

# UPMC Hamot Heart and Vascular Institute

**Stop the Clot! – Making Sense of the Thrombosis  
Clinic Model and Protocols  
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## What is Vascular Medicine?

- Niche specialty with emphasis is on clinical approaches to vascular disorders by physicians with special expertise and training in treating vascular disease
- Includes the non-invasive treatment of medical issues involving the circulatory system outside the heart including arterial, venous, and lymphatic disorders
- Entails a collegial interaction with a community of vascular professionals including Vascular Surgery, IC, IR, Vascular Ultrasound, Primary Care Physicians and other disciplines

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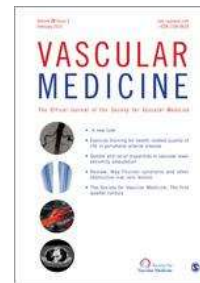
## What is Vascular Medicine?

- Increased awareness last 20 years, ? Vascular Renaissance?
- Lead Public and Professional Educational Programs through the Society for Vascular Medicine
- Pioneer bench to bedside medical advances
- Educate health professionals about Vascular Medicine
- Team Based Vascular Care
- History of the Field/History of Interest

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## What is Vascular Medicine?

- **Society of Vascular Medicine**



- **American Board of Vascular Medicine**



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## What Makes Vascular Medicine Unique?



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## We Have A Unique Skill Set!



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## So What Do We Do?

- The goal of the Vascular Medicine specialist is to improve the care of the patient with undiagnosed or complicated vascular disease
- Non-operative specialty, not a lot of us out there
- We specifically try to manage vascular disease “comprehensively”, including before and after interventions

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## So What Do We Do?

- **Medical Treatment of Vascular Disorders**
- **Follow Aneurysms and Stenosis**
- **Follow patients with unusual vascular disorders**
- **Focus on Primary/Secondary Prevention**
- **Optimize and prepare patients for vascular intervention procedures**

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## So What Do We Do?

- **Vasculitis and CTD**
- **Venous Thromboembolism and Thrombophilias**
- **Upper and Lower Extremity Venous and Arterial Disease**
- **Perioperative Management of Vascular Surgery**
- **Arterial and Venous Testing in the Vascular Lab**
- **Wound Care**
- **Atherosclerosis –Early Detection, Standardized Therapies, Surveillance and Outcomes**

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## Common Vascular Medicine Consultations

- Carotid Artery Disease
- Peripheral Artery Disease
- Aneurysms
- The Swollen Limb
- VTE and Chronic Venous Insufficiency
- Thrombophilias
- Risk Factor Modification in Vascular Patients
- Diagnostic Testing Abnormalities
- Unusual Vascular Disorders

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## Unusual Vascular Disorders

- Hypercoagulable States/Thrombophilias
- Thermal Disorders including Frostbite, Pernio, Cryoglobulinemia, Raynauds, Erythromelalgia
- Non-Atherosclerotic Vascular Disorders such as Fibromuscular Dysplasia, Popliteal Artery Entrapment Syndrome, Cystic Adventitial Disease, External Iliac Artery Endofibrosis, TAO/Buerger's disease (think of these disorders in young patients with no risk factors for ASO)
- Uncommon arteriopathies such as Vasculitis

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## What We Don't Do!



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## Vascular Medicine at the UPMC Heart and Vascular Institute

- Comprehensive Program and Model
- Eclectic Outpatient Consultative Services
- Inpatient Consultations
- Multidisciplinary Comprehensive ASO Clinic
- Thrombosis Clinic and Coagulation Clinic
- Stroke Bridge Clinic/Post Stroke Risk Factor Management Clinic
- Centralized Outpatient Vascular Lab
- Screening Programs/Pilot Programs

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## UPMC Hamot Vascular Program

- **Multidisciplinary**
- **Team Approach with Resources/Support**
- **Center of Excellence**
- **Vascular Medicine as the Front Door**
- **Complementary and Cooperative**
- **Screening**
- **Atherosclerosis Clinic Model**
- **RFM Model and Report Cards**

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## Multidisciplinary Comprehensive Atherosclerosis Clinic

- **Early Detection/Treatment**
- **Surveillance and Outcomes**
- **Screening Programs and Expos**
- **Data Mining Excursions with EMRs (Epic)**
- **Medical Nutrition/Smoking Cessation/NA**
- **Exercise Programs/Walking Programs**
- **Patient Education and Branded Handouts**
- **Report Cards/Outcomes**

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## Vascular Lab

- Centralized Outpatient Lab at the HHVI
- ICAVL Accredited
- SVU Signature Lab
- Reading Panel
- Expectations and Oversight
- Potential for Core Lab Research

