Venous thromboembolism (VTE) represents a major cause of morbidity and mortality in the United States. The incidence of VTE exceeds 1 per 1000 with approximately 250,000 first lifetime cases diagnosed annually in the United States. The 30-day mortality of patients suffering a thrombotic event exceeds 25%. Of those individuals surviving the thrombotic event, 30% will develop recurrent VTE within 10 years. Furthermore, the risk of postthrombotic syndrome is approximately 30% at 10 years, with 10% of the patients suffering venous stasis ulceration.

Venous thromboembolism is typically a disease of older people, with the incidence of thrombotic events increasing substantially beyond the age of 60 years. Despite advances in radiographic detection, expanded knowledge of risk factors, and anticoagulant development, the incidence of VTE has been relatively constant over the past several decades.

The American College of Chest Physicians Antithrombotic Guidelines, 9th edition (AT9), were published in 2012 and include a large number of clinical practice recommendations encapsulated in an executive summary with 21 separate supporting articles that explore the available evidence used to formulate the guidelines. Expert panelists reviewed the literature underlying each clinical question, evaluated the quality of evidence (A = high, B = moderate, and C = low), and provided a strength of the resulting recommendation (strong = grade 1 and weak = grade 2).

The intent of the article is to provide clinicians with a focused review of the AT9 VTE guidelines in a concise, easily digested format. Furthermore, controversial areas of clinical practice in which the guidelines are not straightforward are discussed. In particular, we endeavor to clarify issues surrounding debated treatment decisions and offer our current practice patterns for these situations.

VTE EVALUATION

Pretest Probability of Disease Assessment

The AT9 guidelines for the evaluation of patients with suspected VTE are based on several important considerations. First, the clinical evaluation is an essential component to initial evaluation and must incorporate a pretest probability of disease assessment. The clinical assessment alone, however, is unreliable and must include further objective testing whereby only a minority of patients with suspected deep vein thrombosis (DVT) actually have the diagnosis (20%). Moreover, the consequences of misdiagnosis are serious. Patients may be placed at an unnecessary risk of major hemorrhage if anticoagulation therapy is initiated for a thrombotic event that is not present. In contrast, patients with VTE who were left untreated may suffer thrombus propagation, embolization, or death. According to the AT9 guidelines, diagnostic strategies are deemed acceptable if 2% or less of the patients with VTE are missed during the evaluation process including the ensuing 3 to 6 months.

The Wells criteria for the clinical pretest probability assessment of DVT classify patients into low, moderate, or high probability categories depending on the underlying risk,
physical findings on examination, and the probability of a more likely alternate diagnosis. On the basis of this tool, the prevalence of DVT ranges from 5% (low), 17% (moderate), to 53% (high) pretest probability of disease. Similar tools are available for pulmonary embolism (PE) prediction. For patients with suspected VTE, the AT9 guidelines recommend performing a thorough clinical assessment to determine the pretest probability of disease to guide further evaluation (grade 2B; Figures 1 and 2).

**Fibrin D-Dimer**

Fibrin D-dimer provides a measure of ongoing fibrinolysis. When combined with clinical pretest probability of disease assessment, fibrin D-dimer can be useful in the diagnostic evaluation of patients suspected to have VTE (Figure 1). These assays can be generally classified as quantitative or qualitative. The quantitative assays use ELISA, enzyme-linked or latex platforms for plasma-based testing, and take longer to perform but are highly sensitive for circulating fibrin D-dimer. The qualitative assays are typically performed on whole blood samples and are available at point of care. With the use of the highly sensitive quantitative assays, a negative test result when combined with low or moderate pretest probability of disease clinical assessment excludes the diagnosis of venous thrombosis with 99% negative predictive value. The high sensitivity fibrin D-dimer assay is therefore the first line of testing for patients with low-to-moderate pretest likelihood of disease (grade 2B). Similar negative predictive values can be achieved with the moderate sensitivity qualitative assays, however, only for patients at low pretest probability of disease. The moderate sensitivity assay is the first line of testing for patients with low pretest likelihood of disease only (grade 2C). For either type of assay, a positive test result is not helpful and carries a low positive predictive value. For patients likely to have a positive assay result owing to comorbid conditions such as recent surgery or major trauma, fibrin D-dimer assessment would have little utility.

**Compression Duplex Ultrasound**

Imaging with compression duplex ultrasound is warranted for patients with a high pretest probability of DVT, for patients with a moderate pretest probability of disease in which the available fibrin D-dimer assay has moderate sensitivity, or for patients with a positive fibrin D-dimer test result (Figure 1). Compression ultrasound testing of the lower extremity for venous thrombosis assessment is defined as “proximal,” where the popliteal, femoral, and common femoral veins are imaged, or as “whole leg,” which includes an assessment of venous segments of the calf. Whole leg compression ultrasound is useful in the assessment of symptomatic patients with an overall specificity of 96% and a 3-month VTE rate of 0.57% with a

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**ARTICLE HIGHLIGHTS**

- Practical recommendations for peri-operative venous thromboembolism prophylaxis.
- Discussion of available antithrombotic agents used in venous thromboembolism treatment.
- Brief review of clinical evaluation for deep vein thrombosis and pulmonary embolism.
- Recommendations for controversial areas of venous thromboembolism management.
When an ultrasound assessment is limited to proximal veins, a negative study result does not adequately exclude DVT. For patients with high pretest probability of disease and a negative "proximal" ultrasound, the AT9 guidelines recommend further testing to include either a high sensitivity fibrin D-dimer, whole leg ultrasound, or repeat proximal ultrasound in 1 week (all grade 1B) or venography (grade 2B).6

