work-Up of Hypertension

- Complete Medical, Family, Social, and Medication History
- Assessment of CV risk
- Signs and symptoms (blurred vision, palpitations, chest pain, edema)
- Secondary HTN symptoms (snoring at night, headaches, muscle weakness/muscle cramps/arrhythmias)
- Physical Examination
- Laboratory evaluation (CMP, lipid panel) & urinalysis
- ECG

Drug/Class	Comments on Specific Drug and Mechanism
Systemic anti-infective drugs (MAOIs)	No difference or an increase of up to 3/15 during and 2 weeks after treatment. MAOIs can antagonize the effects of beta-blockers and beta-receptors.
Combined oral contraceptive pill	6.3 mmHg increase with high doses of ethinyl estradiol (up to 50 mcg and 1-4 mcg progestin).
Antidepressants	SSRIs increase with 200 mg fluoxetine and venlafaxine (uptake inhibitor). Increased risk of up to 1/4 of hypertension with typical antidepressant use.
Antidiabetics	Increased risk of up to 3/4 of hypertension with insulin therapy.
Antiepileptics	Increased risk of up to 1/4 of hypertension with phenytoin, carbamazepine, and sodium valproate.
Other medications	Diuretics: Furosemide, bumetanide, torsemide. Beta-blockers: Propranolol, carvedilol, nebivolol. Calcium channel blockers: Amlodipine, nifedipine. Vasopressors: Norepinephrine, epinephrine. Sympathomimetics: Pseudoephedrine, cocaine, amphetamine. Stimulants: Methylphenidate, amphetamine, cocaine. Anesthetics: Propofol, sevoflurane, isoflurane, desflurane. Anticholinergics: Atropine, glycopyrronium, scopolamine. Opioids: Fentanyl, oxycodone, morphine, codeine. Alpha-1 antagonists: Alfentanil, remifentanyl.
Hypertensive emergencies	Diuretics, beta-blockers, calcium channel blockers, vasodilators, and ACE inhibitors.

5

Hypertension: White Coat or Masked?

Hypertension: White Coat or Masked?

Determine of White Coat or Masked Hypertension with patients not on drug therapy

- Office BP < 130/80 mm Hg and > 130/80 mm Hg after 3 mo trial of therapy modification and suspected white coat hypertension
- Office BP > 130/80 mm Hg and < 130/80 mm Hg after 3 mo trial of therapy modification and suspected masked hypertension

White Coat Hypertension: Office BP < 130/80 mm Hg and > 130/80 mm Hg after 3 mo trial of therapy modification and suspected white coat hypertension. Recommendation: Lifestyle modifications, Annual AFBM or HBEM, Home blood pressure monitoring (HBPM).

Masked Hypertension: Office BP > 130/80 mm Hg and < 130/80 mm Hg after 3 mo trial of therapy modification and suspected masked hypertension. Recommendation: Lifestyle modifications, Annual AFBM or HBEM, Home blood pressure monitoring (HBPM).

Elevated BP: Office BP > 130/80 mm Hg and > 130/80 mm Hg after 3 mo trial of therapy modification and suspected elevated BP. Recommendation: Lifestyle modifications, Annual AFBM or HBEM, Home blood pressure monitoring (HBPM).

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Causes of Hypertension

Environmental Risk Factors	Non Environmental Risk Factors
1. Sodium intake	1. Genetics
2. Sedentary lifestyle	
3. Alcohol	
4. Medications	

Screen for CVD Risk Factors

Recommendation for Coexistence of Hypertension and Related Chronic Conditions

References that support the recommendation are summarized in Tables 6 and 7 (Supplement 1)

Class	Class	Recommendation
1	B-III	1. Screening for and management of other modifiable CVD risk factors are recommended in adults with hypertension. ^{1,2,3,4,5,6,7,8}

Modifiable Risk Factors*	Relatively Fixed Risk Factors†
Current cigarette smoking, secondhand smoking	CKD
Diabetes mellitus	Family history
Dyslipidemia/hypercholesterolemia	Increased age
Overweight/obesity	Low socioeconomic/educational status
Physical inactivity/low fitness	Male sex
Unhealthy diet	Obstructive sleep apnea
	Psychosocial stress

ACC/AHA Guidelines for Prevention and Management of High Blood Pressure in Adults. 2017. In: ACC/AHA Guidelines for Prevention and Management of High Blood Pressure in Adults. 2017. © 2017 American Heart Association, Inc.

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Management of Hypertension: Lifestyle Modifications

Lifestyle Modifications

Salt reduction	There is strong evidence for a relationship between high salt intake and increased blood pressure. ¹⁻⁵ Reduce salt intake when preparing foods, and at the table. Avoid or limit consumption of high salt foods such as soy sauce, fish foods and processed food including breads and cereals high in salt.
Healthy diet	Eating a diet that is rich in whole grains, fruits, vegetables, polyunsaturated fats and dairy products and reducing total fat in eggs, including fat and trans fats, such as the DASH diet (http://www.healthline.com/health/diet-dash) is associated with lower blood pressure. ^{6,7} Salt in many vegetables and meats. Other beneficial foods and nutrients include those high in magnesium, calcium and potassium such as avocados, nuts, seeds, legumes and soy. ^{8,9}
Healthy drinks	Moderate consumption of coffee, green and black tea. ¹⁰ Other beverages that can be beneficial include kombu (bibulous tea), polyphenols juice, fermented juice and cocoa. ¹¹
Moderation of alcohol consumption	Positive linear association exists between alcohol consumption, blood pressure, the prevalence of hypertension, and CVD risk. ¹² The recommended daily limit for alcohol consumption is 2 standard drinks for men and 1 to 2 for women (12 g alcohol/standard drink, about 5 oz of wine). ¹³
Weight reduction	Body weight control is indicated to avoid obesity. Particularly abdominal obesity should be managed. Ethnic-specific cut-offs for BMI and waist circumference should be used. ¹⁴ Abnormally, a waist-to-height ratio <0.5 is recommended for all populations. ^{15,16}
Smoking cessation	Smoking is a major risk factor for CVD, CVDPS and cancer. Smoking cessation and referral to smoking cessation programs are advised. ¹⁷
Regular physical activity	Studies suggest that regular aerobic and resistance exercise may be beneficial for both the prevention and treatment of hypertension. ^{18,19} Moderate intensity aerobic exercise (walking, jogging, cycling, yoga, or swimming) for 30 minutes 5-7 days per week or 100 high intensity interval training which involves alternating short bursts of intense activity with subsequent recovery periods of lighter activity. Strength training also can help reduce blood pressure. Performance of resistance/strength exercises on 2-3 days per week.
Reduce stress and induce relaxation	Chronic stress has been associated to high blood pressure later in life. ²⁰ Although more research is needed to determine the effects of chronic stress on blood pressure, relaxation clinical trials exploring the effects of transcranial meditation/meditation on blood pressure suggest that this practice lowers blood pressure. ²¹ Stress should be reduced and mindfulness or meditation introduced into the daily routine.
Complementary, alternative or traditional medicines	Large proportions of hypertensive patients use complementary, alternative or traditional medicines (in regions such as Africa and traditional medicines). ²² Evidence from studies suggest a negative effect of air pollution on blood pressure in the long term. ²³

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Management of Hypertension: Pharmacological Treatment

Established Diagnosis of Hypertension

```

    graph TD
      A[Established Diagnosis of Hypertension] --> B[Grade 1  
SBP 140-159/DBP 90-109 mmHg]
      A --> C[Grade 2  
SBP ≥160/DBP ≥100 mmHg]
      B --> D[Likely advice]
      B --> E[Immediate drug treatment in high-risk patients or those with CVD, CKD, DM or HMOCD]
      C --> F[Immediate drug treatment in all patients]
      E --> G{Limited drug availability?}
      G -- Yes --> H[Target BP reduction by at least 20/10 mmHg, ideally to <140/90 mmHg]
      G -- No --> I[Drug treatment in line to moderate risk patients without CVD, CKD, DM or HMOCD after 2-4 months of lifestyle intervention. If BP still not controlled, drug treatment is indicated]
      I --> J[Target BP reduction by at least 20/10 mmHg, ideally to <140/90 mmHg]
      J --> K[Aim for BP control within 3 months]
      subgraph Legend
        L[RECOMMENDED]
        M[CONSIDER]
        N[STRONGLY CONSIDER]
      end
  
```

Target BP reduction by at least 20/10 mmHg, ideally to <140/90 mmHg

Aim for BP control within 3 months

Legend:
 RECOMMENDED (Green)
 CONSIDER (Blue)
 STRONGLY CONSIDER (Red)

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The Future of Hypertension: Renal Denervation

Catheter-based renal denervation - 9 year follow up data on safety and blood pressure reduction in patients with resistant hypertension

Participants
66 RDN trial participants
9 years follow-up

Research Questions
Is BP reduced long term?
Are there detrimental effects on renal function?

Key Findings
BP significantly reduced by -22.1/8.8 mmHg on ARB
No evidence of detrimental effects on eGFR

Conclusions: Blood Pressure is significantly reduced at nine year follow up after renal denervation without adverse renal consequences

Time (years)	Systolic BP (mmHg)	Diastolic BP (mmHg)
0	160	95
1	145	85
2	140	80
3	145	80
4	140	80
5	140	80
6	140	80
7	140	80
8	140	80
9	140	80

Renal denervation

- Renal nerves
- Renal artery
- Radiofrequency energy
- Kidney
- Catheter

Hypertension/ARJ Guidelines, 2023

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PAD Non-pharmacological Therapies: Supervised Exercise Therapy

Diane Treat-Jacobson, PhD, RN, FAHA, MSVM
Professor, School of Nursing
University of Minnesota

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1




Data related to access to SET in PAD are limited

- SET coverage by Medicare began in October of 2017
- Few programs existed prior to that coverage decision
- Objectives:
 - Describe the evidence we have about SET uptake thus far.
 - Discuss the Minnesota experience with developing SET for PAD in one health system
 - Discuss barriers and potential solutions

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Current evidence for SET referral

- 1,735 (1.3%) of 129,699 patients diagnosed with claudication were referred for SET in the first year of Medicare coverage (June 2017-Dec, 2018)
- Median # sessions attended 16; 5.1% Completed 36 sessions (required within 12 weeks)
- Those enrolled more likely to be older, white, male, and not dually enrolled in Medicare/Medicaid
- Majority from the Midwest (48.1%) and Northeast (16.1%). Those from the South and Middle Atlantic areas were proportionately under-enrolled.

