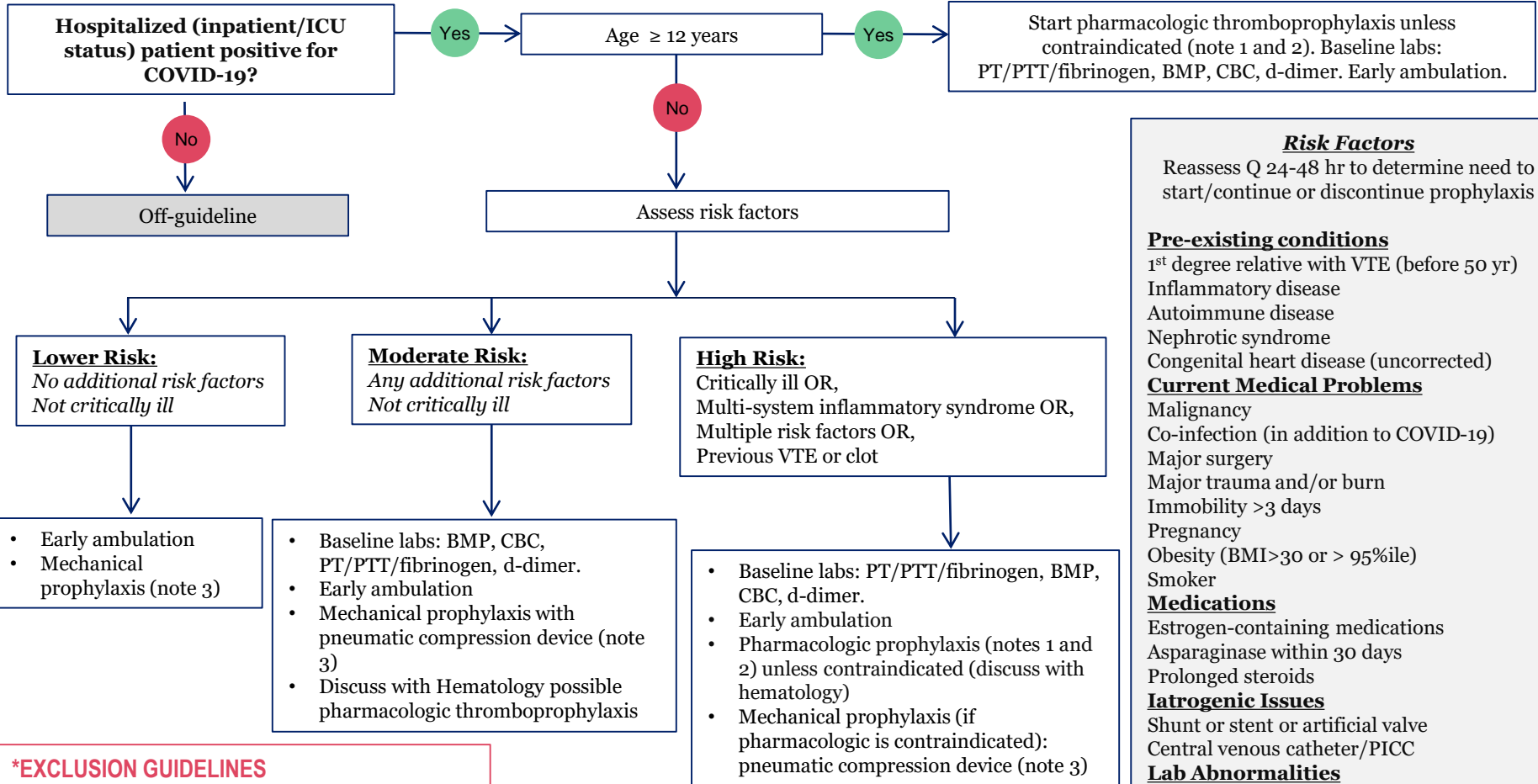


Aim: To standardize risk screening and prophylaxis for VTE in patients hospitalized with COVID-19.