Computed Tomography and Magnetic Resonance Venography
Neither computed tomography venography nor magnetic resonance venography is recommended for routine use in the evaluation of DVT.6

Specific Evaluation for PE
Signs and symptoms of PE are often nonspecific11; therefore, a well-structured evaluation is needed when the diagnosis of PE is considered. The Wells8 and Geneva9 criteria provide a framework for risk-stratifying individuals on the basis of the pretest probability of disease. The Christopher study12 categorized patients into “PE unlikely” for a Wells score of 4 or less or “PE likely” for a Wells score of more than 4, with subsequent evaluation dependent on the likelihood of disease (Figure 2). Online calculators are readily available to calculate the Wells score for DVT (http://www.mdcalc.com/wells-criteria-for-dvt/) and PE (http://www.mdcalc.com/wells-criteria-for-pulmonary-embolism-pe/). Additional imaging is necessary for patients with “PE likely” or an abnormal D-dimer test result. Computed tomography pulmonary angiography is the most common imaging modality used to diagnose PE with a high specificity and sensitivity.13 With high pretest probability of disease, a negative study result is associated with a 3-month cumulative VTE rate of 1.5%. If a negative fibrin D-dimer test result is added to the testing algorithm, the 3 months event rate is 0.5% or less. The use of iodinated contrast dye is problematic for individuals with advanced renal disease, substantial risk factors for contrast-induced nephropathy, or allergic reactions. For these individuals, a ventilation-perfusion scan can be useful if interpreted as showing either definitely “positive” or definitely “negative” test results.13 However, a substantive proportion of ventilation-perfusion scans are “indeterminate” and require additional testing. The gold standard diagnostic tool for PE, invasive pulmonary angiography, has largely been supplanted by using computed tomography pulmonary angiography.

VTE TREATMENT

Medications for Acute VTE
In patients with acute DVT or PE, initial therapy can include either parenteral anticoagulation (grade 1B) or oral rivaroxaban.14 For patients bridged to warfarin, the use of either low-molecular-weight heparin (LMWH) or fondaparinux is preferred to unfractionated heparin (UFH) (grade 2C).14

Unfractionated Heparin
Unfractionated heparin is typically given intravenously by using a weight-based dosing algorithm (80 U/kg bolus followed by 18 U/kg per hour).15,16 Dose adjustment nomograms using the activated partial thromboplastin time should be validated at one’s institution, with an activated partial thromboplastin time ratio of 1.5 to 2.5 approximating heparin levels of 0.3 to 0.7 IU/mL measured by using an anti–factor Xa assay.15,17 Weight-based unmonitored subcutaneous UFH has been found to provide an
effective treatment of acute VTE (Table 1). In the FIDO trial, subcutaneous heparin (333 U/kg followed by 250 U/kg twice daily) was as safe and effective as weight-based low-molecular-weight UFH for the treatment of VTE.

An important complication of heparin therapy is heparin-induced thrombocytopenia (HIT), which may be associated with both arterial and venous thrombosis. With UFH, the risk of HIT may be as high as 5%, depending on the patient population studied. The development of HIT appears to be particularly high in patients recuperating from orthopedic surgery and receiving UFH. Immune-mediated HIT is associated with a decline of 50% or more in platelet count typically within 5 to 10 days after beginning heparin (earlier with previous exposure). In patients with a risk of HIT of 1% or more, the AT9 guidelines recommend platelet count monitoring every 2 to 3 days from day 4 to day 14 while receiving heparin to proactively screen for thrombocytopenia (grade 2C).

**Low-Molecular-Weight Heparin**

Food and Drug Administration (FDA)-approved LMWHs include enoxaparin and dalteparin. For patients with acute lower extremity DVT with or without PE, the AT9 guidelines prefer LMWH over intravenous (grade 2C) or subcutaneous (grade 2B) UFH. Low-molecular-weight heparin should overlap warfarin initiation for a minimum of 5 days or until the international normalized ratio (INR) exceeds 2.0 for at least 24 hours (grade 1B).

Once daily injection is preferred to twice daily injections (grade 2C). For once daily dosing, however, the AT9 guidelines recommend using the full anticoagulant LMWH dose of enoxaparin (2 mg/kg once daily) or dalteparin (200 IU/kg once daily) (no grading provided). Many clinicians however, use enoxaparin 1.5 mg/kg once daily on the basis of a single randomized trial. This trial compared intravenous UFH with enoxaparin 1 mg/kg twice daily or enoxaparin 1.5 mg/kg once daily in 900 patients with symptomatic leg DVT, including approximately one-third with PE. In this study, neither symptomatic VTE recurrence nor major bleeding differed by treatment arm. On the basis of this study, enoxaparin 1.5 mg/kg once daily has therefore become the clinical standard for outpatient treatment of DVT. It is our practice to treat patients with either acute DVT or small PE with enoxaparin 1.5 mg/kg once daily. For

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**TABLE 1. VTE Treatment Options (Presently FDA Approved)**

