

## VTE Prevention: Medical patients

**If using pharmacological VTE prophylaxis for patients - aim to start it as soon as possible. Ideally within 14 hours of admission where possible, unless contraindicated or otherwise stated in population-specific recommendations**

<p><b>ALL Acutely ill patients:</b></p> <p><i>If the likely benefit from reducing the risk of symptomatic VTE is greater than the risk of harm from major bleeding caused by anticoagulation:</i></p> <ul style="list-style-type: none"> <li>• Add LMWH (prophylactic weight based dosing) for a minimum of 7 days if mobility not fully restored to baseline on discharge; or according to senior clinician discretion.</li> </ul>	<p><b>Patients with renal impairment:</b> <i>If the likely benefit from reducing the risk of symptomatic VTE is greater than the risk of harm from major bleeding caused by anticoagulation:</i></p> <ul style="list-style-type: none"> <li>• If using pharmacological VTE prophylaxis for people with renal impairment, use LMWH or UFH</li> <li>• If needed, reduce the dose of LMWH (see Appendix 3 for local protocol)</li> </ul>	<p><b>Ambulatory patients with cancer (Out-patients):</b></p> <p>Unless patient at increased risk of VTE due to risk factors other than cancer, do not offer VTE prophylaxis to patients with cancer /receiving cancer-modifying treatments (radio-/ chemo-/ immunotherapy) <b>who are mobile, consider in:</b></p> <ul style="list-style-type: none"> <li>• <b>Pancreatic cancer receiving chemotherapy:</b> consider LMWH for duration of chemotherapy</li> </ul>
<p><b>Acute stroke patients:</b></p> <ul style="list-style-type: none"> <li>• Do not offer anti-embolism stockings (AES)</li> <li>• Consider intermittent pneumatic compression (IPC) if immobile (start within first 3 days of stroke)</li> <li>• Continue IPC for 30 days or until mobile or discharged</li> <li>• Do not offer pharmacological thromboprophylaxis in acute stroke (ischaemic stroke within 2 weeks, reassess risk of VTE at 2 weeks; haemorrhagic stroke, consider after 1 month following discussion with senior clinician)</li> </ul>	<p><b>Acute coronary syndromes (ACS):</b></p> <p>If receiving anticoagulation for ACS management, patients do not usually require VTE prophylaxis acutely, but should be risk assessed and prescribed LMWH once ACS resolved, if still an in-patient.</p>	<ul style="list-style-type: none"> <li>• <b>Myeloma currently treated with thalidomide, pomalidomide or lenalidomide in combination with steroids:</b> LMWH (weight-based dosing), or aspirin (75mg or 150mg) or DOAC using locally agreed protocols</li> </ul>
<p><b>Patients receiving palliative care:</b></p> <p>If the risk assessment of VTE outweighs the risk of bleeding:</p> <ul style="list-style-type: none"> <li>• Use LMWH (weight-based dosing)</li> <li>• If life expectancy short, consider views of patient and family/ carers (document clearly).</li> <li>• Review thromboprophylaxis daily</li> <li>• Do not offer VTE prophylaxis to patients in the last few days of life.</li> </ul>	<p><b>Patients admitted for psychiatric illness:</b></p> <p>If the risk of VTE outweighs the risk of bleeding:</p> <ul style="list-style-type: none"> <li>• LMWH (weight-based dosing)</li> <li>• Continue VTE prophylaxis until patient no longer at risk for VTE.</li> </ul>	<p><b>Patients admitted to critical care:</b></p> <p>If the risk assessment of VTE outweighs the risk of bleeding:</p> <ul style="list-style-type: none"> <li>• LMWH (weight based dosing)</li> <li>• Risk assess at least daily</li> <li>• If pharmacological prophylaxis is contraindicated offer AES/ IPC until mobility returned to normal baseline</li> <li>• Refer to BSUH Covid VTE guideline for specific guidance on this group of patients</li> </ul>

Fondaparinux can be considered as an alternative to LMWH in certain circumstances – see Appendix 3