Appendix 1 Adult Inpatient Drug Chart

Risk Assessment for Venous Thromboembolism (VTE) for ALL adult patients admitted to hospital

(For pregnancy related admission see separate guidelines)	Risk assessment con	npleted at Pre-assessment
---	---------------------	---------------------------

	Assess mobility for all patients admitted to hospital - tick one box	✓	Action	
tep	Surgical patient		Assess for thrombosis risk and	
	Medical patient expected to have ongoing reduced mobility relative to normal state		bleeding risk (Step 2 and 3)	
	Medical patient NOT expected to have ongoing reduced mobility relative to normal state		VTE prophylaxis not required Go to Step 4	

	Assess thrombosis risk - tick all that apply (✓) Any tick should prompt thromboprophylaxis								
	Patient related	First	72hr		First	72hr			
	Active cancer or cancer treatment			Use of hormone replacement therapy					
	Age >60			Use of oestrogen-containing contraceptive therapy					
2	Dehydration			Varicose veins with phlebitis					
a	Known thrombophilias			Critical Care admission					
te	Obesity (BMI >30kg/m²)			Significant medical comorbidities (eg. heart disease,					
	Personal history or first-degree relative with a history of VTE			metabolic, endocrine or respiratory pathologies; acute infectious disease; inflammatory conditions)					
	Admission related								
	If total anaesthetic + surgery time > 90 minutes			Significant reduction in mobility for 3 days or more					
	If surgery involving pelvis or lower limb with a total anaesthetic + surgical time > 60 minutes			If acute surgical admission with inflammatory or intra-abdominal condition					

	Assess bleeding risk (re-assess regularly) - tick all that apply () If bleeding risks identified pharmacological prophylaxis may be contra-indicated, discuss with senior doctor before prescribing</th									
l	Patient related	First	72hr		First	72hr				
	Active bleeding			Thrombocytopenia (platelets < 50x10°/l)						
3	Acquired bleeding disorders (e.g. acute liver failure)			Uncontrolled systolic hypertension (≥ 230 mmHg)						
tep	Concurrent use of anticoagulants (e.g. warfarin with INR > 2, therapeutic dose of enoxaparin or rivaroxaban)			Untreated inherited bleeding disorders (such as haemophilia or Von Willebrand's disease)						
S	Acute stroke (< 3 months ago or risk of CNS bleeding)									
	Admission related									
	Neurosurgery, spinal surgery or eye surgery			Lumbar puncture/epidural/spinal anaesthesia within						
	(except cataract surgery)			the previous 4hrs or expected within the next 12hrs		ĺ				
	Other procedure with high bleeding risk			(If pharmacological prophylaxis is indicated clearly document timing of dose and/or start date)						

4	r	Assessmer	nt completed by (can be any	trained healthcare worker)			
2	7	Assessment	Signature	Name (PRINT)	Grade	Bleep	Date
15	וצ	First					
U	,	72 hour					

	Consider contra-indications tick if present (✓)								
5	Contra-indications to pharmacological prophylaxis (consider anti-embolism stockings)								
l e	Contra-indications to anti-embolism stockings (see below)								
S	Peripheral arterial disease	Peripheral arterial bypass grafting	Recent skin graft	Cardiac failure	Known Allergy		i l		
	Peripheral neuropathy	Severe dermatitis or gangrene	Severe leg oedema	Leg deformity					

		Guidelines used to risk assess: Surgical Orthopaedic Obstetric Other(specify)						
	9	Record treatment decisions tick all that apply (✓)		72hr		First	72hr	
e	ē	None - cross off enoxaparin from chart (drug 1)			Unfractionated Heparin (UFH)			
	Š	Enoxaparin			Rivaroxaban			
1		Intermittent pneumatic compression or footpumps			Anti-embolism stockings			

Write prescription

Ensure selected thromboprophylaxis is prescribed at an appropriate dose for the patient Consider eGFR and body weight if applicable (see 'Enoxaparin VTE Dosing Guidance' on opposite page)

Monitoring patients on enoxaparin

Heparin Induced Thrombocytopenia (HIT): For inpatients check baseline platelet count and then every 2-4 days until day 14. As a minimum for extended prophylaxis post-discharge check once between 4-7 days and again after 10-14 days of enoxaparin treatment. If platelets fall by more than 50% of baseline stop enoxaparin and discuss with haematology.

Hyperkalaemia: monitor serum potassium in at risk patients (those with diabetes mellitus, chronic renal failure, pre-existing metabolic acidosis or

taking potassium sparing drugs) before starting treatment and recheck regularly

 $\textbf{All patients to be periodically reassessed during inpatient stay as risk factors or contraindications \ may \ change.}$ Review assessment at 24 hrs then reassess every 72 - 96 hrs or, as or when patient's condition changes.