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<tr>
<th>Medication</th>
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| Unfractionated heparin | Intravenous: 80 U/kg bolus and then 18 U/kg per hour infusion  
Subcutaneous: 333 U/kg and then 250 U/kg twice daily | • HIT monitoring  
• Nomogram aPTT ratio of 1.5-2.5 with anti--factor Xa level (0.3-0.7 IU/mL) |
| Enoxaparin       | Subcutaneous: 1 mg/kg twice daily or 1.5 mg/kg once daily              | • If CrCl < 30 mL/min, caution is advised because of LMWH accumulation       |
| Dalteparin       | Subcutaneous: 200 U/kg once daily                                      | • If CrCl < 30 mL/min, caution is advised because of LMWH accumulation       |
| Fondaparinux     | Subcutaneous: Based on weight:                                         | • Contraindicated if CrCl < 30 mL/min                                         |
|                  | <50 kg = 5 mg once daily                                               |                                                                                |
|                  | 50-100 kg = 7.5 mg once daily                                          |                                                                                |
|                  | >100 kg = 10 mg once daily                                             |                                                                                |
| Warfarin         | Oral: Initial dosing typically 5 mg once daily titrated for a goal of INR 2-3 | • Lower starting dose in elderly patients, poor nutritional status, concurrent medications affecting metabolism, or underlying liver disease |
| Rivaroxaban      | Oral: 15 mg twice daily for 21 days and then 20 mg once daily          | • Avoid if CrCl < 30 mL/min  
• Avoid if moderate-to-severe liver disease or hepatic coagulopathy  
• Avoid for pregnant or nursing patients |

aPTT = activated partial thromboplastin time; FDA = Food and Drug Administration; HIT = heparin-induced thrombocytopenia; INR = international normalized ratio; LMWH = low-molecular-weight heparin; VTE = venous thromboembolism.
patients with large PE, we recommend enoxaparin 1 mg/kg twice daily.

Owing to the more neutral charge of LMWHs, there is a lower risk of HIT such that platelet count monitoring is not indicated for most patients (grade 2C).19

**Fondaparinux**

Fondaparinux, a synthetic pentasaccharide, binds antithrombin with high affinity and has a greater specificity for factor Xa inhibition.14 Given once daily weight-based dosing, it has excellent bioavailability after subcutaneous injection with a long half-life (~17 hours). Fondaparinux is contraindicated in severe renal insufficiency (CrCl < 30 mL/min). For patients with acute VTE, the AT9 guidelines prefer fondaparinux over intravenous heparin (grade 2C) or subcutaneous UFH (grade 2C).14 There is insufficient data for use in pregnancy.

The risk of HIT with fondaparinux is insignificantly low such that platelet count monitoring is not indicated (grade 2C).19

**Vitamin K Antagonist**

The warfarin-specific AT9 guidelines include the following: (1) early initiation is preferred to delayed initiation (grade 1B); (2) stable outpatients begun on warfarin at a dose of 10 mg for the first 2 days followed by anticipated maintenance dosing, monitored with INR measurements (grade 2C); (3) for patients with consistently stable INRs, INR testing intervals may be lengthened to 12 weeks instead of a month (grade 2B); (4) for patients with previously stable INRs who have a single out of range result ±0.5, continue the present dose and recheck the INR within 1 to 2 weeks (grade 2C); (5) home INR monitoring may be implemented for patients who demonstrate competency in self-testing and who have stable INR measurements (grade 2B); and (6) nonsteroidal anti-inflammatory drugs and/or antiplatelet drugs should be avoided unless specifically indicated (eg, mechanical heart valves and acute coronary syndrome; grade 2C). Of these guidelines, we have not adopted the practice of extending INR interval testing beyond 4 weeks. Our initiation dose of warfarin is more conservative than outlined especially for elderly patients and those with confounding medications and coexisting comorbidities known to increase warfarin sensitivity.

The use of vitamin K for lowering excessive INR values depends on both the hemostatic assessment of the patient and the level of INR excess.21 For INR values 4.5 to 10 with no bleeding, the routine use of vitamin K is not recommended (grade 2B).21 In such patients, it is our practice to reassess the INR the following day to ensure patient safety and determine the INR trend. For INR values greater than 10 and no bleeding, vitamin K may be orally administered (grade 2C).21 Typically 1 to 2 mg administered orally will result in INR reduction within 18 to 24 hours. With major bleeding, prothrombin complex concentrates are recommended (instead of plasma) in addition to parenteral vitamin K (grade 2C).21 It has been our experience that parenteral vitamin K is typically well tolerated.22 When indicated, we suggest 1 mg aliquots adequately diluted and administered intravenously (using an infusion pump) over 20 to 30 minutes followed by serial INR assessment.

For patients not receiving warfarin therapy, an increased INR in the setting of liver dysfunction (so-called auto-anticoagulation) is insufficient to prevent thromboembolism.23 Warfarin is metabolized by the highly polymorphic hepatic cytochrome p450 CYP2C9 system, which affects the rapidity of warfarin elimination. Enzymes responsible for vitamin K metabolism are also polymorphic, an important example being vitamin K epoxide reductase complex subunit 1.24,25 There are presently insufficient randomized controlled trial data to warrant universal pharmacogenetic testing before warfarin initiation. The AT9 guidelines do not endorse pharmacogenomic-guided warfarin dosing (grade 1B).21

**Direct Thrombin Inhibitor**

Dabigatran, an oral direct thrombin inhibitor, was assessed in the RECOVER trial which randomized 2539 patients with acute VTE to receive either dabigatran 150 mg twice daily or warfarin26 after a median of 9 days of the initial parenteral anticoagulant therapy. In this trial, dabigatran was found to be noninferior to warfarin from a safety and efficacy standpoint. The REMEDY and RESONATE trials27 assessed dabigatran 150 mg twice daily compared with either warfarin or placebo for the secondary prevention of VTE in patients already completing at least 3 months of anticoagulant therapy. Venous
thromboembolism recurrence rates were similarly improved with either dabigatran or warfarin. Bleeding rates were lower in patients treated with dabigatran compared with warfarin. At present, dabigatran is not FDA approved for either the treatment or the prevention of VTE.

