2023 MID-ATLANTIC CONFERENCE 11th ANNUAL CURRENT CONCEPTS IN VASCULAR THERAPIES



Hilton Virginia Beach Oceanfront Virginia Beach, Virginia





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I was diagnosed with a DVT

What NOW?

Justin Milligan MD FACS RPVI Sentara Vascular Specialists

No Disclosures



Goals of Presentation

Introduction to DVTDiagnosis of DVTManagement of DVT

Goals of Presentation

Introduction to DVT

Diagnosis of DVTManagement of DVT

DEEP VENOUS THROMBOSIS (DVT)



- There are over 200,000 new cases of DVT each year in the U.S.
- The incidence of PE in patients with DVT ranges from 5 20% and can be fatal
- After an episode of DVT, 20 50% of patients develop <u>Post Thrombotic</u> <u>Syndrome</u> within the first 2 years



DVT Pathophysiology



Virchow triad:

Hypercoagulability Stasis Intimal injury

DVT Pathophysiology

Balance



Coagulation, Fibrinolysis

When balance is tipped to coagulation...

RT CFV normal blood flow

Thrombosis



RT CFV thrombosis



Impaired fibrinolysis



Venous thrombosis

Common Etiologies:

- Injury / trauma
- Surgery
- Cancer
- Prolonged inactivity
- Bed rest post-op or post-partum
- Extended plane or car travel (longer than 4 hrs)
- Contraceptives/estrogen therapy
- Severe infection
- Undiagnosed anticoagulation disorder

Venous thrombosis

Symptoms:

- Sudden swelling in affected limb
- Limb pain/tenderness
- Pain on dorsiflexion of the foot
- Dilated superficial collateral veins
- Cyanosis or pallor
- Warm skin over area of thrombosis
- Lack of distal pulses



When to consider surgical options







Extreme presentation

Phlegmasia alba dolens

Phlegmasia cerulea dolens

(One precedes the other in 50-60% of cases.)

- Phlegmasia alba thrombosis involves major deep venous channels but spares the collateral veins
 - Venous drainage is significantly decreased but still present.
- Phlegmasia cerulea thrombosis extends to collateral veins resulting in:
 - Severe venous congestion, massive fluid sequestration and edema
 - Compromise of arterial circulation due to shock, increase venous outflow resistance and collapse arterioles due to increased interstitial pressure.



Goals of Presentation

Introduction to DVT Diagnosis of DVT

Management of DVT

DVT Probability Score/Wells Score

Clinical Characteristic	Score
Active cancer (patient receiving treatment for cancer within the previous 6 months or currently receiving palliative treatment)	1
Paralysis, paresis, or recent plaster immobilization of the lower extremities	1
Recently bedridden for 3 or more days or major surgery within the previous 12 weeks requiring general or regional anesthesia	1
Localized tenderness along the distribution of the deep venous system	1
Entire leg swollen	1
Calf swelling at least 3 cm larger than on the asymptomatic side (measured 10 cm below the tibial tuberosity)	1
Pitting edema confined to the symptomatic leg	1
Collateral superficial veins (nonvaricose)	1
Previously documented deep venous thrombosis	1
Alternative diagnosis at least as likely as deep venous thrombosis	<mark>-</mark> 2

Note: A score of 2 or higher indicates that the probability of deep venous thrombosis is high; a score of less than 2 indicates that the probability of deep venous thrombosis is low. In patients with symptoms in both legs, the more symptomatic leg is used.

Diagnosis

- Patient risk factors/medical history.
- Physical exam
 - Specific limb symptoms (edema, pallor)
- Clinical predictive scores
- D-dimer: used to exclude PE in patients with low clinical suspicion (99% NPV)
- Duplex ultrasound dominant diagnostic test.



Diagnostic Duplex Criteria DVT

- Major:
 - Venous incompressibility
 - Thrombus visualization (less echogenic)
 - Absent or diminished spontaneous venous flow
 - Absence of respiratory variation
 - Absent or incomplete color filling of lumen
 - Increased venous diameter, Immobile venous valves



Diagnosis cont.

- <u>Venography</u> "the gold standard" for diagnosis of DVT
- <u>Computer Tomographic Venography</u>—has not been studied in calf DVT and has some false positives
- <u>Magnetic Resonance Venography</u> may be considered with anatomic limitations in imaging inferior vena cava and iliac vessels

Goals of Presentation

- Introduction to DVTDiagnosis of DVT
- Management of DVT

DVT: Treatment Options

- Observation/Compression
- High-dose Heparin Therapy
- Surgical Treatment
- Thrombolytic Therapy
- Mechanical Thrombectomy
- Pharmacomechanical thrombectomy

DVT: Treatment Options

Duration of Clinical Symptoms:

- Acute <14 days
- Sub-acute >14 days
- Chronic Changes Months to Years
- Acute on chronic

Expected technical success is highest in patients with acute DVT (<14 days)



DVT: Treatment Options

Anticoagulation is Standard of Care

(heparin, warfarin, LMWH)

Advantages:

- Reduces occurrence of a PE
- Can decrease symptoms
- Easy to administer

Disadvantages:

- Bleeding complications from long-term use of anticoagulants
- Does not decrease thrombus burden
- Has not been shown to reduce incidence of valve damage



Anticoagulation for DVT



- Only one controlled trial performed
- 1960 Barritt and Jordan
- Anticoagulation vs no anticoagulation for acute PE
- Heparin 1.5 days/ nicoumalone 14 days

14 Day Follow-up

- Treated
 - 16 patients
 - No recurrent PEs
 - No fatal PEs
 - One death (bleeding)

- Untreated
 - 19 patients
 - 10 recurrent PEs
 - 5 fatal PEs
 - 5 deaths

Historical anticoagulation options (pre-2010)

<u>Oral</u>

• Warfarin

Intravenous/subcutaneous

- Heparin
- Fondaparinux
- Argatroban
- Lepirudin



Direct Factor Targeting



reed.