Divakaran, et al, Circ Cardiovasc Qual Outcomes, 2021

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Structure of CMS-Reimbursed SET program

- Symptomatic PAD patients eligible
 - 36 sessions within 12 weeks
 - Patients can be referred for an additional 36 sessions if they continue to be symptomatic
 - 72 session lifetime cap
 - Asymptomatic patients (including post revascularization) are not eligible despite significant continued disability
 - Primary mode is treadmill walking but other modes are acceptable including total body recumbent stepping and aerobic arm or leg cycling

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SET for PAD in Minnesota

- SET for symptomatic PAD has been available in the Twin Cities within the MHealth Fairview health system since 2013
- Prior to the Medicare coverage determination, SET occurred through a hybrid phase III cardiac rehabilitation Wellness and Exercise for Life programs across 7 hospitals in one health system
- Since October 2017 SET has been implemented through a stand-alone Medicare reimbursed SET program
- Early adoption allowed for the development of infrastructure to quickly launch the CMS-reimbursed program in 2017
 - This included creating a SET referral order in the EHR
 - Training cardiac rehabilitation therapists how to implement SET
 - Increasing provider awareness of program availability

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Uptake and outcomes of supervised exercise therapy for peripheral artery disease: The importance of vascular medicine specialists at a large midwestern health care system during the first 5 years of CMS reimbursement

Mary O Whipple, Marsha A Burt, Aaron L Pergolski, Paige McArthur, Diane Treat-Jacobson, and Dereck L Salisbury;
VMJ, 2013

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Structure of MN SET program

- Intake evaluation completed to obtain a medical history, set exercise goals, do treadmill familiarization to ensure safety and set initial speed and grade of exercise
- Outcome measures are not required by CMS but are recommended to document effectiveness of programs
- Evaluation is performed at initial evaluation and after completion of 12 weeks of therapy
 - 6-minute walk test (6MWT)
 - Timed up and go (TUG test)
 - Vascular Quality of Life Questionnaire (VASCUQOL-6)
 - Treadmill metabolic equivalent of task (MET) achieved during exercise using grade and speed initially and after 12 weeks

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Outcomes for MHealth Fairview from 2017-2022

Whipple, et al, VMJ, 2023

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Characteristics

Patients who completed an intake visit (n=415)

<ul style="list-style-type: none"> • White - 88% • Male - 63% • Mean ABI - 0.67 • Prior revascularization - 24% • Ever smoker - 83% • Diabetes - 44% • Hypertension - 89% • Dyslipidemia - 86% • Coronary artery disease - 57% 	<ul style="list-style-type: none"> • Cardiology - 12% • Vascular medicine - 31% • Vascular Surgery/IR - 52% • Other - 6%
---	--

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Attendance

Supervised exercise therapy session attendance and adherence among all patients (N = 415) and those who completed the supervised exercise therapy (n = 207)

Characteristic	Overall (N = 415)			Completed SET (n=207)		
	Mean (SD)	Range	n (%)	Mean (SD)	Range	n (%)
No. of sessions completed	17.6 (11.3)	1-36		25.8 (7.4)	7-36	
Completed >= 24 sessions			136 (32.8)			125 (59.8)
Completed >= 30 sessions			87 (21.0)			82 (39.2)

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Changes in physical function outcomes following 12 weeks of supervised exercise therapy (n = 207)

Characteristic	Mean (SD)		t value	p
	Baseline	12 Weeks		
Claudication onset distance, m (n = 167)	118.1 (73.8)	174.4 (83.3)	9.70	< 0.001
Claudication onset time, sec (n = 75)	127 (68)	172 (68)	5.02	< 0.001
Total distance, m (n = 204)	298.6 (99.3)	352.1 (91.8)	11.97	< 0.001
TUG, sec (n = 171)	9.1 (2.8)	8.6 (2.6)	-4.69	< 0.001
Treadmill METs (n = 184)	2.7 (0.9)	3.9 (1.5)	17.16	< 0.001
VasculQoL-6 (n = 112)	14.0 (3.3)	16.9 (3.1)	9.46	< 0.001

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Takeaways from the MN experience

- Referral rates are quite low overall (14.5%), although higher than in the earlier national sample
- Enrollment of those referred was 56%
- Completion of those referred was 27%
- So, despite getting a 'head start', we still have much work to do.

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Barriers to access and maximization of SET

- **Provider factors:**
 - Lack of awareness of this therapeutic option and of available programs
 - Lack of ease of referral
 - Need to change practice patterns
- **Patient factors:**
 - Work conflicts
 - Distance
 - \$11.00 co-pay/session (\$396 for 12 weeks)
 - Lack of understanding of etiology of PAD, preventing activity avoidance due to fear of pain.
 - Lack of access in underserved rural and urban communities

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Barriers to access and maximization of SET

- **System factors- Opportunities to influence policy**
 - Slow development of new programs
 - Requirement to complete 36 sessions in 12 weeks – You do the math!
 - Exclusion of SET following revascularization despite robust evidence supporting benefit of combination therapy
 - Lack of mechanisms to track programs and referrals
 - No reimbursement for hybrid or community-based structured exercise programs

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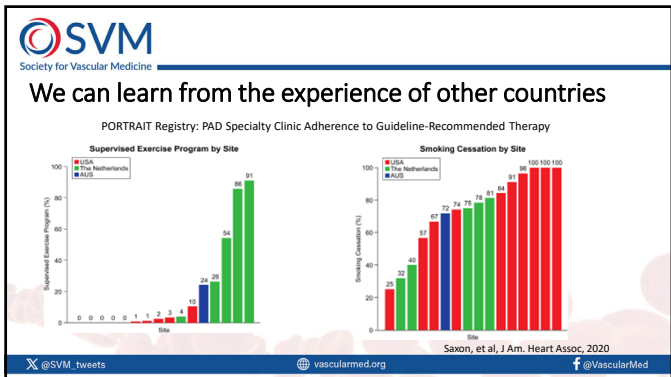


Addressing the barriers to access to SET programs

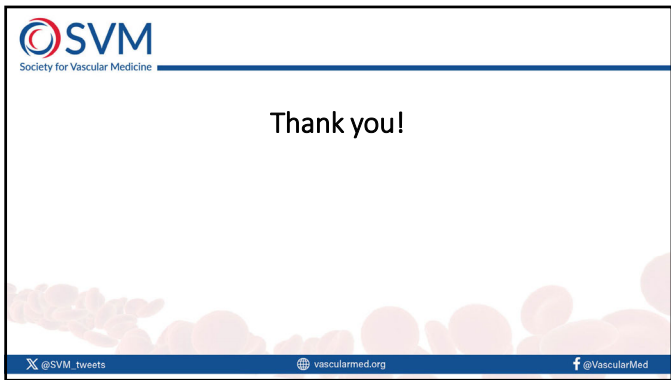
- Track and publicize available programs
- Improve awareness of program availability to providers and patients
- Adapt EHR to make referral to SET a therapeutic option – Make it easy for providers to prescribe
- Develop guidelines for hybrid models that combine some in-person sessions with remote coaching and monitoring using technology developed and refined over the past 2 years – Capitalize on our creative adaptation.
- Minimize the co-pay burden on our patients
- Model post-revascularization program options after cardiac rehabilitation programs.
- Get creative about how to offer SET in rural communities.

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17



Post-thrombotic and Post-Intervention Imaging

Eri Fukaya MD, PhD
Vascular Medicine, Division of Vascular Surgery
Stanford University School of Medicine

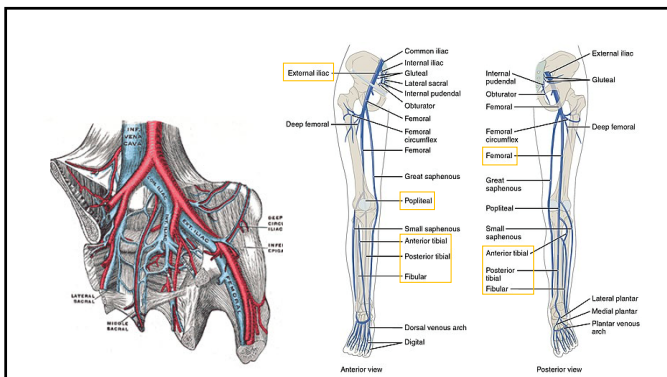
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1

Post Thrombotic Imaging

GOAL:
 Detection of the anatomical cause of post thrombotic syndrome symptoms
 Detect intravascular vs extravascular causes of PTS.
 Detect the flow irregularity causing venous hypertension (reflux, stenosis, obstruction)

2



3

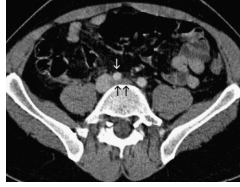
Case 1: 57F

CC: left leg swelling

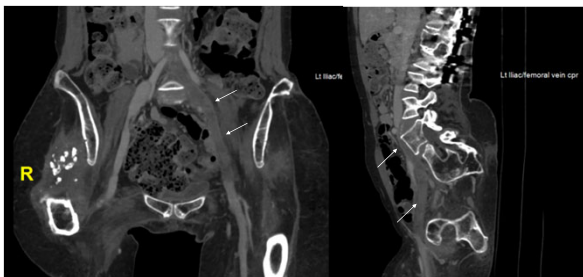
HPI: Developed leg swelling s/p 5 hrs sit ski.
Acute occlusive DVT in proximal left CIV, EIV, CFV and FV.
May Turner Syndrome

PMH: T11 Paraplegia due to MVA (1992)
Breast cancer (2014)- no recurrence

FH: Father DVTx3



4

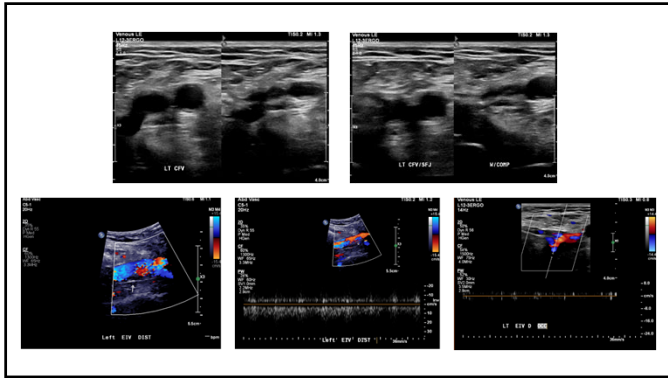


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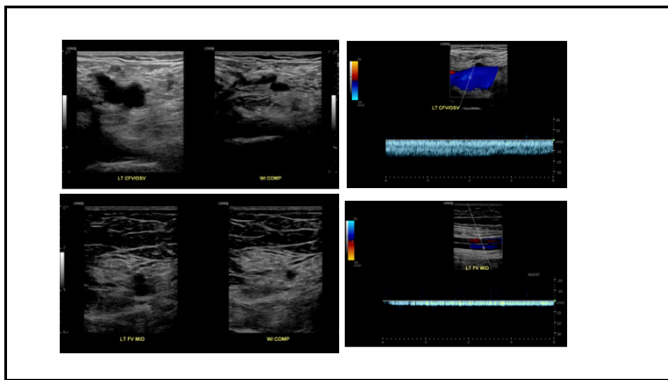
Case 1 (continued)

- Patient returns after 8 months.
- Completed anticoagulation with rivaroxaban 20mg daily.
- Swelling much improved.
- Wearing 30-40mmHg compression daily.
- *Chronic changes in CFV and EIV.*

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Case 2: 60M

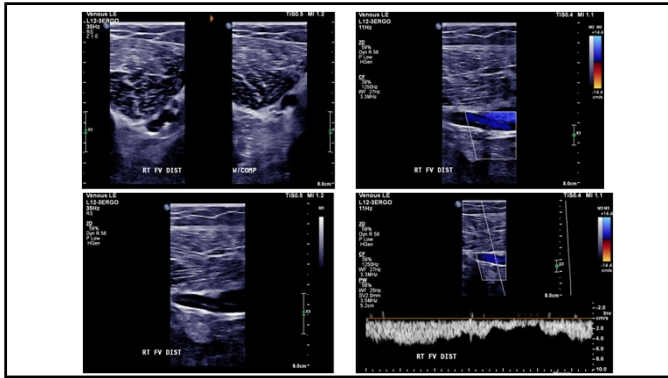
CC: consult regarding duration of anticoagulation therapy

HPI: 6/2018: Acute PE during Afghanistan convoy. Sitting in vehicle with heavy gear.
 Completed 3 months of anticoagulation with rivaroxaban.
 12/2019: Returned to Afghanistan. Developed right foot swelling.
Occlusive DVT in right POPV, PTV and peroneal V.
Non occlusive thrombus in right FV.
 Currently taking rivaroxaban 20mg daily