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## Thrombosis Clinic

- VTE Usual and Unusual
- Low Risk DVT Disposition Program with Admit/Readmit Prevention
- PERT Program with CDT and mechanical thrombectomy
- Vascular Services Council and HHVI Leadership
- System Wide Anticoagulation Committees and Pathways
- Anticoagulation Clinic Model

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## The Burden of Vascular Disease

- Greater than 25 million people living in the US suffer from non coronary vascular disease
- Represents the single most important cause of death and disability in our nation and will remain so for decades ahead. Examples include:
  - PAD - affects 1 in 5 males, 1 in 6 females age greater than 65, > 8-11 million Americans
  - DVT/PE – most preventable cause of hospital death
  - Aneurysms
  - CVA - Unusual Vascular Disorders
  - “Orphan Disorders” – Lymphedema/CVI

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## Venous Thromboembolism (VTE)

- Deep Venous Thrombosis
- Pulmonary Embolism
- VTE in Unusual Sites

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## Venous Thromboembolism

- Pathophysiology
- Risk Factors
- Diagnosis/Clinical Prediction Rules
- Treatment and Recurrence Risk
- New and Emerging Therapies
- Updated ACCP Guidelines
- Outpatient VTE Therapies
- Hospital Outcomes

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## Virchow's Triad Rudolf Virchow c. 19<sup>th</sup> Century

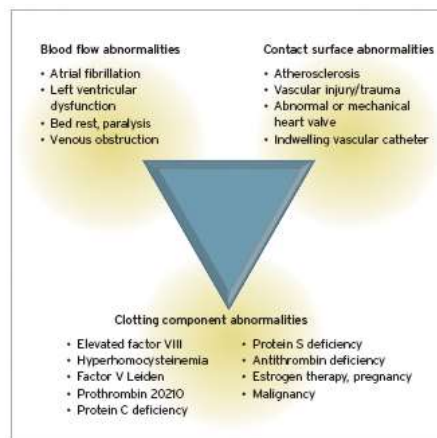


FIGURE 2. Virchow's triad

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## Virchow's Triad

- **Damage to the Lining of Vein**
  - Permits clots to attach themselves to the damaged portion (e.g. hip/knee surgery, CHF)
- **Slowing of Blood Flow**
  - Allows clumping of blood coagulation factors that would normally be washed away (e.g. bedrest)
- **Increased Tendency to Clot**
  - Encourages rapid clot formation (e.g. cancer/ID)

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## Epidemiology of DVT/PE

- **At least 600,000 Americans suffer PE and over 1 million suffer DVT annually, some estimates 3X higher**
- **100,000-180,000 US deaths per year, kills more people than traffic accidents, HIV and breast cancer per year**
- **PE is the #1 preventable cause of death among hospitalized patients**
- **Negative impact on QOL of survivors: CTEPH and PTS**
- **Health care costs over 10 billion dollars in 2011**

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## Prevention

- **Significant medical/financial impact of VTE**
- **System wide approach to Prevention**
- **Over 2 million people develop DVT annually**
- **DVT progress to PE in 600,000 cases with 60,000 fatalities**
- **Morbidity of the debilitating post thrombotic syndrome can arise in 1/3, esp. in patients with extensive or recurrent DVT**

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## My Approach to VTE

- **First Event or Recurrent**
- **Location Usual or Unusual**
- **Provoked or Unprovoked/Idiopathic**
- **Unusual Historical Features or Factors**
- **Site Confirmation – Distal/Proximal/Iliacs**
- **Massive/Submassive/Low Risk for PE**
- **Pick Treatment**
- **Pick Duration**
- **Prevent Complications**

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## My Approach to VTE

- **Provoked vs Unprovoked Recurrence Risk**
- **DVT Cephalad Propagation Risk**
- **Renal/Hepatic Function**
- **Access to Laboratory Monitoring**
- **Patient Compliance**
- **Compression Regimens**
- **ACCP/CHEST 2016**

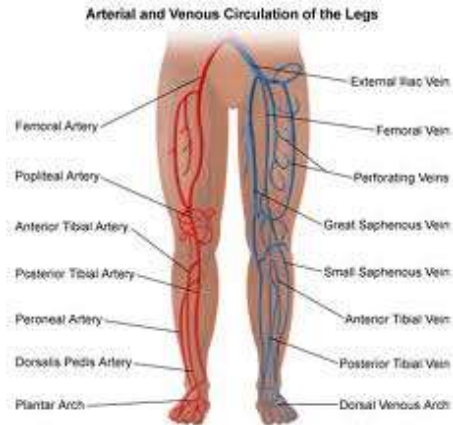
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## DVT