Direct Factor Xa Inhibitors
Rivaroxaban, an oral direct factor Xa inhibitor, has a time to peak concentration of 2 to 4 hours with an elimination half-life of 7 to 11 hours. Metabolized in the liver and excreted by the kidneys, rivaroxaban is contraindicated in patients with severe liver disease, any degree of hepatic coagulopathy, or severe renal insufficiency (CrCl < 30 mL/min) for the VTE indication. Rivaroxaban is FDA approved for the treatment of VTE (grade 2B). In our practice, the use of rivaroxaban for the treatment of VTE is steadily increasing.

Antiplatelet Agents
Although aspirin is the cornerstone antiplatelet therapy for the prevention and treatment of coronary and cerebral atherothrombotic events, recent attention has focused on its role in VTE. The WARFASA trial found that aspirin 100 mg once daily reduced the risk of recurrence (6.6% vs 11.2% per year) in patients with unprovoked VTE after an initial treatment period of 6 to 18 months with a vitamin K antagonist. In the similarly designed ASPIRE trial, low-dose aspirin resulted in a trend toward reduced VTE recurrence rates (hazard ratio, 0.74; 95% CI, 0.52-1.05; P=.09). With regard to VTE secondary prevention, the AT9 guidelines make no specific recommendations. A recent study found that aspirin was noninferior to LMWH for extended VTE prophylaxis for patients after total hip arthroplasty who were initially treated with 10 days of LMWH.

Outpatient vs Inpatient Treatment
The decision to treat suspected VTE while awaiting confirmatory testing depends on the pretest probability of finding an acute DVT or PE and the anticipated delay to acquiring test results. If the pretest probability is high, prompt parenteral anticoagulation is recommended while over rivaroxaban for VTE treatment (grade 2B).
awaiting test results (grade 2C). For those at moderate pretest probability, early treatment is recommended if there is an anticipated delay in test results exceeding 4 hours (grade 2C). With a low clinical suspicion of DVT or PE using the Wells criteria, empiric therapy is not recommended, provided that confirmatory testing is available within 24 hours (grade 2C).

For stable patients without significant risk of bleeding, outpatient therapy is acceptable for acute DVT treatment (grade 1B). A number of studies have compared outpatient LMWH with inpatient intravenous UFH for the initial treatment of DVT. One study assessed the importance of hospitalization by randomizing 201 patients with DVT to outpatient vs inpatient LMWH for this initial therapy. The rates of thrombus extension (1% vs 2%), major bleeding (2% vs 2%), and mortality (0% vs 2%) were almost identical for outpatient vs inpatient LMWH delivery. For patients with DVT, threatened venous gangrene, or extensive iliofemoral involvement, inpatient admission for mechanical and pharmacologic thrombolytic therapy should be considered.

For appropriately selected patients with PE, outpatient treatment may also be acceptable. Criteria include hemodynamic and clinical stability, strong social support and access to medical care, and good adherence to medically recommended treatment. The simplified PE Severity Index (PESI) is an easy-to-use risk stratification tool to help discern which patients with PE may be treated as outpatients. This tool assigns 1 point each for age greater than 80 years, history of cancer, history of chronic lung disease or heart failure, pulse exceeding 110 per minute, systolic blood pressure less than 100 mm Hg, and oxygen saturation less than 90%. Risk is classified as low (0 points) or high (≥1 point) risk. In both the derivation and validation cohorts, nearly one-third of the patients who were deemed “low risk” by these criteria had a very low 30-day mortality rate of 1.1% or less. This result is comparable to a 30-day mortality rate approaching 10% for high-risk patients.

**Duration of Anticoagulation**

For a DVT or PE that is provoked by either surgery or a transient risk factor, the recommended duration of treatment is 3 months (grade 1B). In the setting of an unprovoked proximal DVT or PE, the duration of anticoagulation depends on the risk of bleeding for the individual patient. For patients at low-to-moderate risk for bleeding, extended therapy is recommended (grade 2B). For those with high risk of bleeding, treatment may be discontinued after 3 months (grade 1B).

Bleeding risk assessment may be complicated, relies on the summation of a number of variables, and risk calculation tools have not been well validated. Factors used in this prediction may vary by extent and severity (eg, thrombocytopenia extent), temporal relation to a specific provocation (eg, major trauma or surgery), and the extent to which previous causes have been modified. In general, bleeding rates vary from low (0.8%/y) to moderate (1.6%/y) to high (≥6.5%/y) risk. A number of bleeding prediction tools have been proposed, including online calculators.

Factors deemed inadequate to affect anticoagulation duration include fibrin D-dimer assessment after the initial treatment of VTE, antiphospholipid antibodies, hereditary thrombophilias, male sex, and the presence of residual thrombus on follow-up ultrasound imaging (no grade provided). It is our practice to provide prolonged secondary prophylaxis to patients with high-risk hereditary (homozygous or compound heterozygous factor V Leiden, prothrombin G20210A sequence variations, or congenital deficiencies of protein C, protein S, or antithrombin) or acquired (antiphospholipid antibody syndrome) thrombophilias.