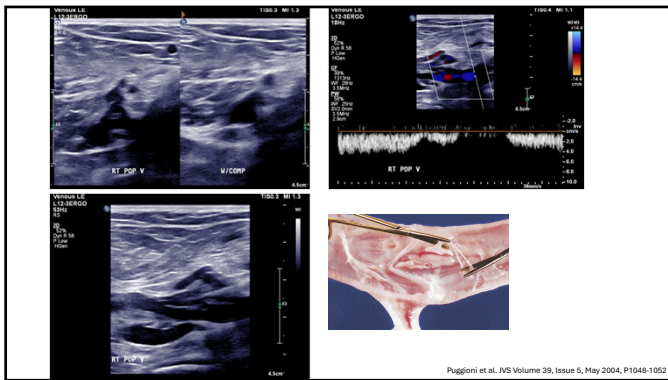
PMH: DVT, PE

FH: No VTE

9



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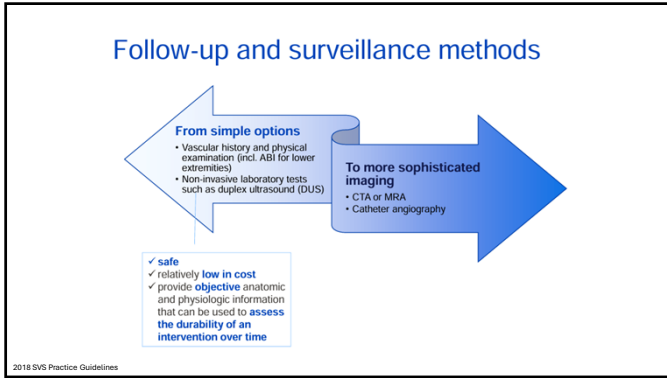


11

Post Intervention Imaging

GOAL:
Detection of recurrent disease and other complications (stenosis, occlusion).
Detect significant problems at an early stage when they can be managed most safely and effectively, even before clinical signs and symptoms are evident.

12



13

Post Intervention Imaging

Open surgical and endovascular arterial revascularization procedures:

- extracranial carotid artery
- thoracic and abdominal aorta → *Christine Owen, ACNP*
- mesenteric and renal arteries
- lower extremity artery

14

Modes of Failure

- neointimal hyperplasia
- distal embolization
- graft thrombosis
- anatomic stenosis
- progressive atherosclerosis

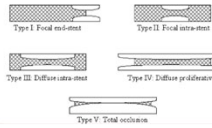
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Extracranial Carotid Artery

Recommendation	Strength of recommendation	Quality of evidence
1. After CEA or CAS, we recommend surveillance with DUS at baseline and every 6 months for 2 years and annually thereafter until stable (i.e., until no restenosis or ISR is observed in 2 consecutive annual scans). The first or baseline DUS should occur soon after the procedure, preferably within 3 months, with the goal of establishing a post-treatment baseline. Considering the small risk of delayed restenosis or ISR, some interval of regular surveillance (e.g. every 2 years) should be maintained for the life of the patient.	1 (Strong)	B (Moderate)
2. For patients undergoing CAS with diabetes, aggressive patterns of ISR (Type IV), prior treatment for ISR, prior cervical radiation, or heavy calcification, in addition to the baseline DUS we recommend surveillance with DUS every 6 months until a stable clinical pattern is established and annually thereafter.	1 (Strong)	B (Moderate)
3. We recommend that DUS after CAS include at least the following assessments: A. Doppler measurement of PSV and EDV in the native CCA in the proximal, mid, and distal stent, and in the distal native CCA. Modified threshold velocity criteria should be used to interpret the significance of these velocity measurements after CAS. B. B-mode imaging should be used to supplement and to enhance the accuracy of velocity criteria to estimate the severity of luminal narrowing.	1 (Strong)	C (Low)

Mode of failure:
Early: neointimal hyperplasia
Late: progressive atherosclerosis

Incidence:
CEA- 7.6% (3-18 months, >60%)
CAS- 6.4 % (5 year, >80%)



Morphological patterns of in-stent stenosis

2018 SVS Practice Guidelines

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Mesenteric and Renal Arteries

Recommendation	Strength of recommendation	Quality of evidence
1. There are no prospective reports documenting the efficacy of a surveillance protocol after mesenteric artery stenting or bypass grafts. However, recurrent mesenteric ischemia is potentially life-threatening. Therefore, after mesenteric artery (celiac, superior mesenteric, and inferior mesenteric) angioplasty with or without stenting, or mesenteric artery bypass grafting, we recommend the following: A. Clinical follow up and baseline DUS within 1 month of the procedure. B. Clinical follow up and DUS at 6 months, 12 months, and then annually thereafter.	1 (Strong)	C (Low)
2. We suggest contrast imaging for patients with symptoms of recurrent mesenteric ischemia after mesenteric artery stents or bypass grafts, or for the following duplex findings: A. Celiac axis: PSV >70 cm/s or a substantial increase from the post-treatment baseline PSV (what constitutes a substantial increase has not been defined). B. Superior mesenteric artery: PSV >40 cm/s, or a substantial increase from the post-treatment baseline PSV (what constitutes a substantial increase has not been defined). C. Inferior mesenteric artery: substantial increase from the post-treatment baseline PSV (what constitutes a substantial increase has not been defined).	2 (Weak)	C (Low)

Recommendation	Strength of recommendation	Quality of evidence
1. There are no prospective reports documenting the efficacy of a surveillance protocol after renal artery interventions. After renal artery angioplasty with or without stenting, or renal artery bypass or endarterectomy, we suggest the following: A. Clinical follow up and baseline DUS within 1 month of the procedure. B. Clinical follow up and DUS at 6 months, 12 months, and then annually thereafter.	2 (Weak)	C (Low)
2. We suggest contrast imaging for loss of renal parenchyma (a decrease in kidney length of >1 cm) or for the following duplex findings: A. Renal artery: PSV >280 cm/s, or a substantial increase from the post-treatment baseline PSV (what constitutes a substantial increase has not been defined). B. RAR of 14:5	2 (Weak)	B (Moderate)

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Lower Extremity Arteries (Open)

Recommendation	Strength of recommendation	Quality of evidence
1. We recommend clinical examination and ABI, with or without the addition of DUS, in the early postoperative period to provide a baseline for further follow-up after aorto-bi-femoral bypass. This evaluation should be repeated at 6 and 12 months and then annually as long as there are no new signs or symptoms.	1 (Strong)	C (Low)
2. We recommend clinical examination and ABI, with or without the addition of DUS, in the early postoperative period to provide a baseline for further follow-up after femoral-femoral bypass. This evaluation should be repeated at 6 and 12 months and then annually as long as there are no new signs or symptoms.	1 (Strong)	C (Low)
3. We recommend clinical examination and ABI, with or without the addition of DUS, in the early postoperative period to provide a baseline for further follow-up after femoral-femoral bypass. This evaluation should be repeated at 6 and 12 months and then annually as long as there are no new signs or symptoms.	1 (Strong)	C (Low)
4. We recommend clinical examination and ABI, with or without the addition of DUS, in the early postoperative period to provide a baseline for further follow-up after aorto-bi-femoral bypass. This evaluation should be repeated at 6 and 12 months and then annually as long as there are no new signs or symptoms.	1 (Strong)	C (Low)
5. Based on the high prevalence of abnormalities detected by DUS, as well as the relatively low associated cost and risk, we recommend clinical examination, ABI and DUS for infrainguinal vein graft surveillance. This should include an early postoperative baseline evaluation and follow up at 3, 6, and 12 months, and at least annually thereafter. More frequent surveillance may be considered when uncorrected abnormalities are identified on DUS or when alternative vein conduits (other than great saphenous vein) are used.	1 (Strong)	B (Moderate)
6. After prosthetic infrainguinal bypass grafts, we recommend clinical examination and ABI, with or without the addition of DUS, in the early postoperative period to provide a baseline for further follow-up. This evaluation should be repeated at 6 and 12 months, and then annually as long as there are no new signs or symptoms.	1 (Strong)	B (Moderate)

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Lower Extremity Arteries (Endovascular)

Recommendation	Strength of recommendation	Quality of evidence
1. We recommend clinical examination, ABI, and DUS within the first month following aorto-iliac segment EVT to provide a post-treatment baseline and evaluate for residual stenosis. Clinical examination and ABI, with or without the addition of DUS, should be performed at 6 and 12 months, and then annually as long as there are no new signs or symptoms.	1 (Strong)	C (Low)
2. We suggest clinical examination, ABI, and DUS within the first month following femoropopliteal artery EVT to provide a post-treatment baseline and evaluate for residual stenosis. Continued surveillance at 3 months and then every 6 months is indicated for the following: A. Patients with interventions utilizing stents, due to the potential increased difficulty of treating an occlusive versus stenotic lesion. B. Patients undergoing angioplasty or atherectomy for critical limb ischemia, due to increased risk of recurrent critical limb ischemia should the intervention fail.	2 (Weak)	C (Low)
3. We suggest clinical examination, ABI, and DUS within the first month following tibial artery EVT to provide a post-treatment baseline and evaluate for residual stenosis. Continued surveillance at 3 months and then every 6 months should be considered. These patients, with a deteriorating clinical vascular examination, return of rest pain, non-healing wounds, or new tissue loss, should undergo repeat DUS.	2 (Weak)	C (Low)

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Case 3: 78F

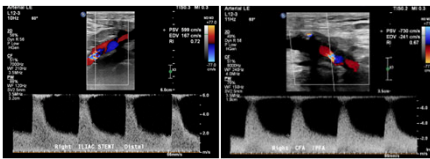
CC: Right foot gangrene

HPI: Polyvascular disease patient. DM, HTN, HL
6/2018: Right iliac stenting for peripheral artery disease
Lost to follow up fo 3 years.

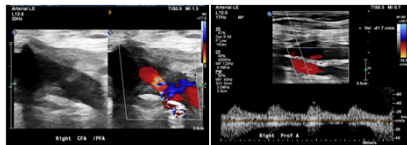
Presents to ER with right foot gangrene.

20

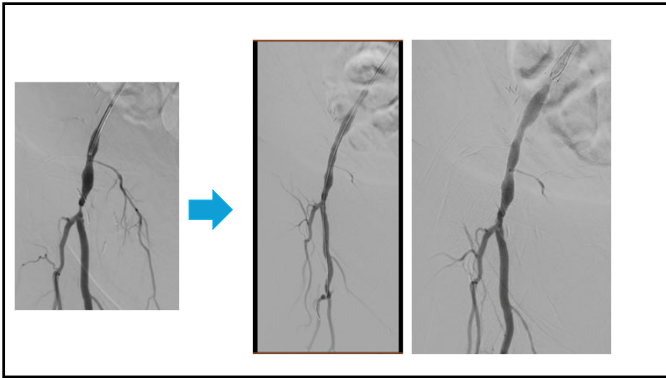
High grade in-stent restenosis of the RIGHT common and external iliac artery stents, common femoral endarterectomy site and proximal profunda.



