

## **Pediatric Venous Thromboembolism (VTE) Prophylaxis Guideline**

**Purpose:** This guideline provides the minimum standard of care for any patient on the Pediatric Trauma Service with suspected or confirmed injury and those at risk for deep vein thrombosis.

### **Definitions:**

1. Altered mobility: A permanent or temporary state in which the child has a limitation in independent, purposeful physical movement of the body or of one or more extremities.
2. Deep Vein Thrombosis (DVT): A blood clot (thrombus) that was initially formed in a deep (non-peripheral) vein.
3. Thrombo-Emboloc Deterrent Hose (TED): Elastic stockings, either knee- or thigh-high.
4. Risk category: Refer to VTE Risk Factors algorithm
  - Low risk: No VTE risk factors
  - Moderate risk: Multiple risk factors for VTE in the absence of altered mobility or has altered mobility with one or fewer additional risk factors.
  - High risk: Altered mobility plus two or more additional risk factors
5. Sequential Compression Device (SCD): A device designed to intermittently squeeze blood from underlying deep veins in the leg upon compression of an inflatable sleeve, and to allow the blood to flow again when it decompresses.
6. Venous Thromboembolism (VTE): A blood clot (thrombus) in a vein or one that has broken free and is carried in the bloodstream (embolus).

### **Guideline:**

- It is recommended that patients age 10–17 years be assessed for VTE risk factors at the time of inpatient admission and reassessed at 48 – 72 hours of hospitalization.
- Based on that assessment, the patient is assigned a risk category (low, moderate, high).
- This should be documented in the patient's medical record.
- It is recommended that VTE prophylaxis be administered based on risk category as soon as feasible, but within 24 hours of assessment, unless there are contraindications (See algorithm).
- Refer to the algorithm for Risk Category Assessment and Prophylaxis for VTE. Scope:

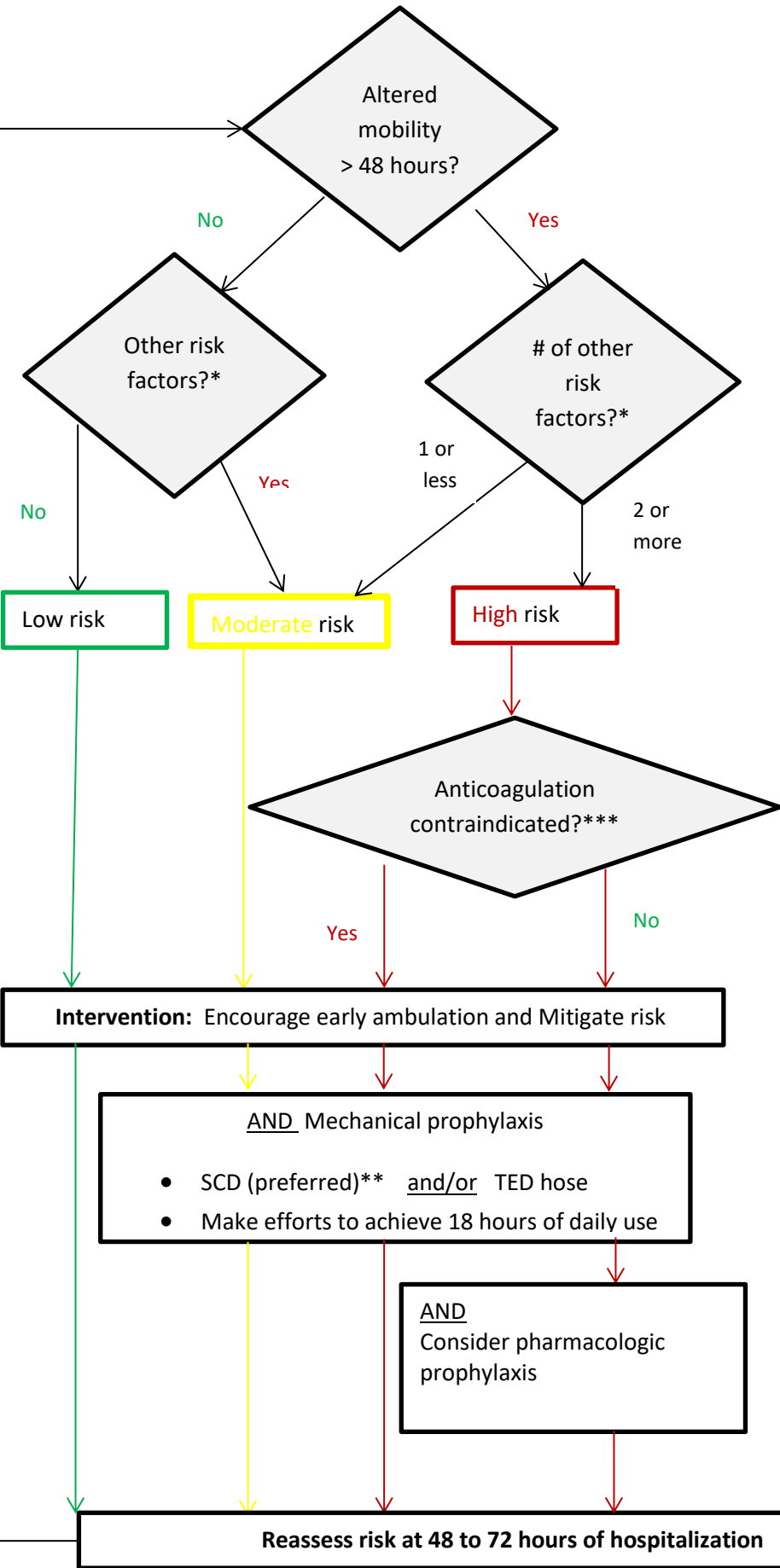
## RISK ASSESSMENT FOR THROMBOSIS 10-17 years old (to be completed by MD, RN or APN)