**VTE in the Setting of Underlying Malignant Disease**

For patients with VTE in the setting of an active malignant disease, extended anticoagulation is recommended regardless of the bleeding risk (grade 2B) and LMWH is the preferred anticoagulant (grade 2B). Anticoagulants are continued until there is no evidence of active malignant disease defined as any evidence of cancer on cross-sectional imaging or any cancer-related treatment (surgery, radiation, or chemotherapy) within the past 6 months.

The CLOT Investigators randomized 672 patients with active cancer and acute VTE to receive either LMWH (dalteparin 200 IU/kg once daily for 1 month followed by 150 IU/kg once daily for 5 months) or adjusted dose warfarin. At the end of the 6-month study
period, recurrent thromboembolism was 17% in the oral anticoagulant group and 9% in the dalteparin group (hazard ratio, 0.48; \( P = .002 \)). There was no significant difference between groups in the rate of major bleeding (dalteparin 6% vs warfarin 4%). Enoxaparin is likely a reasonable alternative LMWH.49-51

For patients with cancer who have suffered recurrent VTE despite LMWH, “LMWH failure,” there are several entities to consider. First, it is helpful to obtain a heparin level by using anti–factor Xa assay, which should be performed before additional heparin is added. This finding can help determine whether LMWH was within the therapeutic range as well as whether the patient adhered to therapy. Second, it is important to exclude HIT as a potential therapeutic adverse reaction. Third, it is important to determine whether cancer recurrence or progression has occurred as a contributing factor. Augmenting the LMWH dose and transitioning to twice daily dosing may be helpful. In general, however, LMWH failure in the setting of cancer portends a poor prognosis.

**Acute Isolated Distal DVT**

Distal DVT is defined as DVT confined to the deep veins of the calf (posterior or anterior tibial, peroneal, soleal, or gastrocnemius veins). Initial anticoagulation is recommended for acute isolated distal leg DVT if patients have risk factors for extension or severe symptoms (grade 2C).14 Risk factors for extension include thrombus length greater than 5 cm, multiple veins involved, unprovoked event, cancer, previous VTE, hospitalization, or recent surgery.14 In addition, in the discussion section of the AT9 guidelines, D-dimer is noted as a possible risk factor for extension; however, there are no recommendations for how this is to be used clinically. In the absence of risk factors or severe symptoms, serial imaging can be performed at 1 and 2 weeks while withholding anticoagulation (grade 1B).14 Most of the thrombus propagation occurs within the first 2 weeks. If there is thrombus propagation, then anticoagulation should be initiated (grade 1B).14 It is our practice to offer anticoagulation for patients with calf DVT unless there is an obvious contraindication such as major bleeding. Even with an unprovoked calf DVT, 3 months of anticoagulation is recommended over extended therapy (grade 1B).14

**VTE in Pregnant Patients**

Anticoagulation in pregnancy must take both the fetus and the mother into consideration.52 Low-molecular-weight heparin is recommended over warfarin for both prevention and treatment of VTE in pregnant patients (grade 1A).52 If acute VTE is diagnosed during pregnancy, anticoagulation should be continued until delivery and reinitiated for at least 6 weeks postpartum with a total duration of at least 3 months (grade 2C).52 The use of fondaparinux or parenteral direct thrombin inhibitors should be reserved for women with HIT (grade 2C).52 The oral direct thrombin inhibitors and factor Xa inhibitors should also be avoided in pregnancy owing to the lack of safety data (grade 1C).52

Antepartum prophylaxis is suggested for pregnant women at increased risk: previous unprovoked VTE, pregnancy or estrogen-related VTE, homozygous factor V Leiden or prothrombin G20210A, or antiphospholipid antibody syndrome (grade 2C).52 For these patients, prophylactic or intermediate dose LMWH (enoxaparin 40 mg once daily, dalteparin 5000 U once daily, or LMWH with dose-adjusted anti–factor Xa levels 0.2-0.6 U/mL) is recommended. For pregnant women with single previous VTE associated with a transient risk factor (not pregnancy or estrogen related), no antepartum prophylaxis is recommended (grade 2C).52 Pregnant women with previous VTE should receive postpartum prophylaxis for 6 weeks with prophylactic dose LMWH or warfarin with a goal INR of 2.0 to 3.0 (grade 2B).52 Neither LMWH nor warfarin is secreted in breast milk.

**Catheter-Related Upper Extremity DVT**

For patients with a catheter-related upper extremity DVT, therapeutic anticoagulation is indicated, typically for 3 months.14 Catheter retrieval is not necessary as long as it remains functional and required for clinical care (grade 2C).14 Anticoagulation for upper extremity DVT in the setting of a central venous catheter should be continued as long as the catheter is in place (grade 1C).14 The deep veins of the upper extremity include the brachial, axillary, subclavian, and innominate veins. Superficial thrombosis of the cephalic and basilic veins does not require anticoagulant therapy. It is not our practice to provide primary prophylaxis.
to patients with indwelling central venous catheters.

