

## Inclusion Criteria for this Guideline:

- Meets definition for clinical suspicion for Pulmonary Embolism (Clinical Symptoms: dyspnea, chest pain, hypoxemia, hemoptysis, shock or cardiac arrest)
- Increased suspicion in patients with Risk Factors (see below) Exclusion Criteria for this Guideline:
- > 18 yo of age Apply Wells Criteria to assess risk and activate the PERT team at NH.

This guideline is intended to be used for patients with pulmonary embolism regardless of location at NCH who meet the below inclusion criteria.

# Key to using guideline

- This is a guideline, not a policy. Patient variation and other factors may impact management decisions. Patients must meet inclusion criteria and not meet one or more of the exclusion criteria.
- Blue boxes at bottom of page will contain hyperlinks to other pages of the guidelines. Clicking on the underlined word or phrase will take you to the page.
- Green boxes represent steps in an algorithm
- Yellow shapes represent decision branch points or key points of concern/caution

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### If questions about this guideline, contact

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Version 2: June 2025





<sup>b</sup>When intubation is indicated caution should be taken given the high risk of cardiac arrest in acute right heart failure. IV fluids, vasoactive infusions, resuscitation medications, equipment, and appropriate personnel should be present at the bedside. Induction medications with favorable hemodynamic profiles should be used.

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Heparin and Alteplase dosing

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Medication	Dosing	Considerations	
Heparin	<1 yr old Bolus: 75 units/kg over 10 min Infusion: 28 units/kg/hr ≥ 1 yr old AND <30 kg Bolus: 75 units/kg over 10 min Infusion: 20 units/kg/hr ≥ 1 yr old AND ≥≥30 kg Bolus: 75 units/kg over 10 min Infusion: 18 units/kg/hr	<ul> <li>Monitor PTT</li> <li>PTT goal 59-97</li> <li>Use actual body weight</li> </ul>	
Alteplase	<ul> <li><u>High dose</u> (High Risk PE)</li> <li>Patients &lt;33.3 kg: 0.5 mg/kg/hr, Intravenous, Continuous, For 6 hours</li> <li>Patients &gt;33.3 kg: 16.7 mg/hr, Intravenous, Continuous, For 6 hours</li> <li>Reduce heparin infusion by 50% during Alteplase infusion and then increase to prior dose.</li> <li>Patients may receive one to two additional doses of alteplase in cases of ongoing instability.</li> <li><u>Low dose</u> (Intermediate Risk PE)</li> <li>Patients &lt;27.7 kg: 0.3 mg/kg/hr, Intravenous, Continuous, For 6 Hours</li> <li>Patients &gt;27.7 kg: 8.33 mg/hr, Intravenous, Continuous, For 6 Hours <u>Bolus dose</u> (Arresting Patient)</li> <li>0.5 mg/kg over 2 min</li> <li>Maximum bolus dose = 50 mg</li> </ul>	• Monitor CBC, platelets, fibrinogen q6h	

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- · Painful leg swelling recent history of DVT
- Family history of DVT/PE
- Personal history of DVT/PE
- Known thrombophilia predisposing to DVT/PE
- Recent or current central venous catheter
- Pregnancy or OCP use
- Recent or current immobility
- Recent orthopedic surgery/trauma
- Acute of chronic inflammatory conditions

PE Category	Description
High Risk (Massive) PE	Acute PE causing cardiopulmonary arrest, sustained hypotension (systolic BP < 5th percentile by age for at least 15 min or requiring vasoactive support), or normotension with signs or symptoms of shock
Intermediate Risk (Submassive) PE	Acute PE without hypotension or compensated shock, but evidence of RV strain by imaging (CTA or echo), myocardial necrosis by elevated cardiac troponin levels, or both
Low Risk PE	Acute PE not meeting criteria for high-risk or intermediate-risk PE

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PE Huddle				
Huddle attendees				
CICU and/or PICU Attending				
CV Surgery Attending				
Interventiona Cardiology Attending				
ECMO Primer oncall				
Hematology Attending				
Pharmacist				
<ul> <li>Huddle location: patient bedside (usually the PICU or CICU) or by phone/digital technology if all</li> </ul>				
attendees not present in the hospital				
Huddle occurs before delivery of systemic hrombolysis				
Goal of huddle is to determine best treatment method: systemic thrombolysis vs surgical				
intervention vs catheter-based embolectomy.				