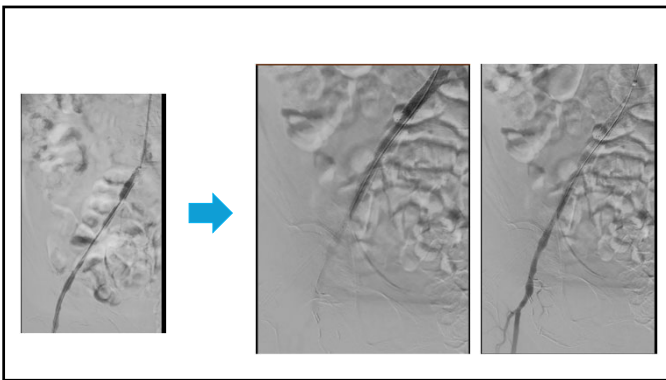
Distal iliac stent: 599cm/s within hyperplasia
PFA 730cm/s (prior 347cm/s)
PFA 42cm/s



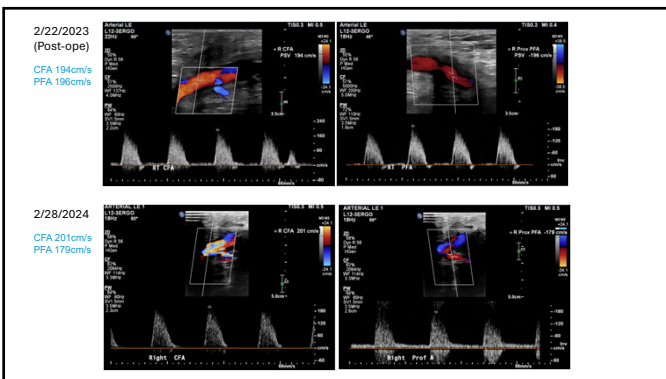
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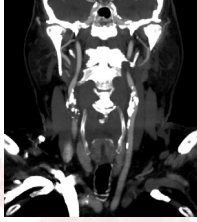


Thank you!



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Asymptomatic High Grade Carotid Lesion




SVM APP Course – Imaging & Vascular Cases
March 17, 2024

Deborah Hornacek, MD, FSVM
Cleveland Clinic, Section of Vascular Medicine

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Asymptomatic Carotid Stenosis


- **“Asymptomatic”**: adults without history of ischemic stroke, transient ischemic attack, or neurologic signs
- **Screening Guidelines** for the asymptomatic patient -or- “No, No, and Sometimes Yes”
- Carotid duplex ultrasound assessment
- Deciding revascularization -> What is a high grade lesion, high risk features

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All Clinical Recommendations

Choosing Wisely®

Screening for Carotid Artery Stenosis in Asymptomatic Adult Patients

Recommendation

Don't screen for carotid artery stenosis (CAS) in asymptomatic adult patients.

- There is good evidence that for adult patients with no symptoms of carotid artery stenosis, the harms of screening outweigh the benefits.
- Screening could lead to non-indicated surgeries that result in serious harms, including death, stroke and myocardial infarction.

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SVM Society for Vascular Medicine **U.S. Preventive Services TASK FORCE** 40 YEARS OF IMPROVING HEALTH

2021 USPSTF – "No"

Recommendation Summary

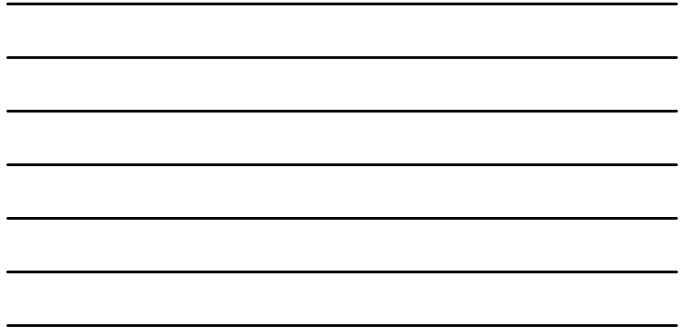
Population	Recommendation	Grade
Asymptomatic adults	The USPSTF recommends against screening for asymptomatic carotid artery stenosis in the general adult population. See the Practice Considerations section for a description of adults at increased risk.	D

"Although screening for asymptomatic carotid artery stenosis is not recommended for the general adult population, several factors increase risk for carotid artery stenosis, including older age, male sex, hypertension, smoking, hypercholesterolemia, diabetes, and heart disease. However, there are no externally validated, reliable methods to determine who is at increased risk for carotid artery stenosis or who is at increased risk of stroke when carotid artery stenosis is present."

...More on that later...

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ACC AHA guidelines classification scheme

- **Class:** benefit > risk > harm
- **Level:** strength of data available, RCT >> expert opinion

EVIDENCE OF CLINICAL BENEFIT OF TREATMENT EFFECT	SIZE OF TREATMENT EFFECT			
	CLASS I >>> Risk Reduction/Treatment Benefit is well established	CLASS IIa Additional studies with favorable outcome IT IS REASONABLE to perform preventive/treatment	CLASS IIb Additional studies with neutral outcome without harm IT IS REASONABLE to perform preventive/treatment MAY BE CONSIDERED	CLASS III An Inconclusive or Harmful Effect
LEVEL A Multiple randomised controlled trials or meta-analyses	Recommendation that treatment is strongly supported by multiple randomized trials or meta-analyses	Recommendation in favor of treatment or prevention being useful effective	Recommendation's usefulness/utility less well established	Recommendation that treatment is not recommended and may be harmful
LEVEL B Limited randomised evidence	Recommendation that treatment is supported by multiple randomized trials or meta-analyses	Recommendation in favor of treatment or prevention being useful effective	Recommendation's usefulness/utility less well established	Recommendation that treatment is not recommended and may be harmful
LEVEL C Very limited randomised evidence	Recommendation that treatment is supported by multiple randomized trials or meta-analyses	Recommendation in favor of treatment or prevention being useful effective	Recommendation's usefulness/utility less well established	Recommendation that treatment is not recommended and may be harmful
Support/Pharm for specific recommendations	Should be supported	Should be supported	May/should be considered	Should not be recommended
Considerable effectiveness proven?	Yes	Yes	Yes	No

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SVM Society for Vascular Medicine **2011 Circulation – "Sometimes Yes"**

- **Class IIa:** It is reasonable to perform duplex ultrasonography to detect hemodynamically significant carotid stenosis in asymptomatic patients with carotid bruit. (Level of Evidence: C)
- **Class IIb**
 - Consider screening asymptomatic patients with **symptomatic peripheral arterial disease (PAD), coronary artery disease, or atherosclerotic aortic aneurysm**, but because such patients already have an indication for medical therapy to prevent ischemic symptoms, it is unclear whether establishing the additional diagnosis of ECVD in those without carotid bruit would justify actions that affect clinical outcomes. (Level of Evidence: C)
 - Consider screening asymptomatic patients without clinical evidence of atherosclerosis **who have 2 or more of the following risk factors: hypertension, hyperlipidemia, tobacco smoking, a family history of ischemic stroke**. However, it is unclear whether establishing a diagnosis of ECVD would justify actions that affect clinical outcomes. (Level of Evidence: C)
- **Class III (No Benefit):** Carotid duplex ultrasonography is not recommended for routine screening of asymptomatic patients who have no clinical manifestations of or risk factors for atherosclerosis. (Level of Evidence: C)

2011 ASA/ACC/AHA/AANN/AANS/ACR/ASNR/ASNA/SNIR/SNIS/SVM/SVS Guideline on the Management of Patients With Extracranial Carotid and Vertebral Artery Disease: Executive Summary
Circulation. 2011;124:489-532

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SVM 2011 Circulation – “Sometimes Yes”
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- First line screening test: carotid duplex ultrasound (Class I, LOE: C)
- Note: auscultation for carotid bruit has poor sensitivity, not recommended as a screening exam (ie do not substitute for an ultrasound).

2011 ASA/ACCF/AHA/AANN/AAAS/ACR/ASNR/CNS/SAIP/SCAI/SIR/SNIS/SVM/SVS
Guideline on the Management of Patients With Extracranial Carotid and Vertebral Artery
Disease: Executive Summary
Circulation. 2011;124:489–532

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SVM Ultrasound Basics: 4 Characteristics
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- **Grayscale visualization of lesion:** does it look narrow? Can you see what is causing the flow disturbance?
- **Color aliasing:** loss of smooth laminar flow, creates speckling color that “wraps around” the color Doppler map
- **Velocity elevations:** going from an area of normal velocities to high
 - Velocity criteria: Society of Radiologists in Ultrasound (SRU) Consensus
- **Turbulent waveforms:** change from a normal Doppler waveform to abnormal “spiky” waveforms. Correlates with areas of elevated velocities.
- **Note on types of arterial waveforms:**
 - High resistive: little to no forward flow in diastole, normal for vessels supplying muscle tissue or the fasting gut
 - Low resistive: continuous flow through diastole, normal and essential for organs like the brain, kidneys, post-prandial gut

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SVM Determining Degree of Stenosis
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Ultrasound = Velocity Criteria

Degree of Stenosis, %	Primary Parameters		Additional Parameters	
	ICA PSV, cm/sec	Plaque Estimate, %*	ICA/CCA PSV Ratio	ICA/EDA, cm/sec
Normal	<180	None	<2.0	<40
1/3	180-200	<5%	2.0-4.0	40-100
2/3**	>200	>5%	>4.0	>100
>70 has less than near occlusion	High, low, or undetectable	Visible	Variable	Variable
Near occlusion	High, low, or undetectable	Visible	Variable	Variable
Total occlusion	Undetectable	Visible, no detectable lumen	Not applicable	Not applicable

*Plaque estimate (diameter reduction) is 0% proximal and 100% distal. †SRU Consensus.
**PSV >180 cm/sec and ICA/CCA PSV Ratio >2.0 also consistent with 2/3-40% stenosis.
Modified by updated criteria proposed by Society of Radiologists in Ultrasound (SRU) Consensus Conference.

CT angio = Diameter

	NASCET	ECST
30	65	
40	70	
50	75	
60	80	
70	85	
80	90	
90	97	

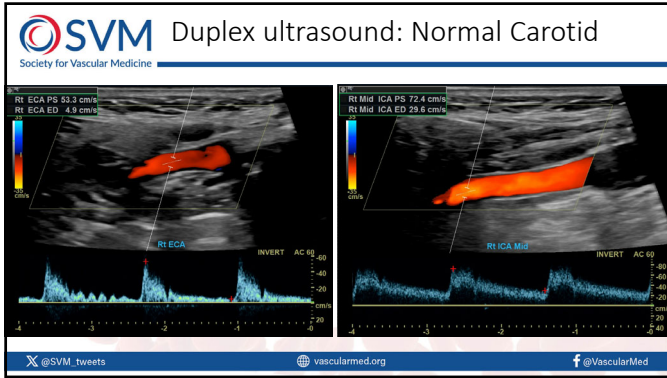
Approximate equivalent degrees of internal carotid artery stenosis used in NASCET and ECST according to recent direct comparisons

Common carotid artery: NASCET $\frac{A-B}{A}$, ECST $\frac{C-B}{C}$

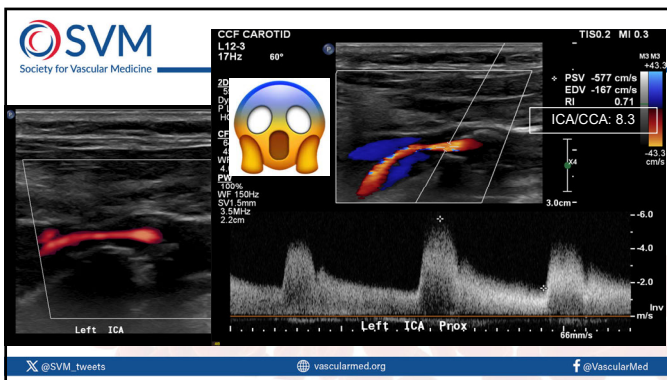
North American Symptomatic Carotid Endarterectomy Trial (NASCET) study group 1991

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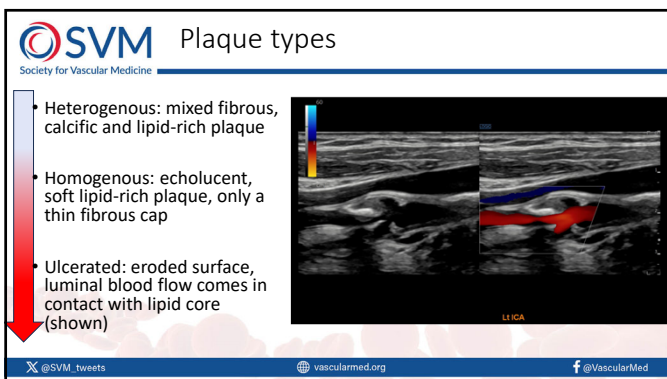
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12

SVM Case 1: "Mr. T"
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- 73yo M referred after routine eye doctor exam with right eye finding (shown).
- No change in visual acuity.
- PMHx: DM type 2 complicated by neuropathy and proteinuria, HTN, PAD, CKD 3b. Remote tobacco use.
- Normal exam. BP 126/78.
- A1c 7.2%, LDL 63 on atorvastatin 80mg, on aspirin 81mg and losartan 50mg.