Factor Xa inhibitors: Rivaroxaban

- AKA XARELTO
- High bioavailability (80%)
- Rapid onset 2-4 h
- Half life 5-9h
- 66% renal excretion
- Also hepatic and fecal elimination
- Substrate for P-glycoprotein transporters
 - Contraindicated for patients taking HIV protease inhibitors and azole drugs
- May be taken with food

Factor Xa inhibitors: Apixaban

- AKA ELIQUIS
- Peak plasma concentration: 3h
- Half life: 8-11h (once daily) and 12-15h (twice daily)
- Main metabolism involves CYP3A4
- Apixaban presence can be determined by
 - Drug specific Factor Xa assays
 - Mild prolongation of PT and aPTT

When to admit to hospital





When to (







al options



Surgical Treatment Options

Endovascular Option

Mechanical Thrombectomy for the removal of venous thrombus

Mechanical Thrombectomy Advantages

- On-label
- Quick reperfusion
- Removal of thrombus vs. dissolving
- Can be used with other treatment strategies

Potential risks:

- Renal insufficiency/failure due to hemolysis
- Embolization including PE

Thrombolysis for DVT

Indications

- Iliofemoral location
- Symptoms <14 days
- Good functional status
- >1 year life expectancy
- Low risk of bleeding

Goals of treatment

- Prevent post-thrombotic syndrome
- Prevent recurrence
- Prevent death from PE

Thrombolysis

- Non-operative
- Identification of underlying stenoses
- May allow for more directed operation
- Clears clot in smaller, non-accessible branch vessels
- Avoid intimal injury associated with balloons or thrombectomy catheters
- Preserves endothelial function and valve competence
- Distal valve incompetence develops with anticoagulation alone
- Clot resolution within 90 days usually preserves valve function
- Disadvantages:
 - Potential for bleeding complications
 - Embolization risks
 - Higher doses and longer infusions increase risk of potential complications



- Removal of distal thrombus mechanically with Esmark compression or massage
- AVF between GSV branch and SFA



• Consider fasciotomy in phlegmasia cases





Figure 1. Preprocedure venograms demonstrate thrombus in the left popliteal, femoral, common femoral, external iliac, and common iliac segments (A, B). No contrast flowed into the IVC (C). Extracted thrombus (D). Repeat venogram demonstrates patent popliteal, femoral, common femoral, and iliac vein segments (E, F). Completion venogram confirms good flow into the IVC and resolved stenosis (G).











ATTRACT Trial

- Addition of catheter-based intervention to anticoagulation <u>failed to</u> reduce occurrence of post-thrombotic syndrome compared to anticoagulation alone
- Above-knee DVT evaluated both <u>ileo-femoral and fem-pop DVT</u>
- <u>Not</u> sufficiently powered to eval ileo-femoral DVT group
- Catheter-based therapy group had significantly higher major bleeding within 10 days
- The CaVenT trial had shown reduction (ARR 28%) in PTs with catheterbased interventions but no improvement in quality of life

National Venous Registry

- Registry comparing patients treated with thrombolysis to cohort of anticoagulation alone
- Quality of life evaluated at 16 and 22 months
- Significant improvement in lysis group vs anticoagulation



Duration of Anticoagulation Dur **for DVT or PE*** DVT)

•	Reve	Event	Duration	Strength of Recommendation
•	– Unp	First Time event of Reversible cause (surgery/trauma)	At least 3 mos	A til cancer resolved
	-	First episode of idiopathic VTE	At least 6 mos	A 🥂
•	– Secc –	Recurrent idiopathic VTE or continuing risk factor (e.g., thrombophilia, cancer)	At least 12 mos	B
		Symptomatic isolated calf-vein thrombosis	6 to 12 weeks	A
	HIST AND			

*From American College of Chest Physicians

Sequelae of Iliofemoral DVT

- Anticoagulation therapy alone
 - 90% ambulatory venous hypertension
 - 40% venous claudication
 - Up to 15% ulcer development at 5 years

Sequelae of Iliofemoral DVT

- Post-thrombotic syndrome may result in:
 - Chronic pain
 - Swelling
 - Skin ulceration secondary to postphlebitic syndrome
- Chronic condition in 30-75% of DVT patients within 2 years
 - 90% unable to work due to leg symptoms 10 years after iliofemoral DVT
- Irreversible damage to veins & valves
 - Impact on quality of life

Postthrombotic syndrome



Postthrombotic pigmentation



Healed skin ulcer and postthrombotic pigmentation



Chronic (left) leg swelling, skin hardening, and postthrombotic pigmentation

Sequelae of Iliofemoral DVT

Post-Thrombotic Syndrome

- PTS develops in 29% to 74% of patients following DVT
- Affects 5% of US population
- 400,000 to 500,000 individuals have venous stasis ulcers
- Annual direct cost of PTS in the US of \$200 million
- 2 million workdays lost annually in the US



Conclusions

A lot of DVT Patients need our help...



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Thank You!

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