Risk Factors?	<p>History</p> <ul style="list-style-type: none"> <li><input type="checkbox"/> Personal history of blood clot</li> <li><input type="checkbox"/> Family history of clotting disorder</li> </ul> <p>High-risk Medical Conditions</p> <ul style="list-style-type: none"> <li><input type="checkbox"/> Known clotting disorder</li> <li><input type="checkbox"/> Blood stream infection (currently on antibiotics for positive blood culture)</li> <li><input type="checkbox"/> Cancer</li> <li><input type="checkbox"/> Chronic inflammatory condition (ie. Crohn's, Ulcerative Colitis, Lupus)</li> <li><input type="checkbox"/> Nephrotic syndrome</li> <li><input type="checkbox"/> Trauma patient if &gt; 1 lower extremity (LE) fracture, pelvic fracture or spinal cord injury</li> </ul> <p>Medications</p> <ul style="list-style-type: none"> <li><input type="checkbox"/> Estrogen (ie. birth control) in past 2 months (Depo shots don't contain estrogen)</li> </ul> <p>Physical Exam</p> <ul style="list-style-type: none"> <li><input type="checkbox"/> Obesity (BMI &gt;95<sup>th</sup> percentile)</li> <li><input type="checkbox"/> PICC or central line</li> </ul>	Total number of risk factors:
Contraindications to SCD (sequential compression device)?	<ul style="list-style-type: none"> <li><input type="checkbox"/> Current DVT</li> <li><input type="checkbox"/> Fracture of lower extremity (LE)</li> <li><input type="checkbox"/> Skin conditions affecting LE (burn, dermatitis, wound, epidermolysis bullosa)</li> </ul>	<ul style="list-style-type: none"> <li><input type="checkbox"/> Yes (if any checked)</li> <li><input type="checkbox"/> No</li> </ul>
Contraindications to Lovenox?	<ul style="list-style-type: none"> <li><input type="checkbox"/> Active bleeding</li> <li><input type="checkbox"/> Known bleeding disorder</li> <li><input type="checkbox"/> Epidural or lumbar puncture in the last 12 hours</li> <li><input type="checkbox"/> Platelets &lt;50,000/mm or heparin induced thrombocytopenia</li> <li><input type="checkbox"/> Brain tumor</li> <li><input type="checkbox"/> Pelvic fracture in last 48 hours</li> <li><input type="checkbox"/> Recent or scheduled neurosurgical procedure within 48 hours</li> <li><input type="checkbox"/> Uncontrolled hypertension</li> </ul>	<ul style="list-style-type: none"> <li><input type="checkbox"/> Yes (if any checked)</li> <li><input type="checkbox"/> No</li> </ul>
Equal or greater than 2 Risk Factors	<p>Anticipated altered mobility &gt; 48 hours?</p> <ul style="list-style-type: none"> <li><input type="checkbox"/> Bedrest or significant activity restriction</li> <li><input type="checkbox"/> Any line or tube that restricts mobility (id. Epidural, foley, NG to continuous suction, chest tube, EVD, mechanical ventilation)</li> </ul>	<ul style="list-style-type: none"> <li><input type="checkbox"/> Yes = <b>HIGH</b> Risk</li> <li><input type="checkbox"/> No = <b>MODERATE</b> Risk</li> </ul>
0-1 Risk Factor	<p>Anticipated altered mobility &gt; 48 hours?</p> <ul style="list-style-type: none"> <li><input type="checkbox"/> Bedrest or significant activity restriction</li> <li><input type="checkbox"/> Any line or tube that restricts mobility (id. Epidural, foley, NG to continuous suction, chest tube, EVD, mechanical ventilation)</li> </ul>	<ul style="list-style-type: none"> <li><input type="checkbox"/> Yes = <b>MODERATE</b> Risk</li> <li><input type="checkbox"/> No = <b>LOW</b> Risk</li> </ul>