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SVM Right CCA
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Rt Mid CCA PS 95.6 cm/s
Rt Mid CCA ED 12.8 cm/s

Rt Dist CCA PS 84.5 cm/s
Rt Dist CCA ED 12.9 cm/s

Long Rt CCA Distal Long Rt CCA Proximal

INVERT AC 60

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SVM Right ECA
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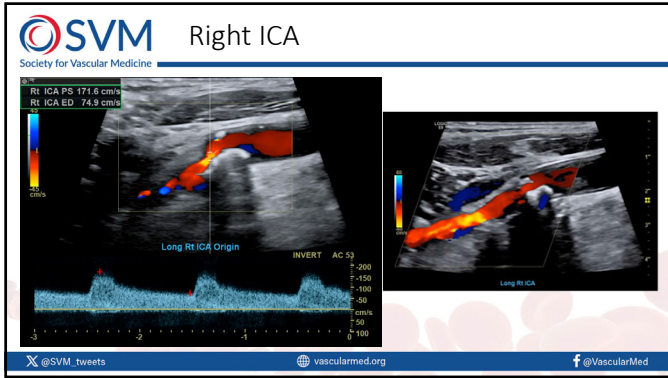
Rt ECA PS 127.9 cm/s
Rt ECA ED 21.5 cm/s

Long Rt ECA Origin

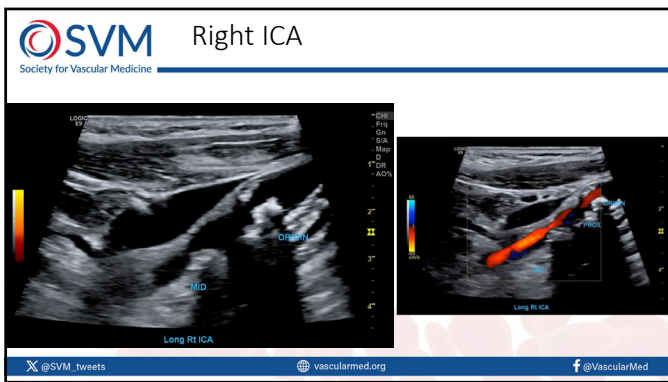
INVERT AC 60

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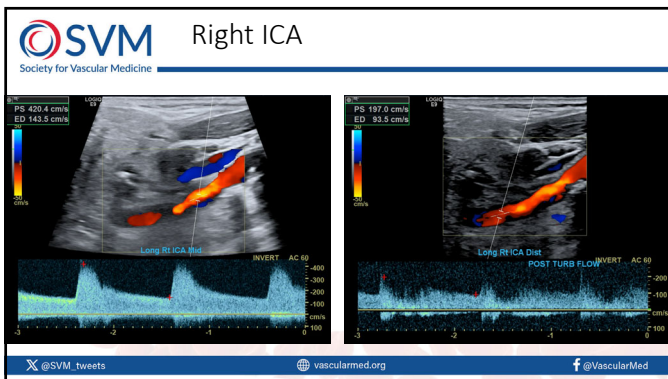
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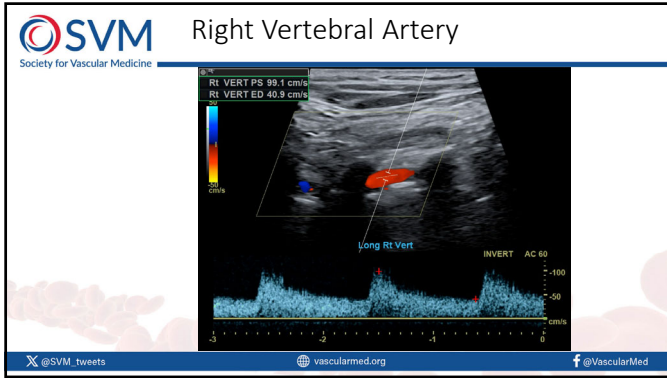
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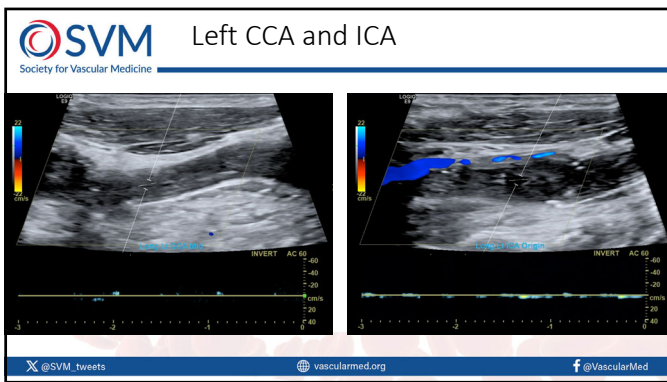
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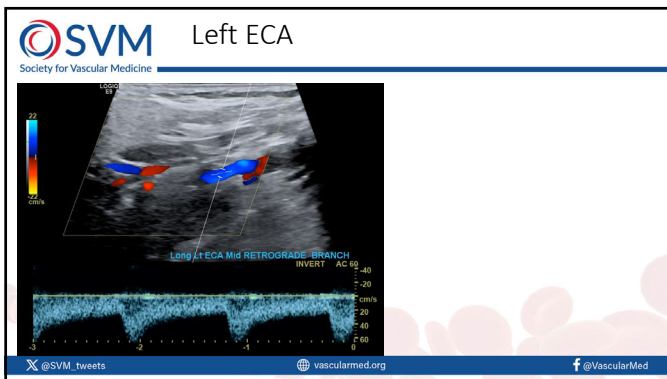
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SVM Left Vertebral Artery
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PS 191.1 cm/s
EDV 43.3 cm/s

PS 20.0 cm/s
EDV 6.1 cm/s

Long Lt Vert PROX

Long Lt Vert MID

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SVM Carotid Ultrasound: final report
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- RIGHT SIDE**
Common carotid artery:
Origin: PSV: 114 cm/s, EDV: 18 cm/s.
Proximal: PSV: 142 cm/s, EDV: 12 cm/s.
Mid: PSV: 96 cm/s, EDV: 11 cm/s.
Distal: PSV: 85 cm/s, EDV: 12 cm/s.
Moderate heterogeneous calcified plaque at dista
- Internal carotid artery:**
Origin: PSV: 172 cm/s, EDV: 75 cm/s.
Proximal: PSV: 119 cm/s, EDV: 57 cm/s.
Mid: PSV: 420 cm/s, EDV: 141 cm/s.
Distal: PSV: 281 cm/s, EDV: 128 cm/s.
Moderate heterogeneous irregular and calcified plaque from origin to proximal.
Severe heterogeneous calcified plaque at mid.
- ICA/CCA Ratio: 5.0
- External carotid artery: PSV: 128 cm/s, EDV: 21 cm/s.
- Subclavian artery: PSV: 128 cm/s, EDV: 6 cm/s.
Moderate heterogeneous calcified plaque from origin to proximal.
- Innominate artery: PSV: 132 cm/s, EDV: 0 cm/s.
Moderate heterogeneous calcified plaque at dista
- Vertebral artery: PSV: 99 cm/s, EDV: 41 cm/s.

- LEFT SIDE**
Common carotid artery:
Proximal: PSV: 0 cm/s, EDV: 0 cm/s.
Mid: PSV: 0 cm/s, EDV: 0 cm/s.
Distal: PSV: 0 cm/s, EDV: 0 cm/s.
Severe heterogeneous plaque from proximal to distal.
- Internal carotid artery:**
Origin: PSV: 0 cm/s, EDV: 0 cm/s.
Proximal: PSV: 0 cm/s, EDV: 0 cm/s.
Mid: PSV: 0 cm/s, EDV: 0 cm/s.
Distal: PSV: 0 cm/s, EDV: 0 cm/s.
Severe heterogeneous plaque throughout.
- External carotid artery:**
Origin: PSV: 0 cm/s, EDV: 0 cm/s.
Mid: PSV: 23 cm/s, EDV: 7 cm/s.
Severe heterogeneous plaque from origin to proximal.
- Subclavian artery:**
Proximal: PSV: 96 cm/s, EDV: 0 cm/s.
- Vertebral artery:**
Proximal: PSV: 197 cm/s, EDV: 44 cm/s.
Mid: PSV: 20 cm/s, EDV: 6 cm/s.

IMPRESSION: Provider notified with results.

- RIGHT SIDE**
Common carotid artery: Plaque visualized without evidence of hemodynamically significant stenosis.
Distal: PSV: 0 cm/s, EDV: 0 cm/s.
Severe heterogeneous plaque from proximal to distal.
- Internal carotid artery:** 80-99% stenosis.
Lesion from origin to proximal : 1.88 cm.
Tandem lesion at mid : 0.87 cm.
80-99% stenosis noted mid vessel.
- Vertebral artery:** Patent and antegrade flow noted.
- Innominate artery:** Plaque visualized without evidence of hemodynamically significant stenosis.
- Subclavian artery:** Plaque visualized without evidence of hemodynamically significant stenosis.

- LEFT SIDE**
Common carotid artery: Occluded.
Internal carotid artery: Occluded.
External carotid artery: Occluded at origin, reconstituted at mid. Reconstituted via retrograde branch.
Vertebral artery: Patent and antegrade flow, evidence of stenosis.

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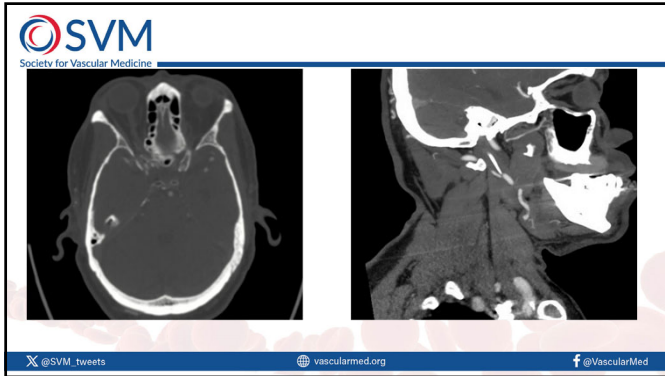
SVM CTA head and neck
Society for Vascular Medicine

Right Common: No significant stenosis.
Right Internal Carotid: Significant calcified and noncalcified atherosclerotic plaques at the bifurcation causing tandem severe stenosis at the bifurcation and just distal to the bifurcation.
Right Internal Carotid Stenosis (% by NASCET Criteria): 80%.