**Splanchnic and Hepatic Vein Thrombosis**

The splanchnic venous system includes the portal, hepatic, splenic, superior, and inferior mesenteric veins. Depending on the extent and rapidity of thrombus propagation, complications may include bowel or splenic infarction or portal hypertension. Anticoagulation initiation depends on symptoms, extent, and acuity of thrombus formation. With symptoms or extensive and acute appearing thrombosis, anticoagulation therapy should be considered (grade 1B). For incidentally found asymptomatic splanchnic vein thrombosis, the AT9 guidelines suggest no anticoagulation (grade 2C). It is our clinical practice to treat acute splanchnic vein thrombosis irrespective of symptoms to prevent thrombus propagation. In the absence of randomized trial data, it has been our practice to treat splanchnic vein thrombosis for 6 months.

**Superficial Venous Thrombosis**

Superficial “phlebitis” is common, with an incidence of approximately 125,000 annually. Whereas approximately 25% of the patients will have an underlying DVT, ultrasound imaging both to confirm the diagnosis and to exclude subclinical DVT is warranted. Most events can be treated conservatively without anticoagulant therapy owing to the low risk of PE (1.3%). For superficial phlebitis exceeding 5 cm in length, prophylactic dose fondaparinux or LMWH is recommended for 45 days (grade 2B). Fondaparinux (2.5 mg daily) is preferred to LMWH (grade 2C). It is our practice to reserve parental anticoagulation for those patients with severe pain, extensive venous involvement, or evidence of thrombus propagation.

**Preventing Postthrombotic Syndrome**

Compression stockings (30-40 mm Hg knee high) worn for 2 years have been found to reduce the risk of postthrombotic syndrome by 50% (grade 2B). (grade 2B). It is our practice to reserve parental anticoagulation for those patients with severe pain, extensive venous involvement, or evidence of thrombus propagation.

**Mechanical Thrombectomy or Catheter-Directed Lysis for Acute DVT**

In patients with proximal DVT, approximately half will develop postthrombotic syndrome despite anticoagulant therapy. Approximately 50% will have disabling venous claudication and will be unable to walk more than 240 m because of severe pain. Moreover, iliac vein involvement is relatively common, affecting 25% of all patients with leg DVT. At present, mechanical thrombectomy and catheter-directed thrombolysis are considered for patients with iliofemoral DVT in whom the risk of postthrombotic syndrome is relatively high as well as for the rare patients with threatened venous gangrene despite treatment with anticoagulant therapy. The CaVenT study compared catheter-directed thrombolysis and anticoagulant therapy in 103 patients with acute proximal venous thrombosis. Eighty-eight percent of the patients treated with catheter-directed therapy achieved either complete or partial (50%-90%) lysis. At 6 months, iliofemoral patency was twice as frequent in the invasive treatment arm as in the conservative arm. At 2 years, iliofemoral patency was significantly increased and postthrombotic syndrome was significantly reduced in the active treatment arm. The risk of major bleeding did not differ between the 2 groups. The AT9 guidelines state that anticoagulant therapy is recommended over catheter-directed thrombolysis (grade 2C). However, for patients at reasonable risk for bleeding who place a high value on preventing postthrombotic syndrome and a low value on treatment complexity, cost, and risk of bleeding, catheter-directed thrombolysis is reasonable. It is our practice to assess all patients with iliac vein thrombosis for thrombolysis and mechanical thrombectomy candidacy.

**Thrombolytic Therapy in the Setting of PE**

For patients with PE and hypotension (systolic blood pressure <90 mm Hg or abrupt decrease by >40 mm Hg persisting for >15 minutes) who are not at high risk for bleeding, systemic thrombolytic therapy is reasonable (grade 1B). Specific recommendations include a short infusion time (2 hours instead of 24 hours) delivered systemically as opposed to directed with a pulmonary artery catheter (grade 2C). Catheter-directed or surgical pulmonary thrombectomy may be pursued for hypotensive patients who have failed thrombolytic therapy, have a contraindication to their use, or are likely to die of shock before realizing any benefit from systemic thrombolytic therapy (grade 2C). The decision to use catheter-directed or surgical thrombectomy depends on available expertise and resources.
Role of Inferior Vena Cava Filters

In general, there is a marked overuse of inferior vena cava (IVC) filters (both temporary and permanent) in the United States.63 Inferior vena cava filters are recommended for patients with acute proximal lower extremity DVT or PE who have a contraindication to anticoagulation (grade 1B).14 Preference is given for a temporary IVC filter that can be removed when the bleeding risk has resolved. Although IVC filters have been associated with an increased DVT recurrence rate,64 this is not an indication for extended secondary prophylaxis with anticoagulants. The duration of anticoagulant therapy is based on patient-specific risk, nature of the VTE event (provoked vs unprovoked), and perceived bleeding risk regardless of whether an IVC filter remains in place.

