OB/GYN VTE SAFETY RECOMMENDATIONS FOR THE PREVENTION OF VTE IN MATERNAL PATIENTS

antepartum intrapartum postpartum

Applies to: Cesarean and Vaginal Delivery

Admission/Transfer of Care

Assess Patient for VTE Risk and Document

Risk Factor(s) (check all that apply)	2 points	3 points	5 points
 □ Minor surgery planned □ Age over 35 years old □ Prior major surgery < 1 month □ Pregnancy or < 1 month postpartum □ Varicose veins (current) □ Inflammatory bowel disease (history/current) □ Overweight (obesity BMI > 30 kg/m²) □ Oral contraceptives or hormone replacement therapy (history) □ Preeclampsia (history/current) □ Smoking (history/current) □ Postpartum hemorrhage (current) 	 □ Major surgery (> 45 min.) □ Laparoscopic surgery (> 45 min.) □ Patient confined to bed > 72 hrs. □ Currently on bedrest / restricted mobility in the antepartum / postpartum period □ Immobilizing plaster cast (current) □ Central venous catheter (current) □ Cesarean-section delivery (current) □ Diabetes (including pre-gestational diabetes) (history/current) □ Malignancy and/or chemotherapy (history/current) □ Parity > 5 	Patient admitted for chronic major illness: myocardial infarction congestive heart failure kidney disease chronic hypertension Severe sepsis/infection (current) VTE (DVT or PE) (history) Factor V Leiden/activated protein C resistance (history/current) Antithrombin III deficiency (history/current) Protein C or S deficiency (history/current) Prothrombin 20210A (history/current) Homocysteinemia (history/current) Other congenital or acquired thrombophilia (history/current)	In last month, patient has had: Major surgery Elective major lower extremity arthroplasty Hip, pelvis or leg fracture Stroke Multiple trauma Acute spinal cord injury (paralysis) Personal or family history of blood clots or clotting disorders
# of Risk Factors x 1 =	# of Risk Factors X 2 =	# of Risk Factors x 3 =	# of Risk Factors x 5 =