- **Deep Veins of the arms and legs most common**
- **Legs**
  - Proximal
  - Distal
- **Definition**
  - Venous thrombi typically form along the valve cusps
  - Propensity to embolize greatest in the first 7 days

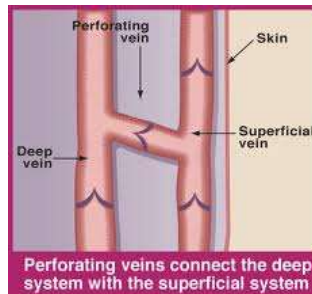
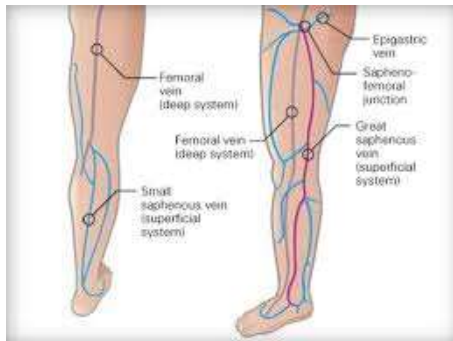
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# LE Venous Anatomy



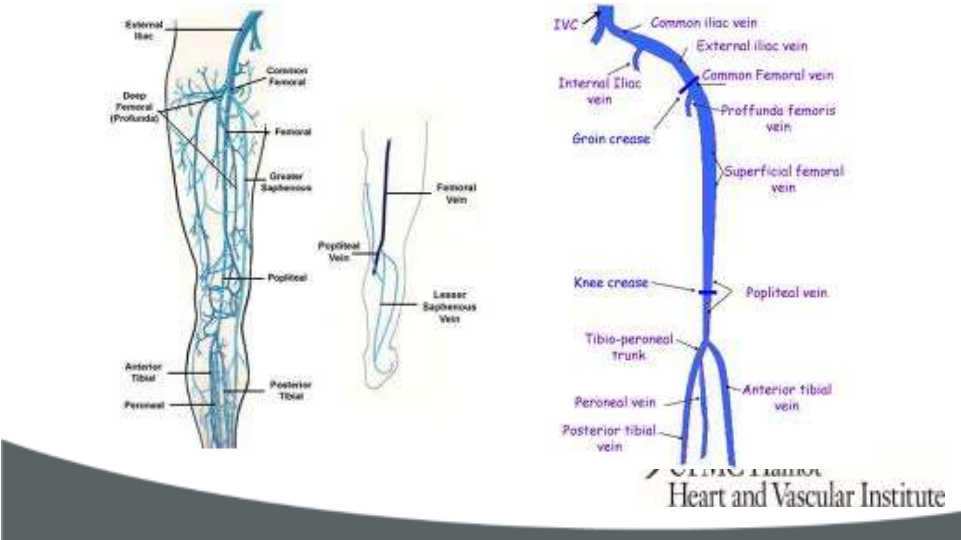
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# Venous Anatomy