Perioperative Management of Anticoagulation

For the perioperative management of anticoagulation, the decision to use bridging therapy depends on the risk of thromboembolism as well as the potential for bleeding complications.55 For patients at moderate-to-high risk for thromboembolism, bridging anticoagulation should be considered in the periprocedural period (grade 2C).55 In our experience, this would include patients with a recent VTE diagnosis (<3 months) or those with active cancer.55 If possible, elective procedures should be delayed until the acute anticoagulation treatment period (3 months) can be completed. For patients requiring more urgent procedures, warfarin is discontinued 5 days before the anticipated procedure and bridging therapy with LMWH is initiated when the INR is less than 2.0 (grade 2C).63 The last dose of LMWH is provided on the morning of the day before the surgery. Warfarin is resumed the evening after surgery, assuming that heparinostasis has been achieved. Resumption of therapeutic LMWH postoperatively should be at 48 hours once heparinostasis is assured. Prophylactic dose LMWH can be initiated 12 hours after surgery. If individuals are at low risk for thromboembolism, anticoagulation can be discontinued 5 days before the procedure without bridging therapy (grade 2C).65

PREVENTION OF VTE

Whereby the adverse consequences of VTE result in considerable morbidity and mortality, prevention of this disease is a high priority. The Agency for Healthcare Research and Quality has deemed VTE prophylaxis as one of the top priorities of improved patient safety practices.57,68 Moreover, the National Quality Forum has endorsed VTE prophylaxis standards and outcomes measures.69 Despite a large volume of high-quality randomized controlled trial data demonstrating safety and efficacy, many hospitalized surgical and nonsurgical patients with an indication fail to receive VTE prophylaxis. For example, the ENDORSE study70 of 358 hospitals in 32 countries enrolling 68,183 patients found that only 58.5% of at-risk surgical patients and 39.5% of at-risk medical patients received American College of Chest Physicians-recommended VTE prophylaxis. Similar results have been reported for US hospitals. In the DVT FREE prospective registry,71 of 2726 hospital inpatients with ultrasound-confirmed DVT, only 42% received prophylaxis within the 30 days before diagnosis. Improving best practices for ensuring appropriate delivery of VTE prophylaxis can be challenging. Information technology using computerized clinical decision support with informed order sets and “opt-in” strategies may improve this delivery.72

General Medical Patients

For patients who are acutely ill and hospitalized, a strategy for VTE prevention may include prophylactic dose LMWH, low-dose UFH (twice or thrice daily), or fondaparinux (Table 2) continued throughout the hospitalization (grade 1B).73 For patients at high risk for bleeding, mechanical thromboprophylaxis with graduated compression stockings and/or intermittent pneumatic compression (IPC) is recommended (grade 2C).73 The AT9 guidelines do not recommend rivaroxaban for VTE prevention in acutely ill general medical patients. The recent MAGELLAN trial74 evaluated the extended use of rivaroxaban (35±4 days) vs enoxaparin (10±4 days) or placebo. Rivaroxaban was associated with fewer VTE events, but at the cost of higher risk of bleeding. Overall, rivaroxaban was noninferior to the standard prevention with enoxaparin.

When determining the likelihood of VTE for general medical patients, the Padua Prediction
Score can help stratify patients into low or high risk. This scoring system requires further validation; however, it is poised to be a useful clinical tool. It is important to identify those general medical patients who will benefit most from pharmacologic thromboprophylaxis.

### Nonorthopedic Surgery Patients

The greatest risk of VTE in nonorthopedic surgery includes general and abdominal pelvic surgery, particularly in the setting of an underlying malignant disease. Decisions regarding pharmacologic (Table 2) or mechanical prophylaxis depend on the patient-specific VTE risk and bleeding propensity (Figure 3). Venous thromboembolism risk assessment can be accomplished by using the Caprini score, which stratifies patient risk into 4 categories of thrombosis risk: very low (0 points; ≤0.5%), low (1-2 points; 1.5%), moderate (3-4 points; 3%), and high (>5 points; 6%) (Figure 2 and Table 3). For patients at very low risk, early and frequent ambulation is recommended only over pharmacologic (grade 1B) or mechanical (grade 2C) interventions.

Patients who undergo cardiac surgery and have an uncomplicated postoperative course can be treated with IPC (grade 2C). With any non-hemorrhagic complications (ie, electromechanical, infectious, pulmonary, neurologic, or renal dysfunction that may prolong hospital stay), LMWH or UFH should be added (grade 2C). For patients who undergo thoracic surgery and have moderate-to-high risk of VTE as assessed by using the Caprini method, without significant risk of bleeding, either LMWH or low-dose UFH should be combined with mechanical prophylaxis (grade 2B). For patients at high risk for bleeding, mechanical prophylaxis should be used until the risk of significant hemorrhage decreases and pharmacologic treatment can be initiated. In the setting of spinal surgery in which a patient is at high risk for VTE, once hemostasis is achieved and the bleeding risk decreases, the addition of pharmacologic prophylaxis is recommended in addition to the use of IPC.

