VANDERBILT VUNIVERSITY				
MEDICAL CI Protocol: Adult DVT Chemoprophylaxis	E NTER Category Protocol Number	Clinical Practice BC-2018-3		
	Approval Date: February 1, 2018 Review Date: February 1, 2019			

Applicable to					
⊠ VUH ☐ Children's ☐ DOT	□ VMG Off-site locations □ VMG	□ VPH □ Other			
Team Members Performing					
 □ All faculty ⊠ Faculty & staff & staff providing direct patient care or □ Other: 	⊠ MD ⊠ House Staff ⊠ APRN/PA	🗆 RN 🔲 LPN			
Content Experts					
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I. Purpose:

Burn patients are at an increased risk of venous thromboembolism (VTE) due to burn induced coagulation changes as well as many of the risk factors native to all hospitalized patients². The purpose of this protocol is to prevent VTE as a complication to any hospitalized burn patient.

II. Population:

Adult patients with cutaneous burns or inhalation injury admitted to Vanderbilt University Medical Center (VUMC)

III. Risk Factors:

Risk Factors	High Risk Factors	Very High Risk Factors
• Age > 40 years	 Inhalation injury 	 Spinal cord injury
Central venous access	 Age > 60 years 	with paraplegia or
 Blood transfusions 	• ISS > 15	quadriplegia
(≥4 units)	• GCS < 9 for > 4 hours	 Complex or multiple
 Surgical procedure 	 Major venous 	(≥ 2) lower extremity
within 72 hrs	injury/repair	fractures
 Immobilization 	• PMH of venous	 Major pelvic fracture
 Malignancy 	thromboembolism	 Multiple (≥ 3) long
• Extensive soft tissue	(VTE)	bone fractures (≥ 1 in
trauma	 Lower extremity 	the lower extremity)
Hormone therapy	fracture	 Age ≥ 75 years with
Obesity	 Multiple spinal 	any high-risk factor
• Burn wound infection	fractures	
	Pregnancy	

IV. Assessment:

- A. Physical Exam Findings
 - 1. PE- tachycardia, tachypnea, mental status changes, diaphoresis
 - 2. DVT- extremity pain, fever, localized edema, warmth/erythema
- B. Lab and Radiology Findings
 - 1. Arterial Blood Gas respiratory alkalosis, hypoxemia
 - 2. Chest X-ray- nonspecific, peripheral wedge defect
 - 3. Extremity Duplex occlusive/non-occlusive thrombus
 - 4. CT Angio Chest filling defect(s)

V. Intervention/Treatment:

- A. All burn patients, unless otherwise specified, should receive VTE prophylaxis with enoxaparin (Lovenox) 30 mg SQ q12 hr within 24 hours of admission.
- B. Obesity: For patients with a BMI of \geq 40 kg/m², starting enoxaparin dose is 40 mg Q 12 hrs.
- C. For patients with > 20% TBSA burn injury with either high risk or very high-risk factors, chemoprophylaxis will be continued until hospital discharge. For patients with impaired mobility who undergo inpatient rehabilitation, chemoprophylaxis is continued.
- D. No doses of enoxaparin will be held for operative procedures unless requested by the operating attending.

VI. Exceptions to VTE Prophylaxis Protocol

- A. Renal Impairment: For patients with a significant rise in SrCr (> 50%) or a creatinine clearance < 30 mL/min, enoxaparin may be renally adjusted to 30 mg daily or subcutaneous heparin 5000 units q8 hours may be substituted.
 - 1. For patients on renal replacement therapy, heparin 5000 units q8 hours is recommended.
- B. Traumatic brain injury and spinal cord injury excluded by the Trauma and Surgical Critical Care VTE Protocol (attached)

VII. LMWH Anti-factor Xa (Anti- Xa) Level Monitoring

- A. An Anti-Xa level should be drawn in patients with the following characteristics:
 - 1. Burn ≥20% TBSA
 - 2. Weight \geq 180 kg and any risk factor (all categories)
 - 3. BMI >40kg/m2 and any HIGH risk factor
 - 4. Anyone with 2 or more high risk or very high risk factors or 3 total of any category of risk factor
 - 5. Patients with concomitant trauma
 - i. Spinal cord injury with paraplegia, quadriplegia
 - ii. Complex or multiple ((≥ 2) lower extremity fractures
 - iii. Major pelvic fracture
 - iv. Multiple (≥ 3) long bond fractures or ≥ 1 lower extremity fracture
- Anti-xa level peaks should be drawn 4 hours after the administration of enoxaparin. These labs should be ordered after the third dose of enoxaparin.

- 1. To order: LMW Heparin Assay (must be timed correctly)
- 2. Goal peak is 0.2-0.4 IU/mL
- 3. Once the goal range is reached, no further monitoring needed.

4. Maximum dosing for enoxaparin is 1mg/kg BID (therapeutic dose). If that dose is reached and assay is still not in goal range, a hematology consult should be considered for heparin resistance and potential alternative therapeutic options.

VIII. Considerations

- A. IVC Filter Placement
 - 1. A prophylactic IVC filter may be considered in high risk burn patients with a contraindication, failure, or complications of anticoagulation.
 - 2. Indications for a therapeutic IVC filter include patients with a known PE or lower extremity DVT and a contraindication, failure, or complication of anticoagulation.

IX. References:

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- 3. Lin H, Faraklas I, Cochran A, Saffle J. Enoxaparin and antifactor Xa levels in acute burn patients. *Journal of burn care & research : official publication of the American Burn Association.* 2011;32(1):1-5.
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- 5. Pannucci CJ, Osborne NH, Wahl WL. Venous thromboembolism in thermally injured patients: analysis of the National Burn Repository. *Journal of burn care & research : official publication of the American Burn Association*. 2011;32(1):6-12.
- 6. Pannucci CJ, Obi AT, Timmins BH, Cochran AL. Venous Thromboembolism in Patients with Thermal Injury: A Review of Risk Assessment Tools and Current Knowledge on the Effectiveness and Risks of Mechanical and Chemical Prophylaxis. *Clinics in plastic surgery*. 2017;44(3):573-581.
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