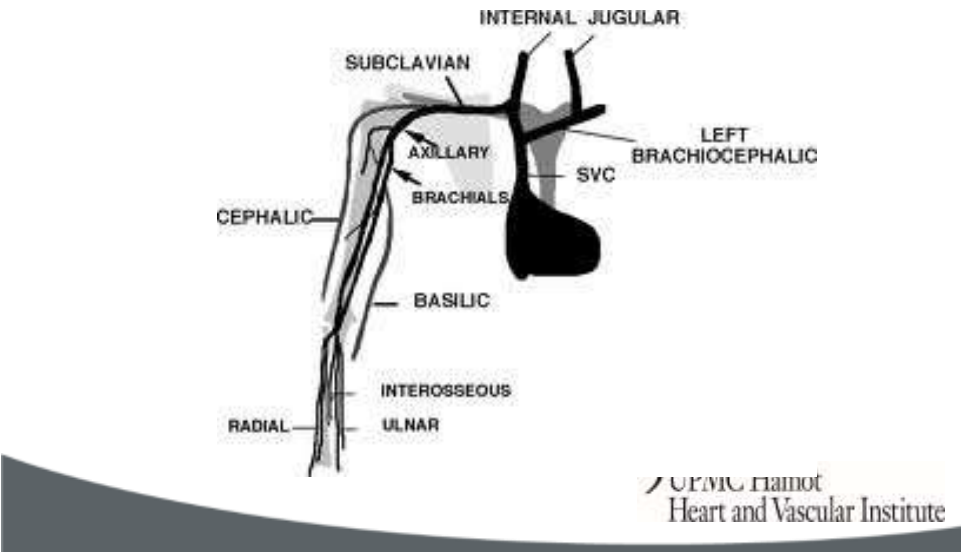


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## LE Venous Nomenclature



## UE Venous Anatomy





## Definitions of Pulmonary Embolism via Guidelines

- **Massive PE (5-10%):** Sustained hypotension, pulselessness, or persistent bradycardia
- **Submassive PE (20-25%):** RV dysfunction or myocardial necrosis, without hypotension
- **Low Risk PE (70%):** no markers of adverse prognosis
  - (Circulation 2011; 123: 1788-1830)

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## Risk Stratification in PE Essential for Management

- **Anticoagulation alone versus anticoagulation plus thrombolysis/pharmacomechanical catheter directed therapy/surgical embolectomy/IVC filter**
- **Triage ICU monitoring vs other**
- **Low Risk get A/C alone, High Risk get A/C plus lysis or embolectomy**
- **Submassive is the grey area**

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## Risk Factors for VTE

- **Surgery/Trauma/Acute Medical Illness**
- **Immobility /LE paresis/ Stroke**
- **Cancer/Cancer Therapy/PNH/Myeloprolif d/o**
- **Previous VTE**
- **Increasing age/Obesity/Smoking**
- **Estrogens/Inherited/Acquired Thrombophilias**
- **Nephrotic Syndrome /Inflammatory Bowel Dis**
- **Central Venous Catheters**
- **Rheumatoid Arthritis**
- **Chronic Liver Disease (up to 1%)**

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## Risk Factors for VTE

- **Strong – Odds Ratio > 10**
  - Fracture – hip/leg
  - Hip or knee replacement
  - Major general surgery
  - Major trauma
  - Spinal cord injury

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## Risk Factors for VTE

- **Moderate – Odds Ratio 2-9**
  - Arthroscopic knee surgery
  - CVL
  - Chemotherapy/Malignancy
  - CHF/Respiratory Failure
  - HRT/BCP/Pregnancy/Post Partum
  - Paralytic stroke
  - Thrombophilia/Previous VTE

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## Risk Factors for VTE

- **Weak – Odds Ratio < 2**
  - Bedrest > 3 days
  - Immobility due to sitting e.g. prolonged car or air travel
  - Increasing age
  - Laparoscopic surgery
  - Obesity
  - Varicose Veins

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## Aggressive Thrombophilias

- **Homozygous/Double Heterozygous Mutations**
  - Factor V Leiden
  - Prothrombin 20210A
- **Antiphospholipid Antibody Syndrome**
  - LA + - Confirm/Repeat
- **Deficiencies:**
  - Antithrombin
  - Protein C
  - Protein S

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## Clinical Features - DVT

- **Non specific**
- **Clinical findings not reliable**
- **Pain/redness/warmth common symptoms**
- **Swelling and tenderness are common signs**
- **Venous distension/palpable cords not specific**
- **Homan's sign not often found**
- **About a 42% chance of making diagnosis on physical exam**
- **About half of DVT patients are asymptomatic**

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## Clinical Features - PE

- Unexplained shortness of breath
- Pleuritic chest pain
- Hemoptysis
- Tachycardia
- Hypotension
- Syncope
- Anxiety

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## Clinical DVT



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## Phlegmasia Cerulea Dolens



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## Differential Diagnosis - DVT

- Muscle Strain or Tear
- Bakers Cyst
- Lymphangitis/Lymphatic Obstruction
- Venous Reflux
- Cellulitis
- Internal abnormality of the knee

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## Clinical Prediction Rules

- Integrate the results of a clinical index with the results of an ultrasound examination
- Enhance the predictive accuracy of a positive ultrasound to 100%, 96%, and 63% in high, moderate, and low probability groups
- Wells' Prediction Index
- DVT and PE