### Table 2. VTE Prophylaxis Strategies.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Acutely Ill General Medical Patients</th>
<th>Patients Who Undergo Nonorthopedic Surgery</th>
<th>Patients Who Undergo Orthopedic Surgery (THA or TKA)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Enoxaparin</td>
<td>Subcutaneous: 40 mg once daily</td>
<td>Subcutaneous: 40 mg once daily</td>
<td>Subcutaneous: 30 mg twice daily</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>≥12 h before or after surgery</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Continued for ≥10-14 d</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Consider up to 35 d</td>
</tr>
<tr>
<td>Dalteparin</td>
<td>Subcutaneous: 5000 IU once daily</td>
<td>Subcutaneous: Low risk: 2500 IU once daily</td>
<td>Subcutaneous: 2500 IU</td>
</tr>
<tr>
<td></td>
<td></td>
<td>High risk: 2500 IU for 12 h after surgery and then 5000 IU once daily</td>
<td>≥12 h after surgery and then 5000 IU once daily</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Continue for ≥10-14 d</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Consider up to 35 d</td>
</tr>
<tr>
<td>Fondaparinux</td>
<td>Subcutaneous: 2.5 mg once daily</td>
<td>Subcutaneous: 2.5 mg once daily continued for up to 10 d</td>
<td>Subcutaneous: 2.5 mg once daily</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Continue for ≥10-14 d</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Consider up to 35 d</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>CrCl 30-50 mL/min: 1.5 mg once daily</td>
</tr>
<tr>
<td>Warfarin</td>
<td>Not recommended</td>
<td>Not recommended</td>
<td>Adjusted dose warfarin for a goal INR of 2.0 to 3.0 for ≥10-14 d</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Consider up to 35 d</td>
</tr>
<tr>
<td>Rivaroxaban</td>
<td>Noninferior to LMWH at a dose of 10 mg once daily (see text discussion)</td>
<td>Not studied</td>
<td>10 mg once daily for ≥6-10 h after surgery</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Continue ≥10-14 d</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Consider up to 35 d</td>
</tr>
<tr>
<td>Unfractionated heparin</td>
<td>Subcutaneous: 5000 U thrice daily in cancer patients</td>
<td>Subcutaneous: 5000 U twice daily or thrice daily until fully ambulatory</td>
<td>Subcutaneous: 5000 U twice daily or thrice daily until fully ambulatory</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Continue for ≥10-14 d</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Consider up to 35 d</td>
</tr>
</tbody>
</table>

THA = total hip arthroplasty; TKA = total knee arthroplasty; VTE = venous thromboembolism.
Patients with major trauma should be treated with both mechanical and pharmacologic prophylaxis unless contraindicated (grade 2C). Inferior vena cava filters are not routinely recommended for primary VTE prevention in patients with major trauma (grade 2C).

**Orthopedic Surgery Patients**

For major orthopedic surgery, there are many options for VTE prevention including LMWH, fondaparinux, dabigatran, apixaban, rivaroxaban, low-dose UFH, adjusted-dose vitamin K, and aspirin (Table 3; grade 1B).

---

**TABLE 3. Caprini Risk Assessment Model**

<table>
<thead>
<tr>
<th>1 Point</th>
<th>2 Points</th>
<th>3 Points</th>
<th>5 Points</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age 41-60 y</td>
<td>Age 61-74 y</td>
<td>Age &gt;75 y</td>
<td>Stroke &lt;1 mo</td>
</tr>
<tr>
<td>Minor surgery</td>
<td>Arthroscopic surgery</td>
<td>History of VTE</td>
<td>Elective lower extremity arthroplasty</td>
</tr>
<tr>
<td>Body mass index &gt;25 kg/m²</td>
<td>Major open surgery &gt;45 min</td>
<td>Family history of VTE</td>
<td>Hip, pelvis, or leg fracture</td>
</tr>
<tr>
<td>Swollen legs</td>
<td>Laparoscopic surgery &gt;45 min</td>
<td>Any thrombophilia</td>
<td>Acute spinal cord injury &lt;1 mo</td>
</tr>
<tr>
<td>Varicose veins</td>
<td>Bed rest &gt;72 h</td>
<td>Elevated serum homocysteine</td>
<td></td>
</tr>
<tr>
<td>Pregnant or postpartum</td>
<td>Immobilizing plaster cast</td>
<td>Heparin-induced thrombocytopenia</td>
<td></td>
</tr>
<tr>
<td>History of miscarriage</td>
<td>Central venous access</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Oral contraceptives or hormone replacement therapy</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sepsis &lt;1 mo</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Serious lung disease &lt;1 mo</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Abnormal pulmonary function</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Acute myocardial infarction</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Congestive heart failure &lt;1 mo</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>History of inflammatory bowel disease</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Medical patient on bed rest</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

VTE = venous thromboembolism.
Low-molecular-weight heparin is preferred with the first dose given at least 12 hours from the time of surgery (either pre- or postoperatively; grade 2C). Rivaroxaban is the only new oral anticoagulant with FDA approval for VTE prophylaxis in orthopedic surgery. Intermittent pneumatic compression for a goal of 18 hours daily is recommended in addition to anticoagulant therapy. When patients have a significant risk of bleeding, IPC may be used alone. A minimum duration of 10 to 14 days of thromboprophylaxis is recommended, with consideration of up to 35 days for patients at higher risk for VTE (grade 2B).

CONCLUSION

Data from an increasing number of published randomized controlled trials exist for VTE prevention and treatment with new and conventional anticoagulants, including mechanical devices. Incorporating a guideline-based approach to VTE evaluation, prevention, and treatment is anticipated to improve patient safety and outcomes, improve satisfaction of health care professionals knowing they have adhered to the latest and highest standards, and reduce health care delivery costs by reducing outcome events.

This is a review for clinicians based on the American College of Chest Physicians Clinical Practice Guidelines 2012 for Antithrombotic Therapy and Prevention of Thrombosis.

Abbreviations and Acronyms: AT9 = American College of Chest Physicians Antithrombotic Guidelines, 9th edition; DVT = deep vein thrombosis; FDA = Food and Drug Administration; HIT = heparin-induced thrombocytopenia; INR = international normalized ratio; IPC = intermittent pneumatic compression; IVC = inferior vena cava; LMWH = low-molecular-weight heparin; PE = pulmonary embolism; UFH = unfractionated heparin; VTE = venous thromboembolism

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REFERENCES