Left Common: Occluded just at the takeoff from the aortic arch.
Left Internal Carotid: Occluded.
Left Internal Carotid Stenosis (% by NASCET Criteria): 100%
Cervical Vertebral Arteries: Scattered calcified plaques along the course of the cervical right vertebral artery.
Patency: Bilateral
Dominance: Right. The left intracranial vertebral artery is markedly diminutive throughout its cervical course and intermittently poorly seen, most likely congenital/developmental.

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Revascularization considerations

Asymptomatic patients with risk factors and/or bruit on exam – consider carotid ultrasound

No or minimal plaque:
Continue best medical treatment.

High grade stenosis $\geq 70\%$
Particularly if high risk features are present:

- microemboli detection on transcranial Doppler
- plaque echolucency on Duplex ultrasound
- progression in the severity of disease
- silent embolic infarcts on brain CT/MRI
- reduced cerebrovascular reserve
- Ulcerated plaque, intra-plaque hemorrhage, or penetrating carotid ulceration.

Best medical treatment & referral for revascularization

Micro embolic signals on TCD.
Eur J Vasc Endovasc Surg 35, 534e540 (2008)

Paraskevas KI, Veith FJ, Spence JD. Stroke and Vascular Neurology 2018;3:e000129

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NASCET: carotid endarterectomy (1998)

Any Ipsilateral Stroke, $< 50\%$ Stenosis
P=0.16

Any Ipsilateral Stroke, 50-69% Stenosis
P=0.048

Any Ipsilateral Stroke, 70-99% Stenosis
P<0.001

Year of Study	Surgical Therapy	Medical Therapy
0	601	614
1	510	502
2	407	406
3	316	300
4	230	207
5	188	142
6	121	101
7	67	65


• Patients with severe $>70\%$ stenosis who underwent CEA demonstrated clear benefit over medical therapy alone.

• 30-day rate of death or disabling ipsilateral stroke at 90 days was 2.1%; this rate increased to only 6.7% at 8 years follow up.

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SVM Carotid Endarterectomy vs Stent
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CENTRAL ILLUSTRATION: Stenting Versus Endarterectomy for Carotid Artery Stenosis: Aggregated Efficacy/Safety Outcome

Study or Subgroup	Carotid Endarterectomy		Carotid Artery Stenting		Risk Ratio, M-H, Random, 95% CI
	Events	Total	Events	Total	
ACI 2006	48	1,060	17	340	12.4%
DECI 2014	303	1,624	37	342	22.9%
EVAS 2014	26	285	27	285	9.5%
CVS-2015	95	813	42	812	5.2%
SAPPAS 2008	16	167	21	167	12.6%
Total (95% CI)	298	3,859	134	3,850	100.0%

Total Events: 298 (CEA), 134 (CAS)
Heterogeneity: $I^2=0.04$, $DIP=1.84$, $IP=1.49$, $P=0.93$, $I^2=0.01$
Test for heterogeneity: $P=1.84$, $DIP=1.84$, $IP=1.49$
Test for publication bias: $P=0.81$

Sardar, P. et al. J Am Coll Cardiol. 2017;69(18):2266-75.

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SVM Carotid Endarterectomy vs Stent
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Carotid Artery Stenting (CAS) vs Carotid Endarterectomy (CEA)

Retrospective, single center study | 1853 CEA, 478 CAS

CEA Symptomatic: 29.1%	Outcomes	CAS Symptomatic: 32.2%	CONCLUSION
at 30 days			CEA and CAS can be performed with excellent outcomes by vascular surgeons at a large-volume hospital
Stroke: 1.1%	Stroke: 1.3%		
Acute MI: 2.2%	Acute MI: 1.7%		
Death: 0.7%	Death: 0.6%		
at 5.4 years			
Stroke: 6.8%	Stroke: 7.7%		
Acute MI: 22.7%	Acute MI: 21.0%		
Death: 28.4%	Death: 28.2%		

CONCLUSION
CEA and CAS can be performed with excellent outcomes by vascular surgeons at a large-volume hospital

JVS Journal of Vascular Surgery | Carvin et al. J Vasc Surg October 2018
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SVM TCAR = the new kid on the block
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Impact of Age on In-Hospital Outcomes After Transcarotid Artery Revascularization (TCAR), Transfemoral Carotid Artery Stenting (TFCAS) and Carotid Endarterectomy (CEA)

Retrospective non-randomized study using the Vascular Quality Initiative database

TCAR (N=3152)	CEA (N=61,650)	TFCAS (N=10,397)
vs:	No significant difference in outcomes except:	In patients ≥ 80 years, TCAR was associated with:
	• TCAR was associated with significant decrease in cranial nerve injury	72% Reduction in stroke risk
		65% Reduction in risk of stroke/death

JVS Journal of Vascular Surgery | Dakour-Arudi et al. J Vasc Surg September 2020
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SVM Patient factors which favor CEA vs CAS
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Table 2: Risk Factors to Consider When Choosing Procedures

High Risk for CEA	Comorbidities	High Risk for CAS	Plaque Characteristics
Surgical Anatomy Previous CEA or neck surgery Presence of tracheostomy Previous radiation Contralateral occlusion Laryngeal nerve palsy Lesion extending above C2 vertebra	Severe CHF Severe CAD Severe pulmonary disease Renal Failure	Vessel Anatomy Type II or III aortic arch Aortic arch disease Tortuosity of ICA or CCA Occlusive disease of access vessels	Lipid rich plaque Intraplaque haemorrhage Calcified plaque Thin fibrous cap Lesion located at a curve Extensive plaque

CAD = coronary artery disease, CAS = carotid artery stenting, CCA = common carotid artery, CEA = carotid endarterectomy, CHF = coronary heart failure, ICA = internal carotid artery.

Vascular & Endovascular Review 2019;2[1]:40-4

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SVM Mr. T: Post-CEA
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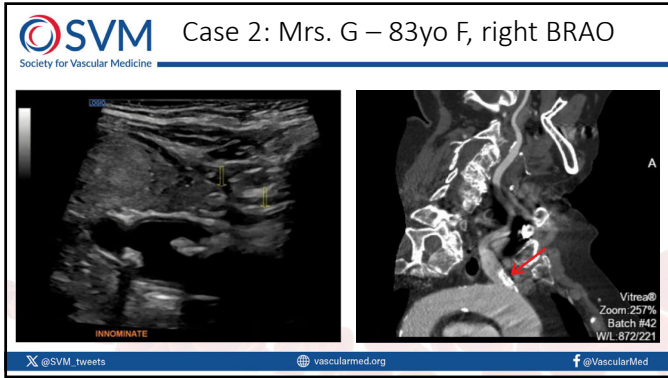
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SVM Case 2: Mrs. G – 83yo F, right BRAO
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Hollenhorst plaque.
Image from JVS. 2007;46(6):1125-1129

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Resistant Hypertension

Daniella Kadian-Dodov, MD
Associate Professor of Medicine
Program Director, Vascular Medicine Fellowship
Mount Sinai Fuster Heart Hospital

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Principles of Accurate BP Measurement

- Bare arm at heart level
- Relax several minutes
- Accurate sized ARM cuff
- If elevated, measure twice and average

Hypertension 2020;75:1334-1357

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Resistant Hypertension

Patients with HTN

21%

13% Blood pressure above goal ($\geq 140/90$ mm Hg) despite maximally tolerated 3 agents of differing mechanisms of action (including a diuretic)
OR patients requiring ≥ 4 agents regardless of level of control

10% may have *pseudoresistance*:
- Improper measurement, medication non-adherence, drug induced or secondary HTN

Hypertension 2018;72:e53-90
Hypertension 2020;75:1334-1357

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SVM The Long Road to Diagnosis
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- 1. Confirm Treatment Resistance**
 - Uncontrolled BP despite 3 agents (including diuretic), or on ≥ 4 agents regardless of control
- 2. Exclude Pseudoresistance**
 - Appropriate measurement (cuff size)
 - Medication non-adherence
 - White-Coat Effect
- 3. Identify and Reverse Contributing Lifestyle Factors**
 - Salt intake, alcohol
 - Drugs: ephedrine, NSAIDs, OCPs, etc
- 4. Screen for Secondary Causes of HTN**
 - OSA
 - Primary aldosteronism
 - Renal artery stenosis
 - Rare : pheochromocytoma, aortic coarctation, Cushing's, thyroid dysfunction

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SVM Renovascular Hypertension
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- Most common cause of secondary HTN (2 – 5%)
- Renal artery stenosis causing activation of the RAAS system
- Most often due to (1) Atherosclerosis (2) Fibromuscular Dysplasia



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SVM When to Consider Renovascular HTN
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- HTN onset < 30 years of age, or severe HTN > age 55 years
- Accelerated, resistant or malignant HTN
- Unexplained atrophy of the kidney or size discrepancy (> 1.5 cm difference)
- Unexplained renal dysfunction, including patients starting renal replacement therapy
- Multivessel coronary or peripheral artery disease
- Unexplained congestive heart failure or refractory angina
- Flash pulmonary edema
- Acute kidney failure with institution of ACEi/ARB
- Abdominal bruit

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SVM Society for Vascular Medicine **Guidelines for Renal Revascularization**

Clinically, consider if renal artery stenosis > 70% (by visual assessment) and one of the following:

1. Resistant or uncontrolled hypertension
2. Acute renal failure with the addition of an ACE inhibitor or ARB
3. Recurrent flash pulmonary edema or CHF with no other identifiable cause
4. CKD with no other identifiable cause and a viable kidney (normal kidney size)

Favorable characteristics for revascularization:

- Absence of proteinuria
- Younger age

Hypertension 2022;79:e128-e143

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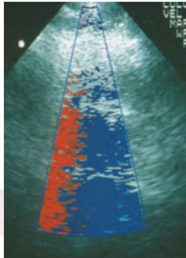
SVM Society for Vascular Medicine **Renal Artery Duplex**

Duplex Criteria	Stenosis
RAR < 3.5 and PSV < 200 cm/sec	0-59%
RAR ≥ 3.5 and PSV > 200 cm/sec	60-99%
RAR > 3.5 and EDV ≥ 150 cm/sec	80-99%
Absence of flow and low amplitude parenchymal signal	Occluded

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SVM Society for Vascular Medicine **Make Sure the Patient is NPO!**



If the patient images are like this:

- STOP
- Do not fight it
- You will get a technically suboptimal study
- This is especially true for hospitalized patients

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Anatomy – Anterior View

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Anatomy – Long View

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Inter-Costal Long View

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SVM Show the Entire Renal Artery
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SVM Make Sure You Get the Origin
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SVM How Our Reports Are Read
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- Normal Study for Atherosclerosis
 - Findings consistent with a 0- 59% stenosis. Patent without evidence for significant renal artery stenosis.
- More than 60% Stenosis
 - Findings consistent with a 60 - 99% stenosis within the proximal vessel.
- More than 80% stenosis
 - Findings consistent with a 60 - 99% stenosis within the proximal vessel. The end diastolic velocity > 150 cm/sec suggest stenosis is greater than 80%.