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## Wells DVT Criteria

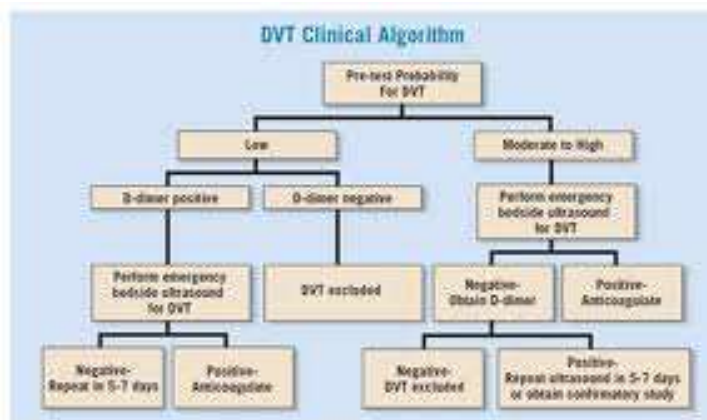
Symptom	Score
Active cancer (treatment ongoing or within previous 6 months or palliative)	1
Paralysis, paresis or recent plaster immobilization of the lower extremities	1
Recently bedridden > 3 days or major surgery within 4 weeks	1
Localized tenderness along the distribution of the deep venous system	1
Entire leg swollen	1
Calf swelling 3 cm > asymptomatic side (measured 10 cm below tibial tuberosity)	1
Pitting oedema confined to the symptomatic leg	1
Collateral superficial veins (non-varicose)	1
Alternative diagnosis as likely or greater than that of DVT	2

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## Wells DVT Criteria

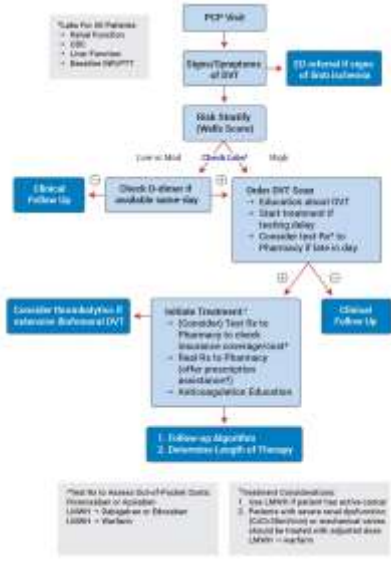
- Score of 3 or greater represents high probability
- Score of 1-2 represents moderate probability
- Score of less than or equal to 0 represents low probability
- Think “Low or Non-Low”

## DVT Clinical Algorithm





# DVT Flow Chart



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# The Vascular Lab



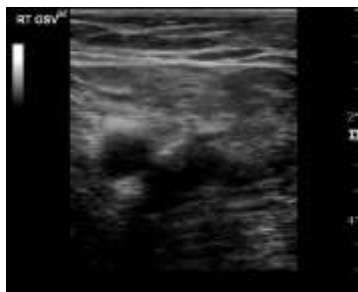
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## The Vascular Lab and Testing/Diagnosis

- Venous Testing – Venous Thrombosis
  - Venous Duplex Testing with DVT Protocol
  
- CTA Chest protocol for PE

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## Venous Compression Images



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## DVT CFV



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## Jugular Vein VTE



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## Treatment of VTE

- **ACCP Guidelines Updated 2016**
- **UFH (1B)**
  - Weight-based nomograms/HIT incidence 5-7%
- **LMWH (1A)**
  - Weight based/Avoid in RF/Preferred IA Rx
  - HIT incidence less than 1%
- **Fondaparinux (1A)**
  - Acceptable Rx/Avoid in RF/One case of HIT
- **VKA/Warfarin/New Agents and New Recommendations**
  - Acceptable Rx/Avoid in Pregnancy

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## Recurrence Rates after Anticoagulation Discontinuation

• Risk Factor	Recurrence Rate 1/5 yr	
• Surgery	1%	3%
• Nonsurgical Reversible/Transient Risk Factor	5%	15%
• Unprovoked	10%	30%

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## Treatment of DVT

- **Provoked**
  - 3 months
- **Recurrent/Unprovoked**
  - 3 months then shared medical decision making discussion regarding recurrence risks, bleeding risks, etc.
- **Thrombophilia-Related**
  - Probably based on clinical factors, patients with APLAS should receive anticoagulation indefinitely

## Unprovoked/Idiopathic DVT

- **Occurs without a clearly identified risk factor i.e. surgery or transient risk factor**
- **Recurrent risk accumulates once anticoagulation is stopped**
- **Three historical strategies to identify high risk:**
  - D-dimer testing
  - Evaluation for residual vein thrombosis
  - Clinical prediction rules

## Treatment of DVT

- **Idiopathic VTE**
  - 3 month duration or more
  - Risk of recurrence 7-10% per year and 30-42% at 5 years
  - D-dimer predictor 3% negative and 10%/yr. +
- **Cancer**
  - Recurrence rate up to 30% per year
  - LWMH preferred treatment for at least 6 months or until cancer is no longer “active” (CLOT trial)

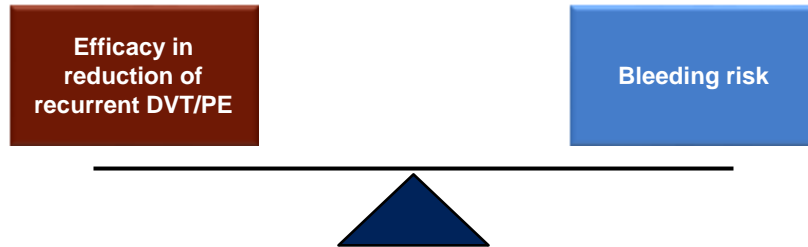
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## Who Get's Secondary Prevention?