LOW RFA 1	MEDIUM RFA 2	HIGH RFA 3-4	HIGHEST RFA 5+
Antepartum	Antepartum	Antepartum	Antepartum
 □ Pharmacological prophylaxis not recommended unless indicated: □ ordered: 	 □ Pharmacological prophylaxis not recommended unless indicated: □ ordered: 	 Pharmacological prophylaxis: Ordered if VTE unprovoked and/or thrombophilia and/or hormonally provoked: 	□ Pharmacological prophylaxis:□ Ordered:□ Prophylactic low-molecular wt. heparin
□ Prophylactic low-molecular weight heparinor □ if LMWH unavailable:	or □ if LMWH unavailable: unfractionated heparin 5000 IU BID □ not ordered (why?	☐ Prophylactic low-molecular wt. heparin or ☐ if LMWH unavailable: unfractionated heparin BID (I tri- mester 5000 IU; II trimester 7500 IU;	or ☐ if LMWH unavailable: unfractionated heparin BID (I tri- mester 5000 IU; II trimester 7500 IU; III trimester 10000 IU)
unfractionated heparin 5000 IU BID not ordered (why?)	☐ Mechanical prophylaxis prescribed: ☐ graduated compression stockings & either: ☐ intermittent pneumatic compression	Ill trimester 10000 IU) not ordered (why?) Mechanical prophylaxis initiated:	□ not ordered (why?) □ Mechanical prophylaxis initiated: □ graduated compression stockings & either:
Postpartum ☐ Early ambulation as prescribed by health provider	or uenous foot pump	☐ graduated compression stockings & either:☐ intermittent pneumatic compression	☐ intermittent pneumatic compression or ☐ venous foot pump
□ Pharmacological prophylaxis not recommended unless indicated (not administered until 12 hours after vaginal delivery/epidural removal or 24 hours after cesarean delivery):	 □ on patient □ properly worn □ patient provided with information on proper use and wearing 	or □ venous foot pump □ Mechanical prophylaxis: □ on patient □ properly worn □ patient provided with information	 □ Mechanical prophylaxis: □ on patient □ properly worn □ patient provided with information on proper use and wearing
 Ordered if previous VTE, thrombophilia BMI>25kg/m² & antepartum immobilization: Prophylactic low-molecular weight 	☐ Pharmacological prophylaxis considered	on proper use and wearing Postpartum	Postpartum Early ambulation as prescribed by health provider
heparin or □UFH 5000 IU BID □ not ordered (<i>why</i> ?)	(not administered until 12 hours after vaginal delivery/epidural removal or 24 hours after cesarean delivery): □ ordered if multiple postpartum VTE Risk	 Early ambulation as prescribed by health provider Pharmacological prophylaxis (not administered until 12 hours after vaginal delivery/epidural removal or 24 hours after cesarean delivery): 	 □ Pharmacological prophylaxis (not administered until 12 hours after vaginal delivery/epidural removal or 24 hours after cesarean delivery): □ ordered:
☐ Mechanical prophylaxis initiated: ☐ graduated compression stockings & either:	Factors Prophylactic low-molecular wt heparin	☐ ordered: ☐ Prophylactic low-molecular wt. heparin or ☐ if LMWH unavailable, unfractionated	☐ Prophylactic low-molecular wt. heparin or ☐ if LMWH unavailable, unfractionated
☐ intermittent pneumatic compression or ☐ venous foot pump	heparin (UFH) 5000 IU BID not ordered (why?)	heparin (UFH) 5000 IU BID not ordered (why?)	heparin 5000 IU TID not ordered (why?)
Mechanical prophylaxis ongoing:on patient	☐ Mechanical prophylaxis initiated: ☐ graduated compression stockings & either:	Mechanical prophylaxis initiated:graduated compression stockings & either:	☐ Mechanical prophylaxis initiated:☐ graduated compression stockings & either:
properly wornpatient provided with information	☐ intermittent pneumatic compression or ☐ venous foot pump	☐ intermittent pneumatic compression or ☐ venous foot pump	☐ intermittent pneumatic compression or ☐ venous foot pump
on proper use and wearing Initiate discharge planning: discussed with patient/family	 □ Mechanical prophylaxis ongoing: □ on patient □ properly worn □ patient provided with information 	□ Mechanical prophylaxis ongoing: □ on patient □ properly worn	 □ Mechanical prophylaxis ongoing: □ on patient □ properly worn □ patient provided with information
 anticipated discharge date determined evaluate patient for home use of: intermittent pneumatic compression 	on proper use and wearing Initiate discharge planning:	 patient provided with information on proper use and wearing Initiate discharge planning: 	on proper use and wearing Initiate discharge planning:
(IPC) or □venous foot pump (VFP)	☐ discussed with patient/family☐ anticipated discharge date determined☐ evaluate patient for home use of:	☐ discussed with patient/family ☐ anticipated discharge date determined	☐ discussed with patient/family ☐ anticipated discharge date determined ☐ content for home was a few
or □no IPC/VFP □ if evaluated for IPC/VFP, initiate availability on discharge	☐ intermittent pneumatic compression (IPC)	 □ evaluate patient for home use of: □ intermittent pneumatic compression (IPC) or □ venous foot pump (VFP) or □ no IPC/VFP 	 □ evaluate patient for home use of: □ intermittent pneumatic compression (IPC) or □ venous foot pump (VFP) or □ no IPC/VFP
_	☐ if evaluated for IPC/VFP, initiate availability on discharge	if evaluated for IPC/VFP, initiate availability on discharge	if evaluated for IPC/VFP, initiate availability on discharge

Patient Reassessment

Patient Discharge

Repeat assessment if Patient hospitalized longer than 24 hrs., before surgery or with any significant change in patient condition.

 □ Discharge instructions include: □ healthcare provider contact information □ signs and symptoms of DVT and PE □ evaluate patient for home use of: □ intermittent pneumatic compression (IPC) or □ venous foot pump (VFP) or □ no IPC/VFP
□ Discharge instructions:□ reviewed with patient and read back□ received by patient
 Patient understands DVT/PE risk factors and how to prevent in postpartum period
☐ Follow up appointment made
☐ If immobility or bedrest required in antepartum period or extending 6 weeks postpartum: ☐ healthcare provider orders completed, including: ☐ evaluated patient for home use of: ☐ intermittent pneumatic compression (IPC) or ☐ venous foot pump (VFP) ☐ length of IPC/VFP treatment
 durable medical equipment unit notified of start date of IPC/VFP treatment
 □ patient provided with information on: □ purpose of IPC/VFP □ proper use and wearing □ importance on maintaining use at home until MD discontinues □ removed for ambulation and skin inspections (every 8 hrs) □ worn minimally 18- 20 hours per day