If **NO** contraindications, the following interventions (and orders) are indicated:

PRE-OP ORDERS	
	Recommended Intervention/Order
ALL patients with surgery scheduled for > 60 minutes	<input type="checkbox"/> Intra-op SCD – apply in pre-op/holding area

POST-OP or ADMISSION ORDERS	
Risk Category	Recommended Intervention/Order
<b>LOW</b> Risk	None-encourage early ambulation
<b>MODERATE</b> Risk	<input type="checkbox"/> SCD – aim for 18 hours of use
<b>HIGH</b> Risk	<input type="checkbox"/> SCD - aim for 18 hours of use <input type="checkbox"/> Lovenox *(first dose 12 hours after surgery and hold 12 hours prior to surgical procedure) <ul style="list-style-type: none"> <li>• &lt; 50 kg= 0.5 mg/kg/dose SQ BID</li> <li>• &gt; 50 kg= 30 mg SQ BID or 40mg SQ daily</li> </ul> <p>*If renal dysfunction, consider decreasing dose and checking LMWH level 4 hours after 2<sup>nd</sup> or 3<sup>rd</sup> dose (goal 0.1-0.3 unit/ml, see BEST statement for management of LMWH in reference for more details)</p>



**\*VTE Risk Factors**

- Blood stream infection
- Central Venous Catheter (including non-tunneled, tunneled and PICS)
- History of venous thrombosis
- Hyperosmolar state (serum osmolality > 320 mOsm/kg)
- Inflammatory disease s(ie. IBD, SLE)
- Medications: asparaginase, estrogen use (within past 2 months)
- Obesity (BMI >95<sup>th</sup> percentile)
- Oncologic diagnosis
- Ortho procedures: hip or knee reconstruction
- Nephrotic syndrome
- Thrombophilia – known, or family history of clots
- Trauma : > 1 lower extremity long bone fracture, complex pelvic fracture or spinal cord injury

**\*\*Contraindications to Mechanical Prophylaxis**

- DVT, suspected or existing (can use TED hose)
- Extremity to be used has acute fracture
- Extremity to be used has PIV access
- Skin conditions affecting extremity
- Unable to achieve correct fit due to patient size

**\*\*\*Contraindications to Anticoagulation**

- Absolute:
- Bleeding disorder, known or tendency
  - Hemorrhage, evidence of or high risk of
  - Platelet count unable to be sustained >50,000

- Relative:
- Intracranial mass
  - Lumbar puncture of epidural catheter removal in prior 12 hours
  - Neurosurgical procedure
  - Pelvic fracture within past 48 hours
  - Uncontrolled hypertension

**Review/Revision History:**

<b>Review/Revision Date:</b>	<b>Approved by:</b>
Created 11/2017	Trauma Services
Revised 02/2022	Trauma Services