- **First unprovoked VTE with low-moderate bleeding risk (2B)**
- **Second unprovoked VTE with low-moderate bleeding risk (1B)**
- **Therapy should be re-assessed annually!**

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## Extended Treatment for DVT and PE: Finding the Balance



- ◆ Clinicians must often balance the long-term risks of recurrent VTE if anticoagulation is stopped against the burden and risks of ongoing therapy

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## ABC's of Bleeding Risk Assessment

- Age > 65, Antiplatelet Therapy, Alcoholism
  - Bleed History
  - Cancer
  - Diabetes
  - Anemia
  - Falls
  - GFR Decrease
  - Hepatic Disease
  - Stroke
  - Surgery (recent)
  - Thrombocytopenia
- Low 0 RF  
Mod 1 RF  
High 2 or more RF

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## Bleeding Risk Assessment

- | <b>Risk Category</b> | <b>Bleeding Rate</b>       |
|----------------------|----------------------------|
| • Low                | 2.4/100 pt. years (3fatal) |
| • Moderate           | 4.9/100 pt. years (5fatal) |
| • High               | 9.8 /100 pt. year(11fatal) |

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## HAS-BLED Score

- Hypertension
- Abnormal Liver/Renal Function
- Stroke History
- Bleeding Presentation/History
- Labile INRs
- “Elderly” Age 65 or greater
- Drugs/Alcohol Use

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## HAS-BLED Score

- **Score of 0 is 0.9% risk of bleeding in one validation study and 1.3 bleeds per 100 patient/years in another validation study**
- **Score of 5 is 9.1% risk of bleeding in one validation study and 12.5 bleeds per 100 patient/years in another validation study**

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## MEN and HERDOO2

- **Risk Factors for Recurrent VTE**
- **MEN and signs of post thrombotic syndrome:**
- **Hyperpigmentation of the lower extremities**
- **Erythema or Redness of the LE**
- **D-dimer level greater than 250 mcg/L**
- **Obesity with BMI greater than 30 kg/m<sup>2</sup>**
- **Older age > 65 years**
- **Two or more risk factors at higher risk**
- **Recent paper regarding women with scores of 0 and 1**

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## ACCP 2016 Updates

- For VTE and no cancer, DOACs over VKA, and VKA over LMWH
- For VTE and cancer, LMWH or VKA and DOACs
- No changes in duration
- Recommend against IVC filters if on A/C
- Recommend thrombolytic therapy with PE and hypotension, and systemic therapy over CDT
- For recurrent VTE on a non-LMWH anticoagulant, recommend LMWH, if on LMWH, then increase dose

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## Preventing Post thrombotic Syndrome

- Chronic burdensome consequence of DVT that occurs despite anticoagulation therapy
- 23-50% of patients and manifests typically in first 2 years
- Leg pain, heaviness, swelling, and cramping
- Severe cases include venous ulcers
- Villalta scale categorizes into mild, moderate or severe – wait 3 months to attribute the diagnosis

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## Post thrombotic Syndrome

- **Compression stockings may reduce risk (of any severity) from 43% to 20% and severe post thrombotic syndrome from 15% to 7%**
- **30-40 mmHg and consider continuing for a minimum of two years if patient has swelling or discomfort**
- **Start as soon as possible after starting anticoagulation therapy**

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## Anticoagulants and Antiplatelets

- **Aspirin - Oral**
- **Warfarin - Oral**
- **UFH/LMWH - Parenteral**
- **Fondaparinux - Parenteral**
- **NOAC/NOAC/TSOAC/DOAC - Oral**
- **Dabigatran –Direct Thrombin Inhibitor**
- **Rivaroxaban – Xa Inhibitor**
- **Apixiban – Xa Inhibitor**
- **Edoxaban – Xa Inhibitor**

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## Treatment of VTE

- Initial anticoagulation used to require overlap “bridging” of a parenteral anticoagulant (UFH, LMWH or fondaparinux) with VKA for a minimum of 5 days and until INR above 2 for at least 24-48 hours
- New agents are now a recommended option
- Determine length of anticoagulation
- Prevent post thrombotic syndrome
- Appropriate screening for occult malignancy prn