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**Treatment considerations by risk category** 

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Favor Embolectomy:	Favor Thrombolysis:
<ul> <li>Suspected tumor embolus (patients with Wilms Tumor, Ewing Sarcoma, osteosarcoma) or other non- thrombotic sources</li> <li>Patients with contraindications to thrombolysis</li> <li>Patients with concomitant intracardiac thrombus</li> <li>Patients with intracardiac communications on echo</li> <li>Patients on ECMO</li> </ul>	<ul> <li>Thrombi extending distally which are not amenable to surgical embolectomy</li> <li>Patients with comorbidities that confer additional surgical or anesthetic risk</li> <li>Patients in whom surgical embolectomy is not readily available (within 2 hours from diagnosis of PE)</li> </ul>

# Intermediate Risk PE Considerations: Admission to PICU or CICU

- Call CICU Attending if needed for help in initiating of PE Huddle.
- Catheter Based Therapy in the Cardiac Cath Lab is preferred for most patients with intermediate-risk PE who require primary reperfusion.
- · If unavailable or patient is a poor candidate, anticoagulation with Heparin ALONE is often preferred over systemic thrombolysis given associated risks and lack of evidence supporting benefits in adult patients with intermediate-risk PE.
- Individual patient considerations to consider include: anticipated bleeding risk, patient's clinical condition, degree of RV strain, presence of elevated troponin.
- If systemic thrombolysis is used, low-dose alteplase may be used to mitigate bleeding risks. Low-dose alteplase = 0.1 to 0.3 mg/kg/h x6 hours (total = 0.6 to 1.8 mg/kg; max dose = 50 mg).
- Monitor patients for risk of decompensation highest 48-72 hours after start of treatment with Heparin alone.

# Other Considerations

· If diagnosed with a high risk or intermediate risk PE at outside facility with CTA, start Heparin drip and transfer to NCH. Do not initiate thrombolysis at outlying facility.

- · If diagnosed with a low risk PE at outside facility transfer to NCH ED for further diagnostic work up
- · If outside facility is a Norton Children's facility consider direct admission to PICU/CICU, if outside facility is not Norton affiliated, admit to NCH ED for further diagnostic work up.

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# Background

## **Background and Rationale**

Acute pulmonary embolism (PE) is a form of venous thromboembolism (VTE) that is common and sometimes fatal. PE refers to obstruction of the pulmonary artery or one of its branches by material (eg. thrombus, tumor, air, or fat) that originated elsewhere in the body. Historically, pediatric PE was thought to be related to infections, though it has become clear that PE is increasingly related to cancer (particularly acute lymphoblastic leukemia), congenital heart disease, acquired and inherited thrombophilias, and central line placements.

## Epidemiology

Studies examining the incidence of PE in children report an incidence of 8.6-57 in 100,000 hospitalized children, and 0.14-0.9 in 100,000 when studying the general population of non-hospitalized children. However, it is likely that these numbers are underestimate due to the silent nature of PE in children. In one study that centered on autopsy data, PE was considered in only 15% of patients that were diagnosed with PE post-mortem.

## Pathophysiology

Similar to any generation of thrombus, Virchow's triad (venous stasis, endothelial injury, and hypercoagulable state) leads to the initial clot. A PE is simply the embolization of said clot to the pulmonary arteries, though there is some growing belief that pediatric PE is not always caused by embolization. Most emboli are thought to arise from lower extremity proximal veins (iliac, femoral, and popliteal), and more than 50% of patients with proximal vein deep thrombosis (DVT) have concurrent PE at presentation.

After a thrombus has broken off and embolized to the pulmonary arteries, infarction, abnormal gas exchange, and cardiovascular compromise can occur. In terms of cardiovascular collapse, hypotension is due to diminished stroke volume and, therefore, lower cardiac output. In these patients, pulmonary vascular resistance (PVR) is increased due to physical obstruction of the vascular bed with thrombus and hypoxic vasoconstriction within the pulmonary arterial system. This process subsequently diminishes output from the RV, reducing left ventricle preload, and finally compromising cardiac output.

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# **Version History**

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Version	Date	Guideline Owner	Summary of Edits	Next Revision Due
1	June 2023	Dr. Jamie Furlong- Dillard	Original	
2	June 2025	Dr. Jamie Furlong- Dillard	<ol> <li>Updated references</li> <li>No changes to content</li> </ol>	June 2029

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