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SVM Case Summary
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73 year old female was transferred from an outside hospital to the CCU

- She was in acute pulmonary edema and on an FiO2 of 50%
- The blood pressure was 180/104 mmHg on:
 - Furosemide 120 mg BID
 - Metolazone 5 mg daily
 - Atenolol 100 mg daily
 - Hydralazine 100 mg TID
 - Clonidine patch 0.3 mg weekly
 - Isosorbide mononitrate 90 mg daily

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SVM Case Summary (continued)
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- Despite large doses of diuretics, her urine output over the last several days was 250 cc/24 hours
- The serum creatinine 4 weeks ago was 1.6 mg/dL and on transfer to the CCU it was 3.9 mg/dL.
 - She underwent hemodialysis and ultrafiltration
- Four weeks ago she had a nuclear stress test that was negative for ischemia and an echocardiogram with LVH, normal systolic LV function and diastolic dysfunction.

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SVM Renal Artery Duplex, Bilateral 60-99%
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RI Prox: RA PSV 123.3 cm/s
RI Prox: RA ED 56.8 cm/s

Juxtarenal Aorta PSV 80 cm/sec
Renal-Aortic Ratio = 5.1

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SVM Case Summary (continued)
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- Post stenting she required no dialysis
- The serum creatinine was 1.2 mg/dL four days after renal artery stent implantation
- The blood pressure was 130/70 mmHg on:
 - HCTZ 25 mg daily
 - Lisinopril 20 mg BID
 - Atenolol 100 mg daily
- She returned in 2 weeks for the first surveillance duplex ultrasound which was normal

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Fibromuscular Dysplasia

- Cannot use same criteria used for atherosclerosis
 - Velocity criteria have never been validated in FMD
 - Multiple sequential stenoses; the velocities are not focused on a single area
 - Affects the mid-distal artery
- We use:
 - There is turbulence/tortuosity/increased velocities in the mid and distal renal artery. This is consistent (or may be seen with) FMD.

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PG: 200.3 cm/s	ED: 154.2 cm/s
Arts. Sup	27
PRV	6.0
SV	46
AC	100
AD%	100
RRA MID	INVEST. AC 45
PRV	2.5
SV	6.5
SVF	1.2
VP	272
AD%	100
PRV	3.1
SV	31
SVF	17.1
VP	47
SVD	5.1
AD%	100

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Gradient 50 mm Hg

Gradient 52 mm Hg

Gradient 50 mm Hg

Gradient 47 mm Hg

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SVM Renal Artery Aneurysm
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RT RA DIST

PR	16
ERI	4.5
CFR	43
RI	8.8
AO%	100
ERI	2.5
CFR	16.0
RI	16
AO%	100
PR	3.2
CFR	40
RI	91.2
AO%	100

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THANK YOU!

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When Walking Isn't Enough

Intermittent Claudication

Sunday, March 17th, 2024
Danielle Vlazny, PA-C, MS, FSVM, CWS

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Case

- A 66 year old male presents with pain and heaviness in the left calf when walking. This is described as tightening and cramping that occurs at less than 1 block.
- Past medical history: T2DM, GERD, HLD, HTN, prostate cancer s/p resection and radiation. Tobacco use, currently 1 pack per day (58 year pack history).
- Medications: ASA 81 mg daily, amlodipine 5mg daily, atorvastatin 40mg daily, losartan 25mg daily, omeprazole 20mg daily, multivitamin

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What is claudication?

- Reproducible discomfort in the buttock, thigh, calf, or foot associated with exertion and resolves within 10 minutes of rest regardless of position. Often recurs at the same distance once walking is resumed.
- Often described in the number of blocks able to ambulate before requiring rest
- With atypical claudication, lower extremity discomfort may occur with walking but may not consistently occur at the same distance walked and may require a longer duration to resolve or require the patient to sit down or change body position to alleviate the symptoms.
- First line management is medical management with supervised exercise programs
- But what do we do when that fails?

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How do we classify PAD?

Rutherford Classification for Chronic Limb Ischemia

- Stage I
- Stage IIa
- Stage IIb
- Stage III
- Stage IV

Fontaine Classification	Clinical Presentation
I	Asymptomatic
IIa	Intermittent claudication after more than 200 meters of pain free walking
IIb	Intermittent claudication after less than 200 meters of walking
III	Ischemic rest pain
IV	Ischemic ulceration or gangrene

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Claudication Evaluation

- Unresolved claudication requires further evaluation to confirm arterial disease as the etiology and determine the level and degree of disease
- Non-invasive imaging is often completed first
 - Ankle Brachial Index and arterial ultrasound imaging
- Second line imaging further delineates disease location and severity
 - CT angiogram or conventional angiography

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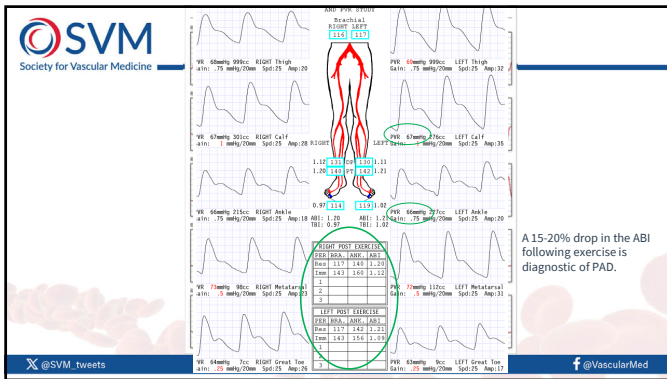
Ankle Brachial Index (ABI)

Measurement of the ankle systolic pressure against the brachial systolic pressure at rest

- ABI Values
 - > 1.00
 - 0.90 – 0.99
 - 0.80 – 0.89
 - 0.50 – 0.79
 - < 0.50
- Classification
 - Normal
 - Borderline
 - Mild
 - Moderate
 - Severe

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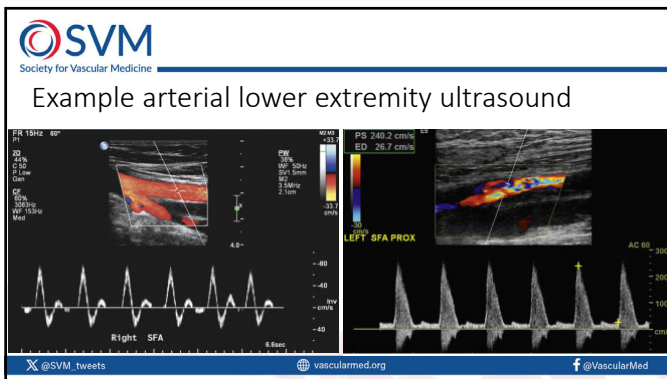
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Lower Extremity Arterial Ultrasound

University of Washington Duplex Criteria for Classification of Lower Extremity Arterial Stenosis

Disease Severity	Spectral Waveform Features	Artery	Peak Velocity \pm SD (cm/s)
Normal	Triphasic waveform No spectral broadening		
1%–19% diameter reduction	Triphasic waveform with minimal spectral broadening Peak reduction Proximal and distal waveforms remain normal	External iliac	119 \pm 22
20%–49% diameter reduction	Triphasic waveform usually maintained (although reverse flow filling in of the clear area under the systolic peak) Peak sys reduction Proximal and distal waveforms remain normal	Common femoral	114 \pm 25
50%–99% diameter reduction	Monophasic waveform with loss of the reverse flow component Spectral broadening Peak systolic velocity is increased >100% relative to normal reduced systolic velocity	Superficial femoral (proximal)	91 \pm 14
		Superficial femoral (distal)	94 \pm 14
		Popliteal	69 \pm 14
Occlusion	No flow is detected within the imaged arterial segment (collateral) waveforms are monophasic with reduced systolic velocities		

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Aortoiliac Ultrasound

- Investigates presence and degree of proximal disease
- Assess internal and external iliac arteries and abdominal aorta
- No pressures but doppler signals, velocities, and patency reported

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Example abdominal aorta/iliac artery ultrasound

Aorta Prox

FR 2MHz 44°
2D
23%
CF 25
P Low
Flow

+ PSV 130 cm/s
EDV 28.7 cm/s
RI 0.78

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Case Imaging – ABI

Doppler

Common femoral: Triphasic / Monophasic
Superficial femoral: Triphasic / Monophasic
Posterior tibial: Triphasic / Monophasic
Dorsalis pedis: Biphasic / Monophasic

Systolic Pressures

	Right		Left	
	Systolic	Index	Systolic	Index
Arm: brachial	136/66		112/56	
Arm Simultaneous	146/71		138/71	
Thigh (lower)	134	0.99	76	0.56
Calf	160	1.18	75	0.55
Ankle PT	157	1.15	65	0.48
Ankle DP	150	1.10	66	0.49
Distal T	111	0.82	57	0.42

Segmental BP

136 (R) / 112 (L) Brachial

134 (R) / 76 (L) Femoral

160 (R) / 75 (L) Popliteal

157 (R) / 65 (L) Tibial

150 (R) / 66 (L) Distal Peroneal

111 (R) / 57 (L) Distal Peroneal

Flowchart

Minutes	Right				Left				Arm (brachial)	
	Systolic	Index	Absent	Pallor	Systolic	Index	Absent	Pallor	Systolic	Diastolic
1	218	1.09			24	0.12		X	200	80
3	210	1.18			30	0.17			178	70
5	180	1.18			36	0.24			152	62
10	172	1.16			70	0.47			148	70

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Case Intervention

- Left femoral endarterectomy of the external iliac, femoral artery and superficial femoral artery with 10-cm bovine pericardial patch angioplasty
- Left common iliac artery stenting (Viabahn VBX 8 x 59 postdilated to 12 mm)
- Left external iliac artery stenting (8 x 60 Innova postdilated to 8 mm)

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Post-revascularization follow-up

- Follow-up with surgical team 3 months after procedure
- Repeat imaging with ABI and LE Duplex US

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Doppler Signal

	Right	Left
Common femoral	Triphasic	Triphasic
Superficial femoral	Triphasic	Triphasic
Popliteal	Triphasic	Triphasic
Posterior tibial	Triphasic	Biphasic
Dorsalis pedis	Biphasic	Biphasic
Post ex CF	Triphasic	Biphasic

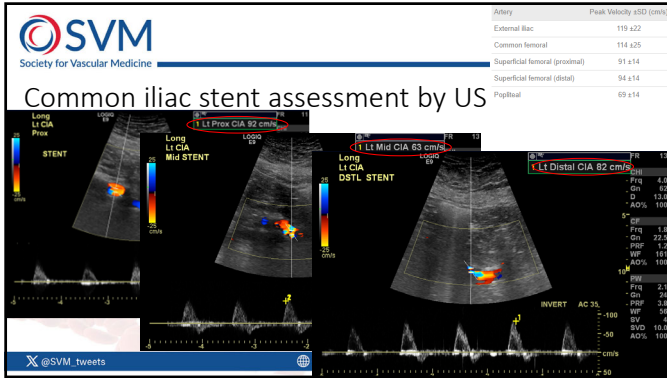
Systolic Pressures

	Right	Index	Left	Index
Arm: brachial	152/76		138/78	
Ankle PT	165	1.09	139	0.91
Ankle DP	175	1.15	142	0.93
Digit 1	136	0.89	131	0.86

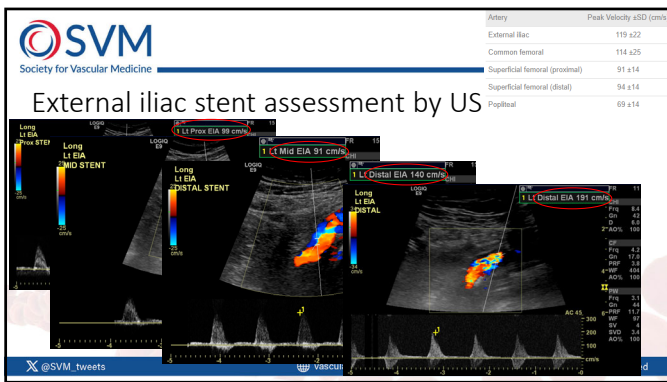
ABI

Minutes	Right Systolic	Right Index	Absent	Pallor	Left Systolic	Left Index	Absent	Pallor	Arm (brachial) Systolic	Diastolic
1	162	0.95			180	1.06			170	74
3	208	1.11			198	1.05			188	78
5	170	1.01			168	1.00			168	78

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- Patent stents in common and external iliac arteries
- Mildly elevated velocities at the distal end point of the external iliac artery stent (191 cm/s) may be due to vessel angulation or change in caliber
- Could be mild stenosis, but there does not appear to be significant luminal narrowing with color Doppler
- Remainder of stents are patent, without stenosis
- Common femoral, upper deep femoral, and superficial femoral arteries are patent, without evidence of stenosis
- Normal waveforms throughout

Aorta/Iliac Arteries

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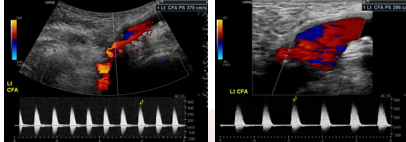
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Education Question

- A patient has been followed by the vascular clinic for yearly follow-up of known lower extremity PAD. This year the arterial ultrasound is different as shown below. What are your next steps?