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## Treatment of VTE

- Idiopathic/Unprovoked VTE
  - 3 month duration, then decide on more
  - Risk of recurrence 6-10% per year
  - D-dimer predictor 3% negative and 10%/yr +
- Cancer
  - Recurrence rate up to 30% per year
  - LMWH preferred treatment for at least 6 months or until cancer is no longer “active” (CLOT trial)

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## Treatment of VTE

- **Rapid initiation of anticoagulation prevents thrombus extension and PE**
- **Extended anticoagulation reduces the risk of recurrent VTE**
- **Don't forget compression stockings if indicated for discomfort and ? prevention of post thrombotic syndrome**
  - 30-40 mmHg for at least 2 years

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## Direct Oral Anticoagulants (aka NOACs/DOACs)

- **Non-Bridged**
  - Rivaroxaban/Xarelto
    - Once a day dosing
    - Starter Pack Available with Free Voucher
  - Apixiban/Eliquis
    - Twice a day dosing
- **Bridged with LMWH or UFH**
  - Edoxaban/Savaysa
  - Dabigatran/Pradaxa

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## Practical Management Issues with Anticoagulants including New Oral Anticoagulants (DOACs)

- Starting
- Switching
- Monitoring
- Stopping/Reversal/Procedural Considerations

### ASA for Preventing VTE Recurrence WARFASA

- **VTE Recurrence**
  - ASA 6.6% per year vs.
  - Placebo 11.2% per year
- **Major bleeding (100 mg/d)**
  - One patient per group

**ASA more effective than placebo for decreasing the risk of recurrent VTE in patients after VKA therapy following first idiopathic VTE**

NEJM 2012;366:1959

## Aspirin for VTE Recurrence Prevention - ASPIRE

- 100 mg dose of ASA reduced by one third the rate of recurrent major vascular events for patients inc VTE, MI, stroke or CV death
- Enrolled patients who had one acute unprovoked VTE and were switched after 3 months of anticoagulant therapy to either ASA or placebo
- Low numbers and power for prediction
- Reasonable “intermediate option”

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## SVM/ABIM Choosing Wisely Campaign

- Don't do a work up for a clotting disorder for patients who develop first episode of DVT in the setting of a known cause
  - Increased testing with no proven benefit
- Don't reimaging DVT in the absence of a clinical change
  - Repeat ultrasound images to evaluate the “response” of a venous clot to therapy does not alter treatment

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## Vena Cava Filters

- **Two primary indications**
  - Absolute contraindication to anticoagulation
  - Failed anticoagulation – i.e. recurrent thromboembolism while receiving therapeutic doses of anticoagulation
- **Do not afford protection from further DVT, rather increase the risk of secondary DVT**
- **Permanent vs Retrievable**

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## New/Emerging Therapies

- **Catheter Directed Thrombolysis**
- **Mechanical Thrombectomy**
- **Treatment of Ilio-Femoral DVT**
- **Anticoagulants and Antiplatelets**
  - DOACs/NOACs
  - Reversal Agents

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## Catheter Directed Thrombolysis

- Delivers thrombolytic agent locally into thrombus using infusion catheters
- Accelerates thrombolysis, reduces dose/duration and decreased bleeding complications as compared to systemic thrombolysis
- EKOS program
- Urokinase and rt-PA have been studied
- Iliofemoral segment DVT results promising for lysis, preservation of valve function and decreased post thrombotic syndrome

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## Percutaneous Mechanical Thrombectomy

- Can be used in combination with CDT
- AngioJet System directs saline jets to macerate and remove thrombus
- Trellis device uses occlusive balloons and dispersion wire to remove thrombus
- Combined with CDT has potential to remove more clot as well as decrease the dose and duration of thrombolytic therapy

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## VTE Prophylaxis in Hospitalized Medical Patients

- **MEDENOX**
- **PREVENT**                      **All q day dosing**
- **ARTEMIS**
  
- **Once daily injected low-dose anticoagulant prophylaxis placebo-controlled trials**
  - Reduced DVT greater than 50% without increasing major bleeding

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## VTE as the First Manifestation of Cancer

- **Strong consideration should be given regarding cancer screening for an idiopathic VTE event**
- **Patients with idiopathic VTE have a significant risk of occult cancer within the first year after diagnosis**
- **History/Physical**
- **Up to date with general health maintenance issues**

