MEDICAL CENTER

# VANDERBILT VUNIVERSITY

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Guideline: Adult Venous Thromboembolism Prophylaxis

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#### I. Purpose

To prevent pulmonary embolism (PE) and deep vein thrombosis (DVT) in burn patients

#### II. Risk Factors

| General Risk Factors   | High Risk Factors   | Very High Risk Factors   |
|--|---|--|
| General Risk FactorsBurn 10-19% TBSAAge > 40 yearsCentral venous accessISS > 9Blood transfusionsSurgical procedure within<br>72 hrsImmobilizationMalignancyExtensive soft tissue<br>traumaHormone therapyObesity | <ul> <li>High Risk Factors</li> <li>Burn 20-39% TBSA</li> <li>Inhalation injury</li> <li>Age &gt; 60 years</li> <li>ISS &gt; 15</li> <li>GCS &lt; 9 for &gt; 4 hours</li> <li>Major venous<br/>injury/repair</li> <li>PMH of venous<br/>thromboembolism<br/>(VTE)</li> <li>Lower extremity<br/>fracture</li> <li>Multiple spinal</li> </ul> | <ul> <li>Very High Risk Factors</li> <li>Burn ≥40% TBSA</li> <li>Spinal cord injury with paraplegia or quadriplegia</li> <li>Complex or multiple (≥2) lower extremity fractures</li> <li>Major pelvic fracture</li> <li>Multiple (≥ 3) long bone fractures (≥ 1 in the lower extremity)</li> <li>Age ≥75 years with any high risk factor)</li> </ul> |
| <ul> <li>AIS ≥ 3 (any region)</li> </ul>   | <ul> <li>Pregnancy</li> </ul>   | • COVID-19 positive  |

# III. Physical Exam Findings

- A. PE tachycardia, tachypnea, decreased oxygen saturation, altered mental status, diaphoresis
- B. DVT extremity pain, fever, localized edema/swelling, erythema

# IV. Lab and Radiology Findings

- A. Blood gas respiratory alkalosis, hypoxemia
- B. CXR nonspecific, peripheral wedge defect
- C. Extremity duplex- occlusive/non-occlusive thrombosis
- D. CT angio chest filling defect(s)

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# V. Pharmacologic Prophylaxis Guideline

A. All patients, unless otherwise specified, should receive VTE prophylaxis with weight-based enoxaparin (Lovenox) upon admission.

| Patient Dry Weight | Initial Enoxaparin Dose | Routine LMWH Monitoring |
|--------------------|-------------------------|-------------------------|
| < 50 kg            | 30 mg q 12h             | Yes                     |
| 50 – 89 kg         | 30 mg q 12h             | No                      |
| 90 – 129 kg        | 40 mg q 12h             | Yes                     |
| 130 – 179 kg       | 60 mg q 12h             | Yes                     |
| ≥ 180 kg           | 80 mg q 12 h            | Yes                     |

- B. Enoxaparin should be ordered via the burn admission order set or via the burn enoxaparin order set (labeled enoxaparin- ADULT BURN USE ONLY) with doses scheduled at 0600/1800
- C. Pharmacologic VTE prophylaxis should be NOT be held for elevated baseline INR due to liver dysfunction.
- D. No doses of VTE prophylaxis will be held for operative procedures unless requested by the operating attending.
  - a. Patients receiving prophylactic enoxaparin doses of 60 mg or 80 mg should instead receive enoxaparin 40 mg for one dose prior to and one dose following the operative procedure and then return to the previous dosing regimen.

#### VI. Exceptions to VTE Prophylaxis Guideline

- A. Renal impairment
  - i. Patients with a significant rise in SCr (≥50%), CrCl < 30 mL/min, or receiving renal replacement therapy should receive renally dosed enoxaparin (dosed once daily) OR be switched to subcutaneous heparin 5000 units q 8 hours.
    - If BMI 40 kg/m<sup>2</sup> AND no epidural catheter or paravertebral block in place, increase subcutaneous heparin dose to 7500 units q 8 hours.
- B. Epidural or Paravertebral Block Placement
  - i. Enoxaparin will not be used 12 hours prior to epidural or paravertebral block placement, while the catheter is indwelling, or for 4 hours after removal.
  - ii. Heparin 5000 units q 8 hours and SCDs should be utilized

#### VII. Routine Monitoring

- A. A low molecular weight heparin (LMWH) level (Anti-Xa level) should be drawn for all patients with the following characteristic
  - i. Burn ≥ 20% TBSA
  - ii. Weight < 50kg or  $\ge$  90 kg
  - iii. Spinal cord injury with paraplegia, quadriplegia
  - iv. Patients with concomitant trauma per trauma division guideline
- B. LMWH level peaks should be drawn 4 hours after at least 3 doses of enoxaparin
  - i. If levels are drawn early or late, the level is not a true peak and cannot be used to adjust enoxaparin doses

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- ii. Goal peak is 0.2 to 0.4 IU/mL
  - If level below goal range, increase dose to the next enoxaparin syringe size
  - If level above goal range, decrease dose to the next syringe size
    - If already receiving enoxaparin 30 mg q 12 hours, decrease to 30 mg q 24 hours
    - If LMWH level remains above goal range despite decreasing to q 24 hour dosing, change to subcutaneous heparin.
- iii. Repeat LMWH levels should be checked every 2 weeks (or sooner if renal function declines and/or there is concern for bleeding).
  - Many patients may require lower enoxaparin doses as burn wounds heal.
- iv. If assistance with interpreting LMWH levels and/or dosing enoxaparin is needed, please contact the burn pharmacist.

#### VIII. IVC Filter Placement

- A. A prophylactic IVC filter may be considered in high risk burn patients with a contraindication, failure, or complication related to anticoagulation.
- B. Indications for a therapeutic IVC filter include known PE or lower extremity DVT and a contraindication, failure, or complication of anticoagulation (not a complete listing).
- C. Placement of IVC filter does not prevent PEs but prevents large clots from traveling to lower extremities. Once patients stabilize, they will still require anticoagulation as long as the filter is in place.

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