- 1 - If worsening symptoms, continue with supervised exercise program
- 2 - If asymptomatic, refer to vascular surgery
- 3 - If worsening symptoms, refer to vascular surgery
- 4 - If asymptomatic, admit patient immediately



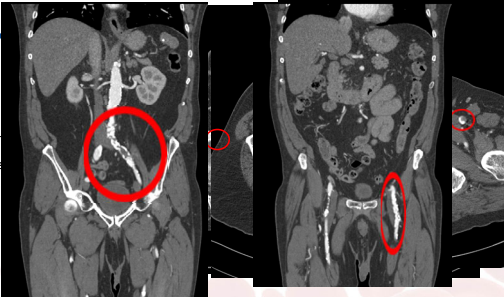
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Case Imaging

- Occlusion left common iliac arteries. Moderate stenosis right common iliac arteries. Two-vascular disease.



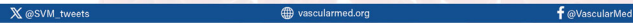
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AAA Pre and Post-EVAR

CT vs. US, post-EVAR, endoleaks
Chris Owen, MS, ACNP, RNFA
MedStar Heart and Vascular Institute
Annapolis, MD




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Disclosures

- I am not a diagnostic or interventional radiologist.
- I am Irish, **Sláinte!**


 Everyone's Irish
On March 17th.







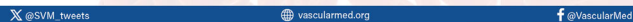
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Objectives

At the end of this presentation, you will have received tools to understand:

- Screening, risk factors and routine surveillance for AAA
- When and what treatment is appropriate for AAA
- Vascular lab studies associated with AAA
- Endovascular leaks post EVAR



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Screening, risk factors and routine surveillance for AAA

Screening	Risk Factors	Surveillance
Men & women age 65-75 h/o smoking	Age	3.0 - 3.9cm 3 - year interval US
Men & women w/family history	Sex	4.0 - 4.9cm annual US
US w/ aortic diameter measuring >2.5 cm but <3.0 cm again in 10 years	Race	5.0 - 5.4cm 6 - month US
	Family history	
	Smoking	

SOCIETY FOR VASCULAR SURGERY DOCUMENT
The Society for Vascular Surgery practice guidelines on the care of patients with an abdominal aortic aneurysm

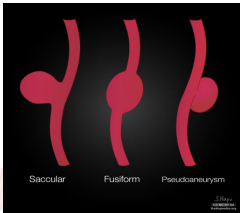
Elliot L. Chaouf, MD, PhD, Prakash L. Dalman, MD, Mark E. Eskandar, MD, Benjamin M. Jackson, MD, W. Anthony Lee, MD, M. Ashraf Mansour, MD, Tara M. Mastracci, MD, Matthew Mel, MD, M. Hassan Miral, MD, MPH, Ch. Nigam, MD, MBA, MPH, G. Gourevic, Odonir, MD, Braden S. Patel, MD, MBA, Scott A. Miller, MD, MPH, L. Schlemmer, MD, MPH, and Benjamin W. Starnes, MD, MPH, Miami, PA, Ala, Calif, Chicago, IL, Philadelphia, PA, Boca Raton, Fla, Grand Rapids, Mich, London, United Kingdom, Rochester, Minn, and Seattle, Wash.

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AAA



True Aneurysm
Saccular – sack-like, unilateral outpouching
Fusiform – circumferential dilatation (most common)

False Aneurysm
Pseudoaneurysm – hematoma that communicates with an artery but forms outside the arterial wall

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When and what treatment is appropriate for AAA

- Symptomatic AAA: abdominal/back pain or rupture
- Fusiform AAA >5.4 cm in otherwise healthy patient
- Saccular AAA repair recommended at smaller diameter
- Young, women w/AAA 5.0- 5.4cm
- Rapid expansion of small fusiform AAA
- Open
- Endovascular
 - 80% of all AAAs are treated by EVAR in USA

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Vascular lab studies associated with AAA

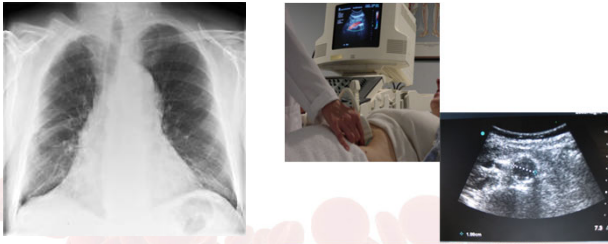
- **Non-invasive**
 - X-ray
 - Ultrasound
- **Invasive**
 - CT/CTA
 - MRI/A
 - Angiography
 - IVUS

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Chest X-Ray Ultrasound



The slide displays two medical images. On the left is a standard chest X-ray showing the lungs and heart silhouette. On the right is an ultrasound image showing a cross-section of a vessel with a bright, echogenic area, likely representing a calcified aortic aneurysm. The text 'Chest X-Ray' and 'Ultrasound' are placed above their respective images.

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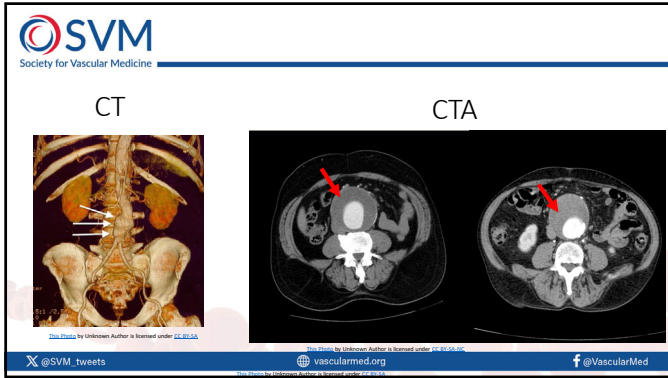
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Invasive Testing

- CT Scan or CTA are imaging studies utilizing non-contrast and/or IV contrast to identify specific anatomy.
- MRI/MRA create images of blood vessels using strong magnets that sends out radio waves to create the images with or without contrast.
- Angiogram/Arteriogram is an x-ray of the arteries using contrast to highlight the blood vessel.
- IVUS (Intravascular Ultrasound) sends high frequency sound waves into the blood vessel from a catheter inside the artery forming an image.

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EVAR: Endovascular Aneurysm Repair

- Minimally invasive procedure used to treat pathology of the aorta
 - Femoral cut down or percutaneous
 - Contrast, fluoroscopy
 - 24 hours stay
 - Follow up with US/CTA

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TYPES OF ENDOLEAKS

ENDOLEAK		
TYPE I Type I: gap between graft and vessel	1A (Proximal) 1B (Distal) 1C (Mid)	Contrast extravasation in continuity with the site of the graft attachment
TYPE II Type II: most common, flow through branch vessels into aneurysm sac	2A (Femoral) 2B (2 or more vessels)	Retrograde flow through branch vessels (lumbar arteries or inferior mesenteric artery)
TYPE III Type III: defect/misalignment of endograft materials	3A (Distal separation of the modular components) 3B (Fractures or holes involving the endograft)	Contrast extravasation central or distal to the graft attachment
TYPE IV Type IV: porous graft material	4 (Graft porosity)	Contrast extravasation anywhere of the aneurysmal sac without evidence of clear leak origins
TYPE V Type V ("endotension"): no evidence of leak; but continued enlargement of aneurysm	5 (Endotension)	Continued expansion of aneurysm sac without demonstrable leak on imaging

Case courtesy of Gerard Carbo, Radiopaedia.org, rID: 32126

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Case Study

79 yoF with finding of infrarenal fusiform AAA on screening, BLE popliteal arterial duplex negative for aneurysm

- PMH significant for:
 - HLD, HTN, CVA, CKD, Contact dermatitis, Sinusitis
- Meds:
 - ASA, Amlodipine, Atorvastatin, Levetiracetam, losartan, multivitamin
- Allergies: No known Drug Allergies
- Social Hx: previous smoker
- PSH: Colonoscopy x3, tubal ligation, hysterectomy, right knee meniscus tear, s/p EVAR 3/2022 known Type II endoleak

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Proximal Aorta: PDI 10 cm, 2.2 cm, 2.5 cm

Distal Aorta: PDI 14 cm, 2.7 cm, 2.9 cm

Common Iliac: +10%, PDI 12 cm

External Iliac: +10%, PDI 10 cm

1 L 4.49 cm
2 L 4.43 cm

AO MID

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Growth

Proximal Aorta: PDI 10 cm, 2.2 cm, 2.5 cm

Distal Aorta: PDI 14 cm, 2.7 cm, 2.9 cm

Common Iliac: +10%, PDI 12 cm

External Iliac: +10%, PDI 10 cm

1 Dist: Aorta AP 5.08 cm
2 Dist: Aorta AP 4.99 cm

AO DIST

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Video of initial CTA

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Post EVAR


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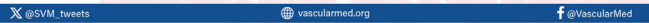
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Video of endoleak on CTA

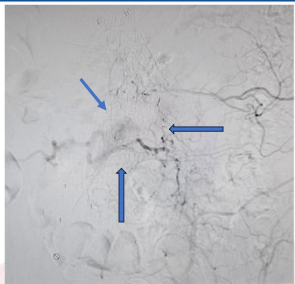



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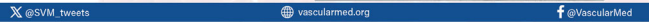
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Blushing on angiogram demonstrating leak







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Video of Angiogram & Intervention Attempt




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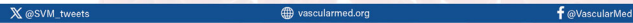
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Routine Surveillance Post

- CTA/US 1 month post EVAR
- 6 months thereafter with the most reliable and safe study
- Once stability and SAC shrinkage is demonstrated over time may move to annual surveillance




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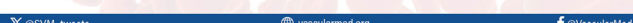
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Question

- After AAA has been identified what is the best way to follow and why?



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Question

- When we are following Post EVAR what are we looking for and why? When would we use US and when would we use CTA?



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Summary

- Identification** AAA
 - CTA/US
- Size or rate of growth**
 - EVAR vs. Open
- Routine surveillance** CTA/US
 - Goals – sack shrinkage

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Hartung M. How to read a CT of the abdomen and pelvis. Case study. Radiopaedia.org (Accessed on 05 Feb 2024)
<https://doi.org/10.53347/rID-66174> DOI: <https://doi.org/10.53347/rID-66174>

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