Risk Factors

Reassess Q 24-48 hr to determine need to start/continue or discontinue prophylaxis

Pre-existing conditions

- 1st degree relative with VTE (before 50 yr)
- Inflammatory disease
- Autoimmune disease
- Nephrotic syndrome
- Congenital heart disease (uncorrected)

Current Medical Problems

- Malignancy
- Co-infection (in addition to COVID-19)
- Major surgery
- Major trauma and/or burn
- Immobility >3 days
- Pregnancy
- Obesity (BMI>30 or > 95thile)
- Smoker

Medications

- Estrogen-containing medications
- Asparaginase within 30 days
- Prolonged steroids

Iatrogenic Issues

- Shunt or stent or artificial valve
- Central venous catheter/PICC

Lab Abnormalities

Elevated D-Dimer may indicate increased risk-Discuss with Hematology if no other risk factors to prompt prophylaxis

***EXCLUSION GUIDELINES**

Patients **excluded** from this guideline:

- Patients in OBSERVATION status with mild illness
- Patients with identified thrombus or VTE
- Young infants are also at risk for VTE, however, this is out of scope for guideline due to lack of studies on pharmacologic interventions

Note: Information on COVID-19 management is rapidly evolving. Please refer to www.CDC.gov, www.who.int. Due to the dynamic situation, this guideline was not reviewed by the Guideline Governance Council but is updated regularly by clinical leadership.

Venous Thromboembolism (VTE) Prophylaxis in COVID-19+ patients

(Age 6 months*)

Aim: To standardize risk screening and prophylaxis for VTE in patients hospitalized with COVID-19.

Note 1. Pharmacologic prophylaxis options:

LMWH-enoxaparin, or heparin, see hospital policy 322.

- < 60 kg: Enoxaparin 0.5 mg/kg/dose subQ q12 h. Pharmacy to titrate.
- ≥ 60 kg, give 30 mg subQ BID. No routine monitoring needed.
- >60 kg actual weight (not dosing weight) and BMI greater than 40, give 40 mg subQ Q12H
- Avoid LMWH-enoxaparin if creatinine clearance < 30, consider heparin.
- Patients with MIS-C may receive aspirin AND LMWH-enoxaparin. In other patients with COVID-19 recommend holding aspirin while giving chemical thromboprophylaxis.

Note 3. Contraindications to mechanical prophylaxis: Affected extremity has acute fracture or peripheral IV, or skin/other condition (dermatitis, burn, tumor), OR unable to achieve correct fit due to patient size, OR lower extremity peripheral arterial insufficiency.

References:

<https://www.hematology.org/covid-19/covid-19-and-vte-anticoagulation>
CHOP VTE-COVID Prophylaxis Pathway

Monitor for signs and symptoms of bleeding if on pharmacologic prophylaxis

Oozing at sites (IV, surgical wounds, etc.)

Gross hematuria

Severe epistaxis (requiring intervention)

Bleeding causing a drop in hgb by 2 g/dL

Lower GI bleeding (black tarry stools, frank blood)

Upper GI bleeding (hemoptysis)

Ecchymosis or petechiae

Note 2. Pharmacologic VTE Prophylaxis Contraindications

Discuss thromboprophylaxis with any involved surgical services.

Absolute contraindications:

Active hemorrhage

Diagnosed bleeding disorder, known or tendency

Thrombocytopenia (platelets < 25 k)

Neurosurgery, TBI, or major solid organ injury in last 72 hr

Recent intracranial hemorrhage or acute stroke

Thrombolytic therapy within last 24 hr

Epidural or paraspinal hematoma

Epidural catheter in place (may use heparin)

Lumbar puncture or epidural catheter removed in last 6 hr

Significant uncorrected coagulopathy (e.g. INR > 2 or fibrinogen < 100 or PTT > 40): *Consult hematology in this scenario*

Heparin-induced thrombocytopenia, or other hypersensitivity to heparin or LMWH-enoxaparin

Relative contraindications:

For LMWH-enoxaparin, renal dysfunction (may need dose adjustment)

Significant uncontrolled hypertension > 99thile

Pelvic fracture in last 24–48 hr

Intracranial/spinal lesion at high risk of bleed

Anti-platelet therapy (discuss management with primary service, e.g. cardiology)

*For invasive procedures: hold heparin x 6 hours, hold LMWH-enoxaparin x 12-24 hours

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Note: Information on COVID-19 management is rapidly evolving. Please refer to www.CDC.gov, www.who.int. Due to the dynamic situation, this guideline is not reviewed by the Guideline Governance Council but is updated regularly by clinical leadership.

Disclaimer: This guideline is designed for general use with most patients; each clinician should use his or her own independent judgment to meet the needs of each individual patient. This guideline is not a substitute for professional medical advice, diagnosis or treatment.

Reviewer: Workgroup
Revised 05/2020