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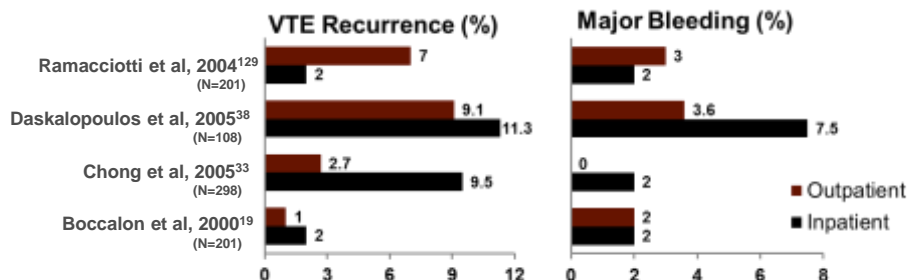
## Outpatient Treatment of DVT and PE

- **ACCP Guidelines 1B for Acute DVT**
  - Initial therapy at home over treatment in the hospital
- **ACCP Guidelines 2B for Low-Risk PE**
  - Initial treatment at home or early discharge over standard discharge (PE is a tougher call due to litigious culture)
- **Contingent on adequate home circumstances**
  - Well maintained living conditions/phone access
  - Strong support network/patient feels well enough
  - Patient has ability to promptly hospitalized if necessary

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## Acute DVT May Be Managed in the Outpatient Setting

- **Controlled clinical trials suggest that outpatient management is at least as effective as inpatient management for acute DVT**



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## Considerations for Patient Selection for Outpatient Therapy

- **60%-95% of patients with acute, proximal DVT may be eligible for outpatient therapy**
- **Exclusion criteria on institutional protocols include:**
  - Comorbid illness requiring hospitalization
  - Active or high risk for bleeding
  - Severe hypertension
  - Catheter-associated DVT
  - Recent surgery
  - Morbid obesity
  - Hypercoagulable state
  - Pregnancy

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## Considerations for Identifying Patients With Low-Risk PE

- ◆ **Risk stratification tools may help to identify patients with low-risk PE who may be candidates for outpatient therapy**
- ◆ **Potential candidates include patients with acute PE who are:**
  - Clinically stable with good cardiopulmonary reserve
  - No recent bleeding
  - No severe thrombocytopenia (ie, platelet count  $\geq 70,000/\text{mm}^3$ )
  - No severe liver or renal disease
  - Expected to be compliant with treatment
  - Feeling well enough to be treated at home

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## The PESI and Simplified PESI Are Validated Tools Used to Identify Low-Risk Patients

Variable	Score	
	PESI	sPESI
Age >80 years	Age in years	1
Male sex	10	0
History of cancer	30	1
History of heart failure	10	
History of chronic lung disease	10	1*
Pulse $\geq$ 110 bpm	20	1
Systolic BP <100 mm Hg	30	1
Respiratory rate $\geq$ 30 breaths/min	20	0
Temperature <36° C	20	0
Altered mental status <sup>†</sup>	60	0
SaO <sub>2</sub> <90% <sup>‡</sup>	20	1



Classification by Total Score		
	PESI	sPESI
Class I	$\leq$ 65	Low risk=0
Class II	66-85	
Class III	86-105	High risk $\geq$ 1
Class IV	106-125	
Class V	>125	

\*Heart failure or history of chronic lung disease combined into a single category of chronic cardiopulmonary disease.

<sup>†</sup>Disorientation, lethargy, stupor, or coma. <sup>‡</sup>With or without the administration of supplemental oxygen.

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## Who May Be Candidates for Outpatient Therapy?

- The decision to treat a patient with DVT or low-risk PE in the outpatient setting is based on numerous factors:
  - Adequate home circumstances
  - Risk factors for recurrence and/or bleeding
- Risk stratification tools may aid in patient selection
- The actual decision depends on the clinical judgment of the treating clinician

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## Outpatient Treatment of VTE

- Despite outpatient options for DVT and low-risk PE, hospital admissions for VTE remain high
- Average cost in 2005 for admitted DVT was \$10K and for PE about \$15K
- If you aren't admitted, you can't be readmitted!
- Hospital admissions can lead to complications
- New oral drugs can decrease LOS, in one study with Xarelto resulted in 3 day decrease in LOS

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## UPMC System Wide Approach

- System Wide Anticoagulation Committee
- System Wide P&T Committee
- Pilot Programs including Xarelto To Go
- Trifold Handouts for Clinicians
- Xa level (rather than aPTT) monitoring for UFH
- HIT protocols

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# The Real Value of Vascular Medicine?



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# Thank You!



If Women Ran the World

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